Donor 92015

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Blank fields are intentional and indicate that the information is not available.

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General Information

Donor ID:	92015
Year of Birth:	1995
Marital Status:	Single
Number of Children:	0
Religion:	None
Occupation:	Lighting Technician
Blood Type:	O +
Nationality:	Maternal: Chinese, Filipino, Irish, Paternal: French, Irish
Race:	
CMV Status: Learn More	NEG
Reported Pregnancy:	Yes
Canadian Compliant:	Yes
UK Compliant:	Yes
Identity Disclosure:	Yes
Last Medical History Update:	12/04/2019
Audio File:	Not Available

Child Photo

Adult Photo

Marital Status: Single
Number of Children: 0
Religion: None
Occupation: Lighting Technician **Blood Type:** O + **Ethnicity:** Maternal: Chinese, Filipino, Irish, Paternal: French, Irish
Audio File:

Donor Photos

Number of photos available: Not available at this time.

Physical Attributes

Height:	5'10" (177.80cm)
Weight:	170 lbs (77kg)
Eye Color:	Brown, Medium
Hair Color:	Black
Hair Texture:	Thick
Hair Loss:	None
Hair Type:	Wavy
Dominant Hand:	Right
Hairy Chest:	No
Hairy:	No
Ear Lobes:	Attached
Beard Color:	Black

Eyebrows:	Medium
Dimples:	Yes
Acne:	Yes
Acne Information:	Age 12/13, middle school, not severe, did not need treatment
Shoe Size:	10
Body Build:	Medium
Freckles:	None
Skin Tone:	Brown, Medium
Face Shape:	Oval
Lips:	Normal
Nose Shape:	Normal
Long Eyelashes:	No

In His Own Words

Hey! I made the decision to donate to help parents like you have a child of their own. I'm sure you want to know a little about me. I grew up playing sports with my friends - basketball, football, soccer, baseball, golf, you name it! Playing these sports as I grew up, I learned a lot of lessons. Mostly they taught me how to treat other people with respect. You never know what someone else is going through, and some people use sports as an outlet

to forget about their problems. I have applied the lessons I learned from sports to my entire life. On and off the field, you have to lookout for one another, always try help when you can, and stand up for people when they are scared to stand up for themselves.
This type of thinking shaped me as a person and helped me become who I am today. I am lucky to have wonderful parents. They didn't give me handouts but they gave me love, support, and guidance on how to be a strong, independent person. From a young age they insisted that I worked so I would know how to balance a job with regular life. I hated it at the time, but as I grew up, I realized how much it helped me get my priorities in order.

After I graduated from high school my sports career began to fade and I really started diving into the art of music production. I attended a university for a business degree but after a year of doing that I realized I was unmotivated in that area of study. Instead of continuing in that major, I took a break. The whole time I continued to produce music and learn as much as possible about sound design and engineering. After a couple years of just working a day job and learning how to produce, I finally got to a point where I realized what I was really good at it. I knew I wanted music production to be my career. I'm currently enrolled in an audio engineering program, enjoying every second of it. My point is, don't do something you don't want because everyone else is doing it. The people who have my utmost respect are the people who are passionate about what they do and don't let others influence their decisions. Follow your passions. It will only inspire the others around you to do the same thing.

To sum it up, all of the above has gotten me to where I am today. I live in an awesome home with a great group of roommates who are my best friends. We're all working towards the future. It's important to surround yourself with people who you trust and who have pure intentions. There's a quote "Who you're with is who you become" basically, if you hang out with people that steal and lie, there's a chance it could rub off on you… So find some good friends, have fun, stay safe, be logical, be respectful, listen to good music and be the person you would want your own kid to be!

Much love.

Genetic Testing

Over the years, Xytex has expanded genetic testing of donors as genetic knowledge and technology have advanced. As a result, not all donors have been tested for the same conditions or with the same methods. For more information about genetic testing view our "**Genetics Video and FAQs**."

This donor has been evaluated for carrier status of the listed conditions. A negative result reduces, but does not eliminate, the likelihood of the donor being a carrier. If you are a known carrier of a genetic condition please **contact us**, and we will provide a copy of the donor's test reports to your physician and/or genetic counselor.

This donor was tested for **283** conditions.

Dystrophic Epidermolysis Bullosa (COL7A1)	No disease-causing mutations detected
Emery-Dreifuss Myopathy 1 (EMD)	No disease-causing mutations detected
Enhanced S-Cone Syndrome (NR2E3)	No disease-causing mutations detected
Factor IX Deficiency (F9)	No disease-causing mutations detected
Familial Autosomal Recessive Hypercholesterolemia (LDLRAP1)	No disease-causing mutations detected
Familial Hypercholesterolemia (LDLR)	No disease-causing mutations detected
Fanconi Anemia, Group G (FANCG)	No disease-causing mutations detected
Fumarase Deficiency (FH)	No disease-causing mutations detected
Gitelman Syndrome (SLC12A3)	No disease-causing mutations detected
Glutaric Acidemia, Type IIa (ETFA)	No disease-causing mutations detected
Glutaric Acidemia, Type IIc (ETFDH)	No disease-causing mutations detected
Glycogen Storage Disease, Type IV also known as Adult Polyglucosan Body Disease (GBE1)	No disease-causing mutations detected
Glycogen Storage Disease, Type VII (PFKM)	No disease-causing mutations detected
Hemochromatosis, Type 2A (HFE2)	No disease-causing mutations detected
Hemochromatosis, Type 3 (TFR2)	No disease-causing mutations detected

Hereditary Spastic Paraparesis 49 (TECPR2)	No disease-causing mutations detected
Hermansky-Pudlak Syndrome, Type 1 (HPS1)	No disease-causing mutations detected
Hermansky-Pudlak Syndrome, Type 3 (HPS3)	No disease-causing mutations detected
Homocystinuria due to MTHFR Deficiency (MTHFR)	No disease-causing mutations detected
Homocystinuria, cblE Type (MTRR)	No disease-causing mutations detected
Hyperornithinemia-Hyperammonemia-Homocitrullinuria Syndrome (SLC25A15)	No disease-causing mutations detected
Hypohidrotic Ectodermal Dysplasia 1 (EDA)	No disease-causing mutations detected
Infantile Cerebral and Cerebellar Atrophy (MED17)	No disease-causing mutations detected
Leber Congenital Amaurosis 10 and Other CEP290-Related Ciliopathies (CEP290)	No disease-causing mutations detected
Leber Congenital Amaurosis 13 (RDH12)	No disease-causing mutations detected
Leber Congenital Amaurosis 2 also known as Retinitis Pigmentosa 20 (RPE65)	No disease-causing mutations detected
Leber Congenital Amaurosis 5 (LCA5)	No disease-causing mutations detected
Leber Congenital Amaurosis 8 also known as Retinitis Pigmentosa 12 (CRB1)	No disease-causing mutations detected
Lethal Congenital Contracture Syndrome 1 (GLE1)	No disease-causing mutations detected
Lipoprotein Lipase Deficiency (LPL)	No disease-causing mutations detected
Lysinuric Protein Intolerance (SLC7A7)	No disease-causing mutations detected

Microphthalmia and Anophthalmia (VSX2)	No disease-causing mutations detected
Mitochondrial Complex I Deficiency, ACAD9-Related (ACAD9)	No disease-causing mutations detected
Mitochondrial Complex I Deficiency, NDUFAF5-Related (NDUFAF5)	No disease-causing mutations detected
Mitochondrial Complex I Deficiency, NDUFS6-Related (NDUFS6)	No disease-causing mutations detected
Mitochondrial DNA Depletion Syndrome 6 also known as Navajo Neurohepatopathy (MPV17)	No disease-causing mutations detected
Mitochondrial Myopathy and Sideroblastic Anemia 1 (PUS1)	No disease-causing mutations detected
Mucopolysaccharidosis Type IIID (GNS)	No disease-causing mutations detected
Mucopolysaccharidosis Type VI (ARSB)	No disease-causing mutations detected
Mucopolysaccharidosistype IX (HYAL1)	No disease-causing mutations detected
Myoneurogastrointestinal Encephalopathy (TYMP)	No disease-causing mutations detected
N-Acetylglutamate Synthase Deficiency (NAGS)	No disease-causing mutations detected
Nephrogenic Diabetes Insipidus, Type II (AQP2)	No disease-causing mutations detected
Neuronal Ceroid-Lipofuscinosis, MFSD8-Related (MFSD8)	No disease-causing mutations detected
Odonto-Onycho-Dermal Dysplasia also known as Schopf-Schulz-Passarge Syndrome (WNT10A)	No disease-causing mutations detected
Omenn Syndrome and Severe Combined Immunodeficiency, Athabaskan-Type (DCLRE1C)	No disease-causing mutations detected
Omenn Syndrome, RAG2-Related (RAG2)	No disease-causing mutations detected

Ornithine Aminotransferase Deficiency (OAT)	No disease-causing mutations detected
Pontocerebellar Hypoplasia Type 1A (VRK1)	No disease-causing mutations detected
Pontocerebellar Hypoplasia Type 6 (RARS2)	No disease-causing mutations detected
Primary Ciliary Dyskinesia, DNAI1-Related (DNAI1)	No disease-causing mutations detected
Primary Ciliary Dyskinesia, DNAI2-Related (DNAI2)	No disease-causing mutations detected
Progressive Cerebello-Cerebral Atrophy (SEPSECS)	No disease-causing mutations detected
Progressive Familial Intrahepatic Cholestasis Type 2 (ABCB11)	No disease-causing mutations detected
Pyruvate Dehydrogenase E1-Alpha Deficiency (PDHA1)	No disease-causing mutations detected
Pyruvate Dehydrogenase E1-Beta Deficiency (PDHB)	No disease-causing mutations detected
Renal Tubular Acidosis and Deafness (ATP6V1B1)	No disease-causing mutations detected
Retinitis Pigmentosa 25 (EYS)	No disease-causing mutations detected
Retinitis Pigmentosa 26 (CERKL)	No disease-causing mutations detected
Retinitis Pigmentosa 28 (FAM161A)	No disease-causing mutations detected
Rhizomelic Chondrodysplasia Punctata Type 3 (AGPS)	No disease-causing mutations detected
Roberts Syndrome (ESCO2)	No disease-causing mutations detected
Schimke Immunoosseous Dysplasia (SMARCAL1)	No disease-causing mutations detected

Steel Syndrome (COL27A1)	No disease-causing mutations detected
Stuve-Wiedemann Syndrome (LIFR)	No disease-causing mutations detected
Usher Syndrome Type ID (CDH23)	No disease-causing mutations detected
Zellweger Syndrome Spectrum, PEX10- Related (PEX10)	No disease-causing mutations detected
Zellweger Syndrome Spectrum, PEX6-Related (PEX6)	No disease-causing mutations detected
17-Alpha-Hydroxylase-Deficient Congenital Adrenal Hyperplasia (CYP17A1)	No disease-causing mutations detected
21-Alpha-Hydroxylase-Deficient Congenital Adrenal Hyperplasia (CYP21A2)	No disease-causing mutations detected
3-Beta-Hydroxysteroid Dehydrogenase Type II Deficiency (HSD3B2)	No disease-causing mutations detected
3-Methylcrotonyl-CoA Carboxylase Deficiency, MCCC1-Related (MCCC1)	No disease-causing mutations detected
3-Methylcrotonyl-CoA Carboxylase Deficiency, MCCC2-Related (MCCC2)	No disease-causing mutations detected
3-Methylglutaconic Aciduria, Type III also known as Costeff Optic Atrophy Syndrome (OPA3)	No disease-causing mutations detected
3-Phosphoglycerate Dehydrogenase Deficiency, PHGDH-Related (PHGDH)	No disease-causing mutations detected
6-Pyruvoyl-Tetrahydropterin Synthase Deficiency (PTS)	No disease-causing mutations detected
Abetalipoproteinemia (MTTP)	No disease-causing mutations detected
Achromatopsia (CNGB3)	No disease-causing mutations detected
Acrodermatitis Enteropathica (SLC39A4)	No disease-causing mutations detected

Acute Infantile Liver Failure (TRMU)	No disease-causing mutations detected
Acyl-CoA Oxidase I Deficiency (ACOX1)	No disease-causing mutations detected
Adenosine Deaminase Deficiency (ADA)	No disease-causing mutations detected
Adrenoleukodystrophy, X-Linked (ABCD1)	No disease-causing mutations detected
Aicardi-Goutieres Syndrome (SAMHD1)	No disease-causing mutations detected
Alpha-Mannosidosis (MAN2B1)	No disease-causing mutations detected
Alpha-Thalassemia (HBA1/HBA2)	No disease-causing mutations detected; normal red blood cell indices and hemoglobin analysis
Alpha-Thalassemia Mental Retardation Syndrome (ATRX)	No disease-causing mutations detected
Alport Syndrome, COL4A3-Related (COL4A3)	No disease-causing mutations detected
Alport Syndrome, COL4A4-Related (COL4A4)	No disease-causing mutations detected
Alport Syndrome, X-Linked (COL4A5)	No disease-causing mutations detected
Alstrom Syndrome (ALMS1)	No disease-causing mutations detected
Andermann Syndrome (SLC12A6)	No disease-causing mutations detected
Argininosuccinic Aciduria (ASL)	No disease-causing mutations detected
Aromatase Deficiency (CYP19A1)	No disease-causing mutations detected

Arthrogryposis, Mental Retardation, and Seizures (SLC35A3)	No disease-causing mutations detected
Asparagine Synthetase Deficiency (ASNS)	No disease-causing mutations detected
Aspartylglucosaminuria (AGA)	No disease-causing mutations detected
Ataxia With Vitamin E Deficiency (TTPA)	No disease-causing mutations detected
Ataxia-Telangiectasia (ATM)	No disease-causing mutations detected
ATP7A-Related Disorders, X-Linked (ATP7A)	No disease-causing mutations detected
Bardet-Biedl Syndrome 13 also known as Meckel Syndrome 1 (MKS1)	No disease-causing mutations detected
Bardet-Biedl Syndrome, BBS1-Related (BBS1)	No disease-causing mutations detected
Bardet-Biedl Syndrome, BBS10-Related (BBS10)	No disease-causing mutations detected
Bardet-Biedl Syndrome, BBS12-Related (BBS12)	No disease-causing mutations detected
Bardet-Biedl Syndrome, BBS2-Related (BBS2)	No disease-causing mutations detected
Bare Lymphocyte Syndrome, Type II (CIITA)	No disease-causing mutations detected
Bartter Syndrome, Type 4A (BSND)	No disease-causing mutations detected
Bernard-Soulier Syndrome, Type A1 (GP1BA)	No disease-causing mutations detected
Bernard-Soulier Syndrome, Type C (GP9)	No disease-causing mutations detected
Beta-Ketothiolase Deficiency (ACAT1)	No disease-causing mutations detected

Beta-Thalassemia, Sickle Cell Disease, Beta-Globin-Related Hemoglobinopathies (HBB)	No disease-causing mutations detected; normal red blood cell indices and hemoglobin analysis
Bilateral Frontoparietal Polymicrogyria (GPR56)	No disease-causing mutations detected
Biotinidase Deficiency (BTD)	No disease-causing mutations detected
Bloom Syndrome (BLM)	No disease-causing mutations detected
Canavan Disease (ASPA)	No disease-causing mutations detected
Carbamoylphosphate Synthetase I Deficiency (CPS1)	No disease-causing mutations detected
Carnitine Palmitoyltransferase IA Deficiency (CPT1A)	No disease-causing mutations detected
Carnitine Palmitoyltransferase II Deficiency (CPT2)	No disease-causing mutations detected
Carpenter Syndrome (RAB23)	No disease-causing mutations detected
Cartilage-Hair Hypoplasia (RMRP)	No disease-causing mutations detected
Cerebral Creatine Deficiency Syndrome 1 (SLC6A8)	No disease-causing mutations detected
Cerebral Creatine Deficiency Syndrome 2 (GAMT)	No disease-causing mutations detected
Cerebrotendinous Xanthomatosis (CYP27A1)	No disease-causing mutations detected
Charcot-Marie-Tooth Disease, Type 4D (NDRG1)	No disease-causing mutations detected
Charcot-Marie-Tooth Disease, Type 5 / Arts Syndrome (PRPS1)	No disease-causing mutations detected

Charcot-Marie-Tooth Disease, X-Linked (GJB1)	No disease-causing mutations detected
Choreoacanthocytosis (VPS13A)	No disease-causing mutations detected
Choroidemia (CHM)	No disease-causing mutations detected
Chromosomal Analysis	Normal male karyotype
Chronic Granulomatous Disease, CYBA-Related (CYBA)	No disease-causing mutations detected
Chronic Granulomatous Disease, CYBB-Related (CYBB)	No disease-causing mutations detected
Citrin Deficiency (SLC25A13)	No disease-causing mutations detected
Citrullinemia Type 1 (ASS1)	No disease-causing mutations detected
Cohen Syndrome (VPS13B)	No disease-causing mutations detected
Combined Malonic and Methylmalonic Aciduria (ACSF3)	No disease-causing mutations detected
Combined Oxidative Phosphorylation Deficiency 1 (GFM1)	No disease-causing mutations detected
Combined Oxidative Phosphorylation Deficiency 3 (TSFM)	No disease-causing mutations detected
Combined Pituitary Hormone Deficiency 2 (PROP1)	No disease-causing mutations detected
Combined Pituitary Hormone Deficiency 3 (LHX3)	No disease-causing mutations detected
Combined SAP Deficiency (PSAP)	No disease-causing mutations detected
Congenital Amegakaryocytic Thrombocytopenia (MPL)	No disease-causing mutations detected
Congenital Disorder of Glycosylation Type 1a (PMM2)	No disease-causing mutations detected

Congenital Disorder of Glycosylation Type 1b (MPI)	No disease-causing mutations detected
Congenital Disorder of Glycosylation, Type Ic (ALG6)	No disease-causing mutations detected
Congenital Insensitivity to Pain with Anhidrosis (NTRK1)	No disease-causing mutations detected
Congenital Myasthenic Syndrome, CHRNE-Related (CHRNE)	No disease-causing mutations detected
Congenital Myasthenic Syndrome, RAPSN-Related (RAPSN)	No disease-causing mutations detected
Congenital Neutropenia, HAX1-Related (HAX1)	No disease-causing mutations detected
Congenital Neutropenia, VPS45-Related (VPS45)	No disease-causing mutations detected
Corneal Dystrophy and Perceptive Deafness (SLC4A11)	No disease-causing mutations detected
Corticosterone Methyloxidase Deficiency (CYP11B2)	No disease-causing mutations detected
Cystic Fibrosis (CFTR)	No disease-causing mutations detected
Cystinosis (CTNS)	No disease-causing mutations detected
D-Bifunctional Protein Deficiency (HSD17B4)	No disease-causing mutations detected
Deafness, Autosomal Recessive 77 (LOXHD1)	No disease-causing mutations detected
Dyskeratosis Congenita, RTEL1-Related (RTEL1)	No disease-causing mutations detected
Dystrophinopathies, X-Linked (DMD)	No disease-causing mutations detected
Ehlers-Danlos Syndrome, Type VIIC (ADAMTS2)	No disease-causing mutations detected

Ellis-Van Creveld Syndrome, EVC-Related (EVC)	No disease-causing mutations detected
Ethylmalonic Encephalopathy (ETHE1)	No disease-causing mutations detected
Fabry Disease, X-Linked (GLA)	No disease-causing mutations detected
Factor XI Deficiency (F11)	No disease-causing mutations detected
Familial Dysautonomia (IKBKAP)	No disease-causing mutations detected
Familial Hyperinsulinism, ABCC8-Related (ABCC8)	No disease-causing mutations detected
Familial Hyperinsulinism, KCNJ11-Related (KCNJ11)	No disease-causing mutations detected
Familial Mediterranean Fever (MEFV)	No disease-causing mutations detected
Fanconi Anemia Group A (FANCA)	No disease-causing mutations detected
Fanconi Anemia Group C (FANCC)	No disease-causing mutations detected
Fragile X Syndrome (FMR1)	No disease-causing mutations detected; CGG repeat number within normal range
Galactokinase Deficiency (GALK1)	No disease-causing mutations detected
Galactosemia, GALT-Related (GALT)	No disease-causing mutations detected
Gaucher Disease (GBA)	No disease-causing mutations detected
GJB2-Related DFNB1 Nonsyndromic Hearing Loss and Deafness (GJB2)	No disease-causing mutations detected
Glutaric Acidemia Type 1 (GCDH)	No disease-causing mutations detected

Glycine Encephalopathy, AMT-Related (AMT)	No disease-causing mutations detected
Glycine Encephalopathy, GLDC-Related (GLDC)	No disease-causing mutations detected
Glycogen Storage Disease Type Ia (G6PC)	No disease-causing mutations detected
Glycogen Storage Disease Type Ib (SLC37A4)	No disease-causing mutations detected
Glycogen Storage Disease Type III (AGL)	No disease-causing mutations detected
Glycogen Storage Disease Type V (PYGM)	No disease-causing mutations detected
GRACILE Syndrome (BCS1L)	No disease-causing mutations detected
Hereditary Fructose Intolerance (ALDOB)	No disease-causing mutations detected
HMG-CoA Lyase Deficiency (HMGCL)	No disease-causing mutations detected
Holocarboxylase Synthetase Deficiency (HLCS)	No disease-causing mutations detected
Homocystinuria, CBS-Related (CBS)	No disease-causing mutations detected
Hydrolethalus Syndrome (HYLS1)	No disease-causing mutations detected
Hypophosphatasia, Autosomal Recessive (ALPL)	No disease-causing mutations detected
Inclusion Body Myopathy 2 (GNE)	No disease-causing mutations detected
Isovaleric Acidemia (IVD)	No disease-causing mutations detected
Joubert Syndrome 2 (TMEM216)	No disease-causing mutations detected

Joubert Syndrome 7 also known as Meckel Syndrome 5 (RPGRIP1L)	No disease-causing mutations detected
Junctional Epidermolysis Bullosa, LAMA3-Related (LAMA3)	No disease-causing mutations detected
Junctional Epidermolysis Bullosa, LAMB3-Related (LAMB3)	No disease-causing mutations detected
Junctional Epidermolysis Bullosa, LAMC2-Related (LAMC2)	No disease-causing mutations detected
Juvenile Retinoschisis, X-Linked (RS1)	No disease-causing mutations detected
Krabbe Disease (GALC)	No disease-causing mutations detected
Leigh Syndrome, French-Canadian Type (LRPPRC)	No disease-causing mutations detected
Limb-Girdle Muscular Dystrophy Type 2A also known as Calpainopathy (CAPN3)	No disease-causing mutations detected
Limb-Girdle Muscular Dystrophy Type 2B also known as Dysferlinopathy (DYSF)	No disease-causing mutations detected
Limb-Girdle Muscular Dystrophy Type 2C also known as Gamma-Sarcoglycanopathy (SGCG)	No disease-causing mutations detected
Limb-Girdle Muscular Dystrophy Type 2D also known as Alpha-Sarcoglycanopathy (SGCA)	No disease-causing mutations detected
Limb-Girdle Muscular Dystrophy Type 2E also known as Beta-Sarcoglycanopathy (SGCB)	No disease-causing mutations detected
Limb-Girdle Muscular Dystrophy Type 2I (FKRP)	No disease-causing mutations detected
Lipoid Adrenal Hyperplasia (STAR)	No disease-causing mutations detected
Long-Chain 3-Hydroxyacyl-CoA Dehydrogenase Deficiency (HADHA)	No disease-causing mutations detected
Maple Syrup Urine Disease Type 1A (BCKDHA)	No disease-causing mutations detected

Maple Syrup Urine Disease Type 1B (BCKDHB)	No disease-causing mutations detected
Maple Syrup Urine Disease Type 3 also known as Lipoamide Dehydrogenase Deficiency (DLD)	No disease-causing mutations detected
Medium Chain Acyl-CoA Dehydrogenase Deficiency (ACADM)	No disease-causing mutations detected
Megalencephalic Leukoencephalopathy With Subcortical Cysts (MLC1)	No disease-causing mutations detected
Metachromatic Leukodystrophy (ARSA)	No disease-causing mutations detected
Methylmalonic Acidemia, MMAA-Related (MMAA)	No disease-causing mutations detected
Methylmalonic Acidemia, MMAB-Related (MMAB)	No disease-causing mutations detected
Methylmalonic Acidemia, MUT-Related (MUT)	No disease-causing mutations detected
Methylmalonic Aciduria and Homocystinuria, Cobalamin C Type (MMACHC)	No disease-causing mutations detected
Methylmalonic Aciduria and Homocystinuria, Cobalamin D Type (MMADHC)	No disease-causing mutations detected
Mucolipidosis II and IIIA (GNPTAB)	No disease-causing mutations detected
Mucolipidosis III Gamma (GNPTG)	No disease-causing mutations detected
Mucolipidosis Type IV (MCOLN1)	No disease-causing mutations detected
Mucopolysaccharidosis Type I also known as Hurler Syndrome (IDUA)	No disease-causing mutations detected
Mucopolysaccharidosis Type IVb also known as GM1 Gangliosidosis (GLB1)	No disease-causing mutations detected
Mucopolysaccharidosis, Type II, X-Linked (IDS)	No disease-causing mutations detected

Mucopolysaccharidosis, Type IIIA (SGSH)	No disease-causing mutations detected
Mucopolysaccharidosis, Type IIIB (NAGLU)	No disease-causing mutations detected
Mucopolysaccharidosis, Type IIIC (HGSNAT)	No disease-causing mutations detected
Multiple Sulphatase Deficiency (SUMF1)	No disease-causing mutations detected
Muscle-Eye-Brain Disease (POMGNT1)	No disease-causing mutations detected
Myotubular Myopathy, X-Linked (MTM1)	No disease-causing mutations detected
Nemaline Myopathy 2 (NEB)	No disease-causing mutations detected
Nephrotic Syndrome, NPHS1-Related also known as Congenital Finnish Nephrosis (NPHS1)	No disease-causing mutations detected
Nephrotic Syndrome, NPHS2-Related also known as Steroid-Resistant Nephrotic Syndrome (NPHS2)	No disease-causing mutations detected
Neuronal Ceroid-Lipofuscinosis, CLN3-Related (CLN3)	No disease-causing mutations detected
Neuronal Ceroid-Lipofuscinosis, CLN5-Related (CLN5)	No disease-causing mutations detected
Neuronal Ceroid-Lipofuscinosis, CLN6-Related (CLN6)	No disease-causing mutations detected
Neuronal Ceroid-Lipofuscinosis, CLN8-Related also known as Northern Epilepsy (CLN8)	No disease-causing mutations detected
Neuronal Ceroid-Lipofuscinosis, PPT1-Related (PPT1)	No disease-causing mutations detected
Neuronal Ceroid-Lipofuscinosis, TPP1-Related (TPP1)	No disease-causing mutations detected
Niemann-Pick Disease Type C, NPC1-Related (NPC1)	No disease-causing mutations detected

Niemann-Pick Disease Type C, NPC2-Related (NPC2)	No disease-causing mutations detected
Niemann-Pick Disease, SMPD1-Related (SMPD1)	No disease-causing mutations detected
Nijmegen Breakage Syndrome (NBN)	No disease-causing mutations detected
Ornithine Transcarbamylase Deficiency, X-Linked (OTC)	No disease-causing mutations detected
Osteochondrodysplasias, Sulfate Transporter-Related (SLC26A2)	No disease-causing mutations detected
Osteopetrosis Type 1 (TCIRG1)	No disease-causing mutations detected
Pendred Syndrome (SLC26A4)	No disease-causing mutations detected
Phenylalanine Hydroxylase Deficiency also known as Phenylketonuria (PAH)	No disease-causing mutations detected
Polycystic Kidney Disease, Autosomal Recessive (PKHD1)	No disease-causing mutations detected
Polyglandular Autoimmune Syndrome Type 1 (AIRE)	No disease-causing mutations detected
Pompe Disease also known as Glycogen Storage Disease Type II (GAA)	No disease-causing mutations detected
Primary Carnitine Deficiency (SLC22A5)	No disease-causing mutations detected
Primary Ciliary Dyskinesia, DNAH5-Related (DNAH5)	No disease-causing mutations detected
Primary Hyperoxaluria Type 1 (AGXT)	No disease-causing mutations detected
Primary Hyperoxaluria Type 2 (GRHPR)	No disease-causing mutations detected
Primary Hyperoxaluria, Type 3 (HOGA1)	No disease-causing mutations detected

Propionic Acidemia, PCCA-Related (PCCA)	No disease-causing mutations detected
Propionic Acidemia, PCCB-Related (PCCB)	No disease-causing mutations detected
Pycnodysostosis (CTSK)	No disease-causing mutations detected
Retinitis Pigmentosa 59 (DHDDS)	No disease-causing mutations detected
Rhizomelic Chondrodysplasia Punctata Type 1 (PEX7)	No disease-causing mutations detected
Salla Disease (SLC17A5)	No disease-causing mutations detected
Sandhoff Disease (HEXB)	No disease-causing mutations detected
Segawa Syndrome (TH)	No disease-causing mutations detected
Severe Combined Immunodeficiency, X-Linked (IL2RG)	No disease-causing mutations detected
Sjogren-Larsson Syndrome (ALDH3A2)	No disease-causing mutations detected
Smith-Lemli-Opitz Syndrome (DHCR7)	No disease-causing mutations detected
Spastic Ataxia of Charlevoix-Saguenay, Autosomal Recessive (SACS)	No disease-causing mutations detected
Spinal Muscular Atrophy (SMN1)	No disease-causing mutations detected (2 copies detected)
Spondylothoracic Dysostosis (MESP2)	No disease-causing mutations detected
Tay-Sachs Disease (HEXA)	No disease-causing mutations detected; normal enzyme analysis

TGM1-Related Autosomal Recessive Congenital Ichthyosis (TGM1)	No disease-causing mutations detected
Tyrosinemia Type 1 (FAH)	No disease-causing mutations detected
Usher Syndrome Type IB (MYO7A)	No disease-causing mutations detected
Usher Syndrome Type IC (USH1C)	No disease-causing mutations detected
Usher Syndrome Type IF (PCDH15)	No disease-causing mutations detected
Usher Syndrome Type IIA (USH2A)	No disease-causing mutations detected
Usher Syndrome Type III (CLRN1)	No disease-causing mutations detected
Vanishing White Matter Disease (EIF2B5)	No disease-causing mutations detected
Very Long Chain Acyl-CoA Dehydrogenase Deficiency (ACADVL)	No disease-causing mutations detected
Walker-Warburg Syndrome, FKTN-Related (FKTN)	No disease-causing mutations detected
Wilson Disease (ATP7B)	No disease-causing mutations detected
Wolman Disease also known as Cholesteryl Ester Storage Disease (LIPA)	No disease-causing mutations detected
Zellweger Spectrum Disorder, PEX1-Related (PEX1)	No disease-causing mutations detected
Zellweger Spectrum Disorder, PEX2-Related (PEX2)	No disease-causing mutations detected

Health Information

Medication Allergy:	No	**Eyesight Correction:**	No
Food Allergy:	No	**Near or Far Sighted:**	
Pet Allergy:	No	**Skin Infection:**	No
Hay Fever Allergy:	No	**Gallstones:**	No
Insect Allergy:	No	**Removed Gall Bladder:**	No
Vaccine Allergy:	No	**Hernia:**	No
Healthy Teeth:	Yes	**Mumps:**	No
Braces:	No	**Measles:**	No
Back Problems:	No	**Measles Age:**	
Bronchitis:	Yes	**German Measles:**	No
Chicken Pox:	No	**German Measles Age:**	
Chicken Pox Age:		**Sinus Infection:**	Yes
Vertigo:	Yes	**Stomach Ulcers:**	No

Note: The medical and social history was provided by the donor and cannot be verified for accuracy.

Personality and Interest Attributes

Athletic	1	2	3	4	5	**Couch Potato**
Optimistic	1	2	3	4	5	**Pessimistic**
Assertive	1	2	3	4	5	**Passive**
Leader	1	2	3	4	5	**Follower**

Easy Going	1	2	**3**	4	5	Controlling, Rigid
Artistic	**1**	2	3	4	5	Michelangelo Who?
Extrovert	1	2	**3**	4	5	Introvert

Favorite Hero:	Clint Eastwood. My dad showed me his movies when I was young, and I loved how tough he was & how he wasn't afraid of anyone.
Awards:	Honor student in high school
Perfect Day:	Wake up early, make breakfast & coffee, start working on a song/project, finish the night off seeing my favorite band.
Personality:	Easy going, to the point, love being around friends/people, uplifting/positive, sometimes can be shy if I'm by myself around many people.
Accomplishments:	One of my first gigs, my buddy and I opened up for a renowned artist. It was an amazing experience getting to play before him and meet him.
Character Flaws:	Can be too nice/forgiving, sometimes I overload my schedule, I remember everything but also forget things.
Favorite Music:	Hard To Say
Favorite Music Explanation:	I am a producer of electronic music, but I also love jam music just as much.
Favorite Book:	The Raven
Favorite Author:	Edgar Allan Poe
Favorite Author/Book Explanation:	I've always been into horror movies and his books/poems were the only literature I thought was interesting in high school.
Celebrity Look Alike:	Johnny Tsunami
Vegetarian:	No
Adjectives:	Dedicated, Understanding, Passionate

Fine Art Skills:	Music production
Technical/Mechanical Ability:	In school for audio engineering, working as an electrician in training (lighting)
Skills, Hobbies and Interest:	I Produce And Write Music

Keirsey Profile Assessment

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Birth & Education

Carried To Term:	No	**Birth Length:**	Not Available
Pregnancy Complications:	No	**Twin:**	No
Birth Weight:	8 lbs	**Twin Type:**	
Childhood Health:	Excellent		

In School:	Yes
Did the donor graduate from high school?	Yes
Did the donor receive a diploma?	Yes
Is the donor currently enrolled in or does the donor hold an undergraduate degree?	Yes
Degrees Earned:	Audio Engineer
Degree Status:	Currently Enrolled
Major: Audio Technology	**Minor:** N/a
Is the donor currently enrolled in or does the donor hold a graduate degree?	No

Is the donor currently enrolled in or does the donor hold a post graduate degree?	No
Is the donor currently enrolled in or does the donor have any specialized training (Including Military, Police, Firefighter, EMS, Real Estate, etc.)?	No

Health & Diseases

The following medical conditions apply to the donor and his BLOOD RELATIVES ONLY (grandparents, parents, aunts, uncles, cousins, brothers, sisters, nieces, nephews, and children of the donor).

Autoimmune Diseases	Yes	No	Which relative	Father's side	Mother's side	Age of onset
Crest Syndrome		●				
Scleroderma		●				
Lupus		●				
Sjogren's Syndrome		●				

Blood Diseases	Yes	No	Which relative	Father's side	Mother's side	Age of onset
Anemia		●				
Fanconi anemia		●				
Hemophilia		●				
Immunodeficiency		●				
Leukemia		●				
Sickle cell anemia		●				
Thalassemia		●				
VonWillebrand's disease		●				

Cancer	Yes	No	Which relative	Father's side	Mother's side	Age of onset
Breast		●				
Colon		●				
Leukemia		●				
Lung		●				
Lymphoma		●				
Melanoma		●				
Skin		●				
Cardiovascular Diseases	**Yes**	**No**	**Which relative**	**Father's side**	**Mother's side**	**Age of onset**
Heart Attack		●				
Heart disease(congenital or other)		●				
High Blood Pressure		●				
High Cholesterol/Triglycerides		●				
Stroke		●				
Congenital Malformations	**Yes**	**No**	**Which relative**	**Father's side**	**Mother's side**	**Age of onset**
Cleft palate		●				
Cleft lip		●				
Club foot		●				
Hypospadia		●				
Polydactyly		●				
Undescended testicles		●				

Gastro-Intestinal Diseases	Yes	No	Which relative	Father's side	Mother's side	Age of onset
Crohns disease		●				
Diverticulitis		●				
Gall Stones		●				
Hemochromatosis		●				
Hepatitis		●				
Pyloric Stenosis		●				
Ulcers		●				
Ulcerative Colitis		●				

Kidney Disease	Yes	No	Which relative	Father's side	Mother's side	Age of onset
Adrenal hyperplasia		●				
Born with only 1 kidney		●				
Kidney disease or urinary tract dis.		●				
Polycystic kidney disease		●				
Progressive kidney disease		●				

Mental Health	Yes	No	Which relative	Father's side	Mother's side	Age of onset
Anxiety disorder		●				
Depressions(severe		●				
Manic-depressive (bipolar) disorder		●				
Obsessive-compulsive disorder		●				
Schizophrenia		●				

Metabolic/Endrocrine Dis.	Yes	No	Which relative	Father's side	Mother's side	Age of onset
Diabetes(Insulin Dependent)		●				
Diabetes(Non-Insulin Dependent)		●				
Galactosemia		●				
Gauchers disease		●				
Goiter		●				
Hypoglycemia		●				
Maple Syrup Disease		●				
PKU		●				
Tay-Sachs		●				
Thyroid disease		●				

Muscular/Bones/Joint Disease	Yes	No	Which relative	Father's side	Mother's side	Age of onset
Achondroplasia (Dwarfism)		●				
Arthritis: osteoarthritis		●				
Arthritis: Rheumatoid		●				
Arthritis: Other		●				
Congenital hip disease		●				
Gout		●				
Loss of muscle coordination		●				
Marfan's syndrome		●				
Muscular dystrophy		●				
Osteoporosis		●				

Scoliosis		●				
Spinal Muscular Atrophy		●				

Neurological Disease	Yes	No	Which relative	Father's side	Mother's side	Age of onset
Alzheimers		●				
Autism/Aspergers		●				
Canavan's disease		●				
Cerebral palsy		●				
CJD (Creutzfeld-Jacob Disease)		●				
Dementia		●				
Disorders of the spinal cord		●				
Epilepsy		●				
Familial dysautonomia		●				
Guillain-Barre		●				
Huntington's		●				
JC virus		●				
Learning disorder:ADD		●				
Lesch-Nyhan		●				
Lou Gehrig's disease		●				
Mental Retardation		●				
Migraines		●				
Mucolipidosis type IV		●				
Multiple Sclerosis		●				

Nieman-Pick		●				
Neurofibromatosis		●				
Neural tube disorder(spinal bifida)		●				
Parkinson's disease		●				
Spongiform encephalopathy/prion disease		●				
Subacute sclerosing panencephalitis		●				
Tourette Syndrome		●				
Respiratory Disease	**Yes**	**No**	**Which relative**	**Father's side**	**Mother's side**	**Age of onset**
Allergies: Hay fever		●				
Asthma		●				
Emphysema		●				
SARS		●				
TB		●				
Sight/Sound/Smell	**Yes**	**No**	**Which relative**	**Father's side**	**Mother's side**	**Age of onset**
Blindness		●				
Cataracts		●				
Color Blindness		●				
Deafness (before age 50)		●				
Ear deformity		●				
Glaucoma		●				
Macular degeneration		●				

Retinoblastoma		●				

Skin	Yes	No	Which relative	Father's side	Mother's side	Age of onset
Albinism		●				
Eczema		●				
Psoriasis		●				
Pigmentation disorders(including Vitiligo)		●				

Other	Yes	No	Which relative	Father's side	Mother's side	Age of onset
Alcoholism		●				
Bloom syndrome		●				
Cystic fibrosis		●				
Down syndrome		●				
Drug abuse		●				
Encephalitis: viral or of unknown origin		●				
Exposure to: radiation		●				
Explosuer To: Toxic chemicals		●				
Fragile X		●				
Klinefelter		●				
Meningitis		●				
Noonan syndrome		●				
SIDS		●				
Turner		●				

West Nile(suspected or confirmed by lab testing)		•				
Other genetic condition		•				
Premature degeneration of any organ		•				

Immediate Family Medical History

Family Member:	Father	**Hair Color:**	Brown, Dark
Eyesight:		**Freckles:**	None
Hair Texture:	Thin	**Health Status:**	Good
Skin Tone:	Medium	**Dominant Hand:**	
Year of Birth:	1969	**Cause of Death:**	
Hair Loss:		**Height:**	6' 02" (187.96 cm)
Age At Death:		**Body Hair:**	
Weight:	230 lbs. (104 kg)	**Occupation:**	Corporate
Eye Color:	Brown, Light	**Half Siblings:**	
Education:	College (Some)	**Body Build:**	Large
Half Siblings Parent:		**Eyesight Correction:**	
Near or Far Sighted:			

Comments: Has corrected vision. He's a great father who raised me to speak with respect and how to logically breakdown a situation. He's a handyman with cars and motors in general. Enjoys fishing and golf.

Family Member:	Mother	**Hair Color:**	Black
Eyesight:		**Freckles:**	None
Hair Texture:	Thick	**Health Status:**	Good
Skin Tone:	Brown, Medium	**Dominant Hand:**	
Year of Birth:	1965	**Cause of Death:**	
Hair Loss:		**Height:**	5' " (152.40 cm)
Age At Death:		**Body Hair:**	Wavy
Weight:	120 lbs. (54 kg)	**Occupation:**	Manager
Eye Color:	Brown, Dark	**Half Siblings:**	
Education:	College (Some)	**Body Build:**	Small
Half Siblings Parent:		**Eyesight Correction:**	
Near or Far Sighted:			

Comments: Has corrected vision and dimples. She’s the sweetest lady I know, always had my back, even when I said I was going to leave home and produce music. Enjoys golf, sewing, and crafts.

Family Member:	Brother
Eyesight:	
Hair Texture:	Thick
Skin Tone:	Brown, Dark
Year of Birth:	1985
Hair Loss:	
Age At Death:	
Weight:	190 lbs. (86 kg)
Eye Color:	Brown, Dark
Education:	College (Some)
Half Siblings Parent:	Maternal
Near or Far Sighted:	

Hair Color:	Black
Freckles:	None
Health Status:	Good
Dominant Hand:	
Cause of Death:	
Height:	5' 10" (177.80 cm)
Body Hair:	Wavy
Occupation:	Bar Manager
Half Siblings:	Yes
Body Build:	Medium
Eyesight Correction:	

Comments: I get my witty sarcasm from him, he has the biggest heart known to man. Enjoys surfing, construction, and agriculture cultivation.

Family Member:	Sister
Eyesight:	
Hair Texture:	Thick
Skin Tone:	Brown, Medium
Year of Birth:	1989
Hair Loss:	
Age At Death:	
Weight:	135 lbs. (61 kg)
Eye Color:	Brown, Dark
Education:	Masters
Half Siblings Parent:	Maternal
Near or Far Sighted:	

Hair Color:	Black
Freckles:	None
Health Status:	Good
Dominant Hand:	
Cause of Death:	
Height:	5' 02" (157.48 cm)
Body Hair:	Wavy
Occupation:	Teacher
Half Siblings:	Yes
Body Build:	Medium
Eyesight Correction:	

Comments: Has corrected vision. A wonderful mother, toughest person I know. Enjoys math and coaches cheerleading (she cheered her entire life).

♂ Paternal Family Medical History

Family Member:	Grandfather
Eyesight:	
Hair Texture:	Thin
Skin Tone:	Fair
Year of Birth:	1948
Hair Loss:	
Age At Death:	
Weight:	200 lbs. (90 kg)
Eye Color:	Brown, Light
Education:	College
Half Siblings Parent:	
Near or Far Sighted:	

Hair Color:	Brown, Dark
Freckles:	None
Health Status:	Fair
Dominant Hand:	
Cause of Death:	
Height:	6' 01" (185.42 cm)
Body Hair:	
Occupation:	Title Loan Establishments-Owner, Retired
Half Siblings:	
Body Build:	Medium
Eyesight Correction:	

Comments: Has corrected vision. Raised on respect and fairness.

Family Member:	Grandmother
Eyesight:	
Hair Texture:	
Skin Tone:	Fair
Year of Birth:	1948
Hair Loss:	
Age At Death:	
Weight:	180 lbs. (81 kg)
Eye Color:	Hazel/Green
Education:	College (Some)
Half Siblings Parent:	
Near or Far Sighted:	

Hair Color:	Brown, Dark
Freckles:	Numerous
Health Status:	Good
Dominant Hand:	
Cause of Death:	
Height:	6' " (182.88 cm)
Body Hair:	Wavy
Occupation:	Partial Owner of Title Loan Establishment, Retired
Half Siblings:	
Body Build:	Medium
Eyesight Correction:	

Comments: Has corrected vision. Always willing to help. Loves cooking.

Family Member:	Aunt	**Hair Color:**	
Eyesight:		**Freckles:**	
Hair Texture:		**Health Status:**	Good
Skin Tone:		**Dominant Hand:**	
Year of Birth:	1980	**Cause of Death:**	
Hair Loss:		**Height:**	
Age At Death:		**Body Hair:**	
Weight:		**Occupation:**	
Eye Color:		**Half Siblings:**	
Education:		**Body Build:**	
Half Siblings Parent:		**Eyesight Correction:**	
Near or Far Sighted:			

Comments:

Maternal Family Medical History

Family Member:	Grandfather	**Hair Color:**	Black
Eyesight:		**Freckles:**	None
Hair Texture:		**Health Status:**	Fair
Skin Tone:	Brown, Dark	**Dominant Hand:**	
Year of Birth:	1944	**Cause of Death:**	
Hair Loss:		**Height:**	6' " (182.88 cm)
Age At Death:		**Body Hair:**	
Weight:	200 lbs. (90 kg)	**Occupation:**	
Eye Color:	Brown, Dark	**Half Siblings:**	
Education:		**Body Build:**	Medium
Half Siblings Parent:		**Eyesight Correction:**	
Near or Far Sighted:			

Comments: Raised my mom to be loving.

Family Member:	Grandmother	**Hair Color:**	Black
Eyesight:		**Freckles:**	None
Hair Texture:		**Health Status:**	Fair
Skin Tone:	Brown, Dark	**Dominant Hand:**	
Year of Birth:	1946	**Cause of Death:**	
Hair Loss:		**Height:**	
Age At Death:		**Body Hair:**	
Weight:		**Occupation:**	
Eye Color:	Brown, Dark	**Half Siblings:**	
Education:		**Body Build:**	Small
Half Siblings Parent:		**Eyesight Correction:**	
Near or Far Sighted:			

Comments: Has corrected vision. Taught my mom how to be a very strong woman.