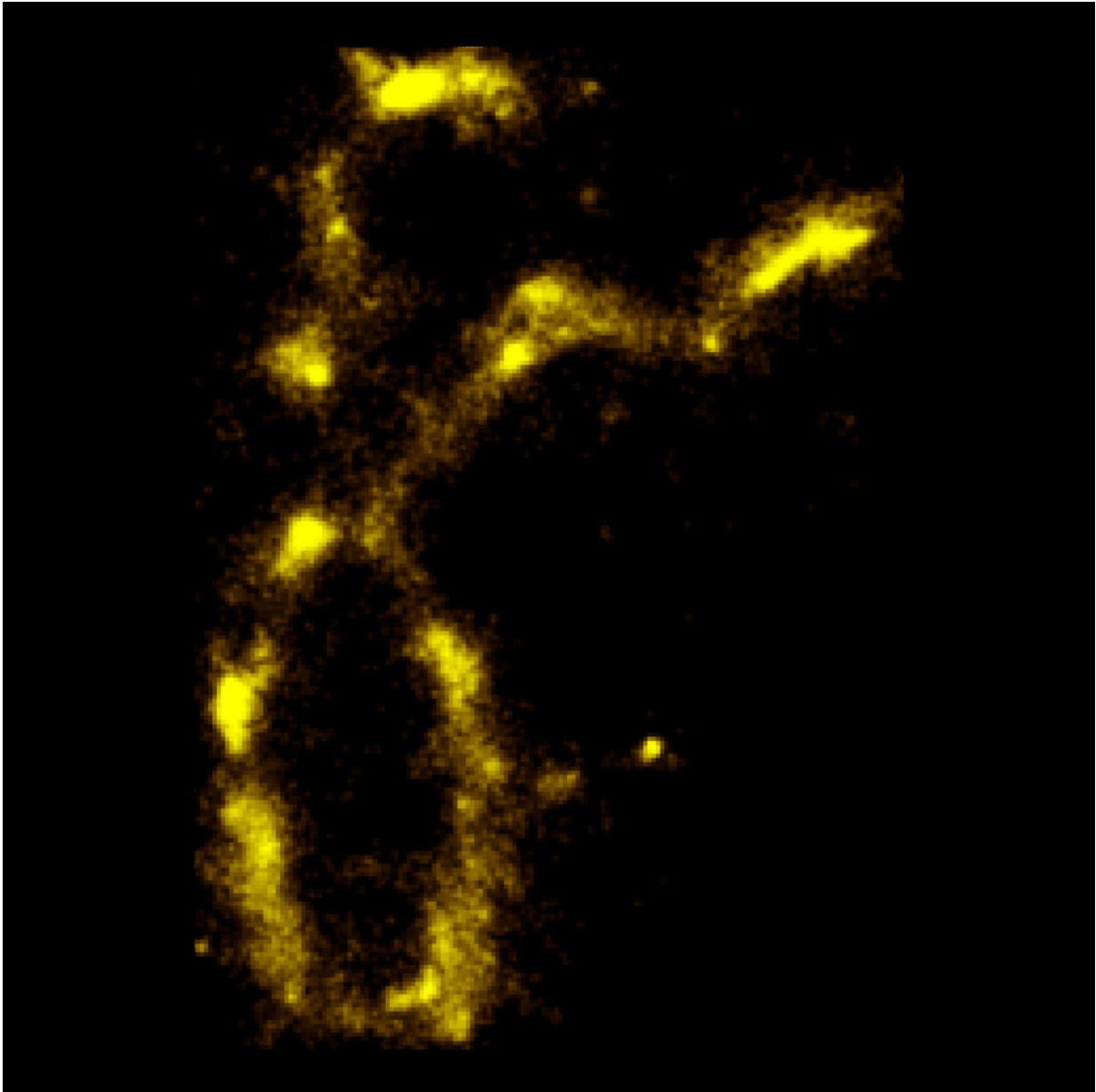


Researchers are first to see DNA 'blink'

February 17 2017



A powerful Northwestern University imaging tool is the first to measure the

structure of isolated chromosomes without the use of fluorescent labels. Credit: Northwestern University

Many of the secrets of cancer and other diseases lie in the cell's nucleus. But getting way down to that level—to see and investigate the important genetic material housed there—requires creative thinking and extremely powerful imaging techniques.

Vadim Backman and Hao Zhang, nanoscale imaging experts at Northwestern University, have developed a new imaging technology that is the first to see DNA "blink," or fluoresce. The tool enables the researchers to study individual biomolecules as well as important global patterns of gene expression, which could yield insights into cancer.

Backman will discuss the tool and its applications—including the new concept of macrogenomics, a technology that aims to regulate the global patterns of gene expression without gene editing—Friday (Feb. 17) at the American Association for the Advancement of Science (AAAS) annual meeting in Boston.

The talk, "Label-Free Super-Resolution Imaging of Chromatin Structure and Dynamics," is part of the symposium "Optical Nanoscale Imaging: Unraveling the Chromatin Structure-Function Relationship," which will be held from 1 to 2:30 p.m. Eastern Time Feb. 17 in Room 206, Hynes Convention Center.

The Northwestern tool features six-nanometer resolution and is the first to break the 10-nanometer resolution threshold. It can image DNA, chromatin and proteins in cells in their native states, without the need for labels.

For decades, textbooks have stated that macromolecules within living cells, such as DNA, RNA and proteins, do not have visible fluorescence on their own.

"People have overlooked this natural effect because they didn't question conventional wisdom," said Backman, the Walter Dill Professor of Biomedical Engineering in the McCormick School of Engineering. "With our super-resolution imaging, we found that DNA and other biomolecules do fluoresce, but only for a very short time. Then they rest for a very long time, in a 'dark' state. The natural fluorescence was beautiful to see."

Backman, Zhang and collaborators now are using the label-free technique to study chromatin—the bundle of [genetic material](#) in the cell nucleus—to see how it is organized. Zhang is an associate professor of biomedical engineering at McCormick.

"Insights into the workings of the chromatin folding code, which regulates patterns of [gene expression](#), will help us better understand cancer and its ability to adapt to changing environments," Backman said. "Cancer is not a single-gene disease."

Current technology for imaging DNA and other genetic material relies on special fluorescent dyes to enhance contrast when macromolecules are imaged. These dyes may perturb cell function, and some eventually kill the cells—undesirable effects in scientific studies.

In contrast, the Northwestern technique, called spectroscopic intrinsic-contrast photon-localization optical nanoscopy (SICLON), allows researchers to study biomolecules in their natural environment, without the need for these fluorescent labels.

Backman, Zhang and Cheng Sun, an associate professor of mechanical

engineering at McCormick, discovered that when illuminated with visible light, the biomolecules get excited and light up well enough to be imaged without fluorescent stains. When excited with the right wavelength, the biomolecules even light up better than they would with the best, most powerful fluorescent labels.

"Our technology will allow us and the broader research community to push the boundaries of nanoscopic imaging and molecular biology even further," Backman said.

Provided by Northwestern University

Citation: Researchers are first to see DNA 'blink' (2017, February 17) retrieved 19 April 2024 from https://phys.org/news/2017-02-dna_1.html

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.