

Prior Authorization Form

GEHA FEDERAL - STANDARD OPTION
Growth Hormones (FA-PA)

This fax machine is located in a secure location as required by HIPAA regulations.
Complete/review information, sign and date. Fax signed forms to CVS/Caremark at **1-888-836-0730**.
Please contact CVS/Caremark at **1-855-240-0536** with questions regarding the prior authorization process.
When conditions are met, we will authorize the coverage of Growth Hormones (FA-PA).

Drug Name (select from list of drugs shown)

Genotropin (somatropin)	Nutropin (somatropin)	Nutropin AQ (somatropin)
Omnitrope (somatropin)	Saizen (somatropin)	Zomacton (somatropin)

Quantity	Frequency	Strength
Route of Administration	Expected Length of Therapy	

Patient Information

Patient Name: _____
Patient ID: _____
Patient Group No.: _____
Patient DOB: _____
Patient Phone: _____

Prescribing Physician

Physician Name: _____
Physician Phone: _____
Physician Fax: _____
Physician Address: _____
City, State, Zip: _____

Diagnosis: _____ **ICD Code:** _____

Comments: _____

Please circle the appropriate answer for each question.

- | | |
|---|---|
| 1. Is the requested product being prescribed for a patient with chronic kidney disease? | <input type="checkbox"/> Y <input type="checkbox"/> N |
| 2. Is the requested product Nutropin AQ? | <input type="checkbox"/> Y <input type="checkbox"/> N |
| 3. Is the requested product being prescribed for a patient with Prader-Willi syndrome? | <input type="checkbox"/> Y <input type="checkbox"/> N |
| 4. Is the requested product Genotropin or Omnitrope? | <input type="checkbox"/> Y <input type="checkbox"/> N |
| 5. The preferred products for your patient's plan are Humatrope and Norditropin. Can the patient's treatment be switched to Humatrope or Norditropin? | <input type="checkbox"/> Y <input type="checkbox"/> N |

6. Did the patient experience documented inadequate treatment response(s) and/or intolerable adverse event(s) to previous trials with BOTH of the preferred products (Humatrope and Norditropin)? ACTION REQUIRED: IF 'YES', ATTACH CHART NOTES DETAILING THE OUTCOME(S) OF TREATMENT AND/OR INTOLERABLE ADVERSE EVENT(S) EXPERIENCED FROM TREATMENT WITH BOTH OF THE PREFERRED PRODUCTS.

 Y N

7. Does the patient have a documented contraindication to any of the preferred products (Humatrope or Norditropin) or any of their components? ACTION REQUIRED: IF 'YES', ATTACH DOCUMENTATION DETAILING THE CONTRAINDICATION(S) TO TREATMENT WITH ANY OF THE PREFERRED PRODUCTS.

 Y N

I affirm that the information given on this form is true and accurate as of this date.

Prescriber (Or Authorized) Signature and Date