



STANDARD COMMERCIAL DRUG FORMULARY
PRIOR AUTHORIZATION GUIDELINES

PACRITINIB

Generic	Brand	HICL	GCN	Medi-Span	Exception/Other
PACRITINIB CITRATE	VONJO	47850		GPI-10 (2153755010)	

GUIDELINES FOR USE

INITIAL CRITERIA (NOTE: FOR RENEWAL CRITERIA SEE BELOW)

1. Does the patient have a diagnosis of intermediate- or high-risk primary or secondary (post-polycythemia vera or post-essential thrombocytopenia) myelofibrosis and meet **ALL** of the following criteria?
 - The patient is 18 years of age or older
 - The patient has a platelet count below 50,000/uL

If yes, **approve for 6 months by HICL or GPI-10 with a quantity limit of #4 per day.**

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 **PACRITINIB (Vonjo)** requires the following rule(s) be met for approval:

- A. You have intermediate- or high-risk primary or secondary (post-polycythemia vera [type of blood cell disorder] or post-essential thrombocytopenia [type of blood cell disorder]) myelofibrosis (type of bone marrow cancer)
- B. You are 18 years of age or older
- C. You have a platelet count below 50,000/uL

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.

RENEWAL CRITERIA

1. Does the patient have a diagnosis of intermediate- or high-risk primary or secondary (post-polycythemia vera or post-essential thrombocytopenia) myelofibrosis (MF)?

If yes, continue to #2.

If no, do not approve.

DENIAL TEXT: See the renewal denial text at the end of the guideline.

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PACRITINIB

RENEWAL CRITERIA (CONTINUED)

- 2. Has the patient shown symptom improvement by meeting **ONE** of the following criteria?
 - The patient has a spleen volume reduction of 35% or greater from baseline
 - The patient has a 50% or greater reduction in total symptom score (e.g., Myeloproliferative Neoplasm Symptom Assessment Form [MPN-SAF TSS], modified Myelofibrosis Symptom Assessment Form [MFSAF] v2.0)
 - The patient has a 50% or greater reduction in palpable spleen length

If yes, **approve for 12 months by HICL or GPI-10 with a quantity limit of #4 per day.**
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 **PACRITINIB (Vonjo)** requires the following rule(s) be met for renewal:

- A. You have intermediate- or high-risk primary or secondary (post-polycythemia vera [type of blood cell disorder] or post-essential thrombocythemia [type of blood cell disorder]) myelofibrosis (type of bone marrow cancer)
- B. You have shown symptom improvement by meeting ONE of the following:
 - 1. You have a spleen volume reduction of 35% or greater from baseline
 - 2. You have a 50% or greater reduction in total symptom score (such as Myeloproliferative Neoplasm Symptom Assessment Form [MPN-SAF TSS], modified Myelofibrosis Symptom Assessment Form [MFSAF] v2.0)
 - 3. You have a 50% or greater reduction in palpable (can be felt by external examination) spleen length

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 Vonjo.

REFERENCES

- Vonjo [Prescribing Information]. Seattle, WA: CTI BioPharma Corp.; February 2022.

Library	Commercial	NSA
Yes	Yes	No

Part D Effective: N/A

Commercial Effective: 04/01/22

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