

Department of Obstetrics and Gynecology
Division of Ultrasound and Genetics
Prenatal/Preconceptual Counseling Questionnaire

FEMALE PARTNER

Name _____ Date of Birth _____ Age _____

How many times have you been pregnant? _____

How many living children do you have? _____

Have you ever had an elective abortion? Yes No

a. If yes, please indicate what year(s) and approximate stage of pregnancy _____

b. If yes, for what reason? Medical Personal

Have you ever had a miscarriage? Yes No If yes, how many _____

Have you ever had a stillborn baby or a baby who died within a month after birth? Yes No

If yes, please indicate year, sex of baby, and cause of death _____

How old were you at the time of your first period? _____

Are your menstrual periods:

- Regular, every 28 – 31 days Regular, less than 28 days apart
 Regular, more than 31 days apart Irregular

Do you have any major or chronic medical conditions (i.e., asthma, hypertension, diabetes, cancer, etc.)? Yes No If yes:

Condition	Date of Diagnosis	Condition	Date of Diagnosis
_____	_____	_____	_____
_____	_____	_____	_____
_____	_____	_____	_____
_____	_____	_____	_____

Please list any medications taken on a regular basis and specify approximate dosage:

Medication	Approximate Dose	Medication	Approximate Dose
_____	_____	_____	_____
_____	_____	_____	_____
_____	_____	_____	_____
_____	_____	_____	_____
_____	_____	_____	_____

What is your occupation? _____

Are you exposed to radiation or toxic substances on any basis? Yes No

If yes, please indicate the type and amount of exposure _____

Have you had any alcohol or recreational drug exposure during this pregnancy? Yes No

MALE PARTNER

Name _____ Date of Birth _____ Age _____

How many living children do you have (include children by previous unions)? _____

Have you ever fathered a child with a birth defect or genetic disease? Yes No

If yes, please describe the condition and status of the child _____

Do you have any chronic medical conditions (i.e., asthma, hypertension, diabetes, etc.)? Yes No If yes:

Condition	Date of Diagnosis	Condition	Date of Diagnosis
_____	_____	_____	_____
_____	_____	_____	_____
_____	_____	_____	_____
_____	_____	_____	_____

Please list any medications taken on a regular basis and specify approximate dosage:

Medication	Approximate Dose	Medication	Approximate Dose
_____	_____	_____	_____
_____	_____	_____	_____
_____	_____	_____	_____
_____	_____	_____	_____
_____	_____	_____	_____
_____	_____	_____	_____
_____	_____	_____	_____

What is your occupation? _____

Are you exposed to radiation or toxic substances on any basis? Yes No

If yes, please indicate the type and amount of exposure _____

CONSANGUINITY

Are the two of you related in any way other than marriage (e.g., first cousins, uncle and niece)? Yes No

If yes, please describe the relationship _____

Where did your family come from originally (e.g., England, Italy, etc.)?

Female partner _____

Male partner _____

Patient Questionnaire

Are You at Increased Risk to Carry a Genetic Disorder for which Carrier Testing is Available?

Each of us carry at least 6 disease genes, which, if inherited in a double dose, can cause a disease. In the absence of a family history, the only clue to which disorders we might be carrying is knowing our ancestry or ethnic background. In many cases, genetic testing by a blood test can be done to determine if a woman and/or her partner is a carrier for these disorders. If you would like to know if you or your partner is at increased risk to carry any of these diseases, look through the list below and check any and all boxes that match your ancestry or ethnic background (include your Parents and Grandparents, if known).

MOTHER OF BABY

- Northern European (England, Ireland, etc.)
- Asian Indian
- Ashkenazi Jewish
- African American
- Latino/Hispanic
- French Canadian
- Italian
- Greek
- Mediterranean
- Middle Eastern
- Asian
- Caribbean

FATHER OF BABY

- Northern European (England, Ireland, etc.)
- Asian Indian
- Ashkenazi Jewish
- African American
- Latino/Hispanic
- French Canadian
- Italian
- Greek
- Mediterranean
- Middle Eastern
- Asian
- Caribbean

You may also be a carrier if you or your partner have a family history of any of the following diseases:

- Cystic Fibrosis
- Tay-Sachs Disease
- Sickle Cell Disease
- Canavan Disease
- Alpha Thalassemia
- Familial Dysautonomia
- Beta Thalassemia

Have you or your partner had genetic carrier screening for CF, Sickle Cell, Thalassemia, Tay Sachs or any other genetic conditions? Yes No

If yes,

Condition Tested For

Result

Please give this form to the genetic counselor. She will review your information, and identify those genetic disorders which you are at increased risk to carry. She can also provide you with specific information about how we inherit and pass on our genetic material AND a description of the disorder(s) you are at increased risk to carry. If you and/or your partner would like to be tested, please contact your physician or healthcare provider's office.

Prenatal Genetic Screening Questionnaire

Patient's Name _____ Date of Birth _____

The following questions will enable us to determine whether certain tests may be appropriate in helping to evaluate the health of your unborn baby. Please complete this questionnaire and bring it with you at the time of your appointment. All information will be kept confidential.

This questionnaire is designed to identify potential genetic issues that have immediate relevancy to fetal health. If you have particular concerns about other conditions in your family (cancer, heart disease, diabetes, etc.), please make the genetic counselor aware of your concerns.

1. Have you OR the baby's father had any children (living or dead) with a birth defect, intellectual disability or serious health problem (include any children from you and/or your partner's previous relationships/marriages)? Yes No

If yes, please explain _____

2. Were you OR the baby's father born with any birth defects (congenital heart defect, cleft palate, etc.) or do either of you have any serious health problems? Yes No

If yes, please explain _____

3. Does/did anyone in either your or the baby's father's family have any of the following?

Please include yourself, the baby's father, your children, your parents, brothers, sisters, nieces, nephews, aunts, uncles, and grandparents.

Intellectual disability, learning disability, or autism? Yes No

Down syndrome or other chromosome abnormality? Yes No

Born with a heart defect? Yes No

Born with cleft lip or palate? Yes No

Born with a neural tube defect (open spine, spina bifida or anencephaly)? Yes No

Born with extra/missing fingers or toes or abnormality of arms, legs, hands or feet? Yes No

Hearing problems or deafness (before age 60)? Yes No

Serious eye problems or blindness? Yes No

Hemophilia or a bleeding disorder? Yes No

Neuromuscular disease or muscular dystrophy? Yes No

Huntington disease? Yes No

Three or more miscarriages and/or stillbirths? Yes No

Seizures or epilepsy? Yes No

4. Do you, the baby's father, or anyone in either of your families have a birth defect, inherited disorder, or chromosome abnormality not listed above? Yes No

If yes, indicate condition(s) and person(s) affected _____

5. Have you taken any medications, including alcohol or recreational drugs, or had any X-rays since your last menstrual period? Yes No

If yes, please list and the dates taken _____

6. Do you or the baby's father have concerns about any other conditions in either of your families? Yes No

If yes, please explain _____