



Please place collection kit barcode here.

123456-2-X

PANORAMA INTERNATIONAL REQUISITION

PLEASE COMPLETE ALL FIELDS IN ENGLISH. REQUISITION FORMS SUBMITTED WITH MISSING INFORMATION MAY CAUSE A DELAY IN TURNAROUND TIME OF THE TEST.

1

PATIENT INFORMATION

First Name _____

Last Name _____

Date of Birth (DD/MM/YYYY) _____

Address _____

City _____

Country _____

Telephone _____

Email _____

Weight (kg) _____ Height (cm) _____

Expected Due Date (DD/MM/YYYY) _____

Patient must be at least 9 weeks gestation. 22q is not available for dizygotic twins or egg donors. Extended panel is not available for twins or egg donors.

For twin or surrogate pregnancies, check all that apply. We do NOT accept vanished twin, multiple gestation with more than 2 fetuses, or twins conceived using a surrogate or egg donor.

- IVF conceived pregnancy:** Age of mother at egg retrieval: _____
- Ongoing **twin pregnancy:** Monochorionic Dichorionic Don't Know
- Surrogate** or **egg donor** pregnancy

2

CLINIC INFORMATION

Clinic Name _____

Telephone _____

Natera® LIMS ID _____

Ordering Clinician Name _____

3

SCREENING OPTIONS

SELECT SCREENING OPTION BELOW

- PANORAMA PRENATAL PANEL** Chromosomes 13, 18, 21, X & Y; Triploidy
- PANORAMA PRENATAL PANEL + 22q11.2 DELETION** Chromosomes 13, 18, 21, X & Y; Triploidy; 22q11.2 deletion
- PANORAMA EXTENDED PANEL** Chromosomes 13, 18, 21, X & Y; Triploidy; 22q11.2 deletion PLUS four microdeletions
- Check to add fetal sex to report (available with any screening option above)

Date of Sample Collection (DD/MM/YYYY)

4

DISPOSITION OR RETENTION OF SAMPLES

Laboratory (Reseller) represents and confirms that the patient has given informed consent compliance with applicable law to Natera's following sample disposition or retention policy: PATIENT UNDERSTANDS AND CONSENTS THAT: (i) her/his sample will be sent to the United States for performance of the test; (ii) Natera may retain the patient's leftover, de-identified samples to use for medical and technology advancement, research & development, product validation and quality assurance, independently or collaboration with third-party partners, either in or outside the United States; and (iii) patient and patient's heirs will not receive any payments, benefits, or rights to any resulting products or discoveries.