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Signatera™ Residual disease test (MRD)





Signatera™ for early detection of molecular residual disease using ctDNA

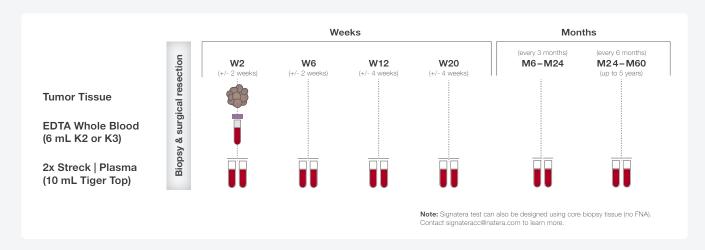
Personalized for each patient's tumor

Ordering Signatera for patients with solid tumors

Signatera is custom-designed for each patient using their own tumor tissue.

	RECURRING ORDER PROGRAM MRD & Recurrence Monitoring Program	IMMUNOTHERAPY & TREATMENT Monitoring Program
Clinical Use Case	Order before or after surgery to help inform surgical and/or therapeutic intervention	Use Signatera to determine treatment effectiveness, or to help rule out disease progression
Medicare Coverage	Stage II-III colorectal cancer and Stage IV oligometastatic colorectal cancer	Pan-cancer immunotherapy monitoring
Turn around time: 2–3 weeks for initial test setup		

Recommended ordering schedule



Ordering Altera Comprehensive Genomic Profiling Assay

Clinical use case

- Somatic profiling includes: RNA sequencing (call fusions with established clinical reference, detect novel fusions), introns, promoters; reporting TMB, MSI, and genes related to HRD
- \triangleright For tumor profiling analysis and to help guide therapy selection
- > Turn around time is approximately 2 weeks from the day all required samples for testing have been received in our laboratory.

Flexible Ordering

Choose to have your patient's blood draw managed by the clinic on-site or by Natera's mobile phlebotomy service.

For clinic-managed Signatera blood draws:

- On the Requisition Form, fill in the "Date of Blood Collection" and write "Drawn at Clinic" above the date
- On the Provider's portal, in the section "Blood draw managed by," select "Clinic"

- On the Requisition Form, leave "Date of Blood Collection" blank and write "Natera manage" above the line
- On the Provider's portal, in the section "Blood draw managed by," select "Natera"

For Natera-managed blood draws, provider intervention is not required.



Natera Customer Care will reach out to patient for availability for blood draw, and ship blood sample kit with pre-filled Requisition Form to patients

Phlebotomy services will reach out to patients to confirm date, time, and blood draw location



Patients will be notified when phlebotomist is on the way



Phlebotomist will ship kit with blood samples and completed Requisition Form

Natera's phlebotomy services will become the default for subsequent draws, unless updated in the portal or notifying Customer Care at **1.650.489.9050** or **signateracc@natera.com**.

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The tests described have been developed and their performance characteristics determined by the CLIA-certified laboratory performing the test. The tests have not been cleared or approved by the US Food and Drug Administration (FDA). Although FDA is exercising enforcement discretion of premarket review and other regulations for laboratory-developed tests in the US, certification of the laboratory is required under CLIA to ensure the quality and validity of the tests. CAP accredited, ISO 13485 certified, and CLIA certified. © 2022 Natera, Inc. All Rights Reserved. SGN_STANDING_ORDER_OS_20220125_NAT-8020082

