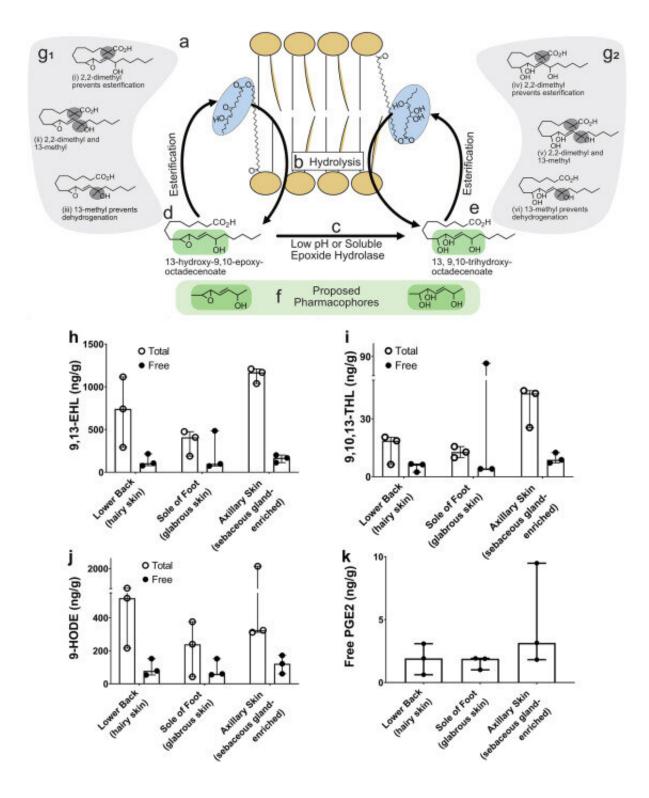


Fatty acid affects sensitivity of psoriasis plaques to heat, pain

January 3 2023, by Tracey Peake





Linoleic acid (LA) metabolites are found in both the free and total pools of fatty acids in human skin. A box represents the schematic depiction of a) LA or oxylipins (i.e., 9,13-EHL and 9,10,13-THL) derived from LA are esterified into lipid membranes. b) Hydrolysis by enzymes such as phospholipases, can release esterified oxylipins, as free acids, from the membrane where they can bind to



receptors or participate in molecular processes. c) Epoxide hydrolysis in a low pH environment or by epoxide hydrolases can convert 9,13-EHL to 9,10,13-THL. d and e) Free acid oxylipins can be esterified into the phospholipid bilayer. f) Proposed active sites (i.e., pharmacophores) of oxylipins are highlighted. g) Stable analogs of 9,13-EHL and 9,10,13-THL can be synthesized to prevent esterification, and/or prevent degradation of the pharmacophore. Concentrations of oxylipins measured h) 9,13-EHL, i) 9,10,13-THL, j) 9-HODE, and k) PGE₂ in human skin. Data in panels h—k are presented as Mean ± Standard Deviation (SD); no statistical tests were run on these data. Each data point corresponds to one biological replicate, n=3. Credit: *JID Innovations* (2022). DOI: 10.1016/j.xjidi.2022.100177

A common fatty acid found in the Western diet breaks down into compounds that contribute to increased temperature and pain—but not itch—sensitivity in psoriatic lesions. The finding could lead to better understanding of how lipids communicate with sensory neurons, and potentially to improved pain and sensitivity treatments for psoriasis patients.

Linoleic acid is a fatty acid found in vegetable oils, nuts and seeds, and is one of the predominant fatty acids found in the Western diet.

Metabolites from linoleic acid—the products formed when the body breaks it down through digestion—play a role in skin barrier function.

"We noticed high levels of two types of lipids derived from linoleic acid in psoriatic lesions," says Santosh Mishra, associate professor of neuroscience at North Carolina State University and corresponding author of the research. "That led us to wonder whether the lipids might affect how sensory neurons in these lesions communicate. We decided to investigate whether their presence could be related to the temperature or pain hypersensitivity that many psoriasis patients report."



The work appears in *JID Innovations*. Josh Wheeler, a postdoctoral researcher at NC State, is first author.

The research team used mass spectrometry to create <u>lipid</u> profiles of skin from psoriatic lesions. They focused on two types of linoleic acidderived lipids, or oxylipids: 13-hydroxy-9,10-epoxy octadecenoate (9,13-EHL) and 9,10,13-trihydroxy-octadecenoate (9,10,13-THL). The first form, 9,13-EHL, can convert into the more stable 9,10,13-THL form via interaction with certain enzymes.

The researchers found that while both forms bind to receptors on <u>sensory neurons</u> within the skin, the more stable form—9,10,13-THL—had a longer lasting effect than 9,13-EHL.

They also found that once the lipids bind to the neuronal receptor, they activate the neurons expressing TRPA1 and TRPV1 receptors that are involved in temperature and pain hypersensitivity, opening communications channels to the central nervous system.

Interestingly, the lipids did not have any effect on itch.

"It was surprising that these lipids could create hypersensitivity but not impact <u>itch</u> sensation, which is usually the most troublesome symptom associated with psoriasis," Mishra says. "This most likely has to do with how the neuron is activated—a mechanism we still haven't uncovered."

Now that an association between linoleic acid and <u>hypersensitivity</u> to temperature and pain has been established, the researchers want to further explore exactly how this response is being created. They hope that the answers may lead to solutions that can relieve these symptoms in psoriasis patients.

"We know that this lipid moves from one form to another, but don't yet



know what causes that," Mishra says. "We also know what protein the lipids are binding to, but not where the bond occurs. Answering these questions may hopefully lead to new therapies—or dietary solutions—for some psoriasis sufferers."

More information: Joshua J. Wheeler et al, Endogenous Derivatives of Linoleic Acid and Their Stable Analogues are Potential Pain Mediators, *JID Innovations* (2022). DOI: 10.1016/j.xjidi.2022.100177

Provided by North Carolina State University

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