



STANDARD COMMERCIAL DRUG FORMULARY
PRIOR AUTHORIZATION GUIDELINES

BELZUTIFAN

| Generic | Brand | HICL | GCN | Medi-Span | Exception/Other |
|------------|---------|-------|-----|------------------------|-----------------|
| BELZUTIFAN | WELIREG | 47546 | | GPI-10 (2142102000) | |

GUIDELINES FOR USE

1. Does the patient have a diagnosis of von Hippel-Lindau (VHL) disease and meet **ALL** of the following criteria?
 - The patient is 18 years of age or older
 - The patient requires therapy for associated renal cell carcinoma (RCC), central nervous system (CNS) hemangioblastomas, or pancreatic neuroendocrine tumors (pNET)
 - The patient does NOT require immediate surgery

If yes, **approve for 12 months by HICL or GPI-10 with a quantity limit of #3 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 **BELZUTIFAN (Welireg)** requires the following rule(s) be met for approval:

- A. You have von Hippel-Lindau (VHL) disease (genetic disorder that causes tumors to grow in the body)
- B. You are 18 years of age or older
- C. You require therapy for associated renal cell carcinoma (RCC: a type of kidney cancer), central nervous system (CNS) hemangioblastomas (tumor in the brain or spinal cord), or pancreatic neuroendocrine tumors (pNET: tumor in the pancreas)
- D. You do NOT require immediate surgery

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 Welireg.

REFERENCES

- Welireg [Prescribing Information]. Whitehouse Station, NJ: Merck & Co, Inc.; August 2021.

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

Part D Effective: N/A

Commercial Effective:01/01/22

Created: 10/21

Client Approval: 11/21

P&T Approval: 10/21