

The processes for anabolic and catabolic activities that regulate the maintenance of DNA repair for genes are outlined in the following:

<http://www.mcfip.net/upload/Epigenetics%20-%20DNA%20Repair%20-%20Copy%20Errors.pdf>

Anabolic activity encompassing the nine epigenetic signaling molecules can be designated as NUP98, FOXA9 or cGAS. In addition to PDCK9, the following provides designation(s) that can be used to identify the catabolic activity; i.e. autophagy.



Having identified the near certain elemental constituents of the SOD1 and SOD3 based epigenetic signaling molecules for autophagy, when PAH-1 was mentioned in the following article relative to fat (lipid) metabolism, we were able to correlate it with our findings.

Further review identified the existence of two other forms of PAH; i.e. PAH2 and PAH3.

Our bioidentic modeling identified PAD1 - 3 as being designations for the three “enzymes” (epigenetic signaling molecules that function to perform autophagy).

Details of the amino acids that are formed when IL-18 is subjected to catabolic activity as described below can be shared with interested parties as part of DNA repair activities.

How the signaling molecules are formed:

<http://www.mcfip.net/upload/Cell%20Surface%20Signaling%20Molecule%20Formation%207-2017.pdf>

How they function for DNA repair:

<http://www.mcfip.net/upload/Epigenetics%20-%20DNA%20Repair%20-%20Copy%20Errors.pdf>

# A fat-regulating enzyme could hold the key to obesity, diabetes, cancer, other diseases

September 18, 2017

It had already been known that the enzyme known as phosphatidic acid phosphatase plays a crucial role in regulating the amount of fat in the human body. Controlling it is therefore of interest in the fight against obesity.

But scientists at Rutgers University-New Brunswick have now found that getting rid of the enzyme entirely can increase the risk of cancer, inflammation and other ills. Their findings were published online in the *Journal of Biological Chemistry* last month.

"The goal of our lab is to understand how we can tweak and control this enzyme," said George M. Carman, Board of Governors professor in the Department of Food Science in the School of Environmental and Biological Sciences. "For years, we have been trying to find out how to fine-tune the enzyme's activity so it's not too active, and creating too much fat, but it's active enough to keep the body healthy."

The enzyme was discovered in 1957 and Gil-Soo Han, research assistant professor in the Rutgers Center for Lipid Research, discovered the gene encoding the enzyme in 2006. The enzyme determines whether the body's phosphatidic acid will be used to create fat, or create the lipids in cell membranes.

The study used baker's yeast as a model organism, since it also contains the key enzyme. Han, study lead author, deleted a gene in yeast to eliminate the enzyme. That led to accumulations of phosphatidic acid, with cells making far more membrane lipids than necessary, said Carman, who founded the center in Rutgers' New Jersey Institute for Food, Nutrition, and Health a decade ago.

"We have found that maybe a more critical role for the enzyme is to make sure that cells are not making too much membrane lipid," Carman said. "If you make too much membrane lipid, you make too much membrane and the cells are permitted to grow uncontrollably, a condition characteristic of cancer."

Since the discovery of the gene encoding the enzyme, people worldwide have studied the enzyme because of its relation to obesity, lipodystrophy, inflammation, diabetes and other conditions, Carman said.

Lately, the Rutgers scientists have been trying to understand the enzyme's structure and function. The next step is to figure out how to control it, Carman said.

"The key take-home message is that things have to be balanced," he said. "To keep the balance between making storage fat and membrane lipid, you have to have balanced diet."

**Explore further: Researchers find fat gene**

**More information:** Gil-Soo Han et al, Yeast PAH1 -encoded phosphatidate phosphatase controls the expression of CHO1 -encoded phosphatidylserine synthase for membrane phospholipid synthesis, *Journal of Biological Chemistry* (2017). **DOI:** **10.1074/jbc.M117.801720**

**Journal reference:** Journal of Biological Chemistry

Read more at: <https://phys.org/news/2017-09-fat-regulating-enzyme-key-obesity-diabetes.html#jCp>