



STANDARD COMMERCIAL DRUG FORMULARY
PRIOR AUTHORIZATION GUIDELINES

MOBOCERTINIB

Generic	Brand	HICL	GCN	Medi-Span	Exception/Other
MOBOCERTINIB SUCCINATE	EXKIVITY	47578		GPI-10 (2136005060)	

GUIDELINES FOR USE

1. Does the patient have a diagnosis of locally advanced or metastatic non-small cell lung cancer (NSCLC) and meet **ALL** of the following criteria?
 - The patient is 18 years of age or older
 - The patient has epidermal growth factor receptor (EGFR) exon 20 insertion mutations, as detected by an FDA-approved test
 - The patient's disease progressed on or after platinum-based chemotherapy (e.g., cisplatin, carboplatin, oxaliplatin)

If yes, **approve for 12 months by HICL or GPI-10 with a quantity limit of #4 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 **MOBOCERTINIB (Exkivity)** requires the following rule(s) be met for approval:

- A. You have locally advanced or metastatic (cancer that has spread from where it started to nearby tissue or has spread to other parts of the body) non-small cell lung cancer (NSCLC: type of lung cancer)
- B. You are 18 years of age or older
- C. You have epidermal growth factor receptor (EGFR) exon 20 insertion mutations (type of gene mutation), as detected by a Food and Drug Administration (FDA)-approved test
- D. Your disease progressed (disease has gotten worse) on or after platinum-based chemotherapy such as cisplatin, carboplatin, oxaliplatin

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 Exkivity.

REFERENCES

- Exkivity [Prescribing Information]. Lexington, MA: Takeda Pharmaceuticals America, Inc., September 2021.

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Library	Commercial	NSA
Yes	Yes	No

Part D Effective: N/A

Commercial Effective:01/01/22

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Client Approval: 11/21

P&T Approval:10/21