



**UNCOVER WHAT'S POSSIBLE
WITH TREMFYA®**



RESULTS



FDA APPROVED FOR ADULTS WITH MODERATE TO SEVERE **PLAQUE PSORIASIS** SINCE 2017

ALSO APPROVED FOR ADULTS WITH **ACTIVE PSORIATIC ARTHRITIS**

TREMFYA® is the first FDA-approved medication of its kind to selectively block IL-23.

TREMFYA® is a prescription medicine used to treat adults:

- with moderate to severe plaque psoriasis who may benefit from taking injections or pills (systemic therapy) or phototherapy (treatment using ultraviolet light or UV light)
- with active psoriatic arthritis (PsA).

UNCOVER CLEARER SKIN THAT CAN LAST

In clinical studies, at week 16, at least **7 out of 10 patients** with moderate to severe plaque psoriasis saw at least **90% clearer skin**, and more than 8 out of 10 were rated clear or almost clear.

In a clinical study, nearly 9 out of 10 people who saw 90% clearer skin at 28 weeks continued to see 90% clearer skin at 48 weeks.

UNCOVER 4-YEAR* RESULTS

In another study, the majority of TREMFYA® patients saw 90% clearer skin at nearly 4 years. At 1 year or thereafter, patients knew they were getting TREMFYA®, which may affect results. Results may vary.

*At week 204.

UNCOVER COMPLETELY CLEAR SKIN

In one clinical study, 5 out of 10 patients saw completely clear skin or were rated as clear at 16 weeks.

UNCOVER RESULTS YOU CAN SEE



Photos are of a real patient with moderate to severe plaque psoriasis who received TREMFYA® 100 mg at weeks 0, 4, and then every 8 weeks.

Individual results may vary.

UNCOVER JOINT RELIEF, AND CLEARER SKIN TOO

TREMFYA® can help reduce the **joint pain, stiffness, and swelling** of active psoriatic arthritis (PsA).

In clinical studies, more than 5 out of 10 patients treated with TREMFYA® had at least a 20% improvement in active PsA symptoms (joint pain, swelling, stiffness) at 24 weeks.

In clinical studies at 24 weeks, an improvement in psoriasis skin plaques was seen in psoriatic arthritis patients treated with TREMFYA®.

FEWER JOINT SYMPTOMS—MORE OF WHAT YOU USED TO DO

TREMFYA® can help reduce the joint pain, stiffness, and swelling that make everyday tasks harder to do. In clinical studies, patients' ability to perform everyday activities was assessed in a questionnaire. At 24 weeks, people taking TREMFYA® showed an overall improvement in their ability to perform everyday activities such as:



Individual results may vary.

FATIGUE WAS REDUCED IN SOME ACTIVE PSA PATIENTS TREATED WITH TREMFYA®

In two medical studies at 24 weeks, the level of fatigue was assessed using a questionnaire to measure self-reported tiredness, weakness, and difficulty conducting usual activities due to fatigue. Patients were asked to indicate their level of fatigue in the seven days prior.

SELECTED IMPORTANT SAFETY INFORMATION

TREMFYA® is not for everyone; only your doctor can decide if it's right for you. Do not use if you are allergic to TREMFYA®. TREMFYA® is a prescription medicine that may cause serious side effects, including serious allergic reactions and infections. TREMFYA® affects your immune system. It may increase your risk of infections and lower your ability to fight them. Please read the [Important Safety Information](#) on page 5 and the [Medication Guide](#) for TREMFYA® to learn more about these and other risks for TREMFYA®. Discuss any questions you have with your doctor.

UNCOVER TREMFYA® DOSING

TREMFYA® is a 100 mg injection given under your skin at weeks 0 and 4, and then every 8 weeks. Following the 2 starter doses, that's 6 doses a year.

FIRST 52 WEEKS OF THERAPY

TREMFYA®
2 STARTER DOSES AT
WEEK 0 AND WEEK 4

6

MAINTENANCE DOSES
1 dose every 8 weeks

INJECTION SUPPORT



You may enroll in the **TREMFYA® Injection Training Support Program** after your doctor has provided you with initial self-injection training. The program provides additional live training with an Injection Training Specialist, either in person* or over the phone.

Visit [TREMFYA.com/Patient-Support](https://www.tremfya.com/Patient-Support) or ask your doctor for more information.

This program is limited to education about your Janssen medication, its administration, and/or the condition it treats. It is not intended to provide medical advice, replace a treatment plan you receive from your doctor or nurse, or serve as a reason for you to start or stay on treatment.

*In-person live training may not be available in all areas.

SELECTED IMPORTANT SAFETY INFORMATION

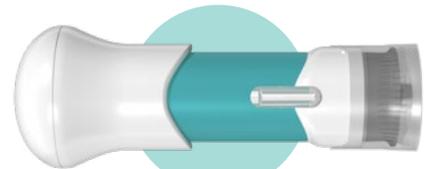
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UNCOVER THE ONE-PRESS IN 3D

A patient-controlled self-injection device that was created with you in mind. The One-Press device features an easy-to-hold design. And if you're not a fan of seeing needles, you're in luck: One-Press has a hidden needle feature so you won't see the needle unless you go looking for it.



The One-Press Self-Injector was awarded the Arthritis Foundation's Ease of Use seal after being tested by patients with psoriatic arthritis.



Not actual size

Your first self-injection should be performed at your doctor's office so he or she can show you the right way to give yourself the injections. After this training, and with your healthcare provider's approval, you may be able to inject at home.

Please read the Instructions for Use in the back pocket before using your TREMFYA® One-Press injector and each time you get a refill.

SCAN THE CODE WITH YOUR SMARTPHONE TO BEGIN



1. Point your smartphone camera at the marker to get started
2. Rotate and zoom in on the One-Press with your fingers
3. Tap "See It Live" and experience the One-Press in 3D

UNCOVER SUPPORT OPTIONS

Once you and your doctor have decided TREMFYA® is right for you, Janssen CarePath will help you find the resources you may need to get started and stay on track. We will give you information on your insurance coverage, potential out-of-pocket costs, and treatment support, and identify options that may help make your treatment more affordable.

CALL A JANSSEN CAREPATH CARE COORDINATOR TODAY

Visit [JanssenCarePath.com/TREMFYA](https://www.JanssenCarePath.com/TREMFYA). Create a Janssen CarePath Account at [MyJanssenCarePath.com](https://www.MyJanssenCarePath.com) where you can learn about your insurance coverage for TREMFYA®; if eligible, enroll in the Janssen CarePath Savings Program and manage program benefits; and sign up for treatment support.

Need help? Call a Janssen CarePath Care Coordinator today at **877-CarePath (877-227-3728)**, Monday–Friday, 8:00 AM to 8:00 PM ET.

START WITH TREMFYA®



The **TREMFYA® So Simple Trial Program** provides you with your first dose of TREMFYA® so you and your doctor can determine if it is right for you.

- \$0 COST
- Shipment can be authorized within 3 business days after a prescription*
- This trial program is open to patients who have commercial insurance, government coverage, or no insurance coverage. However, there is no guarantee of continuous accessibility after the program ends.

This program is for medication only. Terms expire at the end of each calendar year and may change.

*Shipment date is dependent on you opting in to the program and scheduling shipment.

SAFE RETURNS®

If you've received approval from your doctor to inject at home and have been properly trained, you can use Safe Returns® as a simple, safe, and convenient way to dispose of your used syringes or the One-Press injector—at no cost to you. When you sign up, you'll receive an FDA-approved Safe Returns® plastic disposal container in the mail, along with easy-to-follow mail-back instructions.

STAY ON TREMFYA®

➤ **If you have commercial insurance and your insurance coverage for TREMFYA® is approved**

JANSSEN CAREPATH SAVINGS PROGRAM*

- Eligible patients **pay \$5 per injection**
- \$20,000 maximum program benefit per calendar year
- See full program requirements at [TREMFYA.JanssenCarePathSavings.com](https://www.TREMFYA.JanssenCarePathSavings.com)

ENROLL TODAY

Text "Savings" to 56011

Enroll and receive an electronic Savings Program card that can be saved to your digital wallet on your iPhone or Android device

- OR -

Check your eligibility and enroll at

[MyJanssenCarePath.com/express](https://www.MyJanssenCarePath.com/express)

You will not have a Janssen CarePath account and will not be able to view and manage your Savings Program benefits.

➤ **If your commercial insurance coverage is delayed (>5 business days) or denied**

JANSSEN LINK†

- You can receive TREMFYA® at no cost until you receive insurance coverage approval
- See full program requirements at [JanssenCarePath.com/Patient/TREMFYA/Starting-Treatment](https://www.JanssenCarePath.com/Patient/TREMFYA/Starting-Treatment)

†Both the Janssen CarePath Savings Program and Janssen Link are unavailable to individuals who use any state or federal government-funded healthcare program to cover a portion of medication cost, such as Medicare, Medicaid, TRICARE, Department of Defense, or Veterans Administration. These programs are for medication only. Terms expire at the end of each program year and may change.

➤ **If you are using a government-funded healthcare program or have no insurance coverage**

- Janssen CarePath can provide information about other resources that may be able to help with your out-of-pocket medication costs
- Visit [JanssenPrescriptionAssistance.com](https://www.JanssenPrescriptionAssistance.com) for more information about affordability programs

Please read the [Important Safety Information for TREMFYA®](#) on page 5.



IMPORTANT SAFETY INFORMATION

What is the most important information I should know about TREMFYA®?

TREMFYA® is a prescription medicine that may cause serious side effects, including:

- **Serious Allergic Reactions.** Stop using TREMFYA® and get emergency medical help right away if you develop any of the following symptoms of a serious allergic reaction:
 - fainting, dizziness, feeling lightheaded (low blood pressure)
 - trouble breathing or throat tightness
 - chest tightness
 - swelling of your face, eyelids, lips, mouth, tongue or throat
 - skin rash, hives
 - itching
- **Infections.** TREMFYA® may lower the ability of your immune system to fight infections and may increase your risk of infections. Your healthcare provider should check you for infections and tuberculosis (TB) before starting treatment with TREMFYA® and may treat you for TB before you begin treatment with TREMFYA® if you have a history of TB or have active TB. Your healthcare provider should watch you closely for signs and symptoms of TB during and after treatment with TREMFYA®.

Tell your healthcare provider right away if you have an infection or have symptoms of an infection, including:

- fever, sweats, or chills
- muscle aches
- weight loss
- cough
- warm, red, or painful skin or sores on your body different from your psoriasis
- diarrhea or stomach pain
- shortness of breath
- blood in your phlegm (mucus)
- burning when you urinate or urinating more often than normal

Do not take TREMFYA® if you have had a serious allergic reaction to guselkumab or any of the ingredients in TREMFYA®.

Before using TREMFYA®, tell your healthcare provider about all of your medical conditions, including if you:

- have any of the conditions or symptoms listed in the section **“What is the most important information I should know about TREMFYA®?”**

- have an infection that does not go away or that keeps coming back.
- have TB or have been in close contact with someone with TB.
- have recently received or are scheduled to receive an immunization (vaccine). You should avoid receiving live vaccines during treatment with TREMFYA®.
- are pregnant or plan to become pregnant. It is not known if TREMFYA® can harm your unborn baby.
- are breastfeeding or plan to breastfeed. It is not known if TREMFYA® 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.

What are the possible side effects of TREMFYA®?

TREMFYA® may cause serious side effects. See “What is the most important information I should know about TREMFYA®?”

The most common side effects of TREMFYA® include:

upper respiratory infections, headache, injection site reactions, joint pain (arthralgia), diarrhea, stomach flu (gastroenteritis), fungal skin infections, herpes simplex infections, and bronchitis.

These are not all the possible side effects of TREMFYA®. Call your doctor for medical advice about side effects.

Use TREMFYA® exactly as your healthcare provider tells you to use it.

Please read the full [Prescribing Information](#), including [Medication Guide for TREMFYA®](#), and discuss any questions that you have with your doctor.

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.

**UNCOVER MORE
ABOUT TREMFYA®**

**IN MODERATE TO SEVERE
PLAQUE PSORIASIS**





START WITH A CONFIDENT CONVERSATION

Speaking up for yourself can be hard. That's why we want to help you have a bolder, clearer conversation with your doctor. Don't forget, you and your doctor have the same goal. Watch Kate and Joe have their confident conversation about their moderate to severe plaque psoriasis.



"My doctor and I decided TREMFYA® was right for me and I haven't looked back since."

- Kate, TREMFYA® patient

Individual results may vary.

Visit [TREMFYA.com/straighttalk](https://www.tremfya.com/straighttalk) to learn more about Kate and other patients' stories.

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