

DUPIXENT[®]
(dupilumab) Injection 300mg

**WELCOME TO DUPIXENT, THE FIRST AND ONLY
FDA-APPROVED TREATMENT FOR ADULTS WITH**

BULLOUS PEMPHIGOID (BP)

Patient portrayal

INDICATION

DUPIXENT is a prescription medicine used to treat adults with bullous pemphigoid (BP). It is not known if DUPIXENT is safe and effective in children with BP under 18 years of age.

IMPORTANT SAFETY INFORMATION: Do not use if you are allergic to dupilumab or to any of the ingredients in DUPIXENT[®].

Please see Important Safety Information throughout and full Prescribing Information and Patient Information.

WHAT IS BULLOUS PEMPHIGOID (BP)?

Bullous pemphigoid is a rare chronic-relapsing autoimmune skin condition that causes fluid-filled blisters and other inflammatory lesions. It is caused by the immune system attacking a layer of tissue in the skin. These outbreaks could be triggered by certain medications. The reason why this happens is not fully understood.

SIGNS AND SYMPTOMS OF BP INCLUDE:

- ▶ Recurring phases of:
 - ▶ Blisters and painful sores
 - ▶ Rashes
- ▶ Intense itchiness

While blisters may occur anywhere, they commonly form near creases in the skin.



Patient portrayal

IMPORTANT SAFETY INFORMATION: Before using DUPIXENT, tell your healthcare provider about all your medical conditions, including if you: have eye problems; have a parasitic (helminth) infection; are scheduled to receive any vaccinations. You should not receive a “live vaccine” right before and during treatment with DUPIXENT.

HOW DO YOU TREAT BP?

DUPIXENT is approved by the FDA to treat bullous pemphigoid in adults 18 years of age and older. It targets a source of underlying inflammation that may be a cause of BP. DUPIXENT works continuously between doses to help manage symptoms.

DUPIXENT was studied in a 52-week trial with 106 adults with BP. Some patients on DUPIXENT achieved sustained remission at week 36. Sustained remission means there are no new blisters, old ones are healing, and steroids are tapered off by week 16, with no relapses from week 16 to week 36, and no rescue treatment through week 36.

AT 36 WEEKS

MORE PATIENTS HAD SUSTAINED REMISSION
(18% on DUPIXENT vs 6% on placebo)

MORE PATIENTS EXPERIENCED LESS ITCH
(38% on DUPIXENT vs 11% on placebo)

LESS TOTAL ORAL STEROIDS WERE USED
(2.8 g for patients on DUPIXENT vs 4.1 g on placebo)

This analysis was not statistically significant.

The most common side effects observed in clinical trials were: joint pain (arthralgia), eye problems, including eye and eyelid inflammation, redness, swelling, itching, and blurred vision, and herpes virus infections.

Please see Important Safety Information throughout and full [Prescribing Information](#) and [Patient Information](#).

ABOUT DUPIXENT[®] (dupilumab)

- ▶ First and only FDA-approved treatment for adults with BP
- ▶ Works continuously between doses to help manage symptoms
- ▶ Not a steroid or immunosuppressant
- ▶ Doesn't require initial or routine blood work, according to the Prescribing Information
- ▶ Has no known drug-to-drug interactions

DUPIXENT IS A TREATMENT THAT

- » TARGETS A KEY SOURCE OF UNDERLYING INFLAMMATION THAT MAY CONTRIBUTE TO BP
- » WAS STUDIED IN A CLINICAL TRIAL THAT EVALUATED TREATMENT IN ADULTS WITH BP

IMPORTANT SAFETY INFORMATION: Before using DUPIXENT, tell your healthcare provider about all your medical conditions, including if you (cont'd): are pregnant or plan to become pregnant. It is not known whether DUPIXENT will harm your unborn baby. A pregnancy registry for women who take DUPIXENT during pregnancy collects information about the health of you and your baby; are breastfeeding or plan to breastfeed. It is not known whether DUPIXENT passes into your breast milk.

Tell your healthcare provider about all the medicines you take, including prescription and over-the-counter medicines, vitamins, and herbal supplements.

Especially tell your healthcare provider if you are taking oral, topical, or inhaled corticosteroid medicines or if you have BP and asthma and use an asthma medicine. **Do not** change or stop your other medicines, including corticosteroid medicine or other asthma medicine, without talking to your healthcare provider. This may cause other symptoms that were controlled by those medicines to come back.

Please see [Important Safety Information](#) throughout and full [Prescribing Information](#) and [Patient Information](#).

HOW DUPIXENT IS TAKEN

DUPIXENT is a biologic medicine given under the skin (subcutaneous injection). It can be:

- ▶ Self-administered or given by your caregiver at home
- ▶ Administered by your healthcare provider at their office

DUPIXENT is taken in combination with a taper regimen of oral steroids. If necessary, your doctor may reinstitute steroids.



PRE-FILLED PEN



PRE-FILLED SYRINGE

Adult patients begin DUPIXENT with an initial dose of 600 mg (two 300-mg injections) followed by 300 mg every other week. Use DUPIXENT exactly as prescribed by your healthcare provider. Your healthcare provider will tell you how much DUPIXENT to inject and how often.

Do not try to prepare and inject DUPIXENT until being properly trained by your doctor.



USE THE CAMERA ON YOUR PHONE TO SCAN OR [CLICK HERE](#) TO WATCH SUPPLEMENTAL TRAINING VIDEOS ON HOW TO ADMINISTER DUPIXENT

HELPFUL RESOURCES FOR YOU, EVERY STEP OF THE WAY

DUPIXENT MyWay is a patient support program designed to help you start DUPIXENT as quickly as possible and stay on track with your treatment.

ENROLLING IN **DUPIXENT MyWay**[®] IS FREE AND SIMPLE

Contact us directly at 1-844-DUPIXENT (1-844-387-4936) Option 1, Monday–Friday 8 PM–9 PM ET.

Once enrolled in the program, you'll have a dedicated Case Manager who can provide:



Help with understanding your insurance coverage for DUPIXENT



Financial assistance, including copay support, if eligible



Connection to a **DUPIXENT MyWay** Nurse Educator who can offer supplemental injection training in person, virtually, or by the phone



Tips and tools to help you manage treatment

IMPORTANT SAFETY INFORMATION: DUPIXENT can cause serious side effects, including: Allergic reactions. DUPIXENT can cause allergic reactions, including skin reactions, that can sometimes be severe. Stop using DUPIXENT and tell your healthcare provider or get emergency help right away if you get any of the following signs or symptoms: breathing problems or wheezing, swelling of the face, lips, mouth, tongue, or throat, fainting, dizziness, feeling lightheaded, fast pulse, fever, hives, skin rash, including rash that looks like a bullseye, painful red or blue bumps under the skin, or red pus-filled spots on the skin, general ill feeling, itching, swollen lymph nodes, nausea or vomiting, joint pain, or cramps in your stomach area.

OPTIONS FOR FINANCIAL SUPPORT



The **DUPIXENT MyWay Copay Card Program** may help eligible patients with commercial health insurance cover out-of-pocket costs for DUPIXENT.*

The **DUPIXENT MyWay Patient Assistance Program**[†] may be able to help you if you do not have health insurance, if you are experiencing difficulty paying for your DUPIXENT treatment, or have Medicare Part D.

Find out if you qualify by speaking with a **DUPIXENT MyWay Case Manager** at 1-844-DUPIXENT (1-844-387-4936) Option 1.



USE THE CAMERA ON YOUR PHONE TO SCAN OR CLICK HERE TO LEARN MORE ABOUT **DUPIXENT MyWay**

**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.*

[†]Eligible commercial patients and Medicare patients without the Part D (pharmacy) benefit may receive free shipments of up to an 84-day supply of DUPIXENT (for up to 12 months). Patients may reapply after 12 months if they still meet the eligibility criteria. Patients with Medicare Part D who meet eligibility criteria must reapply for the Patient Assistance Program each calendar year, or by December 31st for eligibility consideration for the following year. The assistance period for Medicaid patients varies based on the eligibility criteria. Please note that DUPIXENT MyWay reserves the right to make eligibility determinations, monitor participation, ensure equitable product availability, and modify or discontinue the DUPIXENT MyWay Patient Assistance Program at any time without notice.

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(dupilumab) Injection 300mg

**HELP
HEAL
YOUR
SKIN
FROM
WITHIN**[™]



SCAN WITH YOUR PHONE CAMERA,
CLICK HERE, OR VISIT
[DUPIXENT.COM](https://www.dupilumab.com) TO LEARN MORE

» TALK TO YOUR DOCTOR ABOUT DUPIXENT TODAY

IMPORTANT SAFETY INFORMATION: DUPIXENT can cause serious side effects, including (cont'd): Eye problems. Tell your healthcare provider right away if you have any new or worsening eye problems, including eye pain or changes in vision, such as blurred vision. Your healthcare provider may send you to an ophthalmologist for an eye exam if needed. **Joint aches and pain.** Some people who use DUPIXENT have had trouble walking or moving due to their joint symptoms, and in some cases needed to be hospitalized. Tell your healthcare provider about any new or worsening joint symptoms. Your healthcare provider may stop DUPIXENT if you develop joint symptoms.

The most common side effects in patients with bullous pemphigoid include joint pain (arthralgia); eye problems, including eye and eyelid inflammation, redness, swelling, itching, and blurred vision; and herpes virus infections.

Tell your healthcare provider if you have any side effect that bothers you or that does not go away. These are not all the possible side effects of DUPIXENT. Call your doctor for medical advice about side effects. You are encouraged to report negative side effects of prescription drugs to the FDA. Visit www.fda.gov/medwatch, or call 1-800-FDA-1088.

Use DUPIXENT exactly as prescribed by your healthcare provider. It's an injection given under the skin (subcutaneous injection). Your healthcare provider will decide if you or your caregiver can inject DUPIXENT. **Do not** try to prepare and inject DUPIXENT until you or your caregiver have been trained by your healthcare provider.

Please see full [Prescribing Information](#) and [Patient Information](#).

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