



Please complete ALL information below and fax your request to 1-888-671-5285

Omnitrope® Prior Authorization Request Form (Page 1 of 4)

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Member Information (required)			Provider Information (required)		
Member Name:			Provider Name:		
Insurance ID#:			NPI#:		Specialty:
Date of Birth:			Office Phone:		
Street Address:			Office Fax:		
City:	State:	Zip:	Office Street Address:		
Phone:			City:	State:	Zip:
Medication Information (required)					
Medication Name:			Strength:		Dosage Form:
<input type="checkbox"/> Check if generic substitution is acceptable			Directions for Use:		
<input type="checkbox"/> Check if request is for continuation of therapy					
Clinical Information (required)					
Select the diagnosis below: <input type="checkbox"/> Pediatric growth hormone deficiency <input type="checkbox"/> Growth hormone deficiency in adults <input type="checkbox"/> Growth hormone deficiency in transition phase adolescents <input type="checkbox"/> Isolated growth hormone deficiency in adults <input type="checkbox"/> Pediatric growth failure associated with chronic renal insufficiency <input type="checkbox"/> Prader-Willi Syndrome <input type="checkbox"/> Short-stature homeobox (SHOX) gene deficiency <input type="checkbox"/> Small for gestational age (SGA) <input type="checkbox"/> Turner syndrome or Noonan syndrome <input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____					
Prescriber's Specialty: Select if the requested medication is prescribed by or in consultation with one of the following specialists: <input type="checkbox"/> Endocrinologist <input type="checkbox"/> Nephrologist					
For pediatric growth hormone deficiency, also answer the following: Is the patient an infant < 4 months of age? <input type="checkbox"/> Yes <input type="checkbox"/> No Does the infant have growth deficiency? <input type="checkbox"/> Yes <input type="checkbox"/> No Does the patient have history of neonatal hypoglycemia associated with pituitary disease? <input type="checkbox"/> Yes <input type="checkbox"/> No Does the patient have panhypopituitarism? <input type="checkbox"/> Yes <input type="checkbox"/> No Select if the diagnosis of pediatric GH deficiency is confirmed by the patient's height as documented by the following (utilizing age and gender growth charts related to height): <input type="checkbox"/> Height > 2.0 standard deviations [SD] below mid-parental height <input type="checkbox"/> Height > 2.25 SD below population mean (below the 1.2 percentile for age and gender) Is the patient's growth velocity > 2 SD below mean for age and gender? <input type="checkbox"/> Yes <input type="checkbox"/> No Does the patient have delayed skeletal maturation of > 2 SD below mean for age and gender (e.g., delayed > 2 years compared with chronological age)? <input type="checkbox"/> Yes <input type="checkbox"/> No Is there documentation the patient's bone age is < 16 years for males or < 14 years for females? <input type="checkbox"/> Yes <input type="checkbox"/> No					
<continued on the next page>					

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<continuation of pediatric growth hormone deficiency>

Select if the patient has undergone provocative GH stimulation tests with the following: **(Document the GH response)**

- Arginine Peak value: _____ mcg/L
- Clonidine Peak value: _____ mcg/L
- Glucagon Peak value: _____ mcg/L
- Insulin Peak value: _____ mcg/L
- Levodopa Peak value: _____ mcg/L
- Growth hormone releasing hormone Peak value: _____ mcg/L

For patients less than 1 year of age, select if the following is below the age and gender adjusted normal range as provided by the physician's lab: **(Document the specified lab value and reference range)**

- Insulin-like growth factor 1 (IGF-1/Somatomedin-C) IGF-1/Somatomedin-C level: _____ Reference range: _____
- Insulin growth factor binding protein-3 (IGFBP-3) IGFBP-3 level: _____ Reference range: _____

Reauthorization:

Please document that the patient has had a height increase of at least 2 cm/year over the previous year of treatment below:

Previous height: _____ Date obtained: _____
Current height: _____ Date obtained: _____

Has the expected adult height been reached? Yes No

Document the expected adult height goal: _____

For growth hormone (GH) deficiency in adults, also answer the following:

Are there clinical records supporting a diagnosis of childhood-onset GH deficiency? Yes No

Does the patient have adult-onset GH deficiency? Yes No

Are there clinical records documenting that hormone deficiency is a result of hypothalamic-pituitary disease from organic or known causes (e.g., damage from surgery, cranial irradiation, head trauma, or subarachnoid hemorrhage)? Yes No

Select if the patient has undergone one of the following GH stimulation tests to confirm adult GH deficiency and the peak GH value is as follows:

- Insulin tolerance test (ITT) ≤ 5 mcg/L
- Arginine & GH-releasing hormone (GHRH+ARG) ≤ 11 mcg/L if body mass index (BMI) is < 25 kg/m²; ≤ 8 mcg/L if BMI is ≥ 25 and < 30 kg/m²; ≤ 4 mcg/L if BMI is ≥ 30 kg/m²
- Glucagon ≤ 3 mcg/L
- Arginine (ARG) ≤ 0.4 mcg/L

Select if there is documentation the patient has deficiency of the following anterior pituitary hormones:

- Adrenocorticotrophic hormone (ACTH)
- Prolactin
- Follicle-stimulating hormone/luteinizing hormone (FSH/LH)
- Thyroid stimulating hormone (TSH)

Does the patient have an IGF-1/Somatomedin-C level below the age and gender adjusted normal range as provided by the physician's lab? Yes No

Reauthorization:

Is there evidence of ongoing monitoring as demonstrated by documentation within the past 12 months of an IGF-1/Somatomedin-C level? Yes No

For isolated growth hormone deficiency in adults, also answer the following:

Is there documentation the patient has deficiency of GH defined by a failure to produce a peak serum GH level of > 5 mcg/L after provocative pharmacologic stimulation by two of the following tests: Insulin, L-arginine, and/or glucagon? Yes No

Reauthorization:

Is there evidence of ongoing monitoring as demonstrated by documentation within the past 12 months of an IGF-1/Somatomedin-C level? Yes No



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For growth hormone (GH) deficiency in transition phase adolescents, also answer the following:

Has the expected adult height been reached? Yes No

Are the patient's epiphyses closed on bone radiograph? Yes No

Select if there is documentation the patient has high risk of GH deficiency due to GH deficiency in childhood from one of the following:

- Embryopathic/congenital defects
- Irreversible structural hypothalamic-pituitary disease
- Genetic mutations
- Panhypopituitarism
- Deficiency of three or more of the following anterior pituitary hormones: ACTH, TSH, Prolactin, FSH/LH

Does the patient have an IGF-1/Somatomedin-C level below the age and gender adjusted normal range as provided by the physician's lab? Yes No

Is the patient at low risk of severe GH deficiency (e.g., due to isolated and/or idiopathic deficiency)? Yes No

Has GH therapy been discontinued for at least 1 month? Yes No

Select if the patient has undergone one of the following GH stimulation tests after discontinuation of therapy for at least 1 month and the peak GH value is as follows:

- Insulin tolerance test (ITT) ≤ 5 mcg/L
- Arginine & GH-releasing hormone (GHRH+ARG) ≤ 11 mcg/L if body mass index (BMI) is < 25 kg/m²; ≤ 8 mcg/L if BMI is ≥ 25 and < 30 kg/m²; ≤ 4 mcg/L if BMI is ≥ 30 kg/m²
- Glucagon ≤ 3 mcg/L
- Arginine (ARG) ≤ 0.4 mcg/L

Reauthorization:

Is there evidence the patient has had a positive response to therapy (e.g., increase in total lean body mass, exercise capacity or IGF-1 and IGFBP-3 levels)? Yes No

For pediatric growth failure associated with chronic renal insufficiency, also answer the following:

Is there documentation the patient's bone age is < 16 years for males or < 14 years for females? Yes No

Reauthorization:

Please document that the patient has had a height increase of at least 2 cm/year over the previous year of treatment below:

Previous height: _____ Date obtained: _____
Current height: _____ Date obtained: _____

Has the expected adult height been reached? Yes No

Document the expected adult height goal: _____

For Prader-Willi syndrome, also answer the following:

Reauthorization:

Is there evidence the patient has had a positive response to therapy (e.g., increase in total lean body mass, decrease in fat mass)? Yes No

Please document that the patient has had a height increase of at least 2 cm/year over the previous year of treatment below:

Previous height: _____ Date obtained: _____
Current height: _____ Date obtained: _____

Has the expected adult height been reached? Yes No

Document the expected adult height goal: _____

For short-stature homeobox (SHOX) gene deficiency, also answer the following:

Does the patient have a diagnosis of pediatric growth failure with short stature homeobox (SHOX) gene deficiency as confirmed by genetic testing? Yes No

Is there documentation the patient's bone age is < 16 years for males or < 14 years for females? Yes No

Reauthorization:

Please document that the patient has had a height increase of at least 2 cm/year over the previous year of treatment below:

Previous height: _____ Date obtained: _____
Current height: _____ Date obtained: _____

Has the expected adult height been reached? Yes No

Document the expected adult height goal: _____



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For small for gestational age (SGA), also answer the following:

Select if the diagnosis of SGA is based on demonstration of catch up growth failure in the first 24 months of life using a 0-36 month growth chart as confirmed by one of the following:

- Patient's **birth weight** was below the 3rd percentile for gestational age (> 2 SD below population mean)
- Patient's **birth length** was below the 3rd percentile for gestational age (> 2 SD below population mean)

Does patient's height remain \leq the 3rd percentile (> 2 SD below population mean)? Yes No

Reauthorization:

Please document that the patient has had a height increase of at least 2 cm/year over the previous year of treatment below:

Previous height: _____ Date obtained: _____

Current height: _____ Date obtained: _____

Has the expected adult height been reached? Yes No

Document the expected adult height goal: _____

For Turner syndrome (gonadal dysgenesis) or Noonan syndrome, also answer the following:

Is there documentation the patient's bone age is < 16 years for males or < 14 years for females? Yes No

Is the patient's height below the 5th percentile on growth charts for age and gender? Yes No

Reauthorization:

Please document that the patient has had a height increase of at least 2 cm/year over the previous year of treatment below:

Previous height: _____ Date obtained: _____

Current height: _____ Date obtained: _____

Has the expected adult height been reached? Yes No

Document the expected adult height goal: _____

Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?

Please note: This request may be denied unless all required information is received.