

Stick patient's label here

## **Informed Consent Form for**

## Prenatal and Postnatal Chromosomal Karyotyping/FISH

The following points have been explained to me:

- (i) The purpose of performing this type of chromosomal karyotyping/fluorescence in situ hybridization (FISH) is to determine if a person has been born with a chromosomal problem
- (ii) Common symptoms that may suggest the presence of a chromosomal problem include
  - Abnormal ultrasound findings, abnormal serum screening, NIPS, advanced maternal age, prior pregnancy with chromosomal abnormality and/or parental concern
  - Dysmorphic features
  - Multiple congenital anomalies
  - Intellectual disability
  - · Failure to thrive; short stature
  - Features of a known chromosomal disorder
  - Parent or other family member carrying a known chromosomal abnormality
  - Recurrent pregnancy losses; infertility;
  - Other reason\_\_\_\_\_(please state)
- (iii) Chromosome karyotyping/FISH analyses can pick up conditions involving abnormal chromosomal numbers, large rearrangements, large deletions and duplications.
- (iv) However, it has a limited resolution, and cannot detect other genetic changes such as small deletions, duplications, and point mutations
- (v) Chromosomal karyotyping may have unexpected test results that are not directly related to the clinical reason for ordering the test.
- (vi) Chromosomal karyotyping requires mitotic (dividing) cells from a live specimen; the test may not be successful if the cells are not healthy.
- (vii) Fluorescence in situ hybridization (FISH) does not require mitotic cells and can detect presence/absence of specific genes/gene regions, but it cannot detect point mutations.
- (viii) The specimen required may be amniotic fluid, chorionic villi, abortus tissue, a blood sample, a skin sample, or any other living specimen.
- (ix) There may be other relevant alternative tests to chromosomal karyotyping/FISH. This should have been explained to me by my doctor.
- (x) The test results are strictly confidential, and the test results will only be released to the doctor who ordered the test and not directly to the patient/parent. The patient/parent will then have to meet this doctor/ designate/genetic counsellor to get the test results.
- (xi) After the test results have been released to the doctor who ordered the test, the test results will be part of the patient's medical records and will be protected as required under the Personal Data Protection Act of Singapore and other relevant legislations such as the Private Hospitals and Medical Clinics Act.
- (xii) The chromosomal karyotyping/FISH results may affect health insurance coverage or claims.
- (xiii) The access, collection, use, transfer and storage of the above-named patient's personal data (and, where applicable, the disclosure of the above-named patient's personal data to authorized and relevant third parties) will be carried out by NUH for the purpose of the chromosomal karyotyping/FISH on the above-named patient;

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## Restricted, Sensitive (Normal)



I confirm that I have had the opportunity to clarify any queries I have in relation to the chromosomal karyotyping/FISH (including any queries in relation to the contents of this form) and these have been clarified to my satisfaction;

I confirm that the nature, effect, purpose, limitations		
karyotyping/FISH have been fully explained to me b	by Dr	and I fully
understand the explanation; I consent to chromosomal karyotyping/FISH to b	be done on	
Name of Patient:		
NRIC / Birth Certificate / Passport No.*:		
I consent/do not consent* to a small amount of my		
personal information and retained for the use of med		
karyotyping/FISH testing is completed. I understand	·	•
any tests conducted on the DNA sample will not be	reported to me/my child. I understand that	if I do not consent to this,
the quality of my test results will not be affected.		
Patient/Parent/Legal Guardian's* signature	Date	
Name of Parent/Legal Guardian *:		
NRIC of Parent/Legal Guardian *:		
Relationship to Patient:		
I have explained the nature, effect, purpose, limitation	ons, potential benefits and risks of, and alte	ernatives to,
chromosomal karyotyping/FISH to the patient/paren	nt and/or legal guardian*, and answered all	questions satisfactorily.
Doctor's signature	 Date	
Name of Doctor obtaining consent:		
MCR number of Doctor obtaining consent:		
If consent was obtained via an interpreter:		
Language used to obtain consent:		
	Interpreter's signature	Date
Name of Interpreter obtaining consent:		
NRIC of Interpreter obtaining consent:		
*Delete accordingly		

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