REPUBLIEK VAN SUID-AFRIKA

REPUBLIC OF SOUTH AFRICA

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PROKLAMASIE

van die

Staatspresident

van die Republiek van Suid-Afrika

No. 119, 1993

AANSTELLING VAN LEDE VAN 'N SPESIALE HOF KRAGTENS DIE WET OP DIE HANDHAWING EN BEVORDERING VAN MEDEDINGING, 1979 (WET No. 96 VAN 1979)

Kragtens die bevoegdheid my verleen by artikel 15 van die Wet op die Handhawing en Bevordering van Mededinging, 1979 (Wet No. 96 van 1979), stel ek hiermee regter Douglas Lennox Lyall Shearer as president en professor Mihkel Lemmit Truu en doktor Hans-Hugo Snyckers as lede van die spesiale hof aan wat ek by Proklamasie No. 13 in *Staatskoerant* No. 14641 van 12 Maart 1993 ingestel het.

Gegee onder my Hand en die Seël van die Republiek van Suid-Afrika te Kaapstad, op hede die Twintigste dag van September Eenduisend Negehonderd Drie-en-negentig.

F. W. DE KLERK,
Staatspresident.
Op las van die Staatspresident-in-Kabinet:

D. J. DE VILLIERS,
Minister van die Kabinet.

PROCLAMATION

by the

State President

of the Republic of South Africa

No. 119, 1993

APPOINTMENT OF MEMBERS OF A SPECIAL COURT IN TERMS OF THE MAINTENANCE AND PROMOTION OF COMPETITION ACT, 1979 (ACT No. 96 OF 1979)

By virtue of the powers vested in me by section 15 of the Maintenance and Promotion of Competition Act, 1979 (Act No. 96 of 1979), I hereby appoint Judge Douglas Lennox Lyall Shearer as president and Professor Mihkel Lemmit Truu and Doctor Hans-Hugo Snyckers as members of the special court which I established by Proclamation No. 13 in *Government Gazette* No. 14641 of 12 March 1993.

Given under my Hand and the Seal of the Republic of South Africa at Cape Town this Twentieth day of September, One thousand Nine hundred and Ninety-three.

F. W. DE KLERK,
State President.
By Order of the President-in-Cabinet:

D. J. DE VILLIERS,
Minister of the Cabinet.

GOEWERMENTSKENNISGEWINGS

ADMINISTRASIE: RAAD VAN VERTEENWOORDIGERS

DEPARTEMENT VAN ONDERWYS EN KULTUUR

No. 2273 3 Desember 1993

OORDRAG VAN STAATSONDERSTEUNDE SKOOL

Ek, Pieter Willem Saaiman, Minister van Onderwys en Kultuur: Raad van Verteenwoordigers, dra hierby kragtens artikel 5 (1) van die Wet op Onderwys vir

GOVERNMENT NOTICES

ADMINISTRATION: HOUSE OF REPRESENTATIVES

DEPARTMENT OF EDUCATION AND CULTURE

No. 2273 3 December 1993

TRANSFER OF STATE-AIDED SCHOOL

I, Pieter Willem Saaiman, Minister of Education and Culture: House of Representatives, hereby under section 5 (1) of the Coloured Persons Education Act, 1963

Kleurlinge, 1963 (Wet No. 47 van 1963), met ingang van die datum van publikasie van hierdie kennisgewing, die bestuur van en die beheer oor die Primêre Skool Doornbaai (NGK), distrik Springbok, aan die Administrasie: Raad van Verteenwoordigers oor.

P. W. SAAIMAN,
Minister van Onderwys en Kultuur:
Raad van Verteenwoordigers.

ADMINISTRASIE: VOLKSRAAD

DEPARTEMENT VAN BEHUISING EN WERKE

No. 2293 **3 Desember 1993**

WET OP HUURBEHEER, 1976

INTREKKING VAN GOEWERMENTSKENNISGEWING No. 3007 VAN 30 OKTOBER 1992 EN No. 3115 VAN 13 NOVEMBER 1992 INGEVOLGE WAARVAN SEKERE WONINGS, MOTORHUISE, MOTORSTAANPLEKKE EN BEDIENDEKAMERS VAN HUURBEHEER VOORWAARDELIK VRYGESTEL IS

Kragtens die bevoegdheid my verleen by artikel 51 (g) van die Wet op Huurbeheer, 1976 (Wet No. 80 van 1976), word Goewermentskennisgewing No. 3007 van 30 Oktober 1992 en No. 3115 van 13 November 1992 hierby ingetrek.

A. J. VLOK,
Minister van Behuising en Werke.

DEPARTEMENT VAN BUITELANDSE SAKE

No. 2271 **3 Desember 1993**

ERKENNING VERLEEN AS VISEKONSUL

Hierby word bekendgemaak dat aan mnr. James Baxter Hunt III met ingang van 24 September 1993 erkenning verleen is as Visekonsul van die Verenigde State van Amerika in Durban, met die provinsie Natal as sy regsgebied.

Mnr. Hunt is die opvolger van Me. L. Peterson.

(72/33/4)

No. 2305 **3 Desember 1993**

ERKENNING VERLEEN AS EREKONSUL

Hierby word bekendgemaak dat aan mev. Marie-May Kolsch met ingang van 1 September 1993 erkenning verleen is as Erekonsul van die Republiek Seychelle in Johannesburg, met die Republiek van Suid-Afrika as haar regsgebied.

(72/194/1)

(Act No. 47 of 1963), transfer the management and control of the Doornbaai (NGK) Primary School, District of Springbok, to the Administration: House of Representatives with effect from the date of publication of this notice.

P. W. SAAIMAN,
Minister of Education and Culture:
House of Representatives.

ADMINISTRATION: HOUSE OF ASSEMBLY

DEPARTMENT OF HOUSING AND WORKS

No. 2293 **3 December 1993**

RENT CONTROL ACT, 1976

WITHDRAWAL OF GOVERNMENT NOTICE No. 3007 OF 30 OCTOBER 1992 AND No. 3115 OF 13 NOVEMBER 1992 IN TERMS OF WHICH CERTAIN DWELLINGS, GARAGES, PARKING SPACES AND SERVANTS' ROOMS WERE CONDITIONALLY EXEMPTED FROM RENT CONTROL

Under the powers vested in me by section 51 (g) of the Rent Control Act, 1976 (Act No. 80 of 1976), Government Notice No. 3007 of 30 October 1992 and No. 3115 of 13 November 1992 is hereby withdrawn.

A. J. VLOK,
Minister of Housing and Works.

DEPARTMENT OF FOREIGN AFFAIRS

No. 2271 **3 December 1993**

RECOGNITION GRANTED AS VICE-CONSUL

It is hereby notified that Mr James Baxter Hunt III has, with effect from 24 September 1993, been granted recognition as Vice-Consul of the United States of America in Durban, with the Province of Natal as his area of jurisdiction.

Mr Hunt is the successor to Ms L. Peterson.

(72/33/4)

No. 2305 **3 December 1993**

RECOGNITION GRANTED AS HONORARY CONSUL

It is hereby notified that Mrs Marie-May Kolsch has, with effect from 1 September 1993, been granted recognition as Honorary Consul of the Republic of Seychelles in Johannesburg, with the Republic of South Africa as her area of jurisdiction.

(72/194/1)

DEPARTEMENT VAN NASIONALE OPVOEDING

No. 2290 3 Desember 1993

BURO VIR HERALDIEK

KENNISGEWING VAN DIE WYSIGING VAN REGISTRASIES VAN HERALDIESE VOORSTELLINGS, 'N NAAM EN UNIFORM

Die Buro vir Heraldiek gee hierby ingevolge artikel 10 van die Heraldiekwet, 1962 (Wet No. 18 van 1962), kennis van die wysiging van registrasies van heraldiese voorstellings, 'n naam en uniform soos hieronder uiteengesit:

1. Aangesien die naam en uniform (rugbytrui en -kouse) van die Transvaalse Rugbyvoetbalunie, soos geregistreer by Goewermentskennisgewings Nos. 953 van 3 Julie 1936 en No. 1400 van 22 Julie 1977, verander is, sal die betrokke aspekte van die registrasie soos volg gewysig word:

Nuwe naam van die aansoeker: **Transvaalse Rugby Unie.**

Nuwe beskrywing van die uniform:

Rugbytrui: 'n Wit trui met 'n 12 cm breë horisontale rooi streep om die bors, rug en moue heen; die kraag, mansjette en ribwerk van die moue ook rooi.

Rugbykous: 'n Swart kous met drie rooi strepe, elk 2,5 cm breed, op 'n wit omslag.

2. Aangesien die naam van die Veldskool Schoemansdal, wie se wapen by Goewermentskennisgewing No. 42 van 12 Januarie 1979 geregistreer is, verander is, sal die betrokke aspek van die registrasie soos volg gewysig word:

Nuwe naam van die aansoeker: **Schoemansdal Omgewingsopvoedingsentrum.**

3. Aangesien die wapen van die Umzingisi-Skool, soos gepubliseer by Goewermentskennisgewing No. 2652 van 30 Desember 1988, verander is, sal die betrokke aspek van die registrasie soos volg gewysig word:

Nuwe blasoen van die wapen: In bruin, 'n goue verkorte pylkruis, oor die middel heen 'n silwer tandrat belaai met 'n blou vierblad.

Wapenspreuk: MAWU KHANYE

No. 2291 3 Desember 1993

BURO VIR HERALDIEK

REGISTRASIE VAN HERALDIESE VOORSTELLINGS

Hierby word ingevolge artikel 5 (b) van die Heraldiekwet, 1962 (Wet No. 18 van 1962), kennis gegee dat die ondergenoemde by die Buro vir Heraldiek, Privaat Sak X236, Pretoria, 0001, geregistreer is:

1. Die wapen van die **Munisipaliteit van Carletonville,** soos gepubliseer in die *Offisiële Koerant van die Provinsie Transvaal* van 20 Junie 1962, te wete:

Wapen: Deursnee in drie, groen, goud en blou, in die skildhoof twee goue koringgerwe, op die middelpunt 'n voorstelling van 'n atoom

DEPARTMENT OF NATIONAL EDUCATION

No. 2290 3 December 1993

BUREAU OF HERALDRY

NOTICE OF THE AMENDMENT OF REGISTRATIONS OF HERALDIC REPRESENTATIONS, A NAME AND A UNIFORM

The Bureau of Heraldry hereby gives notice in terms of section 10 of the Heraldy Act, 1962 (Act No. 18 of 1962), of the amendment of registrations of heraldic representations, a name and a uniform as set out below:

1. Since the name and uniform (rugby jersey and stockings) of the Transvaal Rugby Football Union, as registered under Government Notices Nos. 953 of 3 July 1936 and No. 1400 of 22 July 1977, have been changed, the relevant aspects of the registration will be amended as follows:

New name of the applicant: **Transvaal Rugby Union.**

New description of the uniform:

Rugby jersey: A white jersey with a 12 cm wide horizontal red stripe round the chest, back and sleeves; the collar, cuffs and ribbing of the sleeves also red.

Rugby stocking: A black stocking with three red stripes, each 2,5 cm wide, on a white turnover.

2. Since the name of the Schoemansdal Veld School, whose arms were registered under Government Notice No. 42 of 12 January 1979, has been changed, the relevant aspect of the registration will be amended as follows:

New name of the applicant: **Schoemansdal Enviromental Education Centre.**

3. Since the arms of the Umzimgisi School, as published under Government Notice No. 2652 of 30 December 1988, have been changed, the relevant aspect of the registration will be amended as follows:

New blazon of the arms: Brunâtre, a cross humetty barbed or, surmounted in the centre by a cogwheel Argent charged with a quatrefoil Azure.

Motto: MAWU KHANYE

No. 2291 3 December 1993

BURO OF HERALDRY

REGISTRATION OF HERALDIC REPRESENTATIONS

Notice is hereby given in terms of section 5 (b) of the Heraldry Act, 1962 (Act No. 18 of 1962), that the undermentioned have been registered with the Bureau of Heraldry, Private Bag X236, Pretoria, 0001:

1. The arms of the **Municipality of Carletonville,** as published in the *Official Gazette of the Province of the Transvaal* of 20 June 1962, to wit:

Arms: Tierced per fess, Vert, Or and Azure, in chief two garbs Or, at fess point a representation of an atom consisting of three elliptical

bestaande uit drie elliptiese loopkringe, elk belaai met 'n elektron, wat mekaar rondom 'n kernpunt oorkruis, alles rooi, en in die skildvoet twee silwer golwende streepbalke.

Helmteken: 'n Geboë regterarm van natuurlike kleur, bokant die elmboog rooi gekleed, silwer omsoom, in die hand 'n skuinslinks geplaaste pyl van natuurlike kleur, punt na onder.

Wrong en dekklede: Rooi en silwer

Skildhouers: Regs 'n eland en links 'n buffel, albei van natuurlike kleur.

Wapenspreuk: EX DUBIO PROVENIAT SPES

2. Die wapen van die **Munisipaliteit van Warrenton,** soos gepubliseer in die *Offisiële Koerant van die Provinsie die Kaap die Goeie Hoop* van 23 Junie 1961, te wete:

Wapen: Deursnede, silwer en groen, in die skildhoof 'n blou golwende dwarsbalk en in die skildvoet 'n silwer paal belaai met swart tweelingbalke, regs vergesel van 'n aansiende bulkop van natuurlike kleur en links van 'n goue geopende boek; 'n skildhoof geskaak van drie rye van tien vierkante elk, silwer en blou.

Helmteken: 'n Goue halwe uitkomende leeu, rooi getong, wat 'n silwer ruitvormig geslypte diamant vashou.

Wrong en dekklede: Groen en silwer

Wapenspreuk: DEUR HOOP EN VLYT

No. 2304 3 Desember 1993

WET OP DIE SUID-AFRIKAANSE SERTIFISERINGSRAAD, 1986

AANSTELLING VAN LEDE VAN DIE SUID-AFRIKAANSE SERTIFISERINGSRAAD

Aangesien daar vyf vakatures bestaan in die Suid-Afrikaanse Sertifiseringsraad, ingestel by artikel 2 van die Wet op die Suid-Afrikaanse Sertifiseringsraad, 1986 (Wet No. 85 van 1986), versoek ek, Pieter Gariel Marais, Minister van Nasionale Opvoeding, enige liggaam, vereniging of organisasie om die name van persone wat vanweë hul opvoedkundige kwalifikasies en kundigheid in sake rakende die werksaamhede van genoemde Raad moontlik geskikte kandidate kan wees, voor 10 Januarie 1994 aan my voor te lê ten einde my in staat te stel om vyf lede vir genoemde Raad te kies.

Bedoelde voorleggings, vergesel van die *curriculum vitae* van elke betrokke persoon, moet gestuur word aan:

Die Direkteur-generaal: Nasionale Opvoeding
Posbus X122
PRETORIA
0001.

P. G. MARAIS,
Minister van Nasionale Opvoeding.

orbits, each showing an electron, intersecting one another about a central nuclear point, Gules, and in base two barrulets wavy Argent.

Crest: A dexter arm embowed proper, vested above the elbow Gules, edged Argent, the hand grasping an arrow in bend sinister, point downwards, proper.

Wreath and mantling: Gules and Argent

Supporters: Dexter an eland and sinister a buffalo, proper.

Motto: EX DUBIO PROVENIAT SPES

2. The arms of the **Municipality of Warrenton,** as published in the *Official Gazette of the Province of the Cape of Good Hope* of 23 June 1961, to wit:

Arms: Per fess, Argent and Vert, in chief a bar wavy Azure and in base on a pale Argent, between dexter a bull's head caboshed proper and sinister an open book Or, bars gemelles Sable; a chief chequy of three rows of ten squares each, Argent and Azure.

Crest: A demi-lion issuant Or, langued Gules, holding a lozenge-shaped cut diamond Argent.

Wreath and mantling: Vert and Argent

Motto: BY HOPE AND DILIGENCE

No. 2304 3 December 1993

SOUTH AFRICAN CERTIFICATION COUNCIL ACT, 1986

APPOINTMENT OF MEMBERS OF THE SOUTH AFRICAN CERTIFICATION COUNCIL

Since five vacancies exist in the South African Certification Council, established by section 2 of the South African Certification Council Act, 1986 (Act No. 85 of 1986), I, Pieter Gabriel Marais, Minister of National Education, hereby, request any body, society or organisation to submit to me before 10 January 1994 the names of persons who on account of their educational qualifications and expertise in matters affecting the functions of the said Council may be suitable candidates in order to enable me to select five members for the said Council.

The said submissions, accompanied by the *curriculum vitae* of each person concerned, must be sent to:

The Director-General: National Education
P.O. Box X122
PRETORIA
0001.

P. G. MARAIS,
Minister of National Education.

No. 2306 3 Desember 1993

ARGIEFWET, 1962 (WET No. 6 VAN 1962)

VERKLARING VAN DIE KANTOOR VAN DIE OUDITEUR-GENERAAL KRAGTENS ARTIKEL 2A TOT 'N STAATSKANTOOR

Kragtens artikel 2A van die Argiefwet, 1962 (Wet No. 6 van 1962), verklaar ek, Pieter Gabriel Marais, Minister van Nasionale Opvoeding, in oorleg met die Minister van Staatsbesteding, hierby die Kantoor van die Ouditeur-generaal, ingestel by artikel 3 van die Ouditreëlingswet, 1992 (Wet No. 122 van 1992), tot 'n Staatskantoor vir die doeleindes van eersgenoemde Wet.

P. G. MARAIS,
Minister van Nasionale Opvoeding.

No. 2306 3 December 1993

ARCHIVES ACT, 1962 (ACT No. 6 OF 1962)

DECLARATION OF THE OFFICE OF THE AUDITOR-GENERAL TO BE A GOVERNMENT OFFICE IN TERMS OF SECTION 2A

In terms of section 2A of the Archives Act, 1962 (Act No. 6 of 1962), I, Pieter Gabriel Marais, Minister of National Education, hereby declare the Office of the Auditor-General, established in consultation with the Minister of State Expenditure, by section 3 of the Audit Arrangements Act, 1992 (Act No. 122 of 1992), to be a Government Office for the purposes of the first-mentioned Act.

P. G. MARAIS,
Minister of National Education.

DEPARTEMENT VAN STREEK- EN GRONDSAKE

No. 2292 3 Desember 1993

AANWYSING VAN SEKERE GROND GELEË IN DIE DISTRIK PIETERMARITZBURG, PROVINSIE NATAL, VIR DOELEINDES VAN DIE WET OP REËLING VAN GRONDTITELS, 1993 (WET No. 111 VAN 1993)

Kragtens die bevoegdheid my verleen by artikel 2 (1) van die Wet op Reëling van Grondtitels, 1993 (Wet No. 111 van 1993), wys ek, Anthon Tobias Meyer, Adjunkminister van Grondsake, hierby die grond in Bylae vermeld, aan as grond waarmee ooreenkomstig die bepalings van genoemde Wet gehandel kan word.

A. T. MEYER,
Adjunkminister van Grondsake.

BYLAE

1. Perseel 761 van die plaas Edendale 775, distrik Pietermaritzburg.
2. Onderverdeling 1 van Perseel 766 van die plaas Edendale 775, distrik Pietermaritzburg.
3. Perseel 655 van die plaas Edendale 775, distrik Pietermaritzburg.
4. Restant van Perseel 283 van die plaas Edendale 775, distrik Pietermaritzburg.
5. Onderverdeling 2 van Perseel 328 van die plaas Edendale 775, distrik Pietermaritzburg.
6. Perseel 132 van die plaas Edendale 775, distrik Pietermaritzburg.
7. Restant van Perseel 131 van die plaas Edendale 775, distrik Pietermaritzburg.
8. Restant van Perseel 730 van die plaas Edendale 775, distrik Pietermaritzburg.
9. Restant van Perseel 735 van die plaas Edendale 775, distrik Pietermaritzburg.
10. Onderverdeling 28 van Perseel 283 van die plaas Edendale 775, distrik Pietermaritzburg.

DEPARTMENT OF REGIONAL AND LAND AFFAIRS

No. 2292 3 December 1993

DESIGNATION OF CERTAIN LAND SITUATED IN THE DISTRICT OF PIETERMARITZBURG, PROVINCE OF NATAL, AS LAND FOR PURPOSES OF THE LAND TITLES ADJUSTMENT ACT, 1993 (ACT No. 111 OF 1993)

Under the powers vested in me by section 2 (1) of the Land Titles Adjustment Act, 1993 (Act No. 111 of 1993), I, Anthon Tobias Meyer, Deputy Minister of Land Affairs, hereby designate the land mentioned in the Schedule as land which may be dealt with in accordance with the provisions of the said Act.

A. T. MEYER,
Deputy Minister of Land Affairs.

SCHEDULE

1. Lot 761 of the farm Edendale, District of Pietermaritzburg.
2. Subdivision 1 of Lot 766 of the farm Edendale 775, District of Pietermaritzburg.
3. Lot 655 of the farm Edendale 775, District of Pietermaritzburg.
4. Remainder of Lot 283 of the farm Edendale 775, District of Pietermaritzburg.
5. Subdivision 2 of Lot 328 of the farm Edendale 775, District of Pietermaritzburg.
6. Lot 132 of the farm Edendale 775, District of Pietermaritzburg.
7. Remainder of Lot 131 of the farm Edendale 775, District of Pietermaritzburg.
8. Remainder of Lot 730 of the farm Edendale 775, District of Pietermaritzburg.
9. Remainder of Lot 735 of the farm Edendale 775, District of Pietermaritzburg.
10. Subdivision 28 of Lot 283 of the farm Edendale 775, District of Pietermaritzburg.

11. Restant van Perseel 755 van die plaas Edendale 775, distrik Pietermaritzburg.
12. Perseel 658 van die plaas Edendale 775, distrik Pietermaritzburg.
13. Restant van Perseel 116 van die plaas Edendale 775, distrik Pietermaritzburg.
14. Perseel 134 van die plaas Edendale 775, distrik Pietermaritzburg.
15. Perseel 653 van die plaas Edendale 775, distrik Pietermaritzburg.
16. Perseel 734 van die plaas Edendale 775, distrik Pietermaritzburg.
17. Restant van Perseel 736 van die plaas Edendale 775, distrik Pietermaritzburg.
18. Onderverdeling 1 van Perseel 755 van die plaas Edendale 775, distrik Pietermaritzburg.
19. Onderverdeling 3 van Perseel 131 van die plaas Edendale 775, distrik Pietermaritzburg.
20. Onderverdeling 2 van Perseel 131 van die plaas Edendale 775, distrik Pietermaritzburg.
21. Onderverdeling 1 van Perseel 137 van die plaas Edendale 775, distrik Pietermaritzburg.
22. Onderverdeling 2 van Perseel 137 van die plaas Edendale 775, distrik Pietermaritzburg.
23. Onderverdeling 3 van Perseel 136 van die plaas Edendale 775, distrik Pietermaritzburg.
24. Perseel 754 van die plaas Edendale 775, distrik Pietermaritzburg.
25. Onderverdeling 24 van Perseel 283 van die plaas Edendale 775, distrik Pietermaritzburg.
26. Restant van Onderverdeling 3 van Perseel 253 van die plaas Edendale 775, distrik Pietermaritzburg.
27. Perseel 143 van die plaas Edendale 775, distrik Pietermaritzburg.
28. Perseel 744 van die plaas Edendale 775, distrik Pietermaritzburg.
29. Perseel 745 van die plaas Edendale 775, distrik Pietermaritzburg.
30. Restant van Perseel 766 van die plaas Edendale 775, distrik Pietermaritzburg.
31. Perseel 693 van die plaas Edendale 775, distrik Pietermaritzburg.
32. Perseel 211 van die plaas Edendale 775, distrik Pietermaritzburg.
33. Perseel 130 van die plaas Edendale 775, distrik Pietermaritzburg.
34. Onderverdeling 1 van Perseel 253 van die plaas Edendale 775, distrik Pietermaritzburg.
35. Perseel 725 van die plaas Edendale 775, distrik Pietermaritzburg.
36. Perseel 657 van die plaas Edendale 775, distrik Pietermaritzburg.
37. Onderverdeling 13 van Perseel 283 van die plaas Edendale 775, distrik Pietermaritzburg.
38. Perseel 746 van die plaas Edendale 775, distrik Pietermaritzburg.
39. Perseel 296 van die plaas Edendale 775, distrik Pietermaritzburg.

11. Remainder of Lot 755 of the farm Edendale 775, District of Pietermaritzburg.
12. Lot 658 of the farm Edendale 775, District of Pietermaritzburg.
13. Remainder of Lot 116 of the farm Edendale 775, District of Pietermaritzburg.
14. Lot 134 of the farm Edendale 775, District of Pietermaritzburg.
15. Lot 653 of the farm Edendale 775, District of Pietermaritzburg.
16. Lot 734 of the farm Edendale 775, District of Pietermaritzburg.
17. Remainder of Lot 736 of the farm Edendale 775, District of Pietermaritzburg.
18. Subdivision 1 of Lot 755 of the farm Edendale 775, District of Pietermaritzburg.
19. Subdivision 3 of Lot 131 of the farm Edendale 775, District of Pietermaritzburg.
20. Subdivision 2 of Lot 131 of the farm Edendale 775, District of Pietermaritzburg.
21. Subdivision 1 of Lot 137 of the farm Edendale 775, District of Pietermaritzburg.
22. Subdivision 2 of Lot 137 of the farm Edendale 775, District of Pietermaritzburg.
23. Subdivision 3 of Lot 136 of the farm Edendale 775, District of Pietermaritzburg.
24. Lot 754 of the farm Edendale 775, District of Pietermaritzburg.
25. Subdivision 24 of Lot 283 of the farm Edendale 775, District of Pietermaritzburg.
26. Remainder of Subdivision 3 of Lot 253 of the farm Edendale 775, District of Pietermaritzburg.
27. Lot 143 of the farm Edendale 775, District of Pietermaritzburg.
28. Lot 744 of the farm Edendale 775, District of Pietermaritzburg.
29. Lot 745 of the farm Edendale 775, District of Pietermaritzburg.
30. Remainder of Lot 766 of the farm Edendale 775, District of Pietermaritzburg.
31. Lot 693 of the farm Edendale 775, District of Pietermaritzburg.
32. Lot 211 of the farm Edendale 775, District of Pietermaritzburg.
33. Lot 130 of the farm Edendale 775, District of Pietermaritzburg.
34. Subdivision 1 of Lot 253 of the farm Edendale 775, District of Pietermaritzburg.
35. Lot 725 of the farm Edendale 775, District of Pietermaritzburg.
36. Lot 657 of the farm Edendale 775, District of Pietermaritzburg.
37. Subdivision 13 of Lot 283 of the farm Edendale 775, District of Pietermaritzburg.
38. Lot 746 of the farm Edendale 775, District of Pietermaritzburg.
39. Lot 296 of the farm Edendale 775, District of Pietermaritzburg.

40. Restant van Perseel 297 van die plaas Edendale 775, distrik Pietermaritzburg.
41. Restant van Perseel 316 van die plaas Edendale 775, distrik Pietermaritzburg.
42. Onderverdeling 10 van Perseel 212 van die plaas Edendale 775, distrik Pietermaritzburg.
43. Onderverdeling 1 van Perseel 316 van die plaas Edendale 775, distrik Pietermaritzburg.
44. Onderverdeling 2 van Perseel 316 van die plaas Edendale 775, distrik Pietermaritzburg.
45. Restant van Perseel 255 van die plaas Edendale 775, distrik Pietermaritzburg.
46. Onderverdeling 11 van Perseel 256 van die plaas Edendale 775, distrik Pietermaritzburg.
47. Restant van Perseel 350 van die plaas Edendale 775, distrik Peitermaritzburg.
48. Restant van Perseel 213 van die plaas Edendale 775, distrik Pietermaritzburg.
49. Restant van Onderverdeling 3 van Perseel 297 van die plaas Edendale 775, distrik Pietermaritzburg.
50. Perseel 150 van die plaas Edendale 775, distrik Pietermaritzburg.
51. Onderverdeling 2 van Perseel 346 van die plaas Edendale 775, distrik Pietermaritzburg.
52. Restant van Perseel 346 van die plaas Edendale 775, distrik Pietermaritzburg.
53. Perseel 239 van die plaas Edendale 775, distrik Pietermaritzburg.
54. Onderverdeling 1 van Perseel 350 van die plaas Edendale 775, distrik Pietermaritzburg.
55. Onderverdeling 18 van Perseel 342 van die plaas Edendale 775, distrik Pietermaritzburg.
56. Perseel 284 van die plaas Edendale 775, distrik Pietermaritzburg.
57. Perseel 294 van die plaas Edendale 775, distrik Pietermaritzburg.
58. Restant van Perseel 235 van die plaas Edendale 775, distrik Pietermaritzburg.
59. Onderverdeling 3 van Perseel 235 van die plaas Edendale 775, distrik Pietermaritzburg.
60. Onderverdeling 5 van Perseel 235 van die plaas Edendale 775, distrik Pietermaritzburg.
61. Restant van Onderverdeling 4 van Perseel 8 van die plaas Edendale 775, distrik Pietermaritzburg.
62. Perseel 141 van die plaas Edendale 775, distrik Pietermaritzburg.
63. Onderverdeling 9 van Perseel 255 van die plaas Edendale 775, distrik Pietermaritzburg.
64. Perseel 240 van die plaas Edendale 775, distrik Pietermaritzburg.
65. Restant van Perseel 52 van die plaas Edendale 775, distrik Pietermaritzburg.
66. Perseel 53 van die plaas Edendale 775, distrik Pietermaritzburg.
67. Restant van Perseel 98 van die plaas Edendale 775, distrik Pietermaritzburg.

40. Remainder of Lot 297 of the farm Edendale 775, District of Pietermaritzburg.
41. Remainder of Lot 316 of the farm Edendale 775, District of Pietermaritzburg.
42. Subdivision 10 of Lot 212 of the farm Edendale 775, District of Pietermaritzburg.
43. Subdivision 1 of Lot 316 of the farm Edendale 775, District of Pietermaritzburg.
44. Subdivision 2 of Lot 316 of the farm Edendale 775, District of Pietermaritzburg.
45. Remainder of Lot 255 of the farm Edendale 775, District of Pietermaritzburg.
46. Subdivision 11 of Lot 256 of the farm Edendale 775, District of Pietermaritzburg.
47. Remainder of Lot 350 of the farm Edendale 775, District of Pietermaritzburg.
48. Remainder of Lot 213 of the farm Edendale 775, District of Pietermaritzburg.
49. Remainder of Subdivision 3 of Lot 297 of the farm Edendale 775, District of Pietermaritzburg.
50. Lot 150 of the farm Edendale 775, District of Pietermaritzburg.
51. Subdivision 2 of Lot 346 of the farm Edendale 775, District of Pietermaritzburg.
52. Remainder of Lot 346 of the farm Edendale 775, District of Pietermaritzburg.
53. Lot 239 of the farm Edendale 775, District of Pietermaritzburg.
54. Subdivision 1 of Lot 350 of the farm Edendale 775, District of Pietermaritzburg.
55. Subdivision 18 of Lot 342 of the farm Edendale 775, District of Pietermaritzburg.
56. Lot 284 of the farm Edendale 775, District of Pietermaritzburg.
57. Lot 294 of the farm Edendale 775, District of Pietermaritzburg.
58. Remainder of Lot 235 of the farm Edendale 775, District of Pietermaritzburg.
59. Subdivision 3 of Lot 235 of the farm Edendale 775, District of Pietermaritzburg.
60. Subdivision 5 of Lot 235 of the farm Edendale 775, District of Pietermaritzburg.
61. Remainder of Subdivision 4 of Lot 8 of the farm Edendale 775, District of Pietermaritzburg.
62. Lot 141 of the farm Edendale 775, District of Pietermaritzburg.
63. Subdivision 9 of Lot 255 of the farm Edendale 775, District of Pietermaritzburg.
64. Lot 240 of the farm Edendale 775, District of Pietermaritzburg.
65. Remainder of Lot 52 of the farm Edendale 775, District of Pietermaritzburg.
66. Lot 53 of the farm Edendale 775, District of Pietermaritzburg.
67. Remainder of Lot 98 of the farm Edendale 775, District of Pietermaritzburg.

68. Onderverdeling 1 van Perseel 266 van die plaas Edendale 775, distrik Pietermaritzburg.
69. Restant van Onderverdeling 1 van Perseel 59 van die plaas Edendale 775, distrik Pietermaritzburg.
70. Onderverdeling 1 van Perseel 306 van die plaas Edendale 775, distrik Pietermaritzburg.
71. Onderverdeling 16 van Perseel 522 van die plaas Edendale 775, distrik Pietermaritzburg.
72. Onderverdeling 11 van Perseel 522 van die plaas Edendale 775, distrik Pietermaritzburg.
73. Onderverdeling 10 van Perseel 522 van die plaas Edendale 775, distrik Pietermaritzburg.
74. Perseel 339 van die plaas Edendale 775, distrik Pietermaritzburg.
75. Restant van Perseel 246 van die plaas Edendale 775, distrik Pietermaritzburg.
76. Onderverdeling 2 van Perseel 101 van die plaas Edendale 775, distrik Pietermaritzburg.
77. Onderverdeling 1 van Perseel 101 van die plaas Edendale 775, distrik Pietermaritzburg.
78. Onderverdeling 4 van Perseel 522 van die plaas Edendale 775, distrik Pietermaritzburg.
79. Perseel 340 van die plaas Edendale 775, distrik Pietermaritzburg.
80. Restant van Perseel 101 van die plaas Edendale 775, distrik Pietermaritzburg.
81. Perseel 299 van die plaas Edendale 775, distrik Pietermaritzburg.
82. Restant van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.
83. Restant van Perseel 266 van die plaas Edendale 775, distrik Pietermaritzburg.
84. Restant van Perseel 244 van die plaas Edendale 775, distrik Pietermaritzburg.
85. Onderverdeling 8 van Perseel 300 van die plaas Edendale 775, distrik Pietermaritzburg.
86. Onderverdeling 5 van Perseel 244 van die plaas Edendale 775, distrik Pietermaritzburg.
87. Onderverdeling 6 van Perseel 244 van die plaas Edendale 775, distrik Pietermaritzburg.
88. Restant van Perseel 215 van die plaas Edendale 775, distrik Pietermaritzburg.
89. Onderverdeling 1 van Perseel 300 van die plaas Edendale 775, distrik Pietermaritzburg.
90. Onderverdeling 23 van Perseel 300 van die plaas Edendale 775, distrik Pietermaritzburg.
91. Onderverdeling 5 van Perseel 300 van die plaas Edendale 775, distrik Pietermaritzburg.
92. Onderverdeling 35 van Perseel 300 van die plaas Edendale 775, distrik Pietermaritzburg.
93. Perseel 245 van die plaas Edendale 775, distrik Pietermaritzburg.
94. Onderverdeling 20 van Perseel 300 van die plaas Edendale 775, distrik Pietermaritzburg.
95. Onderverdeling 29 van Perseel 300 van die plaas Edendale 775, distrik Pietermaritzburg.

68. Subdivision 1 of Lot 266 of the farm Edendale 775, District of Pietermaritzburg.
69. Remainder of Subdivision 1 of Lot 59 of the farm Edendale 775, District of Pietermaritzburg.
70. Subdivision 1 of Lot 306 of the farm Edendale 775, District of Pietermaritzburg.
71. Subdivision 16 of Lot 522 of the farm Edendale 775, District of Pietermaritzburg.
72. Subdivision 11 of Lot 522 of the farm Edendale 775, District of Pietermaritzburg.
73. Subdivision 10 of Lot 522 of the farm Edendale 775, District of Pietermaritzburg.
74. Lot 339 of the farm Edendale 775, District of Pietermaritzburg.
75. Remainder of Lot 246 of the farm Edendale 775, District of Pietermaritzburg.
76. Subdivision 2 of Lot 101 of the farm Edendale 775, District of Pietemaritzburg.
77. Subdivision 1 of Lot 101 of the farm Edendale 775, District of Pietermartizburg.
78. Subdivision 4 of Lot 522 of the farm Edendale 775, District of Pietermaritzburg.
79. Lot 340 of the farm Edendale 775, District of Pietermaritzburg.
80. Remainder of Lot 101 of the farm Edendale 775, District of Pietermaritzburg.
81. Lot 299 of the farm Edendale 775, District of Pietermaritzburg.
82. Remainder of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.
83. Remainder of Lot 266 of the farm Edendale 775, District of Pietermaritzburg.
84. Remainder of Lot 244 of the farm Edendale 775, District of Pietermaritzburg.
85. Subdivision 8 of Lot 300 of the farm Edendale 775, District of Pietermaritzburg.
86. Subdivision 5 of Lot 244 of the farm Edendale 775, District of Pietermaritzburg.
87. Subdivision 6 of Lot 244 of the farm Edendale 775, District of Pietermaritzburg.
88. Remainder of Lot 215 of the farm Edendale 775, District of Pietermaritzburg.
89. Subdivision 1 of Lot 300 of the farm Edendale 775, District of Pietermaritzburg.
90. Subdivision 23 of Lot 300 of the farm Edendale 775, District of Pietermaritzburg.
91. Subdivision 5 of Lot 300 of the farm Edendale 775, District of Pietermaritzburg.
92. Subdivision 35 of Lot 300 of the farm Edendale 775, District of Pietermaritzburg.
93. Lot 245 of the farm Edendale 775, District of Pietermaritzburg.
94. Subdivision 20 of Lot 300 of the farm Edendale 775, District of Pietermaritzburg.
95. Subdivision 29 of Lot 300 of the farm Edendale 775, District of Pietermaritzburg.

96. Onderverdeling 2 van Perseel 522 van die plaas Edendale 775, distrik Pietermaritzburg.
97. Onderverdeling 8 van Perseel 305 van die plaas Edendale 775, distrik Pietermaritzburg.
98. Onderverdeling 3 van Perseel 799 van die plaas Edendale 775, distrik Pietermaritzburg.
99. Onderverdeling 1 van Perseel 218 van die plaas Edendale 775, distrik Pietermaritzburg.
100. Perseel 789 van die plaas Edendale 775, distrik Pietermaritzburg.
101. Onderverdeling 7 van Perseel 214 van die plaas Edendale 775, distrik Pietermaritzburg.
102. Perseel 149 van die plaas Edendale 775, distrik Pietermaritzburg.
103. Onderverdeling 23 van Perseel 217 van die plaas Edendale 775, distrik Pietermaritzburg.
104. Onderverdeling 6 van Perseel 214 van die plaas Edendale 775, distrik Pietermaritzburg.
105. Perseel 139 van die plaas Edendale 775, distrik Pietermaritzburg.
106. Perseel 148 van die plaas Edendale 775, distrik Pietermaritzburg.
107. Onderverdeling 7 van Perseel 799 van die plaas Edendale 775, distrik Pietermaritzburg.
108. Perseel 248 van die plaas Edendale 775, distrik Pietermaritzburg.
109. Perseel 138 van die plaas Edendale 775, distrik Pietermaritzburg.
110. Perseel 799 van die plaas Edendale 775, distrik Pietermaritzburg.
111. Perseel 222 van die plaas Edendale 775, distrik Pietermaritzburg.
112. Onderverdeling 2 van Perseel 214 van die plaas Edendale 775, distrik Pietermaritzburg.
113. Onderverdeling 96 van Onderverdeling 79 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.
114. Perseel 313 van die plaas Edendale 775, distrik Pietermaritzburg.
115. Onderverdeling 4 van Perseel 217 van die plaas Edendale 775, distrik Pietermaritzburg.
116. Onderverdeling 1 van Perseel 799 van die plaas Edendale 775, distrik Pietermaritzburg.
117. Onderverdeling 15 van Perseel 217 van die plaas Edendale 775, distrik Pietermaritzburg.
118. Onderverdeling 6 van Perseel 799 van die plaas Edendale 775, distrik Pietermaritzburg.
119. Restant van Perseel 356 van die plaas Edendale 775, distrik Pietermaritzburg.
120. Perseel 242 van die plaas Edendale 775, distrik Pietermaritzburg.
121. Perseel 65 van die plaas Edendale 775, distrik Pietermaritzburg.
122. Onderverdeling 10 van Perseel 799 van die plaas Edendale 775, distrik Pietermaritzburg.
123. Onderverdeling 17 van Perseel 217 van die plaas Edendale 775, distrik Pietermaritzburg.
124. Perseel 250 van die plaas Edendale 775, distrik Pietermaritzburg.

96. Subdivision 2 of Lot 522 of the farm Edendale 775, District of Pietermaritzburg.
97. Subdivision 8 of Lot 305 of the farm Edendale 775, District of Pietermaritzburg.
98. Subdivision 3 of Lot 799 of the farm Edendale 775, District of Pietermaritzburg.
99. Subdivision 1 of Lot 218 of the farm Edendale 775, District of Pietermaritzburg.
100. Lot 789 of the farm Edendale 775, District of Pietermaritzburg.
101. Subdivision 7 of Lot 214 of the farm Edendale 775, District of Pietermaritzburg.
102. Lot 149 of the farm Edendale 775, District of Pietermaritzburg.
103. Subdivision 23 of Lot 217 of the farm Edendale 775, District of Pietermaritzburg.
104. Subdivision 6 of Lot 214 of the farm Edendale 775, District of Pietermaritzburg.
105. Lot 139 of the farm Edendale 775, District of Pietermaritzburg.
106. Lot 148 of the farm Edendale 775, District of Pietermaritzburg.
107. Subdivision 7 of Lot 799 of the farm Edendale 775, District of Pietermaritzburg.
108. Lot 248 of the farm Edendale 775, District of Pietermaritzburg.
109. Lot 138 of the farm Edendale 775, District of Pietermaritzburg.
110. Lot 799 of the farm Edendale 775, District of Pietermaritzburg.
111. Lot 222 of the farm Edendale 775, District of Pietermaritzburg.
112. Subdivision 2 of Lot 214 of the farm Edendale 775, District of Pietermaritzburg.
113. Subdivision 96 of Subdivision 79 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.
114. Lot 313 of the farm Edendale 775, District of Pietermaritzburg.
115. Subdivision 4 of Lot 217 of the farm Edendale 775, District of Pietermaritzburg.
116. Subdivision 1 of Lot 799 of the farm Edendale 775, District of Pietermaritzburg.
117. Subdivision 15 of Lot 217 of the farm Edendale 775, District of Pietermaritzburg.
118. Subdivision 6 of Lot 799 of the farm Edendale 775, District of Pietermaritzburg.
119. Remainder of Lot 356 of the farm Edendale 775, District of Pietermaritzburg.
120. Lot 242 of the farm Edendale 775, District of Pietermaritzburg.
121. Lot 65 of the farm Edendale 775, District of Pietermaritzburg.
122. Subdivision 10 of Lot 799 of the farm Edendale 775, District of Pietermaritzburg.
123. Subdivision 17 of Lot 217 of the farm Edendale 775, District of Pietermaritzburg.
124. Lot 250 of the farm Edendale 775, District of Pietermaritzburg.

125. Onderverdeling 6 van Perseel 217 van die plaas Edendale 775, distrik Pietermaritzburg.
126. Restant van Perseel 217 van die plaas Edendale 775, distrik Pietermaritzburg.
127. Onderverdeling 2 van Perseel 799 van die plaas Edendale 775, distrik Pietermaritzburg.
128. Onderverdeling 1 van Perseel 214 van die plaas Edendale 775, distrik Pietermaritzburg.
129. Perseel 146 van die plaas Edendale 775, distrik Pietermaritzburg.
130. Onderverdeling 1 van Perseel 93 van die plaas Edendale 775, distrik Pietermaritzburg.
131. Perseel 96 van die plaas Edendale 775, distrik Pietermaritzburg.
132. Onderverdeling 11 van Perseel 214 van die plaas Edendale 775, distrik Pietermaritzburg.
133. Onderverdeling 26 van Perseel 217 van die plaas Edendale 775, distrik Pietermaritzburg.
134. Onderverdeling 12 van Perseel 214 van die plaas Edendale 775, distrik Pietermaritzburg.
135. Onderverdeling 25 van Perseel 217 van die plaas Edendale 775, distrik Pietermaritzburg.
136. Onderverdeling 27 van Perseel 217 van die plaas Edendale 775, distrik Pietermaritzburg.
137. Onderverdeling 77 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.
138. Onderverdeling 78 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.
139. Onderverdeling 121 van Onderverdeling 69 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.
140. Onderverdeling 118 van Onderverdeling 69 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.
141. Onderverdeling 84 van Onderverdeling 75 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.
142. Onderverdeling 6 van Perseel 97 van die plaas Edendale 775, distrik Pietermaritzburg.
143 .Onderverdeling 3 van Perseel 97 van die plaas Edendale 775, distrik Pietermaritzburg.
144. Onderverdeling 2 van Perseel 97 van die plaas Edendale 775, distrik Pietermaritzburg.
145. Onderverdeling 9 van Perseel 799 van die plaas Edendale 775, distrik Pietermaritzburg.
146. Perseel 94 van die plaas Edendale 775, distrik Pietermaritzburg.
147. Onderverdeling 6 van Onderverdeling 4 van Perseel 101 van die plaas Edendale 775, distrik Pietermaritzburg.
148. Perseel 332 van die plaas Edendale 775, distrik Pietermaritzburg.
149. Perseel 331 van die plaas Edendale 775, distrik Pietermaritzburg.
150. Onderverdeling 10 van Perseel 214 van die plaas Edendale 775, distrik Pietermaritzburg.

125. Subdivision 6 of Lot 217 of the farm Edendale 775, District of Pietermaritzburg.
126. Remainder of Lot 217 of the farm Edendale 775, District of Pietermaritzburg.
127. Subdivision 2 of Lot 799 of the farm Edendale 775, District of Pietermaritzburg.
128. Subdivision 1 of Lot 214 of the farm Edendale 775, District of Pietermaritzburg.
129. Lot 146 of the farm Edendale 775, District of Pietermaritzburg.
130. Subdivision 1 of Lot 93 of the farm Edendale 775, District of Pietermaritzburg.
131. Lot 96 of the farm Edendale 775, District of Pietermaritzburg.
132. Subdivision 11 of Lot 214 of the farm Edendale 775, District of Pietermaritzburg.
133. Subdivision 26 of Lot 217 of the farm Edendale 775, District of Pietermaritzburg.
134. Subdivision 12 of Lot 214 of the farm Edendale 775, District of Pietermaritzburg.
135. Subdivision 25 of Lot 217 of the farm Edendale 775, District of Pietermaritzburg.
136. Subdivision 27 of Lot 217 of the farm Edendale 775, District of Pietermaritzburg.
137. Subdivision 77 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.
138. Subdivision 78 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.
139. Subdivision 121 of Subdivision 69 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.
140. Subdivision 118 of Subdivision 69 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.
141. Subdivision 84 of Subdivision 75 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.
142. Subdivision 6 of Lot 97 of the farm Edendale 775, District of Pietermaritzburg.
143. Subdivision 3 of Lot 97 of the farm Edendale 775, District of Pietermaritzburg.
144. Subdivision 2 of Lot 97 of the farm Edendale 775, District of Pietermaritzburg.
145. Subdivision 9 of Lot 799 of the farm Edendale 775, District of Pietermaritzburg.
146. Lot 94 of the farm Edendale 775, District of Pietermaritzburg.
147. Subdivision 6 of Subdivision 4 of Lot 101 of the farm Edendale 775, District of Pietermaritzburg.
148. Lot 332 of the farm Edendale 775, District of Pietermaritzburg.
149. Lot 331 of the farm Edendale 775, District of Pietermaritzburg.
150. Subdivision 10 of Lot 214 of the farm Edendale 775, District of Pietermaritzburg.

151. Onderverdeling 2 van Perseel 188 van die plaas Edendale 775, distrik Pietermaritzburg.

152. Restant van Perseel 97 van die plaas Edendale 775, distrik Pietermaritzburg.

153. Onderverdeling 19 van Perseel 217 van die plaas Edendale 775, distrik Pietermaritzburg.

154. Onderverdeling 20 van Perseel 217 van die plaas Edendale 775, distrik Pietermaritzburg.

155. Onderverdeling 112 van Onderverdeling 69 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.

156. Restant van Perseel 188 van die plaas Edendale 775, distrik Pietermaritzburg.

157. Perseel 92 van die plaas Edendale 775, distrik Pietermaritzburg.

158. Perseel 696 van die plaas Edendale 775, distrik Pietermaritzburg.

159. Onderverdeling 82 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.

160. Perseel 701 van die plaas Edendale 775, distrik Pietermaritzburg.

161. Onderverdeling 68 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.

162. Restant van Perseel 685 van die plaas Edendale 775, distrik Pietermaritzburg.

163. Restant van Perseel 717 van die plaas Edendale 775, distrik Pietermaritzburg.

164. Restant van Perseel 687 van die plaas Edendale 775, distrik Pietermaritzburg.

165. Perseel 765 van die plaas Edendale 775, distrik Pietermaritzburg.

166. Perseel 676 van die plaas Edendale 775, distrik Pietermaritzburg.

167. Restant van Onderverdeling 60 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.

168. Restant van Perseel 769 van die plaas Edendale 775, distrik Pietermaritzburg.

169. Onderverdeling 107 van Onderverdeling 55 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.

170. Onderverdeling 44 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.

171. Onderverdeling 63 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.

172. Restant van Perseel 680 van die plaas Edendale 775, distrik Pietermaritzburg.

173. Onderverdeling 43 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.

174. Onderverdeling 41 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.

175. Onderverdeling 1 van Perseel 685 van die plaas Edendale 775, distrik Pietermaritzburg.

176. Onderverdeling 58 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.

177. Onderverdeling 1 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.

151. Subdivision 2 of Lot 188 of the farm Edendale 775, District of Pietermaritzburg.

152. Remainder of Lot 97 of the farm Edendale 775, District of Pietermaritzburg.

153. Subdivision 19 of Lot 217 of the farm Edendale 775, District of Pietermaritzburg.

154. Subdivision 20 of Lot 217 of the farm Edendale 775, District of Pietermaritzburg.

155. Subdivision 112 of Subdivision 69 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.

156. Remaider of Lot 188 of the farm Edendale 775, District of Pietermaritzburg.

157. Lot 92 of the farm Edendale 775, District of Pietermaritzburg.

158. Lot 696 of the farm Edendale 775, District of Pietermaritzburg.

159. Subdivision 82 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.

160. Lot 701 of the farm Edendale 775, District of Pietermaritzburg.

161. Subdivision 68 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.

162. Remainder of Lot 685 of the farm Edendale 775, District of Pietermaritzburg.

163. Remainder of Lot 717 of the farm Edendale 775, District of Pietermaritzburg.

164. Remainder of Lot 687 of the farm Edendale 775, District of Pietermaritzburg.

165. Lot 765 of the farm Edendale 775, District of Pietermaritzburg.

166. Lot 676 of the farm Edendale 775, District of Pietermaritzburg.

167. Remainder of Subdivision 60 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.

168. Remainder of Lot 769 of the farm Edendale 775, District of Pietermaritzburg.

169. Subdivision 107 of Subdivision 55 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.

170. Subdivision 44 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.

171. Subdivision 63 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.

172. Remainder of Lot 680 of the farm Edendale 775, District of Pietermaritzburg.

173. Subdivision 43 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.

174. Subdivision 41 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.

175. Subdivision 1 of Lot 685 of the farm Edendale 775, District of Pietermaritzburg.

176. Subdivision 58 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.

177. Subdivision 1 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.

178. Onderverdeling 2 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.
179. Perseel 671 van die plaas Edendale 775, distrik Pietermaritzburg.
180. Perseel 694 van die plaas Edendale 775, distrik Pietermaritzburg.
181. Restant van Perseel 768 van die plaas Edendale 775, distrik Pietermaritzburg.
182. Perseel 698 van die plaas Edendale 775, distrik Pietermaritzburg.
183. Onderverdeling 38 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.
184. Onderverdeling 67 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.
185. Perseel 695 van die plaas Edendale 775, distrik Pietermaritzburg.
186. Onderverdeling 42 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.
187. Perseel 709 van die plaas Edendale 775, distrik Pietermaritzburg.
188. Perseel 708 van die plaas Edendale 775, distrik Pietermaritzburg.
189. Onderverdeling 30 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.
190. Onderverdeling 83 van Onderverdeling 55 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.
191. Onderverdeling 66 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.
192. Perseel 660 van die plaas Edendale 775, distrik Pietermaritzburg.
193. Perseel 667 van die plaas Edendale 775, distrik Pietermaritzburg.
194. Perseel 699 van die plaas Edendale 775, distrik Pietermaritzburg.
195. Onderverdeling 13 van Perseel 803 van die plaas Edendale 775, distrik Pietermaritzburg.
196. Onderverdeling 9 van Perseel 803 van die plaas Edendale 775, distrik Pietermaritzburg.
197. Onderverdeling 178 van Onderverdeling 20 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.
198. Onderverdeling 20 van Perseel 803 van die plaas Edendale 775, distrik Pietermaritzburg.
199. Perseel 663 van die plaas Edendale 775, distrik Pietermaritzburg.
200. Restant van Onderverdeling 55 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.
201. Onderverdeling 43 van Perseel 803 van die plaas Edendale 775, distrik Pietermaritzburg.
202. Onderverdeling 30 van Perseel 803 van die plaas Edendale 775, distrik Pietermaritzburg.
203. Perseel 665 van die plaas Edendale 775, distrik Pietermaritzburg.
204. Perseel 762 van die plaas Edendale 775, distrik Pietermaritzburg.

178. Subdivision 2 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.
179. Lot 671 of the farm Edendale 775, District of Pietermaritzburg.
180. Lot 694 of the farm Edendale 775, District of Pietermaritzburg.
181. Remainder of Lot 768 of the farm Edendale 775, District of Pietermaritzburg.
182. Lot 698 of the farm Edendale 775, District of Pietermaritzburg.
183. Subdivision 38 of Lot 796 of the farm Edendale 755, District of Pietermaritzburg.
184. Subdivision 67 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.
185. Lot 695 of the farm Edendale 775, District of Pietermaritzburg.
186. Subdivision 42 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.
187. Lot 709 of the farm Edendale 775, District of Pietermaritzburg.
188. Lot 708 of the farm Edendale 775, District of Pietermaritzburg.
189. Subdivision 30 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.
190. Subdivision 83 of Subdivision 55 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.
191. Subdivision 66 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.
192. Lot 660 of the farm Edendale 775, District of Pietermaritzburg.
193. Lot 667 of the farm Edendale 775, District of Pietermaritzburg.
194. Lot 699 of the farm Edendale 775, District of Pietermaritzburg.
195. Subdivision 13 of Lot 803 of the farm Edendale 775, District of Pietermaritzburg.
196. Subdivision 9 of Lot 803 of the farm Edendale 775, District of Pietermaritzburg.
197. Subdivision 178 of Subdivision 20 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.
198. Subdivision 20 of Lot 803 of the farm Edendale 775, District of Pietermaritzburg.
199. Lot 663 of the farm Edendale 775, District of Pietermaritzburg.
200. Remainder of Subdivision 55 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.
201. Subdivision 43 of Lot 803 of the farm Edendale 775, District of Pietermaritzburg.
202. Subdivision 30 of Lot 803 of the farm Edendale 775, District of Pietermaritzburg.
203. Lot 665 of the farm Edendale 775, District of Pietermaritzburg.
204. Lot 762 of the farm Edendale 775, District of Pietermaritzburg.

205. Restant van Onderverdeling 5 van Perseel 210 van die plaas Edendale 775, distrik Pietermaritzburg.
206. Onderverdeling 55 van Onderverdeling 1 van Perseel 797 van die plaas Edendale 775, distrik Pietermaritzburg.
207. Perseel 686 van die plaas Edendale 775, distrik Pietermaritzburg.
208. Onderverdeling 72 van Onderverdeling 3 van Perseel 797 van die plaas Edendale 775, distrik Pietermaritzburg.
209. Onderverdeling 73 van Onderverdeling 3 van Perseel 797 van die plaas Edendale 775, distrik Pietermaritzburg.
210. Onderverdeling 78 van Onderverdeling 3 van Perseel 797 van die plaas Edendale 775, distrik Pietermaritzburg.
211. Onderverdeling 79 van Onderverdeling 3 van Perseel 797 van die plaas Edendale 775, distrik Pietermaritzburg.
212. Onderverdeling 85 van Onderverdeling 3 van Perseel 797 van die plaas Edendale 775, distrik Pietermaritzburg.
213. Restant van Onderverdeling 70 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.
214. Onderverdeling 39 van Perseel 803 van die plaas Edendale 775, distrik Pietermaritzburg.
215. Onderverdeling 41 van Perseel 803 van die plaas Edendale 775, distrik Pietermaritzburg.
216. Onderverdeling 45 van Perseel 803 van die plaas Edendale 775, distrik Pietermaritzburg.
217. Onderverdeling 69 van Onderverdeling 3 van Perseel 797 van die plaas Edendale 775, distrik Pietermaritzburg.
218. Onderverdeling 46 van Perseel 803 van die plaas Edendale 775, distrik Pietermaritzburg.
219. Onderverdeling 54 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.
220. Onderverdeling 7 van Perseel 803 van die plaas Edendale 775, distrik Pietermaritzburg.
221. Onderverdeling 32 van Perseel 803 van die plaas Edendale 775, distrik Pietermaritzburg.
222. Onderverdeling 97 van Onderverdeling 79 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.
223. Onderverdeling 17 van Perseel 803 van die plaas Edendale 775, distrik Pietermaritzburg.
224. Onderverdeling 6 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.
225. Onderverdeling 1 van Perseel 687 van die plaas Edendale 775, distrik Pietermaritzburg.
226. Onderverdeling 21 van Perseel 803 van die plaas Edendale 775, distrik Pietermaritzburg.
227. Onderverdeling 4 van Perseel 803 van die plaas Edendale 775, distrik Pietermaritzburg.
228. Onderverdeling 82 van Onderverdeling 3 van Perseel 797 van die plaas Edendale 775, distrik Pietermaritzburg.

205. Remainder of Subdivision 5 of Lot 210 of the farm Edendale 775, District of Pietermaritzburg.
206. Subdivision 55 of Subdivision 1 of Lot 797 of the farm Edendale 775, District of Pietermaritzburg.
207. Lot 686 of the farm Edendale 775, District of Pietermaritzburg.
208. Subdivision 72 of Subdivision 3 of Lot 797 of the farm Edendale 775, District of Pietermaritzburg.
209. Subdivision 73 of Subdivision 3 of Lot 797 of the farm Edendale 775, District of Pietermaritzburg.
210. Subdivision 78 of Subdivision 3 of Lot 797 of the farm Edendale 775, District of Pietermaritzburg.
211. Subdivision 79 of Subdivision 3 of Lot 797 of the farm Edendale 775, District of Pietermaritzburg.
212. Subdivision 85 of Subdivision 3 of Lot 797 of the farm Edendale 775, District of Pietermaritzburg.
213. Remainder of Subdivision 70 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.
214. Subdivision 39 of Lot 803 of the farm Edendale 775, District of Pietermaritzburg.
215. Subdivision 41 of Lot 803 of the farm Edendale 775, District of Pietermaritzburg.
216. Subdivision 45 of Lot 803 of the farm Edendale 775, District of Pietermaritzburg.
217. Subdivision 69 of Subdivision 3 of Lot 797 of the farm Edendale 775, District of Pietermaritzburg.
218. Subdivision 46 of Lot 803 of the farm Edendale 775, District of Pietermaritzburg.
219. Subdivision 54 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.
220. Subdivision 7 of Lot 803 of the farm Edendale 775, District of Pietermaritzburg.
221. Subdivision 32 of Lot 803 of the farm Edendale 775, District of Pietermaritzburg.
222. Subdivision 97 of Subdivision 79 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.
223. Subdivision 17 of Lot 803 of the farm Edendale 775, District of Pietermaritzburg.
224. Subdivision 6 of Lot 796 of the farm Edendae 775, District of Pietermaritzburg.
225. Subdivision 1 of Lot 687 of the farm Edendale 775, District of Pietermaritzburg.
226. Subdivision 21 of Lot 803 of the farm Edendale 775, District of Pietermaritzburg.
227. Subdivision 4 of Lot 803 of the farm Edendale 775, District of Pietermaritzburg.
228. Subdivision 82 of Subdivision 3 of Lot 797 of the farm Edendale 775, District of Pietermaritzburg.

229. Subdivision 81 of Subdivision 3 of Lot 797 of the farm Edendale 775, District of Pietermaritzburg.

230. Subdivision 57 of Subdivision 1 of Lot 797 of the farm Edendale 775, District of Pietermaritzburg.

231. Subdivision 77 of Subdivision 3 of Lot 797 of the farm Edendale 775, District of Pietermaritzburg.

232. Subdivision 58 of Subdivision 1 of Lot 797 of the farm Edendale 775, District of Pietermaritzburg.

233. Subdivision 60 of Subdivision 1 of Lot 797 of the farm Edendale 775, District of Pietermaritzburg.

234. Subdivision 3 of Subdivision 1 of Lot 680 of the farm Edendale 775, District of Pietermaritzburg.

235. Subdivision 37 of Lot 803 of the farm Edendale 775, District of Pietermaritzburg.

236. Subdivision 35 of Lot 803 of the farm Edendale 775, District of Pietermaritzburg.

237. Remainder of Subdivision 1 of Lot 210 of the farm Edendale 775, District of Pietermaritzburg.

238. Subdivision 98 of Subdivision 79 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.

239. Subdivision 80 of Subdivision 3 of Lot 797 of the farm Edendale 775, District of Pietermaritzburg.

240. Subdivision 1 of Lot 307 of the farm Edendale 775, District of Pietermaritzburg.

241. Subdivision 1 of Lot 221 of the farm Edendale 775, District of Pietermartizburg.

242. Subdivision 2 of Lot 307 of the farm Edendale 775, District of Pietermaritzburg.

243. Subdivision 49 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.

244. Subdivision 11 of Lot 221 of the farm Edendale 775, District of Pietermaritzburg.

245. Subdivision 10 of Lot 221 of the farm Edendale 775, District of Pietermaritzburg.

246. Subdivision 54 of Subdivision 1 of Lot 797 of the farm Edendale 775, District of Pietermaritzburg.

247. Subdivision 53 of Subdivision 1 of Lot 797 of the farm Edendale 775, District of Pietermaritzburg.

248. Subdivision 166 of Subdivision 7 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.

249. Subdivision 6 of Lot 208 of the farm Edendale 775, District of Pietermaritzburg.

250. Subdivision 2 of Lot 797 of the farm Edendale 775, District of Pietermaritzburg.

251. Lot 123 of the farm Edendale 775, District of Pietermaritzburg.

229. Onderverdeling 81 van Onderverdeling 3 van Perseel 797 van die plaas Edendale 775, distrik Pietermaritzburg.

230. Onderverdeling 57 van Onderverdeling 1 van Perseel 797 van die plaas Edendale 775, distrik Pietermaritzburg.

231. Onderverdeling 77 van Onderverdeling 3 van Perseel 797 van die plaas Edendale 775, distrik Pietermaritzburg.

232. Onderverdeling 58 van Onderverdeling 1 van Perseel 797 van die plaas Edendale 775, distrik Pietermaritzburg.

233. Onderverdeling 60 van Onderverdeling 1 van Perseel 797 van die plaas Edendale 775, distrik Pietermaritzburg.

234. Onderverdeling 3 van Onderverdeling 1 van Perseel 680 van die plaas Edendale 775, distrik Pietermaritzburg.

235. Onderverdeling 37 van Perseel 803 van die plaas Edendale 775, distrik Pietermaritzburg.

236. Onderverdeling 35 van Perseel 803 van die plaas Edendale 775, distrik Pietermaritzburg.

237. Restant van Onderverdeling 1 van Perseel 210 van die plaas Edendale 775, distrik Pietermaritzburg.

238. Onderverdeling 98 van Onderverdeling 79 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.

239. Onderverdeling 80 van Onderverdeling 3 van Perseel 797 van die plaas Edendale 775, distrik Pietermaritzburg.

240. Onderverdeling 1 van Perseel 307 van die plaas Edendale 775, distrik Pietermaritzburg.

241. Onderverdeling 1 van Perseel 221 van die plaas Edendale 775, distrik Pietermartizburg.

242. Onderverdeling 2 van Perseel 307 van die plaas Edendale 775, distrik Pietermaritzburg.

243. Onderverdeling 49 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.

244. Onderverdeling 11 van Perseel 221 van die plaas Edendale 775, distrik Pietermaritzburg.

245. Onderverdeling 10 van Perseel 221 van die plaas Edendale 775, distrik Pietermaritzburg.

246. Onderverdeling 54 van Onderverdeling 1 van Perseel 797 van die plaas Edendale 775, distrik Pietermaritzburg.

247. Onderverdeling 53 van Onderverdeling 1 van Perseel 797 van die plaas Edendale 775, distrik Pietermaritzburg.

248. Onderverdeling 166 van Onderverdeling 7 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.

249. Onderverdeling 6 van Perseel 208 van die plaas Edendale 775, distrik Pietermaritzburg.

250. Onderverdeling 2 van Perseel 797 van die plaas Edendale 775, distrik Pietermaritzburg.

251. Perseel 123 van die plaas Edendale 775, distrik Pietermaritzburg.

252. Onderverdeling 37 van Perseel 796 van die plaas Edendale 775, distrik Pietermaritzburg.
253. Restant van Perseel 209 van die plaas Edendale 775, distrik Pietermaritzburg.
254. Onderverdeling 21 van Perseel 125 van die plaas Edendale 775, distrik Pietermaritzburg.
255. Onderverdeling 20 van Perseel 125 van die plaas Edendale 775, distrik Pietermaritzburg.
256. Onderverdeling 5 van Perseel 221 van die plaas Edendale 775, distrik Pietermaritzburg.
257. Onderverdeling 4 van Perseel 253 van die plaas Edendale 775, distrik Pietermaritzburg.
258. Onderverdeling 1 van Perseel 246 van die plaas Edendale 775, distrik Pietermaritzburg.

DEPARTEMENT VAN VERVOER

No. 2270 **3 Desember 1993**

WYSIGING VAN PROKLAMASIE No. 163 VAN 1980: NASIONALE ROETE 3, SEKSIE 3: PIETERMARITZBURG-VERBYPAD: PROVINSIE NATAL

Kragtens die bevoegdheid my verleen by artikel 4 (1) (c) van die Wet op Nasionale Paaie, 1971 (Wet No. 54 1971), soos gewysig, wysig ek hierby, op aanbeveling van die Suid-Afrikaanse Padraad, Proklamasie No. 163 van 1980 deur daarin velle 9, 10, 15A en 15B van Plan P310/79 deur bygaande velle 9A, 10A en 15C respektiewelik te vervang.

P. J. WELGEMOED,
Minister van Vervoer.

252. Subdivision 37 of Lot 796 of the farm Edendale 775, District of Pietermaritzburg.
253. Remainder of Lot 209 of the farm Edendale 775, District of Pietermaritzburg.
254. Subdivision 21 of Lot 125 of the farm Edendale 775, District of Pietermaritzburg.
255. Subdivision 20 of Lot 125 of the farm Edendale 775, District of Pietermaritzburg.
256. Subdivision 5 of Lot 221 of the farm Edendale 775, District of Pietermaritzburg.
257. Subdivision 4 of Lot 253 of the farm Edendale 775, District of Pietermaritzburg.
258. Subdivision 1 of Lot 246 of the farm Edendale 775, District of Pietermaritzburg.

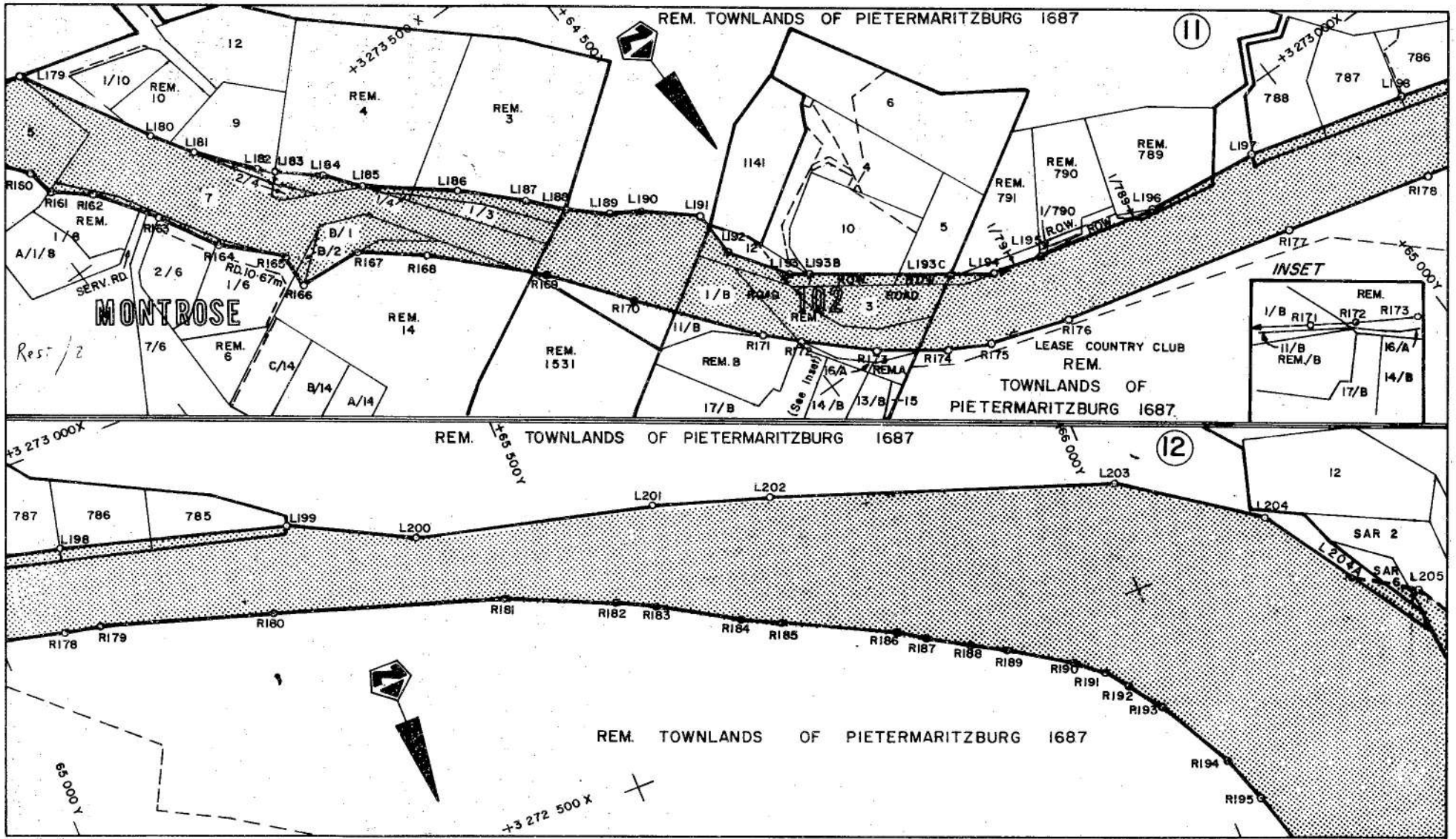
DEPARTMENT OF TRANSPORT

No. 2270 **3 December 1993**

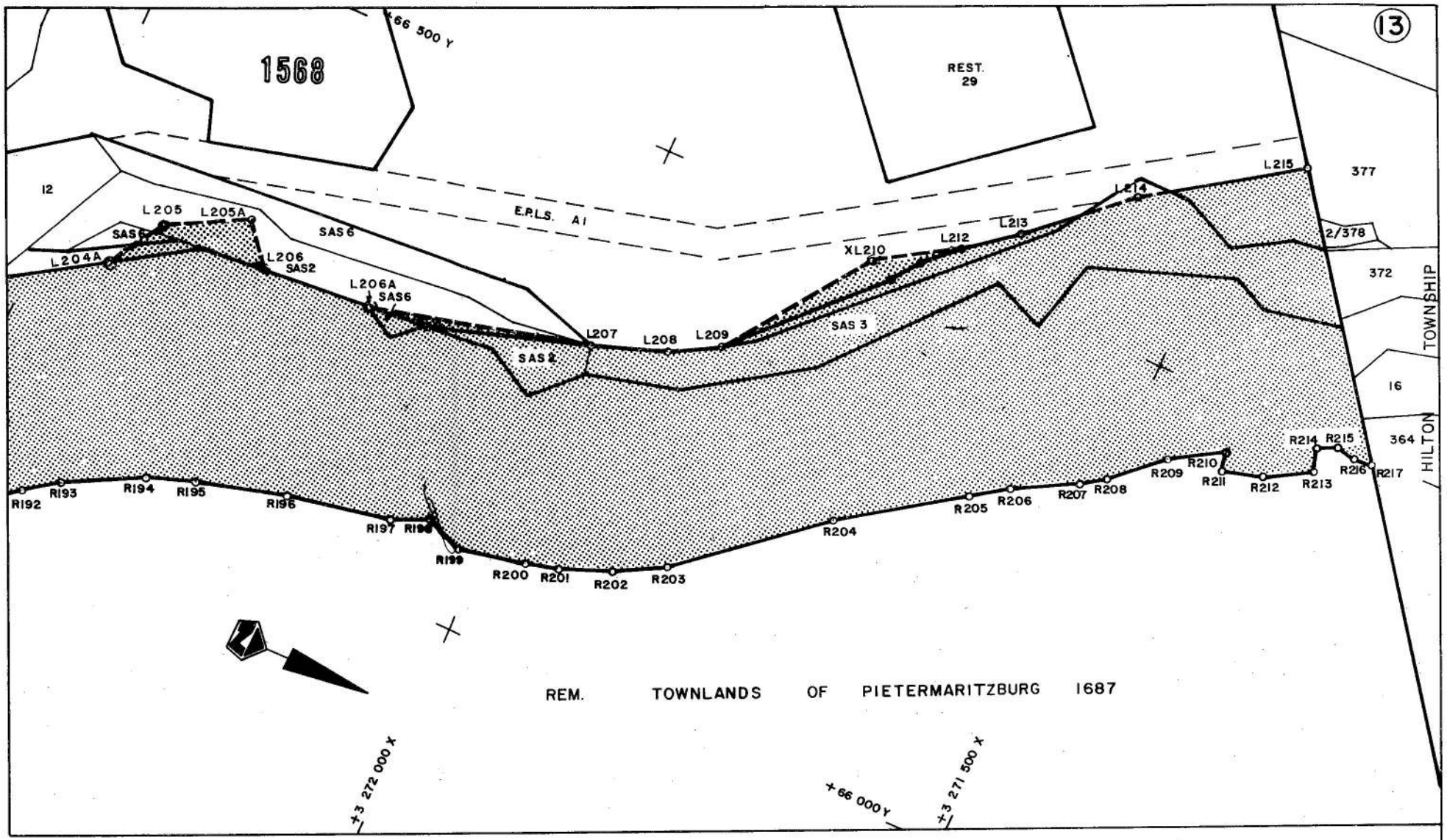
AMENDMENT OF PROCLAMATION No. 163 OF 1980: NATIONAL ROUTE 3, SECTION 3: PIETERMARITZBURG BYPASS: PROVINCE OF NATAL

By virtue of the powers vested in me by section 4 (1) (c) of the National Roads Act, 1971 (Act No. 54 of 1971), as amended, I hereby, on the recommendation of the South African Roads Board, amend Proclamation No. 163 of 1980, by substituting the subjoined sheets 9A, 10A and 15C for sheets 9, 10, 15A and 15B of Plan P310/79 respectively.

P. J. WELGEMOED,
Minister of Transport.



Nasionale Vervoerkommissie National Transport Commission	Die figuur getoon The figure shown		stel die padreserwe voor van 'n gedeelte represents the road reserve of a portion van Nasionale Roete 3 Seksie 3 of National Route Section	Vel 9A van 15 Sheet of P 310/79-



Nasionale Vervoerkommissie National Transport Commission	Die figuur getoon The figure shown		stel die padreserwe voor van 'n gedeelte represents the road reserve of a portion van Nasionale Roete 3 Seksie 3 of National Route Section	Vel 10A van 15 Sheet of P 310/79

PADRESERWE KOÖRDINATE / ROAD RESERVE CO-ORDINATES

LINKERKANT/LEFT HAND SIDE

	Y	X Lo 31°
L168	+63 469.93	+3 274 074.89
L169	+63 525.83	+3 274 067.26
L170	+63 591.27	+3 274 036.90
L171	+63 622.65	+3 274 027.44
L172	+63 684.08	+3 274 033.36
L173	+63 720.39	+3 274 000.50
L174	+63 725.81	+3 273 943.14
L175	+63 825.32	+3 273 944.63
L176	+63 887.49	+3 273 908.87
L177	+63 973.29	+3 273 804.88
L178	+63 955.88	+3 273 782.13
L179	+64 066.87	+3 273 683.82
L180	+64 129.69	+3 273 566.63
L181	+64 152.14	+3 273 529.98
L182	+64 189.64	+3 273 483.55
L183	+64 203.41	+3 273 470.72
L184	+64 235.42	+3 273 442.69
L185	+64 259.01	+3 273 412.22
L186	+64 326.83	+3 273 356.41
L187	+64 374.22	+3 273 310.81
L188	+64 400.42	+3 273 281.45
L189	+64 429.54	+3 273 254.05
L190	+64 455.13	+3 273 238.22
L191	+64 495.71	+3 273 201.92
L192	+64 497.77	+3 273 159.40
L193	+64 530.92	+3 273 108.93
L193B	+64 548.68	+3 273 096.04
L193C	+64 651.82	+3 273 021.15
L194	+64 684.22	+3 272 998.02
L195	+64 727.39	+3 272 985.96
L196	+64 836.49	+3 272 958.61
L197	+64 942.20	+3 272 945.98
L198	+65 084.23	+3 272 908.80
L199	+65 287.79	+3 272 851.30
L200	+65 394.98	+3 272 795.45
L201	+65 612.42	+3 272 741.00
L202	+65 716.98	+3 272 707.70
L203	+66 017.39	+3 272 600.05
L204	+66 134.29	+3 272 520.24
L204A	+66 188.16	+3 272 435.99
L205	+66 240.00	+3 272 402.00
L205A	+66 278.00	+3 272 330.00
L206	+66 241.65	+3 272 297.66
L206A	+66 250.25	+3 272 196.03
L207	+66 303.19	+3 271 989.58
L208	+66 325.91	+3 271 923.05
L209	+66 351.38	+3 271 879.31
XL210	+66 482.00	+3 271 782.00
L212	+66 527.93	+3 271 713.02
L213	+66 563.14	+3 271 669.72
L214	+66 639.71	+3 271 585.55
L215	+66 729.99	+3 271 451.10

REGTERKANT/RIGHT HAND SIDE

	Y	X
R171	+64 476.99	+3 273 078.16
R172	+64 699.28	+3 273.054.73
R173	+64 553.77	+3 273 002.12
R174	+64 606.76	+3 272 964.94
R175	+64 640.33	+3 272 947.12
R176	+64 712.03	+3 272 922.53
R177	+64 927.27	+3 272 869.31
R178	+65 058.76	+3 272 834.07
R179	+65 092.09	+3 272 826.60
R180	+65 245.28	+3 272 778.96
R181	+65 450.74	+3 272 711.07
R182	+65 545.93	+3 272 667.76
R183	+65 580.29	+3 272 650.95
R184	+65 648.42	+3 272 610.58
R185	+65 681.73	+3 272 592.09
R186	+65 777.91	+3 272 545.32
R187	+65 799.64	+3 272 529.08
R188	+65 834.63	+3 272 507.74
R189	+65 864.86	+3 272 491.13
R190	+65 918.72	+3 272 455.72
R191	+65 941.19	+3 272 437.79
R192	+65 957.23	+3 272 418.20
R193	+65 978.78	+3 272 386.55
R194	+66 015.94	+3 272 318.47
R195	+66 031.14	+3 272 274.45
R196	+66 053.10	+3 272 192.75
R197	+66 073.29	+3 272 094.92
R198	+66 089.17	+3 272 061.12
R199	+66 074.45	+3 272 023.42
R200	+66 087.18	+3 271 960.11
R201	+66 095.84	+3 271 930.70
R202	+66 113.41	+3 271 884.72
R203	+66 138.70	+3 271 839.13
R204	+66 242.24	+3 271 716.39
R205	+66 314.84	+3 271 610.78
R206	+66 337.34	+3 271 579.64
R207	+66 368.21	+3 271 523.77
R208	+66 382.52	+3 271 502.49
R209	+66 422.03	+3 271 457.32
R210	+66 450.37	+3 271 409.54
R211	+66 433.64	+3 271 406.74
R212	+66 443.81	+3 271 370.16
R213	+66 467.46	+3 271 329.70
R214	+66 488.77	+3 271 335.85
R215	+66 498.01	+3 271 318.03
R216	+66 494.85	+3 271 298.97
R217	+66 496.36	+3 271 281.85

Vel Sheet	15C	van 15 of	P 310/79

No. 2295 **3 Desember 1993**

WYSIGING VAN PROKLAMASIE No. 200 VAN 1974: NASIONALE ROETE 2, SEKSIE 23: UMZINTO—AMAHLONGWA: DISTRIK UMZINTO, PROVINSIE NATAL

Kragtens die bevoegdheid my verleen by artikel 4 (1) (c) van die Wet op Nasionale Paaie, 1971 (Wet No. 54 van 1971), soos gewysig, wysig ek hierby, op aanbeveling van die Suid-Afrikaanse Padraad, Proklamasie No. 200 van 1974 deur daarin velle 3 en 8 van Plan P230/74 respektiewelik te vervang deur bygaande velle 3A en 8A.

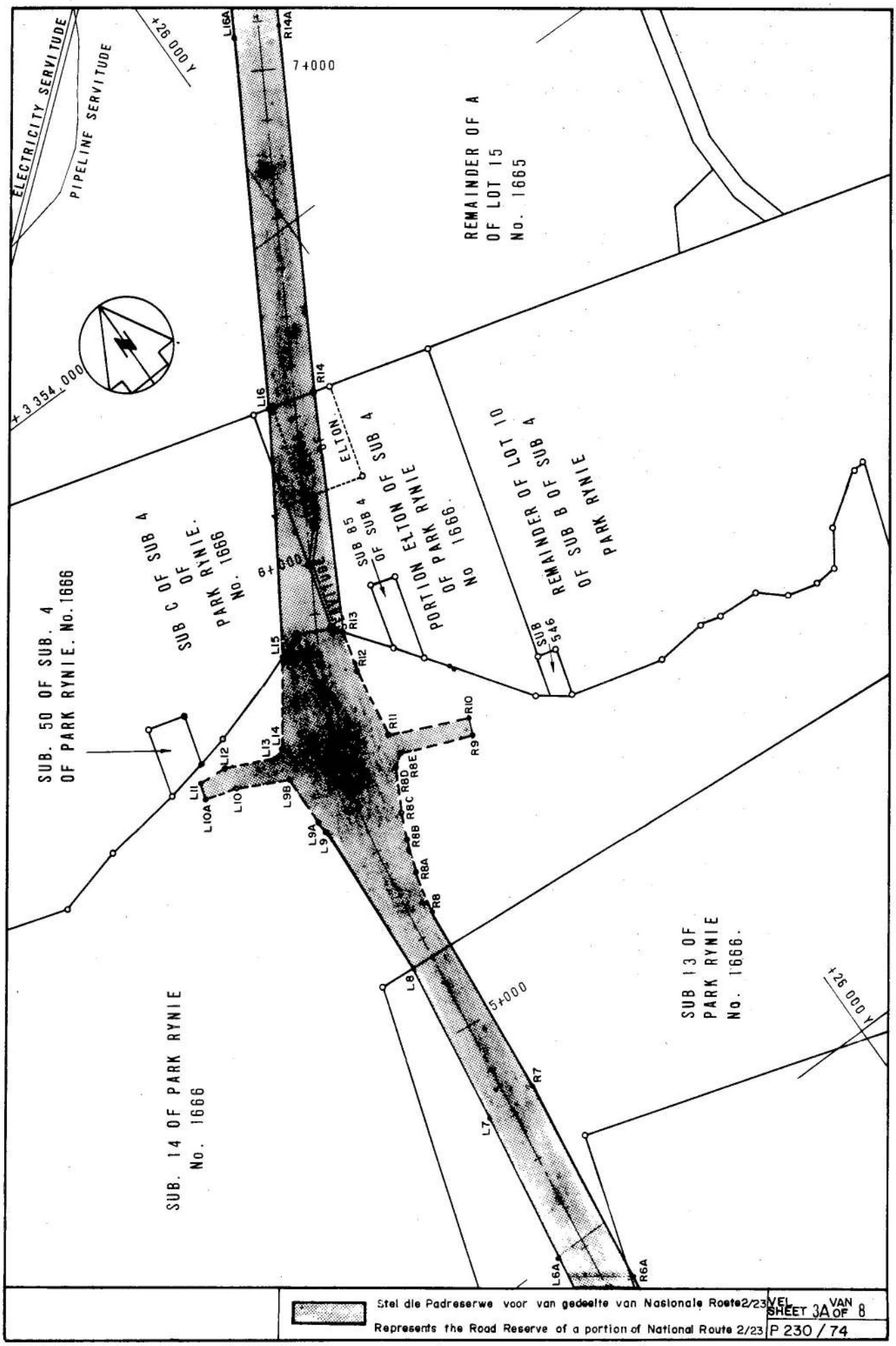
P. J. WELGEMOED,
Minister van Vervoer.

No. 2295 **3 December 1993**

AMENDMENT OF PROCLAMATION No. 200 OF 1974: NATIONAL ROUTE 2, SECTION 23: UMZINTO—AMAHLONGWA: DISTRICT OF UMZINTO, PROVINCE OF NATAL

By virtue of the powers vested in me by section 4 (1) (c) of the National Roads Act, 1971 (Act No. 54 of 1971), as amended, I hereby, on the recommendation of the South African Roads Board, amend Proclamation No. 200 of 1974, by substituting the subjoined sheets 3A and 8A for sheets 3 and 8 of Plan P230/74 respectively.

P. J. WELGEMOED,
Minister of Transport.


ELECTRICITY SERVITUDE
PIPELINE SERVITUDE
+26 000 Y
+3 354 000
7+000
6+000
5+000
REMAINDER OF A
OF LOT 15
No. 1665
SUB C OF SUB 4
OF
PARK RYNIE.
No. 1666
SUB. 50 OF SUB. 4
OF PARK RYNIE. No. 1666
SUB 85
OF SUB 4
OF
PORTION ELTON OF SUB 4
OF PARK RYNIE
No 1666.
ELTON
REMAINDER OF LOT 10
OF SUB 4
OF SUB B
PARK RYNIE
SUB
546
SUB. 14 OF PARK RYNIE
No. 1666
SUB 13 OF
PARK RYNIE
No. 1666.
L16A
R14A
L16
R14
L15
R13
R12
L14
L13
L12
L11
L10A
L10
L9B
L9A
L9
R11
R10
R9
R8E
R8D
R8C
R8B
R8A
R8
L8
L7
R7
L6A
R6A
Stel die Padreserwe voor van gedeelte van Nasionale Roete 2/23
Represents the Road Reserve of a portion of National Route 2/23
SHEET 3A OF 8
P 230 / 74

PADRESERWE KOÖRDINATE / ROAD RESERVE CO-ORDINATES

LINKERKANT/LEFT HAND SIDE			REGTERKANT/RIGHT HAND SIDE		
Lo 31°	Y	X		Y	X
L1	+29 096,92	+3 359 494,02	R1	+28 983,74	+3 359 547,14
L2	+28 965,50	+3 359 245,00	R2	+28 765,00	+3 359 217,50
L2A	+27 974,54	+3 358 105,66	R3	+28 482,10	+3 358 888,89
L3	+27 646,50	+3 357 728,50	R3A	+27 836,94	+3 358 164,55
L4	+27 362,30	+3 357 391,50	R4	+27 172,00	+3 357 418,00
L5	+27 183,78	+3 357 223,94	R5	+26 874,89	+3 356 948,63
L6	+26 900,80	+3 356 794,33	R6	+26 742,50	+3 356 627,00
L6A	+26 768,23	+3 356 005,39	R6A	+26 671,90	+3 356 132,01
L7	+26 716,50	+3 355 697,50	R7	+26 610,00	+3 355 698,00
L8	+26 667,79	+3 355 361,54	R8	+26 564,00	+3 355 294,50
L9	+26 650,70	+3 355 044,75	R8A	+26 549,00	+3 355 208,50
L9A	+26 652,00	+3 355 008,50	R8B	+26 526,50	+3 355 143,00
L9B	+26 650,00	+3 354 903,00	R8C	+26 502,00	+3 355 090,00
L10	+26 747,50	+3 354 852,50	R8D	+26 459,00	+3 355 009,50
L10A	+26 808,55	+3 354 825,33	R8E	+26 435,50	+3 354 991,50
L11	+26 797,80	+3 354 791,50	R9	+26 285,25	+3 355 055,50
L12	+26 745,00	+3 354 804,00	R10	+26 268,00	+3 355 019,50
L13	+26 649,50	+3 354 849,00	R11	+26 432,00	+3 354 944,00
L14	+26 627,00	+3 354 843,50	R12	+26 409,50	+3 354 803,00
L15	+26 513,98	+3 354 694,65	R13	+26 385,43	+3 354 727,20
L16	+26 249,22	+3 354 277,06	R14	+26 153,45	+3 354 304,06
L16A	+25 865,22	+3 353 632,82	R14A	+25 777,74	+3 353 666,53
L17	+25 609,48	+3 353 203,75	R15	+25 674,00	+3 353 490,50
L18	+25 248,42	+3 352 593,06	R16	+25 529,76	+3 353 232,82
L19	+25 207,00	+3 352 481,50	R17	+25 063,09	+3 352 434,56
L20	+25 136,69	+3 352 401,95	R18	+24 905,04	+3 352 212,24
L21	+24 999,72	+3 352 167,66	R19	+24 852,51	+3 352 111,48
L22	+24 953,20	+3 352 075,30	R20	+24 753,60	+3 351 940,72
L23	+24 851,59	+3 351 903,50	R21	+24 579,00	+3 351 596,30
L24	+24 789,32	+3 351 793,94	R22	+24 417,70	+3 350 992,50
L25	+24 674,50	+3 351 515,00	R23	+24 361,30	+3 350 870,00
L26	+24 550,30	+3 351 068,00	R24	+24 363,50	+3 350 676,00
L27	+24 572,50	+3 350 930,50	R25	+24 259,50	+3 350 406,00
L28	+24 396,95	+3 350 407,99	R26	+23 942,50	+3 350 015,00
L29	+23 995,00	+3 349 884,00	R27	+23 716,00	+3 349 787,00
L30	+23 768,00	+3 349 666,50	R27A	+23 196,83	+3 349 178,34
L30A	+23 316,03	+3 349 131,97	R28	+22 985,00	+3 348 930,00
L31	+22 977,00	+3 348 731,00	R29	+22 683,50	+3 348 395,00
L32	+22 816,00	+3 348 379,00			

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No. 2296 3 Desember 1993

VERKLARING VAN NASIONALE PAD TUSSEN MIDDELFONTEIN EN GROOTVALEY: PROVINSIE TRANSVAAL: NASIONALE ROETE 1, SEKSIES 24 EN 25

Kragtens die bevoegdheid my verleen by artikel 4 (1) (a) van die Wet op Nasionale Paaie, 1971 (Wet No. 54 van 1971), soos gewysig, verklaar ek hierby, op aanbeveling van die Suid-Afrikaanse Padraad, dat die opgemete gedeelte van Nasionale Roete 1, Seksies 24 en 25 vanaf Middelfontein tot by Grootvaley, soos aangedui op Plan P395/93, velle 1 tot 7, 'n nasionale pad is.

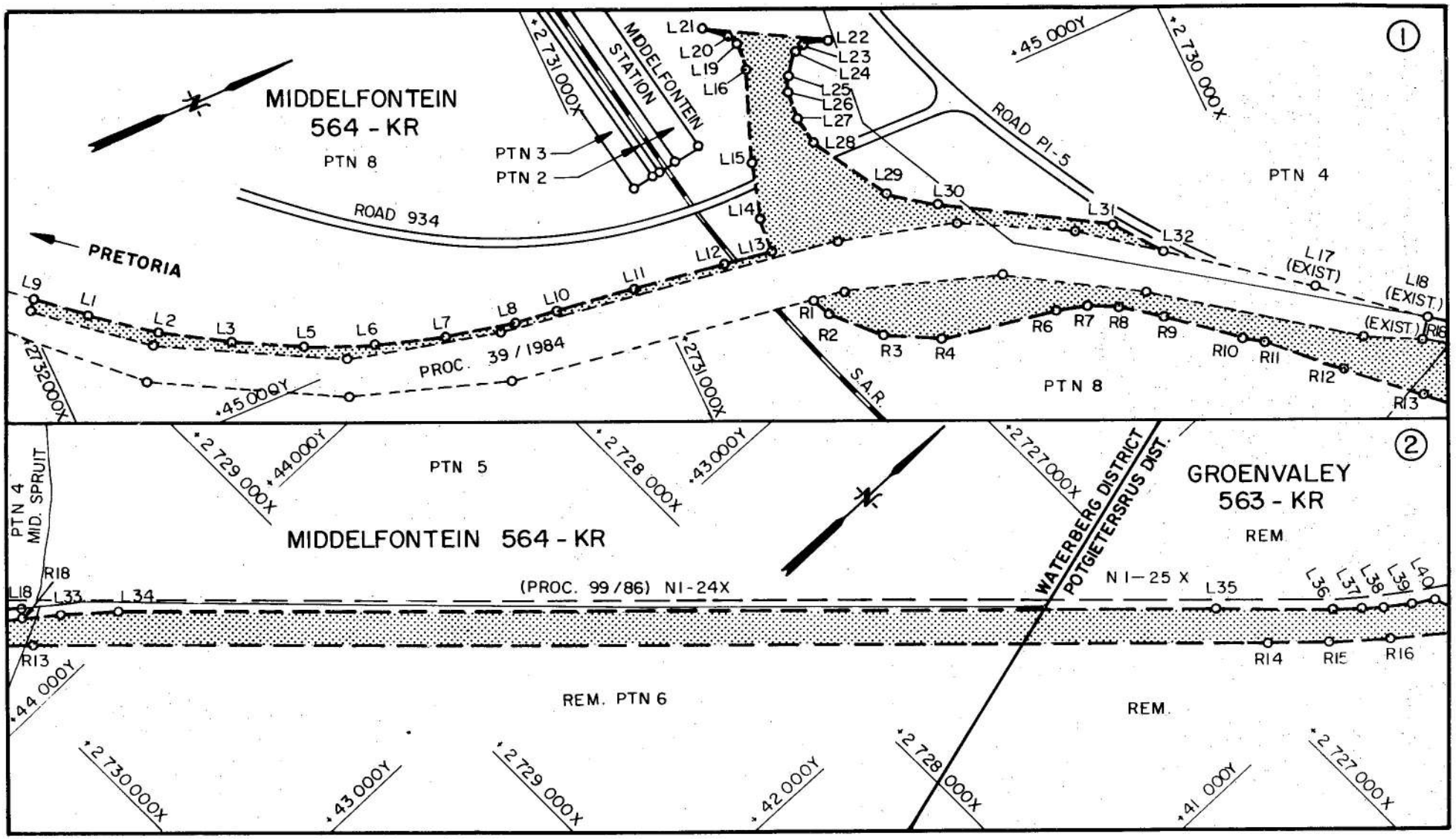
P. J. WELGEMOED,
Minister van Vervoer.

No. 2296 3 December 1993

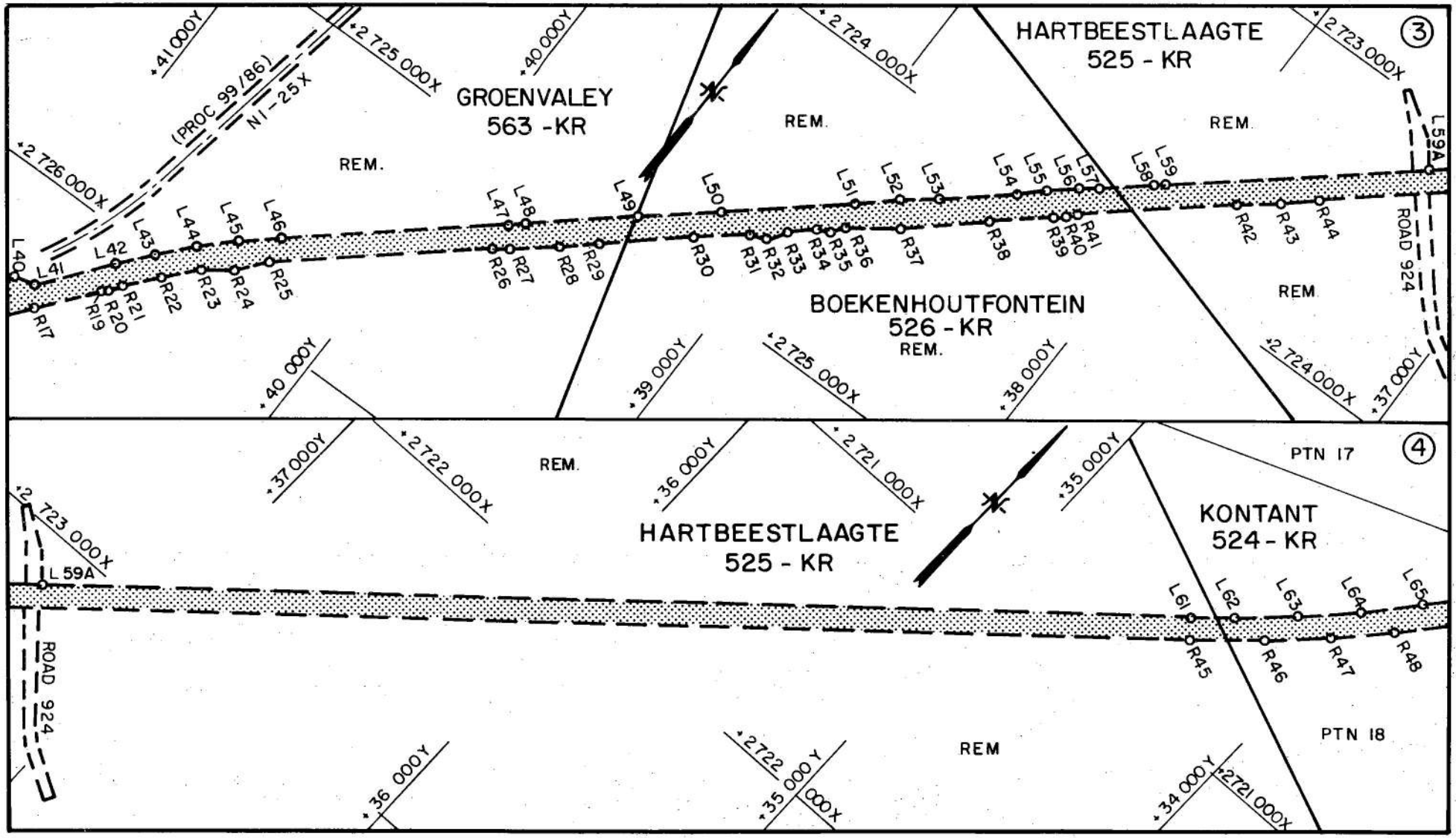
DECLARATION OF NATIONAL ROAD MIDDELFONTEIN AND GROOTVALEY: PROVINCE OF THE TRANSVAAL: NATIONAL ROUTE 1, SECTIONS 24 AND 25

By virtue of the powers vested in my by section 4 (1) (a) of the National Roads Act, 1971 (Act No. 54 of 1971), as amended, I hereby, on the recommendation of the South African Roads Board, declare that the surveyed section of National Route 1, Sections 24 and 25, from Middelfontein to Grootvaley, as indicated on the attached Plan P395/93, sheets 1 to 7, shall be a national road.

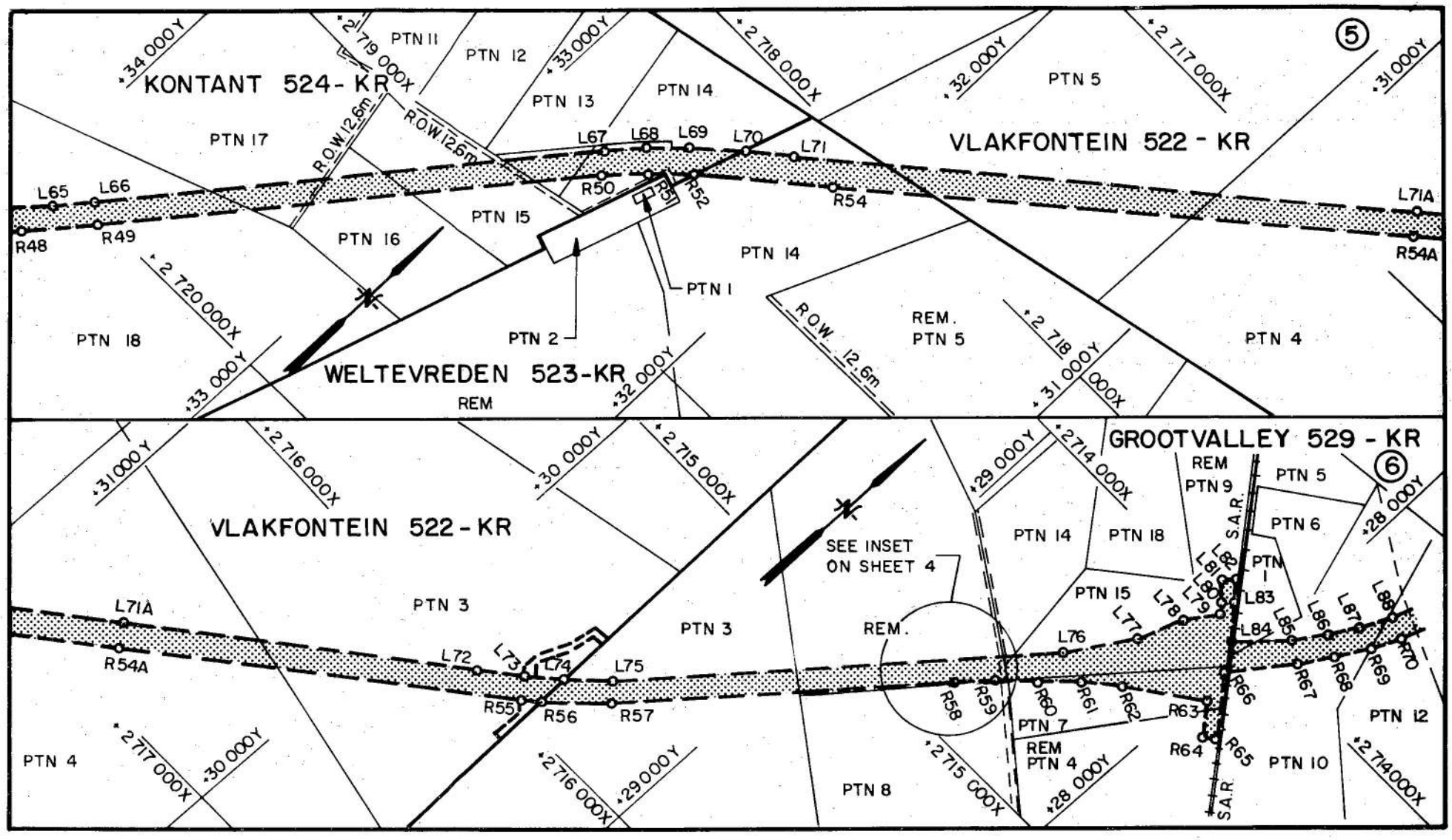
P. J. WELGEMOED,
Minister of Transport.



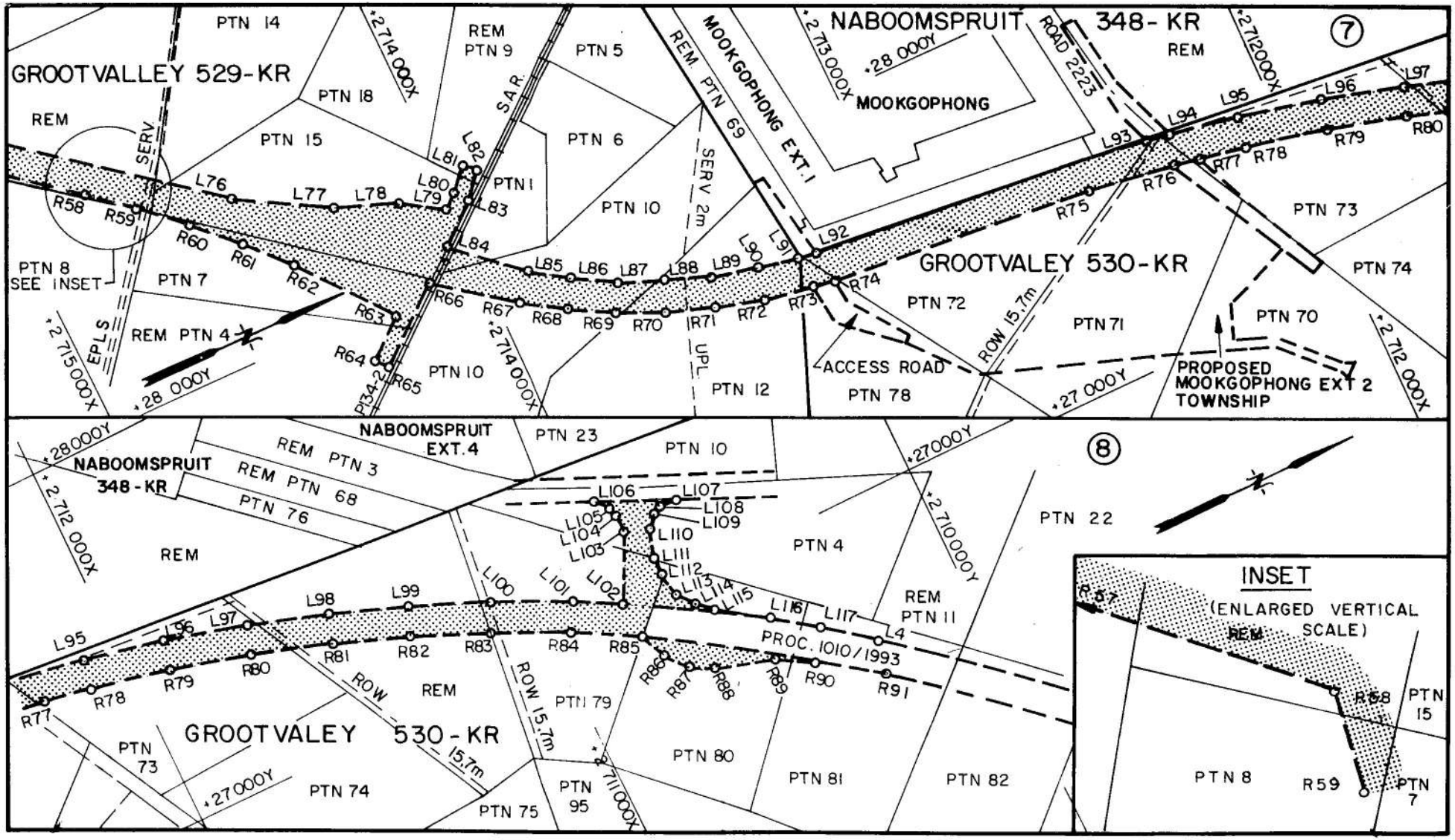
Suid - Afrikaanse Padraad South African Roads Board	Die figuur getoon The figure shown	stel die padreserwe voor van 'n gedeelte represents the road reserve of a portion van Nasionale Roete 1 Seksie 24 & 25 of National Route 1 Section 24 & 25	Vel 1 van 7 Sheet 1 of 7	P 395/93



Suid - Afrikaanse Padraad South African Roads Board	Die figuur getoon The figure shown		stel die padreserwe voor van 'n gedeelte represents the road reserve of a portion van Nasionale Roete 1 Seksie 25 of National Route Section	Vel 2 van 7 Sheet of P 395/93



Suid - Afrikaanse Padraad South African Roads Board	Die figuur getoon The figure shown		stel die padreserwe voor van 'n gedeelte represents the road reserve of a portion van Nasionale Roete 1 Seksie 25 of National Route 1 Section 25	Vel 3 van 7 Sheet 3 of 7 P 395 / 93



Suid - Afrikaanse Padraad South African Roads Board	Die figuur getoon The figure shown	stel die padreserwe voor van 'n gedeelte represents the road reserve of a portion van Nasionale Roete 1 Seksie 25 of National Route 1 Section 25	Vel 4 van 7 Sheet 4 of 7	P 395 / 93

PADRESERWE KOÖRDINATE / ROAD RESERVE CO-ORDINATES

LINKERKANT/LEFT HAND SIDE			Lo 29°	REGTERKANT/RIGHT HAND SIDE		
	Y	X			Y	X
L9	+45 331,73	+2 731 976,64		R1	+44 768,45	+2 730 753,52
L1	+45 265,92	+2 731 903,23		R2	+44 733,66	+2 730 737,80
L2	+45 189,06	+2 731 805,23		R3	+44 661,36	+2 730 669,47
L3	+45 120,91	+2 731 701,00		R4	+44 616,10	+2 730 581,57
L5	+45 061,98	+2 731 591,29		R6	+44 577,20	+2 730 383,01
L6	+45 012,68	+2 731 476,92		R7	+44 563,40	+2 730 330,33
L7	+44 973,39	+2 731 358,75		R8	+44 539,83	+2 730 281,15
L8	+44 944,39	+2 731 237,64		R9	+44 491,64	+2 730 217,61
L10	+44 934,46	+2 731 163,46		R10	+44 401,37	+2 730 110,50
L11	+44 914,36	+2 731 024,89		R11	+44 378,34	+2 730 077,49
L12	+44 890,09	+2 730 866,74		R12	+44 274,06	+2 729 975,51
L13	+44 875,17	+2 730 780,35		R13	+44 176,00	+2 729 867,56
L14	+44 937,09	+2 730 774,86		R14	+41 269,16	+2 726 850,89
L15	+45 032,07	+2 730 747,21		R15	+41 126,88	+2 726 701,93
L16	+45 186,15	+2 730 685,93		R16	+40 990,84	+2 726 546,97
L19	+45 232,83	+2 730 682,49		R17	+40 861,94	+2 726 387,15
L20	+45 251,84	+2 730 690,75		R19	+40 713,74	+2 726 201,65
L21	+45 281,20	+2 730 727,20		R20	+40 697,35	+2 726 189,14
L22	+45 172,45	+2 730 535,72		R21	+40 670,31	+2 726 148,33
L23	+45 185,45	+2 730 578,89		R22	+40 583,85	+2 726 041,84
L24	+45 180,01	+2 730 596,86		R23	+40 491,31	+2 725 940,52
L25	+45 145,28	+2 730 625,60		R24	+40 400,55	+2 725 874,95
L26	+45 119,59	+2 730 638,59		R25	+40 324,12	+2 725 783,60
L27	+45 069,46	+2 730 642,43		R26	+39 747,60	+2 725 291,56
L28	+45 019,74	+2 730 635,58		R27	+39 699,17	+2 725 253,06
L29	+44 886,35	+2 730 560,24		R28	+39 561,96	+2 725 145,19
L30	+44 833,01	+2 730 488,36		R29	+39 467,19	+2 725 057,49
L31	+44 675,34	+2 730 229,70		R30	+39 228,97	+2 724 848,93
L32	+44 595,57	+2 730 170,86		R31	+39 084,94	+2 724 726,00
R18	+44 266,57	+2 729 829,01		R32	+39 026,87	+2 724 702,36
L33	+44 183,69	+2 729 718,83		R33	+38 984,38	+2 724 640,19
L34	+44 058,19	+2 729 573,71		R34	+38 914,77	+2 724 580,77
L35	+41 480,00	+2 726 901,87		R35	+38 872,94	+2 724 558,31
L36	+41 202,47	+2 726 614,01		R36	+38 835,99	+2 724 513,54
L37	+41 137,47	+2 726 544,78		R37	+38 689,38	+2 724 402,88
L38	+41 090,48	+2 726 488,56		R38	+38 463,46	+2 724 204,81
L39	+41 030,68	+2 726 409,44		R39	+38 294,49	+2 724 064,55
L40	+40 984,59	+2 726 339,53		R40	+38 265,96	+2 724 036,36
L41	+40 913,56	+2 726 323,60		R41	+38 239,86	+2 724 004,77
L42	+40 734,47	+2 726 097,07		R42	+37 829,11	+2 723 654,22
L43	+40 644,95	+2 725 988,08		R43	+37 718,06	+2 723 568,64
L44	+40 549,26	+2 725 884,42		R44	+37 620,43	+2 723 476,12
L45	+40 448,75	+2 725 785,42		R45	+34 376,06	+2 720 707,22
L46	+40 342,80	+2 725 693,50		R46	+34 182,50	+2 720 538,25
L47	+39 752,50	+2 725 190,56		R47	+34 028,43	+2 720 386,44
L48	+39 710,11	+2 725 147,80		R48	+33 882,73	+2 720 226,56
L49	+39 421,06	+2 724 901,12				
L50	+39 204,84	+2 724 723,16				
L51	+38 861,47	+2 724 430,10				

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PADRESERWE KOÖRDINATE / ROAD RESERVE CO-ORDINATES

LINKERKANT/LEFT HAND SIDE

	Y	X Lo 29°
L52	+38 752,41	+2 724 329,02
L53	+38 644,47	+2 724 244,92
L54	+38 444,47	+2 724 074,23
L55	+38 372,68	+2 724 003,75
L56	+38 289,00	+2 723 932,34
L57	+38 233,16	+2 723 889,94
L58	+38 096,25	+2 723 773,09
L59	+38 064,15	+2 723 749,64
L59A	+37 391,44	+2 723 175,51
L61	+34 427,96	+2 720 646,34
L62	+34 316,50	+2 720 550,05
L63	+34 161,61	+2 720 405,40
L64	+34 014,71	+2 720 252,62
L65	+33 876,20	+2 720 092,20
L66	+33 783,21	+2 719 975,53
L67	+32 706,88	+2 718 607,65
L68	+32 616,22	+2 718 497,92
L69	+32 518,04	+2 718 394,60
L70	+32 374,88	+2 718 263,78
L71	+32 241,34	+2 718 159,61
L71A	+30 624,02	+2 716 788,97
L72	+29 703,87	+2 716 009,15
L73	+29 578,86	+2 715 900,47
L74	+29 479,57	+2 715 804,48
L75	+29 367,97	+2 715 684,66
L76	+28 409,82	+2 714 501,37
L77	+28 276,66	+2 714 279,69
L78	+28 219,75	+2 714 123,90
L79	+28 145,23	+2 714 019,70
L80	+28 179,50	+2 713 981,90
L81	+28 237,50	+2 713 927,20
L82	+28 211,40	+2 713 896,70
L83	+28 149,46	+2 713 952,47
L84	+28 063,91	+2 714 057,72
L85	+27 920,68	+2 713 893,41
L86	+27 855,94	+2 713 796,89
L87	+27 799,08	+2 713 695,52
L88	+27 750,46	+2 713 589,96
L89	+27 710,40	+2 713 480,86
L90	+27 679,14	+2 713 368,92
L91	+27 658,14	+2 713 264,77
L92	+27 650,42	+2 713 215,87
L93	+27 558,56	+2 712 326,45
L94	+27 544,62	+2 712 267,94
L95	+27 513,60	+2 712 094,84
L96	+27 465,84	+2 711 888,31
L97	+27 405,81	+2 711 684,43
L98	+27 339,58	+2 711 483,73
L99	+27 265,49	+2 711 295,93

REGTERKANT/RIGHT HAND SIDE

	Y	X
R49	+33 720,33	+2 720 024,97
R50	+32 657,32	+2 718 674,03
R51	+32 551,42	+2 718 557,46
R52	+32 426,79	+2 718 436,45
R54	+32 082,81	+2 718 130,13
R54A	+30 572,30	+2 716 850,00
R55	+29 525,07	+2 715 962,49
R56	+29 477,29	+2 715 919,38
R57	+29 307,23	+2 715 736,78
R58	+28 580,33	+2 714 839,07
R59	+28 493,18	+2 714 735,73
R60	+28 397,66	+2 714 629,43
R61	+28 295,67	+2 714 529,31
R62	+28 187,54	+2 714 435,73
R63	+27 959,06	+2 714 250,17
R64	+27 877,06	+2 714 351,07
R65	+27 846,02	+2 714 325,85
R66	+27 996,90	+2 714 140,18
R67	+27 851,47	+2 713 943,94
R68	+27 786,51	+2 713 839,56
R69	+27 726,47	+2 713 732,53
R70	+27 675,14	+2 713 621,08
R71	+27 632,84	+2 713 505,89
R72	+27 599,83	+2 713 387,70
R73	+27 575,70	+2 713 259,15
R74	+27 569,29	+2 713 201,95
R75	+27 498,08	+2 712 519,49
R76	+27 468,46	+2 712 288,50
R77	+27 457,28	+2 712 226,37
R78	+27 435,24	+2 712 110,91
R79	+27 388,37	+2 711 908,28
R80	+27 331,44	+2 711 708,24
R81	+27 264,49	+2 711 511,34
R82	+27 191,81	+2 711 327,08
R83	+27 106,09	+2 711 137,59
R84	+27 010,99	+2 710 952,63
R85	+26 920,44	+2 710 796,26
R86	+26 853,16	+2 710 764,14
R87	+26 799,80	+2 710 721,77
R88	+26 761,73	+2 710 665,84
R89	+26 716,93	+2 710 517,92
R90	+26 665,77	+2 710 430,14
R91	+26 551,92	+2 710 280,06

PADRESERWE KOÖRDINATE / ROAD RESERVE CO-ORDINATES

LINKERKANT/LEFT HAND SIDE			REGTERKANT/RIGHT HAND SIDE		
	Y	X Lo 29°		Y	X
L100	+27 178,13	+2 711 102,80			
L101	+27 081,20	+2 710 914,28			
L102	+27 014,96	+2 710 799,90			
L103	+27 186,68	+2 710 723,00			
L104	+27 234,42	+2 710 718,01			
L105	+27 254,91	+2 710 725,32			
L106	+27 289,85	+2 710 758,97			
L107	+27 197,43	+2 710 562,44			
L108	+27 201,43	+2 710 609,76			
L109	+27 193,59	+2 710 629,50			
L110	+27 158,21	+2 710 661,18			
L111	+27 087,72	+2 710 681,58			
L112	+27 043,83	+2 710 680,96			
L113	+26 980,66	+2 710 669,97			
L114	+26 934,53	+2 710 641,10			
L115	+26 897,53	+2 710 598,56			
L116	+26 815,18	+2 710 485,28			
L117	+26 735,69	+2 710 381,02			
L4	+26 641,93	+2 710 262,13			

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No. 2299 3 Desember 1993

WYSIGING VAN GOEWERMENTSKENNISGEWINGS Nos. 1705, 1706 EN 1711 GEDATEER 10 SEPTEMBER 1993

Die Suid-Afrikaanse Padraad maak hierby kragtens die bepaling van artikel 9 (4) (c) van die Wet op Nasionale Paaie, 1971 (Wet No. 54 van 1971), soos gewysig, bekend dat die Minister van Vervoer en Pos- en Telekommunikasiewese kragtens artikel 9 (4) (a) en 9 (4) (d) daarvan, Goewermentskennisgewings Nos. 1705, 1706 en 1711 gedateer 10 September 1993, gewysig het soos in die Bylae hiervan uiteengesit.

C. F. SCHEEPERS,
Voorsitter: Suid-Afrikaanse Padraad.

BYLAE

(1) Goewermentskennisgewing No. 1705 gedateer 10 September 1993 word hierby gewysig—

(a) deur die invoeging van subparagraaf 4.5:

"4.5 'n Korting soos uiteengesit hieronder, op die tolbedrae soos hierbo vermeld, word toegestaan aan Klas 1: Ligte voertuie, wat deur die Mooi-tolplaza beweeg wat 'n geldige 'Gereelde Gebruikerskaart', goedgekeur deur die Departement van Vervoer, vir betaling aanbied:

(a) Vir 20 tot 40 ritte per enkel Gereelde Gebruikerskaart gedurende 'n kalendermaand—20% afslag.

(b) Vir meer as 40 ritte per enkel Gereelde Gebruikerskaart gedurende 'n kalendermaand—40% afslag.

Die gebruik en korting tree in werking wanneer toerusting vir die prosessering van sodanige betaalmetode in werking gestel word.".

(2) Goewermentskennisgewing No. 1706 gedateer 10 September 1993 word hierby gewysig—

(a) deur subparagraaf (b) van paragraaf 3.2 te skrap met ingang vanaf die datum waarop die gewysigde paragraaf 3.3 (b) en 3.4 geïmplementeer word.

(b) deur subparagraaf (b) van paragraaf 3.3 deur die volgende subparagraaf te vervang:

"(b) gebruikers van die Oribi-tolplaza, wanneer meer as 20 ritte met 'n enkel Vlootbestuurs- of Transkard-kaart, goedgekeur deur die Departement van Vervoer, per kalendermaand onderneem word."; en

(c) deur paragraaf 3.4 deur die volgende paragraaf te vervang:

"3.4 'n Korting soos uitgesit hieronder, op die tolbedrae soos hierbo vermeld, word toegestaan aan Klas 1: Ligte voertuie, wat deur die Oribi-tolplaza beweeg wat 'n geldige 'Gereelde Gebruikerskaart', goedgekeur deur die Departement van Vervoer vir betaling aanbied.

(a) Vir 20 tot 40 ritte per enkel gebruikerskaart gedurende 'n kalendermaand—20% afslag.

No. 2299 3 December 1993

AMENDMENT OF GOVERNMENT NOTICES Nos. 1705, 1706 AND 1711 DATED 10 SEPTEMBER 1993

The South African Roads Board hereby, in terms of section 9 (4) (c) of the National Roads Act, 1971 (Act No. 54 of 1971), as amended, makes known that the Minister of Transport and of Posts and Telecommunications has in terms of section 9 (4) (a) and 9 (4) (d) thereof, amended Government Notices Nos. 1705, 1706 and 1711 dated 10 September 1993 as set out in the Schedule hereto.

C. F. SCHEEPERS,
Chairman: South African Roads Board.

SCHEDULE

(1) Government Notice No. 1705 dated 10 September 1993 is hereby amended—

(a) by the addition of subparagraph 4.5:

"4.5 A rebate on the amounts of toll specified above shall be granted to Class 1: Light vehicles, passing through the Mooi Toll Plaza who tender valid 'Frequent User Cards' approved by the Department of Transport as set out hereunder:

(a) For 20 to 40 trips per single Frequent User Card during a calendar month—20% discount.

(b) For more than 40 trips per single Frequent User Card during a calendar month—40% discount.

This usage and rebate will come into force when equipment for the processing of such payment method becomes operational.".

(2) Government Notice No. 1706 dated 10 September 1993 is hereby amended—

(a) by the deletion of subparagraph (b) of paragraph 3.2 with effect from the date at which the amended paragraphs 3.3 (b) and 3.4 comes into effect;

(b) by the substitution for subparagraph (b) of paragraph 3.3 of the following subparagraph:

"(b) all users of the Oribi Toll Plaza when more than 20 trips are undertaken during a calendar month with a single Fleet Management or a Transkard card which has been approved by the Department of Transport."; and

(c) by the substitution for paragraph 3.4 of the following paragraph:

"3.4 A rebate on the amount of toll specified above shall be granted to Class 1: Light vehicles, passing through the Oribi Toll Plaza who tender valid 'Frequent User Cards' approved by the Department of Transport as set out hereunder:

(a) For 20 to 40 trips per single card during a calendar month—20% discount.

(b) Vir meer as 40 ritte per enkel gebruikerskaart gedurende 'n kalendermaand—40% afslag.

Die konsessietariewe in paragrawe 3.3 (b) en 3.4 bedoel tree in werking sodra die toerusting vir die prossesering van sodanige betaalmetodes in werking gestel word.".

(3) Goewermentskennisgewing No. 1711 gedateer 10 September 1993 word hierby gewysig—

(a) deur die invoeging van subparagraaf 4.3:

"4.3 'n Korting soos uiteengesit hieronder, op die tolbedrae soos hierbo vermeld, word toegestaan aan Klas 1: Ligte voertuie, wat deur die Grasmere Tolplaza beweeg wat 'n geldige 'Gereelde Gebruikerskaart', goedgekeur deur die Departement van Vervoer, vir betaling aanbied:

(a) Vir 20 tot 40 ritte per enkel gereelde gebruikerskaart gedurende 'n kalendermaand—20% afslag.

(b) Vir meer as 40 ritte per enkel gereelde gebruikerskaart gedurende 'n kalendermaand—40% afslag.

Die gebruik en korting tree in werking sodra die toerusting vir die prosessering van sodanige betaalmetodes in werking gestel word.".

(b) For more than 40 trips per single card during a calendar month—40% discount.

The consession tariffs in paragraphs 3.3 (b) and 3.4 will become effective when equipment for the processing of such payment method becomes operational.".

(3) Government Notice No. 1711 dated 10 September 1993 is hereby amended—

(a) by the addition of subparagraph 4.3:

"4.3 A rebate on the amounts of toll specified above shall be granted to Class 1: Light vehicles, passing through the Grasmere Toll Plaza who tender valid 'Frequent User Cards' approved by the Department of Transport as set out hereunder:

(a) For 20 to 40 trips per single Frequent User Card during a calendar month—20% discount.

(b) For more than 40 trips per single Frequent User Card during a calendar month—40% discount.

This usage and rebate will come into force when equipment for the processing of such payment method becomes operational.".

DEPARTEMENT VAN WATERWESE EN BOSBOU

No. 2283 3 Desember 1993

MARICORIVIER EN ALLE SYTAKKE DAARVAN STROOMAF VAN DIE PLAAS WONDERFONTEIN 258 JP, DISTRIK MARICO, TRANSVAAL: WYSIGING VAN DIE PERKE NEERGELÊ IN ARTIKEL 9B (1) (a) VAN DIE WATERWET, 1956 (WET No. 54 VAN 1956), MET BETREKKING TOT DIE OPDAM-, OPGAAR-, UITNEEM- OF UITKEERVERMOË VAN WATERWERKE

Ek, Jacob Albertus van Wyk, Minister van Waterwese, handelende kragtens die bevoegdheid aan my verleen by artikel 9B (1C) (a) van die Waterwet, 1956 (Wet No. 54 van 1956), wysig hierby, met ingang van die datum van publikasie hiervan, die perke in artikel 9B (1) (a) van genoemde Wet neergelê met betrekking tot die opdam-, opgaar-, uitneem- of uitkeervermoë van waterwerke ten opsigte van die Maricorivier en alle sytakke daarvan stroomaf van die plaas Wonderfontein 258 JP, soos aangedui op die bygaande kaart, uitgesonderd daardie gedeeltes van die rivier en sy sytakke wat binne die Marico-Bosveld- en die Klein-Maricopoort-staatswaterbeheergebied geleë is, deur die uitdrukkings "250 000 kubieke meter" en "110 liter per sekonde" waar dit in genoemde artikel voorkom, deur die uitdrukkings "3 000 kubieke meter" en "10 liter per sekonde" te vervang.

DEPARTMENT OF WATER AFFAIRS AND FORESTRY

No. 2283 3 December 1993

MARICO RIVER AND ALL TRIBUTARIES THEREOF DOWNSTREAM OF THE FARM WONDERFONTEIN 258 JP, DISTRICT OF MARICO, TRANSVAAL: AMENDMENT OF THE LIMITS LAID DOWN IN SECTION 9B (1) (a) OF THE WATER ACT, 1956 (ACT No. 54 OF 1956), IN REGARD TO THE IMPOUNDMENT, STORAGE, ABSTRACTION OR DIVERSION CAPACITY OF WATER WORKS

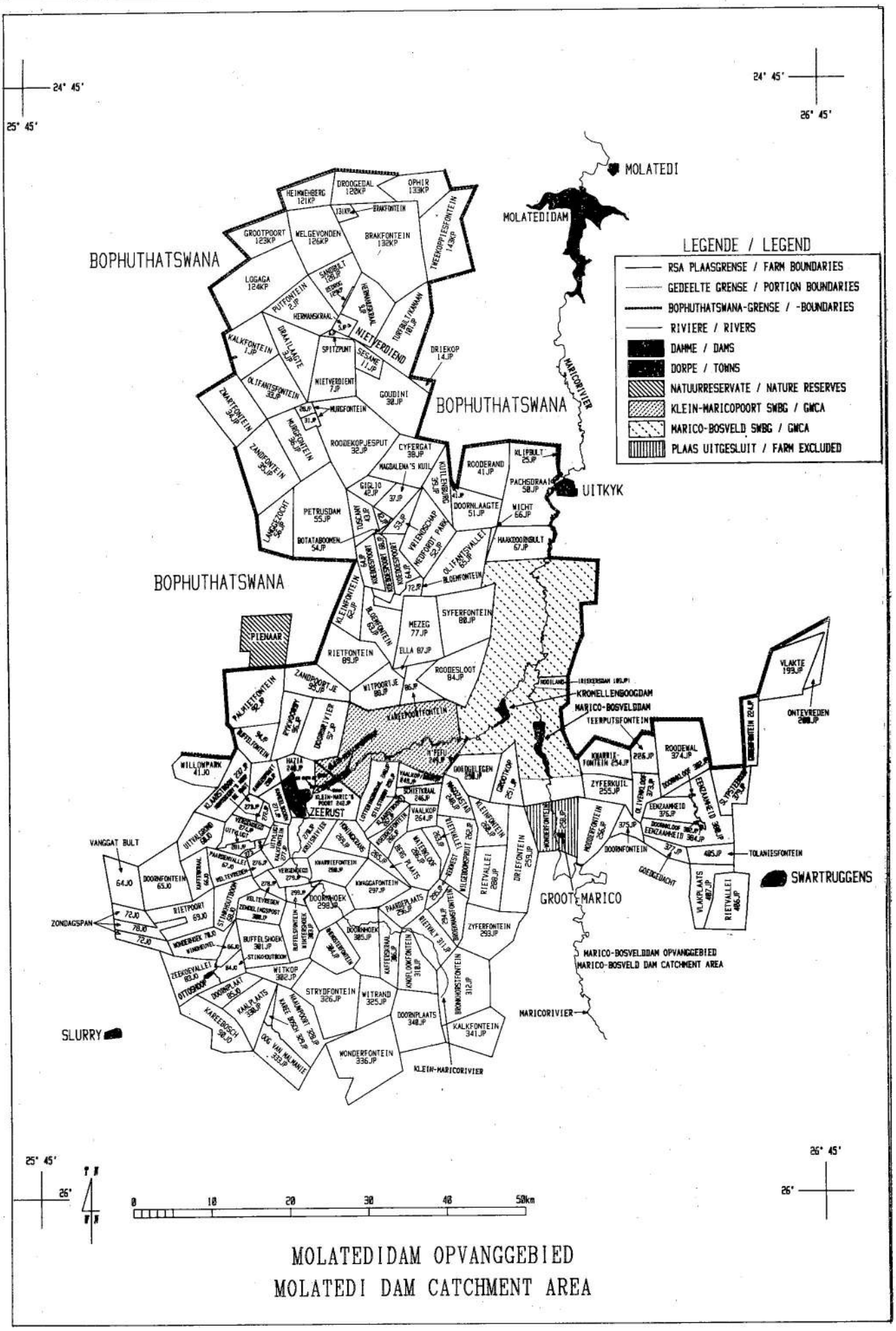
I, Jacob Albertus van Wyk, Minister of Water Affairs, under the powers vested in me by section 9B (1C) (a) of the Water Act, 1956 (Act No. 54 of 1956), hereby, with effect from the date of publication hereof, amend the limits laid down in section 9B (1) (a) of the said Act as far as the impoundment, storage, abstraction or diversion capacity of water works is concerned, in respect of the Marico River and all tributaries thereof, downstream of the farm Wonderfontein 258 JP, as indicated on the accompanying map, excluding those portions of this River and its tributaries situated within the Marico-Bosveld and the Klein Maricopoort Government Water Control Areas, by substituting the expressions "3 000 cubic metres" and "10 litres per second" for the expressions "250 000 cubic metres" and "110 litres per second" where it appears in the said section.

Die uitwerking hiervan is dat geen waterwerk waarin meer as 3 000 (drieduisend) kubieke meter openbare water opgedam of opgegaar of waarmee meer as 10 (tien) liter openbare water per sekonde onttrek, uitgeneem of uitgekeer kan word op 'n eiendom bedoel in genoemde artikel 9B (1) (a) vir sover dit die bedoelde openbare strome betref, opgerig, verander of vergroot mag word nie, behalwe op gesag van 'n permit deur my uitgereik.

J. A. VAN WYK,
Minister van Waterwese.

The effect of this is that no water work in which more than 3 000 (three thousand) cubic metres of public water can be impounded or stored or with which more than 10 (ten) litres of public water per second can be abstracted or diverted on a property contemplated in the said section 9B (1) (a), may be constructed, altered or enlarged in so far as it concerns the intended public streams, except on the authority of a permit issued by me.

J. A. VAN WYK,
Minister of Water Affairs.



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DEPARTEMENT VAN FINANSIES

No. 2300 **3 Desember 1993**

BELASTING OP TOEGEVOEGDE WAARDE

BTW-PRAKTYKNOTA: No. 12

Datum: 24 November 1993

DIE LEWERING VAN BRUINBROOD TEEN DIE NULKOERS

Hierdie praktyknota word uitgereik ten einde interpretasieprobleme met betrekking tot die bepalings van artikel 11 (1) (j) van die Wet op Belasting op Toegevoegde Waarde, 1991, saamgelees met die Tweede Bylae by die Wet, ingevolge waarvan bruinbrood aan BTW teen die nulkoers onderhewig is, uit die weg te ruim.

Bruinbrood, ingevolge die Tweede Bylae, is bruinbrood soos omskryf in Regulasie 1 van die Regulasies ingevolge Goewermentskennisgewing No. R. 577 gepubliseer in *Staatskoerant* No. 13074 van 15 Maart 1991. Hierdie Regulasie bepaal dat bruinbrood, behoudens toelaatbare afwykings, uit 'n deeg gemaak van bruinkoringmeel en water, met of sonder ander bestanddele wat deur gisfermentasie of andersins deursuur is en wat gebak is en enige vorm, grootte of fatsoen, moet bestaan.

Ten einde te verseker dat die lewering van bruinbrood vir die doeleindes van belasting eenvormig hanteer word, word die volgende riglyne verskaf:

Die lewering (anders as in die loop van die verskaffing van maaltye of verversings) van enige soort bruinbrood, sal aan belasting teen die nulkoers onderhewig wees, mits—

(i) dit bemark en verkoop word onder die beskrywing bruinbrood;

(ii) die meelinhoud van die deeg uit ten minste 50 persent bruinkoringmeel bestaan; en

(iii) die massa van die brood 100 gram oorskry.

Produkte soos "volkoringbruinbrood", "hoëveselbruinbrood", "hoëprotiënbruinbrood" en "gesondheidsbruinbrood", wat aan bogenoemde vereistes voldoen, sal gevolglik aan die nulkoers onderhewig wees.

Die lewering van enige gedeelte van 'n bruinbrood sal ook aan die nulkoers onderhewig wees, behalwe waar dit gelewer word in die loop van die verskaffing van 'n maaltyd of verversing (soos 'n toebroodjie).

Bruinbroodrolletjies en ander broodprodukte wat nie aan bogenoemde vereistes voldoen nie, sal aan BTW teen die standaardkoers onderhewig wees.

KOMMISSARIS VAN BINNELANDSE INKOMSTE,
PRETORIA.

DEPARTMENT OF FINANCE

No. 2300 **3 December 1993**

VALUE-ADDED TAX

VAT PRACTICE NOTE: No. 12

Date: 24 November 1993

THE ZERO-RATED SUPPLY OF BROWN BREAD

This practice note is being issued to avoid problems of interpretation in relation to the provisions of section 11 (1) (j) of the Value-Added Tax Act, 1991, read with the Second Schedule to the Act, in terms of which brown bread is subject to VAT at the zero rate.

Brown bread is, in terms of the Second Schedule, brown bread as defined in Regulation 1 of the Regulations in terms of Government Notice No. R. 577 published in *Government Gazette* No. 13074 of 15 March 1991. This Regulation provides that brown bread, subject to the allowable deviations, shall consist of a dough made from brown wheaten meal and water, with or without other ingredients that has been fermented by yeast or otherwise leavened and which has been baked in any form, size or shape.

To ensure that the tax treatment of the supply of brown bread is applied uniformly the following guidelines are furnished:

The supply (other than in the course of the provision of a meal or refreshment) of any type of brown bread shall be subject to tax at the zero rate, provided—

(i) it is marketed and sold under the description brown bread;

(ii) the meal content of the dough consists of at least 50 per cent brown bread meal; and

(iii) the mass of the loaf exceeds 100 grams.

Products such as "whole-wheat brown bread", "high fibre brown bread", "high protein brown bread" and "brown health bread" which satisfy the above requirements will accordingly be subject to the zero rate.

The supply of a portion of a brown bread is also subject to the zero rate, except where supplied in the course of the provision of a meal or refreshment (such as a sandwich).

Brown bread rolls and other bread products which do not comply with the above requirements are subject to VAT at the standard rate.

COMMISSIONER FOR INLAND REVENUE,
PRETORIA.

ALGEMENE KENNISGEWINGS

KENNISGEWING 1175 VAN 1993

DEPARTEMENT VAN VERVOER

"REPORT OF THE COMMITTEE OF INQUIRY INTO A NATIONAL MARITIME POLICY FOR THE RSA"

Die Departement van Vervoer maak hierby bekend dat die Minister van Vervoer en van Pos- en Telekommunikasiewese besluit het om die "Report of the Committee of Inquiry into a National Maritime Policy for the RSA" vir algemene inligting en kommentaar beskikbaar te stel.

GENERAL NOTICES

NOTICE 1175 OF 1993

DEPARTMENT OF TRANSPORT

REPORT OF THE COMMITTEE OF INQUIRY INTO A NATIONAL MARITIME POLICY FOR THE RSA

The Department of Transport hereby makes known that the Minister of Transport and of Posts and Telecommunications has decided to make the Report of the Committee of Inquiry into a National Maritime Policy for the RSA available for general information and comments.

Die Verslag is teen R40 per eksemplaar beskikbaar en verkrygbaar by:

Die Departement van Vervoer, Hoofdirektoraat Skeepvaart, Privaatsak X193, Pretoria, 0001

of

Kamer 1047, Forumgebou, hoek van Bosman- en Strubenstraat, Pretoria.

Navrae rakende die beskikbaarheid van die Verslag kan by mnr. L. J. Fouché by telefoonnommer (012) 290-2361 gedoen word. Die faksimileenommer is (012) 290-2914.

Alle belanghebbende instansies en persone word uitgenooi om voor 15 Februarie 1994 skriftelike kommentaar oor die Verslag te lewer/voorstelle by bogenoemde adres in te dien.

(3 Desember 1993)

KENNISGEWING 1176 VAN 1993

SUID-AFRIKAANSE REGSKOMMISSIE

Die Suid-Afrikaanse Regskommissie stel op versoek van die Minister van Justisie ondersoek in na die vereenvoudiging van die strafprosedure. Dit is 'n omvattende ondersoek wat in fases deur die Kommissie aangepak is. Die Kommissie het reeds 'n werkstuk oor appèlle beskikbaar gestel en 'n konsepverslag sal eersdaags deur die Kommissie oorweeg word. Die Kommissie stel hiermee die tweede werkstuk in sy ondersoek, getiteld **"Vereenvoudiging van die Strafprosedure"**, vry. Die werkstuk handel oor onder andere die redes vir vertragings in die afhandeling van verhore, misbruik wat van die proses gemaak word, besondere bepalings van die Strafproseswet, 1977, wat vertragings in die hand werk en vertragings wat deur die administrasie van die proses veroorsaak word.

Die Kommissie nooi alle belanghebbende persone en instansies uit om kommentaar te lewer op die werkstuk of om voorstelle te doen vir die ontwikkeling, verbetering, modernisering of hervorming van hierdie faset van die reg.

Dit sal waardeer word indien skriftelike kommentaar of voorstelle die Kommissie teen 28 Februarie 1994 by onderstaande adres kan bereik.

Die werkstuk is op aanvraag gratis van die Kommissie verkrygbaar. Die Kommissie se kantore is op die Agste Verdieping, NG Kerk Sinodale Sentrum, Visagiestraat 228, Pretoria. Korrespondensie moet asseblief gerig word aan:

Die Sekretaris
Suid-Afrikaanse Regskommissie
Privaat Sak X668
PRETORIA
0001.

Telefoon: (012) 322-6440 (mev. P. Kotze).

Datum: November 1993.

(3 Desember 1993)

The Report is available at R40 per copy and obtainable from:

The Department of Transport, Chief Directorate Shipping, Private Bag X193, Pretoria, 0001

or

Room 1047, Forum Building, corner of Bosman and Struben Streets, Pretoria.

Enquiries in connection with the availability of the Report can be obtained from Mr L. J. Fouché at telephone number (012) 290-2361. The facsimile number is (012) 290-2914.

All interested parties and persons are invited to submit, in writing, their comments on/make proposals regarding the Report to the above address before 15 February 1994.

(3 December 1993)

NOTICE 1176 OF 1993

SOUTH AFRICAN LAW COMMISSION

The South African Law Commission is investigating the simplification of criminal procedure at the request of the Minister of Justice. This is a comprehensive investigation which the Commission has undertaken in phases. The Commission has already published a working paper on appeals, and a draft report will be considered by the Commission in the near future. The Commission hereby releases the second working paper titled **"Simplification of Criminal Procedure"**. This paper deals with, *inter alia,* the reasons for delays in the finalisation of trials, abuses of the process, particular provisions of the Criminal Procedure Act, 1977, which cause delays and delays resulting from the administration of the process.

The Commission invites all interested persons and bodies to comment on the working paper or to make suggestions for the development, improvement, modernisation or reform of this aspect of the law.

It would be appreciated if written comments or suggestions could reach the Commission by 28 February 1994 at the adress given below.

The working paper is obtainable free of charge from the Commission. The Commission's offices are on the Eighth Floor, NG Kerk Synodal Centre, 228 Visagie Street, Pretoria. Correspondence should be addressed to:

The Secretary
South African Law Commission
Private Bag X668
PRETORIA
0001.

Telephone: (012) 322-6440 (Mrs P. Kotze).

Date: November 1993.

(3 December 1993)

KENNISGEWING 1179 VAN 1993

DEPARTEMENT VAN NASIONALE GESONDHEID EN BEVOLKINGSONTWIKKELING

WET OP BEHEER VAN MEDISYNE EN VERWANTE STOWWE, 1965 (WET No. 101 VAN 1965)

REGISTRASIE VAN MEDISYNE

Hierby word ingevolge artikel 17 van die Wet op Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965), bekendgemaak dat die Registrateur van Medisyne met die goedkeuring van die Medisynebeheerraad ingestel by artikel 2 van genoemde Wet, die volgende medisyne soos in die Bylae hiervan omskryf, geregistreer het.

NOTICE 1179 OF 1993

DEPARTMENT OF NATIONAL HEALTH AND POPULATION DEVELOPMENT

MEDICINES AND RELATED SUBSTANCES CONTROL ACT, 1965 (ACT No. 101 OF 1965)

REGISTRATION OF MEDICINES

It is hereby notified, in terms of section 17 of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965), that the Registrar of Medicines, with the approval of the Medicines Control Council established by section 2 of the said Act, has registered the following medicines described in the Schedule hereto.

Registrasienommer:
Registration Number: **27/17.1/0552.**

Naam van medisyne:
Name of medicine: **Sabax Pancuronium Bromide.**

Doseringsvorm: Inspuiting.
Dosage Form: Injection.

Aktiewe bestanddele: Pankuroniumbromied/
Active ingredients: Pancuronium Bromide . . . 4 mg per 2-mℓ-ampuul/ampoule.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslike vervaardigde produk moet gevalideer word.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.

Applikant:
Applicant: Adcock Ingram Critical Care Ltd.

Rakleeftyd: 24 maande.
Shelf-life: 24 months.

Datum van registrasie: 19 Julie 1993.
Date of registration: 19 July 1993.

Registrasienommer:
Registration Number: **E/11/1695.**

Naam van medisyne:
Name of medicine: **Fams Mist Mag Hydroxide.**

Doseringsvorm: Suspensie.
Dosage Form: Suspension.

Aktiewe bestanddele: Magnesiumoksied akwivalent aan Magnesiumhidroksied/
Active ingredients: Magnesium Oxide equivalent to Magnesium Hydroxide . . . 790 mg per 10-mℓ-suspensie/suspension.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte na registrasie moet gevalideer word, tensy hierdie dokumentasie beskikbaar is.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots after registration must be validated, unless this documentation is available.

Applikant:
Applicant: Genpharm Pharmaceuticals (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 27 Julie 1993.
27 July 1993.

Registrasienommer:
Registration Number: **27/6.2/0467.**

Naam van medisyne:
Name of medicine: **Adenocor.**

Doseringsvorm:
Dosage Form: Inspuiting.
Injection.

Aktiewe bestanddele:
Active ingredients: Adenosien/
Adenosine . . . 6 mg per 2-mℓ-flessie/vial.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of the product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.

Applikant:
Applicant: R & C Pharmaceuticals (Pty) Ltd.

Rakleeftyd:
Shelf-life: 36 maande.
36 months.

Datum van registrasie:
Date of registration: 1 September 1993.
1 September 1993.

Registrasienommer:
Registration Number: **Y/24/9.**

Naam van medisyne:
Name of medicine: **Solumag.**

Doseringsvorm: Bruistablet.
Dosage Form: Effervescent tablet.

Aktiewe bestanddele: Ligte Magnesiumoksied ekwivalent aan Magnesium/
Active ingredients: Light Magnesium Oxide equivalent to Magnesium . . . 150 mg per tablet.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die eerste twee produksielotte van die plaaslike vervaardigde produk moet gevalideer word.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The first two production lots of the locally manufactured product must be validated.

Applikant:
Applicant: Hexal Pharmaceuticals SA (Pty) Ltd.

Rakleeftyd: 24 maande.
Shelf-life: 24 months.

Datum van registrasie: 1 September 1993.
Date of registration: 1 September 1993.

Registrasienommer:
Registration Number: **27/20.1.2/0327.**

Naam van medisyne:
Name of medicine: **Cillagen S-125.**

Doseringsvorm: Poeier vir stroop.
Dosage Form: Powder for syrup.

Aktiewe bestanddele: Amoksisillientrihidraat ekwivalent aan Amoksisillien/
Active ingredients: Amoxycillin Trihydrate equivalent to Amoxycillin . . . 125 mg per 5-mℓ-stroop/syrup.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.
6. 'n Naregistrasie-inspeksie moet op die eerste produksielot van die plaaslike vervaardigde produk uitgevoer word.
7. Bemarking van die produk mag slegs 'n aanvang neem nadat 'n bevredigende na-registrasieinspeksieverslag gedien het.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.

6. A post-registration inspection must be conducted on the first production lot of the locally manufactured product.
7. Marketing of the product may only commence following a satisfactory post-registration inspection report.

Applikant:
Applicant: Lennon Limited.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 1 September 1993.
1 September 1993.

Registrasienommer:
Registration Number: **27/13.9.2/0163.**

Naam van medisyne:
Name of medicine: **Loceryl 5%.**

Doseringsvorm:
Dosage form: Naellak.
Nail lacquer.

Aktiewe bestanddele:
Active ingredients: Amorolfienhidrochloried ekwivalent aan Amorolfien.
Amorolfine Hydrochloride equivalent to Amorolfine . . . 50 mg per 1-mℓ-vloeistof/liquid.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslike vervaardige produk moet gevalideer word.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.

Applikant:
Applicant: Roche Products (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 1 September 1993.
1 September 1993.

Registrasienommer:
Registration Number: **28/13.1/0074.**

Naam van medisyne:
Name of medicine: **Zedchem CHG Obstetric Cream.**

Doseringsvorm:
Dosage form: Room.
Cream.

Aktiewe bestanddele:
Active ingredients: Chloorheksidienglukonaat/
Chlorhexidine Gluconate . . . 1 g per 100-mℓ-room/cream.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslike vervaardigde produk moet gevalideer word.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.

Applikant:
Applicant: Zedchem CC.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 6 September 1993.
6 September 1993.

Registrasienommer:
Registration Number: **28/20.2.6/0064.**

Naam van medisyne:
Name of medicine: **Daramal.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele:
Active ingredients: Chlorokiensulfaat, ekwivalent aan Chlorokien/
Chloroquine Sulphate, equivalent to Chloroquine . . . 150 mg per tablet.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslike vervaardige produk moet gevalideer word.
6. 'n Naregistrasie-inspeksie moet op die eerste produksielot van die plaaslike vervaardigde produk uitgevoer word.
7. Bemarking van die produk mag slegs 'n aanvang neem nadat 'n bevredigende naregistrasie-inspeksieverslag gedien het.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.

6. A post-registration inspection must be conducted on the first production lot of the locally manufactured product.
7. Marketing of the product may only commence following a satisfactory post-registration inspection report.

Applikant:
Applicant: Wellcome (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 10 September 1993.
10 September 1993.

Registrasienommer:
Registration Number: **Z/16/40.**

Naam van medisyne:
Name of medicine: **Andolex Oral Rinse.**

Doseringsvorm:
Dosage form: Mondspoel.
Oral rinse.

Aktiewe bestanddele:
Active ingredients: Bensiedamienhidrochloried/
Benzydamine Hydrochloride . . . 22,5 mg per 15-mℓ-vloeistof/liquid.

Voorwaardes vir registrasie: 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.

Conditions of registration: An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.

Applikant:
Applicant: Riker Laboratories Africa (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 10 September 1993.
10 September 1993.

Registrasienommer:
Registration Number: **28/2.1/0165.**

Naam van medisyne:
Name of medicine: **Isofor.**

Doseringsvorm:
Dosage form: Inhalasie vloeistof.
Inhalation liquid.

Aktiewe bestanddele:
Active ingredients: Isofluraan/
Isoflurane . . . 100 mℓ per 100-mℓ-vloeistof/liquid.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.

Applikant:
Applicant: Rhone-Poulenc Rorer SA (Pty) Ltd.

Rakleeftyd:
Shelf-life: 60 maande.
60 months.

Datum van registrasie:
Date of registration: 10 September 1993.
10 September 1993.

Registrasienommer:
Registration Number: **28/20.2.2/0153.**

Naam van medisyne:
Name of medicine: **Micomisan Vaginal.**

Doseringsvorm: Room.
Dosage Form: Cream.

Aktiewe bestanddele: Klotrimasool.
Active ingredients: Clotrimazole . . . 10 mg per 1-g-room/cream.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with al the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.

Applikant:
Applicant: Zurich Health Care (Pty) Ltd.

Rakleeftyd: 24 maande.
Shelf-life: 24 months.

Datum van registrasie: 10 September 1993.
Date of registration: 10 September 1993.

Registrasienommer:
Registration Number: **W/20.1.2/303.**

Naam van medisyne:
Name of medicine: **Lasmox 1 000 mg Tablet.**

Doseringsvorm: Filmbedekte tablet.
Dosage form: Film-coated tablet.

Aktiewe bestanddele: Amoksisillientrihidraat ekwivalent aan Amoksisillien/
Active ingredients: Amoxycillin Trihydrate equivalent to Amoxycillin . . . 1 000 mg per tablet.

Voorwaardes vir registrasie: 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.

Conditions of registration: An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.

Applikant:
Applicant: Laser Pharmaceuticals (Pty) Ltd.

Rakleeftyd: 36 maande.
Shelf-life: 36 months.

Datum van registrasie: 10 September 1993.
Date of registration: 10 September 1993.

Registrasienommer:
Registration Number: **27/20.1.2/0440.**

Naam van medisyne:
Name of medicine: **Pharmoxin S-250.**

Doseringsvorm: Poeier vir stroop.
Dosage form: Powder for syrup.

Aktiewe bestanddele: Amoksisillientrihidraat ekwivalent aan Amoksisillien/
Active ingredients: Amoxycillin Trihydrate equivalent to Amoxycillin . . . 250 mg per 5-mℓ-stroop-syrup.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of the product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.

Applikant:
Applicant: Lennon Limited.

Rakleeftyd: 24 maande.
Shelf-life: 24 months.

Datum van registrasie: 14 September 1993.
Date of registration: 14 September 1993.

Registrasienommer:
Registration Number: **27/5.2/0268.**

Naam van medisyne:
Name of medicine: **Venapulse-100.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele:
Active ingredients: Atenolol . . . 100 mg per tablet.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslike vervaardigde produk moet gevalideer word.
6. 'n Naregistrasie-inspeksie moet op die eerste produksielot van die plaaslike vervaardigde produk uitgevoer word.
7. Bemarking van die produk mag slegs 'n aanvang neem nadat 'n bevredigende naregistrasie-inspeksieverslag gedien het.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).

3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.
6. A post-registration inspection must be conducted on the first production lot of the locally manufactured product.
7. Marketing of the product may only commence following a satisfactory post-registration inspection report.

Applikant:
Applicant: Lennon Limited.

Rakleeftyd: 24 maande.
Shelf-life: 24 months.

Datum van registrasie: 14 September 1993.
Date of registration: 14 September 1993.

Registrasienommer:
Registration Number: **28/7.5/0245.**

Naam van medisyne:
Name of medicine: **Corebrate Tablets.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele: Besafibraat/
Active ingredients: Bezafibrate . . . 200 mg per tablet.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965.
3. The registration of the product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.

Applikant:
Applicant: Core Pharmaceuticals (Pty) Ltd.

Rakleeftyd: 60 maande.
Shelf-life: 60 months.

Datum van registrasie: 16 September 1993.
Date of registration: 16 September 1993.

Registrasienommer:
Registration Number: **28/7.5/0246.**

Naam van medisyne:
Name of medicine: **Corebrate Retard.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele: Besafibraat/
Active ingredients: Bezafibrate . . . 400 mg per tablet.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of the product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.

Applikant:
Applicant: Core Pharmaceuticals (Pty) Ltd.

Rakleeftyd:
Shelf-life: 60 maande.
60 months.

Datum van registrasie:
Date of registration: 16 September 1993.
16 September 1993.

Registrasienommer:
Registration Number: **28/30.2/0060.**

Naam van medisyne:
Name of medicine: **Intraglobin F 10 mℓ.**

Bereidingsvorm:
Form of preparation: Oplossing.
Solution.

Aktiewe bestanddele:
Active ingredients: Menslike Immunoglobulien/
Human Immunoglobulin . . . 50 mg per 1-mℓ-oplossing/solution.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die eerste twee produksielotte van die plaaslike vervaardigde produk moet gevalideer word.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The first two production lots of the locally manufactured product must be valided.

Applikant:
Applicant: Mednostica CC.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 18 September 1992.
18 September 1992.

Registrasienommer:
Registration Number: **28/30.2/0061.**

Naam van medisyne:
Name of medicine: **Intraglobin F 50 mℓ**, 100 mℓ and 200 mℓ.

Bereidingsvorm:
Form of preparation: Oplossing.
Solution.

Aktiewe bestanddele:
Active ingredients: Menslike Immunoglobulien/
Human Immunoglobulin . . . 50 mg per 1-mℓ-oplossing/solution.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die eerste twee produksielotte van die plaaslike vervaardigde produk moet gevalideer word.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The first two production lots of the locally manufactured product must be valided.

Applikant:
Applicant: Mednostica CC.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 18 September 1992.
18 September 1992.

Registrasienommer:
Registration Number: **28/30.2/0107.**

Naam van medisyne:
Name of medicine: **Intraglobin F 20 mℓ.**

Bereidingsvorm:
Form of preparation: Oplossing.
Solution.

Aktiewe bestanddele:
Active ingredients: Menslike Immunoglobulien/
Human Immunoglobulin . . . 50 mg per 1-mℓ-oplossing/solution.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die eerste twee produksielotte van die plaaslike vervaardige produk moet gevalideer word.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The first two production lots of the locally manufactured product must be valided.

Applikant:
Applicant: Mednostica CC.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 18 September 1992.
18 September 1992.

Registrasienommer:
Registration Number: **27/20.1.2/0262.**

Naam van medisyne:
Name of medicine: **Amoxil Fixtab 125.**

Doseringsvorm:
Dosage form: Bruis-koutablet.
Effervescent chewable tablet.

Aktiewe bestanddele:
Active ingredients: Amoksisillientrihidraat ekwivalent aan Amoksisillien/
Amoxycillin Trihydrate equivalent to Amoxycillin . . . 125 mg per tablet.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.

Applikant:
Applicant: SmithKline Beecham Pharmaceuticals (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 20 September 1993.
20 September 1993.

Registrasienommer:
Registration Number: **27/20.1.2/0263.**

Naam van medisyne:
Name of medicine: **Amoxil Fiztab 250.**

Doseringsvorm:
Dosage form: Bruis-koutablet.
Effervescent chewable tablet.

Aktiewe bestanddele:
Active ingredients: Amoksisillientrihidraat ekwivalent aan Amoksisillien/
Amoxycillin Trihydrate equivalent to Amoxycillin . . . 250 mg per tablet.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goegekeur.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of the product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.

Applikant:
Applicant: SmithKline Beecham Pharmaceuticals (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 20 September 1993.
20 September 1993.

Registrasienommer:
Registration Number: **27/20.1.2/0264.**

Naam van medisyne:
Name of medicine: **Amoxil Fixtab 500.**

Doseringsvorm: Bruis-koutablet.
Dosage form: Effervescent chewable tablet.

Aktiewe bestanddele: Amoksisillientrihidraat ekwivalent aan Amoksisillien/
Active ingredients: Amoxycillin Trihydrate equivalent to Amoxycillin . . . 500 mg per tablet.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.

Applikant:
Applicant: SmithKline Beecham Pharmaceuticals (Pty) Ltd.

Rakleeftyd: 24 maande.
Shelf-life: 24 months.

Datum van registrasie: 20 September 1993.
Date of registration: 20 September 1993.

Registrasienommer:
Registration Number: **27/5.2/0483.**

Naam van medisyne:
Name of medicine: **Salternol.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele: Propranololhidrochloried/
Active ingredients: Propranolol Hydrochloride . . . 40 mg per tablet.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslike vervaardige produk moet gevalideer word.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.

Applikant:
Applicant: Zurich Health Care (Pty) Ltd.

Rakleeftyd: 24 maande.
Shelf-life: 24 months.

Datum van registrasie: 20 September 1993.
Date of registration: 20 September 1993.

Registrasienommer:
Registration Number: **27/10.2.1/0556.**

Naam van medisyne:
Name of medicine: **ZX Salbutamol Inhaler.**

Doseringsvorm: Inhaleerder.
Dosage form: Inhaler.

Aktiewe bestanddele:
Active ingredients: Salbutamol . . . 100 µg per afgemete dosis/metered dose.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.

Applikant:
Applicant: Glaxo South Africa (Pty) Ltd.

Rakleeftyd: 24 maande.
Shelf-life: 24 months.

Datum van registrasie: 20 September 1993.
Date of registration: 20 September 1993.

Registrasienommer:
Registration Number: **27/20.1.2/0475.**

Naam van medisyne:
Name of medicine: **Ampisalt 125 mg.**

Doseringsvorm: Poeier vor suspensie.
Dosage form: Powder for suspension.

Aktiewe bestanddele:
Active ingredients: Ampsillientrihidraat ekwivalent aan Ampisillien/
Ampicillin Trihydrate equivalent to Ampicillin . . . 125 mg per 5-mℓ-suspensie/suspension.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslike vervaardige produk moet gevalideer word.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be valided.

Applikant:
Applicant: Zurich Health Care (Pty) Ltd.

Rakleeftyd: 24 maande.
Shelf-life: 24 months.

Datum van registrasie: 20 September 1993.
Date of registration: 20 September 1993.

Registrasienommer:
Registration Number: **27/5.7.2/0170.**

Naam van medisyne:
Name of medicine: **Clomax.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele:
Active ingredients: Metoklopramiedhidrochloried/
Metoclopramide Hydrochloride . . . 10 mg per tablet.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslike vervaardige produk moet gevalideer word.
6. 'n Naregistrasie-inspeksie moet op die eerste produksielot van die plaaslike vervaardigde produk uitgevoer word.
7. Bemarking van die produk mag slegs 'n aanvang neem nadat 'n bevredigende naregistrasie-inspeksieverslag gedien het.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.
6. A post-registration inspection must be conducted on the first production lot of the locally manufactured product.
7. Marketing of the product may only commence following a satisfactory post-registration inspection report.

Applikant:
Applicant: Amynos Pharmaceuticals (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 20 September 1993.
20 September 1993.

Registrasienommer:
Registration Number: **E/11.4.1/941.**

Naam van medisyne:
Name of medicine: **pH550 Peppermint Antacid Tablets.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele:
Active ingredients: Kalsiumkarbonaat/
Calcium Carbonate . . . 489,9 mg.
Magnesiumkarbonaat/
Magnesium Carbonate . . . 11,4 mg.
Magnesiumtrisilikaat/
Magnesium Trisilicate . . . 5,7 mg per tablet.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte na registrasie moet gevalideer word, tensy hierdie dokumentasie beskikbaar is.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots after registration must be vlidated, unless this documentation is available.

Applikant:
Applicant: The Premier Pharmaceutical Company Limited.

Rakleeftyd:
Shelf-life: 12 maande.
12 months.

Datum van registrasie:
Date of registration: 20 September 1993.
20 September 1993.

Registrasienommer:
Registration Number: **Z/20.1.1/335.**

Naam van medisyne:
Name of medicine: **Tarivid I.V. 100 mg.**

Bereidingsvorm:
Form of preparation: Oplossing vir infuus.
Infusion solution.

Aktiewe bestanddele:
Active ingredients: Ofloksasienhidrochloried ekwivalent aan Ofloksasien/
Ofloxacin Hydrochloride equivalent to Ofloxacin . . . 100 mg per 50-mℓ-flessie/vial.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die eerste twee produksielotte van die plaaslike vervaardige produk moet gevalideer word.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The first two production lots of the locally manufactured product must be validated.

Applikant:
Applicant: Noristan Limited.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 20 September 1993.
20 September 1993.

Registrasienommer:
Registration Number: **Z/20.1.1/336.**

Naam van medisyne:
Name of medicine: **Tarivid I.v. 200 mg.**

Bereidingsvorm:
Form of preparation: Oplossing vir infuus.
Infusion solution.

Aktiewe bestanddele:
Active ingredients: Ofloksasienhidrochloried ekwivalent aan Ofloksasien/
Ofloxacin Hydrochloride equivalent to Ofloxacin . . . 200 mg per 100-mℓ-flessie/vial.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die eerste twee produksielotte van die plaaslike vervaardigde produk moet gevalideer word.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The first two production lots of the locally manufactured product must be validated.

Applikant:
Applicant: Noristan Limited.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 20 September 1993.
20 September 1993.

Registrasienommer:
Registration Number: **27/11/0306.**

Naam van medisyne:
Name of medicine: **Prepulsid 20 mg.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele:
Active ingredients: Sisapriedmonohidraat ekwivalent aan Sisapried/
Cisapride Monohydrate equivalent to Cisapride . . . 20 mg per tablet.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.
6. 'n Naregistrasie-inspeksie moet op die eerste produksielot van die plaaslike vervaardigde produk uitgevoer word.
7. Bemarking van die produk mag slegs 'n aanvang neem nadat 'n bevredigende naregistrasie-inspeksieverslag gedien het.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.

4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.
6. A post-registration inspection must be conducted on the first production lot of the locally manufactured product.
7. Marketing of the product may only commence following a satisfactory post-registration inspection report.

Applikant:
Applicant: Janssen Pharmaceutica (Pty) Ltd.

Rakleeftyd: 24 maande.
Shelf-life: 24 months.

Datum van registrasie: 20 September 1993.
Date of registration: 20 September 1993.

Registrasienommer:
Registration Number: **Z/20.1.1/286.**

Naam van medisyne:
Name of medicine: **Mytobrin Injection 20 mg/2-mℓ.**

Doseringsvorm: Inspuiting.
Dosage form: Injection.

Aktiewe bestanddele: Tobramisiensulfaat ekwivalent aan Tobramisien/
Active ingredients: Tobramycin Sulphate equivalent to Tobramycin . . . 20 mg per 2-mℓ-flessie/vial.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die eerste twee produksielotte moet gevalideer word.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The first two production lots must be validated.

Applikant:
Applicant: Intramed (Pty) Ltd.

Rakleeftyd: 24 maande.
Shelf-life: 24 months.

Datum van registrasie: 20 September 1993.
Date of registration: 20 September 1993.

Registrasienommer:
Registration Number: **E/11.4.1/1271.**

Naam van medisyne:
Name of medicine: **Maalox.**

Doseringsvorm: Suspensie.
Dosage form: Suspension.

Aktiewe bestanddele: Magnesiumhidroksiedpasta ekwivalent aan Magnesiumhidroksied/
Active ingredients: Magnesium Hydroxide Paste equivalent to Magnesium Hydroxide . . . 200 mg.
Nat Aluminiumhidroksiedjel ekwivalent aan Droë Aluminiumhidroksiedjel/
Aluminium Hydroxide Wet Gel equivalent to Dried Aluminium Hydroxide Gel . . . 225 mg per 5-mℓ-suspensie/suspension.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslike vervaardigde produk moet gevalideer word.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.

Applikant:
Applicant: Rhône-Poulenc Rorer SA (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 20 September 1993.
20 September 1993.

Registrasienommer:
Registration Number: **E/11.4.1/1273.**

Naam van medisyne:
Name of medicine: **Extra Strength Maalox.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele:
Active ingredients: Magnesiumhidroksied/
Magnesium Hydroxide . . . 400 mg.
Gedroogde Aluminiumhidroksiedjel/
Dried Aluminium Hydroxide Gel . . . 400 mg per tablet.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.

Applikant:
Applicant: Rhône-Poulenc Rorer SA (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 20 September 1993.
20 September 1993.

Registrasienommer:
Registration Number: **27/24/0538.**

Naam van medisyne:
Name of medicine: **Reach Anti Cavity Fluoride Rinse.**

Doseringsvorm:
Dosage form: Mondspoel.
Mouthwash.

Aktiewe bestanddele:
Active ingredients: Natriumfluoried/
Sodium Fluoride . . . 2,5 mg per 5-mℓ-vloeistof/liquid.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.

Applikant:
Applicant: Pharmedica (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 20 September 1993.
20 September 1993.

Registrasienommer:
Registration Number: **27/20.1.1/0278.**

Naam van medisyne:
Name of medicine: **Zithromax Capsules 250 mg.**

Doseringsvorm:
Dosage form: Kapsuul.
Capsule.

Aktiewe bestanddele:
Active ingredients: Asitromisiendihidraat ekwivalent aan Asitromisien/
Azithromycin Dihydrate equivalent to Azithromycin . . . 250 mg per kapsuul/capsule.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.
6. 'n Naregistrasie-inspeksie moet op die eerste produksielot van die plaaslik vervaardigde produk uitgevoer word.
7. Bemarking van die produk mag slegs 'n aanvang neem nadat 'n bevredigende naregistrasie-inspeksieverslag gedien het.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.

6. A post-registration inspection must be conducted on the first production lot of the locally manufactured product.
7. Marketing of the product may only commence following a satisfactory post-registration inspection report.

Applikant:
Applicant: Pfizer Laboratories (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 20 September 1993.
20 September 1993.

Registrasienommer:
Registration Number: **27/20.1.1/0279.**

Naam van medisyne:
Name of medicine: **Zithromax Powder for Oral Suspension.**

Doseringsvorm:
Dosage form: Poeier vir suspensie.
Powder for suspension.

Aktiewe bestanddele:
Active ingredients: Asitromisiendihidraat ekwivalent aan Asitromisien/
Azithromycin Dihydrate equivalent to Azithromycin . . . 200 mg per 5-mℓ-suspensie/suspension.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.
6. 'n Naregistrasie-inspeksie moet op die eerste produksielot van die plaaslik vervaardigde produk uitgevoer word.
7. Bemarking van die produk mag slegs 'n aanvang neem nadat 'n bevredigende naregistrasie-inspeksieverslag gedien het.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.
6. A post-registration inspection must be conducted on the first production lot of the locally manufactured product.
7. Marketing of the product may only commence following a satisfactory post-registration inspection report.

Applikant:
Applicant: Pfizer Laboratories (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 20 September 1993.
20 September 1993.

Registrasienommer:
Registration Number: **27/2.6.5/0235.**

Naam van medisyne:
Name of medicine: **Risperdal 1 mg.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele:
Active ingredients: Risperidoon/
Risperidone . . . 1 mg per tablet.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.
6. 'n Naregistrasie-inspeksie moet op die eerste produksielot van die plaaslik vervaardigde produk uitgevoer word.
7. Bemarking van die produk mag slegs 'n aanvang neem nadat 'n bevredigende naregistrasie-inspeksieverslag gedien het.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacuture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.
6. A post-registration inspection must be conducted on the first production lot of the locally manufactured product.
7. Marketing of the product may only commence following a satisfactory post-registration inspection report.

Applikant:
Applicant: Janssen Pharmaceutica (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 23 September 1993.
23 September 1993.

Registrasienommer:
Registration Number: **27/2.6.5/0236.**

Naam van medisyne:
Name of medicine: **Risperdal 2 mg.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele:
Active ingredients: Risperidoon/
Risperidone . . . 2 mg per tablet.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.

5. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.
6. 'n Naregistrasie-inspeksie moet op die eerste produksielot van die plaaslik vervaardigde produk uitgevoer word.
7. Bemarking van die produk mag slegs 'n aanvang neem nadat 'n bevredigende naregistrasie-inspeksieverslag gedien het.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacuture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.
6. A post-registration inspection must be conducted on the first production lot of the locally manufactured product.
7. Marketing of the product may only commence following a satisfactory post-registration inspection report.

Applikant:
Applicant: Janssen Pharmaceutica (Pty) Ltd.

Rakleeftyd: 24 maande.
Shelf-life: 24 months.

Datum van registrasie: 23 September 1993.
Date of registration: 23 September 1993.

Registrasienommer:
Registration Number: **27/2.6.5/0237.**

Naam van medisyne:
Name of medicine: **Risperdal 3 mg.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele:
Active ingredients: Risperidoon/
Risperidone . . . 3 mg per tablet.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.
6. 'n Naregistrasie-inspeksie moet op die eerste produksielot van die plaaslike vervaardigde produk uitgevoer word.
7. Bemarking van die produk mag slegs 'n aanvang neem nadat 'n bevredigende naregistrasie-inspeksieverslag gedien het.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacuture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.

6. A post-registration inspection must be conducted on the first production lot of the locally manufactured product.
7. Marketing of the product may only commence following a satisfactory post-registration inspection report.

Applikant:
Applicant: Janssen Pharmaceutica (Pty) Ltd.

Rakleeftyd: 24 maande.
Shelf-life: 24 months.

Datum van registrasie: 23 September 1993.
Date of registration: 23 September 1993.

Registrasienommer:
Registration Number: **27/2.6.5/0238.**

Naam van medisyne:
Name of medicine: **Risperdal 4 mg.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele: Risperidoon/
Active ingredients: Risperidone . . . 4 mg per tablet.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.
6. 'n Naregistrasie-inspeksie moet op die eerste produksielot van die plaaslike vervaardigde produk uitgevoer word.
7. Bemarking van die produk mag slegs 'n aanvang neem nadat 'n bevredigende naregistrasie-inspeksieverslag gedien het.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacuture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.
6. A post-registration inspection must be conducted on the first production lot of the locally manufactured product.
7. Marketing of the product may only commence following a satisfactory post-registration inspection report.

Applikant:
Applicant: Janssen Pharmaceutica (Pty) Ltd

Rakleeftyd: 24 maande.
Shelf-life: 24 months.

Datum van registrasie: 23 September 1993.
Date of registration: 23 September 1993.

Registrasienommer:
Registration Number: **Z/30.2/370.**

Naam van medisyne:
Name of medicine: **Globuman Berna I.V. 20 mℓ.**

Doseringsvorm: Inspuiting.
Dosage form: Injection.

Aktiewe bestanddele: Menslike Immunoglobulien/
Active ingredients: Human Immunoglobulin . . . 1 g per 20-mℓ-flessie/vial.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of the product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.

Applikant:
Applicant: Swisspharm (Pty) Ltd.

Rakleeftyd: 36 maande.
Shelf-life: 36 months.

Datum van registrasie: 24 September 1993.
Date of registration: 24 September 1993.

Registrasienommer:
Registration Number: **Z/30.2/371**

Naam van medisyne:
Name of medicine: **Globuman Berna I.V. 50 mℓ.**

Doseringsvorm: Inspuiting.
Dosage form: Injection.

Aktiewe bestanddele: Menslike Immunoglobulien/
Active ingredients: Human Immunoglobulin . . . 2,5 g per 50-mℓ-flessie/vial.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of the product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.

Applikant:
Applicant: Swisspharm (Pty) Ltd.

Rakleeftyd: 36 maande.
Shelf-life: 36 months.

Datum van registrasie: 24 September 1993.
Date of registration: 24 September 1993.

Registrasienommer:
Registration Number: **Z/30.2/372.**

Naam van medisyne:
Name of medicine: **Globuman Berna I.V. 100 mℓ.**

Doseringsvorm: Inspuiting.
Dosage form: Injection.

Aktiewe bestanddele: Menslike Immunoglobulien/
Active ingredients: Human Immunoglobulin . . . 5 g per 100-mℓ-flessie-vial.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of the product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.

Applikant:
Applicant: Swisspharm (Pty) Ltd.

Rakleeftyd: 36 maande.
Shelf-life: 36 months.

Datum van registrasie: 24 September 1993.
Date of registration: 24 September 1993.

Registrasienommer:
Registration Number: **28/11.9/0146.**

Naam van medisyne:
Name of medicine: **Loperol.**

Doseringsvorm: Kapsuul.
Dosage form: Capsule.

Aktiewe bestanddele: Loperamiedhidrochloried/
Active ingredients: Loperamide Hydrochloride . . . 2 mg per kapsuul/capsule.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.
6. 'n Naregistrasie-inspeksie moet op die eerste produksielot van die plaaslike vervaardigde produk uitgevoer word.
7. Bemarking van die produk mag slegs 'n aanvang neem nadat 'n bevredigende naregistrasie-inspeksieverslag gedien het.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.

4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.
6. A post-registration inspection must be conducted on the first production lot of the locally manufactured product.
7. Marketing of the product may only commence following a satisfactory post-registration inspection report.

Applikant:
Applicant: Rolab (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 24 September 1993.
24 September 1993.

Registrasienommer:
Registration Number: **27/5.8/0272.**

Naam van medisyne:
Name of medicine: **Goldgestant.**

Doseringsvorm:
Dosage form: Vloeistof.
Liquid.

Aktiewe bestanddele:
Active ingredients: Chloorfeniramienmaleaat/
Chlorpheniramine Maleate . . . 2 mg.
Fenielpropanolamienhidrochloried/
Phenylpropanolamine Hydrochloride . . . 2,5 mg.
Fenielefrienhidrochloried/
Phenylephrine Hydrochloride . . . 2,5 mg per 5-mℓ -vloeistof/liquid.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslike vervaardigde produk moet gevalideer word.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lost of the locally manufactured product must be validated.

Applikant:
Applicant: GD Searle (SA) (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 24 September 1993.
24 September 1993.

Registrasienommer:
Registration Number: **28/20.2.2/0119.**

Naam van medisyne:
Name of medicine: **Medaspor Topical.**

Doseringsvorm:
Dosage form: Room.
Cream.

Aktiewe bestanddele:
Active ingredients: Klotrimasool/
Clotrimazole . . . 10 mg per 1-g-room/cream.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslike vervaardigde produk moet gevalideer word.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lost of the locally manufactured product must be validated.

Applikant:
Applicant: Medpro Pharmaceutica (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 24 September 1993.
24 September 1993.

Registrasienommer:
Registration Number: **28/11.4.1/0235.**

Naam van medisyne:
Name of medicine: **Gentrisil.**

Doseringsvorm:
Dosage Form: Suspensie.
Suspension.

Aktiewe bestanddele:
Active ingredients:
Magnesiumtrisilikaat/
Magnesium Trisilicate . . . 500 mg.
Ligte Magnesiumkarbonaat/
Light Magnesium Carbonate . . . 500 mg.
Natriumbikarbonaat/
Sodium Bicarbonate . . . 500 mg per 10-mℓ-suspensie/suspension.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of the product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.

Applikant:
Applicant: Genpharm Pharmaceuticals (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 24 September 1993.
24 September 1993.

Registrasienommer:
Registration Number: **T/30.2/642.**

Naam van medisyne:
Name of medicine: **Hepuman Berna.**

Doseringsvorm: Inspuiting.
Dosage Form: Injection.

Aktiewe bestanddele: Menslike Immunoglobulien/
Active ingredients: Human Immunoglobulin . . . 100–160 mg.
Hepatitis B teenliggaampies/
Hepatitis B antibodies . . . 200 I.E./I.U. per 1-mℓ-flessie/vial.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).

Applikant:
Applicant: Swisspharm (Pty) Ltd.

Rakleeftyd: 36 maande teen 2–8°C.
Shelf-life: 36 months at 2–8°C.

Datum van registrasie: 24 September 1993.
Date of registration: 24 September 1993.

Registrasienommer:
Registration Number: **28/20.1.2/0124.**

Naam van medisyne:
Name of medicine: **Zoxil 250.**

Doseringsvorm: Kapsuul.
Dosage Form: Capsule.

Aktiewe bestanddele: Amoksisillientrihidraat ekwivalent aan Amoksisillien/
Active ingredients: Amoxycillin Trihydrate equivalent to Amoxycillin . . . 250 mg per kapsuul/capsule.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of the product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.

Applikant:
Applicant: GS Pharmaceuticals (Pty) Ltd.

Rakleeftyd: 48 maande.
Shelf-life: 48 months.

Datum van registrasie: 24 September 1993.
Date of registration: 24 September 1993.

Registrasienommer:
Registration Number: **27/3.1/0437.**

Naam van medisyne:
Name of medicine: **Nafasol EC-250.**

Doseringsvorm:
Dosage Form: Tablet.

Aktiewe bestanddele:
Active ingredients: Naproksen/
Naproxen . . . 250 mg per tablet.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.
6. 'n Naregistrasie-inspeksie moet op die eerste produksielot van die plaaslike vervaardigde produk uitgevoer word.
7. Bemarking van die produk mag slegs 'n aanvang neem nadat 'n bevredigende naregistrasie-inspeksieverslag gedien het.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.
6. A post-registration inspection must be conducted on the first production lot of the locally manufactured product.
7. Marketing of the product may only commence following a satisfactory post-registration inspection report.

Applikant:
Applicant: Lennon Limited.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 24 September 1993.
24 September 1993.

Registrasienommer:
Registration Number: **27/3.1/0438.**

Naam van medisyne:
Name of medicine: **Nafasol EC–500.**

Doseringsvorm:
Dosage Form: Tablet.

Aktiewe bestanddele:
Active ingredients: Naproksen/
Naproxen . . . 500 mg per tablet.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).

3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.
6. 'n Naregistrasie-inspeksie moet op die eerste produksielot van die plaaslike vervaardigde produk uitgevoer word.
7. Bemarking van die produk mag slegs 'n aanvang neem nadat 'n bevredigende naregistrasie-inspeksieverslag gedien het.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.
6. A post-registration inspection must be conducted on the first production lot of the locally manufactured product.
7. Marketing of the product may only commence following a satisfactory post-registration inspection report.

Applikant:
Applicant: Lennon Limited.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 24 September 1993.
24 September 1993.

Registrasienommer:
Registration Number: **27/18.1/0482.**

Naam van medisyne:
Name of medicine: **Saltermide.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele:
Active ingredients: Furosemied/
Furosemide . . . 40 mg per tablet.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.

Applikant:
Applicant: Zurich Helath Care (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 24 September 1993.
24 September 1993.

Registrasienommer:
Registration Number: **T/30.1/674.**

Naam van medisyne:
Name of medicine: **Vaxigrip.**

Doseringsvorm: Inspuiting.
Dosage form: Injection.

Aktiewe bestanddele: Geonaktiveerde gekombineerde Griepentstof/
Active ingredients: Inactivated Combined Influenza Vaccine . . .
1 inentingsdosis/
1 vaccinating dose per 0,5-mℓ-vloeistof/liquid.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Een monster van elke lot moet tesame met ses kopieë van die protokolle vir die toets van die finale lot, sowel as ses kopieë van drie vrystellingsertifikaat wat uitgereik is deur die verantwoordelike beheerliggaam in die land waar die produk vervaardig word, ingedien word by die Raad vir lotvrystellings doeleindes.
4. Die stamme van die oorspronklike saadvirusse moet elke jaar deur die Departement van Nasionale Gesondheid en Bevolkingsontwikkeling goedgekeur word.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. One sample of every lot, together with six copies of the protocols for testing of the bulklot and filling lot and six copies of the certificate of release issued by the competent authority in the country in which the product was manufactured, be submitted to Council for lot releasing purposes.
4. The strains of the master seed viruses must be approved by the Department of National Health and Population Development for each year.

Applikant:
Applicant: Rhóne-Poulenc Rorer SA (Pty) Ltd.

Rakleeftyd: 18 maande.
Shelf-life: 18 months.

Datum van registrasie: 24 September 1993.
Date of registration: 24 September 1993.

Registrasienommer:
Registration Number: **28/20.2.2/0056.**

Naam van medisyne:
Name of medicine: **Mycoban Cream.**

Doseringsvorm: Room.
Dosage form: Cream.

Aktiewe bestanddele: Klotrimasool/
Active ingredients: Clotrimazole . . . 10 mg per 1-g-room/cream.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslike vervaardigde produk moet gevalideer word.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.

Applikant:
Applicant: Vesta Medicines (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 24 September 1993.
24 September 1993.

Registrasienommer:
Registration Number: **27/3.1/0205.**

Naam van medisyne:
Name of medicine: **Brexecam.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele:
Active ingredients: Piroksikam Beta-siklodekstrien ekwivalent aan Piroksikam/
Piroxicam Beta-cyclodextrin equivalent to Piroxicam . . . 20 mg per tablet.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslike vervaardige produk moet gevalideer word.
6. 'n Naregistrasie-inspeksie moet op die erste produksielot van die plaaslike vervaardigde produk uitgevoer word.
7. Bemarking van die produk mag slegs 'n aanvang neem nadat 'n bevredigende naregistrasie-inspeksieverslag gedien het.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.
6. A post-registration inspection must be conducted on the first production lot of the locally manufactured product.
7. Marketing of the product may only commence following a satisfactory post-registration inspection report.

Applikant:
Applicant: The Premier Pharmaceutical Company Limited.

Rakleeftyd:
Shelf-life: 36 maande.
36 months.

Datum van registrasie:
Date of registration: 24 September 1993.
24 September 1993.

Registrasienommer:
Registration Number: **T/30.1/662.**

Naam van medisyne:
Name of medicine: **Mérieux Inactivated Rabies Vaccine.**

Dorseringsvorm: Inspuiting.
Dosage form: Injection.

Aktiewe bestanddele: Geonaktiveerde hondsdolheidvirus/
Active ingredients: Inactivated Rabies Virus . . .
ten minste 2,5 I.E./
at least 2,5 I.U. per flessie/vial.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.

Applikant:
Applicant: Rhône-Poulenc Rorer SA (Pty) Ltd.

Rakleeftyd: 36 maande teen 2–8 °C.
Shelf-life: 36 months at 2–8 °C.

Datum van registrasie: 24 September 1993.
Date of registration: 24 September 1993.

Registrasienommer:
Registration Number: **91/26/3.**

Naam van medisyne:
Name of medicine: **Finadyne 20 mg.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele: Fluniksienmeglumien ekwivalent aan Fluniksien/
Active ingredients: Flunixin Meglumine equivalent to Flunixin . . . 20 mg per tablet.

Voorwaardes vir registrasie: 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.

Conditions of registration: An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.

Applikant:
Applicant: The Premier Pharmaceutical Company Ltd.

Rakleeftyd: 24 maande in stulpverpakking.
36 maande in H.D.P.E.
Shelf-life: 24 months in blister packing.
36 months in H.D.P.E.

Datum van registrasie: 30 September 1993.
Date of registration: 30 September 1993.

Registrasienommer:
Registration Number: **27/5.8/0504.**

Naam van medisyne:
Name of medicine: **Borstol Cold and Flu Powders with Vitamin C.**

Doseringsvorm: Poeier.
Dosage form: Powder.

Aktiewe bestanddele: Aspirien/
Active ingredients: Aspirin . . . 650 mg.
Askorbiensuur/
Ascorbic Acid . . . 45 mg per 736,86-mg-poeier/powder.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslike vervaardigde produk moet gevalideer word.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.

Applikant:
Applicant: Group Laboratories SA (Pty) Ltd.

Rakleeftyd: 24 maande.
Shelf-life: 24 months.

Datum van registrasie: 30 September 1993.
Date of registration: 30 September 1993.

Registrasienommer:
Registration Number: **28/21.5.1/0050.**

Naam van medisyne:
Name of medicine: **Inflammide 50.**

Doseringsvorm: Inhaleerder.
Dosage form: Inhaler.

Aktiewe bestanddele: Budesonied/
Active ingredients: Budesonide . . . 50 µg per Afgemete dosis/
Metered dose.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of the product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.

Applikant:
Applicant: Ingelheim Pharmaceuticals (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 30 September 1993.
30 September 1993.

Registrasienommer:
Registration Number: **28/21.5.1/0051.**

Naam van medisyne:
Name of medicine: **Inflammide 100.**

Doseringssvorm:
Dosage form: Inhaleerder.
Inhaler.

Aktiewe bestanddele:
Active ingredients: Budesonied/
Budesonide . . . 100 µg per afgemete dosis/metered dose.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of the product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.

Applikant:
Applicant: Ingelheim Pharmaceuticals (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 30 September 1993.
30 September 1993.

Registrasienommer:
Registration Number: **27/2.7/0567.**

Naam van medisyne:
Name of medicine: **Painkure Paediatric.**

Doseringsvorm:
Dosage Form: Elikser.
Elixir.

Aktiewe bestanddele:
Active ingredients: Parasetamol/
Paracetamol . . . 120 mg per 5-mℓ-elikser/elixir.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).

3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.

Applikant:
Applicant: African Medicines (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 30 September 1993.
30 September 1993.

Registrasienommer:
Registration Number: **27/20.2.6/0394.**

Naam van medisyne:
Name of medicine: **Saltermet 400 mg.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele:
Active ingredients: Metronidasool/
Metronidazole . . . 400 mg per tablet.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.

Applikant:
Applicant: Zurich Health Care (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 30 September 1993.
30 September 1993.

Registrasienommer:
Registration Number: **Z/30.1/239.**

Naam van medisyne:
Name of medicine: **Triviraten Berna.**

Doseringsvorm: Poeier vir inspuiting.
Dosage form: Powder for injection.

Aktiewe bestanddele: Verswakte lewendige maselvirus—Edmonston—Zagreb 19 stam/
Active ingredients: Live attenuated measles virus—Edmonston—Zagreb 19 strain . . . nie minder as/not less than 1000 TCID50.
Verswakte lewendige pampoentjie virus—Rubini stam/
Live attenuated mumps virus—Rubini strain . . . nie minder as/
not less than 5000 TCID50.
Verswakte lewendige rubella virus—Wistar stam/
Live attenuated rubella virus—Wistar strain . . . nie minder as/
not less than 1000 TCID50 per 0,5-mℓ-dosis/dose.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Een monster van elke lot, tesame met vier kopieë van die protokolle vir die toets van die finale lot en die vullot ingedien word by die Raad vir lotvrystelling doeleindes.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. One sample of every lot, together with four copies of the protocols for testing of the bulk lot and filling lot be submitted to Council for lot releasing purposes.

Applikant:
Applicant: Swisspharm (Pty) Ltd.

Rakleeftyd: 24 maande teen 2–8 °C.
Shelf-life: 24 months at 2–8 °C.

Datum van registrasie: 30 September 1993.
Date of registration: 30 September 1993.

Registrasienommer:
Registration Number: **Z/20.2.2/379.**

Naam van medisyne:
Name of medicine: **Mvcoban Vaginal Cream.**

Doseringsvorm: Room.
Dosage form: Cream.

Aktiewe bestanddele: Klotrimasool/
Active ingredients: Clotrimazole . . . 10 mg per 1-g-room/cream.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die eerste twee produksielotte moet gevalideer word.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The first two production lots must be validated.

Applikant:
Applicant: Vesta Medicines (Pty) Ltd.

Rakleeftyd: 24 maande.
Shelf-life: 24 months.

Datum van registrasie: 30 September 1993.
Date of registration: 30 September 1993.

Registrasienommer:
Registration Number: **Z/1.2/323.**

Naam van medisyne:
Name of medicine: **Mianserin Hydrochloride (Lennon) 10 mg Tablets.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele:
Active ingredients: Mianserienhidrochloried/
Mianserin Hydrochloride . . . 10 mg per tablet.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.
4. 'n Naregistrasie-inspeksie moet op die eerste produksielot van die plaaslike vervaardigde produk uitgevoer word.
5. Bemarking van die produk mag slegs 'n aanvang neem nadat 'n bevredigende naregistrasie-inspeksieverslag gedien het.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The first two production lots of the locally manufactured product must be validated.
4. A post-registration inspection must be conducted on the first production lot of the locally manufactured product.
5. Marketing of the product may only commence following a satisfactory post-registration inspection report.

Applikant:
Applicant: Lennon Limited.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 30 September 1993.
30 September 1993.

Registrasienommer:
Registration Number: **T/30.2/635.**

Naam van medisyne:
Name of medicine: **Globuman Berna 1 mℓ.**

Doseringsvorm:
Dosage form: Inspuiting.
Injection.

Aktiewe bestanddele:
Active ingredients: Gammaglobulien/
Gammaglobulin . . . 160 mg per 1-mℓ-oplossing/solution.

Voorwaardes vir registrasie: 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.

Conditions of registration: An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.

Applikant:
Applicant: Swisspharm (Pty) Ltd.

Rakleeftyd:
Shelf-life: 36 maande.
36 months.

Datum van registrasie:
Date of registration: 30 September 1993.
30 September 1993.

Registrasienommer:
Registration Number: **T/30.2/636.**

Naam van medisyne:
Name of medicine: **Globuman Berna 2 mℓ.**

Doseringsvorm: Inspuiting.
Dosage form: Injection.

Aktiewe bestanddele: Gammaglobulien/
Active ingredients: Gammaglobulin . . . 160 mg per 1-mℓ-oplossing/solution.

Voorwaardes vir registrasie: 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.

Conditions of registration: An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.

Applikant:
Applicant: Swisspharm (Pty) Ltd.

Rakleeftyd: 36 maande.
Shelf-life: 36 months.

Datum van registrasie: 30 September 1993.
Date of registration: 30 September 1993.

Registrasienommer:
Registration Number: **T/30.2/637.**

Naam van medisyne:
Name of medicine: **Globuman Berna 5 mℓ.**

Doseringsvorm: Inspuiting.
Dosage form: Injection.

Aktiewe bestanddele: Gammaglobulien/
Active ingredients: Gammaglobulin . . . 160 mg per 1-mℓ-oplossing/solution.

Voorwaardes vir registrasie: 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.

Conditions of registration: An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.

Applikant:
Applicant: Swisspharm (Pty) Ltd.

Rakleeftyd: 36 maande.
Shelf-life: 36 months.

Datum van registrasie: 30 September 1993.
Date of registration: 30 September 1993.

Registrasienommer:
Registration Number: **T/30.2/638.**

Naam van medisyne:
Name of medicine: **Globuman Berna 10 mℓ.**

Doseringsvorm: Inspuiting.
Dosage form: Injection.

Aktiewe bestanddele: Gammaglobulien/
Active ingredients: Gammaglobulin . . . 160 mg per 1-mℓ-oplossing/solution.

Voorwaardes vir registrasie: 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.

Conditions of registration: An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.

Applikant:
Applicant: Swisspharm (Pty) Ltd.

Rakleeftyd: 36 maande.
Shelf-life: 36 months.

Datum van registrasie: 30 September 1993.
Date of registration: 30 September 1993.

Registrasienommer:
Registration Number: **Z/1.2/324.**

Naam van medisyne:
Name of medicine: **Mianserin Hydrochloride (Lennon) 30 mg Tablets.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele: Mianserienhidrochloried/.
Active ingredients: Mianserin Hydrochloride . . . 30 mg per tablet.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.
4. 'n Naregistrasie-inspeksie moet op die eerste produksielot van die plaaslike vervaardigde produk uitgevoer word.
5. Bemarking van die produk mag slegs 'n aanvang neem nadat 'n bevredigende na-registrasie-inspeksieverslag gedien het.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The first two production lots of the locally manufactured product must be validated.
4. A post-registration inspection must be conducted on the first production lot of the locally manufactured product.
5. Marketing of the product may only commence following a satisfactory post-registration inspection report.

Applikant:
Applicant: Lennon Limited.

Rakleeftyd: 24 maande.
Shelf-life: 24 months.

Datum van registrasie: 30 September 1993.
Date of registration: 30 September 1993.

Registrasienommer:
Registration Number: **28/21.5.1/0052.**

Naam van medisyne:
Name of medicine: **Inflammide 200.**

Doseringsvorm: Inhaleerder.
Dosage Form: Inhaler.

Aktiewe bestanddele: Budesonied/
Active ingredients: Budesonide . . . 200 ug per afgemete dosis/metered dose.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardiginspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of the product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.

Applikant:
Applicant: Ingelheim Pharmaceuticals (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 30 September 1993.
30 September 1993.

Registrasienommer:
Registration Number: **27/18.1/0445.**

Naam van medisyne:
Name of medicine: **Dino-Retic.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele:
Active ingredients: Furosemied/
Furosemide . . . 40 mg per tablet.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksieslotte van die plaaslike vervaardigde produk moet gevalideer word.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to the review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.

Applikant:
Applicant: Beige Pharmaceuticals CC.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 30 September 1993.
30 September 1993.

Registrasienommer:
Registration Number: **27/3.1/0315.**

Naam van medisyne:
Name of medicine: **Flexagen-50.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele:
Active ingredients: Natriumdiklofenak
Diclofenac Sodium . . . 50 mg per tablet.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of the product shall be subject to review every three years.
4. The information in the package insert shall be upated on a regular basis to conform to a package insert recently approved by the Council.

Applikant:
Applicant: Lennon Limited.

Rakleeftyd: 24 maande.
Shelf-life: 24 months.

Datum van registrasie: 5 Oktober 1993.
Date of registration: 5 October 1993.

Registrasienommer:
Registration Number: **28/11.4.3/0255.**

Naam van medisyne:
Name of medicine: **Glaxo-Ranitidine 150.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele: Ranitidienhidrochloried ekwivalent aan Ranitidien/
Active ingredients: Ranitidine Hydrochloride equivalent to Ranitidine . . . 150 mg per tablet.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligtingbiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of the product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.

Applikant:
Applicant: Glaxo SA (Pty) Ltd.

Rakleeftyd: 60 maande.
Shelf-life: 60 months.

Datum van registrasie: 5 Oktober 1993.
Date of registration: 5 October 1993.

Registrasienommer:
Registration Number: **28/11.4.3/0256.**

Naam van medisyne:
Name of medicine: **Glaxo-Ranitidine 300.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele:
Active ingredients: Ranitidienhidrochloried ekwivalent aan Ranitidien/
Ranitidine Hydrochloride equivalent to Ranitidine . . . 300 mg per tablet.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of the product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.

Applikant:
Applicant: Glaxo SA (Pty) Ltd.

Rakleeftyd:
Shelf-life: 36 maande.
36 months.

Datum van registrasie:
Date of registration: 5 Oktober 1993.
5 October 1993.

Registrasienommer:
Registration Number: **28/20.2.6/0144.**

Naam van medisyne:
Name of medicine: **Nidatron 200 mg.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele:
Active ingredients: Metronidasool/
Metronidazole . . . 200 mg per tablet.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslike vervaardigde produk moet gevalideer word.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.

Applikant:
Applicant: Zurich Health Care (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 5 Oktober 1993.
5 October 1993.

Registrasienommer:
Registration Number: **27/10.1/0536.**

Naam van medisyne:
Name of medicine: **Trifen Expect Paediatric.**

Doseringsvorm: Oplossing.
Dosage form: Solution.

Aktiewe bestanddele:
Active ingredients:
Triprolidienhidrochloried/
Triprolidine Hydrochloride . . . 0,6 mg.
Pseudoefedrienhidrochloried/
Pseudoephedrine Hydrochloride . . . 12 mg.
Guaifenesien/
Guaiphenesin . . . 50 mg.
Kodeïenfosfaat/
Codeine Phosphate . . . 3 mg per 5 mℓ-oplossing/solution.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksieslotte van die plaaslike vervaardigde produk moet gevalideer word.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two lots of the locally manufactured product must be validated.

Applikant:
Applicant: Be-Tabs Pharmaceuticals (Pty) Ltd.

Rakleeftyd: 24 maande.
Shelf-life: 24 months.

Datum van registrasie: 5 Oktober 1993.
Date of registration: 5 October 1993.

Registrasienommer:
Registration Number: **27/17.1/0569.**

Naam van medisyne:
Name of medicine: **Mivacron 5.**

Doseringsvorm: Inspuiting.
Dosage form: Injection.

Aktiewe bestanddele:
Active ingredients:
Mivakuriumchloried ekwivalent aan Mivakurium/
Mivacurium Chloride equivalent to Mivacurium . . . 10 mg per 5-mℓ-ampuul/ampoule.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van goeie vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.

Conditions of registration:

1. An acceptable standard of good manufacturing practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of the product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.

Applikant:
Applicant: Wellcome (Pty) Ltd.

Rakleeftyd:
Shelf-life: 18 maande.
18 months.

Datum van registrasie:
Date of registration: 5 Oktober 1993.
5 October 1993.

Registrasienommer:
Registration Number: **27/4/0223.**

Naam van medisyne:
Name of medicine: **Denthesen — 2% A.**

Doseringsvorm:
Dosage form: Inspuiting.
Injection.

Aktiewe bestanddele:
Active ingredients: Lignokaienhidrochloried/
Lignocaine Hydrochloride . . . 36 mg.
Adrenaliensuurtartraat/
Adrenaline Acid Tartrate . . . 0,0324 mg per 1,8-mℓ-ampuul/ampoule.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van goeie vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.

Conditions of registration:

1. An acceptable standard of good manufacturing practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of the product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.

Applikant:
Applicant: Wellcome (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 5 Oktober 1993.
5 October 1993.

Registrasienommer:
Registration Number: **27/4/0309.**

Naam van medisyne:
Name of medicine: **Denthesen—3% A.**

Doseringsvorm:
Dosage form: Inspuiting.
Injection.

Aktiewe bestanddele:
Active ingredients: Lignokaienhidrochloried/
Lignocaine Hydrochloride . . . 54 mg.
Adrenaliensuurtartraat/
Adrenaline Acid Tartrate . . . 0,0324 mg per 1,8-mℓ-ampuul/ampoule.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslike vervaardigde produk moet gevalideer word.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.

Applikant:
Applicant: SCP Pharmaceuticals (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 5 Oktober 1993.
5 October 1993.

Registrasienommer:
Registration Number: **27/2.7/0240.**

Naam van medisyne:
Name of medicine: **Maridol Tablets.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele:
Active ingredients: Ibuprofeen.
Ibuprofen . . . 200 mg per tablet.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.
6. 'n Naregistrasie-inspeksie moet op die eerste produksielot van die plaaslike vervaardigde produk uitgevoer word.
7. Bemarking van die produk mag slegs 'n aanvang neem nadat 'n bevredigende na-registrasie-inspeksieverslag gedien het.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.

6. A post-registration inspection must be conducted on the first production lot of the locally manufactured product.
7. Marketing of the product may only commence following a satisfactory post-registration inspection report.

Applikant:
Applicant: Pharmaceutical Enterprises (Pty) Ltd.

Rakleeftyd: 24 maande.
Shelf-life: 24 months.

Datum van registrasie: 5 Oktober 1993.
Date of registration: 5 October 1993.

Registrasienommer:
Registration Number: **Z/1.2/203.**

Naam van medisyne:
Name of medicine: **Plazid.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele: Indapamiedhemihidraat ekwivalent aan Indapamied/
Active ingredients: Indapamide Hemihydrate equivalent to Indapamide . . . 2,5 mg per tablet.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.
3. 'n Naregistrasie-inspeksie moet op die eerste produksielot van die plaaslik vervaardigde produk uitgevoer word.
4. Bemarking van die produk mag slegs 'n aanvang neem nadat 'n bevredigende na-registrasie-inspeksieverslag gedien het.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The first two production lots of the locally manufactured product must be validated.
3. A post-registration inspection must be conducted on the first production lot of the locally manufactured product.
4. Marketing of the product may only commence following a satisfactory post-registration inspection report.

Applikant:
Applicant: Rolab (Pty) Ltd.

Rakleeftyd: 24 maande.
Shelf-life: 24 months.

Datum van registrasie: 11 Oktober 1993.
Date of registration: 11 October 1993.

Registrasienommer:
Registration Number: **Y/7.1/233.**

Naam van medisyne:
Name of medicine: **Lacipil 2 mg.**

Doseringsvorm/
Dosage form: Tablet.

Aktiewe bestanddele: Lasidipien/
Active ingredients: Lacidipine . . . 2 mg per tablet.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.

3. 'n Naregistrasie-inspeksie moet op die eerste produksielot van die plaaslik vervaardigde produk uitgevoer word.
4. Bemarking van die produk mag slegs 'n aanvang neem nadat 'n bevredigende na-registrasie-inspeksieverslag gedien het.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The first two production lots of the locally manufactured product must be validated.
3. A post-registration inspection must be conducted on the first production lot of the locally manufactured product.
4. Marketing of the product may only commence following a satisfactory post-registration inspection report.

Applikant:
Applicant: Glaxo SA (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 11 Oktober 1993.
11 October 1993.

Registrasienommer:
Registration Number: **Y/7.1/234.**

Naam van medisyne:
Name of medicine: **Lacipil 4 mg.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele:
Active ingredients: Lasidipien/
Lacidipine . . . 4 mg per tablet.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van goeie vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.
3. 'n Naregistrasie-inspeksie moet op die eerste produksielot van die plaaslik vervaardigde produk uitgevoer word.
4. Bemarking van die produk mag slegs 'n aanvang neem nadat 'n bevredigende naregistrasie-inspeksieverslag gedien het.

Conditions of registration:

1. An acceptable standard of good manufacturing practice must be maintained in the place of manufacture.
2. The first two production lots of the locally manufactured product must be validated.
3. A post-registration inspection must be conducted on the first production lot of the locally manufactured product.
4. Marketing of the product may only commence following a satisfactory post-registration inspection report.

Applikant:
Applicant: Glaxo SA (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 11 Oktober 1993.
11 October 1993.

Registrasienommer:
Registration Number: **27/2.2/0368.**

Naam van medisyne:
Name of medicine: **ProSom 1 mg.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele:
Active ingredients: Estasolam/
Estazolam . . . 1 mg per tablet.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van goeie vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.
6. 'n Naregistrasie-inspeksie moet op die eerste produksielot van die plaaslike vervaardigde produk uitgevoer word.
7. Bemarking van die produk mag slegs 'n aanvang neem nadat 'n bevredigende naregistrasie-inspeksieverslag gedien het.

Conditions of registration:

1. An acceptable standard of good manufacturing practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of the product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.
6. A post-registration inspection must be conducted on the first production lot of the locally manufactured product.
7. Marketing of the product may only commence following a satisfactory post-registration inspection report.

Applikant:
Applicant: Abbott Laboratories SA (Pty) Ltd.

Rakleeftyd:
Shelf-life: 36 maande.
36 months.

Datum van registrasie:
Date of registration: 11 Oktober 1993.
11 October 1993.

Registrasienommer:
Registration Number: **27/2.2/0369.**

Naam van medisyne:
Name of medicine: **ProSom 2 mg.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele:
Active ingredients: Estasolam/
Estazolam . . . 2 mg per tablet.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van goeie vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.

4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiijet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.
6. 'n Naregistrasie-inspeksie moet op die eerste produksielot van die plaaslike vervaardigde produk uitgevoer word.
7. Bemarking van die produk mag slegs 'n aanvang neem nadat 'n bevredigende naregistrasie-inspeksieverslag gedien het.

Conditions of registration:

1. An acceptable standard of good manufacturing practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.
6. A post-registration inspection must be conducted on the first production lot of the locally manufactured product.
7. Marketing of the product may only commence following a satisfactory post-registration inspection report.

Applikant:
Applicant: Abbott Laboratories SA (Pty) Ltd.

Rakleeftyd:
Shelf-life: 36 maande.
36 months.

Datum van registrasie:
Date of registration: 11 Oktober 1993.
11 October 1993.

Registrasienommer:
Registration Number: **X/11.4.1/55.**

Naam van medisyne:
Name of medicine: **Riopone Plus Chew Tablets.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele:
Active ingredients: Magaldraat/
Magaldrate . . . 480 mg.
Simetikoon/
Simethicone . . . 20 mg per tablet.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van goeie vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.
3. 'n Naregistrasie-inspeksie moet op die eerste produksielot van die plaaslik vervaardigde produk uitgevoer word.
4. Bemarking van die produk mag slegs 'n aanvang neem nadat 'n bevredigende naregistrasie-inspeksieverslag gedien het.

Conditions of registration:

1. An acceptable standard of good manufacturing practice must be maintained in the place of manufacture.
2. The first two production lots of the locally manufactured product must be validated.
3. A post-registration inspection must be conducted on the first production lot of the locally manufactured product.
4. Marketing of the product may only commence following a satisfactory post-registration inspection report.

Applikant:
Applicant: Akromed Products (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 11 Oktober 1993.
11 October 1993.

Registrasienommer:
Registration Number: **27/20.1.1/0277.**

Naam van medisyne:
Name of medicine: **Duracef 1 g Dispersable.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele:
Active ingredients: Kefadroksielmonohidraat ekwivalent aan Kefadroksiel/
Cefadroxil Monohydrate equivalent to Cefadroxil . . . 1 g per tablet.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van goeie vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.

Conditions of registration:
1. An acceptable standard of good manufacturing practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of the product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.

Applikant:
Applicant: Bristol-Myers Squibb (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 11 Oktober 1993.
11 October 1993.

Registrasienommer:
Registration Number: **E/11.4.1/1195.**

Naam van medisyne:
Name of medicine: **Bisma-Rex Powder.**

Doseringsvorm:
Dosage form: Poeier.
Powder.

Aktiewe bestanddele:
Active ingredients: Bismutaluminaat/
Bismuth Aluminate . . . 10 mg.
Aluminiumhidroksied/
Aluminium Hydroxide . . . 19 mg.
Magnesiumtrisilikaat/
Magnesium Trisilicate . . . 32,5 mg.
Kalsiumkarbonaat/
Calcium Carbonate . . . 122 mg.
Natriumbikarbonaat/
Sodium Bicarbonate . . . 656,65 mg.
Magnesiumkarbonaat/
Magnesium Carbonate . . . 153 mg per 1-g-poeier/powder.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.

Applikant:
Applicant: Ciba-Geigy (Pty) Ltd.

Rakleeftyd:
Shelf-life: 36 maande.
36 months.

Datum van registrasie:
Date of registration: 11 Oktober 1993.
11 October 1993.

Registrasienommer:
Registration Number: **27/20.2.2/0150.**

Naam van medisyne:
Name of medicine: **Candizole V.**

Doseringsvorm:
Dosage form: Room.
Cream.

Aktiewe bestanddele:
Active ingredients: Klotrimasool/
Clotrimazole . . . 50 mg per 5-g-room/cream.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslike vervaardigde produk moet gevalideer word.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.

Applikant:
Applicant: Lennon Limited.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 11 Oktober 1993.
11 October 1993.

Registrasienommer:
Registration Number: **27/34/0499.**

Naam van medisyne:
Name of medicine: **Intron-A Solution 5 million I.U./mℓ.**

Doseringsvorm:
Dosage form: Inspuiting.
Injection.

Aktiewe bestanddele:
Active ingredients: Interferon alfa-2b . . . 5 miljoen/million I.E./I.U. per 1-mℓ-oplossing/solution.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.

Applikant:
Applicant: Scherag (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande teen 2–8°C.
24 months at 2–8°C.

Datum van registrasie:
Date of registration: 11 Oktober 1993.
11 October 1993.

Registrasienommer:
Registration Number: **27/5.2/0468.**

Naam van medisyne:
Name of medicine: **Hexa-Blok 50 mg.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele:
Active ingredients: Atenolol . . . 50 mg per tablet.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.
6. 'n Naregistrasie-inspeksie moet op die eerste produksielot van die plaaslike vervaardigde produk uitgevoer word.
7. Bemarking van die produk mag slegs 'n aanvang neem nadat 'n bevredigende naregistrasie-inspeksieverslag gedien het.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.
6. A post-registration inspection must be conducted on the first production lot of the locally manufactured product.
7. Marketing of the product may only commence following a satisfactory post-registration inspection report.

Applikant:
Applicant: Hexal Pharmaceuticals (SA) (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 11 Oktober 1993.
11 October 1993.

Registrasienommer:
Registration Number: **27/5.2/0469.**

Naam van medisyne:
Name of medicine: **Hexa-Blok 100 mg.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele:
Active ingredients: Atenolol . . . 100 mg per tablet.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.
6. 'n Naregistrasie-inspeksie moet op die eerste produksielot van die plaaslike vervaardigde produk uitgevoer word.
7. Bemarking van die produk mag slegs 'n aanvang neem nadat 'n bevredigende naregistrasie-inspeksieverslag gedien het.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.
6. A post-registration inspection must be conducted on the first production lot of the locally manufactured product.
7. Marketing of the product may only commence following a satisfactory post-registration inspection report.

Applikant:
Applicant: Hexal Pharmaceuticals (SA) (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 11 Oktober 1993.
11 October 1993.

Registrasienommer:
Registration Number: **27/10.1/0553.**

Naam van medisyne:
Name of medicine: **Flusin DM.**

Doseringsvorm:
Dosage form: Stroop.
Syrup.

Aktiewe bestanddele:
Active ingredients: Chloorfeniramienmaleaat/
Chlorpheniramine Maleate . . . 2 mg.
Pseudoefedrienhidrochloried/
Pseudoephedrine Hydrochloride . . . 25 mg.
Askorbiensuur/
Ascorbic Acid . . . 50 mg.
Dekstrometorfaanhidrobromied/
Dextromethorphan Hydrobromide . . . 7,5 mg per 5-mℓ-stroop/syrup.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.

Applikant:
Applicant: Columbia Pharmaceuticals (Pty) Ltd.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 19 Oktober 1993.
19 October 1993.

Registrasienommer:
Registration Number: **27/201.2/0439.**

Naam van medisyne:
Name of medicine: **Pharmoxin S-125.**

Doseringsvorm:
Dosage form: Poeier vir stroop.
Powder for syrup.

Aktiewe bestanddele:
Active ingredients: Amoksisillientrihidraat ekwivalent aan Amoksisillien/
Amoxycillin Trihydrate equivalent to Amoxycillin . . . 125 mg per 5-mℓ-stroop/syrup.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.

Applikant:
Applicant: Lennon Limited.

Rakleeftyd:
Shelf-life: 24 maande.
24 months.

Datum van registrasie:
Date of registration: 19 Oktober 1993.
19 October 1993.

Registrasienommer:
Registration Number: **28/20.1.1/0035.**

Naam van medisyne:
Name of medicine: **Klacid P125.**

Doseringsvorm: Granules vir suspensie.
Dosage form: Granules for suspension.

Aktiewe bestanddele: Klaritromisien/
Active ingredients: Clarithromycin . . . Ellipsis 125 mg per 5-mℓ-suspensie/suspension.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.

Applikant:
Applicant: Abbott Laboratories SA (Pty) Ltd.

Rakleeftyd: 24 maande.
Shelf-life: 24 months.

Datum van registrasie: 19 Oktober 1993.
Date of registration: 19 October 1993.

Registrasienommer:
Registration Number: **E/11.4.3/1551.**

Naam van medisyne:
Name of medicine: **Woodward's Celebrated Gripe Water.**

Doseringsvorm: Mengsel.
Dosage form: Mixture.

Aktiewe bestanddele: Natriumbikarbonaat/
Active ingredients: Sodium Bicarbonate . . . 50 mg.
Dillesaadolie/
Dill Seed Oil . . . 2,15 mg per 5-mℓ-vloeistof/Liquid.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
5. Die eerste twee produksielotte van die plaaslik vervaardigde produk moet gevalideer word.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of this product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
5. The first two production lots of the locally manufactured product must be validated.

Applikant:
Applicant: Wrapsa Packaging & Manufacturing (Pty) Ltd.

Rakleeftyd: 12 maande.
Shelf-life: 12 months.

Datum van registrasie: 19 Oktober 1993.
Date of registration: 19 October 1993.

Registrasienommer:
Registration Number: **27/2.6.6/0096.**

Naam van medisyne:
Name of medicine: **Xanor 2,0 mg.**

Doseringsvorm:
Dosage form: Tablet.

Aktiewe bestanddele: Alprasolam/
Active ingredients: Alprazolam . . . 2 mg per tablet.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of the product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.

Applikant:
Applicant: Upjohn (Pty) Ltd.

Rakleeftyd: 60 maande.
Shelf-life: 60 months.

Datum van registrasie: 20 Oktober 1993.
Date of registration: 20 October 1993.

Registrasienommer:
Registration Number: **27/30.1/0127.**

Naam van medisyne:
Name of medicine: **Hepaccine-B Paediatric Injection.**

Doseringsvorm: Inspuiting.
Dosage form: Injection.

Aktiewe bestanddele: Geonaktiveerde Hepatitis B Antigeen/
Active ingredients: Inactivated Hepatitis B Antigen . . . 1,5 µg per 0,5-mℓ-flessie/vial.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of the product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.

Applikant:
Applicant: Mr Selwyn Kahanovitz.

Rakleeftyd:
Shelf-life: 24 maande teen 2–8 °C.
24 months at 2–8 °C.

Datum van registrasie:
Date of registration: 29 Oktober 1993.
29 October 1993.

Registrasienommer:
Registration Number: **27/30.1/0128.**

Naam van medisyne:
Name of medicine: **Hepaccine-B Injection.**

Doseringsvorm:
Dosage form: Inspuiting.
Injection.

Aktiewe bestanddele:
Active ingredients: Geonaktiveerde Hepatitis B Antigeen/
Inactivated Hepatitis B Antigen . . . 3 µg per 1-mℓ-flessie/vial.

Voorwaardes vir registrasie:
1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.

Conditions of registration:
1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of the product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.

Applikant:
Applicant: Mr Selwyn Kahanovitz.

Rakleeftyd:
Shelf-life: 24 maande teen 2–8 °C.
24 months at 2–8 °C.

Datum van registrasie:
Date of registration: 29 Oktober 1993.
29 October 1993.

Registrasienommer:
Registration Number: **27/30.1/0129.**

Naam van medisyne:
Name of medicine: **Hepaccine-B Multidose Injection.**

Doseringsvorm:
Dosage form: Inspuiting.
Injection.

Aktiewe bestanddele:
Active ingredients: Geonaktiveerde Hepatitis B Antigeen/
Inactivated Hepatitis B Antigen . . . 15 µg per 5-mℓ-flessie/vial.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of the product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.

Applikant:
Applicant: Mr Selwyn Kahanovitz.

Rakleeftyd:
Shelf-life: 24 maande teen 2–8 °C.
24 months at 2–8 °C.

Datum van registrasie:
Date of registration: 29 Oktober 1993.
29 October 1993.

Registrasienommer:
Registration Number: **27/30.1/0422.**

Naam van medisyne:
Name of medicine: **Hepaccine-B Paediatric Double Dose Injection.**

Doseringsvorm:
Dosage form: Inspuiting.
Injection.

Aktiewe bestanddele:
Active ingredients: Geonaktiveerde Hepatitis B Antigeen/
Inactivated Hepatitis B Antigen . . . 3 µg per 1-mℓ-flessie/vial.

Voorwaardes vir registrasie:

1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word.
2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965).
3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar.
4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.

Conditions of registration:

1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture.
2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965).
3. The registration of the product shall be subject to review every three years.
4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.

Applikant:
Applicant: Mr Selwyn Kahanovitz.

Rakleeftyd:
Shelf-life: 24 maande teen 2–8 °C.
24 months at 2–8 °C.

Datum van registrasie:
Date of registration: 29 Oktober 1993.
29 October 1993.

Registrasienommer: *Registration Number:*	**27/30.1/0423.**
Naam van medisyne: *Name of medicine:*	**Hepaccine-B Paediatric Multidose Injection.**
Doseringsvorm: *Dosage form:*	Inspuiting. Injection.
Aktiewe bestanddele: *Active ingredients:*	Geonaktiveerde Hepatitis B Antigeen/ Inactivated Hepatitis B Antigen . . . 15 µg per 5-mℓ-flessie/vial.
Voorwaardes vir registrasie:	1. 'n Aanvaarbare standaard van Goeie Vervaardigingspraktyk moet by die plek van vervaardiging gehandhaaf word. 2. Die applikant moet voldoen aan al die wetlike vereistes van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965). 3. Die registrasie van die produk is onderhewig aan hersiening elke drie jaar. 4. Die inligting in die voubiljet moet op 'n gereelde basis opgedateer word in ooreenstemming met 'n voubiljet onlangs deur die Raad goedgekeur.
Conditions of registration:	1. An acceptable standard of Good Manufacturing Practice must be maintained in the place of manufacture. 2. The applicant must comply with all the legal requirements of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965). 3. The registration of the product shall be subject to review every three years. 4. The information in the package insert shall be updated on a regular basis to conform to a package insert recently approved by the Council.
Applikant: *Applicant:*	Mr Selwyn Kahanovitz.
Rakleeftyd: *Shelf-life:*	24 maande teen 2–8 °C. 24 months at 2–8 °C.
Datum van registrasie: *Date of registration:*	29 Oktober 1993. 29 October 1993.

(3 Desember 1993)/(3 December 1993)

KENNISGEWING 1181 VAN 1993

DEPARTEMENT VAN OPENBARE WERKE

AGRÉMENT SUID-AFRIKA

(Goedkeuring van nie-gestandaardiseerde bouprodukte en -stelsels)

Kennis geskied hierby dat die geldigheid van 'n Agrément-sertifikaat, waarvan die besonderhede in die aangehegte Bylae verskyn, verval het.

BYLAE

AGRÉMENT SUID-AFRIKA

Sertifikaathouer: LTA Mac.

Onderwerp: The Conforce House.

Sertifikaat 88/179, waarvan kennisgewing van uitreiking by Kennisgewing 286 van 1988, gedateer 5 Mei 1988, uitgereik is, is gekanselleer.

(3 Desember 1993)

NOTICE 1181 OF 1993

DEPARTMENT OF PUBLIC WORKS

AGRÉMENT SOUTH AFRICA

(Approval of non-standardised building products and systems)

Notice is hereby given that the validity of an Agrément Certificate, details of which appear in the Schedule hereto, has lapsed.

SCHEDULE

AGRÉMENT SOUTH AFRICA

Certificate holder: LTA Mac.

Subject: The Conforce House.

Certificate 88/179, notice of the granting of which was given under Notice 286 of 1988, dated 5 May 1988, has been cancelled.

(3 December 1993)

KENNISGEWING 1182 VAN 1993

DEPARTEMENT VAN OPENBARE WERKE

AGRÉMENT SUID-AFRIKA

(Goedkeuring van nie-gestandaardiseerde bouprodukte en -stelsels)

Kennis geskied hierby dat die geldigheid van 'n MANTAG-sertifikaat, waarvan die besonderhede in die aangehegte Bylae verskyn, verval het.

NOTICE 1182 OF 1993

DEPARTMENT OF PUBLIC WORKS

AGRÉMENT SOUTH AFRICA

(Approval of non-standardised building products and systems)

Notice is hereby given that the validity of a MANTAG Certificate, details of which appear in the Schedule hereto, has lapsed.

BYLAE

AGRÉMENT SUID-AFRIKA

Sertifikaathouer: LTA Mac.

Onderwerp: The Conforce Economic House.

Sertifikaat 1988/M9, waarvan kennisgewing van uitreiking by Kennisgewing 287 van 1988, gedateer 5 Mei 1988, gepubliseer is, is gekanselleer.

(3 Desember 1993)

KENNISGEWING 1183 1993

DEPARTEMENT VAN OPENBARE WERKE

AGRÉMENT SUID-AFRIKA
(Goedkeuring van nie-gestandaardiseerde bouprodukte en -stelsels)

Kennis geskied hierby dat die geldigheid van 'n MANTAG-sertifikaat, waarvan die besonderhede in die aangehegte Bylae verskyn, verval het.

BYLAE

AGRÉMENT SUID-AFRIKA

Sertifikaathouer: ICM Space Frame (Edms.) Beperk.

Onderwerp: Econospace.

Sertifikaat 1988/M20, waarvan kennisgewing van uitreiking by Kennisgewing 190 van 1988, gedateer 25 Maart 1988, gepubliseer is, is gekanselleer.

(3 Desember 1993)

KENNISGEWING 1184 VAN 1993

DEPARTEMENT VAN OPENBARE WERKE

AGRÉMENT SUID-AFRIKA
(Goedkeuring van nie-gestandaardiseerde bouprodukte en -stelsels)

Kennis geskied hierby dat die geldigheid van 'n MANTAG-sertifikaat, waarvan die besonderhede in die aangehegte Bylae verskyn, verval het.

BYLAE

AGRÉMENT SUID-AFRIKA

Sertifikaathouer: Kemclad Natal (Edms.) Beperk.

Onderwerp: Securapane Latrine Superstructure.

Sertifikaat 1988/M10, waarvan kennisgewing van uitreiking by Kennisgewing 527 van 1988, gedateer 29 Julie 1988, gepubliseer is, is gekanselleer.

(3 Desember 1993)

KENNISGEWING 1185 VAN 1993

DEPARTEMENT VAN MANNEKRAG

WET OP ARBEIDSVERHOUDINGE, 1956

INTREKKING VAN REGISTRASIE VAN 'N WERKGEWERSORGANISASIE

Ek, Gerhardus Coenraad Papenfus, Assistentnywerheidsregistrateur, maak hierby kragtens artikel 14 (1) van die Wet op Arbeidsverhoudinge, 1956, bekend dat

SCHEDULE

AGRÉMENT SOUTH AFRICA

Certificate holder: LTA Mac.

Subject: The Conforce Economic House.

Certificate 1988/M9, notice of the granting of which was given under Notice 287 of 1988, dated 5 May 1988, has been cancelled.

(3 December 1993)

NOTICE 1183 OF 1993

DEPARTMENT OF PUBLIC WORKS

AGRÉMENT SOUTH AFRICA
(Approval of non-standardised building products and systems)

Notice is hereby given that the validity of a MANTAG Certificate, details of which appear in the Schedule hereto, has lapsed.

SCHEDULE

AGRÉMENT SOUTH AFRICA

Certificate holder: ICM Space Frame (Pty) Ltd.

Subject: Econospace.

Certificate 1988/M20, notice of the granting of which was given under Notice 190 of 1988 dated 25 March 1988, has been cancelled.

(3 December 1993)

NOTICE 1184 OF 1993

DEPARTMENT OF PUBLIC WORKS

AGRÉMENT SOUTH AFRICA
(Approval of non-standardised building products and systems)

Notice is hereby given that the validity of a MANTAG Certificate, details of which appear in the Schedule hereto, has lapsed.

SCHEDULE

AGRÉMENT SOUTH AFRICA

Certificate holder: Kemclad Natal (Pty) Limited.

Subject: Securapane Latrine Superstructure.

Certificate 1988/M10, notice of the granting of which was given under Notice 527 of 1988 dated 29 July 1988, has been cancelled.

(3 December 1993)

NOTICE 1185 OF 1993

DEPARTMENT OF MANPOWER

LABOUR RELATIONS ACT, 1956

CANCELLATION OF REGISTRATION OF AN EMPLOYERS' ORGANIZATION

I, Gerhardus Coenraad Papenfus, Assistant Industrial Registrar, hereby notify, in terms of section 14 (1) of the Labour Relations Act, 1956, that as I have rea-

aangesien ek rede het om te vermoed dat die Fibre Cement Association nie as werkgewersorganisasie funksioneer nie, sy registrasie ingetrek sal word, tensy redes daarteen binne 'n tydperk van 30 dae vanaf die datum van publikasie van hierdie kennisgewing aangevoer word.

G. C. PAPENFUS,
Assistentnywerheidsregistrateur.
(3 Desember 1993)

son to believe that the Fibre Cement Association is not functioning as an employers' organization, its registration will be cancelled unless cause to the contrary is shown within a period of 30 days from the date of publication of this notice.

G. C. PAPENFUS,
Assistant Industrial Registrar.
(3 December 1993)

KENNISGEWING 1189 VAN 1993

DEPARTEMENT VAN HANDEL EN NYWERHEID

HANDELSWAREMERKE-WET 1941
(WET No. 17 VAN 1941)

VOORGENOME VERBOD OP DIE GEBRUIK VAN 'N SEKERE EMBLEEM

Ooreenkomstig die vereistes van artikel 13 van die Handelswaremerke-wet, 1941, word hierby bekendgemaak dat die Wêreldorganisasie vir Intellektuele Eiendom (WIPO) ingevolge artikel 6*ter* van die Konvensie van Parys vir die Beskerming van Intellektuele Eiendom, 1883, soos gewysig, 'n versoek namens die "EUROPEAN FREE TRADE ASSOCIATION (EFTA)" gerig het dat 'n verbod kragtens artikel 15 (1) van die Wet op die gebruik van die "NEW EMBLEM NO 1" bestaande uit die vlae van die lidlande, nl. Oostenryk, Finland, Ysland, Lichtenstein, Noorweë, Swede en Switserland soos hieroder aangedui, geplaas word, in verband met enige handel, besigheid, beroep of bedryf of in verband met 'n handelsmerk, merk of handelsomskrywing wat op ware aangebring is, uitgesonderd die gebruik daarvan deur die genoemde Vereniging of sy gevolmagtigdes.

Belanhebbendes word versoek om vertoë wat hulle in verband met die aangeleentheid wil rig, skriftelik by die Registrateur van Handelsmerke, Privaatsak X400, Pretoria, 0001, in te dien sodat dit hom binne 30 dae na publikasie van die kennisgewing bereik.
(3 Desember 1993)

NOTICE 1189 OF 1993

DEPARTMENT OF TRADE AND INDUSTRY

MERCHANDISE MARKS ACT, 1941
(ACT No. 17 OF 1941)

PROPOSED PROHIBITION OF THE USE OF A CERTAIN EMBLEM

In pursuance of the requirements of section 13 of the Merchandise Marks Act, 1941, it is hereby notiied that the World Intellectual Property Organisation (WIPO) has, by virtue of article 6*ter* of the Paris Convention for the Protection of Intellectual Property, 1883, as amended, conveyed a request on behalf of the EUROPEAN FREE TRADE ASSOCIATION (EFTA) for the prohibition in terms of section 15 (1) of the said Act, of the use of the NEW EMBLEM NO 1 consisting of the flags of the member countries, i.e. Austria, Finland, Iceland, Lichtenstein, Norway, Sweden and Switzerland as depicted hereunder, in connection with any trade, business, profession or occupation or in connection with a trade mark, mark or trade description applied to goods, other than the use thereof by the said Association or its mandatories.

Interested persons are invited to submit, in writing, such representations as they may care to make in regard to the matter to the Registrar of Trade Marks, Privte Bag X400, Pretoria, 0001, to reach him within 30 days of the publication of this notice.
(3 December 1993)

KENNISGEWING 1190 VAN 1993

RAAD OP FINANSIËLE DIENSTE

WET OP BEHEER VAN EFFEKTE-TRUSTSKEMAS, No. 54 VAN 1981

VRYSTELLING VAN DIE BEPALINGS VAN ARTIKEL 34 (4) (a)

Ek, Petrus Johannes Badenhorst, Registrateur van Effekte-trustmaatskappye, stel hierby, na oorlegpleging met die Advieskomitee oor Effekte-trustskemas, kragtens artikel 45 van die Wet op Beheer van Effekte-trustskemas, 1981 (Wet No. 54 van 1981), Property Fund Managers Beperk, 'n geregistreerde bestuursmaatskappy van 'n effekte-trustskema in eiendomsaandele ingevolge genoemde Wet, vry van die bepalings van paragraaf (a) van artikel 34 (4) van die Wet ten opsigte van die eiendomme omskryf as "Hoewes 7472 (grootte 3 240 vierkante meter), 7 473 (grootte 3 240 vierkante meter), 7 474 (grootte 3 240 vierkante meter), 7 475 (grootte 3 524 vierkante meter) en 7 476 (grootte 2 957 vierkante meter), Pinetown-uitbreiding 72, Westmead, Munisipaliteit van PINETOWN, en in die Port Natal–Ebohdwe Gesamentlike Dienste Raad Area, administratiewe distrik van Natal", tot die mate dat die eiendomme soos hierbo omskryf, ingesluit mag word as bates van Sixway Properties (Edms.) Beperk wie se aandele besit word deur die genoemde Property Fund Managers Beperk.

P. J. BADENHORST,
Registrateur van Effekte-trustmaatskappye.
(3 Desember 1993)

NOTICE 1190 OF 1993

FINANCIAL SERVICES BOARD

UNIT TRUSTS CONTROL ACT, No. 54 OF 1981

EXEMPTION FROM THE PROVISIONS OF SECTION 34 (4) (a)

I, Petrus Johannes Badenhorst, Registrar of Unit Trust Companies, hereby, after consultation with the Unit Trusts Advisory Committee, under section 45 of the Unit Trusts Control Act, 1981 (Act No. 54 of 1981), exempt Property Fund Managers Limited, a registered management company of a unit trust scheme in property shares in terms of the said Act, from the provisions of paragraph (a) of section 34 (4) of the Act in respect of the properties described as "Stands 7472 (measuring 3 240 square metres), 7 473 (measuring 3 240 square metres), 7 474 (measuring 3 240 square metres), 7 475 (measuring 3 524 square metres) and 7 476 (measuring 2 957 square metres), Pinetown Extension 72, Westmead, situate in the Borough of PINETOWN and in the Port Natal–Ebohdwe Joint Services Board Area, Administrative District of Natal", to the extent that the properties as described above, may be included as assets of Sixway Properties (Pty) Limited, whose shares are owned by the said Property Fund Managers Limited.

P. J. BADENHORST,
Registrar of Unit Trust Companies.
(3 December 1993)

KENNISGEWING 1191 VAN 1993 • NOTICE 1191 OF 1993

VOORLOPIGE OPGAWE VAN HANDELSTATISTIEK VAN DIE REPUBLIEK VAN SUID-AFRIKA VRYGESTEL DEUR DIE KOMMISSARIS VAN DOEANE EN AKSYNS

PRELIMINARY STATEMENT OF TRADE STATISTICS OF THE REPUBLIC OF SOUTH AFRICA RELEASED BY THE COMMISSIONER FOR CUSTOMS AND EXCISE

Opmerking: Die in- en uitvoersyfers wat in hierdie opgawe verskyn is grootliks aangepas om dit in ooreenstemming te bring met die vereistes wat gestel word vir die opstel van die betalingsbalans.

Die ondervermelde syfers omsluit die totale buitelandse handelstatistiek van die gemeenskaplike doeanegebied van die Republiek van Suid-Afrika, Botswana, Lesotho, Swaziland, Namibië asook van Transkei, Bophuthatswana, Venda en Ciskei.

L.W.: Die oorskakeling na die Geharmonieerde Tariefstelsel met ingang van 1 Januarie 1988 het die indeling van sekere kommoditeite verander. Wanneer die afdelingstotale vir 1988 en later jare dus met dié van vorige jare vergelyk word, moet die moontlike verskille as gevolg van die oorskakeling nie uit die oog verloor word nie.

Remark: The import and export figures reflected in this statement have been adjusted largely to bring them into line with the requirements for the compilation of the balance of payments.

The undermentioned data entails the total foreign trade statistics of the common customs area of the Republic of South Africa, Botswana, Lesotho, Swaziland, Namibia as well as Transkei, Bophuthatswana, Venda and Ciskei.

N.B.: The change-over to the Harmonized Tariff System with effect from 1 January 1988, altered the classification of certain commodities. When comparing the section totals for 1988 and later years with those of previous years the possible differences due to the change-over should therefore be taken into consideration.

TYDPERK: JANUARIE TOT OKTOBER 1993 • PERIOD: JANUARY TO OCTOBER 1993

TABEL A: TOTALE IN MILJOENE RAND VOLGENS WÊRELDSTREKE EN SKEEPS- EN VLIEGTUIGVOORRADE
TABLE A: TOTALS IN MILLIONS OF RAND ACCORDING TO WORLD ZONES AND SHIPS' AND AIRCRAFT STORES

Wêreldstreke—World Zones	Invoere—Imports		Uitvoere—Exports	
	1993	1992	1993	1992
Afrika—Africa	1 394,7	1 103,4	5 403,3	4 966,1
Europa—Europe	21 846,8	19 549,2	22 033,0	19 322,9
Amerika—America	7 933,5	7 289,9	5 711,2	5 039,0
Asië—Asia	13 017,1	9 898,2	11 336,4	10 054,8
Oseanië—Oceania	612,9	536,6	489,5	317,8
Ander ongeklassifiseerde goedere en betalingsbalansaansuiwerings Other unclassified goods and balance of payments adjustments	4 709,8	5 530,5	19 956,9	15 887,2
Skeeps-/vliegtuigvoorrade—Ships'/Aircraft Stores	—	—	944,0	578,8
GROOTTOTAAL—GRAND TOTAL	49 514,8	43 907,8	65 874,3	56 166,6

TABEL B: TOTALE IN MILJOEN RAND VOLGENS AFDELINGS VAN DIE GEHARMONIEERDE STELSEL
TABLE B: TOTALS IN MILLION RAND ACCORDING TO SECTIONS OF THE HARMONIZED SYSTEM

Afdelings—Sections	Invoere—Imports		Uitvoere—Exports	
	1993	1992	1993	1992
I. Lewende diere; dierlike produkte Live animals; animal products	325,4	344,1	799,6	667,4
II. Plantaardige produkte Vegetable products	1 721,5	1 981,1	2 298,7	2 154,5
III. Dierlike of plantaardige vette en olies en splitsprodukte; voorbereide spysvette; dierlike en plantaardige wasse Animal or vegetable fats and oils and their cleavage products; prepared edible fats; animal and vegetable waxes	416,6	406,7	140,4	121,4
IV. Voorbereide voedsel; dranke, spiritus en asyn; tabak en vervaardigde tabaksurrogate Prepared foodstuffs; beverages, spiritus and vinegar; tobacco and manufactured tobacco substitutes	823,0	962,3	1 459,2	1 599,4
V. Mineraalprodukte Mineral products	453,2	454,3	6 555,2	5 840,9
VI. Produkte van die chemiese of verwante nywerhede Products of the chemical or allied industries	5 562,6	4 828,7	2 674,6	2 756,3
VII. Plastieke en artikels daarvan; rubber en artikels daarvan Plastics and articles thereof; rubber and articles thereof	2 226,3	1 918,7	607,6	612,6
VIII. Ongelooide huide en velle, leer, pelsvelle en artikels daarvan; saal- en tuiemakersware; reisartikels, handsakke en dergelike houers; artikels van diererderm (uitgesonderd sywurmsnaar) Raw hides and skins, leather, furskins and articles thereof; saddlery and harness; travel goods handbags and similar containers; articles of animal gut (other than silk-worm gut)	237,8	224,6	435,6	334,8
IX. Hout en artikels van hout; houtskool; kurk en artikels van kurk; fabrikate van strooi, van esparto of van ander vlegwerkstowwe; mandjiewerk en vlegwerk Wood and articles of wood; wood charcoal; cork and articles of cork; manufacturers of straw; of esparto or of other plaiting materials; basketware and wickerwork	436,6	345,0	514,4	307,1
X. Pulp van hout of van ander veselagtige sellulosiese stof; afval en oorskiet van papier of papierbord; papier en papierbord en artikels daarvan Pulp of wood or of other fibrous cellulosic material; waste and scrap of paper or paperboard; paper and paperboard of paper or paperboard; paper and paperboard and articles thereof	1 436,4	1 224,1	1 578,8	1 577,6
XI. Tekstiele en tekstielartikels Textiles and textile articles	2 267,8	2 059,2	1 490,7	1 516,2

Afdelings—Sections	Invoere—Imports		Uitvoere—Exports	
	1993	1992	1993	1992
XII. Skoeisel, hoofdeksels, sambrele, sonsambrele, wandelstokke, sitstokke, swepe, karwatse en onderdele daarvan; bereide vere en artikels daarvan gemaak; kunsblomme; artikels van mensehaar Footwear, headgear, umbrellas, sun umbrellas, walking-sticks, seat-sticks, whips, riding-crops and parts thereof; prepared feathers and articles made therewith; artificial flowers; articles of human hair	371,6	262,0	54,9	43,2
XIII. Artikels van klip, gips, sement, asbes, mika of dergelike stowwe; keramiese produkte; glas en glasware Articles of stone, plaster, cement, asbestos, mica or similar materials; ceramic products; glass and glassware	657,7	590,8	337,1	293,8
XIV. Natuurlike of gekweekte pêrels, edel- of halfedelstene, edelmetaal, metale met edelmetale bedek, en artikels daarvan; nagemaakte juweliersware; muntstukke Natural or cultured pearls, precious or semi-precious stones, precious metals, metals clad with precious metal and articles thereof; imitation jewellery; coin	1 290,9	275,8	8 815,2	5 939,8
XV. Onedelmetale en artikels van onedelmetaal Base metals and articles of base metal	2 169,1	2 105,7	8 279,8	7 859,5
XVI. Masjinerie en meganiese toestelle; elektriese toerusting; onderdele daarvan; klankopnemers en -weergewers; televisie- beeld- en klankopnemers en -weergewers, en onderdele en bybehoorsels van sodanige artikels Machinery and mechanical appliances; electrical equipment; parts thereof; sound recorders and reproducers, television image and sound recorders and reproducers, and parts and accessories of such articles	14 096,3	12 394,9	2 239,3	1 716,6
XVII. Voertuie, lugvaartuie, vaartuie en verwante vervoertoerusting Vehicles, aircraft, vessels and associated transport equipment	7 572,6	5 733,8	2 133,7	1 934,7
XVIII. Optiese, fotografiese, kinematografiese, meet-, kontroleer-, presisie-, mediese of chirurgiese instrumente en apparate; uurwerke en horlosies; musiekinstrumente; onderdele en bybehoorsels daarvan Optical, photographic, cinematographic, measuring, checking, precision, medical or surgical instruments and apparatus; clocks and watches; musical instruments, parts and accessories thereof	2 177,1	1 818,6	186,7	136,6
XX. Diverse vervaardigde artikels Miscellaneous manufactured articles	583,2	532,1	300,0	243,8
XXI. Kunswerke, versamelaarsstukke, en antieke Works of art, collectors' pieces and antiques	33,9	22,3	20,4	14,7
Ander ongeklassifiseerde goedere en betalingsbalansaansuiwerings Other unclassified goods and balance of payments adjustments	4 655,2	5 423,0	24 952,4	20 495,7
GROOTTOTAAL—GRAND TOTAL	49 514,8	43 907,8	65 874,3	56 166,6

(3 Desember 1993/3 December 1993)

KENNISGEWING 1192 VAN 1993

SENTRALE STATISTIEKDIENS

DIE HOOF: SENTRALE STATISTIEKDIENS maak vir algemene inligting bekend dat die Verbruikersprysindeks soos volg is:

Verbruikersprysindeks, alle items (Basis 1990 = 100)

Oktober 1993 = 147,9.

(3 Desember 1993)

NOTICE 1192 OF 1993

CENTRAL STATISTICAL SERVICE

THE HEAD: CENTRAL STATISTICAL SERVICE notifies for general information that the Consumer Price Index is as follows:

Consumer Price Index, all items (Base 1990 = 100)

October 1993 = 147,9.

(3 December 1993)

KENNISGEWING 1193 VAN 1993

NATALSE PROVINSIALE ADMINISTRASIE

AANSTELLING: HOSPITAALRAAD: MIDDELLANDSE HOSPITAAL, PIETERMARITZBURG

Die Administrateur van die provinsie Natal het kragtens die bevoegdheid hom verleen by artikel 47 van die Wet op Geestesgesondheid, 1973 (Wet No. 18 van 1973), ondergenoemde persone as lede van die Hospitaalraad, Middellandse Hospitaal, Pietermaritzburg, aangestel vir 'n tydperk van drie jaar met ingang van 1 November 1993:

Mnr. B. D. Will (Voorsitter).

Dr. N. T. Karnezos.

Dr. E. P. Ndaba.

Mnr. W. J. van Rooyen.

(3 Desember 1993)

KENNISGEWING 1194 VAN 1993

NATALSE PROVINSIALE ADMINISTRASIE

AANSTELLING: HOSPITAALRAAD: KONING GEORGE V-HOSPITAAL, DURBAN

Die Administrateur van die provinsie Natal het kragtens die bevoegdheid hom verleen by artikel 47 van die Wet op Geestesgesondheid, 1973 (Wet No. 18 van 1973), ondergenoemde persone as lede van die Hospitaalraad, Koning George V-hospitaal, Durban, aangestel vir 'n tydperk van drie jaar met ingang van 1 November 1993:

Mnr. S. K. Sing (Voorsitter).

Mnr. S. B. Garach.

Mnr. C. K. van Eck.

(3 Desember 1993)

KENNISGEWING 1195 VAN 1993

DEPARTEMENT VAN VERVOER

WET OP DIE LISENSIËRING VAN LUGDIENSTE, 1990 (WET No. 115 VAN 1990)

Hierby word ingevolge die bepalings van artikel 15 (1) (b) van Wet 115 van 1990 en regulasie 8 van die Regulasies vir Binnelandse Lugdienste, 1991, vir algemene inligting bekendgemaak dat die Lugdienslisensiëringsraad die aansoeke waarvan besonderhede in die Bylaes hieronder verskyn, sal oorweeg.

Vertoë ingevolge artikel 15 (3) van Wet 115 van 1990 ter ondersteuning of bestryding van 'n aansoek moet die Lugdienslisensiëringsraad, Privaat Sak X193, Pretoria, 0001, binne 21 dae na die datum van publikasie hiervan bereik.

BYLAE 1

AANSOEKE OM DIE TOESTAAN VAN LISENSIES

(A) Volle naam en handelsnaam van aansoeker. (B) Volle besigheids- of woonadres van aansoeker. (C) Klas lisensie waarom aansoek gedoen word. (D) Tipe lugdiens waarop aansoek betrekking het. (E) Kategorie lugvaartuig waarop aansoek betrekking het.

NOTICE 1193 OF 1993

NATAL PROVINCIAL ADMINISTRATION

APPOINTMENT: HOSPITAL BOARD: MIDLANDS HOSPITAL, PIETERMARITZBURG

The Administrator of the Province of Natal has under the powers vested in him by section 47 of the Mental Health Act, 1973 (Act No. 18 of 1973), appointed the following persons to be members of the Hospital Board, Midlands Hospital, Pietermaritzburg, for a three year period with effect from 1 November 1993:

Mr B. D. Will (Chairman).

Dr N. T. Karnezos.

Dr E. P. Ndaba.

Mr W. J. van Rooyen.

(3 December 1993)

NOTICE 1194 OF 1993

NATAL PROVINCIAL ADMINISTRATION

APPOINTMENT: HOSPITAL BOARD: KING GEORGE V HOSPITAL, DURBAN

The Administrator of the Province of Natal has under the powers vested in him by section 47 of the Mental Health Act, 1973 (Act No. 18 of 1973), appointed the following persons to be members of the Hospital Board, King George V Hospital, Durban, for a three year period with effect from 1 November 1993:

Mr S. K. Sing (Chairman).

Mr S. B. Garach.

Mr C. K. van Eck.

(3 December 1993)

NOTICE 1195 OF 1993

DEPARTMENT OF TRANSPORT

AIR SERVICE LICENSING ACT, 1990 (ACT No. 115 OF 1990)

Pursuant to the provisions of section 15 (1) (b) of Act 115 of 1990 and regulation 8 of the Domestic Air Services Regulations, 1991, it is hereby notified for general information that the application(s) details of which appear in the Schedules hereto, will be considered by the Air Service Licensing Council.

Representations in accordance with section 15 (3) of Act 115 of 1990 in support of, or in opposition to, an application, should reach the Air Service Licensing Council, Private Bag X193, Pretoria, 0001, within 21 days of the date of publication hereof.

SCHEDULE 1

APPLICATIONS FOR THE GRANT OF LICENCES

(A) Full name and trade name of applicant. (B) Full business or residential address of applicant. (C) Class of licence applied for. (D) Type of air service to which application applies. (E) Category of aircraft to which application applies.

(A) Berg Flights BK. (B) Posbus 1033, Uvongo, 4270. (C) Klas III. (D) Tipe G9. (E) Kategorie A4.

(A) Wes-Transvaal Vliegskool BK. (B) Posbus 1539, Klerksdorp, 2570. (C) Klas II. (D) Tipe N1. (E) Kategorie A4.

BYLAE 2

AANSOEK OM DIE WYSIGING VAN LISENSIE

(A) Volle naam en handelsnaam van aansoeker. (B) Volle besigheids- of woonadres van aansoeker. (C) Klas lisensie ten opsigte waarvan 'n wysiging gevra word. (D) Tipe lugdiens en die wysiging daarvan waarom aansoek gedoen word. (E) Kategorie lugvaartuig en die wysiging daarvan waarom aansoek gedoen word.

(A) Atair (Edms.) Bpk. (B) Posbus 169, Lanseria, 1748. (C) Klas II. (D) Tipe N1 en N2. (E) Kategorie A2, A3 en A4, voeg by A1, H1 en H2.

(3 Desember 1993)

KENNISGEWING 1196 VAN 1993

DEPARTEMENT VAN VERVOER

WET OP INTERNASIONALE LUGDIENSTE, 1949
(WET No. 51 VAN 1949)

Hierby word ingevolge die bepalings van artikels 5 (a) en (b) van Wet No. 51 van 1949 en regulasie 5 van die Regulasies vir Burgerlugdienste, 1964, vir algemene inligting bekendgemaak dat die Raad op Internasionale Lugdienste die aansoeke waarvan besonderhede in die Bylaes hieronder verskyn, sal aanhoor.

Vertoë ingevolge artikel 6 (1) van Wet No. 51 van 1949 ter ondersteuning of bestryding van 'n aansoek moet die Voorsitter van die Raad op Internasionale Lugdienste, Privaatsak X193, Pretoria, 0001, en die aansoeker binne 21 dae na die datum van publikasie hiervan bereik en daarin moet gemeld word of die persoon of persone wat aldus vertoë rig, van plan is om die verrigtinge by te woon of om daar verteenwoordig te word.

Die Raad op Internasionale Lugdienste sal reël dat kennis van die datum, tyd en plek van die verrigtinge skriftelik gegee word aan die aansoeker en al die persone wat aldus vertoë gerig het en wat verlang om aldus teenwoordig of verteenwoordig te wees.

BYLAE B

LYS VAN AANSOEKE OM DIE HERNUWING VAN LISENSIES

(A) Naam en adres van applikant. (B) Naam waaronder die lugdiens geëksploiteer word. (C) Soort lugdiens ten opsigte waarvan hernuwing aangevra word en die nommer en datum van bestaande lisensie. (D) Besonderhede van lisensie. (i) Gebied wat bedien gaan word. (ii) Roete(s) en frekwensie(s) wat bedien gaan word. (iii) Uitgangsbasis(se). (iv) Soort verkeer wat vervoer gaan word. (v) Soort opleiding wat verskaf gaan word. (vi) Soort werk wat onderneem gaan word. (vii) Tariefskaal. (E) Lugvaartuie wat gebruik gaan word.

(A) Berg Flights CC. (B) P.O. Box 1033, Uvongo, 4270. (C) Clas III. (D) Type G9. (E) Category A4.

(A) Western Transvaal Flying School CC. (B) P.O. Box 1539, Klerksdorp, 2570. (C) Class II. (D) Type N1. (E) Category A4.

SCHEDULE 2

APPLICATION FOR THE AMENDMENT OF LICENCE

(A) Full name and trade name of applicant. (B) Full business or residential address of applicant. (C) The class of licence in respect of which the amendment is sought. (D) Type of air service and the amendment thereto which is being applied for. (E) Category of aircraft and the amendment thereto which is being applied for.

(A) Atair (Pty) Ltd. (B) P.O. Box 169, Lanseria, 1748. (C) Class II. (D) Type N1 and N2. (E) Category A2, A3 and A4, add A1, H1 and H2.

(3 December 1993)

NOTICE 1196 OF 1993

DEPARTMENT OF TRANSPORT

INTERNATIONAL AIR SERVICES ACT, 1949
(ACT No. 51 OF 1949)

Pursuant to the provisions of sections 5 (a) and (b) of Act No. 51 of 1949 and regulation 5 of the Civil Air Services Regulations, 1964, it is hereby notified for general information that the applications, details of which appear in the Schedules hereto, will be heard by the International Air Service Council.

Representations in accordance with section 6 (1) of Act No. 51 of 1949 in support of, or in opposition to, an application, should reach the Chairman of the International Air Service Council, Private Bag X193, Pretoria, 0001, and the applicant within 21 days of the date of publication hereof, stating whether the party or parties making such representation intend to be present or represented at the hearing.

The International Air Service Council will cause notice of the time, date and place of the hearing to be given in writing to the applicant and all parties who have made representations as aforesaid and who desire to be present or represented at the hearing.

SCHEDULE B

LIST OF APPLICATIONS FOR RENEWAL OF LICENCES

(A) Name and address of applicant. (B) Name under which the air service is being operated. (C) Class of air service in respect of which renewal is sought and number and date of existing licence. (D) Particulars of licence. (i) Area to be served. (ii) Route(s) and frequencies to be served. (iii) Base(s). (iv) Types and classes of traffic to be conveyed. (v) Types of training to be provided. (vi) Types of work to be undertaken. (vii) Tariff of charges. (E) Aircraft to be used.

(A) African International Airways (Edms.) Bpk., Posbus 2332, Kempton Park, 1620. (B) African International Airways. (C) Nie-vasgesteldelugvervoerdienslisensie N468 gedateer 28 November 1991. (D) (i) Wêreldwyd, met uitsondering van lugvervoerdienste binne die grense van die Republiek van Suid-Afrika. (iii) Jan Smutslughawe. (iv) Slegs vrag. Spesiale voorwaardes: Tensy die Raad op Internasionale Lugdienste anders besluit, mag die lugdiens slegs bedryf word op roetes tussen die Republiek van Suid-Afrika en lande wat nie deur enige vasgestelde lugdienste bedien word nie. (viii) Vasgestelde koste teen 14,30 sent per ton/km. (E) Twee Douglas DC-8 54F-lugvaartuie.

Mits sodanige lugvaartuig ZS-geregistreer en B-gekategoriseer is.

BYLAE D

LYS VAN AANSOEKE OM DIE VERANDERING OF WYSIGING VAN LISENSIES

(A) Naam en adres van applikant. (B) Naam waaronder die lugdiens geëksploiteer word. (C) Besonderhede betreffende die lisensie en die verandering of wysiging daarvan of die voorwaardes daarvan ten opsigte waarvan aansoek gedoen is.

(A) Transnet (Edms.) Bpk., Posbus 7778, Johannesburg, 2000. (B) Suid-Afrikaanse Lugdiens. (C) Vasgesteldelugvervoerdienslisensies S173 en S704. Nie-vasgesteldelugvervoerdienslisensie N115. Onder "Lugvaartuie wat gebruik gaan word", voeg by: "Airbus A320-231 ZS-SHG en Boeing 747-ZS-SAY".

(3 Desember 1993)

(A) African International Airways (Pty) Ltd, P.O. Box 2332, Kempton Park, 1620. (B) African International Airways. (C) Non-scheduled Air Transport Service Licence N468 dated 28 November 1991. (D) (i) Worldwide with the exclusion of all domestic carriage within the boundaries of the Republic of South Africa. (iii) Jan Smuts Airport. (iv) Cargo only. Special conditions: Unless the International Air Service Council otherwise approves, the air service may only be operated between the Republic of South Africa and countries not served by any scheduled air services. (viii) Fixed costs at 14,3 cents per ton/km. (E) Two Douglas DC-8 54F aircraft.

Provided such aircraft is ZS-registered and categorised B.

SCHEDULE D

LIST OF APPLICATIONS FOR THE ALTERATION, MODIFICATION OR AMENDMENT TO LICENCES

(A) Name and address of applicant. (B) Name under which the air service is operated. (C) Particulars of the licence and of the alteration, modification or amendment thereto or the conditions thereof which has been applied for.

(A) Transnet (Pty) Ltd, P.O. Box 7778, Johannesburg, 2000. (B) South African Airways. (C) Scheduled Air Transport Service Licences S173 and S704. Non-scheduled Air Transport Service Licence N115. Under "Aircraft to be used", add: "Airbus A320-231 ZS-SHG and Boeing 747-ZS-SAY".

(3 December 1993)

KENNISGEWING 1197 VAN 1993

DEPARTEMENT VAN NASIONALE GESONDHEID EN BEVOLKINGSONTWIKKELING

WET OP BEHEER VAN MEDISYNE EN VERWANTE STOWWE (WET No. 101 VAN 1965)

INTREKKING VAN DIE REGISTRASIE VAN MEDISYNE

Hierby word ingevolge artikel 17 van die Wet op die Beheer van Medisyne en Verwante Stowwe, 1965 (Wet No. 101 van 1965), bekendgemaak dat die Registrateur van Medisyne, met die goedkeuring van die Medisynebeheerraad ingestel by artikel 2 van genoemde Wet, die registrasie van die volgende medisyne ingetrek het:

NOTICE 1197 OF 1993

DEPARTMENT OF NATIONAL HEALTH AND POPULATION DEVELOPMENT

MEDICINES AND RELATED SUBSTANCES CONTROL ACT (ACT No. 101 OF 1965)

CANCELLATION OF THE REGISTRATION OF MEDICINES

It is hereby notified, in terms of section 17 of the Medicines and Related Substances Control Act, 1965 (Act No. 101 of 1965), that the Registrar of Medicines, with the approval of the Medicines Control Council established by section 2 of the said Act, has cancelled the registration of the following medicines:

Nommer Number	Naam van produk Name of product	Applikant Applicant	Datum Date
U/30.1/78	Diplovax HDC	African Sub-Equatorial Pharmaceuticals	93-07-13
Y/30.1/247	Diplovax HDC 4.5	African Sub-Equatorial Pharmaceuticals	93-07-13
Y/30.1/248	Diplovax HDC 5.0	African Sub-Equatorial Pharmaceuticals	93-07-13
Z/30.1/220	Diplovax HDC 4.7	African Sub-Equatorial Pharmaceuticals	93-07-13
U/34/97	Isosol Non-Presserved Saline Solution	Allergan Pharmaceuticals (Pty) Ltd	93-06-21
K/11.2/179	Duspatalin	Berlimed (Pty) Ltd	93-09-02
G/3.1/187	Brufen Suppositories	Boots Pharmaceuticals (Pty) Ltd	93-11-05
J/5.4.1/43	Antadine Syrup	Boots Pharmaceuticals (Pty) Ltd	93-10-25
P/7.1.3/86	Tridil 5 mg	Boots Pharmaceuticals (Pty) Ltd	93-11-05
Q/2.7/356	Nubain 10 mg/mℓ	Boots Pharmaceuticals (Pty) Ltd	93-11-05
M/34/249	Chendol Capsules	Fisons Pharmaceuticals (Pty) Ltd	93-06-16
P/8.2/73	Hepsal	Fisons Pharmaceuticals (Pty) Ltd	93-06-16

Nommer Number	Naam van produk Name of product	Applikant Applicant	Datum Date
P/8.2/158	Uniparin	Fisons Pharmaceuticals (Pty) Ltd	93-06-16
Q/34/4	Chendol Tablets	Fisons Pharmaceuticals (Pty) Ltd	93-06-16
Q/8.2/11	Uniparin-CA	Fisons Pharmaceuticals (Pty) Ltd	93-06-16
H/20.2.1/8	Enterocura	Noristan Limited	93-11-11
C/10.1/222	Formula 44 Cough Silencers	Permark International (Pty) Ltd	93-06-18
G/13.12/1417	Skintone Clearasil Pimple and Acne Cream	Permark International (Pty) Ltd	93-06-18
G/13.12/1418	Colourless Clearasil Pimple and Acne Cream	Permark International (Pty) Ltd	93-06-18
K/10.1/301	Vicks Soothing Cough Drops - Lemon Flavour	Permark International (Pty) Ltd	93-06-18
L/13.12/292	Mytolac	Permark International (Pty) Ltd	93-06-18
Q/13.4.1/207	Cortrex Cream	Permark International (Pty) Ltd	93-06-18
K/7.1.4/100	Nitrong 6,5	Rhone-Poulenc Rorer SA (Pty) Ltd	93-08-17
E/13.4.1/238	Purantix Ointment	Sandoz Products (Pty) Ltd	93-11-04
E/13.4.1/239	Purantix Cream	Sandoz Products (Pty) Ltd	93-11-04
F/2.9/125	Fiorinal-P	Sandoz Products (Pty) Ltd	93-11-04
G/5.8/25	Rhinergal Tablets	Sandoz Products (Pty) Ltd	93-11-04
H/22.1.4/175	Vi-De 3 Ampoules	Sandoz Products (Pty) Ltd	93-11-04
J/22.1.4/168	Vi-de 3 Capsules	Sandoz Products (Pty) Ltd	93-11-04
J/3.1/422	Biarison 300 mg Capsules	Sandoz Products (Pty) Limited	93-11-04
J/6.3/423	Digoxin Tablets	Sandoz Products (Pty) Ltd	93-11-04
J/6.3/424	Digoxin Injection	Sandoz Products (Pty) Ltd	93-11-04
N/20.1.1/19	Biophenical Injection	Sandoz Products (Pty) Ltd	93-11-04
Q/3.1/60	Biarison 200 mg Capsules	Sandoz Products (Pty) Ltd	93-11-04
J/2.2/319	Halcion 0,5 mg	Upjohn (Pty) Ltd	93-10-14

(3 Desember 1993)/(3 December 1993)

KENNISGEWING 1198 VAN 1993

DEPARTEMENT VAN LANDBOU

KENNISGEWING VAN VERGADERING VAN SKULDEISERS KRAGTENS ARTIKEL 22 (1) VAN DIE WET OP LANDBOUKREDIET, 1966

Hierby word 'n vergadering van ondergenoemde applikante en hulle skuldeisers op die plek en datum hieronder genoem, belê, met die doel om skuldeisers in staat te stel om hul vorderings teen die applikante te bewys en 'n skikkingsvoorstel van die Landboukredietraad te oorweeg.

J. H. SMIT,
Direkteur: Direktoraat Finansiële Bystand,
Departement van Landbou.

NOTICE 1198 OF 1993

DEPARTMENT OF AGRICULTURE

NOTICE OF MEETING OF CREDITORS IN TERMS OF SECTION 22 (1) OF THE AGRICULTURAL CREDIT ACT, 1966

A meeting of the undermentioned applicants and their creditors is hereby convened at the place and date mentioned hereunder for the purpose of enabling creditors to prove their claims against the applicants and of considering a proposal for a compromise by the Agricultural Credit Board.

J. H. SMIT,
Director: Directorate Financial Assistance,
Department of Agriculture.

Aansoek van Application by	Plek van byeenkoms Place of meeting	Datum en tyd Date and time
Ernst Jacobus Viljoen (Id. No. 380301 5013 00 0), en/and **Paul Philippus Viljoen (Id. No. 660821 5006 08 4)** van die plaas/of the farm Klipfontein; Posbus/P.O. Box 179, Colesberg, 5980	Kantoor van die Landdros/Magistrate's Office, Colesberg	28 Januarie/January 1994 om/at 09:00.

(3 Desember 1993)/(3 December 1993)

KENNISGEWING 1199 VAN 1993

DEPARTEMENT VAN LANDBOU

KENNISGEWING VAN VERGADERING VAN SKULDEISERS KRAGTENS ARTIKEL 22 (1) VAN DIE WET OP LANDBOUKREDIET, 1966

Hierby word 'n vergadering van ondergenoemde applikant en sy skuldeisers op die plek en datum hieronder genoem, belê, met die doel om skuldeisers in staat te stel om hul vorderings teen die applikant te bewys en 'n skikkingsvoorstel van die Landboukredietraad te oorweeg.

J. H. SMIT,
Direkteur: Direktoraat Finansiële Bystand,
Departement van Landbou.

NOTICE 1199 OF 1993

DEPARTMENT OF AGRICULTURE

NOTICE OF MEETING OF CREDITORS IN TERMS OF SECTION 22 (1) OF THE AGRICULTURAL CREDIT ACT, 1966

A meeting of the undermentioned applicant and his creditors is hereby convened at the place and date mentioned hereunder for the purpose of enabling creditors to prove their claims against the applicant and of considering a proposal for a compromise by the Agricultural Credit Board.

J. H. SMIT,
Director: Directorate Financial Assistance,
Department of Agriculture.

Aansoek van Application by	Plek van byeenkoms Place of meeting	Datum en tyd Date and time
Albertus Johannes de Klerk (Id. No. 490210 5016 08 6), van die plaas/of the farm Blesbokpan; Posbus/P.O. Box 23, Theunissen, 9410	Kantoor van die Landdros/Magistrate's Office, Theunissen	14 Januarie/January 1994 om/at 09:00.

(3 Desember 1993)/(3 December 1993)

KENNISGEWING 1200 VAN 1993

KANTOOR VIR OPENBARE ONDERNEMINGS EN PRIVATISERING

RAAD OP MEDEDINGING

GEVOLG VAN ONDERSOEK NA BEPERKENDE PRAKTYKE BY DIE VERVAARDIGING EN BEMARKING VAN PAPIER EN PAPIERPRODUKTE INSLUITENDE VERPAKKINGSMATERIAAL

Die Raad op Mededinging maak hierby vir algemene inligting bekend dat die ondersoek na bogemelde aangeleentheid, wat in Kennisgewing 110 in *Staatskoerant* No. 12291 van 16 Februarie 1990 afgekondig is, afgehandel is.

Die ondersoek, wat in opdrag van die destydse Minister van Administrasie en Privatisering onderneem is, het voortgespruit uit klagtes van beweerde beperkende praktyke in die bedryfstak. Dit het ook verband gehou met die prysvasstelling van papier en pulp na of gelyk aan die pryse van ingevoerde produkte (die sogenaamde invoerpariteitprysvasstelling).

Die Raad het tot die gevolgtrekking gekom dat—

(a) alhoewel prysnavolging endemies is in die bedryfstak, dit nie geag kan word horisontale samespanning te wees soos omskryf in Goewermentskennisgewing No. 801 in *Staatskoerant* No. 10211 van 2 Mei 1986; en

(b) invoerpariteitsprysvasstelling nie *per se* 'n beperkende praktyk soos omskryf in artikel 1 van die Wet op die Handhawing en Bevordering van Mededinging, 1979, daarstel nie.

Ofskoon klagtes aan die Raad voorgelê is dat beperkende praktyke (horisontale samespanning oor pryse en voorwaardes) op sommige vlakke in die bedryfstak plaasvind, kon daar geen voldoende bewyse daarvoor gevind word nie.

Geen optrede deur die Minister vir Openbare Ondernemings is derhalwe nodig nie en die Raad het hom dienooreenkomstig ingelig.

(3 Desember 1993)

NOTICE 1200 OF 1993

OFFICE FOR PUBLIC ENTERPRISES AND PRIVATISATION

COMPETITION BOARD

RESULT OF AN INVESTIGATION INTO RESTRICTIVE PRACTICES IN THE MANUFACTURE AND MARKETING OF PAPER AND PAPER PRODUCTS INCLUDING PACKAGING MATERIAL

The Competition Board hereby makes known for general information that the investigation into the above-mentioned matter, published in Notice 110 in *Government Gazette* No. 12291 of 16 February 1990, has been completed.

The investigation, which was conducted on the instruction of the then Minister for Administration and Privatisation, was undertaken as a result of complaints of alleged restrictive practices in this industry. It also had relevance to the pricing of paper and pulp close or equal to the prices of imported products (the so-called import parity pricing).

The Board concluded that—

(a) although price leadership is endemic in this industry, it could not be regarded as horizontal collusion as defined in Government Notice No. 801 in *Government Gazette* No. 10211 of 2 May 1986; and

(b) import parity pricing is not *per se* a restrictive practice as defined in section 1 of the Maintenance and Promotion of Competition Act, 1979.

Although complaints that restrictive practices (horizontal collusion on prices and conditions) were taking place at certain levels in the industry were laid before the Board, sufficient evidence could not be found to substantiate the complaints.

No action is therefore necessary by the Minister for Public Enterprises and the Board advised him accordingly.

(3 December 1993)

KENNISGEWING 1201 VAN 1993

RAAD OP TARIEWE EN HANDEL

DOEANE- EN AKSYNSTARIEFAANSOEKE: LYS 42/93

Onderstaande aansoeke betreffende die Doeane-en Aksynstarief is deur die Raad op Tariewe en Handel ontvang. Enige beswaar teen of kommentaar op hierdie vertoë moet binne ses weke na die datum van hierdie kennisgewing aan die Voorsitter, Raad op Tariewe en Handel, Privaat Sak X753, Pretoria, 0001, gerig word. Die aandag word daarop gevestig dat die skale van reg wat in die aansoeke genoem word, dié is wat deur die applikante aangevra is en dat die Raad, afhangende van sy bevindinge, hoër of laer skale van reg mag aanbeveel.

Verlaging van die reg op:

Veselvliesstowwe, hetsy geïmpregneer, bestryk, bedek of gelamelleer al dan nie, indeelbaar by tariefpos 56.03, van verskeie skale van reg tot vry.

[RTH-verw. T5/2/11/4/1 (930375) (Me. H. Claassens)]

Applikant:

(i) Raad op Tariewe en Handel, Privaat Sak X753, Pretoria, 0001.

(ii) Vitamed (Pty) Ltd, Posbus 27183, Benrose, 2011.

Korting van die reg (Bylae 3) op:

1. Plate, velle, film, foelie en reep, van polimere van vinielchloried, met 'n dikte van meer as 0,75 mm maar hoogstens 3 mm, indeelbaar by tariefsubpos 3921.12.50, vir die vervaardiging van handsakke.

[RTH-verw. T5/2/8/3/1 (930383) (Mnr. H. P. le Roux)]

Applikant:

Busy Bag and Belt (Pty) Ltd, Posbus 795, Stanger, 4450.

2. Poliëterpoliole, wat twee of meer hidroksielgroepe bevat, vloeistowwe of pastas, met 'n hidrioksielgetal van minstens 20 mg KOH/g of meer maar hoogstens 800 mg KOH/g, indeelbaar by tariefsubpos 3907.20.10, synde gliserien slegs met etileenoksiede volledig gereageer vir die vervaardiging van poliëterpoliole.

[RTH-verw. T5/2/7/21 (930404) (Mnr. G. Bester)]

Applikant:

Polyol Chemicals (Edms.) Bpk., Posbus 246, Umbogintwini, 4120.

3. Poliëterpoliole, wat 2 of meer hidroksielgroepe bevat, vloeistowwe of pastas, met 'n hidroksielgetal van minstens 20 mg KOH/g of meer maar hoogstens 800 mg KOH/g synde poliëterpoliole, gesluit met etileenoksiede met 'n primêre hidroksielinhoud van minstens 20%, indeelbaar by tariefsubpos 3907.20.10 vir die vervaardiging van polioolmengsels.

[RTH-verw. T5/2/7/2/1 (930403) (Mnr. G. Bester)]

NOTICE 1201 OF 1993

BOARD ON TARIFFS AND TRADE

CUSTOMS AND EXCISE TARIFF APPLICATIONS: LIST 42/93

The following applications concerning the Customs and Excise Tariff have been received by the Board on Tariffs and Trade. Any objections to or comments on these representations must be submitted to the Chairman, Board on Tariffs and Trade, Private Bag X753, Pretoria, 0001, within six weeks of the date of this notice. Attention is drawn to the fact that the rates of duty mentioned in the applications are those requested by the applicants and that the Board may, depending on its findings, recommend higher or lower rates of duty.

Reduction in the duty on:

Nonwovens, whether or not impregnated, coated, covered or laminated, classifiable under tariff heading 56.03, from various rates of duty to free.

[BTT Ref. T5/2/11/4/1 (930375) (Ms. H. Claassens)]

Applicant:

(i) Board on Tariffs and Trade, Private Bag X753, Pretoria, 0001.

(ii) Vitamed (Pty) Ltd, P.O. Box 27183, Benrose, 2011.

Rebate of the duty (Schedule 3) on:

1. Plates, sheets, film, foil and strip, of polymers of vinyl chloride, of a thickness exceeding 0,75 mm but not exceeding 3 mm, classifiable under tariff subheading 3921.12.50, for the manufacture of handbags.

[BTT Ref. T5/2/8/3/1 (930383) (Mr H. P. le Roux)]

Applicant:

Busy Bag and Belt (Pty) Ltd, P.O. Box 795, Stanger, 4450.

2. Polyether-polyols, containing two or more hydroxyl groups, liquids or pastes, with a hydroxyl number of 20 mg KOH/g or more but not exceeding 800 mg KOH/g, being glycerine fully reacted with ethylene oxides only, classifiable under tariff subheading 3907.20.10, for the manufacture of polyether polyols.

[BTT Ref. T5/2/7/21 (930404) (Mr G. Bester)]

Applicant:

Polyol Chemicals (Pty) Ltd, P.O. Box 246, Umbogintwini, 4120.

3. Polyether-polyols, containing 2 or more hydroxyl groups, liquids or pastes with a hydroxyl number of 20 mg KOH/g or more but not exceeding 800 mg KOH/g, being polyether-polyols, capped with ethylene oxides and having a primary hydroxyl content of not less than 20%, classifiable under tariff subheading 3907.20.10, for the manufacture of polyol blends.

[BTT Ref. T5/2/7/2/1 (930403) (Mr G. Bester)]

Applikant:

Industrial Urethanes (Edms.) Bpk., Posbus 411, Edenvale, 1610.

Algemeen:

1. Die intrekking van die teruggawevoorsiening vir vis en visprodukte by items: 501.02/03.05/01.00, 501.02/03.06/01.00 en 501.02/48.19/01.00.

[RTH-verw. T5/2/4/2/1 (930390) (Mnr. D. Jonker)]

Applikant:

Kommissaris van Doeane en Aksyns, Privaat Sak X47, Pretoria, 0001.

2. Hersiening van die doeaneregte op vleis en vleisprodukte, indeelbaar by die volgende tariefposte en -subposte 02.01 tot 02.06, 02.10, 15.01 tot 15.06, 1516.10, 15.17.90, 15.22, 16.01 tot 16.03, met die oog op moontlike verhoging/verlaging en die vereenvoudiging van die tariefstruktuur.

[RTH-verw. T5/2/1/3/1 (930405) (Mnr. J. Theron)]

Applikant:

Raad op Tariewe en Handel, Privaat Sak X753, Pretoria, 0001.

(Opmerking: Hierdie regte word hersien na aanleiding van samesprekings met die Vleisraad.)

Algemeen:

Hersiening van die doeane- regte op ru- en geraffineerde plantaardige olies en oliekoek met die oog op verlaging of verhoging van dié regte en die moontlike instelling van 'n outomatiese tariefaanpassingsmeganisme vir die betrokke produkte sowel as die vereenvoudiging van die tarief.

[RTH-verw. T5/2/3/2/1 (930388) (Adri van der Merwe)]

Applikant:

Raad op Tariewe en Handel, Privaat Sak X753, Pretoria, 0001.

(Opmerking: Die Raad het in hierdie verband 'n versoek van die Oliesaderaad ontvang om die moontlikheid van die instelling van 'n outomatiese tariefaanpassing meganisme vir hierdie bedryf te ondersoek.)

Lys 41/93 is by Algemene Kennisgewing 1172 van 26 November 1993 gepubliseer.

(3 Desember 1993)

KENNISGEWING 1202 VAN 1993

DEPARTEMENT VAN BINNELANDSE SAKE

WET OP REGISTRASIE VAN GEBOORTES EN STERFTES, 1992 (WET No. 51 VAN 1992)

KENNISGEWING VAN WYSIGING VAN GELDE BETAALBAAR

Met verwysing na paragraaf (e) van Kennisgewing 1578 van 1993, wat in *Staatskoerant* No. 15081 van 27 Augustus 1993 gepubliseer is, word hiermee ooreenkomstig die bepalings van artikel 8 van die Wet op Registrasie van Geboortes en Sterftes, 1992 (Wet No. 51 van 1992), bekendgemaak dat dienste wat ingevolge die bepalings van artikel 23 van die gemelde Wet met betrekking tot inskrywing van voorname of vanne in geboorteregisters gelewer word, met ingang van 12 November 1993 gratis is.

(3 Desember 1993)

Applicant:

Industrial Urethanes (Pty) Ltd, P.O. Box 411, Edenvale, 1610.

General:

1. Withdrawal of the drawback facilities for fish and fish products under items 501.02/03.05/01.00, 501.02/03.06/01.00 and 501.02/48.19/01.00.

[BTT Ref. T5/2/4/2/1 (930390) (Mr D. Jonker)]

Applicant:

Commissioner for Customs and Excise, Private Bag X47, Pretoria, 0001.

2. Review of the customs duties on meat and meat products, classifiable under the following tariff headings and subheadings 02.01 to 02.06, 02.10, 15.01 to 15.06, 1516.10, 15.17.90, 15.22, 16.01 to 16.03, with a view to the possible increase/decrease and simplification of the tariff structure.

[BTT Ref. T5/2/1/3/1 (930405) (Mr J. Theron)]

Applicant:

Board on Tariffs and Trade, Private Bag X753, Pretoria, 0001.

(Note: These duties are being reviewed following discussions with the Meat Board.)

General:

Review of the customs duties on crude and refined vegetable oils and oilcakes with a view to decreasing or increasing the duties and the possible introduction of an automatic tariff adjustment mechanism on the products concerned as well as the simplification of the tariff.

[BTT Ref. T5/2/3/2/1 (930388) (Adri van der Merwe)]

Applicant:

Board on Tariffs and Trade, Private Bag X753, Pretoria, 0001.

(Note: The Board received a request from the Oilseeds Board in this respect to investigate the possible introduction of an automatic tariff adjustment mechanism for this industry.)

List 41/93 was published under General Notice 1172 of 26 November 1993.

(3 December 1993)

NOTICE 1202 OF 1993

DEPARTMENT OF HOME AFFAIRS

BIRTHS AND DEATHS REGISTRATION ACT, 1992 (ACT No. 51 OF 1992)

NOTICE OF AMENDMENT OF FEES PAYABLE

With reference to paragraph (e) of Notice 1578 of 1993, which was published in *Government Gazette* No. 15081 of 27 August 1993, it is hereby made known in terms of section 8 of the Births and Deaths Registration Act, 1992 (Act No. 51 of 1992), that services which are rendered in terms of section 23 of the said Act in respect of the insertion of forenames or surnames in birth registers, are free of charge with effect from 12 November 1993.

(3 December 1993)

KENNISGEWING 1203 VAN 1993

DEPARTEMENT VAN VERVOER

KONSEP WYSIGINGSWETSONTWERP OP DIE LISENSIËRING VAN LUGDIENSTE

Die Direkteur-generaal: Vervoer publiseer hierby vir algemene inligting en kommentaar die Konsep Wysigingswetsontwerp op die Lisensiëring van Lugdienste, soos in die Bylae uiteengesit. Kommentaar of vertoë op hierdie konsep wetsontwerp moet nie later nie as **31 Januarie 1994** skriftelik by die Direkteur-generaal: Vervoer (Hoofdirektoraat: Burgerlugvaartowerheid), Privaat Sak X193, Pretoria, 0001, ingedien word.

BYLAE

Wysiging van artikel 1 van Wet 115 van 1990

1. Artikel 1 van die Wet op die Lisensiëring van Lugdienste, 1990 (hieronder die Hoofwet genoem), word hierby gewysig—

(a) deur die volgende omskrywings na die omskrywing van "bedryfsertifikaat" in te voeg:

" 'bedryfshandleiding' 'n bedryfshandleiding in artikel 22 (1A) bedoel;

'beslote korporasie' 'n beslote korporasie soos omskryf in artikel 1 van die Wet op Beslote Korporasies, 1984 (Wet No. 69 van 1984);";

(b) deur die volgende omskrywing na die omskrywing van "Direkteur-generaal" in te voeg:

" 'gemagtigde beampte' 'n gemagtigde beampte soos omskryf in artikel 1 van die Lugvaartwet, 1962 (Wet No. 74 van 1962);";

(c) deur die volgende omskrywing na die omskrywing van "inwoner van die Republiek" in te voeg:

" 'inspekteur' 'n inspekteur soos omskryf in artikel 1 van die Lugvaartwet, 1962 (Wet No. 74 van 1962);";

(d) deur die omskrywing van "lugdiens" deur die volgende omskrywing te vervang:

" 'lugdiens' enige diens wat deur middel van 'n lugvaartuig teen vergoeding bedryf word, maar wat geag word nie—

(a) die verhuur van 'n lugvaartuig, tesame met die bemanning, aan 'n lisensiehouer;

(b) 'n diens wat alleenlik tot voordeel van 'n maatskappy of 'n groep maatskappye of 'n filiaal daarvan, in die kommersiële bedrywighede daarvan bedryf word deur 'n persoon wat in diens van daardie maatskappy of groep maatskappye of filiaal is en wat nie teen vergoeding aan die publiek aangebied word nie;

NOTICE 1203 OF 1993

DEPARTMENT OF TRANSPORT

DRAFT AIR SERVICES LICENSING AMENDMENT BILL

The Director-General: Transport hereby publishes the Draft Air Services Licensing Amendment Bill, as set out in the Schedule, for general information and comment. Any comments or representations on this draft bill should be lodged in writing with the Director-General: Transport (Chief Directorate: Civil Aviation Authority), Private Bag X193, Pretoria, 0001, not later than **31 January 1994.**

SCHEDULE

Amendment of section 1 of Act 115 of 1990

1. Section 1 of the Air Services Licensing Act, 1990 (hereinafter referred to as the principal Act), is hereby amended—

(a) by the substitution for the definition of "air service" of the following definition:

" 'air service' means any service operated by means of an aircraft for reward, but shall be deemed not to include—

(a) the hiring out of an aircraft together with a crew to a licensee;

(b) a service operated solely for the benefit of a company or a group of companies or any subsidiary thereof in its commercial activities by a person who is in the employ of such company or group of companies or subsidiary, and which is not offered for reward to the public;

(c) training provided by or on behalf of the seller of an aircraft to the purchaser, where such training is included in the sale agreement and is provided in respect of such aircraft;

(d) the conduction of flight testing or assessment of skills in respect of flying an aircraft;

(e) the conducting in the manner and on the conditions prescribed of any kind of training or instruction in respect of flying an aircraft;";

(c) opleiding wat deur of namens die verkoper van 'n lugvaartuig aan die koper verskaf word, waar daardie opleiding in die koopooreenkoms ingesluit is en ten opsigte van daardie lugvaartuig verskaf word;

(d) die uitvoer van vlugtoetsing of die evaluering van vaardighede ten opsigte van die vlieg van 'n lugvaartuig;

(e) die uitvoer van enige soort opleiding of instruksie ten opsigte van die vlieg van 'n lugvaartuig op die wyse en voorwaardes wat voorgeskryf is,

in te sluit nie;"; en

(e) deur die volgende omskrywing na die omskrywing van "lugdiens" in te voeg:

" 'maatskappy' 'n maatskappy soos omskryf in artikel 1 van die Maatskappyewet, 1973 (Wet No. 61 van 1973);".

Wysiging van artikel 8 van Wet 115 van 1990

2. Artikel 8 van die Hoofwet word hierby gewysig deur subartikel (11) deur die volgende subartikel te vervang:

"(11) Afskrifte van die aantekeninge in subartikel (9) bedoel, of van enige deel daarvan, kan teen betaling van die voorgeskrewe gelde verkry word **[: Met dien verstande dat afskrifte van daardie verrigtinge wat ingevolge artikel 16 (3) (c) *in camera* gehou is, slegs aan persone wat toe-stemming gehad het om by sodanige verrigtinge teenwoordig te wees, beskikbaar gestel word]**.".

Vervanging van artikel 12 van Wet 115 van 1990

3. Artikel 12 van die Hoofwet word hierby deur die volgende artikel vervang:

"12. (1) Behoudens die bepalings van hierdie Wet mag niemand 'n lugdiens bedryf of poog om 'n lugdiens te bedryf nie, tensy dit bedryf word kragtens en ooreenkomstig die bepalings en onderworpe aan die voorwaardes van 'n lugdienslisensie wat aan so iemand ingevolge hierdie Wet uitgereik is of geag word aldus uitgereik te wees.

(2) Die raad kan, op aansoek van 'n persoon, aan hom vrystelling verleen van die bepalings van subartikel (1) of van enige ander bepaling van hierdie Wet, indien na die oordeel van die raad, die betrokke persoon 'n lugdiens op 'n nie-winsgewende grondslag verskaf of voornemens is om dit te verskaf, vir doeleindes wat in verband staan met maatskaplike welsyn, liefdadigheid of ontspanning, of vir doeleindes van reddingswerk uit menslikheidsoorwegings, of dat die toestaan van die vrystelling daartoe sal bydra om lewens te red.

(b) by the insertion after the definition of "air service" of the following definitions:

" 'authorized officer' means an authorized officer as defined in section 1 of the Aviation Act, 1962 (Act No. 74 of 1962);

'close corporation' means a close corporation as defined in section 1 of the Close Corporations Act, 1984 (Act No. 69 of 1984);

'company' means a company as defined in section 1 of the Companies Act, 1973 (Act No. 61 of 1973);";

(c) by the insertion after the definition of "Domestic air service" of the following definition:

" 'inspector' means an inspector as defined in section 1 of the Aviation Act, 1962;"; and

(d) by the insertion after the definition of "operating certificate" of the following definition:

" 'operations manual' means an operations manual referred to in section 22 (1A);".

Amendment of section 8 of Act 115 of 1990

2. Section 8 of the principal Act is hereby amended by the substitution for subsection (11) of the following subsection:

"(11) Copies of the record referred to in subsection (9), or of any part thereof, may be obtained against payment of the prescribed fee **[: Provided that copies of those proceedings which were held *in camera* in terms of section 16 (3) (c), shall be made available only to persons who had permission to attend such proceedings]**.".

Substitution of section 12 of Act 115 of 1990

3. The following section is hereby substituted for section 12 of the principal Act:

"12. (1) Subject to the provisions of this Act, no person shall operate or attempt to operate an air service, unless it is operated under and in accordance with the terms and subject to the conditions of an air service licence issued to that person in terms of this Act or deemed to have been so issued.

(2) The council may on the application of any person, exempt him from the provisions of sub-section (1) or from any other provision of this Act, if in the opinion of the council the person concerned is operating or proposes to operate an air service on a non-profit basis for purposes incidental to social welfare, charity or recreation, or for purposes of salvage on humanitarian grounds, or that the granting of such exemption will assist in saving life.

(3) Enige vrystelling wat kragtens subartikel (2) verleen word, kan beperk word om slegs op een of meer lugvaartuie of een of meer bepaalde roetes, lugreise of transaksies van toepassing te wees, of kan beperk word met betrekking tot tyd, gebied of afstand, of andersins na goeddunke van die raad.

(4) Die raad publiseer die voorgeskrewe besonderhede ten opsigte van elke vrystelling ingevolge subartikel (2) vir 'n tydperk van minstens 90 dae verleen word by kennisgewing in die *Staatskoerant*.".

Wysiging van artikel 14 van Wet 115 van 1990

4. Artikel 14 van die Hoofwet word hierby gewysig—

(a) deur subartikel (2) deur die volgende subartikel te vervang:

"(2) Indien 'n lisensiehouer—

(a) die tipe lugdiens of kategorie lugvaartuig soos op sy lisensie uiteengesit;

(b) sy regstatus;

(c) in die geval van 'n vennootskap of 'n beslote korporasie, die besonderhede van enige vennoot of lid verbonde aan daardie vennootskap of beslote korporasie;

(d) in die geval van 'n maatskappy, die beherende aandeelhouding van daardie maatskappy;

(e) die personeel wat vir veiligheid en betroubaarheid van die lugdiens verantwoordelik en aanspreeklik is en wat deur die lisensiehouer aangewys is; of

(f) die besonderhede op sy lisensie weens 'n beoogde vervreemding van die lugdiens op enige wyse of volgens enige verkoopproses,

wil wysig, doen hy op die voorgeskrewe vorm by die raad vir sodanige wysiging aansoek."; en

(b) deur subartikel (3) deur die volgende subartikel te vervang:

"(3) **['n Aansoek in subartikel (1) of (2) bedoel, word deur die voorgeskrewe gelde ten opsigte van sodanige aansoek vergesel]** Indien 'n lisensiehouer vertoë aan die raad wil rig ten einde vrystelling van die bepalings van artikel 16 (4) (c) of (e) te verkry, moet hy sodanige vertoë op die voorgeskrewe vorm rig.".

Wysiging van artikel 15 van Wet 115 van 1990

5. Artikel 15 van die Hoofwet word hierby gewysig—

(a) deur in subartikel (2) die woorde "en by betaling van die voorgeskrewe gelde" te skrap; en

(3) Any exemption granted under subsection (2) shall be limited so as to apply only in respect of one or more aircraft or one or more particular routes, journeys or transactions, and shall be limited as to time, area or distance, or otherwise as the council may deem fit.

(4) The council shall publish the prescribed particulars in respect of each exemption granted in terms of subsection (2) for a period exceeding 90 days by notice in the *Gazette*.".

Amendment of section 14 of Act 115 of 1990

4. Section 14 of the principal Act is hereby amended—

(a) by the substitution for subsection (2) of the following subsection:

"(2) If a licensee desires to—

(a) amend the type of air service or the category of aircraft specified on his licence;

(b) amend his legal status;

(c) in the case of a partnership or a close corporation, amend the particulars of any partner or member associated in such partnership or close corporation;

(d) in the case of a company, amend the controlling shareholding of such company;

(e) amend the personnel responsible and accountable for the safety and reliability of the air service designated by the licensee; or

(f) amend the particulars on his licence because of an envisaged disposal of the air service by any means or process of sale,

he shall apply to the council on the prescribed form for such amendment."; and

(b) by the substitution for subsection (3) of the following subsection:

"(3) **[An application referred to in subsection (1) or (2) shall be accompanied by the prescribed fee in respect of such application]** If a licensee desires to address representations to the council to obtain an exemption from the provisions of section 16 (4) (c) or (e), he shall make such representations on the prescribed form.".

Amendment of section 15 of Act 115 of 1990

5. Section 15 of the principal Act is hereby amended—

(a) by the deletion in subsection (2) of the words "and on payment of the prescribed fee"; and

(b) deur subartikel (3) deur die volgende subartikel te vervang:

"(3) Enige persoon kan, binne 21 dae nadat die kennisgewing in subartikel (1) bedoel, gepubliseer is, op die voorgeskrewe wyse skriftelik vertoë tot die raad teen of ten gunste van sodanige aansoek rig: Met dien verstande dat daardie vertoë gegrond moet word slegs op die aansoeker se vermoë om aan die vereistes bedoel in artikel 16 (4) te voldoen.".

Wysiging van artikel 16 van Wet 115 van 1990

6. Artikel 16 van die Hoofwet word hierby gewysig—

(a) deur in paragraaf (c) van subartikel (3) die woorde "tensy die raad anders gelas" te skrap;

(b) deur in subartikel (4) paragraaf (b) te skrap;

(c) deur in subartikel (4) paragraaf (d) deur die volgende paragraaf te vervang:

"(d) dat die persoon in paragraaf (c) bedoel, daadwerklik en prakties in beheer van die lugdiens sal wees; en";

(d) deur in subartikel (7) die woorde "by betaling van die voorgeskrewe gelde" te skrap; en

(e) deur die volgende subartikels na subartikel (7) in te voeg:

"(8) Die raad publiseer die voorgeskrewe besonderhede ten opsigte van elke lisensie wat ingevolge subartikel (4) uitgereik of gewysig word by kennisgewing in die *Staatskoerant*.

(9) Indien 'n aansoeker versuim om op 'n skriftelike versoek deur die raad om verdere besonderhede ten opsigte van sy aansoek, binne 90 dae na sodanige versoek te reageer, of indien hy 'n lasgewing in subartikel (3) (a) beoog om voor die raad op 'n vergaderinge verskyn, verontagsaam of by twee opeenvolgende geleenthede versuim om te verskyn sonder om aanvaarbare redes aan die raad te verstrek en daardeur die finale beslissing van sodanige aansoek vertraag, kan die raad na goeddunke die aansoek ingetrek ag, waarna die aansoeker die aansoekgelde wat betaal is, verbeur.".

Wysiging van artikel 18 van Wet 115 van 1990

7. Artikel 18 van die Hoofwet word hierby gewysig deur in subartikel (3) die woorde "by betaling van die voorgeskrewe gelde" te skrap.

Wysiging van artikel 19 van Wet 115 van 1990

8. Artikel 19 van die Hoofwet word hierby gewysig deur die volgende paragrawe na paragraaf (d) in te voeg:

"(e) die lisensiehouer verseker is soos voorgeskryf met betrekking tot die klas en tipe lugdiens, en die kategorie lugvaartuig op sy lisensie uiteengesit, en ten opsigte van die voorgeskrewe aard, klas of soort van versekering; en

(f) die lisensie verval indien die gelde bedoel in artikel 29 (1) (k) nie binne die voorgeskrewe tydperk betaal word nie.".

(b) by the substitution for subsection (3) of the following subsection:

"(3) Any person may address in writing, within 21 days after the publication of the notice referred to in subsection (1), representations in the prescribed manner to the council against or in favour of such application: Provided that those representations shall be founded only on the applicant's ability to comply with the requirements referred to in section 16 (4).".

Amendment of section 16 of Act 115 of 1990

6. Section 16 of the principal Act is hereby amended—

(a) by the deletion in paragraph (c) of subsection (3) of the words "unless the council otherwise determines";

(b) by the deletion in subsection (4) of paragraph (b);

(c) by the substitution in subsection (4) for paragraph (d) of the following paragraph:

"(d) that the person referred to in paragraph (c) will be actively and effectively in control of the air service; and";

(d) by the deletion in subsection (7) of the words "on payment of the prescribed fee"; and

(e) by the insertion after subsection (7) of the following subsections:

"(8) The council shall publish the prescribed particulars in respect of each licence issued or amended in terms of subsection (4) by notice in the *Gazette*.

(9) If an applicant fails to respond to a request in writing by the council for further particulars in respect of his application within 90 days after such request, or if he ignores an order contemplated in subsection (3) (a) to appear before the council at a meeting or fails to appear on two successive occasions without furnishing reasons to the satisfaction of the council and thereby delaying the final adjudication of such application, the council may at its discretion deem the application to be withdrawn upon which the applicant shall forfeit the application fee paid.".

Amendment of section 18 of Act 115 of 1990

7. Section 18 of the principal Act is hereby amended by the deletion in subsection (3) of the words "on payment of the prescribed fee".

Amendment of section 19 of Act 115 of 1990

8. Section 19 of the principal Act is hereby amended by the insertion of the following paragraphs after paragraph (d):

"(e) the licensee is insured as prescribed in relation to the class and type of air service, and the category of aircraft mentioned in his licence, and in respect of the prescribed nature, class or kind of insurance; and

(f) the licence shall lapse if the fees referred to in section 29 (1) (k) are not paid within the prescribed period.".

Wysiging van artikel 22 van Wet 115 van 1990

9. Artikel 22 van die Hoofwet word hierby gewysig—

(a) deur die volgende subartikel na subartikel (1) in te voeg:

"(1A) 'n Lisensiehouer doen op die voorgeskrewe wyse om die goedkeuring van 'n bedryfshandleiding by die Kommissaris van Burgerlugvaart aansoek."; en

(b) deur in subartikel (2) die woorde "die voorgeskrewe gelde, en" te skrap.

Wysiging van artikel 23 van Wet 115 van 1990

10. Artikel 23 van die Hoofwet word hierby gewysig deur in subartikel (3) die woorde "na betaling van die voorgeskrewe gelde" te skrap.

Vervanging van artikel 24 van Wet 115 van 1990

11. Artikel 24 van die Hoofwet word hierby deur die volgende artikel vervang:

"**24.** (1) Die lisensiehouer—

(a) stel die Kommissaris van Burgerlugvaart op die voorgeskrewe wyse in kennis voordat enige verandering van die besonderhede op sy bedryfsertifikaat in werking gestel word;

(b) verskaf binne die voorgeskrewe tydperk die voorgeskrewe statistiese inligting aan die raad;

(c) bewaar sy lisensie en bedryfsertifikaat op 'n veilige plek, en toon sodanige lisensie en bedryfsertifikaat aan 'n gemagtigde beampte of inspekteur vir inspeksie wanneer daar van hom vereis word om dit te doen; en

(d) stel die raad skriftelik **[op die voorgeskrewe wyse—**

(i)] van enige verandering **[betreffende die besonderhede in artikel 14 (4) bedoel; of**

(ii) van enige inkorting, afstanddoening of uitbreiding] van die voorgeskrewe komponente van die bedryf van die betrokke lugdiens of enige gedeelte daarvan,

in kennis, en sodanige kennis moet minstens 14 dae voordat sodanige verandering **[, inkorting, afstanddoening of uitbreiding]** in werking gestel word die raad bereik.

(2)Die raad openbaar nie die statistiese inligting in subartikel (1) (b) bedoel op so 'n wyse dat die bedrywighede van 'n lisensiehouer daardeur geïdentifiseer word nie tensy die skriftelike toestemming van die lisensiehouer verkry is.".

Amendment of section 22 of Act 115 of 1990

9. Section 22 of the principal Act is hereby amended—

(a) by die insertion of the following subsection after subsection (1):

"(1A) A licensee shall apply to the Commissioner for Civil Aviation in the prescribed manner for the approval of an operations manual."; and

(b) by the deletion in subsection (2) of the words "the prescribed fee and".

Amendment of section 23 of Act 115 of 1990

10. Section 23 of the principal Act is hereby amended by the deletion in subsection (3) of the words "on payment of the prescribed fee".

Substitution of section 24 of Act 115 of 1990

11. The following section is hereby stituted for section 24 of the principal Act:

"**24.** (1) The licensee shall—

(a) notify the Commissioner for Civil Aviation, in the prescribed manner, before any change is effected to the particulars on his operating certificate;

(b) furnish the council within the prescribed period with the prescribed statistical information;

(c) keep his licence and operating certificate in a safe place and produce such licence and operating certificate to an authorized officer or inspector for inspection when required to do so; and

(d) notify the council in **[the prescribed manner—**

(i)] writing of any change **[regarding the particulars referred to in section 14 (4); or**

(ii) of any curtailment, abandonment or extension] to the prescribed components of the operation of the air service concerned or any part thereof,

and such notice shall reach the council at least 14 days before such change **[, curtailment, abandonment or extension]** is effected.

(2) The council shall not disclose the statistical information referred to in subsection (1) (b) in such a manner that the activities of the licensee are identified thereby unless the written permission of the licensee has been obtained.".

Wysiging van artikel 25 van Wet 115 van 1990

12. Artikel 25 van die Hoofwet word hierby gewysig deur subartikel (1) deur die volgende subartikel te vervang:

"(1) Iemand wat hom veronreg voel—

(a) deur die weiering van die raad of die Kommissaris van Burgerlugvaart om aan hom 'n lisensie of 'n bedryfsertifikaat, na gelang van die geval, uit te reik;

(b) deur 'n beslissing van die raad ingevolge artikel 20 (1) (b) of (c); of

(c) deur 'n beslissing van die Kommissaris van Burgerlugvaart ingevolge artikel 22 (8) (b),

kan op die voorgeskrewe wyse teen sodanige weiering of beslissing appelleer na die provinsiale of plaaslike afdeling van die Hooggeregshof van Suid-Afrika wat jurisdiksie het oor die gebied waarin sodanige persoon woonagtig is, binne 30 dae nadat hy van sodanige weiering of beslissing bewus geword het, of binne die verdere tydperk, wat nie twee maande oorskry nie, wat bedoelde hof by die aanvoer van goeie gronde toelaat.".

Wysiging van artikel 26 van Wet 115 van 1990

13. Artikel 26 van die Hoofwet word hierby gewysig deur subartikel (1) deur die volgende subartikel te vervang:

"(1) Iemand wat opsetlik of nalatiglik—

(a) versuim om aan 'n lasgewing in artikel 11 (1) of (3) bedoel, te voldoen, of versuim om, nadat hy ingevolge artikel 11 (1) verskyn het, aanwesig te bly of weier om as getuie beëdig te word of te bevestig of om deur die raad ingevolge artikel 11 (3), of 'n persoon ingevolge artikel 16 (3) (d), ondervra te word of om vrae tydens sodanige ondervraging te beantwoord;

(b) artikel 12 of enige bepaling of voorwaarde in artikel 12 bedoel of 'n voorwaarde in artikel 19 bedoel, oortree of versuim om daaraan te voldoen;

(c) weier of versuim om 'n lisensie ingevolge artikel 21 terug te besorg;

(d) weier of versuim om die pligte in artikel 24 bedoel, na te kom;

(e) 'n lisensie, bedryfsertifikaat of ander dokument kragtens hierdie Wet uitgereik, vervals, namaak, verander, ontsier of skend, of 'n byvoeging daartoe maak, of in besit is van 'n lisensie, bedryfsertifikaat of ander dokument wat aldus vervals, nagemaak, verander, ontsier of geskend is, of waartoe 'n byvoeging gemaak is;

Amendment of section 25 of Act 115 of 1990

12. Section 25 of the principal Act is hereby amended by the substitution for subsection (1) of the following subsection:

"(1) Any person who feels aggrieved—

(a) by the refusal of the council or the Commissioner for Civil Aviation to issue to him a licence or an operating certificate, as the case may be;

(b) by a decision of the council in terms of section 20 (1) (b) or (c); or

(c) by a decision of the Commissioner for Civil Aviation in terms of section 22 (8) (b),

may in the prescribed manner appeal against such refusal or decision to the provincial or local division of the Supreme Court of South Africa having jurisdiction in the area within which such person is resident, within 30 days after he became aware of such refusal or decision, or within such further period, not exceeding two months, as the said court may allow on good cause shown.".

Amendment of section 26 of Act 115 of 1990

13. Section 26 of the principal Act is hereby amended by the substitution for subsection (1) of the following subsection:

"(1) Any person who wilfully or negligently—

(a) fails to comply with a direction referred to in section 11 (1) or (3), or fails to remain in attendance after appearing in terms of section 11 (1), or refuses to be sworn or to affirm as a witness or to be interrogated by the council in terms of section 11 (3), or any person in terms of section 16 (3) (d), or to answer questions during such interrogation;

(b) contravenes or fails to comply with section 12 or a term or condition referred to in section 12 or a condition referred to in section 19;

(c) refuses or fails to return a licence in terms of section 21;

(d) refuses or fails to fulfil the duties referred to in section 24;

(e) falsifies, counterfeits, alters, defaces or mutilates, or adds anything to, a licence, operating certificate or other document issued under this Act, or is in possession of a licence, operating certificate or other document which has been thus falsified, counterfeited, altered, defaced or mutilated, or to which an addition has been made;

(f) 'n lisensie, bedryfsertifikaat of ander dokument kragtens hierdie Wet uitgereik waarvan hy nie die houer is nie, gebruik, of toelaat dat dit gebruik word; **[of]**

(g) vir die doeleindes van enige aansoek of vertoë ingevolge hierdie Wet, of tydens ondervraging ingevolge artikel 11 (3) of 16 (3) (d), inligting of besonderhede verstrek wat na sy wete vals of in 'n wesenlike opsig misleidend is; of

(h) 'n lugvaartuig vir die verskaffing van 'n lugdiens gebruik in stryd met die voorwaardes van 'n vrystelling wat kragtens artikel 12 (2) aan hom verleen is,

is aan 'n misdryf skuldig.".

Wysiging van artikel 29 van Wet 115 van 1990

14. Artikel 29 van die Hoofwet word hierby gewysig—

(a) deur subartikel (1) deur die volgende subartikel te vervang:

"(1) Die Minister kan, na oorleg met die raad, regulasies uitvaardig betreffende—

(a) die inligting wat deur 'n aansoeker ten opsigte van die aangeleenthede in artikel 16 (4A) bedoel, verskaf moet word;

(b) die maatstawwe waaraan 'n aansoeker by die toepassing van artikel 16 (4) (a) moet voldoen met betrekking tot die klas lisensie, tipe lugdiens en kategorie lugvaartuig in sy aansoek vermeld;

(c) die uitreiking en bewaring van passasierslugvervoerkaartjies en die besonderhede en endossemente wat in sodanige kaartjies vervat moet wees;

(d) die uitreiking en bewaring van lugvragbriewe en die besonderhede wat in sodanige lugvragbriewe vervat moet wees;

(e) die samestelling en bewaring van passasierslyste en die besonderhede wat in sodanige lyste vervat moet wees;

(f) die uitvoer van vluginspeksies;

(g) die betaling van gelde ten opsigte van enige aansoek wat ingevolge hierdie Wet gedoen word;

(h) die betaling van gelde ten opsigte van die redes vir die goedkeuring of weiering van 'n aansoek om 'n lisensie;

(f) uses, or permits to be used, a licence, operating certificate or other document issued under this Act of which he is not the holder;

(g) for the purposes of any application or representations in terms of this Act or during interrogation in terms of section 11 (3) or 16 (3) (d), furnishes information or particulars which to his knowledge are false or misleading in any material respect; or

(h) uses an aircraft for the provision of an air service in contravention of the terms of an exemption granted to him under section 12 (2),

shall be guilty of an offence.".

Amendment of section 29 of Act 115 of 1990

14. Section 29 of the principal Act is hereby amended—

(a) by the substitution for subsection (1) of the following subsection:

"(1) The Minister may, after consultation with the council, make regulations regarding—

(a) the information to be furnished by an applicant in respect of the matters referred to in section 16 (4A);

(b) the standards to be complied with by an applicant, in relation to the class of licence, type of air service and category of aircraft mentioned in his application;

(c) the issuing and safe-keeping of passenger air transport tickets and the particulars and endorsements to be contained in such tickets;

(d) the issuing and safe-keeping of air waybills and the particulars to be contained in such waybills;

(e) the compilation and safe-keeping of passenger lists and the particulars to be contained in such lists;

(f) the carrying out of inflight inspections;

(g) the payment of fees in respect of any application made in terms of this Act;

(h) the payment of fees in respect of the reasons for the approval or refusal of an application for a licence;

(i) die betaling van gelde ten opsigte van die uitreiking van 'n lisensie, of die wysiging daarvan, met inbegrip van die tydperk waarbinne sodanige gelde betaal moet word;

(j) die betaling van gelde ten opsigte van die uitreiking van 'n bedryfsertifikaat, met inbegrip van die tydperk waarbinne sodanige gelde betaal moet word;

(k) die betaling van jaarlikse gelde ten opsigte van 'n lisensie;

(l) die betaling van gelde ten opsigte van die verskaffing van inligting uit enige register wat ingevolge hierdie Wet gehou word;

[(c)] (m) enige aangeleentheid wat ingevolge hierdie Wet voorgeskryf moet of kan word; en

[(d)] (n) enige ander aangeleentheid waarvan die reëling, volgens die oordeel van die Minister, nodig of wenslik is ten einde die oogmerke van hierdie Wet te bereik of te bevorder."; en

(b) deur die volgende subartikels na subartikel (2) in te voeg:

"(3) Voordat die Minister 'n regulasie kragtens hierdie artikel uitvaardig, moet hy die regulasies wat hy voornemens is om uit te vaardig, by kennisgewing in die *Staatskoerant* publiseer.

(4) Na publikasie van die kennisgewing in subartikel (3) bedoel, kan enige belanghebbende binne 'n in die kennisgewing vermelde tydperk, maar nie minder nie as vier weke vanaf die datum van publikasie van die kennisgewing, skriftelike vertoë tot die Direkteur-generaal: Vervoer vir voorlegging aan die Minister betreffende die betrokke regulasies rig.

(5) Na oorweging van die vertoë in subartikel (4) bedoel, kan die Minister, hetsy hy die betrokke regulasies aangepas het al dan nie, daardie regulasies in die finale vorm daarvan by kennisgewing in die *Staatskoerant* publiseer.

(6) Regulasies in subartikel (1) bedoel, moet binne 14 dae nadat dit kragtens subartikel (5) gepubliseer is, in die Parlement ter Tafel gelê word indien die Parlement dan in gewone sessie is, of, indien die Parlement nie dan in gewone sessie is nie, binne 14 dae na die aanvang van sy eersvolgende gewone sessie.".

(3 Desember 1993)

(i) the payment of fees in respect of the issuance of a licence, or the amendment thereof, including the period within which such fees shall be paid;

(j) the payment of fees in respect of the issuance of an operating certificate, including the period within which such fees shall be paid;

(k) the payment of annual fees in respect of a licence;

(l) the payment of fees in respect of the providing of information from any register which is kept in terms of this Act;

[(c)] (m) any matter which in terms of this Act is required or permitted to be prescribed; and

[(d)] (n) any other matter the regulation of which, in the opinion of the Minister, may be necessary or desirable in order to achieve or promote the objects of this Act."; and

(b) by the insertion after subsection (2) of the following subsections:

"(3) Before the Minister makes any regulation under this section, he shall publish the regulations he intends to make by notice in the *Gazette*.

(4) After publication of the notice referred to in subsection (3) any interested person may, within a period stated in the notice, but not less than four weeks as from the date of publication of the notice, address representations in writing to the Director-General for submission to the Minister regarding the regulations concerned.

(5) After considering the representations referred to in subsection (4), the Minister may, whether or not he has adjusted the regulations concerned, publish those regulations in their final form by notice in the *Gazette*.

(6) Regulations referred to in subsection (1) shall, within 14 days after they have been published under subsection (5), be laid upon the Table in Parliament if Parliament is then in ordinary session, or, if Parliament is not then in ordinary session, within 14 days after the commencement of its next ensuing ordinary session.".

(3 December 1993)

RAADSKENNISGEWING

RAADSKENNISGEWING 138 VAN 1993

KENNISGEWING VAN AANSOEK VIR OMSKRYWING VAN PRODUKSIEGEBIED: HERBERTSDALE

[Ingevolge artikel 6 van die Wyn van Oorsprong-skema ingestel kragtens artikel 14 van die Wet op Drankprodukte, 1989 (Wet No. 60 van 1989), en gepubliseer by Goewermentskennisgewing No. R. 1434 van 29 Junie 1990]

Kennis geskied hiermee dat mnr. J. P. W. Jonker by die Wyn- en Spiritusraad aansoek gedoen het vir die omskrywing in die *Staatskoerant* van die gebied as Herbertsdale, soos beskryf in die Bylae hiertoe, as 'n produksiegebied (wyk) vir die doeleindes van die vervaardiging van Wyn van Oorsprong.

Enigeen wat enige beswaar het teen die aansoek, word hierby aangesê om sy beswaar, met opgaaf van redes, skriftelik in te dien by die Sekretaris, Wyn- en Spiritusraad, Posbus 2176, Dennesig, Stellenbosch, 7599, binne 30 (dertig) dae van publikasie van hierdie kennisgewing.

BYLAE

PRODUKSIEGEBIED (WYK): HERBERTSDALE

Daardie gedeelte grond geleë binne die volgende grense:

Begin by die trigonometriese baken 37 op Grootkop ongeveer 2 km suidoos van die dorp Herbertsdale; daarvandaan in 'n wesnoordwestelike rigting met 'n denkbeeldige reguitlyn wat laasgenoemde baken verbind met die trigonometriese baken 82 op die plaas Die Possie ongeveer 5 km suidwes van die plaas Die Poort waar die Gouritsrivier deur die Langeberg-reeks breek; daarvandaan in 'n noordnoordoostelike rigting met 'n denkbeeldige reguitlyn wat laasgenoemde baken verbind met die trigonometriese baken 94 ongeveer 1,5 km wes van die plaas Die Poort; daarvandaan in 'n oostelike rigting met 'n denkbeeldige reguitlyn wat laasgenoemde baken verbind met die trigonometriese baken 263 ongeveer 4,5 km oos van die plaas Die Poort; daarvandaan in 'n oosnoordoostelike rigting met 'n denkbeeldige reguitlyn wat laasgenoemde baken verbind met die trigonometriese baken 83 ongeveer 9 km noordoos van die dorp Herbertsdale; daarvandaan in 'n suidsuidwestelike rigting met die trigonometriese baken 37 op Grootkop, die beginpunt hierbo genoem.

[Alle aanduidings in hierdie beskrywing verwys na die topografiese kaart van Suid-Afrika 1:50 000, velle 3321 DC Langberg (Tweede Uitgawe), 3321 DD Attakwaskloof (Tweede Uitgawe), 3421 BA Albertinia (Tweede Uitgawe), 3421 BB Herbertsdale (Tweede Uitgawe).]

(3 Desember 1993)

BOARD NOTICE

BOARD NOTICE 138 OF 1993

NOTICE OF APPLICATION FOR DEFINING OF PRODUCTION AREA: HERBERTSDALE

[In terms of section 6 of the Wine of Origin Scheme established under section 14 of the Liquor Products Act, 1989 (Act No. 60 of 1989), and published by Government Notice No. R. 1434 of 29 June 1990]

Notice is hereby given that Mr J. P. W. Jonker has applied to the Wine and Spirit Board to have the area known as Herbertsdale, as described in the Annexure hereto, defined in the *Government Gazette* as an area (ward) for the production of Wine of Origin.

Anyone having any objection against this application is hereby notified to lodge their objections, with motivations, in writing with the Secretary, Wine and Spirit Board, P.O. Box 2176, Dennesig, Stellenbosch, 7599, within 30 (thirty) days of publication of this notice.

ANNEXURE

PRODUCTION AREA (WARD): HERBERTSDALE

That portion of land situated within the following boundaries:

Beginning at the trigonometrical beacon 37 on Grootkop approximately 2 km south-east of the Town of Herbertsdale; thence in a west-north-westerly direction with an imaginary straight line which joins last-mentioned beacon with the trigonometrical beacon 82 on the farm Die Possie approximately 5 km south-west of the farm Die Poort where the Gouritz River passes through the Langeberg Mountain Range; thence in a north-north-easterly direction with an imaginary straight line which joins last-mentioned beacon with the trigonometrical beacon 94 approximately 1,5 km west of the farm Die Poort; thence in an easterly direction with an imaginary straight line which joins last-mentioned beacon with the trigonometrical beacon 263 approximately 4,5 km east of the farm Die Poort; thence in an east-north-easterly direction with an imaginary straight line which joins last-mentioned beacon with the trigonometrical beacon 83 approximately 9 km north-east of the Town of Herbertsdale; thence in a south-south-westerly direction with an imaginary straight line which joins last-mentioned beacon with the trigonometrical beacon 37 on Grootkop, the point of beginning mentioned above.

[All indications in this description refer to the topographical map of South Africa 1:50 000, sheets 3321 DC Langberg (Second Edition) 3321, DD Attakwaskloof (Second Edition), 3421 BA Albertinia (Second Edition), 3421 BB Herbertsdale (Second Edition).]

(3 December 1993)

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AMPTELIKE PUBLIKASIES IN VOORRAAD GENEEM GEDURENDE SEPTEMBER 1993 EN WAT BESKIKBAAR IS IN DIE STAATSDRUKKERY SE BOEKWINKELS TE PRETORIA EN KAAPSTAD

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RP 89/1993—Verslag van die Ouditeur-generaal oor die Appropriasie- en Diverse Rekenings van die Administrasie: Raad van Afgevaardigdes vir 1991–1992. ISBN 0-621-15467-9. Plaaslik **R12,85**; buitelands **R14,10**.

RP 94/1993—Verslag van die Ouditeur-generaal oor die Appropriasie- en Diverse Rekenings ten opsigte van die Administrasie: Volksraad vir 1991–1992. ISBN 0-621-15499-7. Plaaslik **R31,24**; buitelands **R34,25**.

RP 95/1993—Bygewerkte Spesiale Verslag van die Ouditeur-generaal oor die Finansiële Aangeleenthede van die Munisipaliteit van Ladybrand. ISBN 0-621-15504-7. Plaaslik **R1,60**; buitelands **R1,75**.

RP 96/1993—Verslag van die Ouditeur-generaal oor die Rekenings van die Sagtevrugteraad en die Suid-Afrikaanse Plantverbeteringsorganisasie vir die boekjaar 1 Oktober 1988 tot 30 September 1989. ISBN 0-621-15503-9. Plaaslik **R3,55**; buitelands **R3,90**.

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RP 100/1993—Report of the Auditor-General on the Accounts of the Rooibos Tea Board for the financial year 1 January 1991 to 31 December 1991. ISBN 0-621-15808-9. Local **R2,20**; other countries **R2,50**.

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RP 102/1993—Report of the Auditor-General on the Accounts of the Grain Sorghum Board for the financial year 1 January 1992 to 31 December 1992. ISBN 0-621-15807-0. Local **R2,55**; other countries **R2,80**.

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No. 1, Debatte van die Presidentsraad (Hansard): 12 Mei en 15 tot 17 Junie 1993. Plaaslik **R1,14**; buitelands **R1,40**.

No. 2, Debatte van die Presidentsraad (Hansard): 14 en 30 Oktober en 25 November 1992. Plaaslik **R1,14**; buitelands **R1,40**.

No. 22, Debatte van die Parlement (Hansard): Vyfde sessie—Negende Parlement: 21 tot 24 Junie 1993. Plaaslik **R1,14**; buitelands **R1,40**.

DIVERSE VERSLAE

Litostratigrafie van Huis Rivier Formasie (Kango Groep): Suid-Afrikaanse Komitee vir Stratigrafie, Litostratigrafiese Reeks No. 23. ISBN 0-621-15481-4. Plaaslik **R22,00**; buitelands **R25,00**.

Litostratigrafie van die Flamingberg Formasie (Vanrhynsdorp Groep): Suid-Afrikaanse Komitee vir Stratigrafie, Litostratigrafiese Reeks No. 24. ISBN 0-621-15480-6. Plaaslik **R22,00**; buitelands **R25,00**.

Maandelikse Uittreksel van Handelstatistiek: Januarie tot Junie 1993. Plaaslik **R26,95**; buitelands **R29,60**.

Gebinde dele van die *Staatskoerante* vir Mei 1993 (Deel A en B). Plaaslik **R68,40** (per deel); buitelands **R75,00** (per deel).

Patentjoernaal (insluitende Handelsmerk, Modelle en Outeursreg in Rolprente). Vol. 26, September 1993, No. 9. ISBN 0-031-086X. Plaaslik **R1,10**; buitelands **R1,25**.

PRESIDENT'S COUNCIL REPORTS

CR 2/1993—President's Council: Report of the Committee for Economic Affairs on the Promotion of Economic Literacy. ISBN 0-621-15533-0. Local **R36,59**; other countries **R40,15**.

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No. 1, Debates of the President's Council (Hansard): 12 May and 15 to 17 June 1993. Local **R1,14**; other countries **R1,40**.

No. 2, Debates of the President's Council (Hansard): 14 and 30 October and 25 November 1992. Local **R1,14**; other countries **R1,40**.

No. 22, Debates of the Parliament (Hansard): Fifth Session—Ninth Parliament: 21 to 24 June 1993. Local **R1,14**; other countries **R1,40**.

MISCELLANEOUS REPORTS

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Monthly Abstract of Trade Statistics, January to June 1993. Local **R26,95**; other countries **R29,60**.

Bound volumes of the *Government Gazettes* for May 1993 (Part A and B). Local **R68,40** (per part); other countries **R75,00** (per part).

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