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Section: Prescription Drugs Effective Date: October 1, 2024

Subsection: Topical Products Original Policy Date: September 30, 2022

Subject: Spevigo Page: 1 of 6

Last Review Date: September 6, 2024

Spevigo

Description

Spevigo (spesolimab-sbzo)

Background

Spevigo (spesolimab-sbzo) is a humanized monoclonal immunoglobulin G1 antibody that inhibits interleukin-36 (IL-36) signaling by specifically binding to the IL36R. This prevents the subsequent activation of the IL36R and downstream activation of pro-inflammatory and profibrotic pathways (1).

Regulatory Status

FDA-approved indication: Spevigo is an interleukin-36 receptor antagonist indicated for the treatment of generalized pustular psoriasis (GPP) in adults and pediatric patients 12 years of age and older and weighing at least 40 kg (1).

Spevigo may increase the risk of infections. Treatment with Spevigo is not recommended for use in patients with any clinically important active infection until the infection resolves or is adequately treated (1).

Patients should be evaluated for tuberculosis (TB) infection prior to initiating treatment with Spevigo. Spevigo should not be administered to patients with active TB infection. Anti-TB therapy should be considered prior to initiating Spevigo in patients with latent TB or a history of TB in whom an adequate course of treatment cannot be confirmed (1).

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Spevigo-associated hypersensitivity reactions may include immediate reactions such as anaphylaxis and delayed reactions such as drug reaction with eosinophilia and systemic symptoms (DRESS) (1).

The use of live vaccines with Spevigo should be avoided (1).

The safety and effectiveness of Spevigo in pediatric patients less than 12 years of age and weighing less than 40kg have not been established (1).

Related policies

Policy

This policy statement applies to clinical review performed for pre-service (Prior Approval, Precertification, Advanced Benefit Determination, etc.) and/or post-service claims.

Spevigo may be considered **medically necessary** if the conditions indicated below are met.

Spevigo may be considered **investigational** for all other indications.

Prior-Approval Requirements

Age 12 years of age or older

Diagnosis

Patient must have the following:

- 1. Generalized pustular psoriasis (GPP)
 - a. GPP flares are of moderate-to-severe intensity (e.g., at least 5% of body surface area covered with erythema and the presence of pustules)

AND ALL of the following:

- Patient has had an inadequate treatment response, intolerance, or contraindication to **ONE** of the following:
 - a. Methotrexate
 - b. Cyclosporine

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c. Oral retinoid

- 2. Patient weight ≥ 40kg
- 3. Prescriber agrees to monitor for hypersensitivity reactions, including drug reaction with eosinophilia and systemic symptoms (DRESS)
- 4. Result for latent TB infection is negative **OR** result was positive for latent TB and patient completed treatment (or is receiving treatment) for latent TB
- 5. Absence of active infection (including tuberculosis)
- 6. **NOT** used in combination with any other biologic DMARD or targeted synthetic DMARD (see Appendix 1)
- 7. **NOT** given concurrently with live vaccines

Prior – Approval Renewal Requirements

Age 12 years of age or older

Diagnosis

Patient must have the following:

- 1. Generalized pustular psoriasis (GPP)
 - a. Improvement or stabilization of patient's condition (e.g., reduction in the frequency or severity of flares)

AND ALL of the following:

- 1. Patient weight ≥ 40kg
- 2. Prescriber agrees to monitor for hypersensitivity reactions, including drug reaction with eosinophilia and systemic symptoms (DRESS)
- 3. Absence of active infection (including tuberculosis)
- 4. **NOT** used in combination with any other biologic DMARD or targeted synthetic DMARD (see Appendix 1)
- 5. **NOT** given concurrently with live vaccines

Policy Guidelines

Pre - PA Allowance

None

Prior - Approval Limits

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Quantity 4 IV vials + 26 SC syringes

Duration 12 months

Prior - Approval Renewal Limits

Same as above

Rationale

Summary

Spevigo is an interleukin-36 receptor antagonist that is indicated for the treatment of generalized pustular psoriasis (GPP) and for prevention of flares. Spevigo may cause hypersensitivity reactions including DRESS. Spevigo should not be given to patients with clinically important active infections, including TB. The safety and effectiveness of Spevigo in pediatric patients less than 12 years of age and weighing less than 40kg have not been established (1).

Prior approval is required to ensure the safe, clinically appropriate, and cost-effective use of Spevigo while maintaining optimal therapeutic outcomes.

References

- 1. Spevigo [package insert]. Ridgefield, CT: Boehringer Ingelheim Pharmaceuticals, Inc.; March 2024.
- 2. Menter A, Gelfand JM, Connor C, et al. Joint AAD-NPF guidelines of care for the management of psoriasis with systemic nonbiologic therapies. J Am Acad Dermatol. 2020;82:1445-86.

| Policy History | |
|----------------|---|
| Date | Action |
| September 2022 | Addition to PA |
| December 2022 | Annual review |
| September 2023 | Annual review |
| March 2024 | Annual review |
| April 2024 | Per PI update, lowered age requirement to 12 years and older weighing 40 kg or more. Added subcutaneous dosage form. Changed indication to just GPP. Added renewal requirements |
| June 2024 | Annual review |
| September 2024 | Annual review |
| Keywords | |

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This policy was approved by the FEP® Pharmacy and Medical Policy Committee on September 6, 2024 and is effective on October 1, 2024.

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Appendix 1 - List of DMARDs

Biological disease-modifying antirheumatic drugs (DMARDs)

| Diological disease-modifying antimedinatic drugs (DMARDS) | | | |
|---|-------------------------------------|--|--|
| Generic Name | Brand Name | | |
| abatacept | Orencia | | |
| adalimumab | Humira | | |
| anakinra | Kineret | | |
| brodalumab | Siliq | | |
| certolizumab | Cimzia | | |
| etanercept | Enbrel | | |
| golimumab | Simponi/Simponi Aria | | |
| guselkumab | Tremfya | | |
| infliximab | Remicade/Avsola/Inflectra/Renflexis | | |
| ixekizumab | Taltz | | |
| risankizumab-rzaa | Skyrizi | | |
| rituximab | Rituxan/Riabni/Ruxience/Truxima | | |
| sarilumab | Kevzara | | |
| secukinumab | Cosentyx | | |
| spesolimab-sbzo | Spevigo | | |
| tildrakizumab-asmn | Ilumya | | |
| tocilizumab | Actemra | | |
| ustekinumab | Stelara | | |
| vedolizumab | Entyvio | | |

Targeted synthetic disease-modifying antirheumatic drugs (DMARDs)

| Generic Name | Brand Name |
|-----------------|------------|
| apremilast | Otezla |
| baricitinib | Olumiant |
| deucravacitinib | Sotyktu |
| tofacitinib | Xeljanz/XR |
| upadactinib | Rinvoq |