

**AETNA BETTER HEALTH® OF VIRGINIA REQUEST FORM**

**DUR MEDICATION WAYRILZ™ (rilzabrutinib)**

**Fax back to: 1-855-799-2553**

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If the following information is not complete, correct, or legible, the PA process can be delayed. Please use one form per member.

**MEMBER INFORMATION**

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Last Name:

First Name:

Medicaid ID Number:

Date of Birth:

Weight in Kilograms: \_\_\_\_\_

**PRESCRIBER INFORMATION**

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Last Name:

First Name:

NPI Number:

Phone Number:

Fax Number:

**DRUG INFORMATION**

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Drug Name/Form: \_\_\_\_\_

Strength: \_\_\_\_\_

Dosing Frequency: \_\_\_\_\_

Length of Therapy: \_\_\_\_\_

Quantity per Day: \_\_\_\_\_

*(Form continued on next page.)*

Member's Last Name:

Member's First Name:

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**DIAGNOSIS AND MEDICAL INFORMATION**

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**WAYRILZ™ – To obtain a six (6)-months approval for this medication, please complete the following questions.**

1. Is the member  $\geq 18$  years of age?

Yes     No

2. Does the member have a diagnosis of persistent (defined as 3 to 12 months) or chronic (defined as lasting for  $> 12$  months) immune (idiopathic) thrombocytopenia (ITP) and ONE of the following:

- The member has a platelet count  $\leq 30 \times 10^9/L$ ; **OR**
- The member has a platelet count  $> 30 \times 10^9/L$  but  $< 50 \times 10^9/L$  AND has symptomatic bleeding and/or an increased risk of bleeding?

Yes     No

3. Does the member have ONE of the following:

- The member has tried and had an inadequate response to ONE corticosteroid used for the treatment of ITP; **OR**
- The member has an intolerance or hypersensitivity to ONE corticosteroid used for the treatment of ITP; **OR**
- The member has an FDA labeled contraindication to ALL corticosteroids used for the treatment of ITP; **OR**
- The member has tried and had an inadequate response to a thrombopoietin receptor agonist (e.g., avatrombopag [Doptelet], romiplostim [Nplate], eltrombopag [Promacta]); **OR**
- The member has tried and had an inadequate response to immunoglobulins (intravenous immunoglobulin [IVIg] or Anti-D); **OR**
- The member has had an inadequate response to a splenectomy; **OR**
- The member has tried and had an inadequate response to rituximab?

Yes     No

*(Form continued on next page.)*

**Member's Last Name:**

**Member's First Name:**

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4. Does the prescriber attest that the member will NOT use Wayrilz in combination with other kinase inhibitors with similar indications (e.g., fostamatinib [Tavalisse])?

Yes     No

**For renewal, complete the following questions to receive a 1-year approval:**

5. Does the member continue to meet the above criteria (questions 1 through 4)?

Yes     No

6. Does the member continue to experience clinical benefit from the requested treatment?

Yes     No

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**Prescriber Signature (Required)**

**Date**

By signature, the Physician confirms the above information is accurate and verifiable by member records.

**Please include ALL requested information; incomplete forms will delay the PA process.**

Submission of documentation does NOT guarantee coverage.