

# **GPD**<sub>x</sub> Genomic Precision Diagnostic

Comprehensive and Personalized Genetic Testing for Obstetrics, Gynaecology and Feto-Maternal Units



### What are the benefits of Integrating Genomics into Medical Decisions?

- For most of the clinical conditions genetic testing is the only way to make an accurate diagnosis and help avoid additional unnecessary clinical investigations.
- Reduces the uncertainty in a differential diagnosis and provides a directed focus for the specialist to provide appropriate treatment.
- Decreases referrals from specialist to specialist, allowing a high quality patient care.
- Can guide the specialist in choosing the most suitable downstream tests, therapy and support for the patient.

### Diagnostic testing Indications



### **Preconception Prevention**

- Couples and families who want to know if they carry a genetic condition and determine if they are at risk of having a child with a genetic disease.



### **High Risk Pregnancies**

Family history of a genetic condition (hereditary disorder, chromosomal alteration...).
High risk established using screening methods: first and second trimester, cffDNA (NIPT).
Ultrasound findings of a genetic condition: (genetic disorder, chromosomal alteration,...).
Pregnancy with poor evolution or miscarriage.



### Neonatal Screening/Diagnostic

- Identify genetic conditions that could cause disease in newborn babies.

- Prevent complications and increase life expectancy.

## **Diagnostic Technologies**

Get supported by our experienced and high skilled team to choose the best Genetic testing

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#### WHOLE EXOME/GENOME SEQUENCING (WES/WGS)

Complete DNA test to identify mutations in over 24,000 genes related to complex genetic problems.



#### SINGLE GENE ANALYSIS

Many genetic diseases are caused by changes or variants in a single gene. We have a comprehensive range of tests to cover your needs.



#### IGX PRECISION PANELS

Choose your panel and benefit from the highest diagnostic accuracy.

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#### CMA: CHROMOSOMAL MICROARRAY TESTING

Gold standard for the detection of Copy number variations (CNVs).



Our experienced Geneticists guide you throughout the whole diagnostic process



Dr. Garcia-Planells PhD Human Genetics



**Dr. Julio Martín** PhD Molecular Genetics

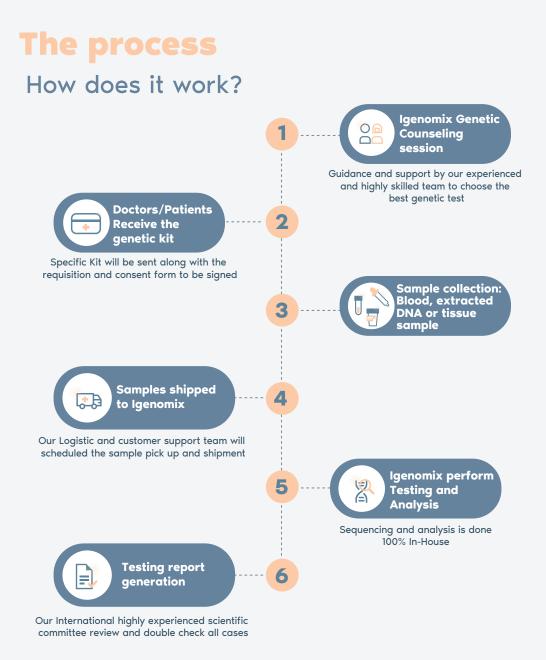


Dr. Lova Satyanarayana PhD Human Genetics



## DIAGNOSTIC SERVICES for all life stages





	Sample type	Container	Transportation Temperature	Volume
	Peripheral blood	EDTA vacutainer	20-25°C	3 – 4 ml
P	Purified genomic DNA	In a sealed Eppendorf tube	20-25°C	A minimum 1 microgram of DNA at a concentration of 50-100ng/µl
2	POC (fetal tissue)	Tissue in sterile container in saline and cardiac or cord blood in vacutainer	20-25°C	3 – 4 mm POC specimen or 50 100 mg of each tissue
J.	Amniotic Fluid	Sterile container	20-25°C	10-15ml
ð	Chorionic villi	Sterile container with culture mediumor saline solution with 1% antibiotic	2-8 °C	300-500mg



Start a New success story with our

### **Genetic Counseling Service**

Certified Genetic Counselors to guide you and your patients in choosing the right genetic testing.



Contact our Genetic Counselors and Geneticists to get guidance and support

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