

# ADVANCED OBSTETRICS & GYNECOLOGY

*“Comprehensive Healthcare for Women”*

## INFORMED CONSENT FOR GENETIC TESTING

- |   |  |
|---|--|
| <input type="checkbox"/> SMA (Spinal Muscular Atrophy)        | <input type="checkbox"/> Beta-Thalassemia      |
| <input type="checkbox"/> Cystic Fibrosis                      | <input type="checkbox"/> Alpha-Thalassemia     |
| <input type="checkbox"/> Fragile X                            | <input type="checkbox"/> Muscular Dystrophy    |
| <input type="checkbox"/> Hemoglobinopathy/Hgb Electrophoresis | <input type="checkbox"/> PKU (Phenylketonuria) |
| <input type="checkbox"/> Inherited Hemochromatosis            | <input type="checkbox"/> Sickle Cell Disease   |
| <input type="checkbox"/> Ashkenazi Jewish Carrier Panel       | <input type="checkbox"/> _____                 |
| <input type="checkbox"/> Thrombophilia Panel                  | <input type="checkbox"/> _____                 |

1. The purpose of the DNA test is to determine whether if a patient has mutation(s) known to be associated with the above genetic disease(s).
2. This testing is done on a small sample of blood.
3. If testing is done to check the genetic status of the infant, testing is done on amniotic fluid or CVS.
4. For most people, if DNA testing shows a mutation, then the person is often a carrier for the disease. Carriers are typically healthy, but are at risk of having a child with significant disease if having a child with another carrier. A discussion with a doctor may be recommended to learn the full meaning of the results and to learn if additional testing might be necessary.
5. When the DNA testing does not show a known mutation, the chance that the person is a carrier or is affected is reduced. There is still a chance to be a carrier or to be affected because the current testing cannot find all the possible changes within a gene.
6. Insurance coverage for the testing noted above is variable depending on insurance carriers and/or policy type. It is important that you understand your insurance coverage, as there may be a significant out-of-pocket expense for some or all of the tests.
7. The decision to have genetic carrier testing is optional. It is the patient's decision whether or not to have the test(s) done.

My signature below indicates that I have read, or had read to me, the above information and I understand it. I have also read or had explained to me the specific disease(s) or condition(s) tested for, and the specific test(s) I am having, including the test descriptions, principles, and limitations. I have had the opportunity to discuss the purposes and possible risks of this testing with a health care professional. I have all the information that I want and all my questions have been answered. I hereby consent to having this genetic testing performed, and I hereby authorize you to retain the results of this genetic test (and any other genetic test performed by any other physician) in my medical record.

\_\_\_\_\_  
Patient's Printed Name

\_\_\_\_\_  
Date

\_\_\_\_\_  
Patient's Signature