LET'S TALK ABOUT CLEARER SKIN FOR ADULTS WITH MODERATE TO SEVERE PLAQUE PSORIASIS

In clinical studies at week 16, at least 7 out of 10 patients saw at least 90% clearer skin, and more than 8 out of 10 were rated cleared or almost cleared.

In a study, 8 out of 10 TREMFYA® patients saw 90% clearer skin at 3 years.

Individual results may vary.

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).

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 affects your immune system and may cause serious side effects, including infections, and may increase your risk of infections. Serious allergic reactions may occur. Please read the Important Safety Information on page 9 and the Medication Guide for TREMFYA® to learn more about these and other risks for TREMFYA®. Discuss any questions you have with your doctor.
RESULTS WITH TREMFYA®

If you’ve tried it all for your moderate to severe plaque psoriasis—like steroids, topical creams, or phototherapy—and still didn’t get the results you wanted, then TREMFYA® may be what you’re waiting for.

In clinical studies at week 16, at least 7 out of 10 patients saw at least 90% clearer skin, and more than 8 out of 10 were rated cleared or almost cleared.

In a study, 8 out of 10 TREMFYA® patients saw 90% clearer skin at 3 years.

GET CLEARER AND STAY CLEARER

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 while on TREMFYA®. Individual results may vary.

TREMFYA® AND SCALP PSORIASIS

In clinical studies at 16 weeks, 4 out of 5 people with scalp psoriasis were rated cleared or almost cleared of their scalp psoriasis after just 3 doses.

Simply put, TREMFYA® can give you results you can see and feel.

RESULTS: WEEK 16 AND WEEK 48

SELECTED IMPORTANT SAFETY INFORMATION

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**IN RESULTS FROM 2 CLINICAL STUDIES**

**TREMFYA® WORKS BETTER THAN HUMIRA®**

In clinical studies, TREMFYA® worked better than Humira® (adalimumab) at providing clearer skin.

HUMIRA® (adalimumab) is a registered trademark of AbbVie Inc. Results reflective of patients studied in the U.S. and Canada.

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### AT WEEK 16

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<thead>
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### AT WEEK 24

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<td><strong>HUMIRA®</strong></td>
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<td><strong>HUMIRA®</strong></td>
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**SELECTED IMPORTANT SAFETY INFORMATION**

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TREMFYA® IS THE 1ST FDA-APPROVED MEDICATION TO SELECTIVELY BLOCK IL-23
TREMFYA® is the first approved biologic to selectively block interleukin-23 (IL-23), one of the key proteins thought to be responsible for plaque psoriasis.

What is a biologic?
Biologics are medicines that work to target an overactive immune system, a possible cause of plaque psoriasis. Most biologics for plaque psoriasis, including TREMFYA®, are given as an injection under your skin. It is administered by a healthcare provider or self-injected only after proper training.

FEWER SYMPTOMS
More people using TREMFYA® were free from symptoms, including itching, skin tightness, burning, stinging, and pain at 24 weeks compared with those using HUMIRA®.

FEWER DOSES
With only 1 maintenance dose every 8 weeks (after 2 starter doses at weeks 0 and 4), that’s about 2 whole months that you don’t have to think about your treatment schedule.

TREMFYA® is administered as a single 100 mg injection under the skin at weeks 0 and 4, and then every 8 weeks. It is administered by a healthcare provider or self-injected only after proper training.

HUMIRA® is a prescription medicine used to treat adults with moderate to severe chronic plaque psoriasis who are ready for systemic therapy or phototherapy, and are under the care of a doctor who will decide if other systemic therapies are less appropriate.

HUMIRA® is a registered trademark of AbbVie Inc.

SELECTED IMPORTANT SAFETY INFORMATION
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INJECTION YOU CAN REALLY CLICK WITH

TREMFYA® is the 1st FDA-approved medication to selectively block interleukin-23 (IL-23) to offer the One-Press patient-controlled injector. Let us tell you about its features:

▷ Don’t love needles? Don’t see needles. One-Press features a hidden needle, so you don’t see it unless you go looking for it.

▷ It comes with an easy-to-hold design and lets you inject at the speed you’re comfortable with.

▷ You’ll know your injection is done when you hear the “click” and then you’ll see the yellow band when the needle guard is locked.

Ask your dermatologist if you’re ready to click with One-Press. (But if you prefer a prefilled injector, don’t worry, that’s still available too.)

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

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

SELECTED IMPORTANT SAFETY INFORMATION

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A SUCCESSFUL CONVERSATION STARTS WITH YOU

Communicating openly with your dermatologist is the first step to better managing your moderate to severe plaque psoriasis.

Your dermatologist is relying on you to tell him or her about your experience with plaque psoriasis. Answer a few simple questions about your symptoms and how they affect you using our Dermatologist Conversation Starter. Then bring it to your provider so you can have a more open discussion about what you’re looking for in your new treatment for moderate to severe plaque psoriasis.

DESCRIBE YOUR EXPERIENCE

How satisfied are you with your current levels of symptom control?

- Very satisfied
- Somewhat satisfied
- Neutral
- Somewhat unsatisfied
- Very unsatisfied

How often do you have symptoms of plaque psoriasis?

- Every other week
- Every month
- Every few months
- Other ________

In your day-to-day life, how bothered are you by your moderate to severe plaque psoriasis symptoms (eg, itching, skin tightness, burning, stinging, and pain)?

- NOT AT ALL BOTHERED
- SLIGHTLY BOTHERED
- BOTHERED

TREMFLYA® is not for everyone. You and your dermatologist can decide if TREMFYA® is right for you. Individual results may vary.

SELECTED IMPORTANT SAFETY INFORMATION

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YOU HAVE A PRESCRIPTION, WHAT’S NEXT?
This may be a good time to ask if you qualify for the PsO Simple Trial Program, offered by Janssen CarePath. Ask your healthcare provider how you can quickly get started to evaluate the safety and efficacy of TREMFYA® with a trial at no cost. At the conclusion of the trial program, you and your healthcare provider can decide whether to continue.

SUPPORT FOR YOU: THE JANSSEN CAREPATH TEAM
Janssen CarePath provides information you may need to help you get started and stay on track once your doctor has prescribed TREMFYA®.

Your Janssen CarePath Care Coordinator can:
• Review your health plan benefits to help answer questions about insurance coverage
• Work with your specialty pharmacy to process and deliver your TREMFYA® medication to your doctor’s office or home
• Connect you with injection support, see below for more information

INSURANCE COVERAGE
If you are starting TREMFYA®, your dermatologist’s office will check to see what your insurance covers. No matter what type of insurance you have, Janssen CarePath can help explain your medication insurance coverage and benefits.

COST SUPPORT OPTIONS

Janssen CarePath Savings Program
Eligible patients using commercial or private insurance pay $5 per injection for their TREMFYA® medication costs, with a $20,000 maximum program benefit each calendar year.
• Not valid for patients using Medicare, Medicaid, or other government-funded programs to pay for their medications.
Terms expire at the end of each calendar year and may change.
There is no income requirement. See full eligibility requirements at Tremfya.JanssenCarepathSavings.com

Enroll at MyJanssenCarePath.com
Information and resources regarding your insurance coverage, affordability options, and treatment support are provided to you by third-party service providers for Janssen CarePath, which is operated by Johnson & Johnson Health Care Systems Inc. on behalf of Janssen Biotech, Inc. Janssen CarePath is not available to patients participating in the Patient Assistance Program offered by Johnson & Johnson Patient Assistance Foundation. Janssen CarePath provides information to you regarding whether the treatment is covered by your insurance or third-party payer. This information and assistance is made available as a convenience to you, and there is no requirement that you or your HCP use any Janssen product in exchange. While Janssen CarePath attempts to provide correct information, they assume no responsibility for and do not guarantee the quality, scope, or availability of the information and assistance provided.

INJECTION SUPPORT
A NURSE IS JUST A PHONE CALL AWAY
Even after you’ve been trained by a healthcare professional to give yourself an injection, you may still have questions. Janssen Nurse Support® can answer questions about giving yourself an injection at home, preparing your injection site prior to self-injecting, and properly disposing of your used syringe or One-Press injector. To contact Janssen Nurse Support, call 877-CarePath (877-227-3728), Monday–Friday, 9 AM–8 PM ET. At all other times, a nurse will typically return your call in about 15 minutes.

CONVENIENT DISPOSAL OF YOUR USED INJECTOR
Dispose of your prefilled syringe or One-Press injector in a puncture-resistant container (FDA-cleared sharps disposal container). Do not reuse needles or syringes. Safe Returns® is a simple, safe, and convenient way to dispose of your used syringes—at no additional cost to you. When you sign up, you’ll receive a Safe Returns® plastic disposal container in the mail, along with simple mail-back instructions.

STORING YOUR PREFILLED SYRINGE OR ONE-PRESS INJECTOR
Whether you’re using One-Press or a prefilled syringe, store it in the original carton to protect it from light and physical damage until time of use. Store TREMFYA® in your refrigerator at 36˚F to 46˚F. Do not freeze. Do not shake. Discard any unused portion. TREMFYA® is sterile and preservative free. Keep out of reach of children.

SUPPORT AND TREATMENT REMINDERS
Support is available by calling Janssen CarePath or visiting JanssenCarePath.com. Also, we understand how important it is for you to take your TREMFYA® medication just as your doctor prescribed. That’s why we created tools to help you stay on track with your TREMFYA® treatment, like tools to remind you when your next dose of TREMFYA® is due.

To sign up for treatment reminders or cost support, log into your personal Janssen CarePath Account at MyJanssenCarePath.com or call: 877-CarePath (877-227-3728), Monday–Friday, 8 AM–8 PM ET.

*Janssen Nurse Support 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.

Please read the Important Safety Information for TREMFYA® on page 9.
THE TREMFYA® TIMELINE: YEAR 1
What can you expect in your first year of treatment with TREMFYA®? From getting your prescription filled, to learning about the possible results, this timeline has the information you need.

TREMFYA® is administered as a single 100 mg injection under the skin at weeks 0 and 4, and then every 8 weeks. It is administered by a healthcare provider or self-injected only after proper training.

I HAVE A PRESCRIPTION. WHAT HAPPENS NOW?
After your healthcare provider prescribes TREMFYA®, his or her office coordinator will confirm your insurance coverage.

MEET JANSSEN CAREPATH
Explore our support services and see everything we can do to help you. For more on Janssen CarePath, see page 7.

THE FIRST STARTER DOSE
Your first injection may be performed at your dermatologist’s office, so he or she can make sure you’re properly trained to self-inject at home.

START

THE SECOND STARTER DOSE
You’re on your way!

THE FIRST MAINTENANCE DOSE
Next one in 8 weeks!

GET CLEAR
In clinical studies, after just 3 doses, 7 out of 10 patients saw at least 90% clearer skin, and more than 8 out of 10 were rated as cleared or almost cleared at 16 weeks in clinical studies. Individual results may vary.

CONTINUED ON THE RIGHT

WEEK 4

WEEK 12

WEEK 16

WEEK 20

WEEK 28

WEEK 36

WEEK 44

WEEK 48

WEEK 52

THE SECOND MAINTENANCE DOSE
Keep up the great work!

THE THIRD MAINTENANCE DOSE
Keep on going!

THE FOURTH MAINTENANCE DOSE
Be sure to keep in touch with your dermatologist or Janssen CarePath with any questions.

THE FIFTH MAINTENANCE DOSE
You’re nearly done with year 1.

STAY CLEARER
In a clinical study, the majority of patients who saw at least 90% clearer skin with TREMFYA® at week 28 continued to see clearer skin all the way through 48 weeks. Individual results may vary.

THE SIXTH MAINTENANCE DOSE
You’ve finished your first year with TREMFYA®!

WE’RE ALWAYS HERE TO HELP
Janssen CarePath Care Coordinators are here to help as you continue your journey with TREMFYA®.

SELECTED IMPORTANT SAFETY INFORMATION
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IMPORTANT SAFETY INFORMATION

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

TREMFYA® may cause serious side effects, including infections.
TREMFYA® is a prescription medicine that 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®?”

Serious Allergic Reactions
Stop using TREMFYA® and get emergency medical help right away if you have any of the following symptoms of a serious allergic reaction: feel faint, swelling of your face, eyelids, lips, mouth, tongue or throat, trouble breathing or throat tightness, chest tightness, or skin rash, hives.

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, and herpes simplex infections.

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.
ASK YOUR DERMATOLOGIST ABOUT RESULTS WITH TREMFYA®

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SELECTED IMPORTANT SAFETY INFORMATION

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