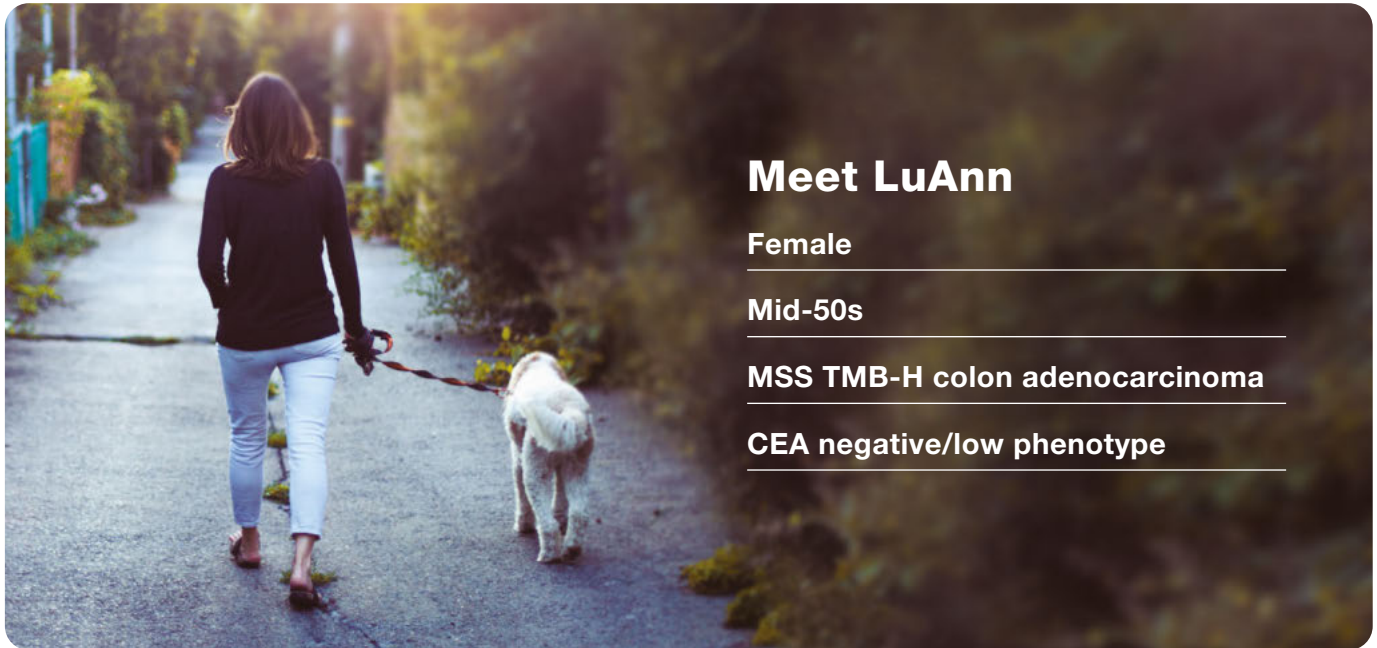


Stage IV colon cancer patient receiving immunotherapy



Meet LuAnn

Female

Mid-50s

MSS TMB-H colon adenocarcinoma

CEA negative/low phenotype



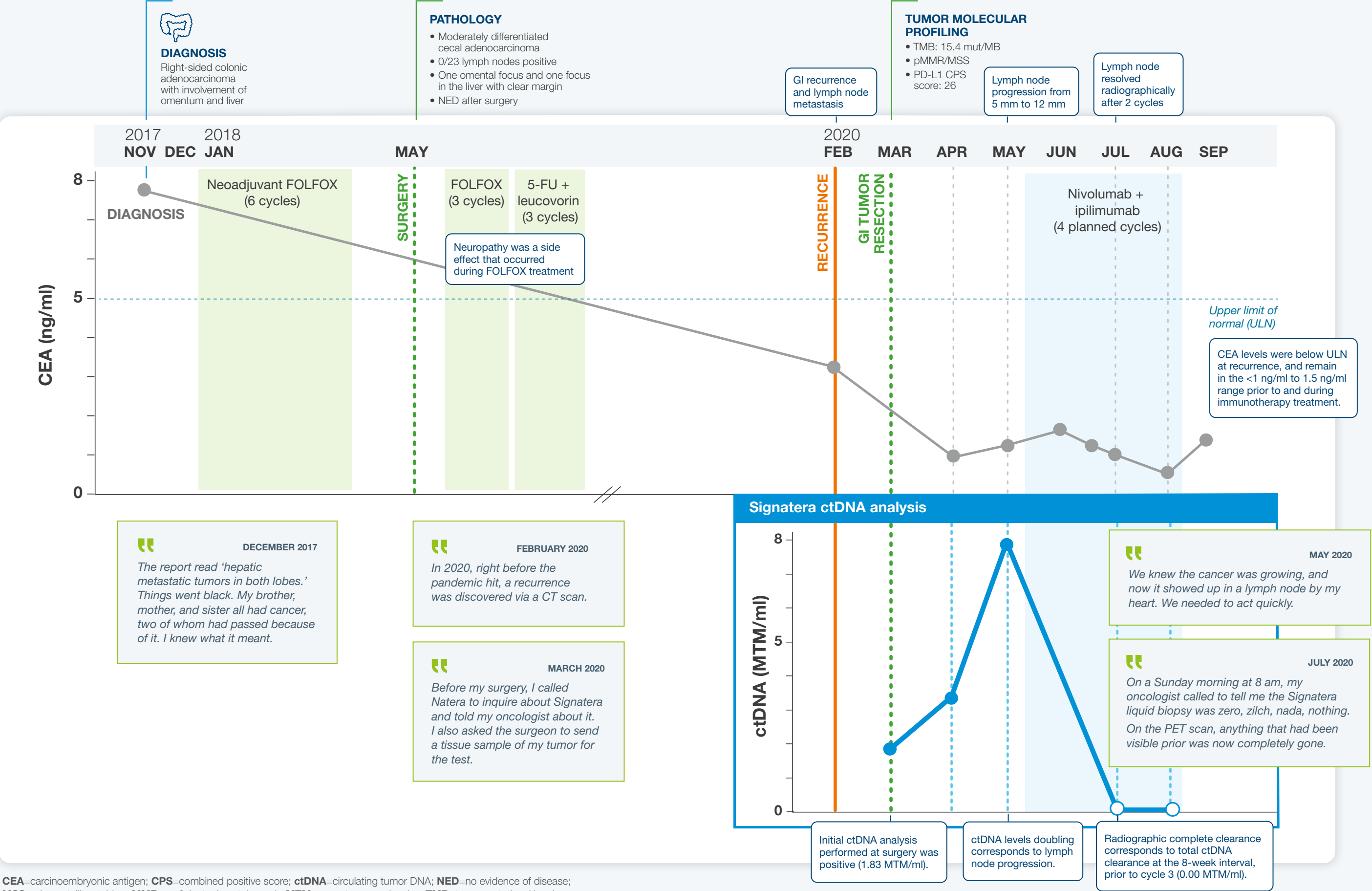
Unfortunately, I never had any symptoms of colon cancer. Perhaps if I had, I would have gotten a colonoscopy at an earlier age. After awakening from the sedation and barely coherent, my doctor told me there was a 7 cm mass in my cecum.

Patient history:

- 55-year-old woman presented with right-sided colonic adenocarcinoma and omental and focal liver metastases
- No evidence of disease after neoadjuvant chemotherapy and surgical resection of primary tumor and metastatic foci
- Despite normal CEA levels, recurrence was detected in the abdominal wall and lymph node after completion of adjuvant chemotherapy

How Signatera was incorporated:

- Serial Signatera ctDNA results were used as alternative indicators of tumor burden
- Signatera ctDNA testing provided an early indication of tumor response to immune-checkpoint inhibition



This case study was presented at the Society for Immunotherapy of Cancer (SITC) 2020 Annual Meeting.

Schneider CJ, Krainock M, Malhotra M, et al. ctDNA clearance and radiographic resolution of lymph node metastasis in a patient with metastatic microsatellite stable colorectal cancer on immunotherapy. Poster presented at SITC 2020 Virtual Meeting; November 10-15, 2020. Poster #26.

The Takeaway



- ctDNA is a dynamic biomarker for monitoring tumor response to immune checkpoint inhibitor treatment, when used in conjunction with radiographic imaging.
- ctDNA can be used as a serologic surrogate to assess systemic disease burden and may outperform standard biomarkers such as CEA.

Value of Signatera



- Signatera is a personalized, tumor-informed assay for ultrasensitive, real-time detection of molecular residual disease.
- Rapid ctDNA clearance correlated with radiographic resolution of metastatic disease provides confidence that Signatera can be used to identify early response during immunotherapy response monitoring.

Learn more at nateraoncology.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. © 2021 Natera, Inc. All Rights Reserved. 20210209_NAT-8020361

