TEST REQUEST FORM: PRENATALSEQ (NON-INVASIVE PRENATAL SCREENING)



PATIENT DETAILS	HEALTH PRACTITIONER DETAILS
Forename*	Account ID*
Surname*	Full Name*
Hospital ID	Phone*
DOB (DD/MM/YYY) *	Email*
	Institution*
CLINICAL INFORMATION	Address 1*
Gestational age at the date of sample draw*1	Address 2
Weeks: Days:	City/town*
Maternal BMI*	County/ State*
Number of Foetuses*2	Post Code*
□1	Country*
☐2 (☐ monozygotic ☐ dizygotic ☐ unknown)	SAMPLE DETAILS
IVF Pregnancy	Sample Type
□Yes (Egg donor is: □ Self □ Non-self) □No	☐ 7-10ml maternal peripheral whole blood collected in a STRECK Cell-Free DNA Blood Collection Tube
¹ Patient must be at least 10 weeks 0 days in gestational age (as	Date Collected (DD/MM/YYY)*
determined by a scan). 2This test is not available for pregnancies with more than two foetuses.	Time Collected (hh:mm)
TEST DETAILS	Is this a repeat sample? □Yes □No
☐ Trisomy 13, 18, and 21	BILLING AND REPORT DETAILS
Please mark any additional test options requested	The invoice for this test will be sent to the default billing address associated with your Account ID. If your
☐ Foetal Sex³	institution requires a PO number, please insert it here. PO Number
□ Sex Chromosome Aneuploidies ⁴ ³ Foetal sex considers the presence or absence of a Y chromosome, and it will be reported as "male" or "female". For twin pregnancies, "male" means that a Y chromosome was detected in at least one of the foetuses, and "female" means that a Y chromosome was not detected in either foetus.	The report for this test will be made available in your account on Genseq's online ordering portal. If you need other health practitioners to have access to the report, please ensure they are registered under your Account to ensure appropriate communication.
⁴ Analysis of sex chromosome aneuploidies is an option available for singleton pregnancies. If analysis of sex chromosomes aneuploidies is performed, and an aneuploidy is detected, the sex of the foetus will also be determined, even if it was not requested Please refer to our Laboratory User Guide for information on the sample requirements for this test, and to confirm eligibility for your patient. This test screens for	*I hereby confirm that I have obtained written informed consent from the patient for this test to be performed, including consent for health practitioners registered under my account to access the report.
Trisomy 13, 18, and 21, and optionally sex chromosome aneuploidies (X, XXX, XXY, XYY) and foetal sex. Aneuploidies involving other chromosomes, polyploidies (such as triploidy), and partial deletions or duplications are not evaluated. If an abnormality is detected, the mosaicism level will not be reported. This test is not clinically indicated for patients that have multiple gestation pregnancies with three or more foetus. In addition, the accuracy of screening results can be adversely affected by certain maternal and foetal factors, including but not limited to: recent maternal blood transfusion; maternal prior bone marrow / organ transplant / stem cell transplant; radiation/ immune/ stem cell therapy; maternal autoimmune disease or cancer unless in remission; maternal	*Please, indicate how long you would like Genseq to store DNA sequencing raw data on your behalf: General Bounds In addition, in order to (a) fulfil your instructions to the requested perform genetic testing, and (b) for us to provide further cascade genetic tests for you, each as undertaken in the context of an accredited genetic testing service, you understand that other data types, such as patient data received on this Test Request Form, laboratory QC data and report data will be stored for a further period of years taking account of applicable law, regulation and industry guidance. In returning this Test Request Form for processing you are

INTERNAL USE ONLY

Sample ID

Genseq, Building 4, Cherrywood Business Park, Dublin, D18 K7W4, Ireland. Document Reference: LF-GEN-0136 (Version 1)

neoplasms (benign and malignant); maternal mosaicism; maternal copy number variations, balanced translocations or whole

chromosomal abnormalities; foetoplacental mosaicism / confined

placental mosaicism and foetal demise / vanishing twin.