

PATIENT DETAILS

Surname: _____

Forename: _____

Date of Birth: ___/___/___

Sex: Male Female

Hospital/Clinic No.: _____

Laboratory No.: _____

Ward: _____

Physician: _____

REQUESTING HOSPITAL/ CLINIC DETAILS

Hospital / Clinic Name: _____

Department: _____

Address: _____

Phone: _____ Fax: _____

PLEASE REMEMBER ALWAYS TO COMPLETE THE INFORMED CONSENT SECTION

SAMPLE DETAILS

Specimen Collection Date: ___/___/___/___ Specimen Type (Amniotic Fluid): _____

Other Sample Type: _____

TEST INFORMATION

PRE-NATAL GENETICS

TEST NAME	CODE	SELECT
QF PCR & Karyotyping (Amniotic Fluid)	ANEUP & KARPN	<input type="checkbox"/>
Chromosome Analysis – Placenta/ CVS	KARPL	<input type="checkbox"/>

OTHER TESTS REQUIRED:

INDICATIONS / SUSPECTED CONDITIONS / PREGNANCY INFORMATION

PLEASE DESCRIBE INDICATIONS/ SUSPECTED CONDITION: _____

PREGNANCY INFORMATION:

- LMP: ___/___/___ EDD: ___/___/___
Day / Month / Year Day / Month / Year

- Scan Date: ___/___/___ i.e. Gestational Age ___/___ of amenorrhea on scan day
Day / Month / Year Weeks / Days

DOWN SYNDROME RISK EVALUATION RESULTS (IF AVAILABLE): _____

FAMILY HISTORY OF CHROMOSOME ABNORMALITIES (DETAIL): _____

PATIENT HISTORY**MEDICAL HISTORY:** _____

_____**SURGICAL HISTORY:** _____

_____**OBSTETRIC HISTORY:** _____

_____**INFORMED CONSENT SECTION****• Patient or Guardian:**

I/we the undersigned confirm that I/we have been fully informed by the Doctor/Pathologist/ Geneticist _____ regarding cytogenetic and/or molecular genetic tests that will be performed on cells and/or DNA extracted from my/our child's blood and/or tissue to:

- confirm or exclude the diagnosis of or a predisposition to a genetic disease.
- determine heterozygote status with a view to obtaining genetic counselling.
- examine gene locus/loci.

I/we give my/our consent to such testing and confirm that I/we have received all the necessary information according to the law.

Patient/Guardian Signature: _____ **Date:** ____/____/____**• Doctor/ Pathologist/Genetic Consultant:**

The Cytogenetic and/or molecular genetic test information is to be given by the Clinical Pathologist prescribing the test, or by the Physician collecting the sample. All relevant issues regarding the involved pathology etiology, development, prognosis and potential treatment must have been raised by the Genetic consultant or the Physician and clearly understood by the patient. All information associated with the patient file will be retained by Eurofins Biomnis. The result must be reported to the Physician only.

Doctor/Pathologist Signature: _____ **Date:** ____/____/____