**INTRODUCTION**

- Growth hormone (GH) affects many of the metabolic processes carried out by somatic cells, most notably increasing body mass. Overall growth is stimulated by GH therapy; however, the effects are not evenly distributed among protein, lipid and carbohydrate compartments. Specifically, body protein content and bone mass increase, total body fat content decreases, and there is an increase in plasma and liver lipid content due to the mobilization of free fatty acids from peripheral fat stores. Another physiological effect of GH is stimulation of cartilage growth (Molitch et al 2011).

- Growth hormone deficiency (GHD) in pediatric patients is a clinical diagnosis that is confirmed by biochemical testing. A patient’s growth patterns are compared to the established norms. The clinical manifestations of GHD vary depending on whether a patient has complete or partial deficiency. In complete deficiency, pediatric patients present with early severe growth failure, delayed bone age, central disposition of body fat and very low serum concentrations of GH, insulin-like growth factor-1 (IGF-1) and IGF binding protein-3. These patients are also more prone to hypoglycemia, prolonged jaundice, microphallus in males and giant cell hepatitis. GHD in pediatric patients with partial deficiency may be more difficult to diagnose, as these manifestations may not be as obvious (Molitch et al 2011).

- Once a diagnosis of GHD is confirmed in pediatric patients, GH therapy should be initiated and continued until cessation of linear growth. Therapy should be initiated as soon as possible, as evidence demonstrates that growth response is more robust when GH therapy is started at a younger age (Molitch et al 2011).

- Several preparations of GH are currently available for use in pediatric patients. Recombinant GH preparations, administered by subcutaneous (SC) injection, are currently the most widely utilized. Due to the variability in individual response to therapy, after initial dosing, the dose of GH is adjusted based on growth response and IGF-1 level. While not universally supported, the therapeutic goal of therapy is to achieve a level of IGF-1 that is slightly higher than average, because growth velocity is typically greatest at these levels. A patient’s growth velocity, as compared to a similar population, should also be monitored to determine if the growth response is adequate (Molitch et al 2011).

- Possible explanations for an inadequate response to GH therapy include poor adherence, incorrect diagnosis of GHD, subtherapeutic dose of GH, or concurrent mild GH insensitivity. In pediatric patients, GH therapy is typically continued at least until linear growth is nearly complete (eg, decreased to less than 2.5 centimeters per year). At this point, retesting for GHD should occur to determine if GH therapy should be continued into adulthood (Molitch et al 2011).

- The majority of pediatric patients with idiopathic, isolated GHD in their childhood have normal GH secretion during late adolescence and young adulthood. In contrast, pediatric patients with genetic GHD, multiple pituitary hormone deficiencies, and/or those with structural defects in the hypothalamic-pituitary region rarely recover the ability to secrete GH as an adult. Therefore, retesting may not be required (Molitch et al 2011).

- GHD may also occur in adult patients. Approximately 15% to 20% of adult-onset GHD represents the continuation of childhood-onset GHD into maturity; the remainder is adult-onset acquired from damage to the pituitary gland or hypothalamus. GHD is associated with increased metabolic syndrome, increased cardiovascular morbidity and mortality rates, reduced lean body mass, increased abdominal adiposity, early atherosclerosis, dyslipidemia, coagulation abnormalities, insulin resistance, decreased bone mineral density, and a decreased quality of life (Reed et al 2013). The role of GH therapy in adults is not as clear as it is in pediatric patients in whom therapy is required for normal growth. There is evidence to demonstrate that when used in adult patients with GHD, GH therapy increases muscle mass and decreases body fat. Evidence of other potential beneficial effects of GH therapy in adults is not as well established and includes improvement in bone mineral density, sense of well-being, muscle strength, and lipid profile. GH therapy can be considered in adult patients with severe clinical manifestations and unequivocal evidence of GHD due to organic disease of childhood- or adult-onset (Molitch et al 2011).

- All of the GH preparations contain somatropin, otherwise known as recombinant human GH. The various preparations are Food and Drug Administration (FDA)-approved for use in a variety of pediatric conditions associated with a failure in growth, including chronic kidney disease (CKD), Turner syndrome, being born small for gestational age, Prader-Willi syndrome, mutations in the Short Stature Homeobox gene, Noonan syndrome, and idiopathic short stature.
The majority of preparations are also indicated for the treatment of GHD in adults. Of note, Serostim is FDA-approved solely for the treatment of human immunodeficiency virus-associated wasting or cachexia in adults, while Zorbtive is approved for the treatment of short bowel syndrome in patients receiving specialized nutritional support. Specific FDA-approved indications for the various GH preparations are outlined in Table 2. All of the available GH preparations are available for SC injection, and there are currently no generics available within the class.

GH preparations are available in various formulations, and several delivery devices are available. The dosing device may be a factor in patient adherence with the prescribed regimen.

Medispan Class: Growth Hormones

### Table 1. Medications Included Within Class Review

<table>
<thead>
<tr>
<th>Drug</th>
<th>Generic Availability</th>
</tr>
</thead>
<tbody>
<tr>
<td>Genotropin (somatropin)</td>
<td>-</td>
</tr>
<tr>
<td>Humatrope (somatropin)</td>
<td>-</td>
</tr>
<tr>
<td>Norditropin Flexpro (somatropin)</td>
<td>-</td>
</tr>
<tr>
<td>Nutropin AQ (somatropin)</td>
<td>-</td>
</tr>
<tr>
<td>Omnitrope (somatropin)</td>
<td>-</td>
</tr>
<tr>
<td>Saizen (somatropin)</td>
<td>-</td>
</tr>
<tr>
<td>Serostim (somatropin)</td>
<td>-</td>
</tr>
<tr>
<td>Zomacton (somatropin)</td>
<td>-</td>
</tr>
<tr>
<td>Zorbtive (somatropin)</td>
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</tr>
</tbody>
</table>

*(Drugs@FDA 2019, Orange Book: Approved Drug Products with Therapeutic Equivalence Evaluations 2019)*

### INDICATIONS

Table 2. Food and Drug Administration Approved Indications

<table>
<thead>
<tr>
<th>Indication</th>
<th>Genotropin</th>
<th>Humatrope</th>
<th>Norditropin Flexpro</th>
<th>Nutropin AQ</th>
<th>Omnitrope</th>
<th>Saizen</th>
<th>Serostim</th>
<th>Zomacton</th>
<th>Zorbtive</th>
</tr>
</thead>
<tbody>
<tr>
<td>Growth failure associated with chronic renal insufficiency before renal transplant</td>
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<td></td>
<td></td>
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<td></td>
<td></td>
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<tr>
<td>Growth failure associated with Noonan syndrome</td>
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<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Growth failure associated with Prader-Willi syndrome</td>
<td>✓</td>
<td>✓</td>
<td></td>
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<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Growth failure associated with short-stature homeobox-containing gene deficiency</td>
<td>✓</td>
<td></td>
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<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Growth failure associated with Turner syndrome</td>
<td>✓</td>
<td>✓</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Growth failure in children born small for gestational age</td>
<td>✓</td>
<td>✓</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Growth hormone deficiency</td>
<td>✓</td>
<td>✓</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Idiopathic short stature</td>
<td>✓</td>
<td>✓</td>
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<td></td>
<td></td>
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</tr>
</tbody>
</table>

Data as of February 20, 2019 JA/U

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Indication | Genotropin | Humatrope | Norditropin Flexpro | Nutropin AQ | Omnitrope | Saizen | Serostim | Zomacton | Zortive |
--- | --- | --- | --- | --- | --- | --- | --- | --- | --- |
Human immunodeficiency virus-associated wasting or cachexia | ✔ | | | | | | | | |
Treatment of short bowel syndrome in patients receiving nutritional support | | | | | | | | | ✔ |


- Information on indications, mechanism of action, pharmacokinetics, dosing, and safety has been obtained from the prescribing information for the individual products, except where noted otherwise.

### CLINICAL EFFICACY SUMMARY

- There are limited head-to-head clinical trials comparing different GH preparations to one another.
- Clinical data support the use of GH for the treatment of growth failure associated with chronic renal insufficiency. A meta-analysis of 16 RCTs (N = 809) evaluating the effects of GH in children with CKD found that patients who were treated with GH had a greater increase in mean height velocity (3.88 cm) than those who received either no treatment or placebo after 1 year (Hodson et al 2012). A retrospective, matched control cohort study found that long-term therapy with GH (mean 4.2 years) reduced linear growth deceleration in children with CKD and improved final height (Bizzarri et al 2018).
- Clinical trials have demonstrated efficacy of GH for the treatment of growth failure in patients with Noonan syndrome. A randomized controlled trial evaluating GH in patients with Noonan syndrome found a positive effect of GH on linear growth. Specifically, there was a significantly greater change in height standard deviation score, and bone maturation was accelerated with GH compared to no treatment. In this trial, data also suggest that once treatment with GH is discontinued, “catch-down” (artificially stimulated growth declines once GH is discontinued) growth can occur (Noordam et al 2001). In a follow-up analysis of 29 patients treated with GH for a median of 6.4 years, a total of 22 children reached an adult height in the normal range (Noordam et al 2008). In a study of 65 patients enrolled in the National Cooperative Growth Study (NCGS) database, it was found that treatment with GH led to gains over predicted height of 9.2 cm in females and 10.9 cm in males (Romano et al 2009).
- Humatrope demonstrated efficacy in increasing first-year height velocity in patients with Short Stature Homeobox-containing gene deficiency when compared to no treatment (p < 0.0001) (Blum et al 2007).
- Several clinical trials have demonstrated that GH significantly increases the growth rate of pediatric patients with Turner syndrome. Overall, various dose ranging trials did not consistently demonstrate a superior weight-based GH dosing regimen over another; all doses of GH were beneficial. In addition, data suggested that increases in height are greatest during the first year of therapy (Baxter et al 2007, Bertrand et al 1996, Massa et al 1995, Nienhuis et al 1993, Sas et al 1999a, Takano et al 1989a, Takano et al 1989b, Takano et al 1993, Takano 1995, van Pareren et al 2003, van Teunenbroek et al 1996). A Cochrane Review of 4 randomized controlled trials demonstrated that GH (0.3 to 0.375 mg/kg/week) increased short-term growth in patients with Turner syndrome by approximately 3 cm during the first year of treatment. Despite the increase, the final height achieved was still below the normal range (Baxter et al 2007).
that response to GH therapy is dose-dependent, and higher doses of GH resulted in additional gain (de Zegher et al 1996, de Zegher et al 2005).


- A systematic review and meta-analysis of 54 placebo-controlled, randomized controlled trials enrolling over 3400 patients found that GH therapy was associated with reduced body fat and increased lean mass in adults with GHD (Hazem et al 2012). Eleven of 16 trials that assessed quality of life outcomes reported positive outcomes, but a meta-analysis was not possible. Furthermore, results from meta-analyses and randomized controlled trials have demonstrated that treatment with GH was associated with improved cardiac function and bone mineral density (Barake et al 2014, Davidson et al 2004, Maison et al 2003). However, there are currently conflicting data with regard to the effect of GH on cognitive function, quality of life, and exercise capacity (Arwert et al 2005, Falleti et al 2006, Rubeck et al 2009, Widdowson, 2010).

- In patients with human immunodeficiency virus-associated wasting, Serostim has been shown to increase body weight, lean body mass, and work output. However, effects on quality of life were variable (Moyle et al 2004, Schambelan et al 1996).

- A meta-analysis assessed the safety and efficacy of GH with or without glutamine supplementation for adult patients with short bowel syndrome; 5 studies were included in the review. Human GH with or without glutamine appeared to provide benefit in terms of increased weight (median [MD] 1.66 kg; 95% confidence interval [CI], 0.69 to 2.63; p = 0.0008), lean body mass (MD 1.93 kg; 95% CI, 0.97 to 2.9; p = 0.0001), energy absorption (MD 4.42 Kcal; 95% CI, 0.26 to 8.58; p = 0.04) and nitrogen absorption (MD 44.85 g; 95% CI, 0.2 to 9.49; p = 0.04) for patients with short bowel syndrome. One randomized controlled trial which focused on parenteral nutrition (PN) requirements demonstrated decreased PN volume, calories, and number of infusions in patients who received GH with or without glutamine supplementation. Only patients who received GH with glutamine maintained statistically significant PN reductions at 3-month follow-up. The results suggested a positive effect of GH on weight gain and energy absorption. However, after cessation of therapy, the effects returned to baseline in the majority of the trials (Wales et al 2010).

**CLINICAL GUIDELINES**


- For adult patients, treatment guidelines recommend the use of GH therapy in patients with clinical features suggestive of adult GHD and biochemically proven evidence of adult GHD (Cook et al 2009). Therapy should be individualized independent of body weight. The dose of GH should be low initially and gradually increased to the minimally effective dose that normalizes IGF-1 levels without side effects (Cook et al 2009, Mollitch et al 2011). The 2009 American Association of Clinical Endocrinologists guidelines state that no evidence exists to support any specific GH product over another (Cook et al 2009).

- Small studies evaluating the use of GH in short bowel syndrome have yielded conflicting results; methodological differences limit definitive conclusions on the efficacy of GH. In carefully selected patients who are candidates for growth factor treatment, the glucagon-like peptide-2 analog, teduglutide, is recommended as first-line therapy (Pironi et al 2016).
SAFETY SUMMARY

- Contraindications to GH products include active malignancy, diabetic retinopathy, hypersensitivity to the agent or any of its excipients, acute critical illness, and use for growth promotion in children with closed epiphyses. Somatropin is also contraindicated in children with Prader-Willi syndrome who are severely obese, have severe respiratory impairment, or have a history of upper airway obstruction or sleep apnea (Genotropin, Humatrope, Norditropin Flexpro, Nutropin AQ, Omnitrope, Saizen, Zomacton).

- Key Warnings/Precautions:
  ○ Somatropin may contribute to the increased mortality in patients with acute critical illness due to complications from open heart surgery, abdominal surgery, accidental trauma, or respiratory failure.
  ○ Somatropin may increase progression or recurrence of intracranial neoplasms, particularly meningiomas in patients treated with radiation to the head for their first neoplasm.
  ○ Undiagnosed or untreated hypothyroidism may impair optimal response to somatropin.
  ○ Somatropin may decrease insulin sensitivity, and previously undiagnosed diabetes mellitus may be unmasked during treatment.
  ○ Intracranial hypertension and pancreatitis have been reported with somatropin treatment.
  ○ Slipped capital femoral epiphyses and scoliosis can occur in pediatric patients.
  ○ Fluid retention has been associated with somatropin in adult patients.
  ○ Increases in serum levels of inorganic phosphorous, alkaline phosphatase, parathyroid hormone and IGF-1 may occur.
  ○ Tissue atrophy may occur when somatropin is SC administered at the same site over a long period of time.
  ○ Somatropin may reduce serum cortisol levels or unmask central hypoadrenalism in patients at risk for pituitary hormone deficiency.

- Adverse Drug Events: Arthralgia, myalgia, edema, carpal tunnel syndrome, paresthesia, hyperglycemia, headaches, lipoatrophy, and injection site reactions.

- Drug Interactions: Estrogens, glucocorticoids, and insulin or other hypoglycemic agents.

DOSING AND ADMINISTRATION

<table>
<thead>
<tr>
<th>Drug</th>
<th>Available Formulations</th>
<th>Route</th>
<th>Usual Recommended Frequency</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>Genotropin (somatropin)</td>
<td>Injection</td>
<td>SC</td>
<td>Weekly dose divided into 6 or 7 injections to Daily</td>
<td>Injections should be rotated to help prevent lipoatrophy.</td>
</tr>
<tr>
<td>Humatrope (somatropin)</td>
<td>Injection</td>
<td>SC</td>
<td>Weekly dose divided into 6 or 7 injections to Daily</td>
<td>Injections should be rotated to help prevent lipoatrophy.</td>
</tr>
<tr>
<td>Norditropin Flexpro (somatropin)</td>
<td>Injection</td>
<td>SC</td>
<td>Weekly dose divided into 6 or 7 injections to Daily</td>
<td>Injections should be rotated to help prevent lipoatrophy.</td>
</tr>
<tr>
<td>Nutropin AQ (somatropin)</td>
<td>Injection</td>
<td>SC</td>
<td>Weekly dose divided into 3 to 7 injections to Daily</td>
<td>Injections should be rotated to help prevent lipoatrophy.</td>
</tr>
<tr>
<td>Omnitrope (somatropin)</td>
<td>Injection</td>
<td>SC</td>
<td>Weekly dose divided into 6 or 7 injections to Daily</td>
<td>Injections should be rotated to help prevent lipoatrophy.</td>
</tr>
<tr>
<td>Saizen (somatropin)</td>
<td>Injection</td>
<td>SC</td>
<td>Weekly dose divided into 3, 6, or 7 injections to Daily</td>
<td>Injections should be rotated to help prevent lipoatrophy.</td>
</tr>
<tr>
<td>Sersotim (somatropin)</td>
<td>Injection</td>
<td>SC</td>
<td>Daily</td>
<td>Injections should be rotated to avoid local irritation.</td>
</tr>
<tr>
<td>Zomacton (somatropin)</td>
<td>Injection</td>
<td>SC</td>
<td>Weekly dose divided into 3, 6, or 7 injections to Daily</td>
<td>Injections should be rotated to help prevent lipoatrophy.</td>
</tr>
<tr>
<td>Zorbtive (somatropin)</td>
<td>Injection</td>
<td>SC</td>
<td>Daily</td>
<td>Injections should be rotated to help prevent lipoatrophy. Dosage titration is recommended for fluid retention</td>
</tr>
</tbody>
</table>

Table 3. Dosing and Administration

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CONCLUSION


- For adult patients, treatment guidelines recommend the use of GH therapy patients with clinical features suggestive of adult GHD and biochemically proven evidence of adult GHD. There should be a careful evaluation of the benefits and risks specific to the individual. The 2009 American Association of Clinical Endocrinologists Guidelines state that no evidence exists to support any specific GH product over another (Cook et al 2009, Flesnerl et al 2016).

- There are several GH preparations currently available, which all contain somatropin (recombinant human GH). The various preparations are equally biopotent and have the same natural sequence structure (Rogol et al 2018). Differences between products such as device features, dose increments, requirement for reconstitution, and requirement for refrigeration may influence individual patient preferences. All of the available GH preparations are available for SC injection, and there are currently no generics available within the class.

- Common adverse reactions that may be observed with GH therapy include arthralgia, myalgia, edema, carpal tunnel syndrome, paresthesia, hyperglycemia, headaches, lipoatrophy, and injection site reactions.

REFERENCES


• Takano K, Shizume K, Hibi I. Treatment of 94 patients with Turner's syndrome with recombinant human growth hormone (SM-9500) for two years—the results of a multicentric study in Japan. Committee for the Treatment of Turner's Syndrome. Endocrinol Jpn. 1989[c];36(4):569-78.


