

# Linear growth and response to GH therapy in children with GHD with normal IGF-I versus those with normal GH secretion associated with low IGFI at presentation.

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## Introduction

Children with idiopathic short stature (ISS) have linear growth impairment despite normal or even high levels of GH (NGH). In some of these children IGFI level is low (NGH + Low IGFI (IGFSDS<-1.5).

It was observed that some children with GHD (Peak GH < 7 ng/dl after provocation) have normal IGFI level (GHD +Normal IGFI).

The linear growth of these two groups at presentation and their response to GH therapy was not clearly studied before.

## Aim

We studied the linear growth at presentation and response to a year of GH therapy in children with low IGFI and NGH secretion compared to those with GHD with normal IGF-I.

## Methods

We conducted a longitudinal study on 78 children presented to pediatric clinic with short stature (January - December 2017).

Children were classified to ISS and GHD based on growth hormone provocation test.

Anthropometrics data (HtSDS, difference from mid-parental height SDS (MPHSD), BMISDS, and weight gain/day (WG) , bone age and insulin-like growth factor 1 (IGF-1) SDS level were studied in all children for 1 year.

We compared the linear growth at presentation and response to a year of GH therapy in children with low IGFI and normal GH secretion (group 1, N= 10) to those with GHD with normal IGF-I (group 2, N = 17).

## Results

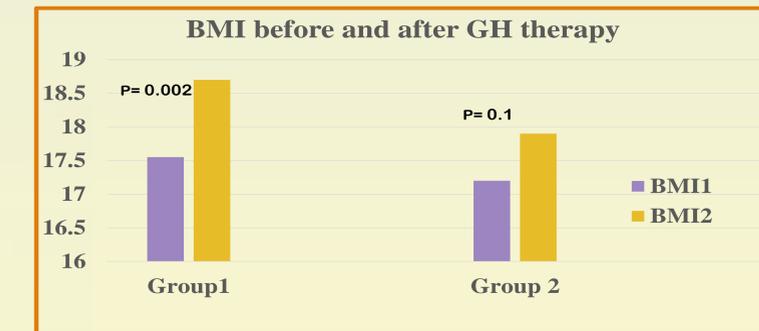
At presentation, the age, HtSDS and BMI did not differ significantly among the 2 groups. The difference between the HtSDS of patients from their MPHSD was bigger in the IGFI deficient group vs GHD group (p =0.04).

		Age (yr.)	HtSDS1	BMISDS1	MPHSDS	HSD1-MPHSD
Group1	Mean	11.18	-2.47	17.55	-0.82**	-1.76
	SD	2.86	0.46	3.7	0.73	0.37
Group 2	Mean	9.64	-2.23	17.2	-1.25*	-0.87 *
	SD	3.67	0.59	3.79	1.16	1.2

After an average of 1 year of GH therapy (0.03 -0.05 mg/kg/day), the HtSDS increased significantly in both groups (P<0.01).

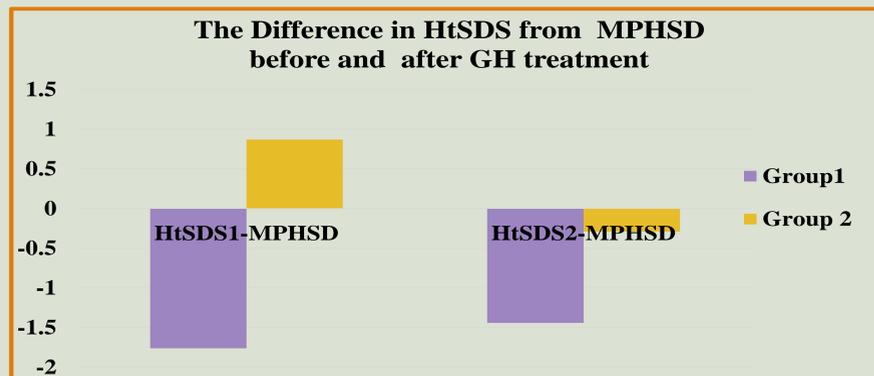
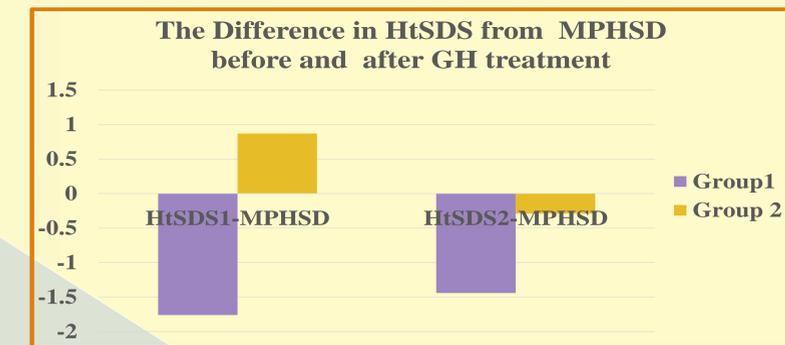
No significant difference in weight gain per day was detected between the two groups.

BMI in group 1 improved after GH therapy (P= 0.002) whereas it didn't differ in group 2.



After a year of GH therapy, there was a significant increase in the IGFSDS in both groups (p<0.05), and the increments of IGFSDS did not differ among them.

The difference in HtSD from their MPHSD after treatment improved significantly in group 2(P=0.04) but did not differ in group1 (P=0.4).



## Growth data after year of Growth hormone therapy

		HtSDS2	BMI 2	HtSDS2 - MPHSD	Delta HtSDS	Delta BMISDS	Delta IGF1SD	Wt. gain g/day
Group 1	Mean	-2.09	18.7	-1.44	0.38**	1.16*	1.23**	16.89
	SD	0.4	4.36	0.38	0.38	0.9	1.38	8.3
Group 2	Mean	-1.87	17.9	-0.29	0.36**	0.29	1.36**	11.53
	SD	0.7	4	1.22	0.44	0.89	0.8	6.16

\*\* Significant P <0.05 within same group  
\* Significant P <0.05 between two group

## Conclusion:

Our data suggests a comparably good growth response to GH therapy and increment in IGF-1SDS in children with GHD with normal IGF-1 and those with NGH with IGF-1 deficiency over the first year of treatment.