



My Characteristics

- *Ethnicity - Chinese*
- *Eye colour - Black*
- *Hair colour - Black*
- *Hair texture - Straight*
- *Skin tone - Light brown*
- *Height - 153cm*
- *Weight - 43.5kg*
- *Vision - Slight blurry vision*
- *Glasses - Yes*
- *Blood group - O Positive*
- *CMV status - Positive*
- *Myriad genetic status - Negative carrier*

My Background

- *Country of Birth - Malaysia*
- *Religion - Buddhist*
- *Education - O level equivalent*
- *Occupation - Admin*
- *Children – Nil to date*
- *Donor children - Yes*

A bit about me

- *Professional Interests - To travel and sing*
- *Personality - Cute and loved by everyone*
- *Attitude - Punctual, being helpful and getting along with people*
- *Why I want to be a donor angel - To help others that cannot get pregnant*
- *As a child - Talkative and playful*
- *How do my friends describe me - Liked by everyone*
- *My relationship with my family - Very good relationship with my parents and thankful they raised me to be a good person*
- *What am I most proud of - Being beautiful within*
- *My strengths - Very positive person*
- *Sports interests - Jogging and badminton*
- *My favourite books - Harry Potter*
- *My favourite movie -Titanic*
- *Do I play a musical instrument- No*

Family Characteristics

	Height	Eye color	Hair color	Skin tone	Occupation
My Mother	155	Black	Black	Light brown	Businesswomen
My Father	165	Black	Black	Light brown	Retired
My Sister	158	Black	Black	Light brown	Clerk
My Brother	158	Black	Black	Light brown	Manager

My families Ancestry and medical history

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

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

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