

# Bacteria that prevent type 1 diabetes

August 6 2015

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Our bodies have ten times more microbes than human cells. This set of bacteria is called microbiota. In some instances, bacteria known as pathogens can cause infectious diseases. However, these micro-organisms can also protect us from certain diseases. Researchers from Inserm, Paris Descartes University and the CNRS (French National Centre for Scientific Research), in collaboration with teams from China and Sweden, have recently shown how microbiota protects against the development of type 1 diabetes in mice. This research is published in the *Immunity* journal.

To combat pathogens, the immune system has developed various mechanisms to detect, fight and even destroy micro-organisms that are harmful to the body. This includes [antimicrobial peptides](#) and natural proteins that destroy pathogenic bacteria by disrupting their cellular membrane. Not only are they produced by immune cells, they are also produced by cells whose functions are not immune-related.

A research team coordinated by Julien Diana, an Inserm Research Fellow at Inserm Unit 1151 "Institut Necker-Enfant Malades" [Necker Institute for Sick Children] (Inserm/CNRS/Université Paris Descartes), is focussing on a category of antimicrobial peptides, i.e. cathelicidins. Apart from their protective function, these peptides have also exhibited immunoregulatory abilities against several autoimmune diseases. As such, scientists hypothesise that cathelicidins may be involved in the control of [type 1 diabetes](#), an autoimmune disease where certain cells in the immune system attack beta cells in the pancreas which secrete insulin.

Firstly, they observed that beta pancreatic cells in healthy mice produce cathelicidins and that, interestingly, this production is impaired in [diabetic mice](#).

To test this hypothesis, they injected diabetic mice with cathelicidins where production was defective.

"Injecting cathelicidins inhibits the development of pancreatic inflammation and, as such, suppresses the development of autoimmune diabetes in these mice" states Julien Diana.

Given that the production of cathelicidins is controlled by short-chain fatty acids produced by gut bacteria, Julien Diana's team are studying the possibility that this may be the cause of the cathelicidin deficiency associated with diabetes. Indeed, researchers have observed that diabetic mice have a lower level of short-chain fatty acids than that found in healthy mice.

By transferring part of the [gut bacteria](#) from healthy to diabetic mice, they are re-establishing a normal level of cathelicidin. Meanwhile, the transfer of micro-organisms reduces the occurrence of diabetes.

For the authors, "this research is further evidence of the undeniable role microbiota plays in [autoimmune diseases](#), particularly in controlling the development of autoimmune diabetes".

Preliminary data, as well as scientific literature, suggest that a similar mechanism may exist in humans, paving the way for new therapies against [autoimmune diabetes](#).

**More information:** Pancreatic beta-cells limit autoimmune diabetes via an immunoregulatory antimicrobial peptide expressed under the influence of the gut microbiota *Immunity* ; 04 August 2015.

Provided by CNRS

Citation: Bacteria that prevent type 1 diabetes (2015, August 6) retrieved 4 May 2024 from <https://medicalxpress.com/news/2015-08-bacteria-diabetes.html>

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