There are Millions of Worries During Pregnancy Have one less with

UNICAD





UNICADNIPT

Unipath Non Invasive Chromosomal Aneuploidy Detection

How does the UNICAD NIPT prenatal screening test work?

During pregnancy, the placenta leaks fetal cell-free DNA which circulates in the maternal bloodstream. As a result, a maternal blood sample contains a mixture of fetal and maternal circulating DNA. The UNICAD NIPT prenatal screening directly measures the amount of the cell-free DNA and can detect small changes in the DNA ratio between the maternal and cell-free DNA to estimate the risk of a fetal chromosomal aneuploidy or a microdeletion being present. The UNICAD NIPT prenatal screening employs the latest advances in Next Generation Sequencing (NGS) to analyze the DNA.

UNICAD NIPT USP

Sensitive

Able to produce valid, Fast & Accurate results in samples with little as 3.5% Fetal Fraction

Quality

Powered by Thermo fisher & Yourgene Health- leading International Service provide of NIPT

Flexible

Customized analysis including all aneuploidies and selected microdeletions

Safe

Non-invasive, with no risk of miscarriage

Accurate

>99% detection of autosomal trisomy conditions & a Low re-draw rate of <1%

Simple

Uses Just one Maternal Blood sample 10ml

Recommendations:

*American College of Obstetrics & Gynecologist (ACOG)

- ACOG recommends that informing all Pregnant Women that NIPS is the most sensitive screening option for traditionally screened Aneuploidies (i.e, patau, Edwards & Down syndromes)
- NIPT can be performed at an earlier gestational age than conventional screening and there is no gestational age upper limit after 10 weeks of gestation. This means that Patients can get the most accurate screening information at an earlier gestational age, this enhances informed decision making

In accordance with PCPNDT ACT We do not reveal Gender



The UNICAD NIPT prenatal screening for?

- >> Autosomal aneuploidies of 23 pairs of chromosomes including Trisomy 21, 18, 13
- >>> Sex chromosome aneuploidies
- Microdeletions

UNICAD NIPT

Autosomal Aneuploidies

- >>> Down's syndrome (Trisomy 21)
- >>> Edwards' syndrome (Trisomy 18)
- Patau's syndrome (Trisomy 13)
- Whole genome-wide an euploidy detection of the whole fetal genome (all 23 pairs of chromosomes)

Sex Chromose Aneuploidies

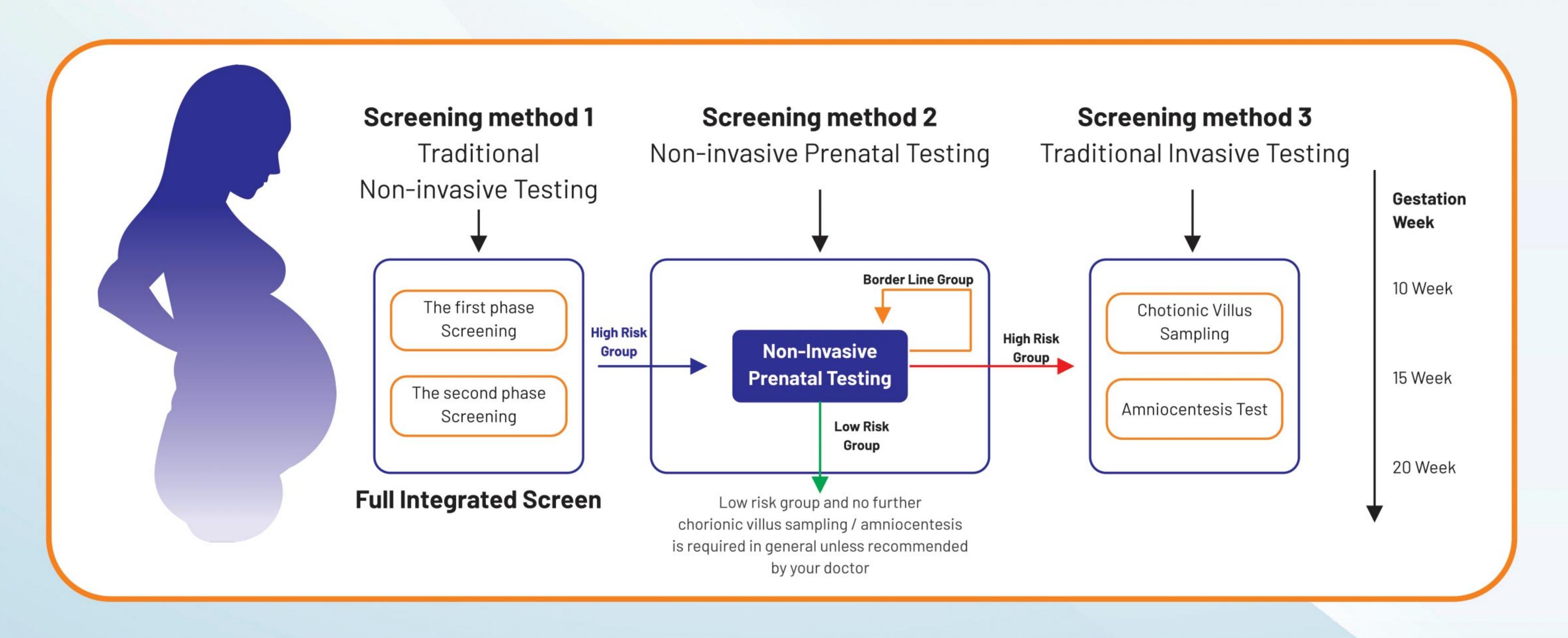
- Monosomy X Turner syndrome
- XXX Triple X syndrome
- >>> XXY Kleinfelter syndrome
- >>> XYY Jacob's syndrome

UNICAD NIPT ADVANCE

Microdeletions

- DiGeorge syndrome
- Angelman syndrome
- >> 1p36 deletion syndrome
- Cri-du-Chat syndrome
- Prader-Willi syndrome
- >> Wolf-Hirschhorn syndrome

UNICAD NIPT SCREENING PATHWAY





Care For Little Ones

NIPT Assay Performance of 10,640 Samples

| | Sensitivity | Specificity | PPV* | NPV* | FPR* | FNR* |
|------------------|-------------|-------------|---------|---------|--------|--------|
| Trisomy 21 (173) | >99.99% | 99.97% | 87.50% | >99.99% | 0.028% | <0.01% |
| Trisomy 18 (48) | >99.99% | 99.96% | 91.67% | >99.99% | 0.038% | <0.01% |
| Trisomy 13 (24) | >99.99% | >99.99% | >99.99% | >99.99% | <0.01% | <0.01% |
| SCAs (107) | >99.99% | 99.91% | 91.59% | >99.99% | 0.085% | <0.01% |

SCAs: Sex Chromosomal Aneuploidies

PPV: Positive Predictive Values* FNP: False Nagative Rate*

FPR : False Positive Rate*

NPV : Negative Predictive Values*

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