

Know Cancer's Next Move

- > Does my patient need adjuvant chemotherapy?
- > Is the treatment working?
- > Is the cancer recurring?

Discover unique insights with molecular residual disease (MRD) detection, informed by ctDNA

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Signatera™
Residual disease test (MRD)

The LifeLabs Genetics logo features the word "LifeLabs" in a large, blue, sans-serif font with a red dot above the "i". Below it, the word "GENETICS" is written in a smaller, blue, all-caps, sans-serif font, with a red vertical bar inside the letter "I".

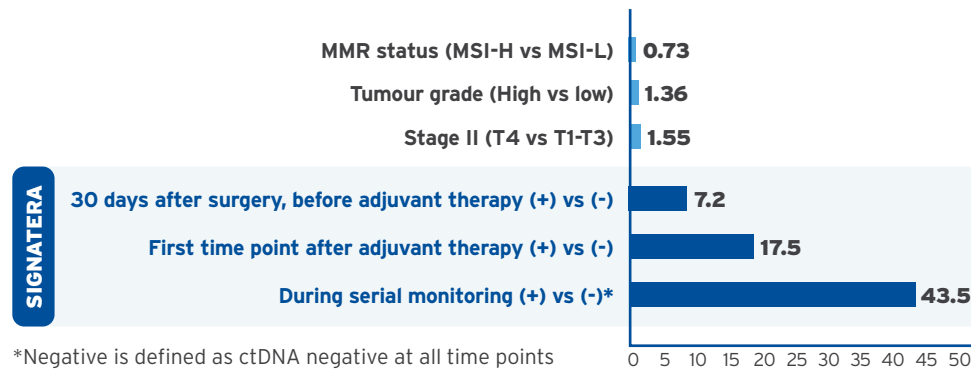
LifeLabs
GENETICS®

When to use Signatera™ for patients with CRC

In the adjuvant setting

Use after surgery to evaluate the need for adjuvant chemotherapy and potentially avoid unnecessary treatment

Signatera™ MRD status outperforms known clinicopathologic risk factors in predicting relapse¹⁻⁴



- > 97% of patients with a positive Signatera™ result relapsed without additional treatment¹
- > Serial testing improved sensitivity and negative predictive value of results

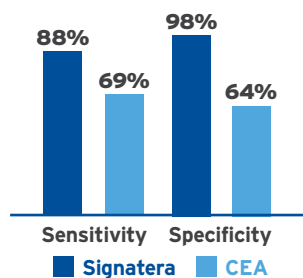
In the surveillance setting

Use along with CEA testing and other surveillance modalities to detect recurrence earlier, to enable surgical resection or other early intervention

Signatera™ was shown to detect relapse more accurately than CEA with clinically meaningful lead times over CT scans¹

- > Get clarity when evaluating patients with indeterminate CEA levels or CT scans
- > Signatera™ facilitates shared decision-making and confident treatment planning

Accuracy in detecting relapse



Average lead time of ctDNA detection before CT scan

8.7
MONTHS

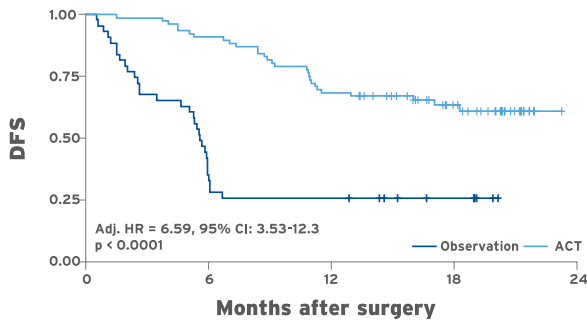
Maximum lead of 16.5 months

CEA = carcinoembryonic antigen; CT = computed tomography; ctDNA = circulating-tumour DNA

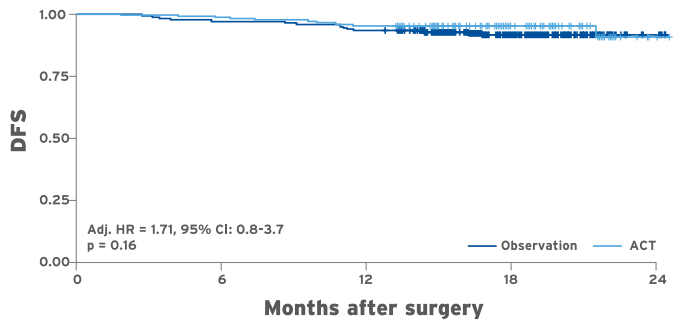
A large-scale prospective, MRD-guided study evaluated the clinical utility of ctDNA analysis in colorectal cancer (CRC)

MRD-positive CRC patients at 4 weeks post-op benefited significantly from chemotherapy while MRD-negative patients at 4 weeks post-op did not demonstrate any significant trend in treatment benefit⁵

MRD-positive patients at 4 weeks post-op



MRD-negative patients at 4 weeks post-op



Track ctDNA dynamics to enable longitudinal monitoring

- > Signatera™ reports presence/absence of ctDNA and ctDNA quantity in terms of MTM/mL for longitudinal assessment

Final Results Summary

Signatera™ Negative

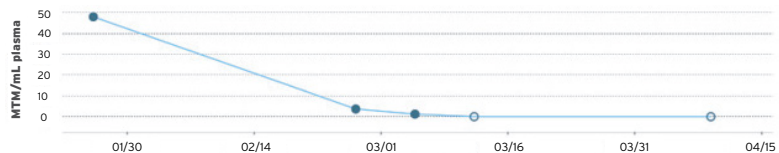


MTM/mL

Not Detected

Mean tumour molecules per mL is calculated based on the mean of ctDNA molecules detected per mL of the patient's plasma

Historical Results



MTM = mean tumour molecules

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Signatera™
Residual disease test (MRD)



Just like no two tumours are alike, Signatera™ is personalized for each patient



Tumour-informed MRD assay for individualized care

- > Customized for each patient's unique tumour signature using Whole Exome Sequencing (WES) to target the top clonal mutations



High sensitivity and specificity for accurate MRD assessment

- > By only tracking tumour-specific variants, sensitivity is optimized with a LOD down to 0.01% VAF⁶
- > Filters out germline and CHIP mutations to reduce background noise and to minimize false positives

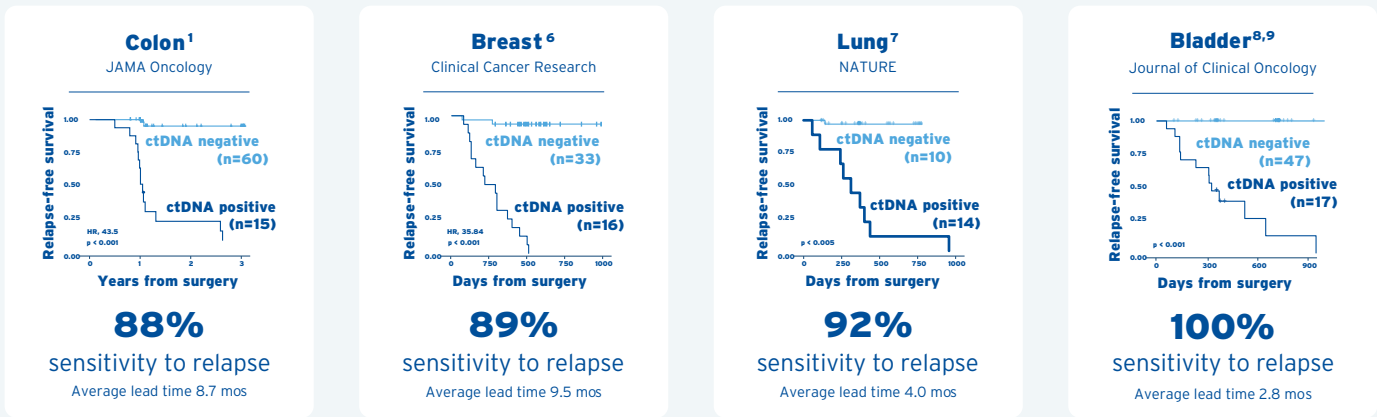


Reliable longitudinal monitoring for confident decision-making

- > Tracks ctDNA dynamics by MTM/mL to enable longitudinal monitoring with a simple blood draw
- > Follows clonal mutations that should persist as the tumour evolves

LOD = limit of detection; CHIP = clonal hematopoiesis of indeterminate potential; VAF = Variant allele frequency; WES = whole exome sequencing

Signatera™ is validated across multiple tumour types^{1,6-8}



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- Data on file

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Signatera™
Residual disease test (MRD)

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