

SENTARA HEALTH PLANS

PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

Directions: The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request. All other information may be filled in by office staff; fax to 1-800-750-9692. No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. If the information provided is not complete, correct, or legible, the authorization process can be delayed.

Non-Preferred Insulins

Drug Requested: Select one from below

Rapid-acting Insulin Products		
<input type="checkbox"/> Admelog® (insulin lispro) vial/SoloStar®	<input type="checkbox"/> Apidra® (insulin glulisine) vial/SoloStar®	<input type="checkbox"/> Fiasp® (insulin aspart) vial/FlexTouch®/PenFill®/PumpCart®
<input type="checkbox"/> insulin aspart vial/cartridge/pen (Novolog ABA)	<input type="checkbox"/> Kirsty™ (insulin aspart-xjhz) vial/pen	<input type="checkbox"/> Lyumjev® (insulin lispro-aabc) vial/KwikPen®
<input type="checkbox"/> Merilog™ (insulin aspart-szjj) vial/SoloStar®	<input type="checkbox"/> Novolog® (insulin aspart) vial/FlexPen®/PenFill®	
Regular or short-acting Insulin Products		
<input type="checkbox"/> Novolin® R (Regular, Human Insulin) vial/FlexPen®		
Intermediate-acting Insulin Products		
<input type="checkbox"/> Novolin® N (NPH, Human Insulin) vial/FlexPen®		
Long-acting Insulin Products		
<input type="checkbox"/> Basaglar® (insulin glargine) KwikPen®	<input type="checkbox"/> insulin degludec vial/pen (Tresiba ABA)	<input type="checkbox"/> insulin glargine vial/SoloStar® (Lantus ABA)
<input type="checkbox"/> insulin glargine SoloStar®/Max SoloStar® (Toujeo SoloStar® ABA)	<input type="checkbox"/> Levemir® (insulin detemir) vial/FlexTouch®	<input type="checkbox"/> Semglee® (insulin glargine-yfgn) vial/pen
<input type="checkbox"/> Tresiba® (insulin degludec) vial/FlexTouch®		
Combination of Insulin Products:		
<input type="checkbox"/> insulin aspart protamine suspension/insulin aspart mix 70/30 vial/FlexPen® (Novolog Mix ABA)	<input type="checkbox"/> Novolin® 70/30 (70% NPH, Human Insulin Isophane Suspension & 30% Regular, Human Insulin) vial/FlexPen®	<input type="checkbox"/> Novolog® Mix 70/30 (70% insulin aspart protamine suspension & 30% insulin aspart) vial/FlexPen®

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MEMBER & PRESCRIBER INFORMATION: Authorization may be delayed if incomplete.

Member Name: _____

Member Sentara #: _____ Date of Birth: _____

Prescriber Name: _____

Prescriber Signature: _____ Date: _____

Office Contact Name: _____

Phone Number: _____ Fax Number: _____

NPI #: _____

DRUG INFORMATION: Authorization may be delayed if incomplete.

Drug Form/Strength: _____

Dosing Schedule: _____ Length of Therapy: _____

Diagnosis: _____ ICD Code, if applicable: _____

Weight (if applicable): _____ Date weight obtained: _____

CLINICAL CRITERIA: Check below all that apply. All criteria must be met for approval. To support each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be provided or request may be denied.

For Novolin® Brand, Mix or ABA products, the following criteria must be met:

Member has tried and failed at least **30 days** of therapy with a Humulin® product

For Admelog®, Apidra®, Fiasp®, insulin aspart, Kirsty™, Lyumjev®, Merilog™ and Novolog® Brand, Mix or ABA products, all the following criteria must be met:

Member must have a **30-day trial** and failure or intolerance to brand Humalog®

Provider must submit clinical chart notes or a completed MedWatch form documenting the experienced treatment failure or intolerance to brand Humalog®

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For Basaglar®, insulin degludec, insulin glargine, Levemir®, Semglee® Brand or ABA products and Tresiba®, the following criteria must be met:

- Member has tried and failed at least **30 days** of therapy with **ONE** of the following
 - Lantus®
 - Toujeo®

Not all drugs may be covered under every Plan.

If a drug is non-formulary on a Plan, documentation of medical necessity will be required.

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.