

Rheumatoid arthritic pain could be caused by antibodies

14 June 2019



Credit: CC0 Public Domain

Antibodies that exist in the joints before the onset of rheumatoid arthritis can cause pain even in the absence of arthritis, researchers from Karolinska Institutet in Sweden report. The researchers believe that the finding, which is published in the *Journal of Experimental Medicine*, can represent a general mechanism in autoimmunity and that the results can facilitate the development of new ways of reducing non-inflammatory pain caused by rheumatoid arthritis and other autoimmune diseases.

"We all know that inflammation is painful," says Camilla Svensson, professor at the Department of Physiology and Pharmacology, Karolinska Institutet. "But pain can appear before any sign of inflammation in the joints and can remain a problem after it has healed. Our aim was to find possible mechanisms to explain that."

Rheumatoid arthritis is an autoimmune disease that occurs when [immune cells](#) attack the cartilage and bone of the joints. The disease affects roughly one percent of the Swedish population.

A common early symptom is [joint pain](#), but even before that, the body has started to produce immune antibodies against proteins in the joint. Researchers at Karolinska Institutet have now studied how these autoantibodies can generate pain.

After injecting cartilage-binding autoantibodies into mice, which served as a model for human rheumatoid arthritis, the researchers found that the mice became more sensitive to pain even before they could observe any signs of inflammation in the joints. Antibodies that had been designed not to activate immune cells and trigger inflammation also induced pain-like behavior in the mice, suggesting increased pain sensitivity in the joints.

The researchers found that the antibodies that caused the behavioral change form so called immune complexes, comprising clusters of antibodies and cartilage proteins in the joints. These complexes activate pain cells via so-called Fc-gamma receptors, which the researchers discovered were present on [pain neurons](#) in the tissue.

When they cultivated pain neurons from the mice, the researchers found that the cells were activated when coming into contact with the antibody complexes. The process was dependent upon the Fc-gamma receptors on the neurons but not on the presence of immune cells. Antibodies in complex can thus act as pain-generating molecules in themselves, independently of the activity of the immune cells, as Camilla Svensson, one of the study's two corresponding authors explains:

"Antibodies in these immune complexes can activate the pain neurons directly, and not, as previously thought, as a result of the destructive joint inflammation," she says. "The antibodies can affect the pain neurons also in conditions without any distinct tissue damage or inflammation."

Although the study was conducted in mice, the researchers show that human pain neurons also have antibody receptors that are functionally similar to those they found on the mouse pain neurons, which leads them to believe that their findings are also relevant to humans.

The results can explain the early pain symptoms in rheumatoid arthritis patients. However, joint and muscle pain are also common symptoms of other autoimmune diseases, and since this newly discovered mechanism operates through the constant part—the shaft—of the autoantibody, the researchers believe that it can explain non-[inflammatory pain](#) caused by other autoimmune diseases too.

"We think that this can be a general pain mechanism in effectively all autoimmune diseases in which these kinds of immune complex form locally in tissue," says Professor Svensson.

More detailed study of what happens in the nerve cell when the antibody complex binds to the receptor could also lead to new targets for reducing the neuronal activity.

"By learning more about the molecular mechanisms of antibody-mediated pain we hope to lay the groundwork for a new way of reducing [pain](#) caused by rheumatoid arthritis and other [autoimmune diseases](#)," says Rikard Holmdahl, professor at the Department of Medical Biochemistry and Biophysics, Karolinska Institutet, and the study's other corresponding author.

The study was financed with grants from the Swedish Research Council, the Swedish Foundation for Strategic Research, the Knut and Alice Wallenberg Foundation, the Ragnar Söderberg Foundation, the Torsten Söderberg Foundation, the Åke Wiberg Foundation, the Alfred Österlund Foundation, the Krapperup Foundation, the King Gustaf V 80-year Foundation, the Swedish Rheumatism Association, Hansa Medical AB, the Royal Physiographic Society, Karolinska Institutet's funds, the Canadian Institutes of Health Research and the Guangdong province of China. The financiers have had no influence on the paper or the decision to publish it.

One of the researchers holds a patent on the use of a type of antibody used in the study and a royalty agreement with Hansa Medical, which also holds a patent on a particular application of the [antibodies](#).

More information: Alex Bersellini Farinotti et al. Cartilage-binding antibodies induce pain through immune complex-mediated activation of neurons, *The Journal of Experimental Medicine* (2019). [DOI: 10.1084/jem.20181657](#)

Provided by Karolinska Institutet

APA citation: Rheumatoid arthritic pain could be caused by antibodies (2019, June 14) retrieved 2 May 2021 from <https://medicalxpress.com/news/2019-06-rheumatoid-arthritic-pain-antibodies.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.