

Little-studied immune cell may be key to activating energy-burning 'beige fat'

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Immune cells called eosinophils could be an essential part of the process that activates "beige fat," a study in mice published this week in *Nature Communications* shows.



If future research confirms this hypothesis, these cells may be a target for anti-obesity therapies down the track.

There are three types of fat, or <u>adipose tissue</u>, in the human body.

"Put simply, white fat—what we generally think of when we think of fat—stores excess energy," says the study's lead author Dr. Kate Quinlan.

"Brown fat, on the other hand, can burn off energy, generating heat in a process called thermogenesis. This phenomenon is particularly important in babies and hibernating animals and, more recently, has also been shown to be present in adult humans.

"Beige fat is a newly discovered type of energy-burning fat that can emerge within white adipose tissue to perform thermogenesis instead of energy storage."

In this paper, the team studied a mutant mouse model which is resistant to obesity. They found it had increased <u>beige fat</u> activation.

"Naturally, we were curious as to why," says Dr. Quinlan. "We also noted that this mutant mouse had more <u>immune cells</u> called eosinophils, and wondered if that was the answer."

In one key experiment, the authors transplanted immune cells from mutant mice into normal mice. The recipient mice ended up being protected from obesity when put on a high-fat, high-sugar 'western' diet. This suggested that immune cells, which are known to reside in and signal to fat, are involved in making the mutant mice beige and obesity-resistant.

"Although many different immune cells could be important, we are



working on the hypothesis that the regulation of metabolism involves eosinophils," says Dr. Quinlan.

Eosinophils are a rare immune cell type important in the immune response to parasitic infection, adversely involved in asthma, Crohn's disease and ulcerative colitis but thought to be beneficial in fat thermogenesis.

"In the lean, beige mouse there were more eosinophils in fat tissue and these eosinophils had unique gene expression profiles," Dr. Quinlan says.

"We also found evidence that eosinophils secrete signaling molecules known to play a role in beige fat activation, providing a possible link between these enigmatic immune <u>cells</u> and their beneficial function in adipose tissue."

Obesity is a worldwide health epidemic. In 2016, more than 1.9 billion adults were overweight, with over 650 million being classified as obese.

"This is a serious problem since obesity is associated with increased risk of chronic conditions, such as diabetes and heart disease," Dr. Quinlan says.

"Turning energy-storing white fat into energy-burning beige fat is an attractive therapeutic strategy for obesity but currently the mechanisms that drive this process within the body are not well understood," Dr. Quinlan says.

"While we need to do more research to further validate our hypothesis, this work suggests that eosinophils may play a larger role in beige fat activation than was previously thought."

Dr. Quinlan and her research team are now focused on uncovering new



factors that eosinophils secrete to stimulate beige fat activation. Such factors could form the basis of future therapeutics to activate beige fat and work towards reversing <u>obesity</u>.

More information: Alexander J. Knights et al. Eosinophil function in adipose tissue is regulated by Krüppel-like factor 3 (KLF3), *Nature Communications* (2020). DOI: 10.1038/s41467-020-16758-9

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