

INDICATION

DUPIXENT is indicated for the treatment of adult and pediatric patients aged 1 year and older, weighing at least 15 kg, with eosinophilic esophagitis (EoE).

IMPORTANT SAFETY INFORMATION

CONTRAINDICATION: DUPIXENT is contraindicated in patients with known hypersensitivity to dupilumab or any of its excipients.

Please see additional Important Safety Information below.



Transform the Way You Manage Eosinophilic Esophagitis (EoE) - The First and Only FDA-Approved Treatment for EoE Patients As Young As 1 Year

PRESENTED BY:



David Poppers, MD, PhD
Professor of Medicine,
Director of GI Quality and
Strategic Initiatives
NYU Grossman School of Medicine,
NYU Langone Health
New York, NY

Saturday, June 6
12:25 PM – 1:00 PM ET
London Ballroom

IMPORTANT SAFETY INFORMATION, CONTINUED

WARNINGS AND PRECAUTIONS

Hypersensitivity: Hypersensitivity reactions, including anaphylaxis, serum sickness or serum sickness-like reactions, angioedema, generalized urticaria, rash, erythema nodosum, and erythema multiforme have been reported. If a clinically significant hypersensitivity reaction occurs, institute appropriate therapy and discontinue DUPIXENT.

Risk Associated with Abrupt Reduction of Corticosteroid Dosage: Do not discontinue systemic, topical, or inhaled corticosteroids abruptly upon initiation of DUPIXENT. Reductions in corticosteroid dose, if appropriate, should be gradual and performed under the direct supervision of a healthcare provider. Reduction in corticosteroid dose may be associated with systemic withdrawal symptoms and/or unmask conditions previously suppressed by systemic corticosteroid therapy.

Patients with Co-morbid Asthma: Advise patients with co-morbid asthma not to adjust or stop their asthma treatments without consultation with their physicians.

Arthralgia and Psoriatic Arthritis: Arthralgia has been reported with the use of DUPIXENT with some patients reporting gait disturbances or decreased mobility associated with joint symptoms; some cases resulted in hospitalization. Cases of new-onset psoriatic arthritis requiring systemic treatment have been reported with the use of DUPIXENT. Advise patients to report new onset or worsening joint symptoms. If symptoms persist or worsen, consider rheumatological evaluation and/or discontinuation of DUPIXENT.

Parasitic (Helminth) Infections: It is unknown if DUPIXENT will influence the immune response against helminth infections. Treat patients with pre-existing helminth infections before initiating therapy with DUPIXENT. If patients become infected while receiving treatment with DUPIXENT and do not respond to anti-helminth treatment, discontinue treatment with DUPIXENT until the infection resolves.

Vaccinations: Consider completing all age-appropriate vaccinations as recommended by current immunization guidelines prior to initiating DUPIXENT. Avoid use of live vaccines during treatment with DUPIXENT.

ADVERSE REACTIONS: The most common adverse reactions (incidence $\geq 2\%$) in patients with EoE are injection site reactions, upper respiratory tract infections, arthralgia, and herpes viral infections.

USE IN SPECIFIC POPULATIONS

- Pregnancy:** A pregnancy exposure registry monitors pregnancy outcomes in women exposed to DUPIXENT during pregnancy. To enroll or obtain information call 1-877-311-8972 or go to <https://mothertobaby.org/ongoing-study/dupilumab/>. Available data from case reports and case series with DUPIXENT use in pregnant women have not identified a drug-associated risk of major birth defects, miscarriage or adverse maternal or fetal outcomes. Human IgG antibodies are known to cross the placental barrier; therefore, DUPIXENT may be transmitted from the mother to the developing fetus.
- Lactation:** There are no data on the presence of DUPIXENT in human milk, the effects on the breastfed infant, or the effects on milk production. Maternal IgG is known to be present in human milk. The developmental and health benefits of breastfeeding should be considered along with the mother's clinical need for DUPIXENT and any potential adverse effects on the breastfed child from DUPIXENT or from the underlying maternal condition.

Please see accompanying full Prescribing Information.

This event is neither sponsored by nor endorsed by ACG.

In accordance with the PhRMA Code on Interactions with Healthcare Professionals, this Program is limited to U.S. Healthcare Professionals and persons with bona fide professional interest in the information presented. Attendance at this Program by guests or spouses is not permitted unless they would qualify as an appropriate attendee on their own. If a meal is provided, actively licensed Minnesota and Vermont prescribers may attend, but not partake in the meal. Full-time Federal Employees may attend and partake in the meal if the Program is considered widely attended (50 or more attendees). If not, they may attend, but not partake in the meal. Part-time Federal Employees acting in their civilian capacity may attend the Program and partake in the meal. Alcohol is not permitted in connection with Speaker Programs and will not be provided. The value of any meal provided in connection with the Program may be reported in accordance with federal and state laws and regulations.

sanofi | **REGENERON**[®]

© 2025 Sanofi and Regeneron Pharmaceuticals, Inc. All Rights Reserved.
DUPIXENT is a registered trademark of Sanofi or an affiliate.
09/2025 US.DUP.25.09.0245

DUPIXENT[®]
(dupilumab) Injection
200mg • 300mg