



STANDARD COMMERCIAL DRUG FORMULARY
PRIOR AUTHORIZATION GUIDELINES

ASPARAGINASE ERWINIA-RYWN

| Generic | Brand | HICL | GCN | Medi-Span | Exception/Other |
|------------------------------|--------|-------|-----|------------------------|-----------------|
| ASPARAGINASE ERWINIA-RYWN | RYLAZE | 47474 | | GPI-10 (2125001060) | |

GUIDELINES FOR USE

1. Does the patient have a diagnosis of acute lymphoblastic leukemia (ALL) or lymphoblastic lymphoma (LBL) and meet **ALL** of the following criteria?

- The patient is 1 month of age or older
- The patient has developed hypersensitivity to E. coli-derived asparaginase
- Rylaze will be used as a component of a multi-agent chemotherapeutic regimen

If yes, **approve for 12 months by HICL or GPI-10.**

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 **ASPARAGINASE ERWINIA-RYWN (Rylaze)** requires the following rule(s) be met for approval:

- A. You have acute lymphoblastic leukemia (ALL: type of blood cancer) or lymphoblastic lymphoma (LBL: type of cancer affecting the immune system)
- B. You are 1 month of age or older
- C. You have developed hypersensitivity to E.coli-derived asparaginase (you are allergic to an enzyme/protein that is from a type of bacteria)
- D. Rylaze will be used as a component of a multi-agent chemotherapeutic regimen

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 Rylaze.

REFERENCES

- Rylaze [Prescribing Information]. Palo Alto, CA: Jazz Pharmaceuticals, Inc.; June 2021.

| Library | Commercial | NSA |
|---------|------------|-----|
| Yes | Yes | No |

Part D Effective: N/A

Commercial Effective:01/01/22

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P&T Approval:10/21