

Modified protein could become first effective treatment for vitiligo

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Mice that have developed vitiligo.

This image shows mice that have developed vitiligo. Credit: Loyola University Chicago Stritch School of Medicine

Loyola University Chicago Stritch School of Medicine researchers have developed a genetically modified protein that dramatically reverses the skin disorder vitiligo in mice, and has similar effects on immune

responses in human skin tissue samples.

The modified protein is potentially the first effective treatment for vitiligo, which causes unsightly white patches on the face, hands and other parts of the body. Loyola University Chicago has submitted a [patent application](#) for the protein, and researchers are seeking regulatory approval and funding for a clinical trial in humans.

I. Caroline Le Poole, PhD, and colleagues describe the modified protein in the journal *Science Translational Medicine*. Le Poole is a professor in Loyola's Oncology Institute and in the departments of Pathology and Microbiology and Immunology.

About 1 million Americans have vitiligo, and the condition affects about 1 in 200 people worldwide. Vitiligo is most noticeable in people of color, but also can be distressing to Caucasians. Vitiligo is an autoimmune disease, in which the immune system goes into overdrive and kills [pigment cells](#), which give skin its color.



Vitiligo mice after vaccination with mutant HSP70 protein.

This image shows mice with vitiligo that have been vaccinated with mutant HSP70 protein. Credit: Loyola University Chicago Stritch School of Medicine.

Previous studies have found that a protein called HSP70i plays a vital role in the [autoimmune response](#) that causes vitiligo. (HSP70i stands for inducible [heat shock protein](#) 70.)

HSP70i consists of 641 building blocks called [amino acids](#). Le Poole and colleagues genetically modified one of these amino acids to create a mutant HSP70i. This [mutant protein](#) supplants normal HSP70i, thereby reversing vitiligo's autoimmune response.

Researchers Jeffrey A. Mosenson and Andrew Zloza gave mutant HSP70i to mice that developed vitiligo, and the results were striking. Mouse fur

– affected by vitiligo—had the coloring of a salt-and-pepper beard. But when the mice were vaccinated with mutant HSP70i, the fur turned black.

"The mice look normal," Le Poole said.

Some of the effects seen in mice also were seen in human skin specimens.

There are no long-term effective treatments for vitiligo. Steroid creams sometimes return some color to affected skin. But this treatment also thins the skin, and can cause streaks or lines. Bright lights, similar to tanning booths, also can return color, but can cause sunburns and other side effects, including vitiligo. Skin grafts transfer skin from unaffected areas to the white patches, but can be painful and expensive. None of the existing treatments effectively prevent vitiligo from progressing.

Le Poole and colleagues wrote that mutant HSP70i "may offer potent treatment opportunities for vitiligo."

Provided by Loyola University Health System

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