



ORIGINAL ARTICLE

Human growth hormone and Turner syndrome[☆]



Silvia Beatriz Sánchez Marco^{a,*}, Antonio de Arriba Muñoz^a, Marta Ferrer Lozano^a, José Ignacio Labarta Aizpún^a, Jesús María Garagorri Otero^b

^a Unidad de Endocrinología Pediátrica, Hospital Universitario Miguel Servet, Zaragoza, Spain

^b Servicio de Pediatría, Hospital Clínico Universitario Lozano Blesa, Zaragoza, Spain

Received 28 November 2015; accepted 24 February 2016

Available online 16 January 2017

KEYWORDS

Turner syndrome;
Short stature;
Growth hormone

Abstract

Objective: The evaluation of clinical and analytical parameters as predictors of the final growth response in Turner syndrome patients treated with growth hormone.

Material and methods: A retrospective study was performed on 25 girls with Turner syndrome (17 treated with growth hormone), followed-up until adult height. Auxological, analytical, genetic and pharmacological parameters were collected. A descriptive and analytical study was conducted to evaluate short (12 months) and long term response to treatment with growth hormone.

Results: A favourable treatment response was shown during the first year of treatment in terms of height velocity gain in 66.6% of cases (height-gain velocity >3 cm/year). A favourable long-term treatment response was also observed in terms of adult height, which increased by 42.82 ± 21.23 cm (1.25 ± 0.76 SDS), with an adult height gain of 9.59 ± 5.39 cm (1.68 ± 1.51 SDS).

Conclusions: Predictors of good response to growth hormone treatment are: (A) initial growth hormone dose, (B) time on growth hormone treatment until starting oestrogen therapy, (C) increased IGF1 and IGFBP-3 levels in the first year of treatment, and (D) height gain velocity in the first year of treatment.

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[☆] Please cite this article as: Sánchez Marco SB, de Arriba Muñoz A, Ferrer Lozano M, Labarta Aizpún JI, Garagorri Otero JM. Hormona de crecimiento y síndrome de Turner. An Pediatr (Barc). 2017;86:81–86.

* Corresponding author.

E-mail address: ssanchezmarco@gmail.com (S.B. Sánchez Marco).

PALABRAS CLAVE

Síndrome de Turner;
Talla baja;
Hormona de
crecimiento

Hormona de crecimiento y síndrome de Turner**Resumen**

Objetivo: Evaluación de parámetros clínicos y analíticos que actúen como predictivos de respuesta al tratamiento con hormona de crecimiento (rhGH) a largo plazo en pacientes con síndrome de Turner.

Material y métodos: Estudio retrospectivo de 25 niñas diagnosticadas de síndrome de Turner, de las cuales 17 recibieron tratamiento con rhGH y fueron controladas hasta alcanzar la talla adulta. Se determinaron diferentes variables auxológicas, analíticas, genéticas y farmacológicas a lo largo de su seguimiento en dichas consultas. Se realizó un estudio descriptivo y analítico mediante regresión lineal, con valoración de la respuesta al tratamiento a corto (12 meses) y a largo plazo.

Resultados: Se observó una respuesta favorable a corto plazo valorada en ganancia de velocidad de crecimiento en el 66,6% de los casos (velocidad de crecimiento > 3 cm/año a los 12 meses de tratamiento respecto a la previa). También se evidenció una respuesta favorable a largo plazo, valorada en una ganancia de talla total de $42,82 \pm 21,23$ cm ($1,25 \pm 0,76$ SDS). Las pacientes ganaron una media de $9,59 \pm 5,39$ cm ($1,68 \pm 1,51$ SDS) respecto a su pronóstico de crecimiento previo al tratamiento.

Conclusiones: El presente estudio evidencia como factores predictivos de buena respuesta al tratamiento con rhGH a largo plazo en orden de importancia: A) dosis de rhGH al inicio del tratamiento, B) tiempo de tratamiento con rhGH hasta inicio de terapia estrogénica, C) incremento en los niveles de IGF1 e IGFBP-3 durante el primer año de tratamiento y D) velocidad de crecimiento en el primer año de tratamiento.

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Introduction

Turner syndrome is a chromosomal disorder that occurs in one of every 2500 live births, characterised by the complete or partial absence of a chromosome X (the complete monosomy [45,X] accounts for more than 50% of cases),^{1,2} although some studies have found evidence of a high prevalence of mosaicism.³ It is associated with a range of phenotypic characteristics, chief of which are short stature, gonadal dysgenesis, hand and foot lymphoedema, pterygium colli, cubitus valgus and cardiovascular malformations, among others.²

Since short stature is a common feature and the sole clinical manifestation in most cases,¹⁻³ several studies have evinced the efficacy of treatment with recombinant human growth hormone (rhGH), with increases in final height of ~7–10 cm.⁴⁻⁶

Treatment with rhGH has to be initiated early when stature is more than two standard deviations (–2 SDS) below that of the general population or height velocity (HV) is below the tenth percentile for the patient's bone age. It should not be delayed past age 4 years nor initiated before age 2 years.^{1,4}

Different studies have identified some predictors of adult height, such as height at initiation of treatment with rhGH, response in the first year of treatment, genetic height potential, age at initiation of treatment or mean weekly dose of rhGH.^{4,6,7}

We present a study conducted in a Spanish cohort of patients with Turner syndrome with the aim of analysing the

association between the response to treatment with rhGH and various factors.

Materials and methods

We conducted a retrospective study of 25 patients with a Turner syndrome diagnosis, 17 of who were treated with rhGH and followed up at the paediatric endocrinology unit of a tertiary level hospital until they reached their adult height. The patients treated with rhGH have been in follow-up from 1995 to present, and having reached adult height was an inclusion criterion. We also retrieved data for older cases that were not treated with rhGH on account of being diagnosed at older ages or the family refusing the treatment.

We reviewed the medical records of patients with Turner syndrome, collecting data for auxological measurements, laboratory tests, karyotyping and pharmacological treatment throughout their followup in the unit. We informed the patients of the purpose of the study and obtained their informed consent.

We assessed the short-term response to treatment with rhGH (12 months) based on changes in HV. We defined response to treatment as an increase in HV of more than three centimetres per year compared to the previous HV or an increase by three SDS at 12 months of treatment. To assess the long-term response to treatment (up to reaching adult height), we used five possible response variables: (1) an increase in height SDS compared to baseline SDS (delta HtSDS: adult height SDS– height at initiation of rhGH)³; (2)

increase in height SDS compared to the baseline predicted adult height SDS (adult height SDS– 10th percentile SDS); (3) increase in height SDS compared to the height at initiation of oestrogen therapy as a response variable (adult height SDS– height at oestrogen initiation); (4) increase in height SDS compared to the difference of the height at initiation of treatment with rhGH and the height at initiation of oestrogen therapy (oestrogen initiation height SDS– rhGH initiation height SDS); (5) increase in height SDS relative to the duration of rhGH therapy prior to initiation of oestrogen therapy.

We conducted a descriptive and inferential analysis with SPSS 18.0 for Windows using nonparametric tests: the Wilcoxon test for quantitative variables, Spearman's correlation test for analysing the linear correlation between quantitative variables, linear regression analysis for quantitative variables that had shown a linear correlation and for quantitative and qualitative variables; the Mann–Whitney *U* test to compare dichotomous qualitative variables with quantitative variables, and the Kruskal–Wallis test to compare nominal qualitative variables with quantitative variables. We defined statistical significance for all tests as a *p*-value of less than .05.

Results

Of the 25 patients under study, 17 (68%) received rhGH treatment, while eight did not.

The most frequent karyotype in these patients was 45,X (42.9%) followed by isochromosome 46,X,i(Xq) (17.8%) and 46,X,i(Xq)/45,X mosaicism (14.3%). Other less frequent karyotypes found in the sample were mosaicisms such as 46,XX(r)/45,X; 47,XXX/45,X/46,XX; 45,X/46,X,der(X) and 45,X/47,XXX; 46,X,i/45,X/46,XX.

When it came to phenotypic expression, we found that the classical presentation (short stature, pterygium colli, cubitus valgus, widely spaced nipples, etc.) was the most frequent phenotype (50%); while six patients did not have characteristic features (27.3%) and received the diagnosis when they underwent evaluation of short stature.

The mean age at initiation of rhGH therapy was 7.90 ± 4.13 years (range, 1.82–16.45 years) with a mean height SDS of -2.49 ± 0.63 .

The mean initial dose of rhGH used for treatment was 0.048 ± 0.01 mg/kg/day and the mean duration of treatment was 7.36 ± 3.88 years. At the time of treatment completion, patients had a mean age of 14.52 ± 1.69 years and had reached a mean final adult height of 156.15 ± 3.66 cm (-1.23 ± 0.62 SDS).

At 12 months of treatment with rhGH, there was an increased HV of 8.21 ± 2.22 cm/year (2.5 ± 2.55 SDS) (Fig. 1). At this time, the predicted adult height for the patients had increased by 4.39 ± 3.14 cm (144.98 ± 7.99 vs 149.37 ± 5.92 cm) and 0.67 ± 1.06 SDS (-2.92 ± 1.21 vs -2.24 ± 0.91 SDS) (Fig. 2). At two years of treatment, the mean predicted adult height increased to 150.56 ± 5.17 cm, with a slight decrease in HV to 6.73 ± 1.63 cm/year.

As for laboratory parameters, there was a progressive increase in the levels of IGF1 and IGFBP3 until the end of treatment, with levels reaching up to three times their

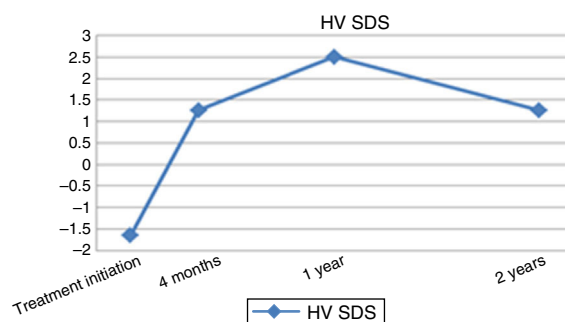


Figure 1 Longitudinal changes in height velocity (SDS).

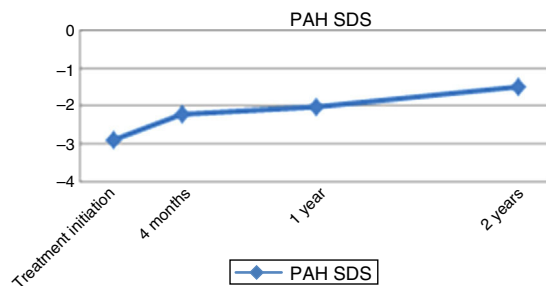


Figure 2 Longitudinal changes in predicted adult height (SDS).

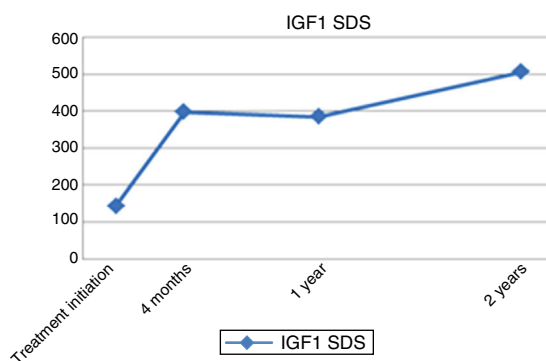


Figure 3 Longitudinal changes in mean IGF1 levels (ng/mL).

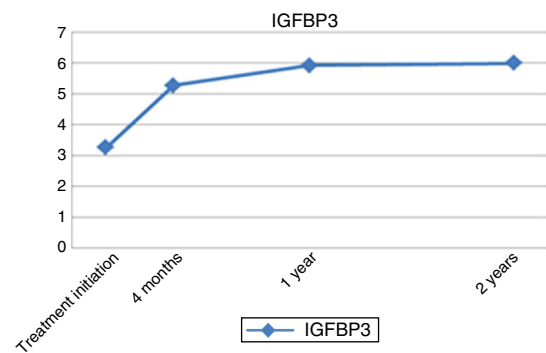


Figure 4 Longitudinal changes in mean IGFBP3 levels (μ g/mL).

baseline values, but always remaining within normal ranges (Figs. 3 and 4).

The mean final height gain compared to baseline height at initiation of treatment was 42.82 ± 21.23 cm (total gain

Table 1 Descriptive analysis of the long-term response to treatment with rhGH.

	Initiation of treatment	2 years of treatment	Adult age	Total gain
Height (cm)	113.32 ± 22.98	124.18 ± 17.74	156.15 ± 3.66	42.82 ± 21.23
Height SDS	-2.49 ± 0.63	-1.93 ± 0.85	-1.23 ± 0.62	1.25 ± 0.76
Predicted adult height (cm)	144.9 ± 7.89	150.57 ± 5.17		9.59 ± 5.39
Predicted adult height SDS	-2.92 ± 1.21	-2.05 ± 0.81		1.68 ± 1.51

Table 2 Analysis of the changes in studied parameters from baseline to one year after initiation of rhGH therapy.

	At treatment initiation, mean ± SDS	At one year of treatment, mean ± SDS	P
Height SDS	-2.49 ± 0.63	-2.09 ± 0.80	.028
Height velocity SDS	-1.64 ± 2.58	2.50 ± 2.55	.004
Predicted adult height SDS	-2.92 ± 1.21	-2.24 ± 0.90	.075
IGF1	143.12 ± 101.02	384.83 ± 111.47	.043
IGFBP3	3.25 ± 1.36	5.77 ± 1.47	.068

Table 3 Correlation between height gain parameters and duration of treatment with rhGH before initiation of oestrogen therapy.

	P	R ²	R	β coefficient
Increase in height velocity at 1 year (cm)	.094	0.75	0.86	0.134
Increase in height velocity SDS at 1 year	.014	0.813	0.91	0.189
Predicted adult height at rhGH initiation	.000	0.546	0.74	0.628
Height gain SDS at 4 months	.050	0.372	0.61	0.685
rhGH dose	.037	0.795	0.891	69.52

of 1.25 ± 0.76 SDS). When it came to the baseline predicted adult height, height gain in patients exceeded it by a mean of 9.59 ± 5.39 cm (1.68 ± 1.51 SDS) (Table 1).

After 12 months of treatment, patients had improvements in height ($P = .011$), HV ($P = .004$) and IGF1 levels ($P = .043$) (Table 2), and we also observed an improvement in the adult heights achieved ($P = .028$).

Our assessment of the short-term response to treatment based on increases in HV showed that in 66.6% of the patients, HV increased by more than 3 cm/year at 12 months of treatment compared to baseline. Furthermore, we found a positive correlation between response to treatment and the dose of rhGH ($P = .037$; $\beta = 69.95$), and response to treatment and increased levels of IGF1 and IGFBP3 ($P = .000$; r^2 [IGF1] = 0.846, r^2 [IGFBP3] = 0.809). When we analysed height gain relative to the interval between initiation of treatment with rhGH and initiation of oestrogen therapy, we found a positive correlation for auxological parameters such as height and HV both in cm and SDS at four months and one year of treatment (Table 3).

The mean difference in final height between patients treated with rhGH and patients not treated with it was 10.69 ± 2.63 cm ($P = .031$; 156.15 ± 3.66 cm vs 145.44 ± 3.69 cm, respectively).

Furthermore, we computed linear regression curves using the predictors for a favourable response identified in previous steps, but since the sample size was small, the results we obtained do not seem to be generalisable.

We did not find any differences in response to treatment based on karyotype ($P = .147$).

Discussion

Short stature is the key feature of Turner syndrome and may occur in the absence of other clinical manifestations.³ Linear growth deceleration starts in infancy and early childhood, and becomes more marked in late childhood and adolescence, resulting in a significantly short stature in adulthood.² The indication of rhGH therapy to increase HV and adult height in patients with Turner syndrome is accepted and supported by various studies.^{1,2,4} There is evidence that this treatment results in increases of five to ten centimetres in the final adult height.^{4,5} Treatment should be initiated when the patient's height is two SDS below the population mean or when HV is below the tenth percentile for the patient's bone age.^{4,8}

Several studies such as the one carried out by the Canadian Growth Hormone Advisory Committee^{9,10} and a few conducted in Spain have demonstrated the importance of rhGH treatment in these patients, as they have found increases in HV of 4.1–8.15 cm at 12 months of treatment.^{11–13} There is great variability in the increases in HV reported in the literature, as they range from -1.69 to 1.97 SDS. The increase is greatest in the first year of treatment, and less significant in subsequent year.¹⁴ In our study,

at 12 months of treatment we found an increased HV of 8.21 ± 2.22 cm/year (2.5 ± 2.55 SDS) with a difference compared to baseline of 3.22 cm, which corresponds to a good initial increase in HV compared to what has been reported in previous studies.^{11–14}

In addition, research in recent years has attempted to both assess response to treatment and determine the factors that promote a good response. Thus, for the purpose of assessing response to treatment in the short term, Quigley et al.¹⁵ noted the increase in HV by ~ 2 cm a year. In our study, we assessed response to treatment with rhGH applying a threshold of 3 cm a year for the increase in HV at 12 months of treatment, which was exceeded by 66.6% of the patients under study. Other authors¹⁶ highlight a mean increase in height SDS in the first year of treatment of $+0.50 \pm 0.03$ SDS. In the short term, these parameters function as predictors of a long-term favourable response to treatment, although this remains controversial.¹⁷ In the study that we present here, we observed that when it came to the longitudinal changes in the main auxological parameters, the largest changes in height, HV and predicted adult height took place in the first year (4–12 months from initiation), followed by lesser increases and an eventual plateau, except in HV.

As to the findings concerning long-term response to treatment, Radetti et al. found increases in final height of up to 9.2 cm, corroborated by more recent studies showing increases in final height of ~ 7 cm¹⁸ and of more than 1 SDS.^{19,20} Our findings were similar, with a mean increase in final height compared to baseline height of 9.59 ± 5.39 cm (1.68 ± 1.51 SDS).

There is disagreement regarding the association between the dosage of rhGH and the improvement in clinical outcomes due to increased levels of growth factors; some authors have found such an association,^{16,20–22} while others, including García et al.²³ or Wetterau et al.,²⁴ believe that it has no impact on the final response to treatment. In our study, we found a positive correlation with the increase of growth factor levels (IGF1 and IGFBP3) at one year of treatment, which is strongly associated with the administered dose of rhGH and better clinical outcomes.^{14,25} Furthermore, studies conducted in recent years have attempted to identify the genes involved in the early response of these patients to rhGH. Some have identified polymorphisms in the *CDK4* gene that may have an impact on IGF1 levels and therefore on the response to treatment with rhGH.²⁶ Ranke et al.^{4,25,27} found positive correlations with the administered dose (greater than 0.27 mg/kg/day), frequency (more days per week) and age at initiation (some authors assert that adult height can reach the normal range if treatment is initiated between 2 and 4 years of age^{6,19}), with no consensus on the subject.²⁸

The duration of treatment with rhGH also seems to influence the final height gain. Chernauek et al.^{19,22,27} described this time interval as an important predictor of final height, and even produced an equation to estimate the approximate height gain: height gain (in cm) = $2.1 \times$ years of rhGH treatment prior to oestrogen initiation.

The timing of initiation of oestrogen therapy continues to be debated; some authors^{27–29} have found greater height gains with initiation of oestrogen therapy at age 14 years; other studies suggest that low doses of oestrogen

administered at earlier ages may lead to an increase of 0.37 SDS in adult height compared to patients in whom initiation of oestrogen therapy is delayed.²⁹ In our opinion, the timing of oestrogen initiation should be decided in agreement with the patient and the family, taking into account not only auxological parameters such as height or bone age, but also bone mineral density and the wishes expressed by the patient in relation to treatment initiation and pubertal development.

Another controversial aspect is the use of anabolic steroids such as oxandrolone, as it seems to increase final height when combined with rhGH. Gault et al.²⁸ found a mean increase in final height of +4.6 cm (range, 1.9–7.2 cm) in a group treated with oxandrolone compared to a group treated with placebo. However, their study did not find a statistically significant positive additive effect for the combination of oxandrolone and late pubertal induction.

Our comparison of patients treated with rhGH and patients not treated with rhGH was consistent with the findings of Paschino et al.³⁰ and Rosenfeld et al.,³¹ with a mean difference in height between both groups of 10.69 ± 2.63 cm ($P = .031$), which evinces the importance of early treatment in these patients. Ranke et al.¹⁸ studied the possibility of patients with specific karyotypes responding better to treatment with rhGH, but we found no differences based on karyotype.

Based on the results of our study, we determined that the predictors for a long-term favourable response are, from most important to least: (A) dose of rhGH at treatment initiation, (B) duration of treatment with rhGH prior to initiation of oestrogen therapy, (C) increased levels of IGF1 and IGFBP3 during the first year of treatment and (D) HV in the first year of treatment.

Conflict of interests

The authors have no conflict of interests to declare.

References

1. Kliegman RM, Stanton BF, St. Geme JW, Schor NF, Behrman RE. *Nelson tratado de pediatría*. 19.ª edición Barcelona: Elsevier; 2012. p. 2024–7.
2. Barreda Bonis AC, Gonzalez Casado I, Gracia Bouthelie R. Síndrome de Turner. *Protoc Diagn ter Pediatr*. 2011;1:218–27.
3. Ríos Orbañanos I, Vela Desojo A, Martínez-Indart L, Grau Bolado G, Rodríguez Estevez A, Rica Echevarría I. Turner syndrome: from birth to adulthood. *Endocrinol Nutr*. 2015;62:499–506.
4. Ranke MB, Lindberg A, KIGS International Board. Observed and predicted growth responses in prepubertal children with growth disorders: guidance of growth hormone treatment by empirical variables. *J Clin Endocrinol Metab*. 2010;95:1229–37.
5. Ross JL, Lee PA, Gut R, Germak J. Increased height standard deviation scores in response to growth hormone therapy to near-adult height in older children with delayed skeletal maturation: results from the ANSWER Program. *Int J Pediatr Endocrinol* [Internet]. 2015;2015:1 [Epub Jan 2015]. Available from: <http://www.ncbi.nlm.nih.gov/pubmed/25904938> [cited 05.01.16].
6. Locher S, Carta L, Ibba A, Guzzetti C. Growth hormone treatment in non-growth hormone-deficient children. *Ann Pediatr Endocrinol Metab*. 2014;19:1–7.

7. Savage MO, Bang P. The variability of responses to growth hormone therapy in children with short stature. *Indian J Endocrinol Metab.* 2012;16:178–84.
8. López-Siguero JP, Jiménez del Valle M. Indicaciones actuales de la hormona de crecimiento en pediatría. *JANO.* 2008;1704:40–1.
9. Rosenfeld RG. Acceleration of growth in Turner syndrome patients treated with growth hormone: summary of three-year results. *J Endocrinol Invest.* 1989;12:49–51.
10. The Canadian Growth Hormone Advisory Committee. Impact of growth hormone supplementation on adult height in Turner syndrome: results of the Canadian randomised controlled trial. *J Clin Endocrinol Metab.* 2005;90:3360–6.
11. Ferrández-Longás A, Mayayo E, Labarta JI, Bagué L, Puga B, Rueda C, et al. Estudio longitudinal de crecimiento y desarrollo. Centro Andrea Prader. Zaragoza 1980–2002. Patrones de crecimiento y desarrollo en España. Atlas de gráficas y tablas. Madrid: Ergon; 2004. p. 61–115.
12. Fernández RM. Patrón de crecimiento humano y su evaluación. In: Pombo M, editor. *Tratado de endocrinología pediátrica.* 3.ª ed. Madrid: McGraw-Hill Interamericana; 2002. p. 244–74.
13. Ferrández A, Labarta JI, Calvo M, Mayayo E, Puga B, Cáncer E, et al. Síndrome de Turner. In: Pombo M, editor. *Tratado de endocrinología pediátrica.* 3.ª ed. Madrid: McGraw-Hill Interamericana; 2002. p. 780–803.
14. Tai S, Tanaka T, Hasegawa T, Ozono K, Tanaka H, Kanzaki S, et al. An observational study of the effectiveness and safety of growth hormone (Humatrope®) treatment in Japanese children with growth hormone deficiency or Turner syndrome. *Endocr J.* 2013;60:57–64.
15. Quigley CA, Crowe BJ, Anglin G, Chipman JJ. Growth hormone and low dose estrogen in Turner syndrome: results of a United States multi-center trial to near-final height. *J Clin Endocrinol Metab.* 2002;87:2033–41.
16. Lee PA, Ross JL, Pedersen BT, Kotnik P, Germak JA, Christesen HT. Noonan syndrome and Turner syndrome patients respond similarly to 4 years' growth-hormone therapy: longitudinal analysis of growth-hormone-naïve patients enrolled in the NordiNet® International Outcome Study and the ANSWER Program. *Int J Pediatr Endocrinol.* 2015;2015:5–2017.
17. Wasniewska M, Aversa T, Mazzanti L, Guarneri MP, Matarazzo P, De Luca F, et al. Adult height in girls with Turner syndrome treated from before 6 years of age with a fixed per kilogram GH dose. *Eur J Endocrinol.* 2013;169:439–43.
18. Ranke MB, Lindberg A, Ferrández Longás A, Darendeliler F. Major determinants of height development in Turner syndrome (TS) patients treated with GH: analysis of 987 patients from KIGS. *Pediatr Res.* 2007;61:105–10.
19. Linglart A, Cabrol S, Berlier P, Stuckens C, Wagner K, de Kerdanet M, et al. Growth hormone treatment before the age of 4 years prevents short stature in young girls with Turner syndrome. *J Clin Endocrinol Metab.* 2010;95:1229–37.
20. Van Pareden YK, de Muinck Keizer-Schrama SM, Stijnen T, Sas TC, Jansen M, Otten BJ, et al. Final height in girls with Turner syndrome after long-term growth hormone treatment in 3 dosages and low dose estrogens. *J Clin Endocrinol Metab.* 2003;88:1119–25.
21. Tillmann V, Patel L, Gill MS, Whatmore AJ, Price DA, Kibirige MS, et al. Monitoring serum insulin-like growth factor-I (IGF-I), IGF binding protein-3 (IGFBP-3), IGF-I/IGFBP-3 molar ratio and leptin during growth hormone treatment for disordered growth. *Clin Endocrinol.* 2000;53:329–36.
22. Cohen P, Rogol AD, Howard C, Bright G, Kappelgaard AM, Rosenfeld R, American Norditropin Study Group. Insulin growth factor-based dosing of growth hormone therapy in children: a randomized, controlled study. *J Clin Endocrinol Metab.* 2007;92:2480–6.
23. García García E. Evidencias en el tratamiento con hormona del crecimiento. Nuevas indicaciones. AEPaped. *Curso de Actualización Pediatría 2010.* Madrid: Exlibris Ediciones; 2010. p. 55–64.
24. Wetterau L, Cohen P. Role of insulin-like growth factor monitoring in optimizing growth hormone therapy. *J Pediatr Endocrinol Metab.* 2000;13:1371–6.
25. Schrier L, de Kam ML, McKinnon R, Che Bakri A, Oostdijk W, Sas TC, et al. Comparison of body surface area versus weight-based growth hormone dosing for girls with Turner syndrome. *Horm Res Paediatr.* 2014;81:319–30.
26. Stevens A, Clayton P, Tatò L, Yoo HW, Rodriguez-Arno MD, Skrodok J, et al. Pharmacogenomics of insulin-like growth factor-I generation during GH treatment in children with GH deficiency or Turner syndrome. *Pharmacogenomics J.* 2014;14:54–62.
27. Chernašek SD, Attie KM, Cara JF, Rosenfeld RG, Frane J. Growth hormone therapy of Turner syndrome: the impact of age of estrogen replacement on final height 1. *J Clin Endocrinol Metab.* 2000;85:2439–45.
28. Gault EJ, Perry RJ, Cole TJ, Casey S, Paterson WF, Hindmarsh PC, et al. Effect of oxandrolone and timing of pubertal induction on final height in Turner's syndrome: randomised, double blind, placebo controlled trial. *BMJ.* 2011;342:d1980.
29. Ross JL, Quigley CA, Cao D, Feuillan P, Kowal K, Chipman JJ, et al. Growth hormone plus childhood low-dose estrogen in Turner's syndrome. *N Engl J Med.* 2011;364:1230–42.
30. Pasquino AM, Pucarelli I, Segni M, Tarani L, Calcaterra V, Larizza D. Adult height in sixty girls with Turner syndrome treated with growth hormone matched with an untreated group. *J Endocrinol Invest.* 2005;28:350–6.
31. Rosenfeld RG, Attie KM, Frane J, Brasel JA, Burstein S, Cara JF, et al. Growth hormone therapy of Turner's syndrome: beneficial effect on adult height. *J Pediatr.* 1998;132:319–24.