

PRIOR AUTHORIZATION REQUEST FORM

BASAL INSULIN

Insulin Glargine, Toujeo[®], Insulin Degludec

For authorization, please answer each question and fax this form PLUS chart notes back to Real Rx at 385-425-4052.

Failure to submit clinical documentation to support this request will result in a dismissal of the request.

If you have prior authorization questions, please call for assistance 385-425-5094.

Disclaimer: Prior authorization request forms are subject to change in accordance with Federal and State notice requirements.

Date:	Member Name:	ID#:
Bate.		
DOB:	Gender:	Physician:
DOD.	Gender.	
Office Phone:	Office Fax:	Office Contact:
Office Filone.	Office Fax.	Office contact.

Height/Weight:

Member must try formulary preferred drugs before a request for a non-preferred drug may be considered. If treatment with preferred products has not been successful, you must submit which preferred products have been tried, dates of treatment, and reason for failure. Reasons for failure must meet the Health Plan medical necessity criteria.

Preferred/Non-Preferred

- 1. Preferred
 - A. Basaglar[®] (Insulin glargine 100 Units/mL), Rezvoglar[™] (insulin glargine-aglr); no prior authorization required
- 2. Non-Preferred Brands with a single step; after trial and failure of Basaglar[®] and in accordance with Prior Authorization Criteria below
 - A. Insulin Degludec (100 Units/mL and 200 Units/mL)
- 3. Non-preferred Brands with a double step; after trial and failure of Basaglar[®] AND Insulin Degludec and in accordance with Prior Authorization Criteria below
 - A. Toujeo® (Insulin glargine 300 Units/mL), Insulin glargine 100 Units/ml
 - B. Levemir® (Insulin detemir): Individual Plans ONLY
- 4. Non-formulary (Commercial Plans Only)
 - A. Levemir[®] (Insulin detemir)

Product being requested: _____

Dosing/Frequency:____

If the request is for reauthorization, proceed to reauthorization section					
Questions		Yes	No	Comments/Notes	
1.	Is this request for an expedited review? By checking the "Yes" box to request an expedited review (24				
hours), you are certifying that applying the standard review time frame (72 hours) may place the member's life, health, or					
	ability to regain maximum function in serious jeopardy.				
Insulin Degludec					
1.	Does the member have a diagnosis of Type 1 or Type 2 diabetes mellitus or gestational diabetes?			Please provide documentation	

2.	Has the member tried Basaglar [®] or Rezvoglar [®] for at least 3 months?			Please provide documentation			
Toujeo and Insulin Glargine							
1.	Does the member have a diagnosis of Type 1 or Type 2 diabetes mellitus or gestational diabetes?			Please provide documentation			
2.	Has the member tried Basaglar® or Rezvoglar® and Insulin Degludec for at least 3 months?			Please provide documentation			
	REAUTHORIZATIO	N					
1.	Is the request for reauthorization of therapy?						
2.	Has the member's therapy been re-evaluated within the past 12 months?						
3.	Does the member show a continued medical need for the therapy?			Please provide documentation			
4.	Has the therapy been tolerable and effective?			Please provide documentation			
	ne of treatment, reason for failure, treatment dates, etc.						
	litional information:						
Physician Signature:							

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Policy: PHARM-011 Origination Date: 03/20/2018 Reviewed/Revised Date: 05/19/2023 Next Review Date: 05/19/2024 Current Effective Date: 06/01/2023

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