

**Revcovi™ Prior Authorization Request Form (Page 1 of 2)**  
 DO NOT COPY FOR FUTURE USE. FORMS ARE UPDATED FREQUENTLY AND MAY BE BARCODED

<b>Member Information</b> (required)			<b>Provider Information</b> (required)		
Member Name:			Provider Name:		
Insurance ID#:			NPI#:	Specialty:	
Date of Birth:			Office Phone:		
Street Address:			Office Fax:		
City:	State:	Zip:	Office Street Address:		
Phone:			City:	State:	Zip:

<b>Medication Information</b> (required)		
Medication Name:	Strength:	Dosage Form:
<input type="checkbox"/> Check if requesting <b>brand</b>	Directions for Use:	
<input type="checkbox"/> Check if request is for <b>continuation of therapy</b>		

<b>Clinical Information</b> (required)
<b>Select the diagnosis below:</b>
<input type="checkbox"/> Adenosine deaminase severe combined immune deficiency (ADA-SCID)
<input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____

<b>Clinical Information:</b>
Will Renvovi be used in combination with Adagen (pegademase-bovine)? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b>
Select if the patient has severe combined immunodeficiency disease (SCID) with a definitive diagnosis of adenosine deaminase deficiency (ADA) as determined by the following:
<input type="checkbox"/> Deficient ADA catalytic activity (< 1% of normal) in hemolysates (in untransfused individuals) or in extracts of other cells (e.g., blood mononuclear cells, fibroblasts)
<input type="checkbox"/> Detection of pathogenic mutations in the ADA gene by molecular genetic testing
Does the patient have a marked elevation of the metabolite deoxyadenosine triphosphate (dATP) or total dAdo nucleotides (the sum of deoxyadenosine monophosphate [dAMP], deoxyadenosine diphosphate [dADP] and dATP) in erythrocytes? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b>
Is the patient <b>NOT</b> a candidate for or has failed hematopoietic cell transplantation (HCT)? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b>
Does the patient have severe thrombocytopenia (< 50,000/mcL)? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b>
Select if baseline laboratory values documenting the following will be submitted:
<input type="checkbox"/> Plasma ADA activity <input type="checkbox"/> Red blood cell dATP <input type="checkbox"/> Trough dAXP levels <input type="checkbox"/> Total lymphocyte counts
<i>Baseline laboratory values must be submitted</i>
Will medical records (e.g., chart notes) documenting patient's current height and weight be submitted? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b>
<i>Medical records (e.g., chart notes) must be submitted</i>
Is the prescribed dose based on patient's ideal body weight? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b>

<b>Reauthorization:</b>
<b>If this is a reauthorization, please answer the following:</b>
Will Renvovi be used in combination with Adagen (pegademase-bovine)? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b>
Does the patient have a marked elevation of the metabolite dATP or total dAdo nucleotides (the sum of dAMP, dADP and dATP) in erythrocytes? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b>
Is the patient <b>NOT</b> a candidate for or has failed hematopoietic cell transplantation (HCT)? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b>
Does the patient have severe thrombocytopenia (< 50,000/mcL)? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b>
<b>&lt; continued on the next page &gt;</b>

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Will medical records (e.g., chart notes) documenting patient's current height and weight be submitted?  Yes  No

*Medical records (e.g., chart notes) must be submitted*

Is the prescribed dose based on patient's ideal body weight?  Yes  No

Has the patient experienced unacceptable toxicity from the drug (e.g., severe injection site reactions, bleeding, severe thrombocytopenia, etc.)?  Yes  No

Select if medical records (e.g., chart notes, laboratory values) documenting patient's disease stability and/or improvement as indicated by the following will be submitted:

- Increase in plasma ADA activity (target trough level greater than or equal to 15 mmol/hr/L)
- Red blood cell dATP level decreased (target less than or equal to 0.005 to 0.015 mmol/L)
- Improvement in immune function with diminished frequency/complications of infection as evidenced in improvement in the ability to produce antibodies
- Improvement in red blood cell dAXP levels (target trough level less than or equal to 0.02 mmol/L)

*Medical records (e.g., chart notes, laboratory values) must be submitted*

**\*\*Use of samples to initiate therapy does not meet step edit/ preauthorization criteria.\*\***

**\*Previous therapies will be verified through pharmacy paid claims or submitted chart notes.\***

**Prescriber Signature:** \_\_\_\_\_ **Date:** \_\_\_\_\_

Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?

Please note:

This request may be denied unless all required information is received.  
For urgent or expedited requests please call 1-800-711-4555.  
This form may be used for non-urgent requests and faxed to 1-800-527-0531.