New drug combos may prevent resistance to melanoma treatments
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Melanoma accounts for about 1% of skin cancer cases, but it causes the vast majority of skin cancer deaths, according to the American Cancer Society. More than half of melanoma patients have cancer-causing mutations in what is known as the BRAF gene. Among those patients, the cancer initially responds well to treatment with vemurafenib, a drug targeting BRAF, but quickly develops resistance, allowing the melanoma to spread.

Misek found that among the resistant melanomas, about half show activation of a signaling pathway involving the protein Rho. Results also showed that combining vemurafenib, a common cancer drug, with other compounds that interfere with Rho signaling, can re-sensitize the cancer cells to treatment.

The study identified one clinically-approved drug and three research compounds that can enhance vemurafenib responses in resistant melanomas. One of the research compounds is being developed by an MSU spinoff company, FibrosIX Inc., with the aim to have it available for eventual clinical use.

Neubig and co-mentor, Kathy Gallo, professor in the Department of Physiology, described Misek as "an extraordinary student and young scientist." This project succeeded due to the "powerful combination of computational and experimental work that Sean was able to bring to the research," Gallo said.

Other MSU researchers involved in the study included Kathryn Appleton, Thomas Dexheimer and Erika Lisabeth, as well as Roger Lo from UCLA and Scott Larsen from the University of Michigan.

"It feels good to make a contribution," Misek said, adding that "We look forward to seeing these results applied to help patients in the future."

Provided by Michigan State University

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