

PATIENT INFORMATION

Name:

Address:

PHN:

Application for Coverage for RITUXIMAB for GN Patients

Rev: Sep/17

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INSTRUCTIONS

1. Ensure the patient is registered in PROMIS under the Provincial Renal Agency program.
 - a. Choose the most appropriate GN diagnosis under the available list of primary renal diagnoses.
 - b. Ensure the patient address and contact information are accurate as these are needed for medication distribution.
2. Complete the information below, **fax this form to the BCPRA at (604) 875-7366.**
3. This application will be reviewed by the BCPRA Pharmacy and Formulary Committee; you will be contacted once approval is decided.
4. If approved, fax approval letter to the hospital pharmacy where RITUXIMAB will be infused, to inform them of BCPRA coverage.
5. Please note that coverage for RITUXIMAB is limited to one course; for subsequent treatment courses this form must be completed again.
6. Coverage is contingent upon the approved medication being entered into PROMIS, with accurate doses (including changes), frequency and start/stop dates.

THE FOLLOWING ARE REQUIRED FOR MEDICATION APPROVAL:

GN Diagnosis with PROMIS codes (pick one):

- ANCA vasculitis / pauci-immune glomerulonephritis (69, 74 or 98)
- Anti-GBM antibody disease / Goodpasture's disease (86)
- FSGS (09 or 11)
- IgA nephropathy (12)
- Minimal change disease (06)
- Membranous nephropathy (14)
- Lupus nephritis (84), provide class _____
- Other: _____
- Additional details about diagnosis, if needed: _____

Weight: _____

Height: _____

Provide body surface area if used for dosing: _____ m²

- 1000 mg
- 375 mg/m² = _____ mg (round to nearest 50 mg)
- Other: _____ mg (round to nearest 50 mg)

Frequency of infusions (pick one):

- Weekly
- Every 2 weeks
- Other: _____

Total number of infusions: _____

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If you are applying for RITUXIMAB for ANCA vasculitis or pauci-immune glomerulonephritis:

Provide details that confirm the patient has renal involvement.

* Note that patients without renal involvement are not eligible for BCPRA coverage and instead you should apply to [BC PharmaCare](#) through the Special Authority Process.

For **induction therapy**: indicate why cyclophosphamide can't be used AND provide details below.

- Disease is resistant to cyclophosphamide
- Cumulative previous exposure to cyclophosphamide exceeds the tolerable limit (exceeding 10 g increases the risk of infertility and exceeding 20 g increases the risk of malignancy)
- Patient has a contraindication to cyclophosphamide

Provide details that justify the above choice:

For **maintenance therapy**: indicate which of the following apply AND provide details below.

- Patient has a contraindication to the use of azathioprine maintenance therapy after a disease flare that included renal involvement.
- Patient is intolerant to at least 1.5 to 2 mg/kg/day of azathioprine maintenance therapy after induction treatment for a disease flare that included renal involvement.
- Disease relapse with renal involvement has occurred while on at least 1.5 to 2 mg/kg/day of azathioprine maintenance therapy.

Provide details that justify the above choice:

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If you are applying for RITUXIMAB for another GN indication (not ANCA vasculitis or pauci-immune glomerulonephritis):

To justify the use of RITUXIMAB, please indicate why the following medications can't be used by checking the appropriate boxes AND briefly providing details:

	Disease Relapsed	Disease Resistant	Patient Intolerant	Contra-indicated	Not Indicated	Provide Details
Prednisone						
Azathioprine						
Mycophenolate/ Myfortic®						
Cyclosporine/ Tacrolimus						
Cyclophosphamide						

Provide any addition details that may support this application:

Name: _____

Signature: _____

Phone: _____

Date: _____

Fax number for approval notification (mandatory): _____

If you do not receive a response to your application within 72 hours, or if you'd like to check on the status of your application, please call (604) 875-7367.