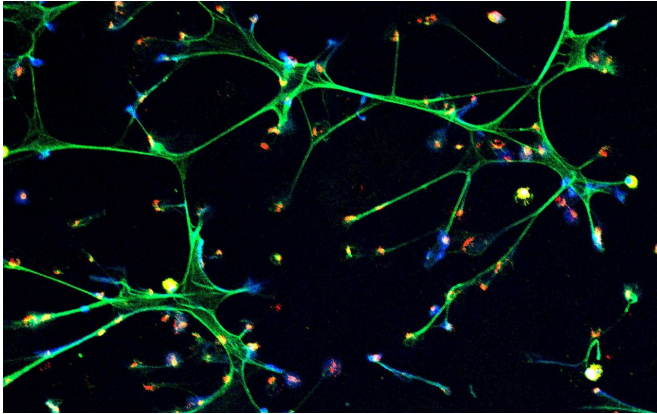


How a sleeping cancer awakens and metastasizes

27 September 2018



When lung tissue is chronically inflamed, dormant cancer cells are able to hijack gauzy "NETs" that spew from nearby white blood cells to re-enter the cell cycle and begin growing again. Dr. Mikala Egeblad and colleagues have figured out how this reawakening can seed metastatic tumors. Credit: Egeblad lab, Cold Spring Harbor Laboratory

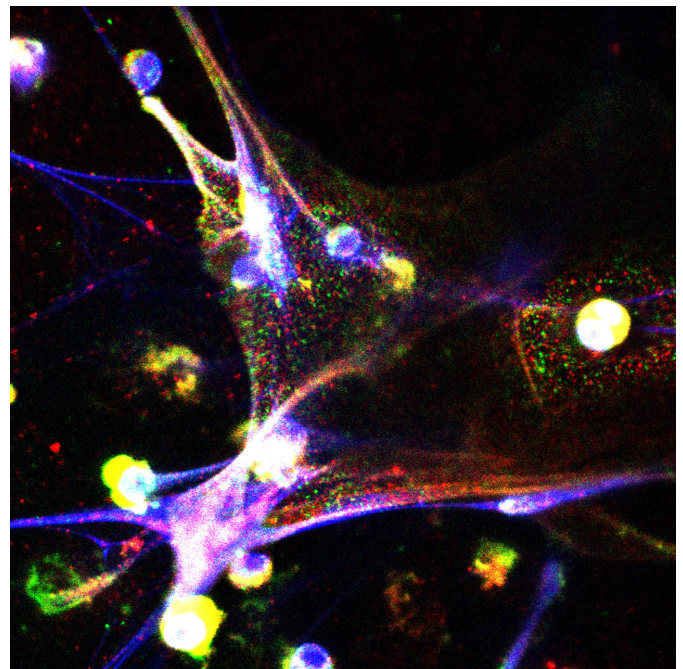
Scientists at Cold Spring Harbor Laboratory (CSHL) have determined one of the ways in which cancers in remission can spring back into action. This knowledge has inspired a new treatment idea designed to prevent cancer recurrence and metastasis.

Even after successful cancer treatment, dormant, non-dividing cancer cells that previously detached from the original tumor may still exist elsewhere in the body. If awakened, these cells can proliferate and grow into metastatic tumors. A CSHL team studying metastasis to the lungs has now identified signals accompanying [inflammation](#) that can awaken dormant cancer cells.

Whether inflammation can directly cause cancer recurrence, and if so how, has not been clear. In their new research, the team demonstrates that sustained [lung](#) inflammation, including that caused

by tobacco smoke exposure, can cause dormant breast and prostate cancer cells that have traveled to the lungs to awaken and begin to divide. These cells can now form a metastasis in the lungs. Metastasis accounts for the bulk of lethality from most common cancers.

Importantly, the team, led by CSHL Associate Professor Mikala Egeblad, and including researchers from the Dana Farber Cancer Institute and UC Davis, also demonstrates a way of blocking the signaling that awakened the dormant cancer cells, a concept that could prevent cancer recurrence or lessen its frequency.

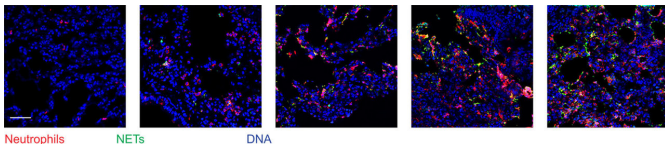


Two enzymes called NE (green dots) and MMP9 (red dots) stud the scaffold of DNA NETs expelled by neutrophils, or white blood cells. These enzymes sequentially cleave a protein called laminin-111 in lung tissue, inducing a signal that causes nearby dormant cancer cells to reawaken and begin proliferating, the seed of a possible metastatic tumor. Credit: Egeblad lab, Cold Spring Harbor Laboratory

Egeblad's team showed that sustained lung inflammation, caused either by exposing mice to tobacco smoke or to a component of bacteria known as endotoxin, induced common white blood cells called neutrophils to awaken nearby dormant cancer cells in an extraordinary way.

Neutrophils, which we normally rely upon to kill invaders like bacteria and yeast, have several ways of vanquishing their prey. One is to expel their DNA into the space beyond the cell membrane. Laced with toxic enzymes, this expelled DNA forms a gauzy, net-like trap (called neutrophil extracellular traps, or NETs) that can kill a pathogen.

The new research shows that sustained lung inflammation causes the formation of NETs in the area around dormant cancer cells. Two enzymes in the NETs, called NE (neutrophil elastase) and MMP9 (matrix metalloproteinase 9), interact with a protein in tissue called laminin. In sequence, first NE then MMP9 make cuts in laminin proteins. This changes the protein's shape, exposing a new surface, called an epitope.



A few scattered white blood cells called neutrophils (red) are seen in normal mouse lung tissue, far left. Four hours after an agent is delivered nasally to promote lung inflammation, the number of neutrophils visibly increases, and web-like DNA "NETs" (green) become visible for the first time. Neutrophils and NETs come to dominate the scene on succeeding days as the inflammatory agent is again delivered. The direct relation of inflammation and NET formation can result in the reawakening of dormant cancer cells if they are in the immediate vicinity--one mechanism of metastasis. Credit: Egeblad lab, Cold Spring Harbor Laboratory

This epitope, when recognized by dormant cancer cells nearby, spurs signaling that awakens the

cancer cells. "The dormant cancer cells recognize that new shape of the laminin and they say, 'we should start growing again,'" Egeblad says.

The team created an antibody to block the epitope exposed on the laminin proteins. In mice, this prevented the reawakening of dormant [cancer cells](#) nearby. Work has begun to optimize the antibody and compare it with other approaches to interfere with NETs, with the hope of eventually conducting trials in people.

Albregues J et al, "Neutrophil extracellular traps produced during inflammation awaken [dormant cancer cells](#) in mice" appears online in *Science*, September 28, 2018.

More information: J. Albregues et al., "Neutrophil extracellular traps produced during inflammation awaken dormant cancer cells in mice," *Science* (2018).

science.sciencemag.org/cgi/doi/10.1126/science.aao4227

"How dormant cancer persists and reawakens," by J.A. Aguirre-Ghiso, *Science* science.sciencemag.org/cgi/doi/10.1126/science.aav0191

Provided by Cold Spring Harbor Laboratory

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