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By [Dr David Moore](#), 23 March 2014

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What are screening tests and why are they done?

Screening tests are tests that try to identify the presence of a problem or risk factor, in a person who does not show obvious signs of that problem (that is, they have no symptoms, or are "asymptomatic"). This means, by definition, screening tests are applied to healthy people with no symptoms of disease or problems. Like any test in medicine, screening tests are not perfect, and the risk of an untrue result needs to be weighed against the benefits of detecting the problem. In essence, this means screening tests should be used only if there is a reasonable chance of improving the health or management of the screened person.



What screening tests are done in pregnancy?

In pregnancy, several screening tests are recommended, and others are offered depending on your individual circumstances. These tests look for conditions that may alter the management of your pregnancy (e.g. anaemia, Rhesus-negative blood group, susceptibility to other infections).

In the absence of specific conditions or complications, the [Royal Australian and New Zealand College of Obstetricians and Gynaecologists](#) recommends the following screening tests, either pre-pregnancy or when pregnancy is diagnosed:

- **Full blood count:** this looks for low blood ("anaemia") and may suggest the underlying cause. Sometimes further tests are needed.

2. **Nuchal translucency or "12 week" scan:** this scan is done as part of Combine First Trimester Screening (CFTS) Down Syndrome. The scan is performed between 11 and 14 weeks (usually around 12-13 weeks), and also assessment. The result is not a "yes" or "no" for Down Syndrome; rather, it gives a personalised assessment of the in a specific pregnancy. A scan at this stage is also highly accurate for estimating your due date (error range of 4 days of baby's development to date). Thus, even couples not wishing to undergo risk assessment for Down Syndrome may blood test or specific measurements required for CFTS.
3. **Morphology or "19 weeks" scan:** during this scan a complete "anatomical survey" of baby is taken, to rule out congenital heart defects. It is usually performed between 18 and 24 weeks. Importantly, while a normal scan is very identified during this scan. This scan also determines the position of the developing placenta, to make sure it is clear

Depending on a woman's history and current pregnancy, additional ultrasounds may be required. For instance, if the placenta (at the morphology scan, a later scan (around 32-34 weeks) will be ordered to ensure it has moved clear. In ultrasounds are required, every 2-4 weeks, to ensure both babies are growing well and without complications.

Other screening tests:

Gestational diabetes screen

Gestational diabetes mellitus (or GDM) is a condition that develops in pregnancy, when the body is not able to make enough insulin levels in the normal range. If untreated, it can result in complications such as blood pressure disorders and large babies. GDM affects around 8% of pregnancies, and usually develops in the third trimester, and so a screening test is recommended. Prompt treatment is needed to limit any risk of complications.

36-week blood test

This is a simple blood test to rule out any late pregnancy anaemia, check your platelet count, and confirm the absence of infection. Negative.

Non-invasive prenatal testing (NIPT)

NIPT is a relatively new technology that aims to accurately predict or exclude Down Syndrome and other chromosomal abnormalities by analysing a baby's DNA in the mother's bloodstream. It simply involves a blood test from the mother, and so is "non-invasive" to the mother. It is the most powerful tool for excluding Down Syndrome. Any positive result, however, still needs to be confirmed with amniocentesis. At the moment, there is no Medicare rebate for this test, and generally costs the patient around \$400-\$500. Please guide the use of these tests, so it remains for your doctor to discuss the potential benefits of this test in your particular case.

David practices evidence-based medicine, and strives to ensure all conditions that may complicate a pregnancy are managed according to the latest literature and published guidelines.

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About Dr David Moore



David is a Fellow of the Royal Australian and New Zealand College of Obstetricians and Gynaecologists, a Specialist in Obstetrics and Gynaecology, Queensland. He is highly skilled in the management of complex and high-risk pregnancies, and has specialised in the management of complex and high-risk pregnancies, and has specialised in the management of complex and high-risk pregnancies.

endometriosis, pelvic floor and incontinence surgery. David has completed a Master of Reproductive Medicine management of fertility problems, and can offer the full range of assisted reproductive treatments. He is a Queensland Medical School, and has published both medical journal and textbook contributions.

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