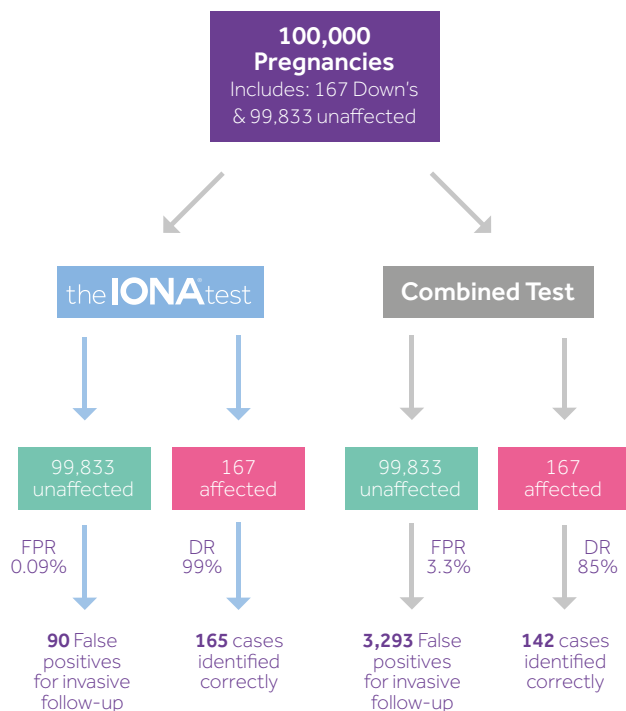


## The IONA® test reduces the need for invasive procedures:

An example scenario showing the difference between the screening tests:



### Where can I get an IONA® test?

Contact the Maternity Screening Department at the Royal United Hospital, Bath.

**Email:** [ruh-tr.maternityscreening@nhs.net](mailto:ruh-tr.maternityscreening@nhs.net)  
**or call:** 01225 825414.

For the latest news and updates about the IONA® test please follow us on:

[f /theIONAtest](https://www.facebook.com/theIONAtest) [t @theIONAtest](https://twitter.com/theIONAtest)



### About Premaitha Health

The IONA® test is developed and manufactured by Premaitha Health, a UK molecular diagnostics company based in Manchester. Premaitha's mission is to develop molecular diagnostic products that will have a positive impact on human health.

[www.premaitha.com](http://www.premaitha.com)

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FOR PREGNANT WOMEN

the IONA® test  
non-invasive prenatal screen: safe, fast, accurate



Non-invasive prenatal screening test for Down's syndrome and other serious genetic conditions

## What is the IONA® test?

The IONA® test is a non-invasive prenatal test (NIPT) for pregnant women which estimates the risk of a fetus having Down's syndrome or some other serious genetic diseases. The IONA® test is an advanced screening test that is carried out on a small maternal blood sample. Pregnant women can expect test results from their healthcare provider within approximately 3-5 days from sample receipt.

## What does IONA® screen for?

The IONA® test estimates the risk of a fetus having Down's syndrome (Trisomy 21), Edwards' syndrome (Trisomy 18) and Patau's syndrome (Trisomy 13). Trisomies occur when three, instead of the usual two, copies of a chromosome are present. Edwards' and Patau's syndromes are much rarer than Down's but are very serious and many affected babies do not survive.

## What are the advantages of the IONA® test?

**Safe:** non-invasive with no risk of miscarriage.

**Fast:** the IONA® test is the fastest NIPT available with results provided within 3-5 working days, from sample receipt.

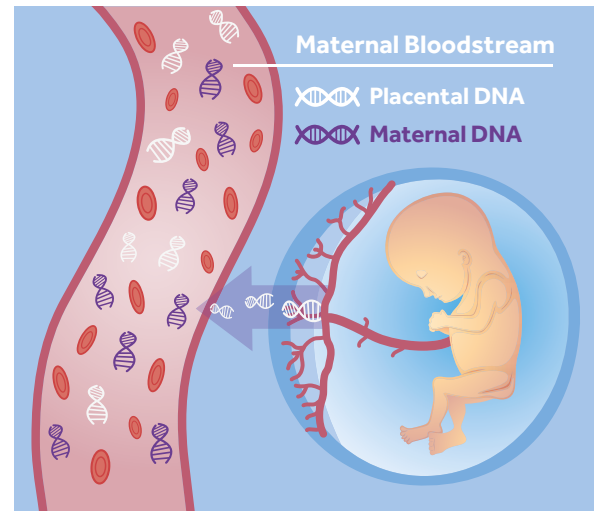
**Accurate:** greater than 99% for detection of trisomy conditions.

**Simple:** uses a simple maternal blood sample. Unlike other NIPT's, the IONA® test is performed in a laboratory local to you. So your blood sample is not shipped to the US or China.

**Quality:** unlike other NIPT's, the IONA® test is a regulated diagnostic, which is CE marked.

## How does it work?

During pregnancy the placenta leaks 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 IONA® test directly measures the amount of this cell-free DNA and can detect small changes in the DNA ratio between the maternal and cell-free DNA when a fetal trisomy 21, 18 or 13 is present.



## Why is IONA® better than the current combined test?

Traditional screening offered during pregnancy is currently called the combined test. This is an ultrasound scan to measure the nuchal translucency (NT) and a blood test. This is much less accurate than NIPT and it only detects around 85% of babies with Down's syndrome.

The IONA® test has a higher detection rate than the current combined test offered to pregnant women. This means that fewer pregnant women will undergo unnecessary invasive follow-up procedures such as amniocentesis or CVS\* which are stressful, painful and can carry a small risk of miscarriage.

## Who can have the IONA® test?

- Suitable for women who are at least 10 weeks pregnant.
- Suitable for all singleton and twin pregnancies.
- Suitable for IVF or surrogate pregnancies.
- Unsuitable for women with cancer or with a trisomy or who have undergone a blood transfusion within the last 12 months.

## How are the IONA® results reported?

- **Low Risk:** means that it is very unlikely your pregnancy is affected by trisomy 21, 18 or 13.
- **High risk:** means that your pregnancy is at increased risk for trisomy 21, 18 or 13 and the result should be confirmed by a follow up invasive procedure such as amniocentesis.
- **No result:** In rare cases there is insufficient fetal DNA in the sample to obtain a result. You may be asked by your healthcare provider for an additional blood sample.

**The IONA® test is a screening test. Suitability for and results from the test must be discussed in detail with your healthcare provider.**

\* Chorionic villus sampling

