

**STELARA® Biosimilars: Food and Drug Administration (FDA)-Approved Ustekinumab Biosimilars for Prescribing**

**Preferred Biosimilar Options**

- PYZCHIVA™ (ustekinumab-twe)
- SELARSDI™ (ustekinumab-aekn)
- YESINTEK™ (ustekinumab-kfck)

These products are FDA-approved biosimilars to STELARA® (ustekinumab) and may be prescribed for eligible patients requiring ustekinumab therapy.

**Why Prescribe a Biosimilar?**

- Same mechanism of action as STELARA® (IL-12/IL-23 inhibitor)
- No clinically meaningful differences in safety, purity, or potency compared with STELARA®
- FDA-approved based on rigorous analytical, pharmacokinetic, and clinical evaluation
- Supports patient access to ustekinumab therapy

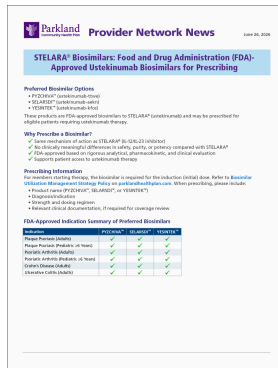
**Prescribing Information**

For members starting therapy, the biosimilar is required for the induction (initial) dose. Refer to [Biosimilar Utilization Management Strategy Policy](#) on [parklandhealthplan.com](http://parklandhealthplan.com). When prescribing, please include:

- Product name (PYZCHIVA™, SELARSDI™, or YESINTEK™)
- Diagnosis/indication
- Strength and dosing regimen
- Relevant clinical documentation, if required for coverage review

**FDA-Approved Indication Summary of Preferred Biosimilars**

Indication	PYZCHIVA™	SELARSDI™	YESINTEK™
Plaque Psoriasis (Adults)	✓	✓	✓
Plaque Psoriasis (Pediatric ≥6 Years)	✓	✓	✓
Psoriatic Arthritis (Adults)	✓	✓	✓
Psoriatic Arthritis (Pediatric ≥6 Years)	✓	✓	✓
Crohn's Disease (Adults)	✓	✓	✓
Ulcerative Colitis (Adults)	✓	✓	✓



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