



NovoLog® Products Prior Authorization Request Form (Page 1 of 2)

DO NOT COPY FOR FUTURE USE. FORMS ARE UPDATED FREQUENTLY AND MAY BE BARCODED

Member Information <small>(required)</small>			Provider Information <small>(required)</small>		
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:

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

Clinical Information <small>(required)</small>	
Please note: A review of the vial will be conducted unless otherwise indicated.	
Select the diagnosis below:	
<input type="checkbox"/> Type 1 diabetes mellitus	
<input type="checkbox"/> Type 2 diabetes mellitus	
<input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____	
The following are the formulary alternatives:	
For Novolog FlexPen and Novolog PenFill- Admelog SoloStar, Fiasp FlexTouch, Humalog cartridge, Humalog Junior KwikPen, Humalog KwikPen	
For Novolog Mix 70/30 FlexPen- Humalog mix 50/50 KwikPen, Humalog mix 75/25 KwikPen	
For Novolog Mix 70/30 vial- Humalog mix 75/25 vial	
For Novolog vial- Humalog vial	
Will the patient be switched to one of the formulary alternatives? <input type="checkbox"/> Yes <input type="checkbox"/> No	
If yes , please specify which alternative the patient will be switched to and notify the pharmacy of the change: _____	
If the patient CANNOT be switched to a formulary alternative, please answer ALL of the following questions:	
1) What previous medication(s) has the patient tried or failed for the diagnosis provided: _____	
2) If the patient has failed, had adverse reactions, or contraindications to the above formulary alternative(s), provide clinical details as to what occurred: _____	

**Please note: Submit chart documentation/medical records to support the information you have provided above	
For NovoLog vials, also answer the following (does NOT apply to NovoLog FlexPen, NovoLog 70/30 Mix, NovoLog 70/30 Mix FlexPen, or NovoLog Penfill):	
Is NovoLog administered using an infusion pump? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Was the infusion pump paid for by Medicare? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Is the patient using a subcutaneous insulin pump [excluding disposable drug delivery systems (e.g., OmniPod, V-Go)]? <input type="checkbox"/> Yes <input type="checkbox"/> No	
Is the patient enrolled in a comprehensive diabetes program with one of the following symptoms? <input type="checkbox"/> Yes <input type="checkbox"/> No	
<ul style="list-style-type: none"> • Dawn phenomenon • Fluctuations in blood glucose • Hemoglobin level (HbA1C) greater than 7 percent • History of recurring hypoglycemia • History of severe glycemic excursions 	
< Continued on next page >	



NovoLog[®] Products Prior Authorization Request Form (Page 2 of 2)

DO NOT COPY FOR FUTURE USE. FORMS ARE UPDATED FREQUENTLY AND MAY BE BARCODED

< Continued from previous page >

Has the patient been on an external insulin infusion pump prior to enrollment in Medicare and has documented frequency of glucose self-testing an average of at least 4 times per day? Yes No

Does the patient have a fasting blood sugar less than or equal to 225mg/dL? Yes No

Does the patient have a Beta cell autoantibody test that is positive? Yes No

Select ONE of the following:

NovoLog is administered at home (not including facility providing skilled nursing care)

The patient is in a long-term care (LTC) facility (e.g., hospital or skilled nursing facility where patient is receiving skilled care)

Prescriber attestation:

Does the prescriber attest that the information provided on this form is true and accurate? Yes No

Prescriber signature: _____ **Date:** _____

(Please note: if a non-formulary exception is approved, the requested drug will process at the **highest** brand tier copay for the plan year)

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.
If the patient is not able to meet the above standard prior authorization requirements, please call 1-800-711-4555.
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.

This document and others if attached contain information that is privileged, confidential and/or may contain protected health information (PHI). The Provider named above is required to safeguard PHI by applicable law. The information in this document is for the sole use of OptumRx. Proper consent to disclose PHI between these parties has been obtained. If you received this document by mistake, please know that sharing, copying, distributing or using information in this document is against the law. **If you are not the intended recipient, please notify the sender immediately.**

Office use only: NovoLogProducts_CMS_2019Jan-W