Growth Hormone Treatment of Girls With Turner Syndrome: The National Cooperative Growth Study Experience

Leslie Plotnick, MD*; Kenneth M. Attie, MD†; Sandra L. Blethen, MD, PhD‡; and Judy P. Sy, PhD‡

ABSTRACT. Objective. To evaluate growth rate and adult height with recombinant growth hormone (GH) treatment in girls with Turner syndrome (TS) and predictors of their growth response.

Methods. Data on girls with TS who were treated with GH in the National Cooperative Growth Study (NCGS) were evaluated. As of January 1997, there were 2798 girls with TS in the NCGS database, 2652 of whom had not previously received GH. Follow-up data on growth were available for 2475 subjects, and data on adult height were available for 622.

Results. The average age of girls with TS at enrollment in the NCGS was 10.1 ± 3.6 years. These patients had severely short stature compared with that of unaffected American girls (height, 118.5 ± 16.5 cm; height standard deviation score [SDS], −3.1 ± 0.9), but their heights were typical of those of American girls with TS (TS-specific height SDS, 0.01 ± 0.9). Treatment with GH for an average duration of 3.2 ± 2.0 years resulted in an increase in height SDS of 0.8 ± 0.7 compared with unaffected girls and of 1.2 ± 0.8 compared with TS standards. Growth rates increased from 4.0 ± 2.3 cm/year before treatment to 7.5 ± 2.0 cm/year after 1 year of treatment. Duration of treatment with GH was the strongest predictor of change in height SDS. After 6 to 7 years of treatment with GH, there was a cumulative change of 2.0 in mean height SDS.

Conclusions. Although the response to treatment with GH varied, it was associated with highly significant gains in growth and adult height in girls with TS. Duration of treatment with GH was the most important variable predicting adult height. Pediatrics 1998;102: 479–481; Turner syndrome, growth hormone, growth.

ABBREVIATIONS. TS, Turner syndrome; GH, growth hormone; NCGS, National Cooperative Growth Study; SDS, standard deviation score(s); NCHS, National Center for Health Statistics; CA, chronologic age; BA, bone age.

Turner syndrome (TS) is a chromosomal disorder that affects about 1 of every 2000 to 2500 live-born girls. More than 95% of girls with TS have short stature. The average difference between adult height of women with TS and that of unaffected women is 20 cm. Gonadal failure (>90%) and infertility (>99%) are other frequent findings. However, as many as 10% to 20% of girls with TS will develop breasts spontaneously, and a small minority (2% to 5%) will have spontaneous menses.

Biosynthetic growth hormone (GH) recently was approved in the United States for use in augmenting height in patients with TS. This study examines the outcomes from a large national database of girls with TS who were treated with GH.

METHODS

The National Cooperative Growth Study (NCGS) began its follow-up of patients treated with Genentech’s GH products in 1985. Methods of enrolling subjects and collecting data have been described previously. As of January 1997, there were 2798 girls with TS enrolled in the NCGS. Of these, 2652 were naïve to treatment with GH at enrollment. Follow-up data on growth were available for 2475 enrollees, and data on adult height were available for 622 enrollees. Data on height are presented as height standard deviation score(s) (SDS) relative to unaffected American girls (National Center for Health Statistics [NCHS]) and to girls with TS. The dosage of GH used is reported as the mean in milligrams per kilogram of body weight per week. Cumulative growth during GH treatment is reported as TS-specific height SDS.

Adult height was considered to have been attained if the chronologic age (CA) and bone age (BA) were older than 14 years at the time of the last reported measurement of height, or if the CA was at least 18 years. If the BA was not available when the height was last measured and the CA was between 14 and 18 years, an extrapolated BA was calculated by using a BA that had been recorded in the previous 3 years. If this BA was at least 14 years, the last height measured was considered the adult height.

The Lyon projection method, which assumes that the adult height SDS of girls with TS will equal their height SDS when first seen, was used to predict what the subjects’ heights would have been had they not been given GH. This method has been validated in the United States for patients with TS.

Frequency data are reported as percentages. Variables are presented as means ± SD.

RESULTS

Baseline data on girls with TS who had not been treated previously with GH are shown in Table 1. These girls were extremely short by NCHS standards (height SDS, −3.1 ± 0.9), but their TS-specific height SDS was 0.01 ± 0.9.

Table 2 shows data on treatment with GH in the 2475 girls for whom follow-up data on growth were available. The average duration of treatment was...
3.2 ± 2.0 years. During this time, treated subjects showed an increase in NCHS height SDS of 0.8 ± 0.7 and an increase in TS-specific height SDS of 1.2 ± 0.8.

On average, the growth rate increased from 4.0 ± 2.3 cm/year before treatment with GH to 7.5 ± 2.0 cm/year during the first year of treatment. As expected, there was some waning in the growth rate during subsequent years of treatment with GH. Figure 1 shows the cumulative change in TS-specific height SDS as a function of duration of treatment with GH.

A number of variables were evaluated to determine predictors of the cumulative change in TS-specific height SDS in patients treated with GH. The variables that had a significant positive association with the change in TS-specific height SDS were the duration of treatment (P < .0001), average number of injections of GH per week (P = .0005), onset of puberty (either spontaneously or after estrogen replacement) during treatment (P = .010), midparental height (P = .0005), and BA delay (n = 457; P = .008)

Variables with a significant negative (ie, inverse) association with the change in TS-specific height SDS were the percentage of injections of GH missed (P = .0005), baseline TS-specific height SDS (P = .0001), and baseline CA (P < .0001) (ie, girls who were younger when they began treatment with GH had greater increases in height SDS). Factors that were not significant predictors were the GH dosage, body mass index, maximum stimulated level of GH, and pretreatment growth rate. Duration of treatment was the strongest predictor.

Data on adult height were available for 622 subjects. Their data before, during, and after GH treatment are shown in Table 3. As a subgroup, these subjects were older (12.9 ± 2.5 years) than the rest of this TS group. Their height SDS (NCHS, −3.3 ± 0.9; TS, 0.3 ± 0.9) and midparental stature (−0.2 ± 0.9) were similar to those for the TS group as a whole. Before treatment with GH, their projected adult height was 141.9 ± 6.2 cm. In contrast, their last measured (hence, adult) height was 148.3 ± 5.6 cm. Thus, the average gain in adult height over pretreatment projected height in this subgroup was 6.4 ± 4.9 cm.

Figure 2 shows the frequency distribution of the gain in adult height in this subgroup. In 60% of these patients, adult height was at least 5 cm greater than the pretreatment projected adult height. Forty percent showed a gain in adult height of at least 7.5 cm.

**TABLE 2.** Data on Treatment With GH in 2475 NCGS Patients With TS

<table>
<thead>
<tr>
<th></th>
<th>Mean Value</th>
<th>SD</th>
</tr>
</thead>
<tbody>
<tr>
<td>Duration of treatment with GH (y)</td>
<td>3.2</td>
<td>2.0</td>
</tr>
<tr>
<td>Change in NCHS height SDS</td>
<td>0.8</td>
<td>0.7</td>
</tr>
<tr>
<td>Change in TS-specific height SDS</td>
<td>1.2</td>
<td>0.8</td>
</tr>
<tr>
<td>GH dose (mg/kg/wk)</td>
<td>0.33</td>
<td>0.06</td>
</tr>
<tr>
<td>Number of injections/wk</td>
<td>6.0</td>
<td>1.4</td>
</tr>
<tr>
<td>Growth rate (cm/y)</td>
<td>4.0</td>
<td>2.3</td>
</tr>
</tbody>
</table>

*At baseline, 88.9% prepubertal; 11.1% pubertal.*

**DISCUSSION**

In general, the girls with TS in the NCGS database are typical of girls with TS in the general population, as indicated by the fact that their mean TS-specific height SDS is nearly 0.0, with an SD of 0.9. Factors that can affect the apparent benefit of GH in TS are the patient’s age at the start of treatment, height deficit at the start of treatment, duration of the treat-

**TABLE 3.** Data on the 622 NCGS Patients With TS Who Attained Adult Height

<table>
<thead>
<tr>
<th></th>
<th>Mean Value</th>
<th>SD</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age at last recorded height (y)</td>
<td>17.1</td>
<td>1.6</td>
</tr>
<tr>
<td>Lyon projection of adult height (cm)</td>
<td>141.9</td>
<td>6.2</td>
</tr>
<tr>
<td>Last recorded height (cm)</td>
<td>148.3</td>
<td>5.6</td>
</tr>
<tr>
<td>Height gain over projected height (cm)</td>
<td>6.4</td>
<td>4.9</td>
</tr>
<tr>
<td>Last recorded NCHS height SDS</td>
<td>−2.4</td>
<td>0.9</td>
</tr>
<tr>
<td>Change in NCHS height SDS</td>
<td>1.0</td>
<td>0.8</td>
</tr>
<tr>
<td>Last recorded TS-specific height SDS</td>
<td>1.2</td>
<td>0.9</td>
</tr>
<tr>
<td>Change in TS-specific height SDS</td>
<td>1.0</td>
<td>0.8</td>
</tr>
<tr>
<td>Duration of treatment with GH (y)</td>
<td>3.7</td>
<td>1.9</td>
</tr>
<tr>
<td>GH dose (mg/kg/wk)</td>
<td>0.33</td>
<td>0.06</td>
</tr>
<tr>
<td>Number of injections of GH/wk</td>
<td>5.8</td>
<td>1.4</td>
</tr>
<tr>
<td>Growth rate (cm/y)</td>
<td>3.5</td>
<td>2.3</td>
</tr>
<tr>
<td>Pretreatment (n = 408)</td>
<td>6.7</td>
<td>2.2</td>
</tr>
<tr>
<td>Year 1 (n = 472)</td>
<td>5.4</td>
<td>1.9</td>
</tr>
<tr>
<td>Year 2 (n = 368)</td>
<td>4.4</td>
<td>1.8</td>
</tr>
<tr>
<td>Year 3 (n = 278)</td>
<td>4.0</td>
<td>1.8</td>
</tr>
<tr>
<td>Year 4 (n = 190)</td>
<td>4.0</td>
<td>1.8</td>
</tr>
</tbody>
</table>
Turner Syndrome and Osteoporosis: Mechanisms and Prognosis

Karen Rubin, MD

ABSTRACT. Despite only limited reports of a greater number of fractures during childhood or adulthood, osteoporosis historically has been described as a feature in Turner syndrome, because of the frequent observation of radiographic osteopenia and the coarse trabecular pattern of the carpal bones on radiographs. The pathogenesis of the skeletal demineralization remains unclear, but the data support the concept of an intrinsic bone defect that is then exacerbated by a number of hormonal factors, including the growth-regulating hormones, the gonadal steroids, and possibly the calcium-regulating hormones. The advent of more refined methods, such as single- and

REFERENCES

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