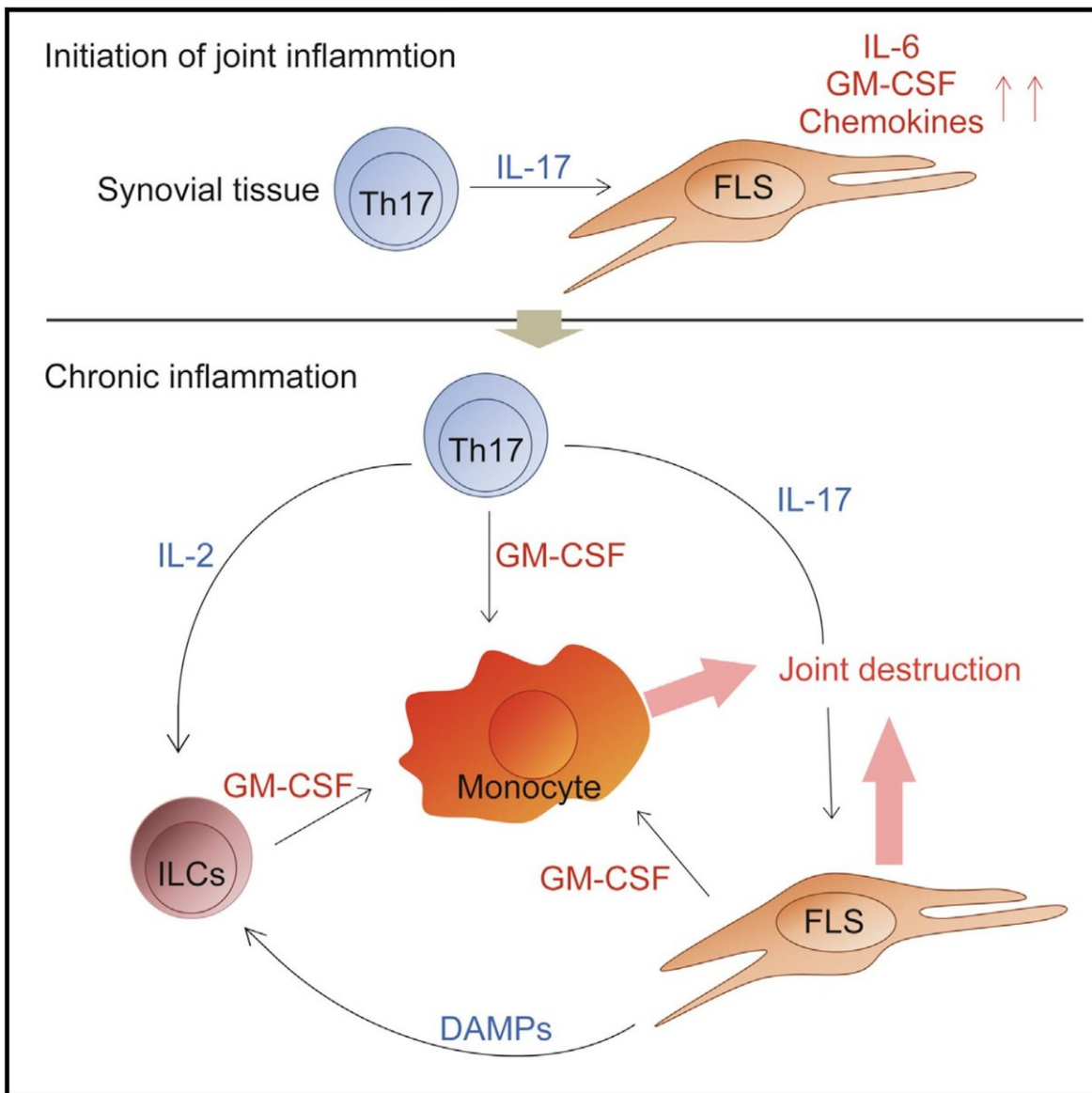


# Igniting the rheumatoid arthritis flame through a cellular cascade

May 30 2018



An inflammatory cellular cascade instigated by an arthritogenic T helper subset and enhanced by GM-CSF-producing synovial-resident innate lymphoid cells.  
Credit: Osaka University

Chronic inflammatory disorders, including autoimmune diseases such as rheumatoid arthritis, involve the action of various inflammatory molecules (cytokines) produced by cells of the immune system. One such cytokine, IL-17, is produced by Th17 cells, which are white blood cells that provide host defense against pathogens, as well as mediating inflammatory reactions. Although Th17 cells are lead players in autoimmune disease, the way in which they control other inflammatory cells had been unclear.

Now, an international team of researchers led by Osaka University has used a mouse model of rheumatoid arthritis to investigate how Th17 [cells](#) interact with other cells at the site of inflammation and influence cytokine production. They reported their findings in the journal *Immunity*.

Several inflammatory cytokines, including IL-6 and TNF $\alpha$ , cause joint inflammation in rheumatoid arthritis. The disease progresses through the interaction of T cells of the immune system with specialized cells in the joint lining (synovium), which promotes the destruction of bone and cartilage. The recent development of new anti-rheumatic drugs such as TNF and IL-6 inhibitors has dramatically improved the quality of life of many patients with rheumatoid arthritis, but treatment for 30% of these patients remains a challenge. Therefore, novel therapies are eagerly awaited.

The inflammatory cytokine GM-CSF activates cells involved in innate and adaptive immune systems, which represent non-specific defense

mechanisms and specific learned responses to foreign substances, respectively. GM-CSF particularly triggers scavenger [white blood cells](#) (macrophages) and immune cell messengers known as [dendritic cells](#).

The team found that GM-CSF was crucial for the development of arthritis in the mouse model. "GM-CSF produced both by stromal cells of the connective tissue and T cells contributed to joint inflammation in the mice, but only stromal cell-derived GM-CSF was needed to initiate arthritis," study first author Keiji Hirota explains. "We also showed that stromal cells secreted GM-CSF in response to stimulation by IL-17 from inflammatory Th17 cells."

GM-CSF was also found to be secreted by a group of innate immune cells, which expanded in number within inflamed joints in response to IL-17 production by Th17 cells and other inflammatory cytokines. This contributed to the maintenance and development of [rheumatoid arthritis](#) in the mice.

"Our findings outline an inflammatory network controlled by autoimmune Th17 cells and involving [stromal cells](#) and innate immune cells, which leads to the onset and development of autoimmune arthritis," corresponding author Shimon Sakaguchi says. "By selectively removing GM-CSF-producing cells from the mouse synovium, we significantly reduced the severity of arthritis. This suggests the usefulness of developing such a novel immunotherapeutic approach that targets the cellular network to reduce chronic joint inflammation."

**More information:** Keiji Hirota et al, Autoimmune Th17 Cells Induced Synovial Stromal and Innate Lymphoid Cell Secretion of the Cytokine GM-CSF to Initiate and Augment Autoimmune Arthritis, *Immunity* (2018). [DOI: 10.1016/j.immuni.2018.04.009](https://doi.org/10.1016/j.immuni.2018.04.009)

Provided by Osaka University

Citation: Igniting the rheumatoid arthritis flame through a cellular cascade (2018, May 30)  
retrieved 23 April 2024 from

<https://medicalxpress.com/news/2018-05-igniting-rheumatoid-arthritis-flame-cellular.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.