

Pharmacy Medical Necessity Guidelines: Tranexamic Acid (Lysteda™)

Effective: January 12, 2021

Prior Authorization Required	√	Type of Review – Care Management	
Not Covered		Type of Review – Clinical Review	√
Pharmacy (RX) or Medical (MED) Benefit	RX	Department to Review	RXUM
<p>These pharmacy medical necessity guidelines apply to the following:</p> <p>Commercial Products</p> <ul style="list-style-type: none"> <input type="checkbox"/> Tufts Health Plan Commercial products – large group plans <input type="checkbox"/> Tufts Health Plan Commercial products – small group and individual plans <input type="checkbox"/> Tufts Health Freedom Plan products – large group plans <input type="checkbox"/> Tufts Health Freedom Plan products – small group plans • CareLinkSM – Refer to CareLink Procedures, Services and Items Requiring Prior Authorization <p>Tufts Health Public Plans Products</p> <ul style="list-style-type: none"> <input type="checkbox"/> Tufts Health Direct – A Massachusetts Qualified Health Plan (QHP) (a commercial product) <input checked="" type="checkbox"/> Tufts Health Together – MassHealth MCO Plan and Accountable Care Partnership Plans <input type="checkbox"/> Tufts Health RITogether – A Rhode Island Medicaid Plan 		<p>Fax Numbers: RXUM: 617.673.0988</p>	

Note: This guideline does not apply to Medicare Members (includes dual eligible Members).

OVERVIEW

FDA-APPROVED INDICATIONS

Tranexamic acid (Lysteda™) is an antifibrinolytic indicated for the treatment of cyclic heavy menstrual bleeding.

COVERAGE GUIDELINES

The plan may authorize coverage of tranexamic acid for Members when **all** the following criteria are met and limitations do not apply:

1. Member was started on tranexamic acid in an inpatient setting
- OR**
1. The Member has a diagnosis of hereditary angioedema (HAE)
- OR**
1. The Member has the diagnosis of heavy menstrual bleeding **AND** the Member tried and failed hormonal therapy, or the provider indicates clinical inappropriateness of hormonal therapy

Hormonal therapy may include:

1. Oral contraceptives
2. Levonorgestrel intrauterine device
3. Progesterone injection

LIMITATIONS

1. Requests for brand-name products, which have AB-rated generics, will be reviewed according to the Brand Name criteria.

CODES

None

REFERENCES

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7. Lethaby A, Augood C, Duckitt K, Farquhar C. Nonsteroidal anti-inflammatory drugs for heavy menstrual bleeding. Cochrane Database Syst Rev. 2007; (4):CD0004000.
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9. Grimes DA, Hubacher D, Lopez LM, Schulz KF. Non-steroidal anti-inflammatory drugs for heavy bleeding or pain associated with intrauterine-device use. Cochrane Database Syst Rev. 2006; (4):CD006034.
10. Wellington K, Wagstaff AJ. Tranexamic acid. A review of its use in the management of menorrhagia. Drugs. 2003; 63(13):1417-1433.
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APPROVAL HISTORY

April 14, 2011: Reviewed by Pharmacy & Therapeutics Committee.

Subsequent endorsement date(s) and changes made:

1. August 12, 2014: No changes
2. August 11, 2015: No changes
3. January 1, 2016: Administrative change to rebranded template.
4. August 9, 2016: Added "Requests for brand-name products, which have AB-rated generics, will be reviewed according to Non-Covered Medications criteria" to the limitations section of the policy.
5. May 9, 2017: Administrative update, Adding Tufts Health RITogether to the template
6. January 9, 2018: No changes.
7. January 8, 2019: Administrative changes made to template.
8. January 14, 2020: No changes.
9. July 14, 2020: Updated criteria to allow for members to be approved for tranexamic acid if Member was started on the agent in an inpatient facility or if the member has a diagnosis of hereditary angioedema.
10. January 12, 2021: No changes.

BACKGROUND, PRODUCT AND DISCLAIMER INFORMATION

Pharmacy Medical Necessity Guidelines have been developed for determining coverage for plan benefits and are published to provide a better understanding of the basis upon which coverage decisions are made. The plan makes coverage decisions on a case-by-case basis considering the individual member's health care needs. Pharmacy Medical Necessity Guidelines are developed for selected therapeutic classes or drugs found to be safe, but proven to be effective in a limited, defined population of patients or clinical circumstances. They include concise clinical coverage criteria based on current literature review, consultation with practicing physicians in the service area who are medical experts in the particular field, FDA and other government agency policies, and standards adopted by national accreditation organizations. The plan revises and updates Pharmacy Medical Necessity Guidelines annually, or more frequently if new evidence becomes available that suggests needed revisions.

For self-insured plans, coverage may vary depending on the terms of the benefit document. If a discrepancy exists between a Pharmacy Medical Necessity Guideline and a self-insured Member's benefit document, the provisions of the benefit document will govern.

Treating providers are solely responsible for the medical advice and treatment of members. The use of this policy is not a guarantee of payment or a final prediction of how specific claim(s) will be adjudicated. Claims payment is subject to member eligibility and benefits on the date of service, coordination of benefits, referral/authorization and utilization management guidelines when applicable, and adherence to plan policies and procedures and claims editing logic.