Donor 5533

Information displayed in this profile is provided by the donor.
Blank fields are intentional and indicate that the information is not available.

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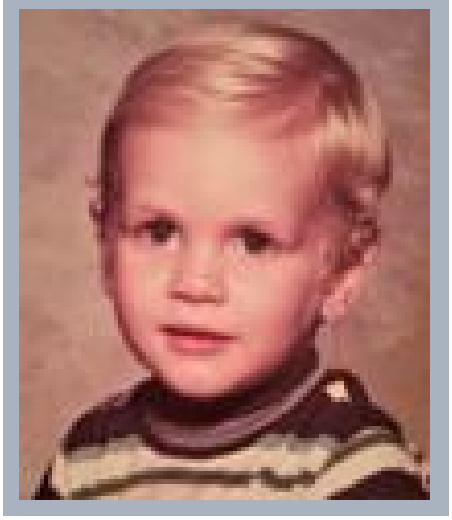
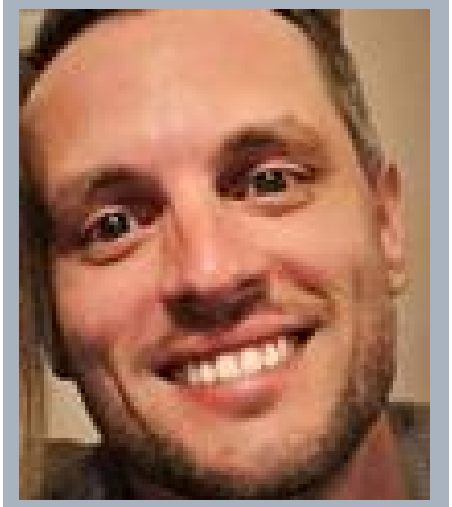
General Information

Donor ID:	5533
Year of Birth:	1981
Marital Status:	Single
Number of Children:	1
Religion:	None
Occupation:	Renewal Specialist
Blood Type:	AB -
Nationality:	Maternal: French, Paternal: English
Race:	
CMV Status:	POS
Reported Pregnancy:	Yes
Canadian Compliant:	Yes
UK Compliant:	Yes
Identity Disclosure:	Yes
Last Medical History Update:	09/23/2019
Audio File:	Not Available

Child Photo



Adult Photo



Marital Status: Single
Number of Children: 1
Religion: None
Occupation: Renewal Specialist **Blood Type:** AB - **Ethnicity:** Maternal: French, Paternal: English
Audio File:

Donor Photos

Number of photos available: Not available at this time.

Physical Attributes

Height:	6'2" (187.96cm)	**Eyebrows:**	Medium
Weight:	178 lbs (80kg)	**Dimples:**	No
Eye Color:	Blue	**Acne:**	No
Hair Color:	Brown	**Acne Information:**	
Hair Texture:	Average	**Shoe Size:**	10.5
Hair Loss:	None	**Body Build:**	Medium
Hair Type:	Average	**Freckles:**	None
Dominant Hand:	Right	**Skin Tone:**	Fair
Hairy Chest:	No	**Face Shape:**	Oval
Hairy:	No	**Lips:**	Normal
Ear Lobes:	Detached	**Nose Shape:**	Normal
Beard Color:	Brown, Dark	**Long Eyelashes:**	Yes

In His Own Words

I was born and raised in the south. As a kid, my family and I moved quite a few times before settling in one place through middle and high school. I got very good at making new friends. It helped me become the extrovert I am today. I had a great childhood growing up with loving parents and family members who have always been there for me.
I have always been athletic and loved to play baseball and basketball. I am also a self-taught sketch artist. I am very patient and kind, and pride myself on being loyal, punctual and genuine to everyone. One of my biggest influences in life has been Michael Jordan. He has been my hero ever since I was old enough to watch basketball. He taught me to always strive to be the best at whatever it was that I wanted to do in life. I have had many accomplishments, but one that I am most proud of is serving in the military. It was an

amazing experience. Some advice that I can give that has always helped me in life and has molded me into the person I am today would be to never put off tomorrow what can be done today. Do what makes you happy and be kind and genuine to everyone, no matter what.

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. "xyGene" donors have undergone the most extensive testing. 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.

Congenital Disorder of Glycosylation, Type Ic (ALG6)	Carrier
Galactosemia, GALT-Related (GALT)	Carrier of Duarte variant
Zellweger Syndrome Spectrum, PEX10-Related (PEX10)	No disease-causing mutations detected
Zellweger Syndrome Spectrum, PEX6-Related (PEX6)	No disease-causing mutations detected
11-Beta-Hydroxylase-Deficient Congenital Adrenal Hyperplasia (CYP11B1)	No disease-causing mutations detected
21-Alpha-Hydroxylase-Deficient Congenital Adrenal Hyperplasia (CYP21A2)	No disease-causing mutations detected
3-Methylglutaconic Aciduria, Type III also known as Costeff Optic Atrophy Syndrome (OPA3)	No disease-causing mutations detected
6-Pyruvoyl-Tetrahydropterin Synthase Deficiency (PTS)	No disease-causing mutations detected
Adenosine Deaminase Deficiency (ADA)	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
Alport Syndrome, COL4A3-Related (COL4A3)	No disease-causing mutations detected
Alport Syndrome, COL4A4-Related (COL4A4)	No disease-causing mutations detected
Alstrom Syndrome (ALMS1)	No disease-causing mutations detected
Andermann Syndrome (SLC12A6)	No disease-causing mutations detected
Argininemia (ARG1)	No disease-causing mutations detected
Argininosuccinic Aciduria (ASL)	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
Beta-Thalassemia, Sickle Cell Disease, Beta-Globin-Related Hemoglobinopathies (HBB)	No disease-causing mutations detected; normal red blood cell indices and hemoglobin analysis
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
Cartilage-Hair Hypoplasia (RMRP)	No disease-causing mutations detected
Cerebrotendinous Xanthomatosis (CYP27A1)	No disease-causing mutations detected
Chromosomal Analysis	Normal male karyotype
Citrullinemia Type 1 (ASS1)	No disease-causing mutations detected
Cohen Syndrome (VPS13B)	No disease-causing mutations detected
Combined Pituitary Hormone Deficiency 2 (PROP1)	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
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
Delta-Sarcoglycanopathy (SGCD)	No disease-causing mutations detected
Dyskeratosis Congenita, RTEL1-Related (RTEL1)	No disease-causing mutations detected
Dystrophinopathies, X-Linked (DMD)	No disease-causing mutations detected
Ellis-Van Creveld Syndrome, EVC-Related (EVC)	No disease-causing mutations detected
Ellis-Van Creveld Syndrome, EVC2-Related (EVC2)	No disease-causing mutations detected
ERCC6-Related Disorders (ERCC6)	No disease-causing mutations detected
ERCC8-Related Disorders (ERCC8)	No disease-causing mutations detected
Fabry Disease, X-Linked (GLA)	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
Galactokinase Deficiency (GALK1)	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
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
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
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
Maple Syrup Urine Disease, Type II (DBT)	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
Merosin-Deficient Muscular Dystrophy (LAMA2)	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
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
Muscle-Eye-Brain Disease (POMGNT1)	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
Peroxisome Biogenesis Disorder, Type 3 (PEX12)	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 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
Pyruvate Carboxylase Deficiency (PC)	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
Short Chain Acyl-CoA Dehydrogenase Deficiency (ACADS)	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
Spastic Paraplegia, Type 15 (ZFYVE26)	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
TGM1-Related Autosomal Recessive Congenital Ichthyosis (TGM1)	No disease-causing mutations detected
Tyrosinemia Type 1 (FAH)	No disease-causing mutations detected
Tyrosinemia, Type II (TAT)	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
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
Xeroderma Pigmentosum, Group A (XPA)	No disease-causing mutations detected
Xeroderma Pigmentosum, Group C (XPC)	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:	No	**German Measles:**	No
Chicken Pox:	Yes	**German Measles Age:**	
Chicken Pox Age:	7	**Sinus Infection:**	No
Vertigo:	No	**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:	Michael Jordan-his story of perseverance & triumph.
Awards:	Individual Navy Award
Perfect Day:	A day at the beach with my family & my dogs
Personality:	I have the patience of Job, but don't let people take advantage of that.
Accomplishments:	Serving our country
Character Flaws:	I can get hyper focused on what I am doing & not pay attention to other things.
Favorite Music:	Classic Rock
Favorite Music Explanation:	It is what I grew up listening to.
Favorite Book:	Everything She Ever Wanted
Favorite Author:	Ann Rule
Favorite Author/Book Explanation:	She writes well-written true crime.
Celebrity Look Alike:	
Vegetarian:	No
Adjectives:	Patient, Kind, Artistic
Fine Art Skills:	Drawing/sketching

Technical/Mechanical Ability:	
Skills, Hobbies and Interest:	Playing Sports, Exercise

Keirsey Profile Assessment

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Birth & Education

Carried To Term:	Yes	**Birth Length:**	22 inches
Pregnancy Complications:	No	**Twin:**	No
Birth Weight:	10 lbs 9 ounces	**Twin Type:**	
Childhood Health:	Excellent		

In School:	No
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?	No
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.)?	Yes

Type Of Training:	Military police	**Training Body:**	Military

Certificate(s) Earned: Training course certificate

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.	●		Grandfather	●		61
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:		**Health Status:**	Good
Skin Tone:	Olive	**Dominant Hand:**	
Year of Birth:	1958	**Cause of Death:**	
Hair Loss:	Receding	**Height:**	5' 11" (180.34 cm)
Age At Death:		**Body Hair:**	Straight
Weight:	220 lbs. (99 kg)	**Occupation:**	Logistics & Distribution
Eye Color:	Blue	**Half Siblings:**	
Education:	High School	**Body Build:**	Medium
Half Siblings Parent:		**Eyesight Correction:**	
Near or Far Sighted:			

Comments: Has full lips and a strong brow line. He is Mr. Fix-it, mechanically inclined, and a hard worker. Enjoys golf, college football, and his grandchildren.

Family Member:	Mother
Eyesight:	
Hair Texture:	Thick
Skin Tone:	Medium
Year of Birth:	1961
Hair Loss:	
Age At Death:	
Weight:	150 lbs. (68 kg)
Eye Color:	Blue
Education:	High School
Half Siblings Parent:	
Near or Far Sighted:	

Hair Color:	Blonde, Strawberry
Freckles:	None
Health Status:	Good
Dominant Hand:	
Cause of Death:	
Height:	5' 08" (172.72 cm)
Body Hair:	Straight
Occupation:	Retired Accountant
Half Siblings:	
Body Build:	Medium
Eyesight Correction:	

Comments: Has dimples (when she smiles) and high cheek bones. She's a pastry chef who is generous and hardworking. Enjoys crafts.

Family Member:	Brother
Eyesight:	
Hair Texture:	Thick
Skin Tone:	Fair
Year of Birth:	1984
Hair Loss:	
Age At Death:	
Weight:	185 lbs. (83 kg)
Eye Color:	Blue
Education:	
Half Siblings Parent:	Maternal
Near or Far Sighted:	

Hair Color:	Red, Dark
Freckles:	Numerous
Health Status:	Fair
Dominant Hand:	
Cause of Death:	
Height:	5' 09" (175.26 cm)
Body Hair:	Straight
Occupation:	Sales
Half Siblings:	Yes
Body Build:	Medium
Eyesight Correction:	

Comments: Enjoys cooking.

Family Member:	Sister
Eyesight:	
Hair Texture:	Thick
Skin Tone:	Fair
Year of Birth:	1990
Hair Loss:	
Age At Death:	
Weight:	150 lbs. (68 kg)
Eye Color:	Blue
Education:	College (Some)
Half Siblings Parent:	Maternal
Near or Far Sighted:	

Hair Color:	Brown, Medium
Freckles:	None
Health Status:	Good
Dominant Hand:	
Cause of Death:	
Height:	5' 07" (170.18 cm)
Body Hair:	Straight
Occupation:	Construction Management
Half Siblings:	Yes
Body Build:	Medium
Eyesight Correction:	

Comments: Has dimples. She was a county Spelling Bee champion when she was younger. Enjoys music, friends, and family.

Family Member:	Brother
Eyesight:	
Hair Texture:	Thick
Skin Tone:	Medium
Year of Birth:	1989
Hair Loss:	
Age At Death:	
Weight:	200 lbs. (90 kg)
Eye Color:	Blue
Education:	High School
Half Siblings Parent:	Paternal
Near or Far Sighted:	

Hair Color:	Brown, Dark
Freckles:	None
Health Status:	Good
Dominant Hand:	
Cause of Death:	
Height:	6' 02" (187.96 cm)
Body Hair:	Straight
Occupation:	Police Officer
Half Siblings:	Yes
Body Build:	Medium
Eyesight Correction:	

Comments: He’s athletic with a strong brow line. Enjoys golf, sports, and volunteering.

Family Member:	Sister
Eyesight:	
Hair Texture:	Thick
Skin Tone:	Fair
Year of Birth:	1986
Hair Loss:	
Age At Death:	
Weight:	145 lbs. (65 kg)
Eye Color:	Blue
Education:	College
Half Siblings Parent:	Paternal
Near or Far Sighted:	

Hair Color:	Brown, Medium
Freckles:	None
Health Status:	Good
Dominant Hand:	
Cause of Death:	
Height:	5' 08" (172.72 cm)
Body Hair:	Straight
Occupation:	Teacher
Half Siblings:	Yes
Body Build:	Medium
Eyesight Correction:	

Comments: She has compassion and leadership skills. She's won multiple beauty pageants.

Family Member:	Son
Eyesight:	
Hair Texture:	
Skin Tone:	Fair
Year of Birth:	2005
Hair Loss:	
Age At Death:	
Weight:	85 lbs. (38 kg)
Eye Color:	Blue
Education:	
Half Siblings Parent:	
Near or Far Sighted:	

Hair Color:	Brown, Medium
Freckles:	None
Health Status:	Good
Dominant Hand:	
Cause of Death:	
Height:	4' 10" (147.32 cm)
Body Hair:	Straight
Occupation:	
Half Siblings:	
Body Build:	Small
Eyesight Correction:	

Comments: Has dimples. He’s polite and quiet. Enjoys basketball.

♂ Paternal Family Medical History

Family Member:	Grandfather	**Hair Color:**	Black
Eyesight:		**Freckles:**	None
Hair Texture:	Thick	**Health Status:**	
Skin Tone:	Fair	**Dominant Hand:**	
Year of Birth:	1931	**Cause of Death:**	Kidney
Hair Loss:		**Height:**	6' " (182.88 cm)
Age At Death:	61	**Body Hair:**	Wavy
Weight:	190 lbs. (86 kg)	**Occupation:**	Barber
Eye Color:	Hazel/Blue	**Half Siblings:**	
Education:	High School	**Body Build:**	Medium
Half Siblings Parent:		**Eyesight Correction:**	
Near or Far Sighted:			

Comments: Served in WWII. Enjoyed history and gem stones.

Family Member:	Grandmother
Eyesight:	
Hair Texture:	Thick
Skin Tone:	Medium
Year of Birth:	1937
Hair Loss:	
Age At Death:	
Weight:	160 lbs. (72 kg)
Eye Color:	Blue
Education:	High School
Half Siblings Parent:	
Near or Far Sighted:	

Hair Color:	Brown, Light
Freckles:	Few
Health Status:	Good
Dominant Hand:	
Cause of Death:	
Height:	5' 06" (167.64 cm)
Body Hair:	Straight
Occupation:	Board of Education
Half Siblings:	
Body Build:	Medium
Eyesight Correction:	

Comments: Has dimples and high cheekbones. She’s a pastry chef. Enjoys church/religion, gems & mining.

Family Member:	Uncle	**Hair Color:**	
Eyesight:		**Freckles:**	
Hair Texture:		**Health Status:**	Good
Skin Tone:		**Dominant Hand:**	
Year of Birth:	1960	**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:

Family Member:	Uncle	**Hair Color:**	
Eyesight:		**Freckles:**	
Hair Texture:		**Health Status:**	Good
Skin Tone:		**Dominant Hand:**	
Year of Birth:	1958	**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:

Family Member:	Aunt	**Hair Color:**	
Eyesight:		**Freckles:**	
Hair Texture:		**Health Status:**	
Skin Tone:		**Dominant Hand:**	
Year of Birth:	1955	**Cause of Death:**	Natural
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:**	Brown, Dark
Eyesight:		**Freckles:**	None
Hair Texture:	Thick	**Health Status:**	Good
Skin Tone:	Olive	**Dominant Hand:**	
Year of Birth:	1942	**Cause of Death:**	
Hair Loss:		**Height:**	5' 11" (180.34 cm)
Age At Death:		**Body Hair:**	Straight
Weight:	175 lbs. (79 kg)	**Occupation:**	Real Estate Broker
Eye Color:	Blue	**Half Siblings:**	
Education:	College (Some)	**Body Build:**	Small
Half Siblings Parent:		**Eyesight Correction:**	
Near or Far Sighted:			

Comments: Has a distinctive jaw line. Received multiple awards including real estate awards. Enjoys ballroom dancing, music, and farming.

Family Member:	Grandmother
Eyesight:	
Hair Texture:	Thick
Skin Tone:	Medium
Year of Birth:	1939
Hair Loss:	
Age At Death:	
Weight:	115 lbs. (52 kg)
Eye Color:	Hazel/Blue
Education:	High School
Half Siblings Parent:	
Near or Far Sighted:	

Hair Color:	Brown, Dark
Freckles:	None
Health Status:	Fair
Dominant Hand:	
Cause of Death:	
Height:	5' 01" (154.94 cm)
Body Hair:	Straight
Occupation:	Housekeeper
Half Siblings:	
Body Build:	Small
Eyesight Correction:	

Comments: Has high cheekbones. She kept up/keeps up with technology well into her ‘old-age’. Enjoys cooking, animals, and grandchildren.

Family Member:	Uncle	**Hair Color:**	
Eyesight:		**Freckles:**	
Hair Texture:		**Health Status:**	Good
Skin Tone:		**Dominant Hand:**	
Year of Birth:	1970	**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:

Family Member:	Aunt	**Hair Color:**	
Eyesight:		**Freckles:**	
Hair Texture:		**Health Status:**	Good
Skin Tone:		**Dominant Hand:**	
Year of Birth:	1966	**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:

Family Member:	Aunt	**Hair Color:**	
Eyesight:		**Freckles:**	
Hair Texture:		**Health Status:**	Good
Skin Tone:		**Dominant Hand:**	
Year of Birth:	1965	**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: