Medimpact

STANDARD COMMERCIAL DRUG FORMULARY PRIOR AUTHORIZATION GUIDELINES

VARENICLINE

Generic	Brand	HICL	GCN	Medi-Span	Exception/Other
VARENICLINE	TYRVAYA		51384	GPI-10	
TARTRATE				(8628008020)	

GUIDELINES FOR USE

INITIAL CRITERIA (NOTE: FOR RENEWAL CRITERIA SEE BELOW)

- 1. Does the patient have a diagnosis of dry eye disease and meet ALL of the following criteria?
 - The patient is 18 years of age or older
 - Therapy is prescribed by or in consultation with an ophthalmologist or optometrist
 - The patient has at least one positive diagnostic test (e.g., tear breakup time, tear film osmolarity, ocular surfacing staining, Schirmer test, etc.)
 - The patient had a trial of or contraindication to ONE ocular lubricant (e.g., carboxymethylcellulose [Refresh, Celluvisc, Thera Tears, Genteal, etc.], polyvinyl alcohol [Liquitears, Refresh Classic, etc.], or wetting agents [Systane, Lacrilube, etc.])
 - The patient had a trial of or contraindication to BOTH of the following preferred agents: Restasis (cyclosporine) AND Xiidra (lifitegrast)

If yes, approve for 3 months by GPID or GPI-10 with a quantity limit of #8.4 mL per 30 days.

If no, do not approve.

INITIAL DENIAL TEXT: *Some terms are already pre-defined in parenthesis. Please use these definitions if the particular text you need to use does not already have definition(s) in it.

Our guideline named **VARENICLINE (Tyrvaya)** requires the following rule(s) be met for approval:

- A. You have dry eye disease
- B. You are 18 years of age or older
- C. Therapy is prescribed by or in consultation with an ophthalmologist or optometrist (types of eye doctor)
- D. You have at least one positive diagnostic test (such as tear breakup time, tear film osmolarity, ocular surfacing staining, Schirmer test)
- E. You had a trial of or contraindication to (harmful for) to ONE ocular lubricant (such as carboxymethylcellulose [Refresh, Celluvisc, Thera Tears, Genteal, etc.], polyvinyl alcohol [Liquitears, Refresh Classic], or wetting agents [Systane, Lacrilube])
- F. You had a trial of or contraindication to (harmful for) BOTH of the following preferred agents: Restasis (cyclosporine) AND Xiidra (lifitegrast)

Your doctor told us [INSERT PT SPECIFIC INFO PROVIDED]. We do not have information showing you [INSERT UNMET CRITERIA]. This is why your request is denied. Please work with your doctor to use a different medication or get us more information if it will allow us to approve this request.

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STANDARD COMMERCIAL DRUG FORMULARY PRIOR AUTHORIZATION GUIDELINES

VARENICLINE

GUIDELINES FOR USE (CONTINUED)

RENEWAL CRITERIA

- 1. Does the patient have a diagnosis of dry eye disease AND meet the following criterion?
 - The patient has demonstrated improvement of dry eye disease

If yes, approve for 12 months by GPID or GPI-10 with a quantity limit of #8.4 mL per 30 days.

If no, do not approve.

RENEWAL DENIAL TEXT: *Some terms are already pre-defined in parenthesis. Please use these definitions if the particular text you need to use does not already have definition(s) in it.

Our guideline named **VARENICLINE (Tyrvaya)** requires the following rule(s) be met for renewal:

- A. You have dry eye disease
- B. You have demonstrated improvement of dry eye disease

Your doctor told us [INSERT PT SPECIFIC INFO PROVIDED]. We do not have information showing you [INSERT UNMET CRITERIA]. This is why your request is denied. Please work with your doctor to use a different medication or get us more information if it will allow us to approve this request.

RATIONALE

For further information, please refer to the Prescribing Information and/or Drug Monograph for Tyrvaya.

REFERENCES

• Tyrvaya [Prescribing Information]. Princeton, NJ: Oyster Point Pharma, Inc., October 2021.

Library	Commercial	NSA
Yes	Yes	No

Part D Effective: N/A Commercial Effective: 07/01/23 Created: 10/21 Client Approval: 05/23

P&T Approval: 04/23

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