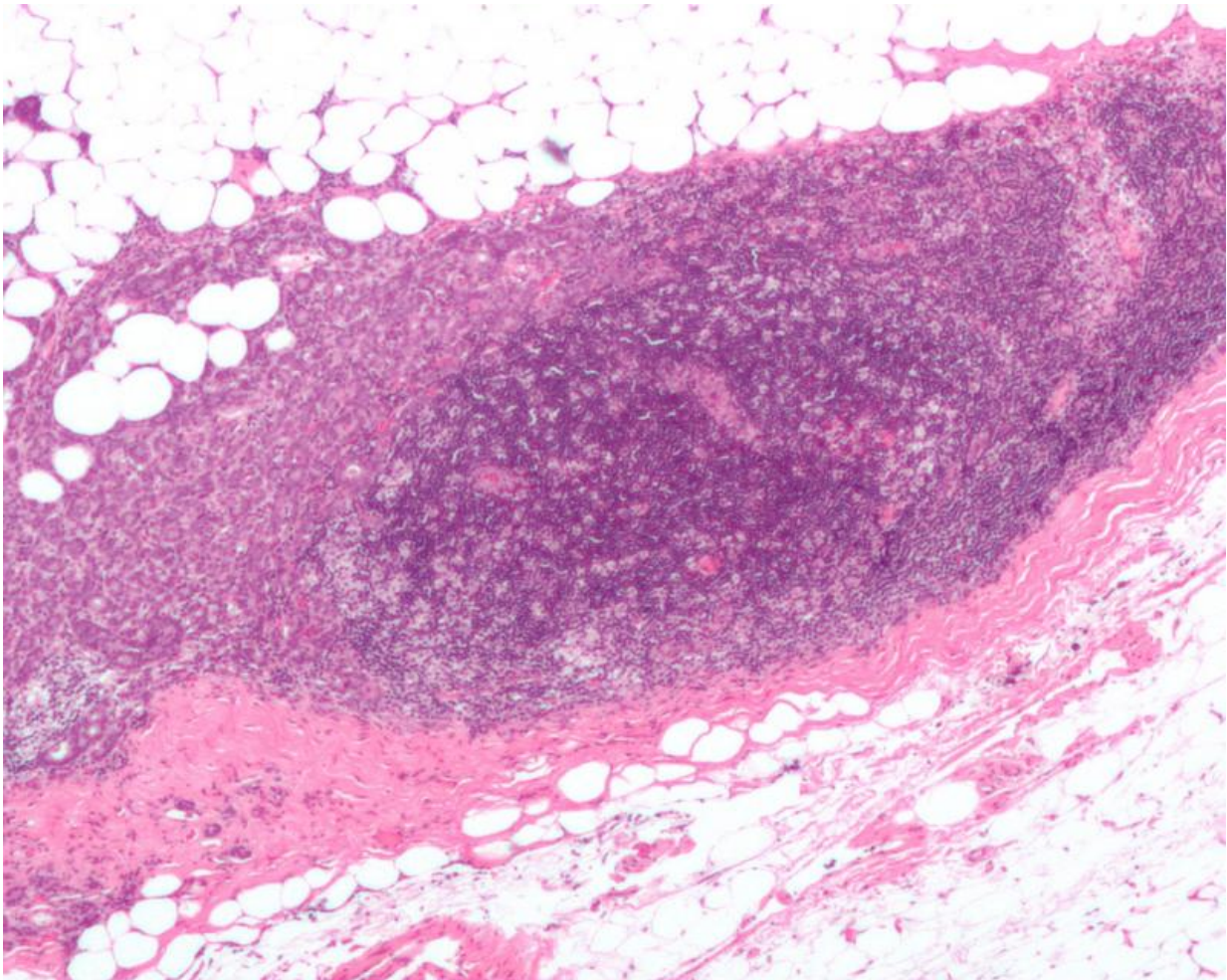


Test could predict whether breast cancer will spread to the brain

October 21 2015



Micrograph showing a lymph node invaded by ductal breast carcinoma, with extension of the tumour beyond the lymph node. Credit: Nephron/Wikipedia

Women with particularly aggressive forms of breast cancer could be identified by a test that predicts whether the disease is likely to spread to the brain.

An analysis of almost 4,000 patients with breast cancer found that testing for high activity in a particular gene called alpha beta (α B)-crystallin could pick out [women](#) who were at greater risk of developing secondary brain tumors compared to women who tested negative.

A team including scientists at The Institute of Cancer Research, London, found that women whose breast cancer had begun to spread and who tested positive in the α B-crystallin test were three times more likely to have disease that spread to the brain than those who tested negative.

The international research effort also included researchers at the University of North Carolina at Chapel Hill, the University of Wisconsin-Madison and the British Columbia Cancer Agency, Canada.

The study was published today in the journal *npj Breast Cancer* and funded by the Breast Cancer Research Foundation, the US National Cancer Institute, Cancer Research UK and the Canadian Breast Cancer Foundation.

Women can live for years with breast cancer that has begun to spread around the body—but the exception is spread to the brain, which usually indicates a woman is in her last few months of life.

In the study, the researchers analysed 969 breast cancer tumors which ultimately spread, or metastasised, to new sites in the body, from a database of almost 4,000 breast cancers from the British Columbia Cancer Agency. 141 had spread first to the brain.

Some 127 scored positive for α B-crystallin, and 842 negative—with

those testing positive three times more likely to have spread first to the brain.

In a further analysis of all 4,000 cases, α B-crystallin was also linked with a significantly higher risk of death—with 36 per cent of women with α B-crystallin positive cancer, compared to 25 per cent with α B-crystallin negative, dying within ten years of diagnosis.

Study co-leader Dr Maggie Cheang, Senior Staff Scientist at The Institute of Cancer Research, London, said:

"Spread of breast cancer to the brain is unfortunately very dangerous, and usually leads to death within months. It's important to find new ways to identify women who are most at risk of their cancer spreading to the brain, so that doctors can work out which women might need more intensive or new treatments to try to keep their cancer at bay for longer.

"Our study linked a positive score in this test with quicker spread to the [brain](#), and importantly showed the factor we were measuring is providing information on patient outcome independently of other biomarkers already measured in the clinic.

"The test needs further development before it will be ready for routine clinical use, but ultimately the first use of this type of test could be to identify opportunities for women with advanced specific types of breast cancer—such as triple negative—to enter clinical trials of new treatments."

Canada study leader Professor Torsten Nielsen, Pathologist at the British Columbia Cancer Agency, said:

"It has taken us years to assemble the clinical outcome database and tissue samples, generate the immunohistochemical biomarkers, gene

expression profiles and analyse the data for this study—and we were delighted to find a definite link between alpha beta crystallin and [breast cancer](#) progression, which we hope will ultimately improve clinical outcomes.

"Our study also definitely demonstrates the importance of team science to deliver a study involving international collaborations".

Provided by Institute of Cancer Research

Citation: Test could predict whether breast cancer will spread to the brain (2015, October 21) retrieved 4 May 2024 from <https://medicalxpress.com/news/2015-10-breast-cancer-brain.html>

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