

5.90.10

Section:	Prescription Drugs	Effective Date:	July 1, 2022
Subsection:	Topical Products	Original Policy Date:	January 1, 2015
Subject:	Luzu	Page:	1 of 4

Last Review Date: June 16, 2022

Luzu

Description

Luzu (Iuliconazole)

Background

Luzu (Iuliconazole) is a topical azole antifungal cream used to treat athlete's foot that is between the toes (interdigital tinea pedis), jock itch (tinea cruris), and ringworm (tinea corporis) caused by the organisms *Trichophyton rubrum* and *Epidermophyton floccosum*. Although the exact mechanism of action of Luzu is unknown, it appears to work by weakening the structure and function of the fungal cell membrane (1).

Regulatory Status

FDA-approved indications: Luzu is an azole antifungal indicated for the topical treatment of interdigital tinea pedis, tinea cruris, and tinea corporis caused by the organisms *Trichophyton rubrum* and *Epidermophyton floccosum* (1).

Safety and effectiveness of Luzu in pediatric patients have been established (1).

Related policies

Ecoza, Ertaczo, Exelderm, Topical Antifungals, Oxistat

Policy

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

Section:	Prescription Drugs	Effective Date:	July 1, 2022
Subsection:	Topical Products	Original Policy Date:	January 1, 2015
Subject:	Luzu	Page:	2 of 4

Luzu may be considered **medically necessary** in patients with interdigital tinea pedis, tinea cruris, or tinea corporis and if the conditions indicated below are met.

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

Prior-Approval Requirements

Diagnoses

Patient must have **ONE** of the following:

1. Interdigital Tinea Pedis
2. Tinea Cruris
3. Tinea Corporis

AND ALL of the following:

1. Laboratory and clinical documentation of **ONE** of the infections:
 - a. *Trichophyton rubrum*
 - b. *Epidermophyton floccosum*
2. Inadequate treatment response, intolerance, or contraindication to a legend topical antifungal therapy

Prior – Approval *Renewal* Requirements

Diagnoses

Patient must have **ONE** of the following:

1. Interdigital Tinea Pedis
2. Tinea Cruris
3. Tinea Corporis

AND ALL of the following:

1. Laboratory and clinical documentation of **ONE** of the infections:
 - a. *Trichophyton rubrum*
 - b. *Epidermophyton floccosum*
2. **NOT** used in a previously treated location within the last 12 months

Section:	Prescription Drugs	Effective Date:	July 1, 2022
Subsection:	Topical Products	Original Policy Date:	January 1, 2015
Subject:	Luzu	Page:	3 of 4

Policy Guidelines

Pre - PA Allowance

None

Prior - Approval Limits

Quantity 60 units

Duration 1 month

Prior – Approval *Renewal* Limits

Same as above

Rationale

Summary

Luzu (luliconazole) is a topical azole antifungal cream used to treat interdigital tinea pedis, tinea cruris, and tinea corporis caused by *Trichophyton rubrum* and *Epidermophyton floccosum*. Although the exact mechanism of action of Luzu is unknown, it appears to work by weakening the structure and function of the fungal cell membrane. Safety and effectiveness of Luzu in pediatric patients have been established (1).

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

References

1. Luzu [package Insert]. Bridgewater, NJ: Bausch Health US, LLC; April 2020.

Policy History

Date	Action
December 2014	Addition to PA and
March 2015	Annual editorial review and reference update
December 2016	Annual editorial review and reference update Added age limit to renewal criteria. Policy number changed from 5.14.10 to 5.90.10
September 2017	Annual editorial review
March 2018	Removal of age from initiation and renewal criteria
June 2018	Annual review

5.90.10

Section:	Prescription Drugs	Effective Date:	July 1, 2022
Subsection:	Topical Products	Original Policy Date:	January 1, 2015
Subject:	Luzu	Page:	4 of 4

September 2019	Annual review
December 2019	Annual review. Addition of quantity limit of 60 units
September 2020	Annual review
June 2021	Annual editorial review and reference update
June 2022	Annual review

Keywords

This policy was approved by the FEP® Pharmacy and Medical Policy Committee on June 16, 2022 and is effective on July 1, 2022.