

Genetic Screening Questionnaire

Fertility and Reproductive Medicine Center
Washington University Physicians and Barnes-Jewish Hospital

Patient's Name _____

Date of Birth _____

Partner's Name _____

Partner's Date of Birth _____

This questionnaire is designed to identify risk factors in your personal or family history that may impact your reproductive risks. The questions will enable us to determine whether you may benefit from additional testing or genetic counseling. All answers will be kept confidential.

Please note that no questionnaire can be comprehensive so if you have specific concerns about your/your partner's personal medical history or family history, please make your physician aware.

1. Please indicate your ancestry/ethnic origin (e.g. German, African, etc.).

Self: _____

Partner: _____

2. Do you or your partner have any Eastern European (Ashkenazi) Jewish ancestry?

☐ Self ☐ Partner ☐ Neither

3. Do you or your partner have any French-Canadian or Cajun ancestry?

☐ Self ☐ Partner ☐ Neither

4. Do you or your partner have any African/African-American, Asian, Caribbean, Hispanic, Mediterranean, Mennonite, Middle Eastern, or Sephardic/Mizrahi Jewish ancestry?

☐ Self ☐ Partner ☐ Neither

5. Have you or your partner ever had genetic testing such as carrier screening or a karyotype (chromosomes)?

☐ Self ☐ Partner ☐ Neither

If yes, explain and please provide a copy of the test report(s) to our office: _____

6. Do you or your partner have a genetic condition or chromosome abnormality such as a translocation?

☐ Self ☐ Partner ☐ Neither

7. Have you or your partner ever had a stillbirth or more than two miscarriages together or with a different partner?

☐ Self ☐ Partner ☐ Neither

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8. Are you and your partner biologically related to one another?

☐ Yes, relationship: _____

☐ No

9. Do you, your partner, or any family member (children, parents, brothers, sisters, nieces, nephews, aunts, uncles, or grandparents) have any of the following? *If yes, please provide details and, if available, genetic test results.*

| Condition | Yes | No | Details (affected individual, age diagnosed, etc) |
|---|--------------------------|--------------------------|---|
| Intellectual disability/developmental delay | <input type="checkbox"/> | <input type="checkbox"/> | |
| Autism | <input type="checkbox"/> | <input type="checkbox"/> | |
| Heart defect present at birth | <input type="checkbox"/> | <input type="checkbox"/> | |
| Cleft lip or palate | <input type="checkbox"/> | <input type="checkbox"/> | |
| Neural tube defect (e.g. spina bifida, anencephaly) | <input type="checkbox"/> | <input type="checkbox"/> | |
| Limb anomaly (e.g. extra/missing fingers, abnormality of arms, legs, hands, feet) | <input type="checkbox"/> | <input type="checkbox"/> | |
| Other birth defect | <input type="checkbox"/> | <input type="checkbox"/> | |
| Hearing loss or deafness diagnosed less than age 60 | <input type="checkbox"/> | <input type="checkbox"/> | |
| Serious eye conditions or blindness | <input type="checkbox"/> | <input type="checkbox"/> | |
| Hemophilia or other bleeding/clotting disorder | <input type="checkbox"/> | <input type="checkbox"/> | |
| Alpha or beta thalassemia | <input type="checkbox"/> | <input type="checkbox"/> | |
| Sickle cell anemia or sickle cell trait | <input type="checkbox"/> | <input type="checkbox"/> | |
| Cystic fibrosis (CF) or CF carrier | <input type="checkbox"/> | <input type="checkbox"/> | |
| Spinal muscular atrophy (SMA) | <input type="checkbox"/> | <input type="checkbox"/> | |
| Tay-Sachs disease | <input type="checkbox"/> | <input type="checkbox"/> | |
| Polycystic kidney disease | <input type="checkbox"/> | <input type="checkbox"/> | |
| Neurofibromatosis | <input type="checkbox"/> | <input type="checkbox"/> | |
| Seizures/epilepsy | <input type="checkbox"/> | <input type="checkbox"/> | |
| Muscular dystrophy (e.g. Duchenne, myotonic) or other neuromuscular disease | <input type="checkbox"/> | <input type="checkbox"/> | |
| Dwarfism or skeletal dysplasia | <input type="checkbox"/> | <input type="checkbox"/> | |
| Huntington's disease | <input type="checkbox"/> | <input type="checkbox"/> | |
| Hereditary cancer syndrome (e.g. BRCA) | <input type="checkbox"/> | <input type="checkbox"/> | |
| Cancer diagnosed less than age 50 | <input type="checkbox"/> | <input type="checkbox"/> | |
| Chromosome translocation or other chromosome condition (e.g. Down syndrome) | <input type="checkbox"/> | <input type="checkbox"/> | |
| Known carrier of a genetic condition | <input type="checkbox"/> | <input type="checkbox"/> | |

10. Do you or your partner have concerns about any other conditions in either of your families not listed above?

☐ Yes, explain: _____

☐ No