CB(2) 1516/98-99(03)

AKWONG: DMA# 7393

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HUMAN REPRODUCTIVE TECHNOLOGY BILL

COMMITTEE STAGE

Amendments to be moved by the secretary for Health and Welfare

Clause		Amendment Proposed
13(3)	(a)	In paragraph (a), by deleting "a severe sex-linked genetic disease" and substituting "any of the sex-linked genetic diseases specified in Schedule 1A".
	(b)	In paragraph (b), by adding "certifies that such disease is sufficiently severe to justify selection and" before "state in writing".
43		By deleting "Schedule 1" and gubstituting "Schedules 1 and 1A".

New By adding -

"SCHEDULE 1A [ss.13(3) (a) & 43]

SEX-LINKED GENETIC DISEASE

Addison's disease with cerebral selerosis
Adrenoleucodystrophy
Adrenal hypoplasis (one type)
Agammagloblinaemia, Bruton type (sometimes also gwiss type)
Albinism, ocular
Albinism-deafness syndrome
Aldrich syndrome
Alport syndrome (some kindreds)
Amelogenesis imperfecta (two types)
Anaemia, heredirary hypochromic
Angiokeratoma (Fabry's disease)
Cataract, congenital (one type)
Cerebellar ataxia (one type)
Cerebral sclerosis, diffuse
Charcot-Marie-Tooth peroneal muscular atrophy (one type)
Choroideraemia
Choroidoretinal degeneration (one rare type)

Coffin-Lowry syndrome

Colour blindness (several types)

Deafness, perceptive (several types)

Diabetes insipidus, nephrogenic

Diabetes insipidus, neurohypophyseal (some families)

Dyskeratosis congenita

Ectodermal dysplasia, anhidrotic

Ehlers-Danlos syndrome, type V

Faciogenital dysplasia, (Aarskog syndrome)

Focal dermal hypoplasia

Glucose 6-phosphatc dehydrogenase deficiency

Glyeogen storage disease, type VIII

Gonadal dysgenesis (XY female type)

Granulomatous disease (chronic)

Haemophilia A

Haemophilia B

Hydroccphalus (aqueduct stenosis, one type)

Hypophosphataemic rickets

Ichthyosis (steriod sulphatase deficiency)

Incontinentia pigmenti

Kallmann syndrome

Keracosis lollicularis spinulosa

Lesch-Nyhan syndrome (hypoxanthine-guanine-phosphoribosyl transferase deficiency)

Lowe (oculocerebrorenal) syndrome

Macular dystrophy of the retina (one type)

Menkes syndrome

Mental remrdation, with or without fragile site (several specific types)

Microphthalmia with multiple anomalies (Lenz syndrome)

Mucopolysaccharidosis II (Hunter syndrome)

Museular dystrophy (Becker, Duchenne and Emery-Drelfuss types)

Myowbular myopathy (one type)

Night blindness, congenital stationary

Norrie's disease (pseudoglioma)

Nystagmus, oculomotor or 'jerky'

Ornithine transearbamylase deficiency (type I hyperammonaemia)

Orofaciodigital syndrome (type I)

Phosphoglycerate kinase deficiency

Phosphoribosylpyrophosphate (PRPP) synthetase deficiency

Reifenstcin syndrome

Retinitis pigmentosa (one type)

Retinoschisis

Spastic paraplegia (one type)

Spinal muscular atrophy (one type)

Spondyloepiphyseal dysplasia tarda

Testicular feminization syndrome

Thrombocytopenia, hereditary (one type)

Thyroxine-binding globulin, absence or
variants of

Xg blood group system".