

DOSAGE & ADMINISTRATION

AD

PN

CSU

BP

MULTIPLE
ADMINISTRATION OPTIONS

ADMINISTERING
DUPIXENT

DUPIXENT MyWay[®]

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DUPIXENT IS FDA-APPROVED FOR USE
IN **4** DERMATOLOGIC CONDITIONS¹



ATOPIC DERMATITIS (AD)

uncontrolled
moderate-to-severe



PRURIGO NODULARIS (PN)



CHRONIC SPONTANEOUS URTICARIA (CSU)

symptomatic despite
H1 antihistamines

NOW APPROVED



BULLOUS PEMPHIGOID (BP)

NOW APPROVED

INDICATIONS

ATOPIC DERMATITIS: DUPIXENT is indicated for the treatment of adult and pediatric patients aged 6 months and older with moderate-to-severe atopic dermatitis whose disease is not adequately controlled with topical prescription therapies or when those therapies are not advisable. DUPIXENT can be used with or without topical corticosteroids.

PRURIGO NODULARIS: DUPIXENT is indicated for the treatment of adult patients with prurigo nodularis (PN).

CHRONIC SPONTANEOUS URTICARIA: DUPIXENT is indicated for the treatment of adult and pediatric patients aged 12 years and older with chronic spontaneous urticaria (CSU) who remain symptomatic despite H1 antihistamine treatment. Limitations of Use: DUPIXENT is not indicated for treatment of other forms of urticaria.

BULLOUS PEMPHIGOID: DUPIXENT is indicated for the treatment of adult patients with bullous pemphigoid (BP).

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 throughout and click [here](#) for full Prescribing Information.



UNCONTROLLED MODERATE-TO-SEVERE ATOPIC DERMATITIS (AD)

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ADULTS (18+ YEARS) ONE DOSAGE REGIMEN¹



18+ YEARS

One dosage
regimen in adults

Initial loading dose:
600 mg 2 x 300 mg

Followed by:
300 mg Q2W 1 x 300 mg

CHILDREN AND ADOLESCENTS (6 TO 17 YEARS) WEIGHT-TIERED DOSAGE REGIMEN^{1,a}



6 TO 17
YEARS

≥60 kg

Initial loading dose:
600 mg 2 x 300 mg

Followed by:
300 mg Q2W 1 x 300 mg

30 to <60 kg

Initial loading dose:
400 mg 2 x 200 mg

Followed by:
200 mg Q2W 1 x 200 mg

15 to <30 kg

Initial loading dose:
600 mg 2 x 300 mg

Followed by:
300 mg Q4W 1 x 300 mg

INFANTS TO PRESCHOOLERS (6 MONTHS TO 5 YEARS) WEIGHT-TIERED DOSAGE REGIMEN^{1,a}



6 MONTHS
TO 5 YEARS

15 to <30 kg

No initial loading
dose recommended

300 mg Q4W 1 x 300 mg

5 to <15 kg

No initial loading
dose recommended

200 mg Q4W 1 x 200 mg

Q2W, once every 2 weeks; Q4W, once every 4 weeks.

^a 5 kg is equal to 11 lb; 15 kg is equal to 33 lb; 30 kg is equal to 66 lb; 60 kg is equal to 132 lb.

HOW DUPIXENT IS SUPPLIED



Choice of administration: Available in a 200 mg and 300 mg single-dose pre-filled pen (for indicated patients 2+ years of age) or pre-filled syringe (for indicated patients 6+ months of age) for subcutaneous injection¹

IMPORTANT SAFETY INFORMATION

WARNINGS AND PRECAUTIONS

Hypersensitivity: Hypersensitivity reactions, including anaphylaxis, acute generalized exanthematous pustulosis (AGEP), serum sickness or serum sickness-like reactions, angioedema, generalized urticaria, rash, erythema nodosum, and erythema multiforme have been reported. A case of AGEP was reported in an adult subject who participated in the bullous pemphigoid development program. If a clinically significant hypersensitivity reaction occurs, institute appropriate therapy and discontinue DUPIXENT.

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PRURIGO NODULARIS (PN)

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ADULTS (18+ YEARS) ONE DOSAGE REGIMEN¹



18+ YEARS

One dosage
regimen in adults

Initial loading dose:
600 mg 2 x 300 mg

Followed by:
300 mg Q2W 1 x 300 mg

Q2W, once every 2 weeks.

HOW DUPIXENT IS SUPPLIED



Choice of administration: Available in a 300 mg single-dose pre-filled pen (for indicated patients 18+ years of age) or pre-filled syringe (for indicated patients 18+ years of age) for subcutaneous injection¹

IMPORTANT SAFETY INFORMATION

WARNINGS AND PRECAUTIONS (cont'd)

Conjunctivitis and Keratitis: Conjunctivitis and keratitis occurred more frequently in AD and BP subjects who received DUPIXENT versus placebo, with conjunctivitis being the most frequently reported eye disorder in AD. Conjunctivitis also occurred more frequently in PN subjects who received DUPIXENT compared to those who received placebo. Conjunctivitis and keratitis have been reported with DUPIXENT in postmarketing settings, predominantly in AD patients. Some patients reported visual disturbances (e.g., blurred vision) associated with conjunctivitis or keratitis. Advise patients or their caregivers to report new-onset or worsening eye symptoms. Consider ophthalmological examination for patients who develop conjunctivitis that does not resolve following standard treatment or signs and symptoms suggestive of keratitis, as appropriate.

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 not to adjust or stop their asthma treatments without consultation with their physicians.

Please see additional Important Safety Information throughout and click [here](#) for full Prescribing Information.



CHRONIC SPONTANEOUS URTICARIA (CSU) SYMPTOMATIC DESPITE H1 ANTIHISTAMINES

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ADULTS (18+ YEARS) ONE DOSAGE REGIMEN¹



18+ YEARS

One dosage
regimen in adults

Initial loading dose:
600 mg 2 x 300 mg

Followed by:
300 mg Q2W 1 x 300 mg

ADOLESCENTS (12 TO 17 YEARS) WEIGHT-TIERED DOSAGE REGIMEN^{1,a}



12 TO 17
YEARS

≥60 kg

Initial loading dose:
600 mg 2 x 300 mg

Followed by:
300 mg Q2W 1 x 300 mg

30 to <60 kg

Initial loading dose:
400 mg 2 x 200 mg

Followed by:
200 mg Q2W 1 x 200 mg

Q2W, once every 2 weeks.

^a 30 kg is equal to 66 lb; 60 kg is equal to 132 lb.

HOW DUPIXENT IS SUPPLIED



Choice of administration: Available in a 200 mg and 300 mg single-dose pre-filled pen (for indicated patients 12+ years of age) or pre-filled syringe (for indicated patients 12+ years of age) for subcutaneous injection¹

IMPORTANT SAFETY INFORMATION

WARNINGS AND PRECAUTIONS (cont'd)

Psoriasis: Cases of new-onset psoriasis have been reported with the use of DUPIXENT for the treatment of atopic dermatitis, including in patients without a family history of psoriasis. In postmarketing reports, these cases resulted in partial or complete resolution of psoriasis with discontinuation of dupilumab, with or without use of supplemental treatment for psoriasis (topical or systemic). Those who continued dupilumab received supplemental treatment for psoriasis to improve associated symptoms. Advise patients to report new-onset psoriasis symptoms. If symptoms persist or worsen, consider dermatologic evaluation and/or discontinuation of DUPIXENT.

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.

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BULLOUS PEMPHIGOID (BP)

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ADULTS (18+ YEARS) ONE DOSAGE REGIMEN¹



18+ YEARS

**One dosage
regimen in adults**

**Initial loading dose:
600 mg** 2 x 300 mg

**Followed by:
300 mg Q2W** 1 x 300 mg

Q2W, once every 2 weeks.

Use DUPIXENT in combination with a tapering course of oral corticosteroids. Once disease control has occurred, gradually taper corticosteroids after which continue DUPIXENT as monotherapy. In case of relapse, corticosteroids may be added if medically advisable.

HOW DUPIXENT IS SUPPLIED



Choice of administration: Available in a 300 mg single-dose pre-filled pen (for indicated patients 18+ years of age) or pre-filled syringe (for indicated patients 18+ years of age) for subcutaneous injection¹

IMPORTANT SAFETY INFORMATION

WARNINGS AND PRECAUTIONS (cont'd)

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.

Please see additional Important Safety Information throughout and click [here](#) for full Prescribing Information.



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PRE-FILLED PEN (FOR AGES 2+ YEARS)^{1,2}

- Subcutaneous autoinjector with hidden needle
- Needle cap is not made with natural rubber latex
- Visual and audible feedback
- Available in 200 mg and 300 mg

PRE-FILLED SYRINGE (FOR AGES 6+ MONTHS)^{1,3}

- Subcutaneous injection with needle shield
- Includes finger grip
- Needle cap is not made with natural rubber latex
- Available in 200 mg and 300 mg

AVAILABILITY

- DUPIXENT is available in cartons containing 2 single-dose pre-filled pens or 2 single-dose pre-filled syringes
 - The pre-filled pen is designed to deliver **300 mg** of DUPIXENT in a **2 mL** solution or **200 mg** in a **1.14 mL** solution
 - The pre-filled syringe is designed to deliver **300 mg** of DUPIXENT in a **2 mL** solution or **200 mg** in a **1.14 mL** solution



IMPORTANT SAFETY INFORMATION

ADVERSE REACTIONS:

Most common adverse reactions are:

- **Atopic Dermatitis** (incidence $\geq 1\%$): injection site reactions, conjunctivitis, blepharitis, oral herpes, keratitis, eye pruritus, other herpes simplex virus infection, dry eye, and eosinophilia. The safety profile in pediatric patients through Week 16 was similar to that of adults with AD. In an open-label extension study, the long-term safety profile of DUPIXENT \pm TCS in pediatric patients observed through Week 52 was consistent with that seen in adults with AD, with hand-foot-and-mouth disease and skin papilloma (incidence $\geq 2\%$) reported in patients 6 months to 5 years of age. These cases did not lead to study drug discontinuation.
- **Prurigo Nodularis** (incidence $\geq 2\%$): nasopharyngitis, conjunctivitis, herpes infection, dizziness, myalgia, and diarrhea.
- **Chronic Spontaneous Urticaria** (incidence $\geq 2\%$): injection site reactions.
- **Bullous Pemphigoid** (incidence $\geq 2\%$): arthralgia, conjunctivitis, vision blurred, herpes viral infections, keratitis.

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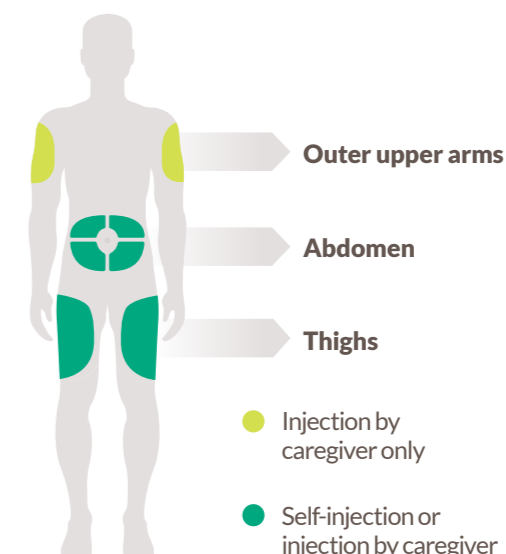
HOW TO TAKE DUPIXENT¹

- DUPIXENT is intended for use under the guidance of a healthcare provider
- A patient may self-inject DUPIXENT after receiving training in subcutaneous injection technique
 - In children 6 months to less than 12 years of age, DUPIXENT should be given by a caregiver
 - In children 12 years of age and older, it is recommended that DUPIXENT be given by or under the supervision of an adult
- DUPIXENT can be administered in the office under the guidance of a healthcare provider if the patient is not an appropriate candidate for at-home administration
- Consider completing all age-appropriate vaccinations as recommended by current immunization guidelines prior to initiating treatment with DUPIXENT

ADMINISTER AT DIFFERENT INJECTION SITES¹⁻³

Remember to provide proper training to patients and/or caregivers on the preparation and administration of DUPIXENT prior to use, according to the Instructions for Use.

- When administering a loading dose (consisting of 2 injections), remember to administer each injection into a different site
- Administer the subcutaneous injection into the thigh or abdomen, except for the 2 inches (5 cm) around the navel
- The upper arm can also be used if a caregiver administers the injection
- Rotate the injection site with each injection. **DO NOT** inject DUPIXENT into skin that is tender, damaged, bruised, or scarred



MISSED DOSE INFORMATION¹

- If an every 2 week dose is missed, administer the injection within 7 days from the missed dose and then resume the patient's original schedule. If the missed dose is not administered within 7 days, administer the dose, starting a new schedule based on this date
- If an every 4 week dose is missed, administer the injection within 7 days from the missed dose and then resume the patient's original schedule. If the missed dose is not administered within 7 days, administer the dose, starting a new schedule based on this date

DUPIXENT prescriptions can only be filled by specialty pharmacies that provide appropriate handling and shipping to home. **DUPIXENT MyWay[®]** offers further patient access and support.

Please see additional Important Safety Information throughout and click [here](#) for full Prescribing Information.



DUPIXENT MyWay[®] CAN HELP ELIGIBLE PATIENTS START AND STAY ON TRACK WITH DUPIXENT

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BP

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HELP TO UNDERSTAND AND NAVIGATE THE INSURANCE PROCESS FOR DUPIXENT



SUPPLEMENTAL INJECTION TRAINING IN PERSON, VIRTUALLY, OR BY PHONE



THE QUICK START PROGRAM may be able to temporarily provide DUPIXENT at no cost to help bridge patients to therapy if there is a coverage delay. Subject to the eligibility requirements listed below^a



TEXT AND EMAIL REMINDERS FOR PATIENTS TO TAKE THEIR NEXT DOSE AND REFILL THEIR PRESCRIPTION



COPAY AND FINANCIAL ASSISTANCE FOR ELIGIBLE PATIENTS^b



TIPS AND TOOLS TO HELP PATIENTS MANAGE THEIR TREATMENT PLANS

DUPIXENT MyWay support begins with the enrollment form. Speak with your field representative, visit [DUPIXENTHCP.COM](https://www.dupilumab.com), or call **1-844-DUPIXENT** to get started

IMPORTANT SAFETY INFORMATION 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.

Please see additional Important Safety Information throughout and click [here](#) for full Prescribing Information.

^a [Patients may be eligible for the Quick Start Program if they: have a valid DUPIXENT prescription for an FDA-approved indication, are new to DUPIXENT, have prescription drug coverage for DUPIXENT through a commercial insurance plan not funded through a government healthcare program, are a resident of the United States, District of Columbia, or Puerto Rico and are treated by a licensed prescriber therein, are not an inpatient, are experiencing a coverage delay, and are actively seeking coverage through their insurance provider. Some patients may be subject to additional diagnostic criteria for eligibility.]

^b [Subject to the program maximum per patient per calendar year. Approval is not guaranteed. **THIS IS NOT INSURANCE.** Not valid for prescriptions paid, in whole or in part, by Medicaid, Medicare, VA, DOD, TRICARE, or other federal or state programs, including any state pharmaceutical assistance programs. This program is not valid where prohibited by law, taxed, or restricted. DUPIXENT MyWay reserves the right to rescind, revoke, terminate, or amend this offer, eligibility, and terms of use at any time without notice. Any savings provided by the program may vary depending on patients' out-of-pocket costs. The program is intended to help patients afford DUPIXENT. Patients may have insurance plans that attempt to dilute the impact of the assistance available under the program. In those situations, the program may change its terms. Additional terms and conditions apply.]



ADMINISTRATION OPTIONS AND OTHER CONSIDERATIONS FOR YOU AND YOUR PATIENTS

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AT-HOME SELF-ADMINISTRATION

- Train patients and/or caregivers on proper administration prior to use



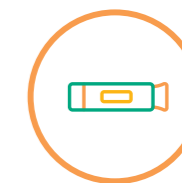
NO INITIAL LAB TESTING OR ONGOING LAB MONITORING

according to the Prescribing Information



PRE-FILLED SYRINGE

(for indicated patients 6+ months of age)



PRE-FILLED PEN

(for indicated patients 2+ years of age)



PRE-FILLED SYRINGE →



PRE-FILLED PEN →

VISIT [DUPIXENT.COM](https://www.dupixent.com) TO VIEW ADMINISTRATION VIDEOS

Patients and caregivers can visit [DUPIXENT.COM](https://www.dupixent.com) or call **1-844-DUPIXENT** to learn more about support available through *DUPIXENT MyWay*

IMPORTANT SAFETY INFORMATION

USE IN SPECIFIC POPULATIONS (cont'd)

- **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 additional Important Safety Information throughout and click [here](#) for full Prescribing Information.

References: 1. DUPIXENT Prescribing Information. 2. DUPIXENT 200 mg/300 mg Pre-filled Pen Instructions for Use. 3. DUPIXENT 200 mg/300 mg Pre-filled Syringe Instructions for Use.

