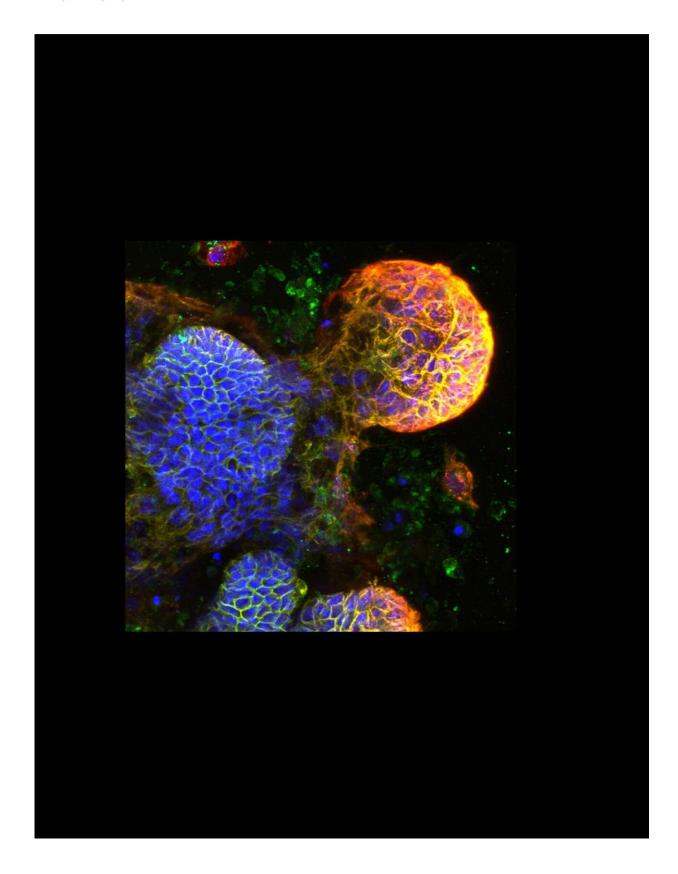


# Biomarker predicts bowel cancer recurrence

June 3 2021





Crypts and buds in the small intestine and colon. Credit: Dr. Maree Faux, WEHI.



A biomarker in the blood of patients with bowel cancer may provide valuable insight into the risk of cancer relapse after surgery and the effectiveness of chemotherapy.

Research published in *PLOS Medicine* found circulating tumor DNA (ctDNA) measured before and after <u>surgery</u> provided a reliable marker for predicting whether the cancer would recur following chemotherapy treatment.

The ctDNA also provided a real-time measure of the effectiveness of chemotherapy, highlighting the potential for this test to provide an early indication of the success of chemotherapy in eradicating microscopic cancer.

### **Prognostic impact of ctDNA**

Led by Associate Professor Jeanne Tie, who is also a medical oncologist at the Peter MacCallum Cancer Centre and Western Health, the research followed a group of patients with metastatic <u>bowel cancer</u> who had secondary cancer in the liver that had been removed by surgery. The study builds on earlier research reported in 2018.

The ctDNA test looks for fragments of tumor DNA in a patient's blood before and after the removal of a cancerous tumor.

The presence of ctDNA in the blood of patients after surgery provides evidence of remaining microscopic tumors, enabling researchers to predict the likelihood of the cancer reoccurring.

Associate Professor Tie said the study once again confirmed the prognostic impact of ctDNA.



"What we found is that if ctDNA is present after surgery, it predicts an almost 100 per cent recurrence rate for these patients," she said.

"In contrast, for patients who were ctDNA-negative after surgery, the likelihood of the cancer reoccurring was far lower, about 25 per cent."

## Measuring chemotherapy in real time

Associate Professor Tie said ctDNA also provided an indication of the effectiveness of chemotherapy.

"This biomarker could also identify whether patients would respond to chemotherapy treatment," she said.

"Until now, we had no way of measuring the effectiveness of chemotherapy in real time. The usual process is to do the surgery to remove the cancer metastases, give the patient chemotherapy, and then follow up with CT scans every six to 12 months, to see if the cancer recurs. And if the cancer does recur, you know the treatment hasn't worked. By measuring the ctDNA in the blood, we could immediately see whether the chemotherapy had cleared the cancer and were therefore able to predict the likelihood of the cancer recurring."

Associate Professor Tie said ctDNA biomarkers might allow clinicians to intervene earlier.

"Cancer that can be detected on a CT scan is unlikely to be curable by chemotherapy. But if we are able to detect microscopic disease, that we can't pick up on a scan, we can intervene earlier and potentially still offer the patient a chance of cure."

### Promising sign for the future of cancer treatment



Associate Professor Tie said while ctDNA technology was already being used in the US, further research was needed before it could be rolled out in Australia.

"The test needs to be very sensitive to be able to pick up microscopic cancer cells. I am hopeful the new technology coming through will have enough sensitivity that we will be able to use this technique in the years ahead to improve patient care and treatment," she said.

"With further development of this technology, this could also mean patients with a low recurrence risk could avoid unnecessary <a href="mailto:chemotherapy">chemotherapy</a>."

**More information:** Jeanne Tie et al, Circulating tumor DNA dynamics and recurrence risk in patients undergoing curative intent resection of colorectal cancer liver metastases: A prospective cohort study, *PLOS Medicine* (2021). DOI: 10.1371/journal.pmed.1003620

#### Provided by Walter and Eliza Hall Institute

Citation: Biomarker predicts bowel cancer recurrence (2021, June 3) retrieved 6 May 2024 from <a href="https://medicalxpress.com/news/2021-06-biomarker-bowel-cancer-recurrence.html">https://medicalxpress.com/news/2021-06-biomarker-bowel-cancer-recurrence.html</a>

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