

If the following information is not complete, correct, or legible, the SA process can be delayed.

Please use one form per member.

MEMBER INFORMATION

Last name:

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First name:

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Medicaid ID number:

--	--	--	--	--	--	--	--	--	--	--	--	--	--	--

Date of birth:

--	--	--	--	--	--	--	--	--	--	--	--	--	--	--

Gender: ☐ Male ☐ Female**PRESCRIBER INFORMATION**

Last name:

--	--	--	--	--	--	--	--	--	--	--	--	--	--	--

First name:

--	--	--	--	--	--	--	--	--	--	--	--	--	--	--

NPI number:

--	--	--	--	--	--	--	--	--	--	--	--	--	--	--

Phone number:

--	--	--	--	--	--	--	--	--	--	--	--	--	--	--

Fax number:

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Is the drug prescribed by or in consultation with a specialty?

☐ Endocrinologist ☐ Nephrologist**DRUG INFORMATION**

Drug name/Form: _____

Strength: _____

Quantity per day: _____

All growth hormone medications require the submission of a Clinical Service Authorization

Preferred medications:

☐ Genotropin[®] ☐ Norditropin FlexPro[®]

Non-Preferred medications:

☐ Humatrope[®] cartridge/vial☐ Nutropin AQ[®] NuSpin[®]☐ Nutropin AQ[®] cartridge/vial☐ Omnitrope[®] cartridge/vial☐ Saizen[®] cartridge/vial☐ Serostim[®] vial☐ Skytrofa[™] Syringe☐ Zomacton[®] vial☐ Zorbtive[®] vial

If requesting a non-preferred agent, please document why a preferred agent cannot be used:

(Form continued on next page.)

Member's last name:

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Member's first name:

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Section B: Pediatric GH Deficiency

9. Did the member have a GH response of less than 10 ng/mL (or otherwise abnormal as determined by the lab) of at least 2 GH stimulation tests LFTs?

☐ Yes ☐ No

Action required: *If YES, please attach documentation of stimulation test results.*

10. Did member have a GH response of less than 15 ng/mL on at least 1 GH stimulation test?

☐ Yes ☐ No

Action required: *Please attach documentation of GH stimulation test result. If YES, indicate results.*

11. Does the member have a defined CNS pathology, history of cranial irradiation or genetic condition associated GH deficiency?

☐ Yes ☐ No

12. Does the member have both IGF-1 and IGFBP-3 levels below normal for age and gender?

☐ Yes ☐ No

Action required: *If YES, please attach documentation from the medical record showing IGF-1 and IGFBP-3 levels below normal.*

13. Does the member have 2 or more documented pituitary hormone deficiencies other than GH?

☐ Yes ☐ No

14. Did the member have an abnormally low GH level in association with neonatal hypoglycemia?

☐ Yes ☐ No

Action required: *If YES, please attach documentation of GH level.*

Section C: Pediatric Chronic Kidney Disease/Chronic Renal Insufficiencies

15. Does the member have any of the following? Indicate any/all the apply:

☐ Creatinine clearance of 75 mL/min/1.73 m² or less ☐ Dialysis dependency
☐ Serum creatinine greater than 3.0 g/dL ☐ None of the above

Section D: Pediatric Chronic Kidney Disease

16. Is this request for a new start, restart (re-initiation) or continuation of GH therapy?

☐ New start, *no further questions* ☐ Restart ☐ Continuation

17. Was GH therapy previously approved for this member?

☐ Yes ☐ No

18. What is the member's current height in inches? _____

Action required: *Please attach documentation from the medical record of current height.
 If Restart, no further questions.*

19. Is the member's growth velocity at least 2 cm per year while on GH therapy?

☐ Yes ☐ No

Action required: *If YES, please attach documentation from medical record supporting growth velocity of at least 2 cm/year. (Form continued on next page.)*

