

## Mobile DNA elements restructure cancer genomes

August 7 2014



In a lung cancer genome, three LINE-1 copies located at human chromosomes 22, 14, and 13 spread non-repetitive DNA by transduction. Credit: DOI: 10.1126/science.1251343



(Medical Xpress)—Researchers have found a novel mechanism that can restructure a cancer's genome and introduce hundreds of mutations. Previously overlooked through misidentification, this mechanism, a common feature of lung and colorectal cancers, raises questions about the effect environmental factors can have on the evolution of a tumour.

Seventeen per cent of the <u>human genome</u> is comprised of 'parasitic', repeated DNA segments called LINE-1 elements that, occasionally, copy themselves and move to new locations in the <u>genome</u>. More rarely, they will aberrantly pick up a piece of adjacent, non-LINE-1 DNA and copy that to a new site: this is known as LINE-1 transduction.

"One in four of all LINE-1 events causing mutations in cancer is due to transduction, a significant contribution to the mutation landscape that has not previously been documented," says Dr Peter Campbell, senior author from the Wellcome Trust Sanger Institute. "Until this work, changes of this type remained mostly ignored because they were misinterpreted; now it's important that we understand this process."

Looking at 290 tumour samples from 12 different cancer types, the team found 3,000 sites where LINE-1 elements were mobilised solely in the tumour. In 24 per cent of the sites, small pieces of non-repetitive DNA were transduced by LINE-1 into another position in the genome, sometimes mobilising exons and even complete genes. These transductions were particularly common in lung and <u>colorectal cancers</u>. Researchers believe this may be caused by changes in methylation levels, which can be caused by <u>environmental factors</u>.

The researchers developed an algorithm that can search through the genome of a cancer sample and identify the origin and subsequent locations of the mobile LINE-1 elements. Strikingly, of the 500,000 LINE-1 elements that are found in the human genome, this study identified just 74 that reproduce themselves in cancer genomes. Of



these, just two gave rise to one-third of all the transduction events identified in the study. Researchers also analysed the activity of LINE-1 elements in a tumour over time. They found that different elements became active and inactive at different points.

"Finding so few LINE-1 elements active in these tumours was surprising, so we looked to see if we could find a molecular explanation for this privileged activity" says Dr Jose Tubio, first author from the Sanger Institute. "We found that methylation of LINE-1 plays a crucial role in switching the elements on and off."

Larger datasets will be required to answer questions raised in this research about the functional consequences for individual genes interrupted by LINE-1 events and the influence these events may have on the development of a cancer. Researchers will now apply their method to 2,000 cancer samples collected by the Pan-Cancer Initiative, a collaboration involving The Cancer Genome Atlas and the International Cancer Genome Consortium.

"We believe that the majority of LINE-1 events are likely to be passenger mutations rather than drivers in cancer evolution," says Dr Campbell. "To understand the functional impact of these events on genes and on the progression of a tumour, we will need to survey the topography of <u>cancer</u> mobile elements in considerably larger scale, across thousands of <u>cancer genomes</u>, integrated with other mutational processes and transcriptional data."

**More information:** Extensive transduction of non-repetitive DNA mediated by L1 retrotransposition in cancer genomes. Tubio, JMC et al *Science* 2014. <u>DOI: 10.1126/science.1251343</u>



## Provided by Wellcome Trust Sanger Institute

Citation: Mobile DNA elements restructure cancer genomes (2014, August 7) retrieved 7 May 2024 from <u>https://medicalxpress.com/news/2014-08-mobile-dna-elements-cancer-genomes.html</u>

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