A Beautiful Beginning
Genesis International
Egg & Sperm Centre

DONOR CODE – 05405



Characteristics

- *Ethnicity - Chinese*
- *Eye colour - Black*
- *Hair colour - Black*
- *Hair texture - Straight*
- *Skin tone - Fair*
- *Height – 153cm*
- *Weight – 45kg*
- *Vision - Clear*
- *Glasses - No*
- *Blood group – B+*
- *CMV status - Positive*
- *Extended genetic status - Positive*

My Background

- *Country of Birth - Malaysia*
- *Religion - Buddhist*
- *Education – High School Diploma*
- *Occupation - Beautician*
- *Children – Nil to date*
- *Donor children – Nil to date*

A bit about me

- *Professional Interests – To work hard*
- *Personality -Always cheerful*
- *Attitude – Approach life with a positive mindset*
- *Why I want to be a donor angel – To help families have their own family*
- *As a child – Playful and active*
- *How do my friends describe me – Always caring and helpful*
- *My relationship with my family – Close with my family*
- *What am I most proud of – Achieving my Diploma*
- *My strengths – Not giving up on my career when times are tough*
- *Sports interests - Badminton*
- *My favourite books - No*
- *My favourite movie - No*
- *Do I play a musical instrument - No*

Family Characteristics

	Height	**Eye color**	**Hair color**	**Skin tone**	**Occupation**
My Mother	158cm	Brown	Black	Fair	Housewife
My Father	168cm	Black	Black	Fair	Farmer
My Sister	153cm	Brown	Black	Fair	Housewife
My Brother	160cm	Brown	Black	Fair	Businessman

My families Ancestry and medical history

	Ethnic origin	Country of birth	Medical History
My Mother	Chinese	Malaysia	Nil known
My Father	Chinese	Malaysia	Nil known
My Maternal Grandmother	Chinese	Malaysia	Nil known
My Maternal Grandfather	Chinese	Malaysia	Nil known
My Paternal Grandmother	Chinese	Malaysia	Nil known
My Paternal Grandfather	Chinese	Malaysia	Nil known

Achondroplasia (Dwarfism)	Nil Known
ADD or ADHD	Nil Known
Allergies Drugs/ Food	Nil Known
Arthritis: Osteoarthritis, Rheumatoid	Nil Known
Asthma	Nil Known
Autism	Nil Known
Auto-immune Disease, CREST Syndrome, Lupus, Scleroderma, Sjorgen's Syndrome	Nil Known
Cancer or Malignant Tumor	Nil Known
Congenital Heart Disease	Nil Known
Congenital Hip Disease	Nil Known
Cystic Fibrosis	Nil Known
Deafness	Nil Known
Diabetes type 1 and 2	Nil Known
Down Syndrome	Nil Known
Dyslexia, learning disabilities	Nil Known
Eczema	Nil Known
Blindness, Colour Blindness, Cataracts, Glaucoma, Retinoblastoma	Nil Known
Fanconi Anaemia	Nil Known
Fragile X	Nil Known
Ulcerative Colitis, Crohns, Pyloric Stenosis	Nil Known
Hepatitis	Nil Known
Heart Attack, Heart Disease (congenital or otherwise)	Nil Known
Haemochromatosis (iron overload)	Nil Known
Haemophilia	Nil Known
High Blood Pressure	Nil Known
Hypoglycaemia	Nil Known
Kidney Disorders including born with 1 kidney	Nil Known
Lesch-Nyhan Syndrome	Nil Known

Loss of muscle coordination	Nil Known
Malformations including: cleft lip or palate, club foot, polydactyly	Nil Known
digits, hypospadias	Nil Known
Marfan's Disease	Nil Known
Mental illness requiring hospitalisation or mental retardation	Nil Known
Manic Depression (bipolar), Schizophrenia	Nil Known
Muscular Dystrophy	Nil Known
Neurological diseases including; Alzheimer's, Epilepsy, Huntington's, Lou Gehrig's, Parkinson's,	Nil Known
Creutzfeld-Jacob Disease (CJD), GullianBarre, JC Virus, Multiple Sclerosis, Sub-Acute Sclerosing Panencephalitis, Spongiform Encephalopathy/Prion Disease	Nil Known
Neurofibromatosis	Nil Known
Neural tube defect	Nil Known
PKU or inherited metabolic disorders	Nil Known
Premature degeneration of any organ	Nil Known
Sickle Cell	Nil Known
Stroke	Nil Known
Sudden Infant Death Syndrome (SIDS)	Nil Known
Skin Disease	Nil Known
Syndrome: Noonan, Bloome, Klinefelters or Turners	Nil Known
Tay Sachs	Nil Known
Thalassaemia	Nil Known
Thyroid disease	Nil Known
Tuberculosis	Nil Known
Xenotransplant	Nil Known