

Card Number:



Please place collection kit barcode here.

## Non Invasive Prenatal Test (NIPT) Pathology Request

PATIENT INFORMATION	ORDERING CLINICIAN
Name (Last, First):	Ordering Clinician:
DOB (DD/MM/YYYY):	Provider Number:
Address:	Address:
City: State: Postcode:	
Country: Phone:	City: State:
Email:	Postcode: Country:
Are you submitting a father's buccal swab sample with this case? ☐ Yes ☐ No	Phone: +61 Fax: +61
If yes, name of father (Last, First):	Copy of Results to:
LMP (DD/MM/YYYY):/ Due Date (DD/MM/YYYY)://	oopy or recount to.
Maternal Weight: (kg): or (lbs):	Clinician Consent
Was an egg donor or a surrogate used?	I confirm that all information on this form is accurate and complete to the best of my knowledge and that I have explained the benefits and limitations of the Natera Panorama NIPT to the patient to the best of my ability. I hereby order the Natera Panorama NIPT for this patient.
Patient Consent I have read or have had read to me the summary information about the Panorama Non-Invasive	Clinician Signature:
Prenatal Test (NIPT) on the reverse of this form. I have had the opportunity to ask questions of my doctor regarding this test, including the reliability of test results, the limitations, and the alternatives.	Date (DD/MM/YYYY):/
I request and authorise Natera to test my sample(s) for the chromosome abnormalities listed below.	INTERNAL USE ONLY:  NATERA LIMS ID 1010
Patient Signature & Date:/ /(DD/MM/YYYY)	Fax copy of results to Melbourne IVF +61 3 9473 4459
3 SCREENING(S) REQUESTED (SELECT ONE PANEL)	
☐ The Panorama™ Prenatal Panel Screening chromosomes 13,18,21, X & Y and Triploidy  I want gender results included in this report. ☐ Y ☐ N OR	The Panorama™ Extended Panel Screening chromosomes 13, 18, 21, X & Y and Triploidy PLUS the microdeletions selected below  I want gender results included in this report.  Y N  I want to screen for 22q11.2 Deletion syndrome  I want to screen for 22q11.2, 1p36, Cri-du-chat, Angelman, & Prader-Willi.
Please select clinical indications  Advanced maternal age, 1st pregnancy Abnormal serum screening  Other:	
4 SAMPLE COLLECTION DETAILS	Melbourne IVF Collection Centres: MelbourneIVF
When calling the clinic to book in for the Panorama Non-Invasive Prenatal Test - your male partner (if he is the father of the pregnancy) will need to accompany you to have a	East Melbourne: 344 Victoria Pde, East Melbourne Ph: (03) 9473 4444 Mon-Thu 8am - 4pm No appointment required
mouth swab. (The father needs to avoid eating or drinking for half an hour before the test)	Mt Waverley: 3 Meadow Cres, Mt Waverley Ph: (03) 8805 7888 By appointment only
Your result will be sent to your referring doctor 2 - 3 weeks after the sample was taken. Please book in to see your doctor to get your test result.	Templestowe 268 Manningham Road, Templestowe Lower Ph: 03 9006 5500 By appointment only
of the patient(s) was verified and the sample(s) labelled immediately after collection.	Werribee: Lvl 1, 242 Hoppers Lne, Wyndham Private Medical Centre, Werribee Ph: (03) 8742 9300 By appointment only
Date of Blood Draw (DD/MM/YYYY):/ Time of Blood Draw:	Number of Blood Sample Tubes Collected: (2 x 10ml required)
Collection Location: Collector's INITIALS:	& Signature:
5 PAYMENT DETAILS	Detach and destroy this portion after processing
Credit card details must be completed below or the test will not be performed.  If the payment fails to complete, you will be contacted to verify that the details below are If payment is not confirmed within 72 hours, the test will not be processed and a new so	
Card type: ☐ Visa ☐ Mastercard ☐ AMEX (excluding NSW and VIC)	Amount to be debited:
	Signature:

CCV:

Expiry Date:

MIVF000 - 21.02.2022





## Non Invasive Prenatal Test [NIPT] Pathology Request Patient Information

This is a non-invasive blood test which studies the fetal DNA in the mother's blood and can be done from 9 to 39 weeks pregnancy. The fetal DNA tested comes from the placenta; this DNA is identical to the DNA found in the actual cells of the fetus in about 98% of all pregnancies. The condition where the DNA in the placenta is different to that of the fetus is known as confined placental mosaicism.

## It is NOT suitable for:

- Multiple pregnancy
- Donor eggs or embryos
- Women with a past bone marrow transfer

The purpose of the Panorama Non-Invasive Prenatal Test (NIPT) is to screen the fetus for the chromosome abnormalities, including the specific whole extra or missing chromosomes 13,18,21, X and Y plus optionally specified microdeletions (small missing sections of specified chromosomes). You have the option of requesting a screen and reporting of the fetal sex as well. This screen will therefore detect common chromosome problems such as Down syndrome that may lead to a child born with degrees of mental impairment.

## It does NOT detect:

- Any other cause for mental impairment in a child
- Disorders of any other chromosomes, that may lead to an early miscarriage
- Any other cause for birth defects

The test takes two to three weeks to perform. At that time you will receive one of the following results:

High Risk: There is an increased likelihood that your fetus has an abnormality of one of the above chromosomes and further

investigation is strongly recommended.

Low Risk: There is a reduced likelihood that your fetus has an abnormality of one of the above chromosomes.

No result: This can happen when there is insufficient fetal DNA to give a clear result. The test would then need to be repeated

[at no cost to you] and this would add 2 to 3 weeks onto your gestation period by the time you receive a new

result. A high BMI will increase the likelihood of a low/insufficient fetal DNA content.

A "No result" may also happen if the parents are related or the mother's parents are related. Also when there is a multiple pregnancy or vanishing twin or for any other reason the DNA pattern is not clear. A repeat test in these

instances is unlikely to generate a result and you will be offered a refund.

The result will be sent back to your referring doctor who will contact you to let you know.

With confined placental mosaicism affecting 1-2% of pregnancies, an incorrect high- or low- risk result can occur. The Panorama screen is not a diagnostic test – it will not confirm any of these chromosome abnormalities. It will only provide the risk for each of these in your current pregnancy. Therefore, **DECISIONS ABOUT YOUR PREGNANCY SHOULD NEVER BE MADE BASED ON THESE SCREENING RESULTS ALONE AS THEY NEITHER CONFIRM NOR RULE OUT THE PRESENCE OF A CHROMOSOME ABNORMALITY IN THE FETUS.** 

It is the responsibility of your doctor ordering this test to understand the reliability of the test results, the limitations and the alternatives and to explain them to you. Before you commence with the test and sign this form, please ask your doctor for more information about the test and the results if required.