

# The hunger games: Uncovering the secret of the hunger switch in the brain

15 April 2021



Credit: CC0 Public Domain

Being constantly hungry, no matter how much you eat, is the daily struggle of people with genetic defects in the brain's appetite controls, and it often ends in severe obesity. In a study published in *Science* on April 15, researchers at the Weizmann Institute of Science, together with colleagues from the Queen Mary University of London and the Hebrew University of Jerusalem, have revealed the mechanism of action of the master switch for hunger in the brain: the melanocortin receptor 4, or MC4 receptor for short. They have also clarified how this switch is activated by setmelanotide (Imcivree), a drug recently approved for the treatment of severe obesity caused by certain genetic changes. These findings shed new light on the way hunger is regulated and may help develop improved anti-obesity medications.

The MC4 receptor is present in a brain region called the hypothalamus—within a cluster of neurons that compute the body's energy balance by processing a variety of energy-related metabolic signals. When the MC4 is activated, or "on"—as it normally is—it sends out commands that cause us to feel full, which means that from the brain's perspective, our default state is satiety. When our

energy levels drop, the hypothalamic cluster produces a "time to eat" hormone that inactivates, or turns off the MC4 receptor, sending out a "become hungry" signal. After we eat, a second, "I'm full" hormone is released. It binds to the same active site on the MC4, replacing the hunger hormone and turning the receptor back on—bringing us back to the satiety default. Mutations that inactivate the MC4 cause people to feel constantly hungry.

MC4 is a prime target for anti-obesity drugs, such as setmelanotide, precisely because it's a master switch: turning it on can control hunger while bypassing all other energy-related signals. But until now it was unknown how exactly this hunger switch works.

The new study began with the predicament of one family, in which at least eight members, plagued by persistent hunger, were severely obese—most of them with a body mass index of over 70, that is, about triple the norm. Their [medical history](#) came to the attention of Hadar Israeli, a [medical student](#) pursuing Ph.D. studies into the mechanisms of obesity under the guidance of Dr. Danny Ben-Zvi at the Hebrew University of Jerusalem. Israeli was struck by the fact that the family's plight was due to a single mutation that ran in the family: one affecting the MC4 receptor. She turned to Dr. Moran Shalev-Benami of Weizmann's Chemical and Structural Biology Department, asking whether new advances in electron microscopy could help explain how this particular mutation could produce such a devastating effect.

Shalev-Benami launched a study into the structure of MC4, inviting Israeli to join her lab as a visiting scientist. Together with Dr. Oksana Degtjarik, a postdoctoral fellow in the lab, Israeli isolated large quantities of pure MC4 receptor from cell membranes, let it bind with setmelanotide and determined its 3D structure using cryogenic electron microscopy. The study was conducted in

collaboration with the teams of Dr. Peter J. McCormick from the Queen Mary University of London and of Prof. Masha Y. Niv from the Hebrew University of Jerusalem.

[science.sciencemag.org/lookup/...  
1126/science.abf7958](https://science.sciencemag.org/lookup/doi/10.1126/science.abf7958)

The 3D structure revealed that setmelanotide activates the MC4 receptor by entering its binding pocket—that is, by directly hitting the molecular switch that signals satiety, even more potently than the natural satiety hormone. It also turned out that the drug has a surprising helper: an ion of calcium that enters the pocket, enhancing the drug's binding to the receptor. In biochemical and computational experiments, the scientists found that similarly to the drug, calcium also assists the natural satiety hormone.

Provided by Weizmann Institute of Science

McCormick: "Calcium helped the satiety hormone activate the MC4 receptor while interfering with the hunger hormone and reducing its activity."

"This was a truly unexpected finding," Shalev-Benami says. "Apparently, the satiety signal can successfully compete with the hunger signal because it benefits from the assistance of calcium, which helps the brain restore the 'I'm full' sensation after we eat."

MC4's structure also revealed that the drug's entry causes structural changes in the receptor; these changes appear to initiate the signals within the neurons that lead to the sensation of fullness. The study has explained how mutations in the MC4 receptor can interfere with this signaling, leading to never-ending [hunger](#) and ultimately obesity.

Moreover, the scientists have identified hotspots that crucially distinguish MC4 from similar receptors in the same family. This should make it possible to design drugs that will bind only to MC4, avoiding side effects that may be caused by interactions with other [receptors](#).

"Our findings can help develop improved and safer anti-obesity drugs that will target MC4R with greater precision," Shalev-Benami says.

**More information:** "Structure Reveals the Activation Mechanism of the MC4 Receptor to Initiate Satiety Signaling" *Science* (2021).

APA citation: The hunger games: Uncovering the secret of the hunger switch in the brain (2021, April 15)  
retrieved 22 April 2021 from  
<https://medicalxpress.com/news/2021-04-hunger-games-uncovering-secret-brain.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.*