



## Donor 2896

### Genetic Testing Summary

Fairfax Cryobank recommends reviewing this genetic testing summary with your healthcare provider to determine suitability.

Last Updated: 08/23/18

Donor Reported Ancestry: Norwegian, English, German, Scottish, Swedish, Dutch

Jewish Ancestry: No

| Genetic Test*   | Result   | Comments/Donor's Residual Risk**  |
|---|--|---|
| Chromosome analysis (karyotype)   | Normal male karyotype                                    | No evidence of clinically significant chromosome abnormalities  |
| Hemoglobin evaluation   | Normal hemoglobin fractionation and MCV/MCH results      | Reduced risk to be a carrier for sickle cell anemia, beta thalassemia, alpha thalassemia trait (aa/-- and a-/a-) and other hemoglobinopathies |
| Cystic Fibrosis (CF) carrier screening  | Negative by genotyping of 108 mutations in the CFTR gene | 1/270   |
| Spinal Muscular Atrophy (SMA) carrier screening   | Negative for deletions of exon 7 in the SMN1 gene        | <1/500  |
| Hb Beta Chain-Related Hemoglobinopathy (including Beta Thalassemia and Sickle Cell Disease) by genotyping | Negative for 37 mutations tested in the HBB gene         | <1/500 for Beta-Thalassemia<br><1/500 for Sickle Cell   |

\*No single test can screen for all genetic disorders. A negative screening result significantly reduces, but cannot eliminate, the risk for these conditions in a pregnancy.

\*\*Donor residual risk is the chance the donor is still a carrier after testing negative.