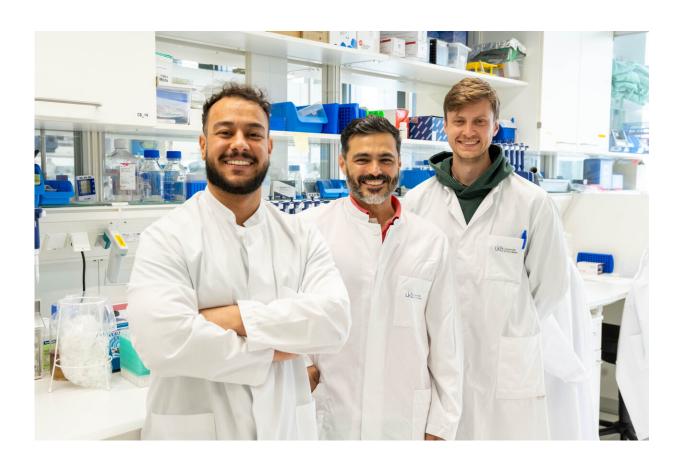


New pathway in immune defense discovered: Interaction of monocytes and platelets decoded

July 8 2024, by Inka Väth



(from left) Dr. Ibrahim Hawwari, Prof. Bernardo Franklin and Lukas Roßnagel discover new intercellular communication mechanism in which blood platelets, so-called thrombocytes, regulate the function of monocytes. Credit: Alessandro Winkler, University Hospital Bonn



Monocytes, a special type of white blood cell, secrete cytokines as inflammatory messengers that are crucial for an appropriate immune response. Researchers at the University Hospital Bonn (UKB) and the University of Bonn have now discovered that platelets, also known as thrombocytes, communicate with monocytes and increase their inflammatory capacity.

By understanding the platelet-monocyte interaction, they hope to improve the treatment of immune disorders and associated diseases.

The results of the study have been <u>published</u> in *EMBO Molecular Medicine*.

Monocytes are white blood cells, known as leukocytes. They are an important part of the innate immune system and contribute to host defense in the blood by secreting large quantities of pro-inflammatory cytokines. Abnormal activity of monocytes leads to hyperinflammation, i.e. very severe inflammation, as well as life-threatening cytokine storms.

On the other hand, disturbed monocyte function is associated with immune paralysis. In this condition, the immune system's ability to fight off invaders such as viruses and bacteria is inhibited. This increases susceptibility to infections.

"It is therefore crucial to understand how the functions of monocytes are regulated," says senior and corresponding author Prof. Dr. Bernardo Franklin from the Institute of Innate Immunity at the UKB and the Cluster of Excellence ImmunoSensation2 at the University of Bonn, explaining the motivation to investigate the role of platelets in the regulation of monocyte-induced inflammation.

Platelets as a central checkpoint in immune defense



Platelets play a central role in <u>blood clotting</u>, but are also thought to perform important functions in the <u>immune system</u>. Prof. Franklin's research team has already identified platelets as an important regulator of inflammation.

They report that a low platelet count in the rare blood disorder immune thrombocytopenia (ITP) or the artificial removal of platelets from healthy monocytes results in "immunoparalysis."

This is characterized by a disturbed cytokine reaction and is an immunological challenge. "Remarkably, supplementing monocytes with fresh platelets reverses this condition and restores the monocyte cytokine response," says corresponding and co-first author Dr. Ibrahim Hawwari, a postdoctoral fellow of the University of Bonn at the Institute of Innate Immunity at the UKB.

The Bonn researchers discovered that the pro-inflammatory signals, including NF-κB and p38 MAPK, propagate from platelets to monocytes and maintain their inflammatory capacity.

"Platelet vesicles as an extended arm of platelets control this intercellular communication," says co-first author Lukas Roßnagel, Ph.D. student ot the University of Bonn at the Institute of Innate Immunity of the UKB.

The results of the study point to a new intercellular communication mechanism in which platelets regulate monocyte function.

"Clinically, this suggests potential therapeutic strategies to counteract monocyte immune paralysis in conditions such as ITP and other <u>inflammatory diseases</u> with the addition of <u>platelets</u>," says Prof. Franklin, who hopes that an understanding of platelet-monocyte interactions will lead to improved treatment of immune disorders and related diseases.



More information: Ibrahim Hawwari et al, Platelet transcription factors license the pro-inflammatory cytokine response of human monocytes, *EMBO Molecular Medicine* (2024). DOI: 10.1038/s44321-024-00093-3

Provided by University Hospital Bonn

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