

Targeting fat-tissue hormone may lead to type 2 diabetes treatment

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A fatty acid-binding protein called aP2 has been linked to obesity, diabetes, and heart disease in humans. New research in mice shows that an antibody therapy blocking aP2 can help reverse some of the metabolic disturbances associated with type 2 diabetes. Credit: V. Altounian / Science Translational Medicine

A new study by researchers from Harvard T.H. Chan School of Public Health and colleagues describes the pre-clinical development of a therapeutic that could potentially be used to treat type 2 diabetes, fatty liver disease, and other metabolic diseases. The researchers developed an antibody that improves glucose regulation and reduces fatty liver in obese mice by targeting a hormone in adipose (fat) tissue called aP2 (also known as FABP4).

The study will be published online December 23, 2015 in *Science Translational Medicine*.

"The importance of this study is two-fold: first, demonstrating the importance of aP2 as a critical hormone in abnormal glucose metabolism, and secondly, showing that aP2 can be effectively targeted to treat diabetes and potentially other immunometabolic diseases," said Gökhan S. Hotamisligil, J.S. Simmons Professor of Genetics and Metabolism and chair of the Department of Genetics and Complex Diseases and the Sabri Ülker Center at Harvard Chan School.

The work is the product of a collaboration on immunometabolism between the biopharmaceutical company UCB and a team of researchers led by Hotamisligil and lead author M. Furkan Burak, a former Hotamisligil lab member and currently a resident in internal medicine at Mount Auburn Hospital, Cambridge, MA. This partnership successfully twins UCB's world-class expertise in monoclonal antibody discovery with Hotamisligil's insight and experience in aP2 biology.

The increase in adipose tissue characteristic of obesity has long been linked to increased risk for [metabolic diseases](#) such as type 2 diabetes and cardiovascular disease. Recently, it has become clear that the tissue itself plays an active role in metabolic disease, in part by releasing hormones which act in distant sites such as the liver, muscle, and brain that affect systemic metabolism. Work from the Hotamisligil lab

previously identified the protein aP2 as a critical hormone mediating communication between adipose tissue and liver. Since aP2 levels are significantly increased in humans with obesity, diabetes, and atherosclerosis, and mutations that reduce aP2 result in significantly reduced risk of diabetes, dyslipidemia, and heart disease, strategies to modify aP2 function carry promise as new lines of therapeutic entities against these common and debilitating chronic diseases.

In the new study, Burak and colleagues describe the development and evaluation of novel monoclonal antibodies targeting aP2. The team found that one of these antibodies effectively improved [glucose regulation](#) in two independent models of obesity. Additionally, beneficial reductions in liver fat were observed.

These monoclonal antibodies have the potential to be transformative first-in-class therapeutics to fight obesity-related metabolic and immunometabolic disease, say the authors. This work is still at the preclinical stage and will require extensive evaluation for safety and effectiveness before being considered for use in humans.

More information: "Development of a therapeutic monoclonal antibody that targets secreted fatty acid binding protein aP2 to treat type 2 diabetes," M. Furkan Burak, Karen E. Inouye, Ariel White, Alexandra Lee, Gurol Tuncman, Ediz S. Calay, Motohiro Sekiya, Amir Tirosh, Kosei Eguchi, Gabriel Birrane, Daniel Lightwood, Louise Howells, Geoffrey Odede, Hanna Hailu, Shauna West, Rachel Garlish, Helen Neale, Carl Doyle, Adrian Moore, Gökhan S. Hotamisligil, *Science Translational Medicine*, online December 23, 2015.
[stm.sciencemag.org/lookup/doi/ ... scitranslmed.aac6336](http://stm.sciencemag.org/lookup/doi/.../scitranslmed.aac6336)

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