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MAART 1985

No. 9621

PROCLAMATION

by the

State President of the Republic of South Africa

No. R. 22, 1985

**RE-APPOINTMENT OF A MEMBER OF THE SPECIAL
COURTS FOR HEARING INCOME TAX APPEALS**

Whereas by Proclamation 190 of 1977 Edgar Alfred Alexander of Cape Town was re-appointed under the provisions of section 83 (5) (a) of the Income Tax Act, 1962 (Act 58 of 1962), as a commercial member of the special courts for hearing income tax appeals, constituted under the provisions of subsection (3) of the said section;

And whereas the appointment of the said Edgar Alfred Alexander expired on the 9th September 1982;

Now, therefore, under the powers vested in me by subsection (5) (b) of the said section I do hereby re-appoint the said Edgar Alfred Alexander as a commercial member of the said special courts for a further period of five years from the date of this Proclamation subject to the provisions of the proviso to subsection (5) (a) of the said section.

Given under my Hand and the Seal of the Republic of South Africa at Cape Town this Twenty-first day of February, One thousand Nine hundred and Eighty-five.

P. W. BOTHA, State President.

By Order of the State President-in-Cabinet:

B. J. DU PLESSIS.

PROKLAMASIE

van die

Staatspresident van die Republiek van Suid-Afrika

No. R. 22, 1985

**HERAANSTELLING VAN 'N LID VAN DIE SPESIALE
HOWE VIR DIE VERHOOR VAN INKOMSTEBELAS-
TINGAPPELSAKE**

Nademaal Edgar Alfred Alexander van Kaapstad kragtens die bepaling van artikel 83 (5) (a) van die Inkomstebelastingwet, 1962 (Wet 58 van 1962), as kommersiële-lid van die spesiale howe vir die verhoor van inkomstebelastingappelsake, ingestel kragtens die bepalings van subartikel (3) van daardie artikel, by Proklamasie 190 van 1977 heraan gestel is;

En nademaal die ampstyd van genoemde Edgar Alfred Alexander op 9 September 1982 verstryk het;

So is dit dat ek hierby, kragtens die bevoegdheid my verleen by subartikel (5) (b) van bovermelde artikel die genoemde Edgar Alfred Alexander vir 'n verdere tydperk van vyf jaar met ingang van die datum van hierdie Proklamasie as kommersiële-lid van die genoemde spesiale howe, behoudens die voorbehoudsbepalings van subartikel (5) (a) van die genoemde artikel, heraan stel.

Gegee onder my Hand en die Seël van die Republiek van Suid-Afrika te Kaapstad, op hede die Een-en-twintigste dag van Februarie Eenduisend Negehoonderd Vyf-en-tagtig.

P. W. BOTHA, Staatspresident.

Op las van die Staatspresident-in-Kabinet:

B. J. DU PLESSIS.

ADMINISTRATION: HOUSE OF ASSEMBLY

DEPARTMENT OF EDUCATION AND CULTURE

No. R. 567

15 March 1985

UNIVERSITIES ACT, 1955

UNIVERSITY OF THE ORANGE FREE STATE.— AMENDMENT OF STATUTE

The Council of the University of the Orange Free State has, with the approval of the Minister of Education, and Culture, under section 17 (1) of the Universities Act, 1955 (Act 61 of 1955), framed the statutes set out in the Schedule hereto.

SCHEDULE

1. In this Schedule, unless inconsistent with the context, the expression "the Statute" means the Statute published under Government Notice R. 429 of 22 March 1963, as amended by Government Notices R. 1418 of 13 September 1963, R. 634 of 23 April 1971, R. 1525 of 25 August 1972, R. 521 of 5 April 1973, R. 348 of 8 March 1974, R. 512 of 1 April 1977, R. 2231 of 10 November 1978, R. 296 of 23 February 1979, R. 283 of 15 February 1980, R. 2405 of 28 November 1980, R. 1253 of 19 June 1981, R. 2836 of 31 December 1981 and R. 175 of 28 January 1983.

2. Paragraph 31 of the Statute is hereby amended by deleting the words "and finance".

3. Chapter VIII of the Statute is hereby amended by adding the following paragraph:

"Amendment of degrees and exchange of degree certificates"

67A. (1) Where the University has conferred a degree on any person and has issued a degree certificate in respect thereof to such person, and the designation of the degree is thereafter changed by the University, such person who wishes to exchange such certificate for a certificate under the new designation, may apply to the University for such exchange.

(2) The University may on the submission by the applicant of the existing certificate, issue a certificate under the new designation to him.

(3) As from the date of any such issue it shall for all purposes be deemed that the degree in question has been conferred on the person concerned under the new designation on the date on which the degree was conferred on him by the University under the previous designation."

DEPARTMENT OF AGRICULTURAL ECONOMICS AND MARKETING

No. R. 568

15 March 1985

MARKETING ACT, 1968 (ACT 59 OF 1968)

SUMMER GRAIN SCHEME.—LEVY AND SPECIAL LEVY ON GRAIN SORGHUM AND GRAIN SORGHUM PRODUCTS

I, Jacob Johannes Greyling Wentzel, Minister of Agricultural Economics, hereby make known in terms of section 79 (a) of the Marketing Act, 1968 (Act 59 of 1968), that—

(a) the Maize Board referred to in section 6 of the Summer Grain Scheme published by Proclamation R. 45, 1979, as amended, has under sections 23 and 24 of the said Scheme imposed the levy and special levy set out in the Schedule;

ADMINISTRASIE: VOLKSRAAD

DEPARTEMENT VAN ONDERWYS EN KULTUUR

No. R. 567

15 Maart 1985

WET OP UNIVERSITEITE, 1955

UNIVERSITEIT VAN DIE ORANJE-VRYSTAAT.— WYSIGING VAN STATUUT

Die Raad van die Universiteit van die Oranje-Vrystaat het kragtens artikel 17 (1) van die Wet op Universiteite (Wet 61 van 1955), met die goedkeuring van die Minister van Onderwys en Kultuur die statute in die Bylae hiervan opgestel.

BYLAE

1. In hierdie Bylae, tensy uit die samehang anders blyk, beteken die uitdrukking "die Statuut" die Statuut gepubliseer by Goewermenskennisgewing R. 429 van 22 Maart 1963, soos gewysig by Goewermenskennisgewings R. 1418 van 13 September 1963, R. 634 van 23 April 1971, R. 1525 van 25 Augustus 1972, R. 521 van 5 April 1973, R. 348 van 8 Maart 1974, R. 512 van 1 April 1977, R. 2231 van 10 November 1978, R. 296 van 23 Februarie 1979, R. 283 van 15 Februarie 1980, R. 2405 van 28 November 1980, R. 1253 van 19 Junie 1981, R. 2836 van 31 Desember 1981 en R. 175 van 28 Januarie 1983.

2. Paragraaf 31 van die Statuut word gewysig deur die woorde "en finansiële" te skrap.

3. Hoofstuk VIII van die Statuut word gewysig deur die volgende paragraaf in te voeg:

"Wysiging van graadbenaming en omruil van graad-sertifikaat"

67A. (1) Waar die Universiteit 'n graad aan iemand toegeken en 'n graadsertifikaat ten opsigte daarvan aan sodanige persoon uitgereik het, en die benaming van die graad daarna deur die Universiteit verander word, kan die betrokke persoon sodanige sertifikaat wil omruil vir 'n sertifikaat onder die nuwe benaming, by die Universiteit aansoek doen om sodanige omruiling.

(2) Die Universiteit kan by inhandiging deur die aansoeker van die bestaande sertifikaat, aan hom 'n sertifikaat onder die nuwe benaming uitreik.

(3) Met ingang van die datum van sodanige omruiling word dit vir alle doeleindes geag dat die betrokke graad onder die nuwe benaming aan die betrokke persoon toegeken is op die datum waarop die graad onder die vorige benaming deur die Universiteit aan hom toegeken is."

DEPARTEMENT VAN LANDBOU- EKONOMIE EN -BEMARKING

No. R. 568

15 Maart 1985

BEMARKINGSWET, 1968 (WET 59 VAN 1968)

SOMERGRAANSKEMA.—HEFFING EN SPESIALE HEFFING OP GRAANSORGHUM EN GRAANSORGHUM- PRODUKTE

Ek, Jacob Johannes Greyling Wentzel, Minister van Landbou-ekonomie, maak hierby ingevolge artikel 79 (a) van die Bemarkingswet, 1968 (Wet 59 van 1968), bekend dat—

(a) die Mielieraad bedoel in artikel 6 van die Somergraanskema gepubliseer by Proklamasie R. 45, 1979, soos gewysig, kragtens artikels 23 en 24 van genoemde Skema die heffing en spesiale heffing in die Bylae uiteengesit, opgelê het;

(b) the said levy and special levy have been approved by me and shall come into operation on the date of publication hereof; and

(c) Government Notice R. 879 of 1 May 1984, as amended by Government Notice R. 1409 of 6 July 1984, and Government Notice R. 1147 of 29 May 1981, as amended by Government Notices R. 918 of 7 May 1982, R. 904 of 29 April 1983 and R. 878 of 1 May 1984, are repealed with effect from the said date of commencement.

J. J. G. WENTZEL, Minister of Agricultural Economics.

SCHEDULE

1. Any word or expression in this Schedule to which a meaning has been assigned in the Scheme shall have that meaning and, unless the context otherwise indicates—

“class and grade”, with regard to grain sorghum, means a class and grade referred to in the regulations relating to the classification and grading of grain sorghum, made under section 89 of the Act;

“controlled area” means the area consisting of—

(a) the provinces of Transvaal and the Orange Free State;

(b) the Magisterial Districts of Alexandria, Aliwal North, East London, Hartswater, Kimberley, King William's Town, Kuruman, Port Elizabeth, Queenstown, Uitenhage, Vryburg and Warrenton in the Cape Province;

(c) those portions of the Magisterial Districts of Gordonia, Hay, Kenhardt and Prieska in the Cape Province, situated within a strip of 48 km wide on both banks of the Orange River between the Boegoeberg Dam and the point on the Orange River directly north of Noudonsies; and

(d) the Magisterial Districts of Bergville, Camperdown, Dannhauser, Dundee, Durban, Estcourt, Ixopo, Klip River, Kranskop, Lions River, Mooi River, Mount Currie, Newcastle, New Hanover, Paulpietersburg, Pietermaritzburg, Pinetown, Port Shepstone, Umvoti, Utrecht, Vryheid and Weenen in the province of Natal;

“seed” means grain sorghum intended for use as propagating material as defined in the Plant Improvement Act, 1976 (Act 53 of 1976); and

“the Scheme” means the Summer Grain Scheme published by Proclamation R. 45, 1979, as amended.

2. A levy and a special levy are hereby imposed on—

(a) all classes and grades of grain sorghum other than seed, that—

(i) is sold within the controlled area to the Board by the producers thereof; and

(ii) is sold within or outside the controlled area by the producers thereof to a person who is registered with the Board under section 36 of the Scheme as a grain sorghum trader, being a person dealing in the course of trade with grain sorghum, or a manufacturer of grain sorghum products, being a person dealing in the course of trade with grain sorghum products; and

(b) all grain sorghum and grain sorghum products that are exported from the Republic, excluding grain sorghum specifically sold by the Board for export.

3. The amount of the levy referred to in clause 2 shall—

(a) in the case of grain sorghum referred to in paragraph (a) of that clause, be R1,85 per ton; and

(b) in the case of grain sorghum and grain sorghum products referred to in paragraph (b) of that clause, be R1,85 per ton.

(b) genoemde heffing en spesiale heffing deur my goedgekeur is en op die datum van publikasie hiervan in werking tree; en

(c) Goewermenskennisgewing R. 879 van 1 Mei 1984, soos gewysig deur Goewermenskennisgewing R. 1409 van 6 Julie 1984, en Goewermenskennisgewing R. 1147 van 29 Mei 1981, soos gewysig deur Goewermenskennisgewings R. 918 van 7 Mei 1982, R. 904 van 29 April 1983 en R. 878 van 1 Mei 1984 met ingang van genoemde datum van inwerkingtreding herroep word.

J. J. G. WENTZEL, Minister van Landbou-ekonomie.

BYLAE

1. In hierdie Bylae het enige woord of uitdrukking waarvan 'n betekenis in die skema geheg is, daardie betekenis en, tensy uit die samehang anders blyk, beteken—

“beheerde gebied” die gebied bestaande uit—

(a) die provinsies Transvaal en die Oranje-Vrystaat;

(b) die landdrostdistrikte Alexandria, Aliwal-Noord, Hartswater, Kimberley, King William's Town, Kuruman, Oos-Londen, Port Elizabeth, Queenstown, Uitenhage, Vryburg en Warrenton in die Kaapprovinsie;

(c) daardie gedeeltes van die landdrostdistrikte Gordonia, Hay, Kenhardt en Prieska in die Kaapprovinsie, wat geleë is binne 'n strook van 48 km wyd aan beide oevers van die Oranjerivier tussen die Boegoebergdam en die punt op die Oranjerivier reg noord van Noudonsies; en

(d) die landdrostdistrikte Bergville, Camperdown, Dannhauser, Dundee, Durban, Estcourt, Ixopo, Kliprivier, Kranskop, Lions River, Mooirivier, Mount Currie, Newcastle, New Hanover, Paulpietersburg, Pietermaritzburg, Pinetown, Port Shepstone, Umvoti, Utrecht, Vryheid en Weenen in die provinsie Natal;

“die Skema” die Somergraanskema gepubliseer by Proklamasie R. 45, 1979, soos gewysig;

“klas en graad”, met betrekking tot graansorghum, 'n klas en graad bedoel in die regulasies met betrekking tot die klassifisering en gradering van graansorghum, kragtens artikel 89 van die Wet uitgevaardig; en

“saad” graansorghum bestem vir gebruik as voortplantingsmateriaal soos omskryf in die Plantverbeteringswet, 1976 (Wet 53 van 1976).

2. 'n Heffing en 'n spesiale heffing word hierby opgelê op—

(a) alle klasse en grade graansorghum anders as saad, wat—

(i) binne die beheerde gebied deur die produsente daarvan aan die Raad verkoop word; en

(ii) binne of buite die beheerde gebied deur die produsente daarvan aan 'n persoon verkoop word wat kragtens artikel 36 van die Skema by die Raad geregistreer is as 'n graansorghumhandelaar, synde 'n persoon wat met graansorghum as 'n besigheid handel, of 'n vervaardiger van graansorghumprodukte synde 'n persoon wat met graansorghumprodukte as 'n besigheid handel; en

(b) alle graansorghum en graansorghumprodukte wat uit die Republiek uitgevoer word, uitgesonderd graansorghum wat spesifiek deur die Raad vir uitvoer verkoop word.

3. Die bedrag van die heffing op klousule 2 bedoel, is—

(a) in die geval van graansorghum in paragraaf (a) van daardie klousule bedoel, R1,85 per ton; en

(b) in die geval van graansorghum en graansorghumprodukte in paragraaf (b) van daardie klousule bedoel, R1,85 per ton.

4. The amount of the special levy referred to in clause 2 shall—

(a) in the case of grain sorghum referred to in paragraph (a) of that clause, be R25,95 per ton; and

(b) in the case of grain sorghum and grain sorghum products referred to in paragraph (b) of that clause, be R50,00 per ton.

If grain sorghum or a grain sorghum product is sold or exported as contemplated in clause 2 in a container, the mass of the container concerned shall for the purposes of the calculation of the amount payable in respect of a levy and a special levy, not be taken into account: Provided that the mass of a bag (Hessian or jute) shall be deemed to be 1 kg.

DEPARTMENT OF FINANCE

No. R. 517

15 March 1985

CUSTOMS AND EXCISE ACT, 1964

AMENDMENT OF SCHEDULE 1 (No. 1/1/1096)

Under section 48 of the Customs and Excise Act, 1964, Part 1 of Schedule 1 to the said Act is hereby amended to the extent set out in the Schedule hereto.

K. D. S. DURR, Deputy Minister of Finance and of Trade and Industry.

4. Die bedrag van die spesiale heffing in klousule 2 bedoel, is—

(a) in die geval van graansorghum in paragraaf (a) van daardie klousule bedoel, R25,95 per ton; en

(b) in die geval van graansorghum en graansorghumprodukte in paragraaf (b) van daardie klousule bedoel, R50,00 per ton.

5. Indien graansorghum of 'n graansorghumprodukt in 'n houer verkoop of uitgevoer word soos in klousule 2 beoog word die massa van die betrokke houer vir die doeleindes van die berekening van die bedrag wat ten opsigte van 'n heffing en 'n spesiale heffing betaalbaar is, buite rekening gelaat: Met dien verstande dat die massa van 'n sak (goiing of jute) geag word 1 kg te wees.

DEPARTEMENT VAN FINANSIES

No. R. 517

15 Maart 1985

DOEANE- EN AKSYNSWET, 1964

WYSIGING VAN BYLAE 1 (No. 1/1/1096)

Kragtens artikel 48 van die Doeane- en Aksynswet, 1964, word Deel 1 van Bylae 1 by genoemde Wet hierby gewysig in die mate in die Bylae hiervan aangetoon.

K. D. S. DURR, Adjunk-minister van Finansies en van Handel en Nywerheid.

SCHEDULE

I Tariff Heading	II Statistical Unit	III Rate of Duty		IV
		General	M.F.N.	
84.06 By the substitution for subheading No. 84.06.99.40 of the following:				
..37 Pistons, machined, whether or not fitted with gudgeon pins, piston rings or cylinder liners or sleeves, with an outside diameter of less than 60 mm, for other motor vehicle engines		20%		
.39 Pistons, machined, whether or not fitted with gudgeon pins, piston rings or cylinder liners or sleeves, with an outside diameter of 60 mm or more but not exceeding 130 mm, for other motor vehicle engines (excluding those for motor cycle engines)		20% or 1 700c per kg less 80%		
.41 Pistons, machined, whether or not fitted with gudgeon pins, piston rings or cylinder liners or sleeves, with an outside diameter of more than 130 mm, for other motor vehicle engines		20%*		

Note.—The rate of duty on machined pistons for certain motor vehicle engines is amended.

BYLAE

I Tariefpos	II Statistiese Eenheid	III Skaal van Reg		IV
		Algemeen	M.B.N.	
84.06 Deur subpos No. 84.06.99.40 deur die volgende te vervang:				
..37 Suiers, gemasjineerd, hetsy met suierpenne, suierringe of silindervoerings of -hulse toegerus al dan nie, met 'n buitedeursnee van minder as 60 mm, vir ander motorvoertuigenjins		20%		
.39 Suiers, gemasjineerd, hetsy met suierpenne, suierringe of silindervoerings of -hulse toegerus al dan nie, met 'n buitedeursnee van minstens 60 mm maar hoogstens 130 mm, vir ander motorvoertuigenjins (uitgesonderd dié vir motorfietsenjins)		20% of 1 700c per kg min 80%		
.41 Suiers, gemasjineerd, hetsy met suierpenne, suierringe of silindervoerings of -hulse toegerus al dan nie, met 'n buitedeursnee van meer as 130 mm, vir ander motorvoertuigenjins		20%*		

Opmerking.—Die skaal van reg op gemasjineerde suiers vir sekere motorvoertuigenjins word gewysig.

No. R. 518

15 March 1985

CUSTOMS AND EXCISE ACT, 1964

AMENDMENT OF SCHEDULE 1 (No. 1/1/1097)

Under section 48 of the Customs and Excise Act, 1964, Part 1 of Schedule 1 to the said Act is hereby amended to the extent set out in the Schedule hereto.

K. D. S. DURR, Deputy Minister of Finance and of Trade and Industry.

No. R. 518

15 Maart 1985

DOEANE- EN AKSYNSWET, 1964

WYSIGING VAN BYLAE 1 (No. 1/1/1097)

Kragtens artikel 48 van die Doeane- en Aksynswet, 1964, word Deel 1 van Bylae 1 by genoemde Wet hierby gewysig in die mate in die Bylae hiervan aangetoon.

K. D. S. DURR, Adjunk-minister van Finansies en van Handel en Nywerheid.

SCHEDULE

I Tariff Heading	II Statistical Unit	III Rate of Duty		IV
		General	M.F.N.	
85.01 By the substitution for subheading No. 85.01.25.30 of the following: ".25 Exceeding 260 kW but not exceeding 5 600 kW	no.	5% or 700c per kg less 95%		
.35 Exceeding 5 600 kW	no.	5%"		

Note.—The rate of duty on certain three-phase electric motors, exceeding 260 kW but not exceeding 5 600 kW, is amended from 5% to 5% or 700c per kg less 95%.

BYLAE

I Tariefpos	II Statistiese Eenheid	III Skaal van Reg		IV
		Algemeen	M.B.N.	
85.01 Deur subpos No. 85.01.25.30 deur die volgende te vervang: ".25 Van meer as 260 kW maar hoogstens 5 600 kW	getal.	5% of 700c per kg min 95%		
.35 Van meer as 5 600 kW	getal.	5%"		

Opmerking.—Die skaal van reg op sekere driefasige elektriese motore van meer as 260 kW maar hoogstens 5 600 kW, word gewysig van 5% na 5% of 700c per kg min 95%.

No. R. 519

15 March 1985

CUSTOMS AND EXCISE ACT, 1964

AMENDMENT OF SCHEDULE 1 (No. 1/1/1098)

Under section 48 of the Customs and Excise Act, 1964, Part 1 of Schedule 1 to the said Act is hereby amended to the extent set out in the Schedule hereto.

K. D. S. DURR, Deputy Minister of Finance and of Trade and Industry.

No. R. 519

15 Maart 1985

DOEANE- EN AKSYNSWET, 1964

WYSIGING VAN BYLAE 1 (No. 1/1/1098)

Kragtens artikel 48 van die Doeane- en Aksynswet, 1964, word Deel 1 van Bylae 1 by genoemde Wet hierby gewysig in die mate in die Bylae hiervan aangetoon.

K. D. S. DURR, Adjunk-minister van Finansies en van Handel en Nywerheid.

SCHEDULE

I Tariff Heading	II Statistical Unit	III Rate of Duty		IV
		General	M.F.N.	
85.19 By the insertion after subheading No. 85.19.80.20 of the following: ".30 Thermo-electric relays incorporating bimetal elements .40 Other, with a value for duty purposes of 25 000c or more each	no. no.	5% 5%"		

Note.—Specific provision, at a rate of duty of 5%, is made for certain types of relays.

BYLAE

I Tariefpos	II Statistiese Eenheid	III Skaal van Reg		IV
		Algemeen	M.B.N.	
85.19 Deur na subpos No. 85.19.80.20 die volgende in te voeg: ".30 Termoëlektriese relê's wat bimetaalelemente inkorporeer .40 Ander, met 'n waarde vir belastingdoeleindes van minstens 25 000c elk	getal getal	5% 5%"		

Opmerking.—Spesifieke voorsiening, teen 'n skaal van reg van 5%, word gemaak vir sekere tipes relê's.

No. R. 525

15 March 1985

CUSTOMS AND EXCISE ACT, 1964

DETERMINATIONS OF TARIFF CLASSIFICATION AND FURNISHING THEREOF ON BILLS OF ENTRY (LIST TAR/121)

The following amendments to determinations are published in terms of section 47 (9) of the Customs and Excise Act, 1964 (Act 91 of 1964).

D. J. COLESKY, Commissioner for Customs and Excise.

Note.—List TAR/120 was published in Government Notice R. 499 of 8 March 1985.

No. R. 525

15 Maart 1985

DOEANE- EN AKSYNSWET, 1964

BEPALINGS VAN TARIEFINDELING EN VERSTREK-KING DAARVAN OP KLARINGSBRIEWE (LYS TAR/121)

Die volgende wysigings van bepalings word kragtens artikel 47 (9) van die Doeane- en Aksynswet, 1964 (Wet 91 van 1964), gepubliseer.

D. J. COLESKY, Kommissaris van Doeane en Aksyns.

Opmerking.—Lys TAR/120 is in Goewermentskennis-gewing R. 499 van 8 Maart 1985 gepubliseer.

AMENDMENTS TO PUBLISHED DETERMINATIONS

<i>Description of goods</i>	<i>Tariff heading/ subheading</i>	<i>Determination no.</i>
Amendments to determinations resulting from amendments to Part 1 of Schedule No. 1 to the Customs and Excise Act (Act 91 of 1964):		
The following are substituted for the existing determinations with effect from 15 March 1985:		
Frequency relays FTF 1123—relays	85.19.80	12
Siemens thermistor tripping units 3UN8 for protecting electric motors against undue temperature rises—relays	85.19.80	22
Siemens thermo flasher unit type RB109—a relay	85.19.80	85
Newman passive infra-red detector unit—a relay	85.19.80	104
Raytek Bispy models 3000 and 3001 infra-red intrusion detectors—relays	85.19.80	114
Ademco self-contained passive infra-red intrusion detectors models Nos. 650 and 651—relays	85.19.80	125
Unwin Doppler microwave detector M1-60—a relay	85.19.80	132
Foxboro 63U series electronic Consotrol difference alarm—a relay	85.19.80	133
Cerberus signal transmitter TUG 2—a relay	85.19.80	134
Cerberus signal receiving module TUP 2—a relay	85.19.80	135
MW 5002 and MW 5003 ultrasonic intrusion detectors not incorporating an alarm—relays	85.19.80	140
Det-Tronics ultra-violet fire detection systems—relays	85.19.80	146
RIS thermocouple alarm model ET-5204—a relay	85.19.80	152
B/W Controls induction type DH relay—a relay	85.19.80	153
Vactec slotted photo switches—relays	85.19.80	154
McGeoch WM 806 movement detector—a relay	85.19.80	159

WYSIGINGS VAN GEPUBLISEERDE BEPALINGS

<i>Beskrywing van goedere</i>	<i>Tariefpos/-subpos</i>	<i>Bepaling no.</i>
Wysigings van bepalings as gevolg van wysigings van Deel 1 van Bylae No. 1 by die Doeane- en Aksynswet (Wet 91 van 1964):		
Die volgende vervang die bestaande bepalings met ingang van 15 Maart 1985:		
Frekwensierelêrs FTF 1123—relêrs	85.19.80	12
Siemens-termister-uitklikeenhede 3UN8 om elektriese motore te beskerm teen oormatige temperatuurstygings—relêrs	85.19.80	22
Siemens-termoflikkereenheid tipe RB109—'n relê	85.19.80	85
Newman passiewe infrarooi verlikkerekereenheid—'n relê	85.19.80	104
Raytek Bispy modelle 3000 en 3001 infrarooi indringingsverlikkers—relêrs	85.19.80	114
Ademco selfstandige passiewe infrarooi indringingsverlikkers modelle Nos. 650 en 651—relêrs	85.19.80	125
Unwin Doppler-mikrogolfverlikker M1-60—'n relê	85.19.80	132
Foxboro 63U-reeks elektroniese Consotrol-differensie-alarm—'n relê	85.19.80	133
Cerberus-seintoestel TUG 2—'n relê	85.19.80	134
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DEPARTMENT OF HEALTH AND WELFARE

No. R. 516

15 March 1985

HUMAN TISSUE ACT, 1983 (ACT 65 OF 1983)

DRAFT REGULATIONS, CODE OF PRACTICE AND RELATED MATTERS REGARDING ARTIFICIAL INSEMINATION

1. The Minister of Health and Welfare intends—

(a) in terms of section 37 of the Human Tissue Act, 1983 (Act 65 of 1983), to make regulations regarding artificial insemination and related matters;

(b) to publish a Code of Practice as intended in section 22 of the said Act.

2. Interested persons are invited within three months of the date of publication of this notice to lodge—

(a) comments;

(b) representations regarding amendments, adjustments or additions pertaining to matters in the Annexures hereto to the Director-General: Health and Welfare, Private Bag X63, Pretoria, 0001, for attention of the Chief Director: Laboratory Services.

F. P. RETIEF, Director-General: Health and Welfare.

ANNEXURE 1

DRAFT REGULATIONS REGARDING THE ARTIFICIAL INSEMINATION OF PERSONS AND RELATED MATTERS

CHAPTER 1

INTERPRETATION AND APPLICATION OF REGULATIONS

Definition

1. In these regulations the "Act" means the Human Tissue Act, 1983 (Act 65 of 1983), and any expression has the same meaning as in the Act unless the context otherwise indicates; and "in vitro insemination" means the bringing together outside the human body of a male and female gamete or gametes with a view to placing the product of a union of such gametes in the womb of a female person;

"recipient" means woman—

(a) in who's internal reproductive organs a male gamete or gametes; or

(b) in who's womb the product of a union of a male and female gamete or gametes which was brought together outside the human body,

as the case may be, has been introduced or placed;

"donation" in regard to gametes means the process by which a gamete or gametes has/have been removed or extracted from a live person for the purpose of artificially inseminating another person;

"donor" in regard to gametes means a person from who's body a gamete or gametes has/have been removed or extracted for the purpose of artificial insemination of another person.

DEPARTEMENT VAN GESONDHEID EN WELSYN

No. R. 516

15 Maart 1985

WET OP MENSLIKE WEEFSEL, 1983 (WET 65 VAN 1983)

KONSEPREGULASIES, GEBRUIKSKODE EN AANVERWANTE AANGELEENTHEDE BETREFFENDE KUNSMATIGE BEVRUGTING

1. Die Minister van Gesondheid en Welsyn is voornemens om—

(a) kragtens artikel 37 van die Wet op Menslike Weefsel, 1983 (Wet 65 van 1983), regulasies betreffende kunsmatige bevrugting en aanverwante aangeleenthede uit te vaardig;

(b) 'n gebruikskode soos bedoel in artikel 22 van genoemde Wet te publiseer.

2. Belanghebbende persone word versoek om binne drie maande na die datum van publikasie van hierdie kennisgewing gemotiveerde—

(a) kommentaar;

(b) voorstelle betreffende aanpassings of byvoegings betreffende die aangeleenthede in die Bylaes hiervan aan die Direkteur-generaal: Gesondheid en Welsyn, Privaatsak X63, Pretoria, 0001, te stuur vir aandag van die Hoofdirekteur: Laboratoriumdienste.

F. P. RETIEF, Direkteur-generaal: Gesondheid en Welsyn.

BYLAE 1

KONSEPREGULASIES BETREFFENDE DIE KUNSMATIGE BEVRUGTING VAN PERSONE EN AANVERWANTE AANGELEENTHEDE

HOOFSTUK 1

UITLEG EN TOEPASSING VAN REGULASIES

Woordskrywing

1. In hierdie regulasies beteken die "Wet" die Wet op Menslike Weefsel, 1983 (Wet 65 van 1983), en het enige uitdrukking waaraan 'n betekenis in die Wet geheg is, daardie betekenis, en, tensy uit die samehang anders blyk, beteken—

"in vitro-bevrugting" die samevoeging buite die menslike liggaam van 'n manlike en 'n vroulike gameet of gamete met die oog op die plasing van die produk van 'n samesmelting van sodanige gamete in die baarmoeder van 'n vroulike persoon;

"ontvanger" 'n vrouspersoon—

(a) in wie se inwendige voortplantingsorgane 'n manlike gameet of gamete op 'n ander wyse as 'n natuurlike wyse; of

(b) in wie se baarmoeder die produk van 'n samesmelting van 'n manlike en 'n vroulike gameet of gamete wat buite die menslike liggaam saamgevoeg

na gelang van die geval, vir die doel van menslike voortplanting, ingevoer of geplaas is;

"skenking" met betrekking tot gamete, die proses waarvolgens 'n gameet of gamete vir die doeleindes van die kunsmatige bevrugting van 'n ander persoon, van die liggaam van 'n lewende persoon verwyder of onttrek word;

"skenker" met betrekking tot gamete, iemand van wie se liggaam 'n gameet of gamete vir doeleindes van die kunsmatige bevrugting van 'n ander persoon verwyder of onttrek word.

Application of regulations

2. These regulations, except regulation 10, are not applicable to the removal or extraction of a gamete or gametes from the body of a married man and the subsequent artificial insemination of the wife of such man with such gamete or gametes.

CHAPTER 2**DONORS, DONATIONS AND RELATED MATTERS***General provision*

3. Unless otherwise provided by these regulations or any other act, no person may remove or extract a gamete or gametes from the body of a living person for the purpose of artificial insemination unless the person from whose body such removal or extraction takes place, has been registered as a donor in terms of these regulations.

Application for registration as a donor

4. (1) (a) An application for the registration of a person as a donor of gametes for the purpose of artificial insemination consists of the following forms, examples of which specimen forms appear in Annexures A and B, compiled by the Director-General and supplied on request:

(i) An application form; and

(ii) An evaluation form.

(b) Such forms are completed in duplicate and—

(i) the original is submitted to the Director-General for consideration; and

(ii) the duplicate is filed in a personal donor file as intended in subregulation (6).

(2) The following medical practitioners and institutions may apply for the registration of a donor:

(a) A State controlled hospital by means of its medical superintendent or a medical practitioner in the service of the State nominated by such superintendent for this purpose;

(b) a hospital under control of a provincial administration by means of its medical superintendent or a medical practitioner in the service of such administration nominated by such superintendent for this purpose;

(c) a partnership of medical practitioners by means of any of the partners concerned;

(d) any sole medical practitioner in private practice;

(e) any other medical practitioner or institution with a medical practitioner in charge and to whom the Director-General, on request, has granted permission to apply for the registration of a donor.

(3) An intended donor in respect of whom registration as a donor is sought, shall, in the presence of the medical practitioner intended in subregulation (2), furnish, date and sign the following particulars and statement, as the case may be, in the evaluation form intended in subregulation (1) (a) (ii):

(a) The intended donor's—

(i) full names, surname, date of birth and identity number;

(ii) other personal particulars such as height, mass, colour of hair and eyes, type of build, population group, sex, marital status, religion and denomination;

(iii) family history;

(iv) replies to other relevant questions with regard to his selection as an appropriate and suitable donor of gametes.

Toepassing van regulasies

2. Hierdie regulasies, behalwe regulasie 10, is nie van toepassing op die verwydering of onttrekking van 'n gameet of gamete van die liggaam van 'n getroude manlike persoon en die daaropvolgende kunsmatige bevrugting van die eegnote van sodanige man met sodanige gameet of gamete nie.

HOOFSTUK 2**SKENKERS, SKENKINGS EN AANVERWANTE AANGELEENTHEDE***Algemene voorbehoud*

3. Tensy, hierdie regulasies of enige ander wet anders bepaal mag niemand 'n gameet of gamete van die liggaam van 'n lewende persoon vir die doeleindes van die kunsmatige bevrugting van 'n persoon verwyder of onttrek nie, tensy die persoon van wie se liggaam sodanige verwydering of onttrekking geskied, as 'n skenker ingevolge hierdie regulasies geregistreer is.

Aansoek om registrasie as 'n skenker

4. (1) (a) 'n Aansoek om die registrasie van 'n persoon as skenker van gamete vir doeleindes van kunsmatige bevrugting bestaan uit die volgende vorms waarvan voorbeeldvorme in Aanhangsels A en B verskyn en wat deur die Direkteur-generaal opgestel en op aanvraag voorsien word:

(i) 'n Aansoekvorm; en

(ii) 'n Evaluasievorm.

(b) Sodanige vorms word minstens in duplikaat ingevul en—

(i) die oorspronklike word na die Direkteur-generaal vir oorweging gestuur; en

(ii) die duplikaat word in 'n persoonlike skenkers lêer, soos bedoel in subregulasie (6) geliasseer.

(2) Die volgende geneeshere en instansies mag aansoek doen vir die registrasie van 'n skenker:

(a) 'n Staatsbeheerde hospitaal deur middel van sy Mediese Superintendent of 'n geneesheer in diens van die Staat wat deur sodanige superintendent vir die doel aangewys is;

(b) 'n hospitaal onder beheer van 'n provinsiale administrasie deur middel van sy Mediese Superintendent of geneesheer in diens van sodanige administrasie wat deur sodanige superintendent vir die doel aangewys is;

(c) 'n vennootskap van geneeshere deur middel van enige van die betrokke vennote;

(d) enige enkel geneesheer wat in privaatpraktyk staan;

(e) enige ander geneesheer of instansie wat 'n geneesheer in beheer het en aan wie die Direkteur-generaal, op aansoek, toestemming verleen het om aansoek om die registrasie van 'n skenker te doen.

(3) 'n Voornemende skenker ten opsigte van wie registrasie as skenker beoog word moet in teenwoordigheid van die betrokke geneesheer bedoel in subregulasie (2) die volgende besonderhede en verklaring, na gelang van die geval in die evaluasievorm bedoel in subregulasie (1) (a) (ii) verstrek, dateer en onderteken:

(a) Die voornemende skenker se—

(i) volle naam, van, geboortedatum en identifikasienommer;

(ii) ander persoonlike besonderhede soos lengte, massa, kleur van hare en oë, tipe liggaamsbou, bevolkingsgroep, geslag, huwelikstaat, geloof en kerkverband;

(iii) familiegeskiedenis;

(iv) antwoorde op ander tersaaklike vrae vir sover dit die keuring van hom as 'n geskikte en gepaste skenker van gamete betref.

(b) A written statement, incorporated in the form, in which the intended donor, for purposes of his evaluation as an appropriate and suitable donor, consents to—

(i) a physical examination and questioning of himself by a medical practitioner;

(ii) a further physical examination and questioning by a person appointed by such medical practitioner in consultation with the intended donor and the reporting back to such medical practitioner by such person;

(iii) the withdrawal of blood samples and the removal or withdrawal of a gamete or gametes from his body by or under supervision of such medical practitioner for the purpose of analysis, testing or other processing which the medical practitioner may deem necessary;

(iv) the reporting back to such medical practitioner of the results of such analysis, testing or processing in cases where the medical practitioner does not himself do the analysis, testing or processing;

(v) the consultation and deliberation amongst all persons mentioned in this regulation;

(vi) perusal of the personal particulars of the donor by the persons mentioned in subregulation (7);

(vii) the submission to the Director-general or any officer in the Department of Health and Welfare so delegated in terms of the Act, of the evaluation form intended in this regulation and other relevant information which the Director-General may require in terms of subregulation (5).

(4) (a) The medical practitioner intended in subregulation (2) or any other medical practitioner appointed in consultation with the intended donor by the former medical practitioner performs, for the purpose of evaluation of an intended donor as an appropriate and suitable person to serve as a donor of gametes on discretion or where the Director-General laid down specific guidelines, a medical examination of such intended donor and provides on the evaluation form intended in subregulation 1 (a) (ii) or as an annexure thereto—

(i) an indication whether diseases and disorders as intimated on the evaluation form by the intended donor do, according to the opinion of such medical practitioner, exist;

(ii) an indication whether diseases and disorders of forefathers and relatives as indicated on the evaluation form, to the best knowledge and opinion of such medical practitioner, exist or existed by the intended donor's forefathers or relatives;

(iii) a motivated report, founded on such medical practitioner's own observation and investigation of the intended donor as well as any relevant information the medical practitioner is aware of, in which the medical practitioner indicates whether the intended donor, to his best knowledge and opinion is an appropriate and suitable person to serve as donor.

(5) The Director-General, when considering the application for registration as a donor, at his discretion, may request further information and details he may require from the applying medical practitioner, the intended donor or any other person.

(6) The medical practitioner intended in subregulation 2 shall—

(a) open or arrange to be opened a separate personal donor file, to which he allocates a sequence number for each person he intends to register as a donor, in which all relevant documents regarding the person after registration as a donor, are filed;

(b) 'n Skriftelike verklaring, geïnkorporeer in die vorm, waarin die voornemende skenker vir doeleindes van die evaluering van homself as 'n geskikte en gepaste skenker, toestemming verleen tot—

(i) 'n liggaamlike ondersoek en ondervraging van homself deur 'n geneesheer;

(ii) 'n verdere liggaamlike ondersoek en ondervraging deur 'n persoon deur sodanige geneesheer in oorleg met die voornemende skenker aangewys en die terugrapportering aan sodanige geneesheer deur sodanige persoon;

(iii) die trek van bloedmonsters en die verwydering of onttrekking van 'n gameet of gamete uit sy liggaam deur of onder toesig van sodanige geneesheer vir doeleindes van ontleding, toetsing of ander prosessering wat die geneesheer nodig mag ag;

(iv) die terugrapportering aan sodanige geneesheer van die resultate van sodanige ontleding, toetsing of ander prosessering in gevalle waar die geneesheer nie self die ontleding, toetsing of ander prosessering doen nie;

(v) die konsultering en oorlegpleging tussen alle persone in hierdie regulasie genoem;

(vi) insae in sy persoonlike besonderhede as skenker deur die persone genoem in subregulasie (7).

(vii) die voorlegging aan die Direkteur-generaal of enige beamppte in die Departement van Gesondheid en Welsyn as sy gedelegeerde ingevolge die Wet, van die evaluasievorm bedoel in hierdie regulasie en ander tersaaklike inligting wat die Direkteur-generaal ingevolge subregulasie (5) mag verlang.

(4) (a) Die geneesheer bedoel in subregulasie (2) of 'n ander geneesheer wat in oorleg met die voornemende skenker deur eersgenoemde geneesheer aangewys is, doen, vir doeleindes van evaluasie van 'n voornemende skenker as 'n geskikte en gepaste persoon om as 'n skenker van gamete te dien na goeddunke of waar die Direkteur-generaal spesifieke riglyne betreffende sodanige ondersoek neergelê het met behoorlike in agneming van sodanige riglyne, 'n geneeskundige ondersoek van sodanige voornemende skenker en verstreke op die evaluasievorm bedoel in subregulasie (1) (a) (ii) of as 'n aanhangsel daarby—

(i) 'n aanduiding of siektetoestande en afwykings soos op die evaluasievorm uiteengesit by die voornemende skenker na die mening van sodanige geneesheer bestaan;

(ii) 'n aanduiding of siektetoestande en afwykings by die voornemende skenker se voorouers of familieleden, soos op die evaluasievorm uiteengesit, na die beste van sodanige geneesheer se wete en oortuiging bestaan of bestaan het;

(iii) 'n gemotiveerde verslag, gegrond op sodanige geneesheer se eie waarneming en ondersoek van die voornemende skenker sowel as enige tersaaklike inligting wat die geneesheer bewus van is, waarin die geneesheer aandui of die voornemende skenker na so 'n geneesheer se beste wete en oortuiging 'n geskikte en gepaste persoon is om as skenker te dien.

(5) Die Direkteur-generaal kan na goeddunke by die ooreweging van 'n aansoek om registrasie as skenker die verdere inligting en besonderhede wat hy nodig ag van die aansoekdoende geneesheer, die voornemende skenker of enige ander persoon aanvra.

(6) Die geneesheer bedoel in subregulasie (2) moet—

(a) 'n aparte persoonlike skenkerslêer, waaraan hy 'n volgnommer toeken, vir elke persoon wat hy beoog om as skenker te registreer oopmaak of laat oopmaak waarin alle tersaaklike stukke betreffende die persoon na registrasie as 'n skenker, geliasseer word;

(b) open or arrange to be opened a donor register in which particulars of every extraction or removal of gametes from bodies of all donors in respect of whom he has made application for registration as a donor and is registered, shall be noted as well as particulars of every artificial insemination of a person with such gametes conforming to these regulations.

(7) Except for the provisions of article 33, the medical practitioner intended in subregulation (2) may—

(a) make a donors register available for perusal to—

(i) a medical practitioner who intends to—

(aa) withdraw or remove a gamete or gametes from the donor concerned or to supervise such withdrawal or removal; or

(bb) artificially inseminate a person with a gamete or gametes which has been removed from such donor or to supervise such insemination;

(ii) an inspector of anatomy; or

(iii) to a person who has been authorised in writing by the Director-General;

(iv) a court of law;

(v) a person acting in accordance with a search warrant issue in terms of the Criminal Procedure Act of 1977;

(b) make available for perusal a donors register to any person or a court of law as stated in paragraph (a) (ii) (iii) (iv) and (v).

(8) A donor file or donor register—

(a) must, when not in use, be kept in a safe, or where a safe is not available, such file or register be protected against fire, theft, unauthorised perusal or destruction.

(b) may not be destroyed or disposed of unless so determined by the Director-General.

Issuing and cancellation of a registration certificate of a donor

5. (1) Where the Director-General, after consideration of an application for registration as a donor together with any further information as intended in regulation 4 (5), has determined that such intended donor be suitable and appropriate to serve as a donor, the Director-General will register such intended donor as a donor of gametes. A registration Certificate in a format determined by the Director-General, an example of which appears in Annexure C, will be issued to the medical practitioner, hospital or institution, as the case may be wherein—

(a) he states that the intended donor concerned is registered as a donor for the purposes of artificial insemination of another person; and

(b) that he has noted all relevant information pertaining to the intended donor with regard to artificial insemination.

(2) The Director-General—

(a) must cancel such registration as soon as he is aware that an artificial insemination or inseminations have resulted in five live births.

(b) may at any time cancel such registration should he consider that valid reasons exist for such cancellation and after such cancellation,

to immediately inform the medical practitioner or authority intended in regulation (4) (2) by registered post of such cancellation.

(b) 'n skenkersregister oopmaak of laat oopmaak waarin besonderhede van iedere onttrekking of verwydering van gamete uit die liggame van alle skenkers ten opsigte waarvan hy aansoek om registrasie gedoen het en wat geregistreer is aanteken of laat aanteken en waarin ook besonderhede van iedere kunsmatige bevrugting van 'n persoon met sodanige gamete ooreenkomstig hierdie regulasies aangeteken word.

(7) Behoudens die bepalings van artikel 33 mag die geneesheer bedoel in subregulasie (2)—

(a) 'n skenkerslêer vir insae beskikbaar stel aan—

(i) 'n geneesheer wat beoog om—

(aa) 'n gameet of gamete van die liggaam van die betrokke skenker te onttrek of te verwyder of toeges oor sodanige onttrekking of verwydering te hou; of

(bb) 'n kunsmatige bevrugting van 'n persoon met 'n gameet of gamete wat van sodanige skenker onttrek of verwyder is doen of toeges oor sodanige bevrugting hou;

(ii) 'n inspekteur van anatomie; of

(iii) 'n persoon deur die Direkteur-generaal skriftelik daartoe gemagtig;

(iv) 'n gereghshof;

(v) iemand behandelende kragtens 'n visiteringslasbrief uitgereik kragtens die Strafproseswet, 1977;

(b) 'n skenkersregister vir insae beskikbaar stel aan persone en 'n gereghshof soos genoem in paragraaf (a) (ii) (iii) (iv) en (v).

(8) 'n Skenkerslêer en skenkersregister—

(a) word deur die geneesheer bedoel in subregulasie (2), wanneer dit nie in gebruik is nie in 'n brandkluis bewaar of, waar 'n brandkluis nie beskikbaar is nie, op 'n ander veilige plek waar sodanige lêer of register beskerm is teen brand, diefstal, ongemagtigde insae of vernietiging;

(b) mag nie vernietig of meer weggedoen word nie behalwe soos deur die Direkteur-generaal bepaal.

Uitreiking en intrekking van registrasiesertifikaat van 'n skenker

5. (1) Waar die Direkteur-generaal, na oorweging van 'n aansoek om registrasie as skenker en enige bykomende inligting bedoel in regulasie 4 (5), hom vergewis het dat die betrokke voornemende skenker 'n geskikte en gepaste persoon is om as skenker te dien registreer die Direkteur-generaal sodanige voornemende skenker as 'n skenker van gamete en reik 'n registrasiesertifikaat in 'n vorm deur die Direkteur-generaal bepaal en waarvan 'n voorbeeldvorm in Aanhangsel C verskyn, aan die aansoekdoende geneesheer, hospitaal of inrigting na gelang van die geval, uit waarin—

(a) hy verklaar dat die betrokke voornemende skenker as 'n skenker vir doeleindes van die kunsmatige bevrugting van 'n ander persoon geregistreer is; en

(b) hy alle tersaaklike inligting wat op die voornemende skenker vir doeleindes van kunsmatige bevrugting betrekking het, aanstip.

(2) Die Direkteur-generaal—

(a) moet sodanige registrasie intrek sodra dit tot sy aandag kom dat 'n kunsmatige bevrugting of bevrugtings gelei het tot vyf lewendige geboortes;

(b) kan sodanige registrasie te eniger tyd intrek indien daar na sy mening goeie redes daarvoor bestaan,

en gee onverwyld nadat hy sodanige intrekking gedoen het die geneesheer of instansie bedoel in regulasie (4) (2) per geregistreerde pos kennis van sodanige intrekking.

(3) The medical practitioner or authority to whom notice has been given in terms of subregulation (2) must—

(a) acknowledge receipt of the notice of cancellation by registered post within three weeks of the date of the notice of cancellation.

(b) immediately make note in the donors register intended in regulation 4 (6) to the effect that registration has been cancelled.

(c) by the words—

“registration cancelled by the Director-General of Health and Welfare’s notice dated” indicate such in red ink on the front of the registration certificate concerned.

(d) inform the donor of such withdrawal; and

(e) immediately—

(i) destroy all gametes of the particular donor still in his possession and inform the Director-General in writing of the date and method of destruction, or

(ii) where he wishes to keep such gametes, to submit motivated representation to the Director-General for the keeping of such gametes,

and to file the registration certificate concerned and the notice of withdrawal in the donors file as intended in regulation 4 (b).

Conditions applicable to donations of gametes

6. (1) (a) No person, excepting a medical practitioner or a person acting under his supervision, may withdraw or remove a gamete or gametes from a donor for the purposes of artificial insemination.

(b) Such donor must at the time of such donation be registered as a donor in terms of these regulations.

(2) Such medical practitioner must, before attending to a donation of a gamete or gametes must—

(a) himself ascertain that—

(i) the intended donor is registered as such in terms of these regulations and that such registration is still noted on the date of donation;

(ii) The donor’s file intended in regulation 4 (6) is available for his inspection and perusal;

(b) ensure that a donor’s form of which an example is shown in Annexure D is available for the intended donation.

(c) ensure that the intended donor concerned submits to—

(i) medical tests for hereditary diseases

(ii) a Sperm analysis in the case of male donor; or

(iii) a gynecological examination in the case of a female donor;

(d) record the results of the tests and examination mentioned in paragraph (c) on the donors form and keep such form in the donor’s file.

(3) Such medical practitioner must immediately after attending to a donation of a gamete or gametes, record at least the following details in the donor’s register intended in regulation 4 (6):

(i) Serial number of the case;

(ii) donor’s name, address, sex, identity number and number of the donor registration certificate;

(3) Die geneesheer of instansie aan wie kennis ingevolge subregulasie (2) gegee is moet—

(a) ontvangs van die kennisgewing van intrekking per geregistreerde pos erken binne drie weke na die datum van kennisgewing;

(b) onverwyld ’n aantekening in die skenkersregister bedoel in regulasie 4 (6) aanbring te dien effekte dat die registrasie van die betrokke skenker ingetrek is;

(c) die woorde—

“registrasie ingetrek by Direkteur-generaal van Gesondheid en Welsyn se kennisgewing gedateer” in rooi ink op die voorkant van die betrokke registrasiesertifikaat aangebring.

(d) die skenker van die intrekking in kennis stel; en

(e) dadelik—

(i) alle gamete van die bepaalde skenker wat hy nog in sy besit het vernietig en die Direkteur-generaal skriftelik in kennis stel van die datum en metode van vernietiging; of

(ii) waar hy sodanige gamete nog wil behou gemotiveerde versoë vir die behoud daarvan aan die Direkteur-generaal rig,

en die betrokke registrasiesertifikaat en kennisgewing van intrekking in die skenkerslêer bedoel in regulasie 4 (6) liasseer.

Voorwaardes van toepassing by skenkings van gamete

6. (1) (a) Niemand, behalwe ’n geneesheer of iemand wat onder sy toesig optree, mag ’n gameet of gamete vir doeleindes van die kunsmatige bevrugting van ’n persoon van ’n skenker verwyder of onttrek nie.

(b) Sodanige skenker moet tydens sodanige skenking as ’n skenker kragtens hierdie regulasies geregistreer wees.

(2) Sodanige geneesheer moet voordat hy ’n skenking van ’n gameet of gamete waarneem—

(a) homself vergewis dat—

(i) die voornemende skenker wel as skenker kragtens hierdie regulasies geregistreer is en dat registrasie op die datum van skenking nog van krag is;

(ii) die skenkingslêer bedoel in regulasie 4 (6) vir sy insae en bestudering beskikbaar is;

(b) toesien dat ’n skenkingsvorm waarvan ’n voorbeeld in Aanhangsel D verskyn vir invulling by die beoogde skenking beskikbaar is;

(c) toesien dat die betrokke voornemende skenker—

(i) geneeskundige toetse vir geslagsoordraagbare siektes; en

(ii) ’n spermanalise in die geval van ’n manlike skenker; of

(iii) ’n ginekologiese ondersoek in die geval van ’n vroulike skenker ondergaan nie langer as sewe dae voor die voorgenome skenking;

(d) die uitslag van die toetse en ondersoek in paragraaf (c) genoem op die skenkingsvorm aanteken en die vorm in die skenker se skenkerslêer liasseer.

(3) Sodanige geneesheer moet direk nadat hy ’n skenking van ’n gameet of gamete waargeneem het minstens die volgende besonderhede in die skenkersregister bedoel in regulasie 4 (6) aanteken:

(i) Volgnommer van geval;

(ii) skenker se naam, adres, geslag, identifikasienommer en nommer van registrasiesertifikaat as skenker;

(iii) note whether a gamete or gametes obtained at the time of donation—

(aa) will be used once for the artificial insemination of a person; or

(bb) whether a portion will be used in the case of gametes, for such insemination with the remainder being stored for further insemination;

(iv) the date of the donation concerned.

CHAPTER 3

ARTIFICIAL INSEMINATIONS

Conditions concerning the artificial insemination of a person

7. (1) An artificial insemination of a person may not be done except in accordance with the conditions of these regulations.

(2) No gamete, which has not originated from a registered donor, may be used for the artificial insemination of a person.

(3) A medical practitioner who intends to perform an artificial insemination on a person, must inform the Director-General in writing prior to such insemination, of—

(a) his intentions to perform such insemination;

(b) the registration number of the intended donor; and

(c) the name and address and identity number (if any) of the intended recipient and in the case of a married woman, her maiden name, on a form drafted by the Director-General of which an example is shown in Annexure E.

(4) A medical practitioner may not perform an artificial insemination of a person or supervise such insemination unless he has ascertained—

(a) that the intended insemination concerned will not lead to the simultaneous existence of more than two artificially produced pregnancies originated from the use of a gamete or gametes originating from the same donor; and

(b) that there are no more than three living births in existence as a result of artificial insemination with such gametes at that time.

Conditions concerning recipients

8. (1) A medical practitioner who intends to carry out an artificial insemination must complete an evaluation form in respect of such person, an example of which appears in Annexure F.

(2) The conditions of regulation 4 (3) are *mutatis mutandis* applicable to the evaluation form referred to in subregulation (1).

(3) The medical practitioner intended in subregulation (1) must fully inform the intended recipient in respect of the intended donor, notwithstanding any other information he might deem necessary to impart to the recipient, all personal details of the intended donor excepting, in the case where such donor has not given written consent that his identity may be revealed to the recipient, any such information that will enable such recipient to identify the donor concerned should the recipient so request.

(iii) notering of 'n gameet of gamete wat tydens die skenking verkry is—

(aa) eenmalig by 'n kunsmatige bevrugting van 'n persoon gebruik sal word; of

(bb) gedeeltelik in die geval van gamete by sodanige bevrugting gebruik sal word met die res bewaar vir 'n toekomstige bevrugting;

(iv) die datum van die betrokke skenking.

HOOFSTUK 3

KUNSMATIGE BEVRUGTINGS

Voorwaardes betreffende die kunsmatige bevrugting van 'n persoon

7. (1) 'n Kunsmatige bevrugting van 'n persoon word nie gedoen behalwe ooreenkomstig die voorwaardes in hierdie regulasie vervat.

(2) Geen gameet wat nie van die liggaam van 'n geregistreerde skenker afkomstig is, mag vir die kunsmatige bevrugting van 'n persoon gebruik word nie.

(3) 'n Geneesheer wat beoog om 'n kunsmatige bevrugting van 'n persoon te doen moet voor sodanige bevrugting die Direkteur-generaal skriftelik kennis gee—

(a) van sy voorneme om sodanige bevrugting te doen;

(b) van die registrasienommer van die beoogde skenker; en

(c) van die naam en adres en identifikasienommer (indien enige) van die voornemende ontvanger met vermelding in die geval van 'n getroude vrou van haar nooiensvan, op 'n vorm deur die Direkteur-generaal opgestel en waarvan 'n voorbeeldvorm 'n Aanhangsel E verskyn.

(4) 'n Geneesheer mag nie 'n kunsmatige bevrugting van 'n persoon doen of toetsig oor sodanige bevrugting hou nie tensy hy hom vergewis het—

(a) dat die betrokke beoogde bevrugting nie kan lei nie tot die gelyktydige bestaan van meer as twee kunsmatige verwekte swangerskappe afkomstig van die gebruik van 'n gameet of gamete afkomstig van een en dieselfde skenker; en

(b) dat daar nie alreeds op daardie tydstip meer as drie lewendige geboortes as gevolg van kunsmatige bevrugtings met sodanige gamete verwek is.

Bepalings betreffende ontvangers

8. (1) 'n Geneesheer wat beoog om 'n kunsmatige bevrugting van 'n persoon te doen moet 'n evaluasievorm waarvan 'n voorbeeldvorm in Aanhangsel F verskyn t.o.v. sodanige persoon opstel.

(2) Die bepalinge van regulasie 4 (3) is *mutatis mutandis* van toepassing op 'n evaluasievorm in subregulasie (1) genoem.

(3) Die geneesheer bedoel in subregulasie (1) moet ten einde die voornemende ontvanger volledig in te lig aangaande die beoogde skenker, afgesien van enige ander inligting wat hy nodig mag ag om aan die ontvanger mee te deel, die volledige persoonlik besonderhede van die beoogde skenker behalwe, in die geval waar sodanige skenker nie skriftelik magtiging verleen het dat sy identiteit aan die ontvanger bekend gemaak mag word, enige sodanige besonderhede wat sodanige ontvanger in staat sal stel om die betrokke skenker te identifiseer, aan die beoogde ontvanger beskikbaar stel, indien daartoe deur die ontvanger versoek.

Specific conditions concerning in vitro inseminations

9. (1) An vitro insemination and the placing of the product of such union of gametes resultant of such insemination may only be carried out—

(a) in or at premises approved by the Director-General; and

(b) subject to the conditions laid down by the Director-General.

(2) (a) A medical practitioner who intends to perform an in vitro insemination must first apply to the Director-General for approval and as part of the application must submit—

(i) full details and address of the premises where the intended in vitro insemination is to take place;

(ii) a brief summary of the procedures to be followed at such insemination.

(b) The Director-General, in considering such approval may, at his discretion require—

(i) further details from the applicant or any other person;

(ii) an inspection of the intended premises by an inspector of anatomy.

Reporting of details of artificially produced infants

10. (1) A medical practitioner who has performed or supervised an artificial insemination on a person and has attended a live birth resulting from such insemination or, in pursuance of subregulation (2) has had details of such birth reported to him, must within 30 days of such birth or on such report of a birth inform the Director-General in writing of—

(a) the name and surname of the child concerned;

(b) the date of birth;

(c) the name of the mother;

(d) the name, surname and registration number of the donor used for the insemination concerned; and

(e) any physical or mental abnormality of such child.

(2) The mother of the child intended in subregulation (1) must, in the event of the medical practitioner intended in that subregulation not attending the birth, inform the medical practitioner in writing of the details stated in subregulation (1) (a) to (e) inclusive within 30 days of such birth.

CHAPTER 4**GENERAL AND SUPPLEMENTARY CONDITIONS***Disclosure of details in respect of an artificial insemination*

11. (1) A medical practitioner or any other person who or was concerned with an artificial insemination of a person may not disclose any fact to another person whereby the identity of—

(a) a child conceived by artificial insemination; or

(b) a recipient can possibly be established, unless—

(i) a court of law rules otherwise;

(ii) a law determines otherwise;

Spesifieke bepalings betreffende in vitro-bevrugtings

9. (1) 'n In vitro-bevrugting en die plasing van die produk van die samesmelting van gamete wat uit sodanige bevrugting ontstaan het, mag net gedoen word—

(a) in of op 'n perseel deur die Direkteur-generaal goedgekeur; en

(b) onderworpe aan voorwaardes deur die Direkteur-generaal bepaal.

(2) (a) 'n Geneesheer wat beoog om 'n in vitro-bevrugting te doen moet vooraf by die Direkteur-generaal om goedkeuring aansoek doen en as deel van sy aansoek—

(i) volledige besonderhede en straatadres van die beoogde perseel waarin of waarop die in vitro-bevrugting beoog word;

(ii) 'n beknopte oorsig van prosedure wat by sodanige bevrugting gevolg sal word, aan die Direkteur-generaal voorlê.

(b) Die Direkteur-generaal kan in die oorweging van sodanige goedkeuring na goeddunke—

(i) verdere besonderhede van die aansoeker of enige ander persoon;

(ii) 'n inspeksie van die beoogde persele deur 'n inspekteur van anatomie, vereis.

Rapportering van besonderhede van kunsmatige verwekte babas

10. (1) Die geneesheer wat 'n kunsmatige bevrugting van 'n persoon gedoen het of onder wie se toesig sodanige bevrugting gedoen was, moet waar hy 'n lewendige geboorte wat die gevolg van sodanige bevrugting is waargeneem het of waar besonderhede van sodanige geboorte ingevolge subregulasie (2) aan sodanige geneesheer gerapporteer is, binne 30 dae na sodanige geboorte of sodanige rapportering, na gelang van die geval, die Direkteur-generaal skriftelik in kennis stel van—

(a) die naam en van van die betrokke kind;

(b) die datum van geboorte;

(c) die naam en van van die moeder;

(d) die naam, van en registrasienommer van die skenker wat by die betrokke bevrugting gebruik is; en

(e) enige liggaamlike of geestesafwyking by sodanige kind.

(2) Die moeder van 'n kind bedoel in subregulasie (1) moet in die geval waar die geboorte van die kind nie deur die geneesheer bedoel in daardie subregulasie waargeneem is nie, die besonderhede genoem in subregulasie (1) (a) tot en met (e) skriftelik binne 30 dae na sodanige geboorte aan daardie geneesheer rapporteer.

HOOFSTUK 4**ALGEMENE EN AANVULLENDE BEPALINGS***Openbaarmaking van besonderhede ten opsigte van 'n kunsmatige bevrugting*

11. (1) 'n Geneesheer of enige ander persoon wat betrokke is of was by die kunsmatige bevrugting van 'n persoon mag nie aan 'n ander persoon 'n feit publiseer waardeur die identiteit van—

(a) 'n kind wat deur kunsmatige bevrugting verwek is; of

(b) 'n ontvanger,

moontlik vasgestel kan word nie, tensy—

(i) 'n geregshof anders gelas;

(ii) 'n wet anders bepaal;

(iii) in the case of a recipient or an artificially conceived child who has legally come of age and is still living as the time of such disclosure, that the recipient or the child who has come of age, as the case may be, has given written consent to such disclosures;

(iv) in the case of an artificially conceived child who has come of age, or a recipient who is deceased at the time of such disclosures—

(aa) that the artificially conceived child of legal age or the recipient as the case may be, has given written consent for such disclosure;

(bb) that the child of legal age or the recipient as the case may be, prior to death, has not in any manner indicated that such consent would not be given and that the spouse, legally of age offspring, parent guardian, legally of age brother or sister of the certifiably conceived child or recipient as the case may be, has given written consent prior to such disclosure.

(2) The Director-General, may subject to the conditions of section 33, request in writing from—

(a) a child conceived by artificial insemination;

(b) a recipient;

(c) the spouse of the recipient,

should he be convinced that the interest of the persons mentioned in paragraphs (a), (b) and (c) will not be harmed, to make available to a named applicant information regarding a donor with regard to a child conceived by artificial insemination.

Reporting of defects

12. (1) A medical practitioner who has performed an artificial insemination on a person or has supervised such insemination must, should it come to his notice that as a result of such insemination, a child displays genetic or psychiatric defects—

(a) carry out or have carried out tests to determine whether the cause of the defects concerned can be attributed to the donor or recipient; and

(b) inform the Director-General in writing of the defect, the tests carried out, the results of such tests and his comments on the matter and to further act in accordance with the Director-General's directives.

(2) A parent of a child intended in subregulation (1) must, where it has come to his notice that such child displays abnormal genetic or psychiatric defects, report such defect to—

(a) the medical practitioner intended in subregulation (1); or

(b) the Director-General if that medical practitioner does not, to the best knowledge and satisfaction bear any knowledge of that defect.

Code of practice

13. The code of practice intended section 22 of the Act in terms of which the person who performs the artificial insemination of a person is the code of practice as published by Government Notice of 19.....

Penalties

14. A person who transgresses a clause of these regulations or fails to adhere thereto shall be guilty of an offence and liable, on conviction to a fine not exceeding R1 000 or to imprisonment for a period not exceeding six months.

(iii) in die geval van 'n ontvanger of meerderjarige kunsmatige verwekte kind wat ten tyde van sodanige publikasie nog in die lewe is, daardie ontvanger of meerderjarige kind na gelang van die geval, voor sodanige publikasie skriftelik toestemming daartoe verleen het;

(iv) in die geval van 'n meerderjarige kunsmatige verwekte kind of ontvanger wat ten tyde van sodanige publikasie oorlede is—

(aa) daardie meerderjarige kunsmatige verwekte kind of ontvanger, so gelang van die geval, voor sy dood skriftelike toestemming tot sodanige publikasie verleen het;

(bb) daardie meerderjarige kind of ontvanger, so gelang van die geval nie voor sy dood op enige wyse aangedui het dat hy nie bereid sou wees om sodanige toestemming te verleen nie en die meerderjarige kind of ontvanger na gelang van die geval se eggenoot, meerderjarige kind, ouer, voog, meerderjarige broer of meerderjarige suster voor sodanige publikasie skriftelike toestemming daartoe verleen het.

(2) Die Direkteur-generaal kan behoudens die bepaling van artikel 33 op skriftelike versoek van—

(a) 'n kind wat deur kunsmatige bevrugting verwek is;

(b) 'n ontvanger;

(c) 'n eggenoot van 'n ontvanger,

indien hy oortuig is dat die belange van persone genoem in paragrawe (a), (b) en (c) nie benadeel sal word nie, besonderhede aangaande die skenker in sover dit betrekking het op die kind wat deur kunsmatige bevrugting verwek is, aan 'n genoemde aansoeker bekend maak.

Rapportering van defekte

12. (1) 'n Geneesheer wat 'n kunsmatige bevrugting van 'n persoon gedoen het of onder wie se toesig sodanige bevrugting gedoen is moet, indien dit onder sy aandag kom dat 'n kind wat as gevolg van sodanige bevrugting verwek is enige abnormale genetiese of psigiatrisiese afwyking toon—

(a) toetse uitvoer of laat uitvoer ten einde vas te stel of die oorsaak van die betrokke afwyking na die skenker of ontvanger herlei kan word; en

(b) die betrokke afwyking, die toetse wat gedoen is, die resultate van sodanige toetse en sy kommentaar oor die aangeleentheid skriftelik aan die Direkteur-generaal rapporteer en verder optree soos deur die Direkteur-generaal gelas.

(2) 'n Ouer van 'n kind bedoel in subregulasie (1) moet waar dit onder sy aandag kom dat sodanige kind enige abnormale genetiese of psigiatrisiese afwyking toon sodanige afwyking aan—

(a) die geneesheer bedoel in subregulasie (1); of

(b) die Direkteur-generaal rapporteer indien daardie geneesheer na die beste van daardie ouer se wete en oortuiging nie kennis van daardie afwyking dra nie.

Gebruikskode

13. Die gebruikskode bedoel in artikel 22 van die Wet waarvolgens die persoon wat die kunsmatige bevrugting van 'n persoon doen moet optree is die gebruikskode soos gepubliseer by Goewermentskennisgewing van 19.....

Strafbepalings

14. Iemand wat 'n bepaling van hierdie regulasies oortree of versuim om daaraan te voldoen is aan 'n misdryf skuldig en by skuldigbevinding strafbaar met 'n boete van hoogstens R1 000 of met gevangenisstraf van hoogstens ses maande.

ANNEXURE 2

DEPARTMENT OF HEALTH AND WELFARE

HUMAN TISSUE ACT, 1983 (ACT 65 OF 1983)

CODE OF PRACTICE IN TERMS OF SECTION 22 OF THE HUMAN TISSUE ACT, 1983, ACCORDING TO WHICH A PERSON, WHO PERFORMS AN ARTIFICIAL INSEMINATION ON A PERSON, SHALL ACT

1. In this Annexure, "the Act" means the Human Tissue Act, 1983 (Act 65 of 1983), and any expression has the same meaning as in the Act and the regulations regarding to artificial insemination promulgated by Government Notice R....., dated

2. Save for the provisions of the Act and the regulations, a gamete removed from the body of a live person shall not be used for the artificial insemination of another person unless—

(a) the medical practitioner who performs such artificial insemination or under who's supervision the insemination is performed, separately or together with a registered geneticist, if available, shall—

(i) give guidance and information to the recipient and her spouse concerning—

(aa) the possibilities of a normal pregnancy;

(bb) all implications of artificial insemination including problems pertaining to the technique of artificial insemination, chances of success, financial aspects, implications for the marriage, psycho-sociological—and educational implications, risks regarding genetic origins and prognosis of the child as well as legal advice obtainable regarding artificial insemination;

(ii) establish that the recipient biologically, physically and mentally is suitable for artificial insemination;

(ii) respect the wishes of the recipient and the spouse, with the regard to the donor's population group and religion;

(iv) where the recipient or the donor originates from a population group where the individual constitutes a high risk to be a genetic carrier (recessive) of a specific defect, e.g. Tay-Sachs, thalassaemia or cystic fibrosis, arrange for the recipient and the donor be tested for the characteristics concerned and not to utilise the gamete of a donor with the same characteristics as that of the recipient for the purposes of artificial insemination;

(v) where, on the grounds of family history a possibility exists that the recipient or the donor be a carrier of a defect which may be transferred by genes, have an examination and tests carried out by a registered geneticist to establish if the recipient or the donor of the defect, as the case may be—

(aa) arrange that the recipient and her spouse be given genetic counselling where it has been established that the recipient be a carrier of the defect concerned; or

(bb) that the gametes of the donor not be used for purposes of artificial insemination where it has been established that the donor be a carrier of the defect or the donor be at high risk as a carrier of the defect.

BYLAE 2

DEPARTEMENT VAN GESONDHEID EN WELSYN

WET OP MENSLIKE WEEFSEL, 1983 (WET 65 VAN 1983)

GEBRUIKSKODE BEDOEL IN ARTIKEL 22 VAN DIE WET OP MENSLIKE WEEFSEL, 1983, WAARVOLGENS 'N PERSOON WAT 'N KUNSMATIGE BEVRUGTING VAN 'N PERSOON DOEN, MOET OPTREE

1. In hierdie Bylae beteken "die Wet" die Wet op Menslike Weefsel, 1983 (Wet 65 van 1983), en het enige uitdrukking waaraan in die Wet en die regulasies betreffende kunsmatige bevrugting van 'n persoon afgekondig by Goewernmentskennisgewing R..... van 'n betekenis geheg is, daardie betekenis.

2. Behoudens die bepalings van die Wet en die regulasies word 'n gameet wat van die liggaam van 'n lewende persoon verwyder is nie vir kunsmatige bevrugting van 'n ander persoon gebruik nie tensy—

(a) die geneesheer wat sodanige kunsmatige bevrugting uitvoer of onder wie se toesig die bevrugting uitgevoer word, afsonderlik of in spanverband waar 'n geregistreerde genetikus beskikbaar is—

(i) die ontvanger en haar eggenoot van raadgewing en inligting bedien betreffende—

(aa) die moontlikhede vir normale swangerskap;

(bb) alle implikasies van kunsmatige bevrugting insluitende probleme rakende die tegniek van kunsmatige bevrugting, kanse vir sukses, finansiële aspekte, gevolge vir die huwelik, psigososiale- en opvoedkundige implikasies risiko's verbonde aan genetiese herkoms en prognose van die kind asook regsadvies wat verkry mag word betreffende kunsmatige bevrugting;

(ii) bepaal het dat die ontvanger biologies, fisies en geestelik geskik is om 'n kunsmatige bevrugting te ontvang;

(iii) die wense van die ontvanger en haar eggenoot met betrekking tot die skenker se bevolkings- en geloofsgroep eerbiedig;

(iv) waar die ontvanger of die skenker afkomstig is uit 'n bevolkingsgroep waar die individu 'n hoë risiko loop om 'n genetiese draer (resessief) van 'n spesifieke afwyking, byvoorbeeld Tay-Sachs, talassemie of sistiese fibrose te wees, die ontvanger en die skenker vir die betrokke eienskappe toets en nie die gameet van 'n skenker met dieselfde eienskappe as dié van die ontvanger vir die doeleindes van kunsmatige bevrugting gebruik nie;

(v) waar op die grond van die familiegeskiedenis 'n moontlikheid bestaan dat die ontvanger of die skenker 'n draer is van 'n afwyking wat deur gene oordraagbaar is, ondersoek en toets deur 'n geregistreerde genetikus laat doen om te bepaal of die ontvanger of die skenker die draer van die afwyking is in welke geval—

(aa) genetiese raadgewing gegee moet word aan die ontvanger en haar eggenoot waar dit bepaal is dat die ontvanger die draer van die betrokke afwyking is; of

(bb) die gameet van die skenker nie vir die doeleindes van kunsmatige bevrugting gebruik word nie waar dit bepaal is dat die skenker die draer van die afwyking is of die skenker 'n hoë risiko as 'n draer van die afwyking is.

ANNEXURE A**APPLICATION FOR REGISTRATION AS A DONOR FOR THE PURPOSE OF ARTIFICIAL INSEMINATION**

Application is hereby sought to register the undermentioned person, of which personal details appear on Annexure B, as a donor for the purpose of artificial insemination in terms of regulation 4, promulgated in terms of the Human Tissue Act, 1983:

Details of intended donor:

Surname
 Full christian names
 Date of birth
 Identity number
 Sex
 Racial group
 Details of applicant:
 Name of medical practitioner/hospital
 Address
 Capacity of applying medical practitioner
 Qualifications
 Date

ANNEXURE B**EVALUATION OF INTENDED DONOR**

Name
 I.D. No.
 Age
 Length
 Mass
 Colour:
 Hair
 Eyes
 Complexion
 Hair form (straight, curly etc.)
 Population group
 Blood group: ABO
 Rh
 Tay-Sachs carrier
 Thalassaemia carrier
 Other genetic conditions (specify)
 Sperm count
 Sexual transmissible diseases (pos./neg.)
 Hepatitis B antigen (pos./neg.)
 Gynaecological abnormalities
 Medical history (briefly)
 Psychiatric history
 Father: Age
 Health condition
 Cause of death, if dead
 Mother: Age
 Health condition
 Cause of death, if dead
 Maternal grandfather: Age
 Health condition
 Cause of death, if dead
 Maternal grandmother: Age
 Health condition
 Cause of death, if dead

AANHANGSEL A**AANSOEK OM REGISTRASIE AS 'N SKENKER VIR DIE DOELEINDES VAN KUNSMATIGE BEVRUGTING**

Aansoek word hiermee gedoen om die ondergenoemde persoon waarvan persoonlike besonderhede op Aanhangsel B verskyn, as 'n skenker vir die doeleindes van kunsmatige bevrugting ingevolge regulasie 4, uitgevaardig kragtens die Wet op Menslike Weefsel, 1983, te registreer:

Besonderhede van voornemende skenker:

Van
 Volle voorname
 Geboortedatum
 Identiteitsnommer
 Geslag
 Bevolkingsgroep
 Besonderhede van aansoeker:
 Naam van geneesheer/hospitaal
 Adres
 Hoedanigheid van aansoekende geneesheer
 Kwalifikasies
 Datum

AANHANGSEL B**EVALUASIE VIR VOORNEMENDE SKENKER****BESONDERHEDE**

Naam
 I.D. No.
 Ouderdom
 Lengte
 Massa
 Kleur: Hare
 Oë
 Gelaat
 Haarvorm (reguit, krullerig ens.)
 Bevolkingsgroep
 Bloedgroep: ABO
 Rh
 Tay-Sachs draer
 Talassemie draer
 Ander genetiese toestande (Spesifiseer)
 Spermteeling
 Geslaggedraagde siektes (pos./neg.)
 Hepatitis B antigeen (pos./neg.)
 Ginekologiese abnormaleite
 Mediese geskiedenis (bepynop)
 Psigiatryse geskiedenis
 Vader: Ouderdom
 Gesondheidstoestand
 Oorsaak van dood indien oorlede
 Moeder: Ouderdom
 Gesondheidstoestand
 Oorsaak van dood indien oorlede
 Oupa (Moederskant): Ouderdom
 Gesondheidstoestand
 Oorsaak van dood indien oorlede

Paternal grandfather: Age.....
 Health condition.....
 Cause of death, if dead.....
 Paternal grandmother: Age.....
 Health condition.....
 Cause of death, if dead.....
 Brothers or sisters:
 1. Age.....
 Health condition.....
 Cause of death, if dead.....
 2. Age.....
 Health condition.....
 Cause of death, if dead.....
 3. Age.....
 Health condition.....
 Cause of death, if dead.....
 4. Age.....
 Health condition.....
 Cause of death, if dead.....
 Other relatives with inherited defects.....
 Form completed by.....
Medical practitioner

Date

I declare that above information is correct and I am prepared to submit to any other further tests/examinations the Director-General may deem necessary.

Intended donor

Dated

ANNEXURE C

Department of Health and Welfare
 Private Bag X63
 PRETORIA

REGISTRATION CERTIFICATE OF DONOR FOR THE PURPOSE OF ARTIFICIAL INSEMINATION.

Registration Certificate No. Date of expiry

This is to certify that undermentioned person has been registered as a donor of gonades for the purpose of artificial insemination in terms of the Human Tissue Act, 1983:

Full names

Date of birth

Identity number

Address

This certificate has been issued with the understanding that artificial insemination as a result of such donations be limited to five (5) live births and with the further understanding that this certificate can be withdrawn at any time should sound reasons exist.

Director-General

Date

Department of Health and Welfare

ANNEXURE D

CONSENT TO ARTIFICIAL INSEMINATION (DONOR)

I.....
 I.D. No.
 Born.....
 hereby state:

1. I authorise Dr.....
 (hereinafter referred to as the doctor) or a suitably trained member of his/her staff to use semen or ova* donated by one of us for artificial insemination of another married woman.

Ouma (Moederskant): Ouderdom.....
 Gesondheidstoestand.....
 Oorsaak van dood indien oorlede.....
 Oupa (Vaderskant): Ouderdom.....
 Gesondheidstoestand.....
 Oorsaak van dood indien oorlede.....
 Ouma (Vaderskant): Ouderdom.....
 Gesondheidstoestand.....
 Oorsaak van dood indien oorlede.....
 Broer of susters:
 1. Ouderdom.....
 Gesondheidstoestand.....
 Oorsaak van dood indien oorlede.....
 2. Ouderdom.....
 Gesondheidstoestand.....
 Oorsaak van dood indien oorlede.....
 3. Ouderdom.....
 Gesondheidstoestand.....
 Oorsaak van dood indien oorlede.....
 4. Ouderdom.....
 Gesondheidstoestand.....
 Oorsaak van dood indien oorlede.....
 Ander familieleden met oorerflike afwykings.....
 Vorm voltooi deur.....

Geneesheer

Datum

Ek verklaar dat bogenoemde inligting korrek is en is bereid om my aan enige verdere toetse/ondersoeke wat die Direkteur-generaal nodig mag ag, te onderwerp.

Voornemende skenker

Datum

AANHANGSEL C

Departement van Gesondheid en Welsyn
 Private Bag X63
 PRETORIA
 0001

REGISTRASIESERTIFIKAAT VAN SKENKER VIR DIE DOELEINDES VAN KUNSMATIGE BEVRUGTING

Registrasiesertifikaat No. Vervaldatum

Hiermee word gesertifiseer dat ondergenoemde persoon in terme van die Wet op Menslike Weefsel, 1983 geregistreer is as 'n skenker van gonade vir die doeleindes van kunsmatige bevrugting:

Volle naam

Geboortedatum

Identiteitsnommer.....

Adres

Hierdie sertifikaat word uitgereik met dien verstande dat kunsmatige bevrugting wat uit sodanige skenkings voortspruit, beperk word tot vyf (5) lewendige geboortes en met dien verstande voorts dat die sertifikaat te eniger tyd ingetrek kan word indien daar goeie redes bestaan.

Direkteur-generaal

Datum

Department van Gesondheid en Welsyn

AANHANGSEL D

TOESTEMMING TOT KUNSMATIGE BEVRUGTING (SKENKER)

Ek

I.D. nommer

Gebore....., verklaar soos volg:

1. Ek magtig dr.....
 (hierna "die geneesheer" genoem) of 'n paslik opgeleide lid van sy/haar personeel om my sperma of ova* wat geskenk word vir kunsmatige bevrugting te gebruik.

2. My wish is that the semen/ova* shall be used for a recipient belonging to the population group who is adherent of the faith.
3. The doctor has explained to me the nature and implications of the procedure.
4. I agree that I shall not be entitled at any stage to be informed as to the identity of the recipient(s).
5. I accept that I have no responsibility to any child/children born as a result of this procedure and I waive any legal rights on such child.
6. I accept that my semen/ova* will be used for five successful inseminations only and agree not again to present myself as a donor once I have been informed that five such inseminations have been performed or if any evidence of genetic disorders or detrimental qualities prevails in any of my descendants.
7. I give permission that my name be recorded as a donor in a highly confidential register kept by the Department of Health and Welfare for this purpose.

Signed at
on this day of 19.....

Donor

Witness

Medical practitioner

* Delete which is not applicable.

ANNEXURE E

NOTICE OF INTENTION TO PERFORM AN ARTIFICIAL INSEMINATION

I, properly registered medical practitioner hereby give notice that I intend to perform an artificial insemination on the undermentioned person:

Name
Identity number
Date of birth
Conjugal state
Maiden surname (if married)
Address

Registration certificate of intended donor
Full details regarding the premises and street address where the artificial insemination is intended.

Signature of medical practitioner	Place
Qualifications	Date
Official designation	

ANNEXURE F

EVALUATION OF INTENDED RECIPIENT

Name
I.D. No.
Age
Length
Mass
Colour:
Hair
Eyes
Complexion
Hair form (straight, curly etc)
Population group
Blood group: ABO
Rh
Tay-Sachs carrier
Thalassaemia carrier

2. Dit is my begeerte dat die sperma/ova* wat gebruik sal word vir 'n ontvanger van die bevolkingsgroep en aanhanger van die geloof, gebruik word.
3. Die geneesheer het die aard, omvang en implikasies van die prosedure aan my verduidelik.
4. Ek aanvaar om in die oordeel van die geneesheer te berus by die keuse van 'n geskikte ontvanger en dat ek op geen tydstip geregtig sal wees om oor die identiteit van die ontvanger/s ingelig te word nie.
5. Ek aanvaar dat ek regtens geen verantwoordelikheid het teenoor enige kinders wat as gevolg van prosedure verwek word nie en doen afstand van enige regte wat ek ten opsigte van sodanige kind sou kon verkry.
6. Ek aanvaar dat my sperma/ova* slegs vir vyf suksesvolle bevrugtings gebruik mag word en onderneem om my nie weer as skenker aan te meld nadat ek meegedeel is dat vyf sodanige bevrugtings plaasgevind het of indien daar enige bewyse van genetiese afwykings/nadelige oorerflike eienskappe by enigeen van my nakomelinge voorkom nie.
7. Ek gee toestemming dat my naam as skenker geplaas mag word in 'n hoogsvertroulike register wat deur die Departement van Gesondheid en Welsyn vir die doel gehou word.

Geteken te
op hierdie dag van 19.....

Skenker

Getuie

Geneesheer

* Skrap wat nie van toepassing is nie.

AANHANGSEL E

KENNISGEWING VAN VOORNEME OM 'N KUNSMATIGE BEVRUGTING TE DOEN

Ek, , behoorlik geregistreerde geneesheer gee hiermee kennis dat ek voornemens is om 'n kunsmatige bevrugting van die ondergenoemde persoon te doen:

Naam
Identifikasienommer
Geboortedatum
Huweliksstaat
Nooiensvan (indien getroud)
Adres

Registrasiesertifikaat van beoogde skenker
Volledige besonderhede oor perseel en straatadres waar kunsmatige bevrugting beoog word

Handtekening van geneesheer	Plek
Kwalifikasies	Datum
Ampstitel	

AANHANGSEL F

EVALUASIE VAN VOORNEMENDE ONTVANGER BESONDERHEDE

Naam
I.D. No.
Ouderdom
Lengte
Massa
Kleur:
Hare
Oë
Gelaat
Haarvorm (reguit, krullerig ens.)
Bevolkingsgroep
Bloedgroep: ABO
Rh
Tay-Sachs draer
Talassemie draer

Other genetic conditions (specify)

Sperm count

Sexual transmissible diseases (pos./neg.)

Hepatitis B antigen (pos./neg.)

Gynaecological abnormalities

Medical history (briefly)

Psychiatric history

Father: Age

Health condition

Cause of death, if dead

Mother: Age

Health condition

Cause of death, if dead

Maternal grandfather: Age

Health condition

Cause of death, if dead

Maternal grandmother: Age

Health condition

Cause of death, if dead

Paternal grandfather: Age

Health condition

Cause of death, if dead

Paternal grandmother: Age

Health condition

Cause of death, if dead

Brothers or sisters:

1. Age

Health condition

Cause of death, if dead

2. Age

Health condition

Cause of death, if dead

3. Age

Health condition

Cause of death, if dead

4. Age

Health condition

Cause of death, if dead

Other relatives with inherited defects

Form completed by

Medical practitioner

Date

I declare that above information is correct and I am prepared to submit to any other further tests/examinations the Director-General may deem necessary.

Intended recipient

Date

DEPARTMENT OF JUSTICE

No. R. 527

15 March 1985

POWERS REGARDING OATH OUTSIDE THE REPUBLIC.—SECTION 8 (1) (a) OF THE JUSTICES OF THE PEACE AND COMMISSIONERS OF OATHS ACT, 1963 (ACT 16 OF 1963)

By virtue of the powers vested in me by section 8 (1) (a) of the Justices of the Peace and Commissioners of Oaths Act, 1963, I, Hendrik Jacobus Coetsee, Minister of Justice, hereby amend Government Notice R. 1872 of 12 September 1980, as amended by Government Notice R. 2828 of 30 December 1983, by the substitution in the Schedule thereto for the words "the holder of an office of Scientific Adviser of the South African Council for Scientific and Industrial Research" of the words "the holder of an office of Science Counsellor or Science Consul of the South African Council for Scientific and Industrial Research".

H. J. COETSEE, Minister of Justice.

Ander genetiese toestande (spesifiseer)

Spermtelling

Geslagsgedraagde siektes (pos./neg.)

Hepatitis B antigeen (pos./neg.)

Ginekologiese abnormaleite

Mediese geskiedenis (bepal)

Vader: Ouderdom

Gesondheidstoestand

Oorsaak van dood indien oorlede

Moeder: Ouderdom

Gesondheidstoestand

Oorsaak van dood indien oorlede

Oupa (Moederskant): Ouderdom

Gesondheidstoestand

Oorsaak van dood indien oorlede

Ouma (Moederskant): Ouderdom

Gesondheidstoestand

Oorsaak van dood indien oorlede

Oupa (Vaderskant): Ouderdom

Gesondheidstoestand

Oorsaak van dood indien oorlede

Ouma (Vaderskant): Ouderdom

Gesondheidstoestand

Oorsaak van dood indien oorlede

Broers of susters:

1. Ouderdom

Gesondheidstoestand

Oorsaak van dood indien oorlede

2. Ouderdom

Gesondheidstoestand

Oorsaak van dood indien oorlede

3. Ouderdom

Gesondheidstoestand

Oorsaak van dood indien oorlede

4. Ouderdom

Gesondheidstoestand

Oorsaak van dood indien oorlede

Ander familieled met oorerflike afwykings

Vorm voltooi deur:

Geneesheer

Datum

Ek verklaar dat bogenoemde inligting korrek is en is bereid om my aan enige verdere toetse/ondersoeke wat die Direkteur-generaal nodig mag ag, te onderwerp.

Voornemende ontvanger

Datum

DEPARTEMENT VAN JUSTISIE

No. R. 527

15 Maart 1985

BEVOEGDHEDE BETREFFENDE EDE BUITE DIE REPUBLIEK.—ARTIKEL 8 (1) (a) VAN DIE WET OP VREDEREGTERS EN KOMMISSARISSE VAN EDE, 1963 (WET 16 VAN 1963)

Kragtens die bevoegdheid my verleen by artikel 8 (1) (a) van die Wet op Vrederegters en Kommissarisse van Ede, 1963, wysig ek, Hendrik Jacobus Coetsee, Minister van Justisie, hierby Goewerménskennisgewing R. 1872 van 12 September 1980, soos gewysig by Goewerménskennisgewing R. 2828 van 30 Desember 1983, deur in die Bylae daarvan die woorde "die bekleër van 'n amp van Wetenskaplike Raadgewer van die Suid-Afrikaanse Wetenskaplike Nywerheidsnavorsingsraad" deur die woorde "die bekleër van 'n amp van Wetenskaplike Raad of Wetenskaplike Konsul van die Suid-Afrikaanse Wetenskaplike en Nywerheidsnavorsingsraad" te vervang.

H. J. COETSEE, Minister van Justisie.

DEPARTMENT OF MANPOWER**No. R. 523****15 March 1985****LABOUR RELATIONS ACT, 1956****BORDER FURNITURE MANUFACTURING INDUSTRY.—EXTENSION OF MAIN AGREEMENT**

I, Jacob Salmon Herselman, Director: Manpower, duly authorised thereto by the Minister of Manpower, hereby, in terms of section 48 (4) (a) (i) of the Labour Relations Act, 1956, extend the periods fixed in Government Notices R. 1227 of 22 June 1984 by a further period ending 20 March 1986.

J. S. HERSELMAN, Director: Manpower.

DEPARTEMENT VAN MANNEKRAG**No. R. 523****15 Maart 1985****WET OP ARBEIDSVERHOUDINGE, 1956****MEUBELNYWERHEID, GRENS.—VERLENGING VAN HOOFOOREENKOMS**

Ek, Jacob Salmon Herselman, Direkteur: Mannekrag, behoorlik daartoe gemagtig deur die Minister van Mannekrag, verleng hierby, kragtens artikel 48 (4) (a) (i) van die Wet op Arbeidsverhoudinge, 1956, die tydperke vasgestel in Goewermentskennisgewings R. 1227 van 22 Junie 1984 met 'n verdere tydperk wat op 20 Maart 1986 eindig.

J. S. HERSELMAN, Direkteur: Mannekrag.

No. R. 576**15 March 1985****LABOUR RELATIONS ACT, 1956****JEWELLERY AND PRECIOUS METAL INDUSTRY (CAPE).—RENEWAL OF TRAINING FUND AGREEMENT**

I, Jacob Salmon Herselman, Director: Manpower, duly authorised thereto by the Minister of Manpower, hereby in terms of section 48 (4) (a) (ii) of the Labour Relations Act, 1956, declare the provisions of Government Notices R. 960 of 3 June 1977 and R. 2156 of 27 October 1978, to be effective from the date of publication of this notice and for the period ending 31 December 1986.

J. S. HERSELMAN, Director: Manpower.

No. R. 576**15 Maart 1985****WET OP ARBEIDSVERHOUDINGE, 1956****JUWELIERSWARE- EN EDELMETAALNYWERHEID (KAAP).—HERNUWING VAN OPLEIDINGSFONDS-OOREENKOMS**

Ek, Jacob Salmon Herselman, Direkteur: Mannekrag, behoorlik daartoe gemagtig deur die Minister van Mannekrag, verklaar hierby, kragtens artikel 48 (4) (a) (ii) van die Wet op Arbeidsverhoudinge, 1956, dat die bepalings van Goewermentskennisgewings R. 960 van 3 Junie 1977 en R. 2156 van 27 Oktober 1978, van krag is vanaf die datum van publikasie van hierdie kennisgewing en vir die tydperk wat op 31 Desember 1986 eindig.

J. S. HERSELMAN, Direkteur: Mannekrag.

DEPARTMENT OF TRADE AND INDUSTRY**No. R. 574****15 March 1985****MAXIMUM INDUSTRIAL PRICES OF SUGAR**

I, Dawid Jacobus de Villiers, Minister of Trade and Industry, after consultation with the South African Sugar Association, hereby—

(a) in terms of section 6 (3) of the Sugar Act, 1978 (Act 9 of 1978), withdraw Government Notice R. 1998 of 14 September 1984 with effect from 15 March 1985;

(b) in terms of section 6 (1) of the Sugar Act, 1978 (Act 9 of 1978), prescribe that, with effect from 15 March 1985, the maximum industrial prices at which the sugar industry products mentioned in the Schedule hereto may be sold shall be as set out in the Schedule.

D. J. DE VILLIERS, Minister of Trade and Industry.

SCHEDULE

The maximum industrial prices at Durban per metric ton of refined and brown sugar packed in 25 kg pockets for sale on the local market shall be:

Refined sugar: R658,00.

Brown sugar: R596,00.

DEPARTEMENT VAN HANDEL EN NYWERHEID**No. R. 574****15 Maart 1985****MAKSIMUM NYWERHEIDSPRYSE VAN SUIKER**

Ek, Dawid Jacobus de Villiers, Minister van Handel en Nywerheid, na oorleg met die Suid-Afrikaanse Suikervereniging—

(a) trek hierby kragtens artikel 6 (3) van die Suikervet, 1978 (Wet 9 van 1978), Goewermentskennisgewing R. 1998 van 14 September 1984 in met ingang van 15 Maart 1985;

(b) Skryf hierby kragtens artikel 6 (1) van die Suikervet, 1978 (Wet 9 van 1978) voor dat met ingang van 15 Maart 1985 die maksimum nywerheidspryse waarteen die suikernywerheidsprodukte wat in die Bylae hiervan gemeld word, verkoop mag word, is soos in die Bylae uiteengesit.

D. J. DE VILLIERS, Minister van Handel en Nywerheid.

BYLAE

Die maksimum nywerheidspryse te Durban per metrieke ton geraffineerde en bruinsuiker verpak in pakkies van 25 kg elk vir verkoop in die plaaslike mark is:

Geraffineerde suiker: R658,00.

Bruinsuiker: R596,00.

DEPARTMENT OF TRANSPORT

No. R. 546

15 March 1985

THIRD AMENDMENT OF THE AERODROME REGULATIONS, 1982

The Minister of Transport Affairs, in terms of section 22 of the Aviation Act, 1962 (Act 74 of 1962), has made the regulations contained in the Schedule hereto.

SCHEDULE

1. In this Schedule "the Regulations" means the Aerodrome Regulations, 1982, promulgated by Government Notice R. 289 of 15 October 1982, as amended by Government Notice R. 130 of 28 January 1983 and R. 1695 of 10 August 1984.

2. Regulation 2 of the Regulations is hereby amended by the substitution, for subregulation 2.9 (1) to (8), of the following:

"2.9 Landing fees and parking fees at Non-State Airports shall be determined by the United Municipal Executive on the recommendation of the operators Committee: Provided that landing fees and parking fees at Non-State Airports shall not be higher than those levied at State Airports."

DEPARTEMENT VAN VERVOER

No. R. 546

15 Maart 1985

DERDE WYSIGING VAN DIE VLEGVELD-REGULASIES, 1982

Die Minister van Vervoerwese het kragtens artikel 22 van die Lugvaartwet, 1962 (Wet 74 van 1962), die regulasies in die Bylae hiervan uitgevaardig.

BYLAE

1. In hierdie Bylae beteken "die Regulasies" die Vliegvelddregulasies, 1982, afgekondig by Goewermentskennisgewing R. 289 van 15 Oktober 1982, soos gewysig by Goewermentskennisgewing R. 130 van 28 Januarie 1983 en R. 1695 van 10 Augustus 1984.

2. Regulasie 2 van die Regulasies word hierby gewysig deur subregulasie 2.9 (1) tot (8) deur die volgende te vervang:

"2.9 Landingsgelde en parkeergelde by die nie-Staatslughawens word op aanbeveling van die Operateurskomitee deur die Verenigde Munisipale Bestuur vasgestel: Met dien verstande dat landingsgelde en parkeergelde by nie-Staatslughawens nie hoër mag wees as dié wat by Staatslughawens gehef word nie."

THE FLOWERING PLANTS OF AFRICA

This publication is issued as an illustrated serial, much on the same lines as Curtis's Botanical Magazine, and for imitating which no apology need be tendered.

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Don't abuse



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water is for everybody

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Ons leef



daarvan.

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