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 **LancetLabSouthAfrica**

 **LancetLab_ZA**

 **lancetlab_za**



PAYMENT PLAN OPTIONS

Should your current medical aid not cover NIPT testing, Lancet Laboratories offers you a four month payment plan option.

4 Month Payment Plan Option

An upfront payment will need to be paid at the time of taking the blood sample. The balance will be payable over 3 consecutive monthly instalments.

	Panorama NIPT	Panorama NIPT + Microdeletions	Panorama NIPT + 22q11.2	Panorama NIPT twins/ Singleton donor
Upfront Payment	R 1, 706.25	R 3,189.75	R 2,658.50	R 1,732.50
Monthly Instalment Amount (3x)	R 1, 706.25	R 3,189.75	R 2,658.50	R 1,732.50
Total	R 6,825.00	R 12,759.00	R 10,634.00	R 6,930.00

The above prices are only valid until 29 February 2020.

For additional information regarding payment plans or Discovery Health NIPT coverage please contact Lancet on 086 163 3337 (Operating Hours: Mon-Fri | 08H00-17H00) or email nipt@lancet.co.za.



MORE INFORMATION

WHO CAN I SPEAK TO FOR MORE INFORMATION?

For Genetic Counselling (pre/post), please contact Dr Karen Milstein, Clinical Geneticist at 086 152 6238 or email: karen.milstein@lancet.co.za.

You can also learn more about Panorama by scheduling a free 15 minute information session with one of Natera's board-certified Genetic Counsellors. They are only available for pre-test counselling. Appointments are available in English only and are limited to certain hours. Simply schedule at:

my.natera.com/services/genetic_information

You can also learn more about this DNA screening test at <http://www.lancet.co.za/prenatal-dna-screening/>

WHERE CAN I TAKE THE TEST?

Lancet Laboratories offers the service of a Travelling Nurse, who is able to visit you at the comfort of your home or office. To book a Travelling Nurse please call 086 163 3337 (Operating Hours: Mon-Fri | 08H00-17H00)

You may also visit one of our many depots nationwide. To find a depot closest to you please go to www.lancet.co.za/locate-a-lab/

This test was developed by Natera, Inc., a laboratory certified under the Clinical Laboratory Improvement Amendments (CLIA). Natera and Panorama are trademarks of Natera Inc., used with permission.

References:

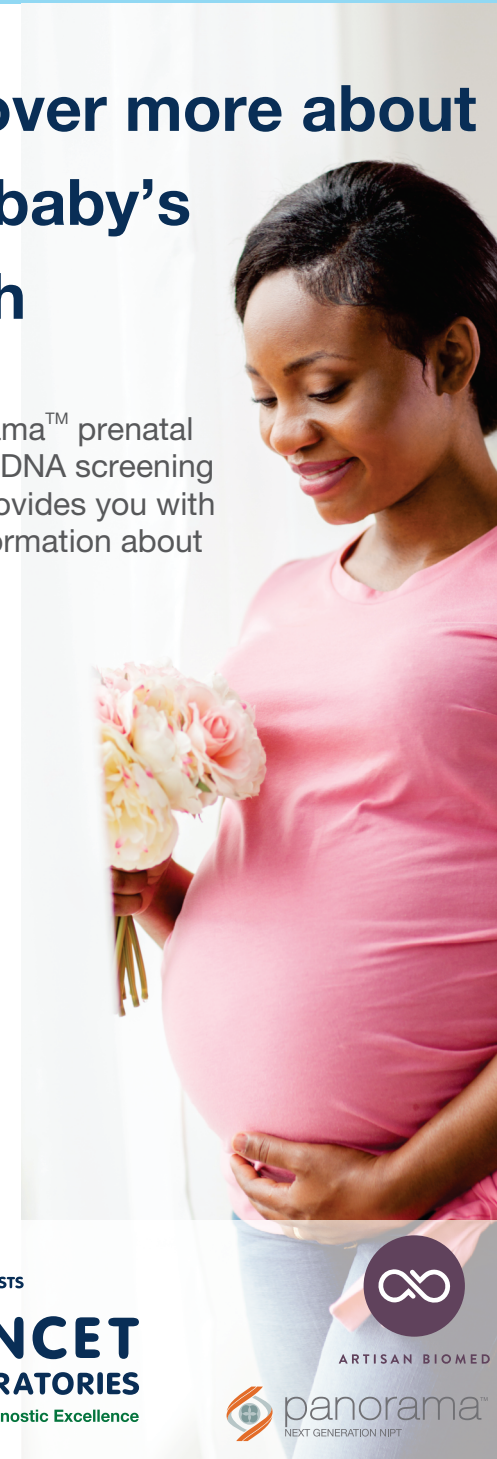
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PANORAMA™ PRENATAL SCREEN

Discover more about your baby's health

The Panorama™ prenatal screen is a DNA screening test that provides you with genetic information about your baby



PATHOLOGISTS
LANCET LABORATORIES
Key to Diagnostic Excellence


ARTISAN BIOMED
 **panorama™**
NEXT GENERATION NIPT

ITEM CODE: B00056

design done and printed by:  **ELECTRONIC LABORATORY SERVICES (PTY) LTD PRINT BUREAU**
corporate branding/brochures/sa/2019/B00056 NIPT 393mmx210mm eng duplex labelnet jan2019.cdr | rev002

WHAT IS NIPT?

Non-invasive prenatal testing (NIPT) uses a blood sample from the mother to analyse DNA from the baby that has crossed into the mother's bloodstream for certain chromosome conditions that could affect your baby's health.

NIPT:

- Screens for genetic abnormalities such as Down syndrome
- Can identify your baby's gender (optional)
- Provides substantially fewer incorrect results than maternal serum screening or other prenatal blood tests
- Can be done as early as nine weeks into your pregnancy
- Poses no additional risk to your baby

WHAT DOES PANORAMA SCREEN FOR?

Singleton pregnancies

- Trisomy 21 (Down syndrome)
- Trisomy 18 (Edwards syndrome)
- Trisomy 13 (Patau syndrome)
- Triploidy
- Monosomy X (Turner syndrome)
- Microdeletions, including 22q11.2 deletion (optional)
- Gender (optional)

Twin pregnancies

- Identical or fraternal twins
- Trisomy 21 (Down syndrome)
- Trisomy 18 (Edward syndrome)
- Trisomy 13 (Patau syndrome)
- Gender of each twin (optional)

If our screening finds that your twins are identical, Panorama can additionally screen for:

- Monosomy X (Turner syndrome)
- 22q11.2 deletion syndrome (optional)

Egg donor or surrogate pregnancies

- Trisomy 21 (Down syndrome)
- Trisomy 18 (Edwards syndrome)
- Trisomy 13 (Patau syndrome)

WHAT ARE MICRODELETIONS?

A small, missing piece of a chromosome is called a microdeletion. Unlike Down syndrome, which occurs more frequently in mothers who are 35 and older, microdeletions occur in pregnancies at the same rate for mothers of any age.

Panorama screens for five microdeletion syndromes associated with serious health problems:

- 22q11.2 deletion (DiGeorge) syndrome
- 1p36 deletion syndrome
- Angelman syndrome
- Prader Willi syndrome
- Cri-du-chat syndrome

IS PANORAMA RIGHT FOR ME?

IF YOU WOULD LIKE TO KNOW whether your baby is at risk for certain genetic conditions, Panorama may be the right option for you. The first step is to talk with your healthcare provider.

Some women have a higher chance of their baby being affected with certain chromosome conditions, like Down syndrome, especially if they:

- Are over the age of 35
- Have certain family histories
- Have abnormal ultrasound findings
- Have abnormal blood test results

Panorama is designed for all pregnant women, regardless of age. We accept samples from:

- Naturally-conceived singleton pregnancies
- Twin pregnancies
- Singleton pregnancies that are using an egg donor or surrogate

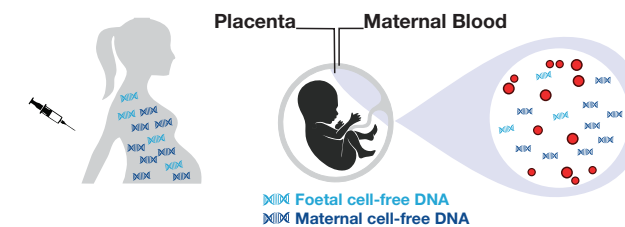
Unfortunately, we cannot accept samples from women in the following categories:

- Bone marrow transplant recipients
- Pregnancy with a vanishing twin
- Pregnancy using an egg donor or surrogate in which there is more than one foetus

HOW IS PANORAMA DIFFERENT?

Other NIPTs cannot tell the difference between mom and baby's DNA. Panorama™ can!

Because of its unique technology, Panorama is the only NIPT that can distinguish between the mom's DNA and the baby's DNA from the placenta. This enables Panorama to be a highly accurate screening test.



FEWER FALSE POSITIVES



Because Panorama analyses the baby's DNA separately, it has a lower false positive and negative rate than other NIPTs.^{1,2,3}

HIGHEST FOETAL SEX ACCURACY



Panorama has the highest reported accuracy in determining the foetal sex, and reporting is optional.^{1,2,3}

TRIPLOIDY



Panorama is the only NIPT that can detect triploidy, a severe chromosomal abnormality that can result in serious pregnancy complications if unmonitored.^{6,7}

ZYGOSITY



Panorama is the only NIPT that can determine zygosity (fraternal or identical twins).

WHAT DO PANORAMA RESULTS TELL ME?

Panorama gives you a personalised risk score and tells you if your baby is at high risk or low risk for the conditions it screens for. Like other screening tests, Panorama does not provide a definitive diagnosis of the condition.

WHAT RESULTS MIGHT I GET WITH PANORAMA?



Low Risk: A Low Risk result indicates that it is unlikely that your baby is affected by one of the conditions on the Panorama panel. Note, however, that a low risk result does not guarantee a healthy pregnancy as Panorama is not a diagnostic test and only screens for certain conditions.



High Risk: A High Risk result means that there is an increased risk that your baby has the condition, but it is not certain. Invasive testing during the pregnancy, such as amniocentesis (amnio) or chorionic villus sampling (CVS), or testing after the baby is born, can tell you for certain if the baby has the condition. Speak with your healthcare provider about your follow-up options. They may recommend that you speak with a genetic chancellor and/or a maternal-foetal medicine specialist.



No Result: In a small percentage of cases, Panorama may not be able to obtain sufficient information from your blood sample to determine an accurate result. If this occurs, a second blood sample may be requested.

WHEN CAN I GET PANORAMA?

You can have this test as early as 9 weeks into the pregnancy. For women who weigh more than 90 kg, it is advised to wait until 12 weeks of pregnancy to improve the likelihood of obtaining a result.

WHEN WILL I GET MY PANORAMA RESULTS?

Most results will be returned to your treating doctor or specialist within 7 - 10 working days.