Tremfya® (guselkumab)

First in Class:

TREMFYA® is the first selective IL-23 inhibitor approved for adults with moderate to severe plaque psoriasis or adults with active psoriatic arthritis

Program Objectives

- Review TREMFYA® clinical efficacy and safety data from pivotal trials in moderate to severe plaque psoriasis, including 252-week open-label extension data from **VOYAGE 1**
- Review TREMFYA® clinical efficacy and safety data from pivotal trials in moderate to severe plaque psoriasis, including data in subpopulations
- Describe clinical efficacy and safety data from **DISCOVER 1** and **DISCOVER 2** trials in patients with active psoriatic arthritis, including 112-week open-label extension data from **DISCOVER 2**

IL = interleukin.

You are cordially invited to attend a promotional speaker program, sponsored by Janssen Biotech, Inc.

PRESENTED BY

G. Michael Lewitt, MD
Dermatologist
Illinois Dermatology Institute
Chicago, IL

Gordon Lam, MD, FACR
Medical Director
Northern Region Research Center
at Atrium Health
Concord, NC

DATE

Tuesday, May 3, 2022
7:00 PM CT - 9:00 PM CT

LOCATION

Virtual Product Theater

Chicago Dermatological Society CDS 1901

REGISTER at www.MyDomeProgramRegistration.com

Enter Meeting Code: 2022-00850

Please note that your e-mail will be required for registration. The information you provide will only be used to facilitate your attendance at this program. If you have questions about this program, please contact your Janssen Biotech, Inc., representative. We look forward to your participation in this informative discussion. SPONSORED BY JANSSEN BIOTECH, INC.

INDICATIONS

TREMFYA® is indicated for the treatment of adults with moderate to severe plaque psoriasis who are candidates for systemic therapy or phototherapy.

TREMFYA® is indicated for the treatment of adults with active psoriatic arthritis.

DOSAGE AND ADMINISTRATION

TREMFYA® is administered as a 100 mg subcutaneous injection once every 8 weeks, after starter doses at weeks 0 and 4. In active psoriatic arthritis, TREMFYA® may be administered alone or in combination with a cDMARD (e.g., methotrexate).

TREMFYA® is intended for use under the guidance and supervision of a physician. Patients may self-inject with TREMFYA® after physician approval and proper training.

cDMARD, conventional disease-modifying antirheumatic drug.

SELECTED IMPORTANT SAFETY INFORMATION

TREMFYA® is contraindicated in patients with a history of serious hypersensitivity reaction to guselkumab or to any of the excipients. Serious hypersensitivity reactions, including anaphylaxis, have been reported. TREMFYA® may increase the risk of infection. Instruct patients to seek medical advice if signs or symptoms of clinically important chronic or acute infection occur. If a clinically important or serious infection develops, discontinue TREMFYA® until infection resolves. Evaluate for tuberculosis before treating with TREMFYA®. Avoid use of live vaccines in patients treated with TREMFYA®. Please see related and other Important Safety Information on the reverse.

IMPORTANT SAFETY INFORMATION

CONTRAINDICATIONS

TREMFYA® is contraindicated in patients with a history of serious hypersensitivity reaction to guselkumab or to any of the excipients.

WARNINGS AND PRECAUTIONS

Hypersensitivity Reactions

Serious hypersensitivity reactions, including anaphylaxis, have been reported with postmarket use of TREMFYA®. Some cases required hospitalization. If a serious hypersensitivity reaction occurs, discontinue TREMFYA® and initiate appropriate therapy.

Infections

TREMFYA® may increase the risk of infection. Treatment with TREMFYA® should not be initiated in patients with a clinically important active infection until the infection resolves or is adequately treated.

Consider the risks and benefits of treatment prior to prescribing TREMFYA® in patients with a chronic infection or a history of recurrent infection. Instruct patients receiving TREMFYA® to seek medical help if signs or symptoms of clinically important chronic or acute infection occur. If a patient develops a clinically important or serious infection, or is not responding to standard therapy, closely monitor and discontinue TREMFYA® until the infection resolves.

Pre-Treatment Evaluation for Tuberculosis (TB)

Evaluate patients for TB infection prior to initiating treatment with TREMFYA®. Initiate treatment of latent TB prior to administering TREMFYA®. Monitor patients for signs and symptoms of active TB during and after TREMFYA® treatment. Do not administer TREMFYA® to patients with active TB infection.

Immunizations

Prior to initiating TREMFYA®, consider completion of all age-appropriate immunizations according to current immunization guidelines. Avoid use of live vaccines in patients treated with TREMFYA®.

ADVERSE REACTIONS

Most common (≥1%) adverse reactions associated with TREMFYA® include upper respiratory infections, headache, injection site reactions, arthralgia, bronchitis, diarrhea, gastroenteritis, tinea infections, and herpes simplex infections.

The overall safety profile observed in patients with psoriatic arthritis is generally consistent with the safety profile in patients with plaque psoriasis, with the addition of bronchitis and neutrophil count decreased.

Please read the enclosed full Prescribing Information and Medication Guide for TREMFYA®. Provide the Medication Guide to your patients and encourage discussion.

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DISCLOSURE

In adherence with PhRMA guidelines, spouses or other guests are not permitted to attend company-sponsored programs.

For all attendees, please be advised that information such as your name and the value and purpose of any educational item, meal, or other items of value you receive may be publicly disclosed. If you are licensed in any state or other jurisdiction, or are an employee or contractor of any organization or governmental entity, that limits or prohibits meals from pharmaceutical companies, please identify yourself so that you (and we) are able to comply with such requirements.

Please note that the company prohibits the offering of gifts, gratuities, or meals to federal government employees/officials. Thank you for your cooperation.

The consultant is a paid speaker for Janssen Biotech, Inc. The speaker is presenting on behalf of Janssen and must present information in compliance with FDA requirements applicable to Janssen.

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