

This appointment is to discuss genetic testing options available to you based on the fertility workup. Information on your personal and family history is collected to see if there are concerns, **other than the reason you were referred to us**, that should be addressed. If you have additional concerns, it may be necessary to arrange a separate appointment.

Today's date: _____

Partner 1:

Name: _____ DOB: _____
(dd/mm/yyyy)

Your doctor : _____

Current gender identity: Male Female Non-binary Other Prefer not to say

Biological Sex Assigned at Birth: Male Female Other (specify) _____
 Prefer not to say

What pronouns do you prefer? He/Him She/Her They/Them
 Other (specify) _____

Please indicate your preferred language of communication:

English French English or French (bilingual) Mandarin Other (specify) _____

Partner 2:

Name: _____ DOB: _____
(dd/mm/yyyy)

Your doctor : _____

Current gender identity: Male Female Non-binary Other Prefer not to say

Biological Sex Assigned at Birth: Male Female Other (specify) _____
 Prefer not to say

What pronouns do you prefer? He/Him She/Her They/Them
 Other (specify) _____

Please indicate your preferred language of communication:

English French English or French (bilingual) Mandarin Other (specify) _____

Reproductive History (Please complete as it pertains to you. Print additional copies as needed)

Female at Birth

Have you had any previous pregnancies?

Yes No

If yes, how many? _____

How many children do you have from:

Current partner? _____ Previous partner(s)? _____

How many miscarriages/losses:

Current partner? _____ Previous partner(s)? _____

Have you ever had a pregnancy in which malformations (physical differences/birth defects) were found in the baby? Yes No

If yes, please describe. _____

How long have you and your present partner been trying to conceive? _____

Have you had difficulty conceiving with a past partner? Yes No

If yes, for how long? _____

How old were you when you started having periods?

Are they regular? Yes No

Has anyone ever told you that you have polycystic ovarian syndrome? Yes No

Has anyone ever told you that you have pre-menopause? Yes No

If yes, what age did it start? _____

Testing Information

Have you been told that you have a low ovarian reserve for your age? Yes No

If yes, please describe _____

Have you received a diagnosis of premature ovarian failure? Yes No

Have you ever had a karyotype (chromosome analysis) done? Yes No

If yes, was an abnormality found? Yes No

If yes, please describe _____

Have you ever had a Fragile X testing done?

Yes No

If yes, were you a carrier? Yes No

Male at Birth

Do you have any children from a previous relationship? Yes No

If yes, how many? _____

How many children do you have? _____

Did you have any miscarriages/losses with previous partner(s)? Yes No

If yes, how many? _____

Have you ever had a pregnancy in which malformations (physical differences/birth defects) were found in the baby? Yes No

If yes, please describe _____

Testing Information

Have you had a spermogram done in the past?

Yes No

If yes, when was the last one done? _____

Were the results abnormal? Yes No

If yes, what was found?

- Azoospermia (no sperm)
- Oligospermia (low sperm count)
- Asthenozoospermia (Abnormal motility)
- Teratozoospermia (Abnormal shape)
- OligoAsthenoteratozoospermia (OAT)

Have you ever had a karyotype (chromosome analysis) done? Yes No

If yes, was an abnormality found?

Yes No

If yes, please describe _____

Have you ever had a Y microdeletion test done?

Yes No

If yes, was an abnormality found?

Yes No

If yes, please describe _____

Have you ever had a diagnosis of congenital bilateral absence of the vas deferens (tubes that carry the sperm out of the testes)? Yes No

If yes, did you have genetic testing for Cystic Fibrosis?

Yes No

Was the result abnormal? Yes No

If yes, please describe _____

Medical History

Do you or your partner have a known medical condition? Yes No

If yes, please describe _____

Were you or your partner born with any physical abnormalities (e.g., a hole in the heart)?

Yes No

If yes, please describe _____

Have you or your partner ever had carrier screening in the past? Yes No

If yes, were one or both of you found carrier of a condition? Yes No

If yes, please describe _____

Do you or your partner take any medication?

Please list: _____

Are you and your partner related by blood?

Yes No If yes, how?

first cousins

third cousins

second cousins

other/ I don't know

Family History

Are you or your partner adopted? Yes No

If yes, who? _____

Please indicate your ancestries (check all that apply):

African	<input type="checkbox"/> Partner 1	<input type="checkbox"/> Partner 2
Ashkenazi Jewish	<input type="checkbox"/> Partner 1	<input type="checkbox"/> Partner 2
Asian	<input type="checkbox"/> Partner 1	<input type="checkbox"/> Partner 2
Arab	<input type="checkbox"/> Partner 1	<input type="checkbox"/> Partner 2
Acadian	<input type="checkbox"/> Partner 1	<input type="checkbox"/> Partner 2
Caribbean	<input type="checkbox"/> Partner 1	<input type="checkbox"/> Partner 2
European	<input type="checkbox"/> Partner 1	<input type="checkbox"/> Partner 2
European (Eastern)	<input type="checkbox"/> Partner 1	<input type="checkbox"/> Partner 2
French Canadian	<input type="checkbox"/> Partner 1	<input type="checkbox"/> Partner 2
Hispanic	<input type="checkbox"/> Partner 1	<input type="checkbox"/> Partner 2
Indigenous Canadian	<input type="checkbox"/> Partner 1	<input type="checkbox"/> Partner 2
Pacific Islander	<input type="checkbox"/> Partner 1	<input type="checkbox"/> Partner 2
Other (Specify):	<input type="checkbox"/> Partner 1: _____	<input type="checkbox"/> Partner 2: _____

Do you or your partner have a family history of any of the following?

If yes, please indicate condition and the relationship with the affected individual.

Muscular dystrophy	<input type="checkbox"/> Yes	<input type="checkbox"/> No	
Excessive bleeding or bruising	<input type="checkbox"/> Yes	<input type="checkbox"/> No	
Neural Tube defect (open spine)	<input type="checkbox"/> Yes	<input type="checkbox"/> No	
A child who was very ill or passed away in infancy or early childhood	<input type="checkbox"/> Yes	<input type="checkbox"/> No	
Childhood cancer or cancer that occurs at a younger age than usual	<input type="checkbox"/> Yes	<input type="checkbox"/> No	
Metabolic conditions (e.g., Gaucher disease, glycogen storage disease)	<input type="checkbox"/> Yes	<input type="checkbox"/> No	
Autism	<input type="checkbox"/> Yes	<input type="checkbox"/> No	
Learning disabilities or mental handicap	<input type="checkbox"/> Yes	<input type="checkbox"/> No	
Malformation at birth (e.g., hole in the heart, cleft lip and/or cleft palate)	<input type="checkbox"/> Yes	<input type="checkbox"/> No	
Hearing or vision loss from an early age	<input type="checkbox"/> Yes	<input type="checkbox"/> No	
Chromosome differences (e.g., Down syndrome)	<input type="checkbox"/> Yes	<input type="checkbox"/> No	

Do you have any other concerns about your family or your partner's family history?
