

# MaatriSeq

**Precision Prenatal Care** 







#### MaatriSeq: Precision Prenatal Care

A cell-free DNA (cfDNA) based screening test designed to provide expectant mothers with early insights into their baby's health.



· Analyses fragments of the baby's DNA circulating in mother's blood



• Performed as early as 10 weeks of pregnancy



• Non-invasive, with a simple blood draw, offers little risk to the mother or fetus



• Comprehensive insights within 10 days



# NGS based NIPS powered by illumina's NovaSeq<sup>™</sup> X Plus

## MaatriSeq screens for



Down Syndrome (Trisomy 21)



Edwards Syndrome (Trisomy 18)



Patau Syndrome (Trisomy 13)



Gonosomal Aneuploidies (X&Y)



All Chromosomal Aneuploidies (22 pairs)



**Turnaround Time** 12 Days



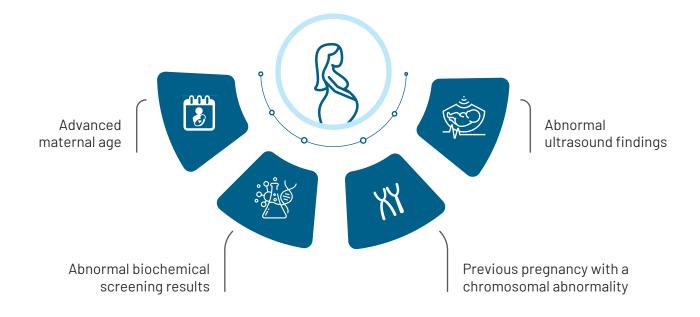
Sample Requirement 10ml Maternal Blood

Disclaimer: This test is exclusively available upon prescription by a certified clinician/clinical geneticist duly registered with PCPNDT guidelines.



### Who should be offered MaatriSeq?

NIPS is offered to all pregnant women, and recommended in cases of:



#### **Proprietary Platforms**



#### **Additional Testing Options**

- Exome Plus: Exome Sequencing
- CNSeq: Low-pass Whole Genome Sequencing
- AneuXpress: Quantitative Fluorescent PCR
- Sanger Based Testing

- Couple Carrier Screening
- Whole Genome Sequencing
- Preimplantation Genetic Testing:
  PGT Aneuploidy



+000,08 Genetic Tests Reported

500+ Projects Executed for Genomics Majors Globally Presence in 20+ Countries **SCAN TO KNOW MORE** 







Reference Sdn Bhd (1278003-P) Wisma MEDINICS, No.2, Jalan Astaka U8/88B, Seksyen U8, Bukit Jelutong, 40150 Shah Alam, Selangor, Malaysia.

