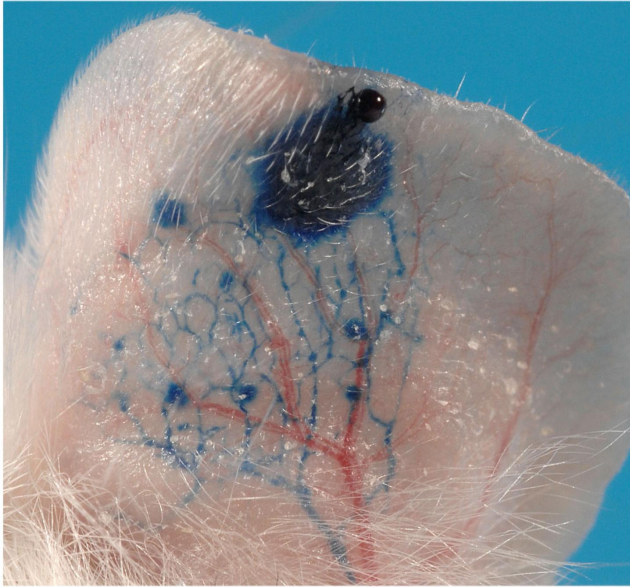


New study links neuropilin 2 deficiency to inflammation-induced edema and lymphedema

14 October 2016



This photograph highlights the structure of both the vascular and lymphatic systems. The ear is injected intradermally with blue dye to visualize the lymphatic vessels (lymphangiography). The blood vessels are visible through the skin and appear red. The blood vessels display a branching pattern; the lymphatic capillary plexus has many looping, interconnected circular structures to promote drainage in the instance of an isolated blockage. In a normal mouse, lymphatic fluid flows in one direction: from the tip of the ear to the base of the ear or from the capillary to the lymph node. The image was taken after the induction of inflammation and shows retrograde flow, pooling, and leakage of the blue lymphatic fluid, conditions that exemplify lymphedema. Credit: Kristin Johnson

Fluid accumulation and swelling (edema) may result from the malfunctioning of regulatory processes controlling vessel permeability in the body. Edema frequently occurs in chronic inflammatory diseases including psoriasis and eczema. Capillaries in the lymphatic system

usually drain the excess fluid but their dysfunction can lead to another serious condition: lymphedema. A new study published in *The American Journal of Pathology* found that deficiency in neuropilin 2 (Nrp2) receptors in vascular endothelial cells results in excessive and prolonged fluid build-up after inflammation. This discovery may guide investigators toward new pharmacological therapies for edema and lymphedema.

The newest research for lead investigator Diane R. Bielenberg, PhD, began with a twist. Bielenberg, an Assistant Professor in the Department of Surgery, Harvard Medical School and Vascular Biology Program, Boston Children's Hospital (Boston), described how previous studies that inhibited vascular endothelial growth factor receptors (VEGFRs, the co-receptors of Nrp2) have shown reduced blood vessel permeability following inflammation. "Therefore, we expected to see less permeability after inflammation in neuropilin 2 (Nrp2) knockout mice," Bielenberg explained. To her team's surprise, the Nrp2 knockout mice showed massively increased and prolonged edema following inflammation. Investigation into this finding led Bielenberg's laboratory to formulate a novel hypothesis - that endogenous Semaphorin 3F (Sema3F) protein acting through Nrp2 inhibits vascular permeability and edema. Additional experiments and data support this new hypothesis.

The investigators measured ear swelling in mice after topically introducing an inflammation-causing substance. Fluid retention was 2.5-fold higher in the Nrp2-deficient mice than controls. The swelling in [control mice](#) began to recede within two days of inflammation and returned to near-normal levels by four days. In contrast, the swelling in the Nrp2-deficient mice remained significantly elevated over control levels for more than 10 days.

The researchers also found that in Nrp2-mutant mice the lack of a superficial lymphatic capillary plexus causes impaired drainage of fluid. Co-investigators Patrick Mucka, MS, and Nicholas Levonyak, MS, of the Vascular Biology Program at Boston Children's Hospital, explain that the prolonged lymphedema seen in Nrp2 [knockout mice](#) may be caused by architectural defects resulting from improper lymphatic vessel sprouting during development.

Interestingly, the enhanced leakage in the Nrp2-deficient mice was associated with the loss of endogenous Sema3F activity. Sema3F competes with VEGFA for binding to Nrp2 and is therefore termed an inhibitory ligand of Nrp2. This competitive inhibition, in turn, prevents [vascular permeability](#). Addition of exogenous Sema3F protein therefore inhibits edema. Dr. Bielenberg encourages the pursuit of future studies on the role of the Sema3F/Nrp2 axis in chronic inflammation or lymphedema. Although these studies were performed in mice, mutations in the human NRP2 gene have been found in patients with primary lymphedema. This gene conservation suggests a high likelihood that humans and mice share a similar molecular mechanism behind this process, an encouraging indication for the translation of these findings.

More information: *The American Journal of Pathology*, [DOI: 10.1016/j.ajpath.2016.07.022](https://doi.org/10.1016/j.ajpath.2016.07.022)

Provided by Elsevier

APA citation: New study links neuropilin 2 deficiency to inflammation-induced edema and lymphedema (2016, October 14) retrieved 18 January 2022 from <https://medicalxpress.com/news/2016-10-links-neuropilin-deficiency-inflammation-induced-edema.html>

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