



STANDARD COMMERCIAL DRUG FORMULARY  
PRIOR AUTHORIZATION GUIDELINES

RANOLAZINE

Generic	Brand	HICL	GCN	Medi-Span	Exception/Other
RANOLAZINE	ASPRUZYO SPRINKLE		52005 52006	GPI-14 (32200040003020, 32200040003040)	

**GUIDELINES FOR USE**

1. Does the patient have a diagnosis of chronic angina and meet **ALL** the following criteria?
  - The patient had a trial of or contraindication to ranolazine ER tablets
  - The patient is unable to swallow ranolazine ER tablets
  - The patient had a trial of or contraindication to a nitrate (e.g., nitroglycerin, isosorbide mononitrate, isosorbide dinitrate)

If yes, **approve for 12 months by GPID or GPI-14 for the requested strength with the following quantity limits:**

- **500mg: #2 per day.**
- **1000mg: #2 per day.**

If no, do not approve.

**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 **RANOLAZINE (Aspruzyo Sprinkle)** requires the following rule(s) be met for approval:

- A. You have chronic angina (a type of heart condition)
- B. You had a trial of or contraindication (harmful for) to ranolazine ER (extended release) tablets
- C. You are unable to swallow ranolazine ER tablets
- D. You had a trial of or contraindication (harmful for) to a nitrate (such as nitroglycerin, isosorbide mononitrate, isosorbide dinitrate)

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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**RATIONALE**

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

**REFERENCES**

- Aspruzyo Sprinkle [Prescribing Information]. Cranbury, NJ: Sun Pharmaceutical Industries, Inc.; March 2022.

Library	Commercial	NSA
Yes	Yes	No

Part D Effective: N/A

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P&T Approval: 07/22