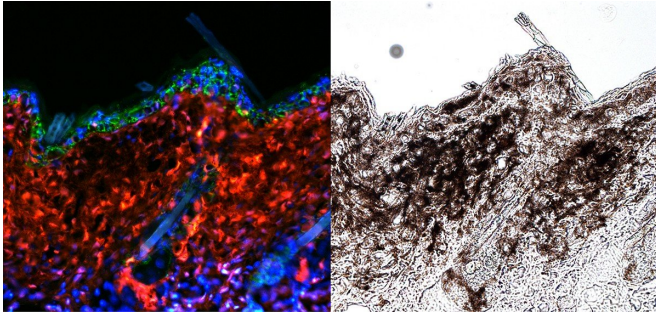


New findings explain how UV rays trigger skin cancer

18 October 2017, by Krishna Ramanujan



Fluorescence microscopy reveals melanoma (red, left; black, right) emerging from melanocyte stem cells. Credit: Hyeongsun Moon and Andrew White, Cornell University

Melanoma, a cancer of skin pigment cells called melanocytes, will strike an estimated 87,110 people in the U.S. in 2017, according to the Centers for Disease Control and Prevention. A fraction of those melanomas come from pre-existing moles, but the majority of them come from sources unknown - until now.

Cornell researchers have discovered that when melanocyte stem cells accumulate a sufficient number of [genetic mutations](#), they can become the cells where these cancers originate. Under normal conditions, ultraviolet (UV) radiation from the sun activates melanocytes to release melanin, a pigment that protects the skin from the sun's rays. But if melanocyte stem cells have surpassed a threshold of genetic [mutations](#), a tumor can start to grow when those [skin stem cells](#) are activated by sun exposure.

"If you had mutations that were sufficient for melanoma, everything would be fine until you went out and got a sunburn," said Andrew White, assistant professor of biomedical sciences at Cornell's College of Veterinary Medicine, and

senior author of a study published this month in the journal *Cell Stem Cell*. Hyeongsun Moon, a postdoctoral researcher in White's lab, is the paper's lead author. "The stimuli that would normally just give you a tanning response could in fact start a [melanoma](#) instead," White said.

The researchers also may have discovered a way to prevent melanomas caused by mutated stem cells. A gene called *Hgma2* was suspected to become expressed in the skin under UV radiation. When expressed, *Hgma2* facilitates melanocyte stem cells to move from the base of skin hair follicles to the skin's surface (the epidermis), where the [cells](#) release melanin. Moon, White and colleagues used [mice](#) engineered with melanocyte stem cell mutations. One set of mice had the mutations, while another set with the mutations had the *Hgma2* gene deleted. They then gave the mice a very low dose of UV radiation, just enough to trigger a tanning response. Mice with tumor-causing mutations and the *Hgma2* gene intact developed melanomas, but the mice with mutations and the deleted gene remained healthy.

More study is needed to better understand the *Hgma2* gene's function.

"We have an actual mechanism, with *Hgma2*, that can be explored in the future and could be a way we can prevent melanomas from happening," White said.

More information: Hyeongsun Moon et al. Melanocyte Stem Cell Activation and Translocation Initiate Cutaneous Melanoma in Response to UV Exposure, *Cell Stem Cell* (2017). [DOI: 10.1016/j.stem.2017.09.001](#)

Provided by Cornell University

APA citation: New findings explain how UV rays trigger skin cancer (2017, October 18) retrieved 4 August 2022 from <https://medicalxpress.com/news/2017-10-uv-rays-trigger-skin-cancer.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.