

PREFERRED SPECIALTY MANAGEMENT POLICY

POLICY: Growth Disorders – Growth Hormone Long-Acting Products Preferred Specialty Management Policy

• Ngenla® (somatrogon-ghla subcutaneous injection – Pfizer)

Skytrofa[®] (Ionapegsomatropin subcutaneous injection – Ascendis)

 Sogroya® (somapacitan-beco subcutaneous injection – Novo Nordisk)

REVIEW DATE: 11/20/2024; selected revision 08/13/2025

INSTRUCTIONS FOR USE

The following Coverage Policy applies to health benefit plans administered by Cigna Companies. Certain Cigna COMPANIES AND/OR LINES OF BUSINESS ONLY PROVIDE UTILIZATION REVIEW SERVICES TO CLIENTS AND DO NOT MAKE COVERAGE DETERMINATIONS. REFERENCES TO STANDARD BENEFIT PLAN LANGUAGE AND COVERAGE DETERMINATIONS DO NOT APPLY TO THOSE CLIENTS. COVERAGE POLICIES ARE INTENDED TO PROVIDE GUIDANCE IN INTERPRETING CERTAIN STANDARD BENEFIT PLANS ADMINISTERED BY CIGNA COMPANIES. PLEASE NOTE, THE TERMS OF A CUSTOMER'S PARTICULAR BENEFIT PLAN DOCUMENT [GROUP SERVICE AGREEMENT, EVIDENCE OF COVERAGE, CERTIFICATE OF COVERAGE, SUMMARY PLAN DESCRIPTION (SPD) OR SIMILAR PLAN DOCUMENT] MAY DIFFER SIGNIFICANTLY FROM THE STANDARD BENEFIT PLANS UPON WHICH THESE COVERAGE POLICIES ARE BASED. FOR EXAMPLE, A CUSTOMER'S BENEFIT PLAN DOCUMENT MAY CONTAIN A SPECIFIC EXCLUSION RELATED TO A TOPIC ADDRESSED IN A COVERAGE POLICY. IN THE EVENT OF A CONFLICT, A CUSTOMER'S BENEFIT PLAN DOCUMENT ALWAYS SUPERSEDES THE INFORMATION IN THE COVERAGE POLICIES. IN THE ABSENCE OF A CONTROLLING FEDERAL OR STATE COVERAGE MANDATE, BENEFITS ARE ULTIMATELY DETERMINED BY THE TERMS OF THE APPLICABLE BENEFIT PLAN DOCUMENT. COVERAGE DETERMINATIONS IN EACH SPECIFIC INSTANCE REQUIRE CONSIDERATION OF 1) THE TERMS OF THE APPLICABLE BENEFIT PLAN DOCUMENT IN EFFECT ON THE DATE OF SERVICE; 2) ANY APPLICABLE LAWS/REGULATIONS; 3) ANY RELEVANT COLLATERAL SOURCE MATERIALS INCLUDING COVERAGE POLICIES AND; 4) THE SPECIFIC FACTS OF THE PARTICULAR SITUATION. EACH COVERAGE REQUEST SHOULD BE REVIEWED ON ITS OWN MERITS. MEDICAL DIRECTORS ARE EXPECTED TO EXERCISE CLINICAL JUDGMENT WHERE APPROPRIATE AND HAVE DISCRETION IN MAKING INDIVIDUAL COVERAGE DETERMINATIONS. WHERE COVERAGE FOR CARE OR SERVICES DOES NOT DEPEND ON SPECIFIC CIRCUMSTANCES, REIMBURSEMENT WILL ONLY BE PROVIDED IF A REQUESTED SERVICE(S) IS SUBMITTED IN ACCORDANCE WITH THE RELEVANT CRITERIA OUTLINED IN THE APPLICABLE COVERAGE POLICY, INCLUDING COVERED DIAGNOSIS AND/OR PROCEDURE CODE(S). REIMBURSEMENT IS NOT ALLOWED FOR SERVICES WHEN BILLED FOR CONDITIONS OR DIAGNOSES THAT ARE NOT COVERED UNDER THIS COVERAGE POLICY (SEE "CODING INFORMATION" BELOW). WHEN BILLING, PROVIDERS MUST USE THE MOST APPROPRIATE CODES AS OF THE EFFECTIVE DATE OF THE SUBMISSION. CLAIMS SUBMITTED FOR SERVICES THAT ARE NOT ACCOMPANIED BY COVERED CODE(S) UNDER THE APPLICABLE COVERAGE POLICY WILL BE DENIED AS NOT COVERED. COVERAGE POLICIES RELATE EXCLUSIVELY TO THE ADMINISTRATION OF HEALTH BENEFIT PLANS. COVERAGE POLICIES ARE NOT RECOMMENDATIONS FOR TREATMENT AND SHOULD NEVER BE USED AS TREATMENT GUIDELINES. IN CERTAIN MARKETS, DELEGATED VENDOR GUIDELINES MAY BE USED TO SUPPORT MEDICAL NECESSITY AND OTHER COVERAGE DETERMINATIONS.

CIGNA NATIONAL FORMULARY COVERAGE:

OVERVIEW

Ngenla, Skytrofa, and Sogroya are the available long-acting (weekly) growth hormone (GH) products. All of these agents are indicated for the treatment of growth failure due to inadequate secretion of endogenous GH in pediatric patients. Ngenla is indicated in patients ≥ 3 years of age, Skytrofa is indicated in patient ≥ 1 year of age (and ≥ 11.5 kg), and Sogroya is indicated in patients ≥ 2.5 years of age. Skytrofa and Sogroya have an additional indication for the replacement of endogenous GH in adults with GH deficiency.

POLICY STATEMENT

This Preferred Specialty Management program has been developed to encourage the use of the Preferred Product. For all medications (Preferred and Non-Preferred), the patient is required to meet the respective standard *Prior Authorization Policy* criteria. The program also directs the patient to try the Preferred Product for 6 months, or have an intolerance, prior to the approval of a Non-Preferred Product. Requests for Non-Preferred Products will also be reviewed using the exception criteria (below). All approvals are provided for the durations noted in the respective standard *Prior Authorization Policy* criteria. If the patient meets the standard *Prior Authorization Policy* criteria but has not met the applicable criteria for the Non-Preferred Product, approval for the Preferred Product will be authorized. All reviews will be directed to a clinician (i.e., pharmacist) for verification of criteria.

National Preferred Formulary

Preferred Product: Ngenla

Non-Preferred Products: Skytrofa, Sogroya

National Preferred Flex Formulary

Preferred Product: Ngenla

Non-Preferred Products: Skytrofa, Sogroya

Basic Formulary

Preferred Products: Ngenla, Skytrofa

Non-Preferred Product: Sogroya

Growth Disorders – Growth Hormone Long-Acting Products Preferred Specialty Management Policy non-preferred product(s) is(are) covered as medically necessary when the following non-preferred product exception criteria is(are) met. Any other exception is considered not medically necessary.

Non-Preferred Product Exception Criteria

Non-	Exception Criteria		
Preferred			
Product			
Skytrofa	1. National Preferred Formulary and National Preferred Flex		
	Formulary. Approve for 1 year if the patient meets BOTH of the		
	following (A <u>and</u> B):		
	A) Patient meets the standard <i>Growth Disorders – Skytrofa</i>		
	Prior Authorization Policy criteria; AND		
	B) Patient meets ONE of the following (i, ii, <u>or</u> iii):		
	i. Patient is < 3 years of age; OR		
	ii. Patient is ≥ 18 years of age; OR		
	iii. Patient meets ONE of the following (a <u>or</u> b):		
	a. Patient has tried Ngenla for 6 months; OR		
	b. Patient has experienced an intolerance with Ngenla.		
	C) If the patient has met the respective standard Prior		
	Authorization criteria (1A), but has <u>not</u> met exception		
	criteria (1B), approve Ngenla.		
	2. Basic Formulary. Approve for 1 year if the patient meets the		
	standard Growth Disorders – Skytrofa Prior Authorization Policy		
	criteria.		

Preferred	Exception Criteria		
Product			
Sogroya 1. I	National Preferred Formulary and National Preferred Flex Formulary. Approve for 1 year if the patient meets BOTH of the following (A and B): A) Patient meets the standard Growth Disorders − Sogroya Prior Authorization Policy criteria; AND B) Patient meets ONE of the following (i, ii, or iii): i. Patient is < 3 years of age; OR ii. Patient is ≥ 18 years of age; OR iii. Patient meets ONE of the following (a or b): a. Patient has tried Ngenla for 6 months; OR b. Patient has experienced an intolerance with Ngenla. C) If the patient has met the respective standard Prior Authorization criteria (1A), but has not met exception criteria (1B), approve Ngenla. Basic Formulary. Approve for 1 year if the patient mees BOTH of the following (A and B): A) Patient meets the standard Growth Disorders − Sogroya Prior Authorization Policy criteria; AND B) Patient meets ONE of the following (i or ii): i. Patient has tried ONE of Ngenla or Skytrofa for 6 months; OR ii. Patient has experienced an intolerance with Ngenla or Skytrofa. C) If the patient has met the respective standard Prior Authorization criteria (2A), but has not met exception criteria (2B), approve Ngenla or Skytrofa.		

REFERENCES

- 1. Ngenla[™] subcutaneous injection [prescribing information]. New York, NY: Pfizer; June 2023
- 2. Skytrofa® subcutaneous injection [prescribing information]. Princeton, NJ: Ascendis Pharma; May 2024.
- 3. Sogroya® subcutaneous injection [prescribing information]. Plainsboro, NJ: Novo Nordisk; April 2023.

HISTORY

Type of	Summary of Changes	Review Date
Revision		
New Policy		11/01/2023
Annual	No criteria changes.	11/20/2024
Revision		
Selected	National Preferred Formulary and National Preferred Flex	08/13/2025
Revision	Formulary:	
	An exception was added to Skytrofa criteria to approve if the patient is \geq 18 years of age (if the PA was met).	
	Basic Formulary: An exception was removed from Sogroya	
	criteria to approve if the patient is ≥ 18 years of age (if the PA was	
	met).	

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