



My Characteristics

- *Ethnicity - Chinese*
- *Eye colour - Brown*
- *Hair colour - Black*
- *Hair texture - Straight*
- *Skin tone - Fair*
- *Height – 170cm*
- *Weight – 66kg*
- *Vision – Short sighted*
- *Glasses - Yes*
- *Blood group – O+*
- *CMV status - Negative*
- *Genetic status - Negative*

My Background

- *Country of Birth - Malaysia*
- *Religion - Buddhist*
- *Education – Degree in Professional Chef training*
- *Occupation – Chef*
- *Children – Nil to date*
- *Donor children – Nil to date*

A bit about me

- *Professional Interests – Cooking, travel and jogging*
- *Personality – Good sense of humour*
- *Attitude – Passionate, hardworking and helpful*
- *Why I want to be a donor- To help infertile couples the chance to have a baby.*
- *As a child – Cute and playful*
- *How do my friends describe me – Friendly and funny*
- *My relationship with my family - Good and close to family*
- *What am I most proud of – All my achievements in life*
- *My strengths – Insistent and passionate*
- *Sports interests - Jogging*
- *My favourite books – Cookbooks that inspire me*
- *My favourite movie -Action/Sci-fi*
- *Do I play a musical instrument - No*

Family Characteristics

	Height	Eye color	Hair color	Skin tone	Occupation
My Mother	160cm	Brown	Black	Fair	Housewife
My Father	165cm	Brown	Black	Fair	Retired
My Sister 1	175cm	Brown	Black	Fair	IT
My Sister 2	165cm	Brown	Black	Fair	Sales

My families Ancestry and medical history

	Ethnic origin	Country of birth	Medical History
My Mother	Chinese	Malaysia	
My Father	Chinese	Malaysia	Heart attack, angiogram required.
My Maternal Grandmother	Chinese	Malaysia	
My Maternal Grandfather	Chinese	Malaysia	
My Paternal Grandmother	Chinese	Malaysia	Alzheimer's age 80 years
My Paternal Grandfather	Chinese	Malaysia	

Achondroplasia (Dwarfism)	Nil known
ADD or ADHD	Nil known
Allergies Drugs/ Food	Yes Donor – Seafood since birth
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)	My father age 64 years old
Haemochromatosis (iron overload)	Nil known
Haemophilia	Nil known
High Blood Pressure	My Father
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,	Grandmother (Fathers side) Age 80 years old
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	Psoriasis Father aged 40 years +
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