



Research Questions

- ☐ Why? Is there a genetic predisposition
- ☐ Mechanism? Co-stimulatory signals? Preventable?
- ☐ How long to avoid? Evidence of re-sensitization from tick and/or meat
- ☐ Treatment (besides avoidance). IT? Anti-IgE?
- ☐ Molecular Mimicry?

Clinical Update



**U.S. FOOD & DRUG
ADMINISTRATION**

[A to Z Index](#) | [Follow FDA](#) | [En Español](#)

Search FDA



[Home](#) [Food](#) [Drugs](#) [Medical Devices](#) [Radiation-Emitting Products](#) [Vaccines, Blood & Biologics](#) [Animal & Veterinary](#) [Cosmetics](#) [Tobacco Products](#)

News & Events

[Home](#) > [News & Events](#) > [Newsroom](#) > [Press Announcements](#)

FDA News Release

FDA approves new eczema drug Dupixent

[f SHARE](#) [t TWEET](#) [in LINKEDIN](#) [p PIN IT](#) [e EMAIL](#) [p PRINT](#)

**For Immediate
Release**

March 28, 2017

Release

[Español](#)

The U.S. Food and Drug Administration today approved Dupixent (dupilumab) injection to treat adults with moderate-to-severe eczema (atopic dermatitis). Dupixent is intended for patients whose eczema is not controlled adequately by topical therapies, or those for whom topical therapies are not advisable. Dupixent can be used with or without topical corticosteroids.

Inquiries

Media

[✉ Sarah Peddicord](#)
[☎ 301-796-2805](#)

[✉ Andrea Fischer](#)
[☎ 301-796-0393](#)

Consumers

[☎ 888-INFO-FDA](#)

Related Information

- [FDA Approved Drugs: Questions and Answers](#)