

Signatera[™] Testing for Circulating Tumor DNA (ctDNA) Detection

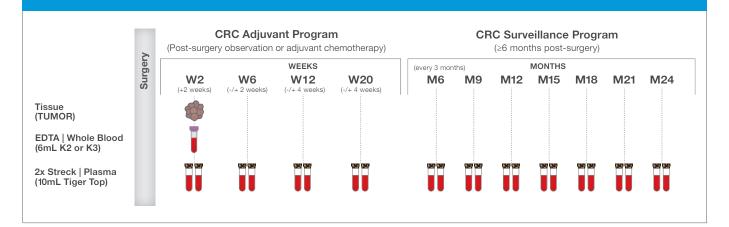
Detect residual disease and recurrence early in colorectal cancer (CRC)

How often should Signatera be ordered for CRC patients?

Depending on where your patients are in their treatment journey, two Signatera testing settings are recommended to detect residual disease.

	Adjuvant Setting	Surveillance Setting
When to Join	Within 6 months after surgery	≥6 months after surgery
Clinical Use Case	After surgery, Signatera can help to evaluate the need for adjuvant chemotherapy or to avoid unnecessary treatment.	Use Signatera in addition to CEA testing to detect recurrence earlier and more accurately while the tumor is potentially resectable.
Medicare Coverage	Stage II-III colorectal cancer	Stage II-III colorectal cancer

Blood and tissue collection timeline

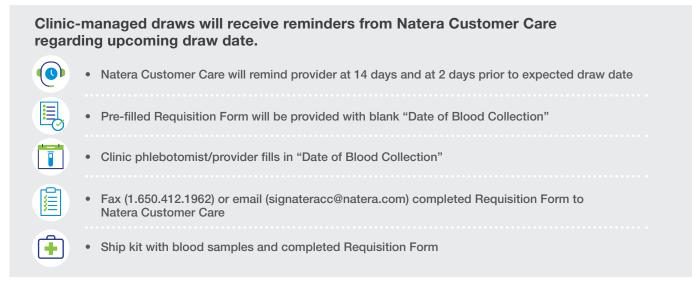


How to place an order for Signatera

To streamline the Signatera testing process, Natera offers the flexibility for blood draws to be managed by the clinic on-site, or by Natera's blood draw services.

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; OR
- On the Provider's portal, in the section "Blood draw managed by," select "Clinic"



The clinic will become the default draw location for subsequent draws, unless updated in the portal or notifying Customer Care at **1.650.489.9050** or **signateracc@natera.com**.

For blood draw managed by Natera's phlebotomy services:

- On the Requisition Form, leave "Date of Blood Collection" blank and write "Natera manage" above the line; OR
- 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 test described has been developed and its performance characteristics determined by the CLIA-certified laboratory performing the test. The test has 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. © 2020 Natera, Inc. All Rights Reserved. SGN_STANDING_ORDER_OS_20201007_NAT-8020082

