

Researchers Discover How Virulent Bacteria Sabotage Immune Response Against It

February 2 2010

Researchers at National Jewish Health have discovered how the virulent food-borne bacteria *Listeria monocytogenes* induces infected immune cells to sabotage their own defensive response. The studies offer insight into host-pathogen interactions and suggest potential therapeutic targets for food poisoning, tuberculosis and autoimmune diseases.

In the Feb. 15 issue of the [Journal of Experimental Medicine](#), Laurel Lenz, PhD, and his colleagues report that macrophages infected by the bacteria *Listeria* release interferon- $\alpha\beta$ (IFN- $\alpha\beta$), which makes them and nearby [immune cells](#) unresponsive to activation signals. This reduces immune resistance to the bacteria, which causes thousands of cases of food poisoning -- and more than 500 deaths -- each year in the United States.

"*Listeria* appears to benefit by triggering an endogenous pathway of the host that dampens its own immune response," said Dr. Lenz. "Our findings suggest that *Listeria* increases its survival in infected individuals by inducing cross-talk between host interferon signaling pathways."

When patrolling immune-system cells encounter non-pathogenic microbes, they normally engulf and destroy them. However, certain pathogens such as *Listeria* can grow within immune cells, which then release alarm signals to other nearby cells. One of these alarms is IFN- $\alpha\beta$. IFN- $\alpha\beta$ protects host cells from viral infection. However, IFN- $\alpha\beta$ also increases growth of *Listeria* and certain other bacteria.

Dr. Lenz and his colleagues showed that IFN- $\alpha\beta$ does this by down-regulating expression of receptors for interferon- γ (IFN- γ). With its receptors down-regulated, IFN- γ cannot drive resting [macrophages](#) into an activated state that is especially effective against bacterial pathogens inside the cell.

"IFN- $\alpha\beta$ acts as a sort of anesthetic to numb the response of immune cells to IFN- γ ," said Dr. Lenz.

The research highlights the crosstalk that exists between the antibacterial and anti-viral arms of the immune response. Dr. Lenz's findings demonstrate that IFN- $\alpha\beta$, well known to stimulate antiviral defenses, dampens anti-bacterial activity as well.

He speculates that this may be a way for the immune system to more efficiently defend against viral pathogens, while avoiding collateral damage caused by an overactive immune cells. In fact, interferon- β , a medication widely used for multiple sclerosis, may do just that. Dr. Lenz speculates that this medication may act in part by down-regulating expression of IFN- γ receptors on myeloid cells, thus reducing the stimulation of autoimmune T cells.

The next step will be to define more precisely how IFN- $\alpha\beta$ mediates down-regulation of the IFN- γ receptors, and to determine whether prevention of these effects improves resistance to infection by *Listeria* and other bacterial pathogens.

Provided by National Jewish Medical and Research Center

Citation: Researchers Discover How Virulent Bacteria Sabotage Immune Response Against It (2010, February 2) retrieved 6 May 2024 from <https://medicalxpress.com/news/2010-02-virulent-bacteria-sabotage-immune-response.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.