Cryos International - USA, LLC
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Summary of Records

05MAR2018

Donor No.	BETZY		
Type:	**Anonymous**	Weight (pounds):	**146**
Race:	**Caucasian**	Psychological profile:	**Qualified**
Ethnicity:	**Dutch,German,Irish**	Education/occupation:	**AS/Lab Tech**
Eye colour:	**Blue**	Blood group:	**0-**
Hair colour:	**Blond**	Extended profile:	**Yes**
Height (feet):	**5-6**		

Infectious Diseases			
HIV 1/2	**Negative**	HBsAg	**Negative**
Chlamydia	**Negative**	Gonorrhea	**Negative**
Syphilis	**Negative**	Anti-HBc	**Negative**
HTLV I/II	**Negative**	Anti HCV	**Negative**
HIV 1/2-NAT	**Negative**	HBV-NAT	**Negative**
HCV-NAT	**Negative**		

Hereditary Diseases			
Karyotype	**Normal(46XX)**	Tay-Sachs Disease (HEXA)	**Negative**
Cystic Fibrosis (CFTR)	**Negative**	Sickle Cell Disease (HBB)	**Negative**
Gaucher's Disease (GBA)	**Negative**	Alfa-thalassemia (HBA)	**Negative**
Beta-thalassemia (HBB)	**Negative**	Canavan's Disease (ASPA)	**Negative**
Familial dysautonomia (IKBKAP)	**Negative**	Carnitine Transporter Deficiency (SLC22A5)	**Negative**
Fragile X Syndrome (FMR1)	**Negative**	Spinal Muscular Atrophy (SMN1)	**Negative**
Fanconi Anemia Type C (FANCC)	**Negative**	Mucolipidosis Type IV (MCOLN1)	**Negative**
Niemann-Pick Type A (SMPD1)	**Negative**	Bloom Syndrome (BLM)	**Negative**
Abetalipoproteinemia (MTTP)	**Negative**	Alport Syndrome (COL4A3)	**Negative**
Alport Syndrome (COL4A4)	**Negative**	Arthrogryposis (SLC35A3)	**Negative**
Bardet-Biedl Syndrome (BBS1)	**Negative**	Bardet-Biedl Syndrome (BBS2)	**Negative**
Bardet-Biedl Syndrome (BBS10)	**Negative**	Carnitine palmitoyltransferase II deficiency (CPT2)	**Negative**
Congenital Amegakaryocytic Thrombocytopenia (MPL)	**Negative**	Congenital Disorder of Glycosylation Type 1a (PMM2)	**Negative**
Dyskeratosis Congenita (RTEL1)	**Negative**	Ehlers-Danlos Type VIIC (ADAMTS2)	**Negative**
Familial Hyperinsulinism (ABCC8)	**Negative**	Galactosemia (GALT)	**Negative**
Glycogen Storage Disease Type Ia (G6PC)	**Negative**	Joubert Syndrome 2 (TMEM216)	**Negative**
Maple Syrup Urine Disease Type 1B (BCKDHB)	**Negative**	Maple Syrup Urine Disease Type 3 (DLD)	**Negative**
Multiple Sulfatase Deficiency (SUMF1)	**Negative**	Nemaline Myopathy (NEB)	**Negative**

Phosphoglycerate Dehydrogenase Deficiency (PHGDH)	**Negative**	Polycystic Kidney Disease (PKHD1)	**Negative**
Retinitis Pigmentosa (DHDDS)	**Negative**	Smith-Lemli-Opitz Syndrome (DHCR7)	**Negative**
Tyrosemia Type 1 (FAH)	**Negative**	Usher Syndrome Type IF (PCDH15)	**Negative**
Usher Syndrome Type III (CLRN1)	**Negative**	Walker-Warburg Syndrome (FKTN)	**Negative**
Wilson Disease (ATP7B)	**Negative**	Zellweger Syndrome (PEX1)	**Negative**
Zellweger Syndrome (PEX2)	**Negative**	Zellweger Syndrome (PEX6)	**Negative**

Quarantine released

Standard	Source	Number
DK	www.cryosinternational.com/standards/dk	1-2
EU	www.cryosinternational.com/standards/eu	1-2
NYS	www.cryosinternational.com/standards/nys	1-2
USA	www.cryosinternational.com/standards/usa	1-2

Released to a Standard in effect at the time of donation.

The donor listed has been screened in accordance with 21 CFR part 1271.55 and has been determined by the establishment mentioned above to be an eligible donor.

Testing was performed by a laboratory certified to perform such testing under the Clinical Laboratory Improvement Amendments of 1988 (42 U.S.C. 263a) and 42 CFR part 493, or that has met equivalent requirements as determined by the Centers of Medicare and Medicare Services in accordance with those provisions.