CONSENT FORM

CONSTITUTIONAL GENETIC TESTING (PRENATAL GENETICS)

ACCOUNT NO. NRIC NO. NAME ADDRESS SEX/BIRTH DATE/RACE DATE AND TIME OF ADMISSION

What is genetic testing?

Genetic testing is the analysis of genetic information of an individual.

Why do I need genetic testing?

You have been offered genetic testing to provide a genetic diagnosis for this pregnancy. The result of the genetic testing may be helpful for the doctors to better care for you and your pregnancy.

In your case, you are offered genetic testing for the following **suspected/ clinical genetic** condition(s):______

Name of the genetic test: _____

What does genetic testing involve?

A chorionic villus / amniotic fluid sample (please delete accordingly) or ______ (sample) will be collected. The sample(s) will only be tested for the gene(s) / condition(s) listed above.

What are the possible results I could receive?

The results you could receive from genetic testing depend on the type of genetic test that was done. Some gene changes can result in a faulty gene and cause certain health problems, whereas other gene changes have no effect on health.

• Genetic variant(s) identified

- A genetic change was identified
- This confirms a genetic diagnosis and may be helpful for medical management.
- This result may have implications for family members.

• Genetic variant(s) NOT identified

- No genetic change was identified
- This result may reduce likelihood of the genetic condition(s) tested for, but does not completely eliminate the possibility of such condition(s), or other genetic condition(s) that were not analysed. Further testing may be required.
- This result could also be due to limitations in current technology and/or knowledge.
- Variant(s) of Uncertain Significance (VUS) identified
 - A genetic change was identified, however, there is insufficient information about the genetic change to associate it with an abnormal outcome, at the time of testing.
 - This result will not be used to direct medical management, unless deemed significant by your medical doctor.
 - Testing other family members may be helpful to clarify if this result is truly associated with a genetic condition / predisposition to disease.

Consent: Constitutional Genetic Testing (Prenatal Genetics) Document Number: 85060-FM-MB-215 Date Issued: October 2020 Inputs from: Neonatal and Developmental Medicine (SGH), Obstetrics and Gy

Inputs from: Neonatal and Developmental Medicine (SGH), Obstetrics and Gynaecology (SGH), Cytogenetics Laboratory (SGH, KKH), Paediatrics Genetics Service (KKH), Antenatal Diagnostics (KKH)

This variant may be reclassified over time, when more information is available. The laboratory
may issue a revised report / addendum, and you will be informed.

Incidental findings

- These are genetic changes that may not be related to the reason for testing. This may or may not have significant implications on the outcome of the pregnancy.
- You have a choice whether you wish to receive such findings, if any.

What precautions must I take for genetic testing?

If a separate blood draw is required from you and/or the biological father, please inform the doctor/genetic counsellor if either of you have had a bone marrow transplant or blood transfusion performed in the past.

What are the potential risks and limitations of genetic testing?

The genetic testing and results may come with some risks and/or limitations:

- The test result can confirm a genetic diagnosis for this pregnancy but cannot determine if, or when, the symptoms will manifest, nor can it provide information on the disease severity or recurrence.
- The test result may not only have implications for the pregnancy, but also other members of the family as it may change their understanding of their genetic risk.
- Genetic test results may result in some forms of discrimination (insurance, employment or other) as they form part of your medical records and may be accessed by and/or disclosed to a third party who has obtained your necessary consent or when such access is allowed or required by law.
- There is a small chance of error in the results due to, but not limited to, limitations in technology, sample contamination, including maternal cell contamination, inconsistencies or differences in classification of variant(s), and/or lack of clinical knowledge and inaccuracies in family history knowledge.
- Genetic testing, in rare cases, may reveal non-paternity/maternity of a presumed parent in your family.
- Occasionally, the laboratory may require additional sample(s) from you and/or family members to clarify the result. In case of insufficient sample, an additional sample would also be required.
- People react differently to receiving genetic test results. You can request for additional support before proceeding with this genetic test and/or after receiving the results.

What can I expect after the test?

- Due to the complexity of the test, your results will only be made available to you by a genetic counsellor or suitably qualified and appointed healthcare professional.
- This genetic test result will be stored in your medical records, which will be accessible by the medical team(s) responsible for your care.
- The results are confidential and will only be released to other medical professionals involved in your care and/or other parties with your written consent or as otherwise allowed by law.
- Any remaining unused portion of the sample may be stored for validation, process development, and/or quality control studies, according to the laboratory's sample retention policies.
- Further testing and/or future re-analysis requested may incur additional charges and/or require an additional sample to be taken and/or may delay the time taken to get a final result.
- To assist with result interpretation, your de-identified genetic results and clinical information may be added to scientific databases (local and/or international).

• The result(s) may be useful to your family members to receive genetic testing. Please inform your doctor/genetic counsellor if you consent to sharing your genetic testing results with your family members.

What are my options?

Genetic testing is voluntary; you may choose not to proceed. You may also withdraw from the genetic testing at any point, before the test is completed. If consent is withdrawn, the sample will be discarded and no report will be issued. However, charges would apply once the test request has been received and processed.

Others (to be filled by Healthcare Professional) [if applicable]

Pa	rt I – Patient's Declaration			
1.	I, (NRIC/Passport No),			
		nat I understand the nature, purpose, risks, limitations,		
2.	,	sted are not intended to be exhaustive. I have had an (i) the above-mentioned risks and limitations; (ii) the relevance to me.		
3.	I hereby consent to undergo the Test.			
4.	I understand and agree that the Test will be performed by the appropriate SingHealth institution (with the involvement of external providers, if necessary) and I will be admitted and/or registered as a patient of that SingHealth institution.			
5.	I understand that I have the option to choose whether to receive incidental findings.			
	\Box I wish to receive incidental findings.			
	\Box I <u>do not</u> wish to receive incidental findings.			
	□ Not applicable			
6.	I understand that my result(s) may be useful for my family members for genetic counseling and testing.			
	\Box I consent to sharing my result(s) with my family members. They will be required to provide my name and NRIC/FIN/Passport number.			
\Box I <u>do not</u> consent to sharing my result(s) with my family members.				
7.	In the event I am uncontactable, the test results may be made known to:			
		NRIC (last 4 digits):		
		Relationship:		
		NRIC (last 4 digits):		
		Relationship:		
	(Signature/[*Left/Right] Thumbprint of Patient)	(Date of Signing)		
	(Name of Witness)	(Designation of Witness)		
	(Signature of Witness)	(Date of Signing)		

* Please delete accordingly

Consent: Constitutional Genetic Testing (Prenatal Genetics) Document Number: 85060-FM-MB-215 Date Issued: October 2020 Inputs from: Neonatal and Developmental Medicine (SGH), Obstetrics and Gynaecology (SGH), Cytogenetics Laboratory (SGH, KKH), Paediatrics Genetics Service (KKH), Antenatal Diagnostics (KKH)

Part II – Parent's / Legal Guardian's / Donee's / Deputy's Declaration (herein referred to as the "Authorised Person") (if applicable)

*Parent / Legal Guardian / Donee / Depart RIC/Passport No	 s) of the Test listed are not intended to be exhaustive. I information about (i) the above-mentioned risks and pecific concern(s) of relevance to the Patient. the Test. performed by the appropriate SingHealth institution (with essary) and the Patient will be admitted and/or registered a whether to receive incidental findings. s. y be useful for the Patient's family members for genetic with Patient's family members. They will be required to 		
nfirm that I understand the nature, pur matitutional Genetic Testing (General C cknowledge that the risks and limitation(s we had an opportunity to ask for more itations; (ii) the risks in general; and (iii) sp ereby consent for the Patient to undergo to inderstand and agree that the Test will be a involvement of external providers if necession a patient of that SingHealth institution. I wish to receive incidental findings. I do not wish to receive incidental finding Not applicable inderstand that the Patient's result(s) may unseling and testing. I consent to sharing Patient's result(s) we provide Patient's name and NRIC/FIN/Passp	 urpose, risks, limitations and options with regard to <u>Senetics</u>) ("Test"). s) of the Test listed are not intended to be exhaustive. I information about (i) the above-mentioned risks and pecific concern(s) of relevance to the Patient. the Test. performed by the appropriate SingHealth institution (with essary) and the Patient will be admitted and/or registered is whether to receive incidental findings. s. y be useful for the Patient's family members for genetic with Patient's family members. They will be required to the Patient to receive to the Patient to receive to the Patient to receive the patient will be required to the patient's family members. 		
enstitutional Genetic Testing (General C cknowledge that the risks and limitation(s we had an opportunity to ask for more litations; (ii) the risks in general; and (iii) sp ereby consent for the Patient to undergo to inderstand and agree that the Test will be a involvement of external providers if neces a patient of that SingHealth institution. Inderstand that I have the option to choose I wish to receive incidental findings. I <u>do not</u> wish to receive incidental finding Not applicable Inderstand that the Patient's result(s) may unseling and testing. I consent to sharing Patient's result(s) we provide Patient's name and NRIC/FIN/Passp	Genetics) ("Test"). ("Test"). (a) of the Test listed are not intended to be exhaustive. I information about (i) the above-mentioned risks and pecific concern(s) of relevance to the Patient. (b) the Test. (c) performed by the appropriate SingHealth institution (with essary) and the Patient will be admitted and/or registered e whether to receive incidental findings. (c) s. (c) y be useful for the Patient's family members for genetic with Patient's family members. They will be required to the Patient to receive to the Patient.		
cknowledge that the risks and limitation(sive had an opportunity to ask for more litations; (ii) the risks in general; and (iii) speceby consent for the Patient to undergot to inderstand and agree that the Test will be a involvement of external providers if nece a patient of that SingHealth institution. Inderstand that I have the option to choose I wish to receive incidental findings. I <u>do not</u> wish to receive incidental finding Not applicable Inderstand that the Patient's result(s) may unseling and testing. I consent to sharing Patient's result(s) we provide Patient's name and NRIC/FIN/Passp	 s) of the Test listed are not intended to be exhaustive. I information about (i) the above-mentioned risks and pecific concern(s) of relevance to the Patient. the Test. performed by the appropriate SingHealth institution (with essary) and the Patient will be admitted and/or registered a whether to receive incidental findings. s. y be useful for the Patient's family members for genetic with Patient's family members. They will be required to 		
ve had an opportunity to ask for more itations; (ii) the risks in general; and (iii) speceed ereby consent for the Patient to undergo to inderstand and agree that the Test will be a involvement of external providers if neceed a patient of that SingHealth institution. Inderstand that I have the option to choose I wish to receive incidental findings. I <u>do not</u> wish to receive incidental finding Not applicable Inderstand that the Patient's result(s) may unseling and testing. I consent to sharing Patient's result(s) we provide Patient's name and NRIC/FIN/Passp	information about (i) the above-mentioned risks and pecific concern(s) of relevance to the Patient. the Test. performed by the appropriate SingHealth institution (with essary) and the Patient will be admitted and/or registered e whether to receive incidental findings. s. y be useful for the Patient's family members for genetic with Patient's family members. They will be required to		
nderstand and agree that the Test will be e involvement of external providers if nece a patient of that SingHealth institution. Inderstand that I have the option to choose I wish to receive incidental findings. I <u>do not</u> wish to receive incidental finding Not applicable Inderstand that the Patient's result(s) may unseling and testing. I consent to sharing Patient's result(s) we provide Patient's name and NRIC/FIN/Passp	performed by the appropriate SingHealth institution (with essary) and the Patient will be admitted and/or registered e whether to receive incidental findings. s. y be useful for the Patient's family members for genetic with Patient's family members. They will be required to		
e involvement of external providers if nece a patient of that SingHealth institution. Inderstand that I have the option to choose I wish to receive incidental findings. I <u>do not</u> wish to receive incidental finding Not applicable Inderstand that the Patient's result(s) may unseling and testing. I consent to sharing Patient's result(s) we ovide Patient's name and NRIC/FIN/Passp	essary) and the Patient will be admitted and/or registered e whether to receive incidental findings. s. y be useful for the Patient's family members for genetic with Patient's family members. They will be required to		
I wish to receive incidental findings. I <u>do not</u> wish to receive incidental finding Not applicable nderstand that the Patient's result(s) may unseling and testing. I consent to sharing Patient's result(s) w ovide Patient's name and NRIC/FIN/Passp	s. y be useful for the Patient's family members for genetic with Patient's family members. They will be required to		
I <u>do not</u> wish to receive incidental finding Not applicable nderstand that the Patient's result(s) may unseling and testing. I consent to sharing Patient's result(s) w ovide Patient's name and NRIC/FIN/Passp	y be useful for the Patient's family members for genetic with Patient's family members. They will be required to		
Not applicable nderstand that the Patient's result(s) may unseling and testing. I consent to sharing Patient's result(s) v ovide Patient's name and NRIC/FIN/Passp	y be useful for the Patient's family members for genetic with Patient's family members. They will be required to		
nderstand that the Patient's result(s) may unseling and testing. I consent to sharing Patient's result(s) v ovide Patient's name and NRIC/FIN/Passp	with Patient's family members. They will be required to		
unseling and testing. I consent to sharing Patient's result(s) v ovide Patient's name and NRIC/FIN/Passp	with Patient's family members. They will be required to		
ovide Patient's name and NRIC/FIN/Passp			
I <u>do not</u> consent to sharing Patient's resu			
	□ I <u>do not</u> consent to sharing Patient's result(s) with Patient's family members.		
In the event I am uncontactable, the test results may be made known to:			
	NRIC (last 4 digits):		
ntact details:	Relationship:		
me:	NRIC (last 4 digits):		
ntact details:	Relationship:		
gnature/ [*Left/Right] Thumbprint of thorised Person)	(Date of Signing)		
ame of Witness)	(Designation of Witness)		
	thorised Person)		

Consent: Constitutional Genetic Testing (Prenatal Genetics) Document Number: 85060-FM-MB-215 Date Issued: October 2020 Inputs from: Neonatal and Developmental Medicine (SGH). Obste

Inputs from: Neonatal and Developmental Medicine (SGH), Obstetrics and Gynaecology (SGH), Cytogenetics Laboratory (SGH, KKH), Paediatrics Genetics Service (KKH), Antenatal Diagnostics (KKH)

Part III – Healthcare Professional's De	claration
---	-----------

I confirm that I have explained to the Patient, or the Authorised Person (if applicable), the Patient's medical condition as well as the nature, purpose, risks, limitations, and alternatives with regard to the Test and have addressed queries of the Patient, or the Authorised Person (if applicable).

(Signature, Full Name, and Professional Registration / *Employee No. of Healthcare Professional) *Only for those without professional registration number (Date of Signing)

Part IV – Interpreter's Declaration (if applicable)

I, ______, confirm that I have interpreted to the Patient, or the Authorised Person (if applicable), the Healthcare Professional's explanation of the Patient's medical condition, nature, purpose, risks, limitations, and alternatives with regard to the Test and the Healthcare Professional's response to the Patient's, or the Authorised Person's (if applicable), queries in ______ (language / dialect).

(Signature of Interpreter)

(Date of Signing)