

Optimizing technologies for discovering cancer cell mutations

September 10 2018



Credit: CC0 Public Domain

Cancer cells often have mutations in their DNA that can give scientists clues about how the cancer started or which treatment may be most effective. Finding these mutations can be difficult, but a new method may offer more complete, comprehensive results.

A team of researchers has developed a new framework that can combine three existing methods of finding these large mutations—or structural variants—into a single, more complete picture.

Feng Yue, assistant professor of biochemistry and molecular biology at Penn State College of Medicine, said the new [method](#)—published today (Sept. 10) in *Nature Genetics*—could help researchers find new structural variations within cancer cell DNA and learn more about how those cancers begin.

"We were able to design and use this computational framework to connect the three methods together, to get the most comprehensive view of the genome," Yue said. "Each method by itself can only review a portion of the structural variations, but when you integrate the results of the three different methods, you can get the most comprehensive view of the cancer genome."

Structural variants are large mutations in DNA that can result in cancer causing [genes](#) being turned on. For example, certain types of brain cancer, such as the types that afflicted senators John McCain and Edward Kennedy, can be caused by structural variants that amplify certain cancer causing genes. In some cancers, knowing a patient has this abnormality helps doctors decide on the best treatment plan.

Yue said finding these structural variants is important for several reasons.

"If you are a cancer patient, knowing about the structural variants that lead to the cancer can help us understand why you got sick and possibly which treatment could be best," Yue said.

The researchers used three existing methods for finding structural variants: optical mapping, high-throughput chromosome conformation

capture (known as Hi-C), and whole genome sequencing, which has been used to discover the majority of structural variants that are already known.

Using their new method, the researchers were able to find structural variations for over 30 types of cancer cells. The team of scientists was also able to use the same methods to begin to learn why certain classes of structural variants may be contributing to cancers.

"Many of the structural variants that are found in human cancers do not appear to directly impact a gene," said Jesse Dixon, a fellow at the Salk Institute in San Diego, and one of the co-authors of the work. "Instead, many structural variants appear in non-coding portions of the genome, what people have historically referred to as junk DNA, and it can be a bit of a mystery as to why these may be contributing to cancer."

What the researchers were able to observe is that some structural variants appear to affect regulatory gene "switches" in noncoding sequences of DNA. Defective switches prevent appropriate turning on or off of specific genes and this can contribute to cancer.

"With many cancers, the gene itself is OK but the 'switch' that controls it is what's causing problems," Yue said. "Using our approach, it's possible that we could find out that the switch was broken and find a cure based on the specific target for that switch. If it's switched off, for example, maybe we could use gene editing technology to turn it back on."

They also used the Hi-C method to explore how structural variations can affect 3-D genome structure—how DNA folds itself within the cell.

"Cells are small, but their DNA is very long. Laid out in a line, all the DNA from one cell would be more than two meters long," said Dr. Job Dekker, professor and co-director of the program in Systems Biology at

University of Massachusetts, investigator of the Howard Hughes Medical Institute and a senior author of the paper. "That's why DNA needs to fold in intricate ways. We have found that genomic alterations in cancer cells can lead to differences in how the genome folds and this can lead to cases where genes become turned on or off by the wrong regulatory switches."

The team of scientists was able to discover that structural variants impact genome folding in cancer [cells](#), and that these changes may be contributing to the cancer.

"One of the discoveries we had made in the past is that our genome is folded up into distinct structures, almost like little neighborhoods," Dixon said. "It appears as though some structural variants cause changes to these neighborhoods, such that a cancer causing gene is moved from a neighborhood where the gene is kept quiet into one where the gene becomes activated."

In the future, Yue and his research team plan to apply the new method in more cancer patients and they are working closely with the Penn State Institute for Personalized Medicine. The scientists suggest that this work could lead to a better ability to predict which structural variants may contribute to cancer, and which genes they may be targeting.

"If we can understand which mutations are driving which genes, this has the potential to suggest that the [cancer](#) would be susceptible to treatment by particular drugs that target those genes," Yue said. "Such an approach has been really challenging in the past for non-coding structural mutations in the [genome](#)."

More information: Integrative detection and analysis of structural variation in cancer genomes, *Nature Genetics* (2018). [DOI: 10.1038/s41588-018-0195-8](#) ,

www.nature.com/articles/s41588-018-0195-8

Provided by Pennsylvania State University

Citation: Optimizing technologies for discovering cancer cell mutations (2018, September 10)
retrieved 1 May 2024 from

<https://medicalxpress.com/news/2018-09-optimizing-technologies-cancer-cell-mutations.html>

<p>This document is subject to copyright. Apart from any fair dealing for the purpose of private study or research, no part may be reproduced without the written permission. The content is provided for information purposes only.</p>
--