



UnitedHealthcare Pharmacy
Clinical Pharmacy Programs

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| Program Number | 2022 P 3050-12 |
| Program | Step Therapy – Insulin |
| Medication | Apidra (insulin glulisine)*, Apidra SoloStar (insulin glulisine)*, Fiasp (insulin aspart)*, Novolin N (NPH, human insulin isophane)*, Novolin R (regular, human insulin)*, Novolin 70/30 (70% NPH, human insulin isophane and 30% regular, human insulin)*, Novolog (insulin aspart)*, Novolog Mix 70/30 (70% insulin aspart protamine and 30% insulin aspart)* |
| P&T Approval Date | 12/2014, 10/2015, 10/2016, 10/2017, 5/2018, 10/2018, 6/2019, 6/2020, 2/2021, 2/2022 |
| Effective Date | 5/1/2022; Oxford: 5/1/2022 |

1. Background:

The American Diabetes Association recommends insulin therapy for Type II diabetes when the appropriate step wise non-insulin approach has failed to lower HbA1c. In Type I diabetes insulin monotherapy is the appropriate treatment. The ADA does not differentiate between brands of insulin but does make recommendations for the initiation of basal insulins or intermediate to short acting insulins.

2. Coverage Criteria^{a,b}:

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| <p>A. Novolin 70/30* will be approved based on the following criteria:</p> <ol style="list-style-type: none">1. History of failure after at least a three month trial^c, contraindication, or intolerance (list reason for therapeutic failure, contraindication, or intolerance) to Humulin 70/30 <p>B. Apidra*, Apidra Solostar*, or Novolog* pens and vials will be approved based on the following criteria:</p> <ol style="list-style-type: none">1. History of failure after at least a three month trial^c, contraindication, or intolerance (list reason for therapeutic failure, contraindication, or intolerance) to Humalog <p>C. Fiasp* pens and vials will be approved based on <u>ONE</u> of the following criteria:</p> <ol style="list-style-type: none">1. For patients 18 years of age and older: |
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a. History of failure after at least a three month trial^c, contraindication, or intolerance (list reason for therapeutic failure, contraindication, or intolerance) to **BOTH** of the following:

(1) Humalog

(2) Lyumjev

2. For patients less than 18 years of age:

a. History of failure after at least a three month trial^c, contraindication, or intolerance (list reason for therapeutic failure, contraindication, or intolerance) to Humalog

D. Novolin N* will be approved based on the following criteria:

1. History of failure after at least a three month trial^c, contraindication, or intolerance (list reason for therapeutic failure, contraindication, or intolerance) to Humulin N

E. Novolin R* will be approved based on the following criteria:

1. History of failure after at least a three month trial^c, contraindication, or intolerance (list reason for therapeutic failure, contraindication, or intolerance) to Humulin R

F. Novolog Mix 70/30* pens and vials will be approved based on the following criteria:

1. History of failure after at least a three month trial^c, contraindication, or intolerance (list reason for therapeutic failure, contraindication, or intolerance) to Humalog 75/25

Authorization will be issued for 12 months

^a State mandates may apply. Any federal regulatory requirements and the member specific benefit plan coverage may also impact coverage criteria. Other policies and utilization management programs may apply.

^b In Florida, Maine, and Tennessee only, diabetes medications may be approved based on both of the following: 1) Provider attests use of this product is medically necessary; and- 2) If applicable, clinical characteristics exist that preclude the use of the covered preferred alternative(s) and use of the covered preferred alternative(s) could result in worsening of patient's condition or inadequate treatment (document alternatives and clinical information related to worsening/inadequate treatment).

^c For Connecticut business, only a 60-day trial will be required. For Kentucky business, only a 30-day trial will be required.

3. Additional Clinical Rules:

- Notwithstanding Coverage Criteria, UnitedHealthcare may approve initial and re-authorization based solely on previous claim/medication history, diagnosis codes (ICD-10) and/or claim logic. Use of automated approval and re-approval processes varies by program and/or therapeutic class
- Supply limits may be in place.

*Typically excluded from coverage.

4. References:

1. Novolog [package insert]. Plainsboro, NJ: Novo Nordisk Inc. October 2021.
2. Novolin 70/30 [package insert]. Plainsboro, NJ: Novo Nordisk Inc. April 2021.
3. Humulin 70/30 [package insert]. Indianapolis, IN: Eli Lilly. November 2019.
4. Humalog [package insert]. Indianapolis, IN: Eli Lilly. November 2019.
5. Apidra [package insert]. Bridgewater, NJ: Sanofi Aventis. December 2020.
6. American Diabetes Association Pharmacologic Approaches to Glycemic Treatment: Standards of Medical Care in Diabetes. Clinical Diabetes. 2021 Jan 44(1): S111-S124.
7. Novolin N [package insert]. Plainsboro, NJ: Novo Nordisk Inc. November 2019.
8. Novolin R [package insert]. Plainsboro, NJ: Novo Nordisk Inc. November 2019.
9. Humulin N [package insert]. Indianapolis, IN: Eli Lilly. November 2019.
10. Humulin R [package insert]. Indianapolis, IN: Eli Lilly. November 2019.
11. Fiasp [package insert]. Plainsboro, NJ: Novo Nordisk Inc. December 2019.

| Program | Step Therapy- Insulin |
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| Change Control | |
| 12/2014 | New program |
| 10/2015 | Added authorization period. Separated out Novolog Mix 70/30 criteria. Added Maryland Continuation of Care |
| 7/2016 | Added Indiana and West Virginia coverage information. |
| 10/2016 | Removed Humulin from step one agents for Novolin 70/30. Administrative changes. |
| 2/2017 | Administrative change. Oxford effective date updated. |
| 10/2017 | Added Fiasp to criteria. State mandate reference language updated. References updated. |
| 5/2018 | Added statement that Fiasp is typically excluded from coverage. |
| 10/2018 | Retire program for 1/1/2019. |
| 6/2019 | Program re-implemented. Updated to note all targeted products are typically excluded from coverage. Updated references. |
| 6/2020 | Annual Review. Updated references. |
| 2/2021 | Updated criteria for Fiasp to require a trial of both Humalog and Lyumjev for patients 18 years old and older. |
| 8/2021 | Administrative change to add Oxford effective date. |



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| 2/2022 | Annual review. Updated references. Added Florida, Maine, and Tennessee mandate language. Updated state mandate language for CT and KY. |
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