

**ustekinumab intravenous****Medical Benefit Drug Policy**

ustekinumab (Stelara)  
ustekinumab-aekn (Selarsdi)  
ustekinumab-aauz (Otulfi)  
ustekinumab-auub (Wezlana)  
ustekinumab-kfce (Yesintek)  
ustekinumab-stba (Steqeyma)  
ustekinumab-srlf (Imuldosa)  
ustekinumab-ttwe (Pyzchiva)

Place of Service

Home Infusion Administration  
Infusion Center Administration  
Office Administration  
Outpatient Facility Infusion Administration

**Drug Details**

**USP Category:** IMMUNOLOGICAL AGENTS

**Mechanism of Action:** Interleukin (IL)-12/IL-23 Inhibitor

**HCPCS:**

J3357:Ustekinumab, for subcutaneous injection, 1 mg  
J3358:Ustekinumab, for intravenous injection, 1 mg  
Q5098:Injection, ustekinumab-srlf (imuldosa), biosimilar, 1 mg  
Q5099:Injection, ustekinumab-stba (steqeyma), biosimilar, 1 mg  
Q5100:Injection, ustekinumab-kfce (yesintek), biosimilar, 1 mg  
Q5137:Injection, ustekinumab-auub (wezlan), biosimilar, subcutaneous, 1 mg  
Q5138:Injection, ustekinumab-auub (wezlan), biosimilar, intravenous, 1 mg  
Q9996:Injection, ustekinumab-ttwe (pyzchiva), subcutaneous, 1 mg  
Q9997:Injection, ustekinumab-ttwe (pyzchiva), intravenous, 1 mg  
Q9998:Injection, ustekinumab-aekn (selarsdi), biosimilar, 1 mg  
Q9999:Injection, ustekinumab-aauz (otulfi), biosimilar, 1 mg

**How Supplied:**

130 mg single-dose vial

**Condition(s) listed in policy** *(see coverage criteria for details)*

- Crohn's Disease, Moderate to Severe
- Ulcerative Colitis, Moderate to Severe

Any condition not listed in this policy requires a review to confirm it is medically necessary. For conditions that have not been approved for intended use by the Food and Drug Administration (i.e., off-label use), the criteria outlined in the California Code of Regulations (CCR), Title 22, Section 51303 and 51313 must be met.

## Special Instructions and Pertinent Information

Provider must submit documentation (such as office chart notes, lab results or other clinical information) to ensure the member has met all medical necessity requirements.

The member's specific benefit may impact drug coverage. Other utilization management processes, and/or legal restrictions may take precedence over the application of this clinical criteria.

## Coverage Criteria

The following condition(s) require Prior Authorization/Preservice.

### Crohn's Disease, Moderate to Severe

Meets medical necessity if all the following are met:

1. Age is consistent with the FDA-approved indication (18 years and older)
2. *Effective 2/1/2026 and after, will require* Prescribed by or in consultation with a gastroenterologist
3. Not being used in combination with other targeted immunomodulators
4. Request for Imuldosa, Otulfi, Pyzchiva, Selarsdi, Stelara, Steqeyma, or Wezlana: Intolerable side effect or contraindication with preferred ustekinumab product (e.g. Yesintek) that is not expected with the requested ustekinumab drug

### Covered Doses:

Initial dosing: A single intravenous infusion using weight-based dosing

Weight Range (kg)	Recommended Dosage
Up to 55 kg	260 mg (2 vials)
Greater than 55 kg to 85 kg	390 mg (3 vials)
Greater than 85 kg	520 mg (4 vials)

Maintenance dosing: Drug is available as a 45 mg/90 mg prefilled syringe and a 45 mg vial that can be self-administered subcutaneously for maintenance. Self-administered products can be requested from your Pharmacy Benefit.

### Coverage Period:

Initial dosing: One-time intravenous dose

### ICD-10:

K50.00-K50.119, K50.80-K50.919

### Ulcerative Colitis, Moderate to Severe

Meets medical necessity if all the following are met:

1. Age is consistent with the FDA-approved indication (18 years and older)

2. **Effective 2/1/2026 and after, will require** Prescribed by or in consultation with a gastroenterologist
3. Not used in combination with a targeted immunomodulator
4. Request for Imuldosa, Otulfi, Pyzchiva, Selarsdi, Stelara, Steqeyma, or Wezlana: Intolerable side effect or contraindication with preferred ustekinumab product (e.g. Yesintek) that is not expected with the requested ustekinumab drug

#### Covered Doses:

Initial dosing: A single intravenous infusion using weight-based dosing

Weight Range (kg)	Recommended Dosage
Up to 55 kg	260 mg (2 vials)
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Maintenance dosing: Drug is available as a 45 mg/90 mg prefilled syringe and a 45 mg vial that can be self-administered subcutaneously for maintenance. Self-administered products can be requested from your Pharmacy Benefit.

#### Coverage Period:

Initial dosing: One-time intravenous dose

#### ICD-10:

K51.0-K51.319, K51.5-K51.519, K51.80-K51.919

#### References

1. Imuldosa (ustekinumab-srlf) Prescribing Information. Accord BioPharma Inc., Raleigh, NC: 10/2024.
2. Otulfi (ustekinumab-aauz) Prescribing Information. Fresenius Kabi USA, LLC, Lake Zurich, IL: 9/2025.
3. Pyzchiva (ustekinumab-ttwe) Prescribing Information. Sandoz Inc. Princeton, NJ: 12/2024.
4. Selarsdi (ustekinumab-aekn) Prescribing Information. Teva Pharmaceuticals, Parsippany, NJ: 2/2025.
5. Stelara (ustekinumab) Prescribing Information. Janssen Biotech, Inc., Horsham, PA: 7/2020.
6. Steqeyma (ustekinumab-stba) Prescribing Information. CELLTRION USA, Inc., Jersey City, NJ: 12/2024.
7. Wezlana (ustekinumab-auub) Prescribing Information. Amgen Inc., Thousand Oaks, CA: 1/2025.
8. Yesintek (ustekinumab-kfce) Prescribing Information. Biocon Biologics Inc., Cambridge, MA: 11/2024.

## Review History

Date of Last Annual Review: 4Q2025

Changes from previous policy version:

- Added specialist requirement for patients with Crohn's disease and patients with ulcerative colitis (UC).

*Blue Shield of California Medication Policy to Determine Medical Necessity  
Reviewed by P&T Committee*