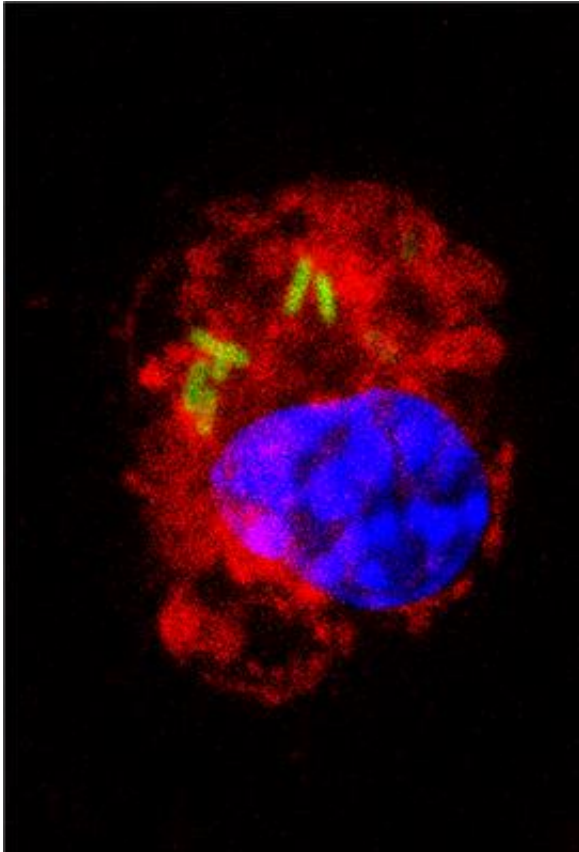


A scientific team describes a new molecular mechanism to fight bacterial infections

2 February 2017



Credit: Universidad de Barcelona

An international team led by the University of Barcelona has identified a new antibacterial mechanism that protects macrophages from infection by the bacteria *Salmonella enterica* serovar Typhimurium, a pathogen associated with several gastrointestinal diseases. This discovery, carried out with mice and published in the journal *Cell Reports*, is led by Professor Annabel Valledor, from the Faculty of Biology of the University of Barcelona, and could open new exploration channels for pharmacological treatments of some bacterial infections.

Macrophages and paradoxes of the immune

response

Macrophages are specialized cells in the immune system with diverse effector functions against pathogens and invasive agents. However, although it seems paradoxical, they become the preferential biological niche in some [bacterial strains](#) to replicate and spread around the body.

This is the case, for instance, with *Salmonella enterica* serovar Typhimurium, an enteric bacterium that causes mild gastroenteritis and severe systemic infections in humans. This strain, in particular, is able to survive and replicate inside the host macrophages as a previous step to spread to other parts of the body.

S. Typhimurium avoids the bactericidal response so that after adhering to the surface of the macrophage, it secretes factors that modify the F-actin cytoskeleton of the host cell. This mechanism allows the bacteria to enter the cell through special phagocytic vacuoles (containing *Salmonella*). Once inside the cell, the bacterium transforms the vacuole's conditions to turn it into a favourable environment to replicate and spread around the infected organism.

In the new study, the scientific team describes a mechanism that limits the capacity of *S. Typhimurium* to infect cells and spread in other tissues. "Our study reveals that pharmacological activation of transcription factors from the family of [nuclear receptors](#)—Liver X receptors or LXR—limits the bacterial infection in macrophages," says Professor Annabel Valledor, who leads the Group Nuclear receptors in Cancer, Metabolism and Inflammation of Nurcainein network.

Nuclear receptors are transcription factors that shape physiological processes through the regulation of the gene expression of several targeted genes. LXR, in particular, can be activated by agonists (molecule activators) and play an

important role in metabolic functions and the regulation of the [immune response](#).

Provided by University of Barcelona

In this case, LXRs regulate the transcriptional activation of the multifunctional enzyme CD38, distributed in cells in the immune system, and which has a potent NAD⁺ activity, which is a metabolite associated with the energy cell mechanism.

"Through this mechanism, the treatment with LXRs agonists is able to lower the intracellular levels of NAD⁺, a process which interferes in a series of morphological and distribution changes in the macrophage's F-actin cytoskeleton and is used for their own benefit in certain invasive bacteria."

The LXRs pharmacological activation, thus, inhibits the *S. Typhimurium* capacity to infect cells and get to other parts of the body, and contributes to improve the clinical symptomatology in laboratory animals infected by this enteric bacterium.

Multiresistant bacteria

According to the World Health Organization (WHO), resistance to antibiotics is one of the biggest threats to world health, food safety, and development. Bacterial multiresistance is speeding up due the excessive and incorrect use of drugs. As a consequence, there are more infections that don't respond to common treatments as antibiotics lose their efficacy.

"Facing the dangers that involve antibiotic-resistant bacterial strains (more common every day), this new research study established the path of LXRs as a new mechanism in the regulation of the host-pathogen interaction which could be pharmacologically exploited as a potential strategy of 'host-directed therapy' in certain bacterial infections," concludes Professor Annabel Valledor

More information: Jonathan Matalonga et al. The Nuclear Receptor LXR Limits Bacterial Infection of Host Macrophages through a Mechanism that Impacts Cellular NAD Metabolism, *Cell Reports* (2017). [DOI: 10.1016/j.celrep.2017.01.007](https://doi.org/10.1016/j.celrep.2017.01.007)

APA citation: A scientific team describes a new molecular mechanism to fight bacterial infections (2017, February 2) retrieved 22 October 2019 from <https://phys.org/news/2017-02-scientific-team-molecular-mechanism-bacterial.html>

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