

This document applies to the following:

Formulary	Applies
Medicare Part B	<input checked="" type="checkbox"/>
Medicare Part B: Advanced Biosimilars First	<input checked="" type="checkbox"/>

Medicare Part B Step Therapy

Rituximab Products

This document informs prescribers of preferred products and provides an exception process for non-preferred products through prior authorization.

These criteria were developed to align with the following: Medicare Part B and Medicare Part B: Advanced Biosimilars First.

Plan Design Summary

This program applies to the rituximab products specified in this document. Coverage for non-preferred products is provided based on clinical circumstances that would exclude the use of the preferred product and may be based on previous use of a product. The coverage review process will ascertain situations where a clinical exception can be made. This program applies to all members requesting treatment with the non-preferred product.

Step therapy is applied in addition to any applicable National Coverage Determination (NCD), Local Coverage Determination (LCD), and Medicare Part B utilization management (UM) programs implemented for the client.

Table. Rituximab Products

Medications considered preferred on your plan may still require a clinical prior authorization review.

	Products
Preferred	<ul style="list-style-type: none">Ruxience (rituximab-pvvr)Truxima (rituximab-abbs)
Non-preferred	<ul style="list-style-type: none">Riabni (rituximab-arrx)Rituxan (rituximab)Rituxan Hycela (rituximab and hyaluronidase human)

Reference number(s)
5328-D

Step Therapy Criteria

Coverage for a non-preferred product is provided when either of the following criteria is met:

- Member has received treatment with the non-preferred product in the past 365 days.
- Member has had a documented intolerable adverse event to both of the preferred products, and the adverse event was not an expected adverse event attributed to the active ingredient as described in the prescribing information (i.e., known adverse reaction for both the reference product and biosimilar products).

References

1. Riabni [package insert]. Thousand Oaks, CA: Amgen, Inc.; February 2023.
2. Rituxan [package insert]. South San Francisco, CA: Genentech, Inc.; December 2021.
3. Rituxan Hycela [package insert]. South San Francisco, CA: Genentech, Inc.; June 2021.
4. Ruxience [package insert]. New York, NY: Pfizer; October 2023.
5. Truxima [package insert]. North Wales, PA: Teva Pharmaceuticals USA, Inc; December 2024.