# Study of Physical Growth Pattern in Thalassemic Children And Adolescent in Hawler Thlassemia Center /Erbil City Nazar Ramadhan Othman \* \*

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الخلاصة

**خلفية البحث :** ثلاسيميا من أكثر الأمراض الوراثية شيوعا في العالم ومن المضاعفات التي تلاحظ المصابين هو ضعف النمو البدني لدى ألاطفال المصابين بالثلاسيميا مر مسبب السمو البداني على المصابيل بالمصابيل بالمرسيمي الهدف : تهدف الدر اسة للتعرف على نمط النمو البدني( الجسمي) لدى الأطفال المصابين بالثلاسيميا ومقارنتها بالأصحاء من طلبة المدارس الثانوية لمدينة اربيل المنهجية : أجربت در اسة في مركز هو لير للثلاسيميا للفترة من 30-2-2012 الى 2012-9-2012 اختبرت عينة من خمسون طفل مصاب بالثلاسيميا ومائتان طالب سليم معافى من طلاب المدارس الثانوية لمدينة اربيل تقنبات الاستبيان البيانات تصميم استمارة استبيان تتضمن جزئين للصمن جريين البيانات تحليل البيانات طريق SPSS البيانات .17 كوسيلة ألنتائج :اظهرت النتائج بوجود فرق في أنماط النمو البدني ( الجسمي) لدى الأطفال المصابين بالثلاسيميا ومقار نتها بالأطفال الأصحاء من طلبة المدارس الاستنتاجات : بينت نتائج الدراسة فيما يخص العمر وعلاقته بالطول بوجد وسط حسابي عالى لدى طلبة المدارس عند الوقوف والجلوس وكذلك عند الوزن والكتلة العضلية لجميع الفئات العمرية عند مقارنتها الأطفال المصابين بالثلاسيميا التوصيات: . اوصَت الدراسة بان مرض الثلاسيميا احدى المشاكل الصحية لدى الاطفال فان التخطيط الصحي لغرض تقليل نسبة حالات الاصابة وبرامج التحري عن المرض اضافة الى التوعية الصحية للأفراد المجتمع

Abstract

Background : Thalassaemia is one of the most common genetic blood disorders in the world. Growth impairment is a common observable complication and usually found a growth retardation in those patients

**Objectives:** The study aimed to identify the Physical growth pattern of Thalassemic( case group) with healthy school( control group) children and adolescents in Erbil city.

Methods: A study was carried out at Hawler Thalassemia Center in Erbil City during the period from the 1-2-2012- to 30-9-2012 . Fifty Thalassemic children( case group) from Hawler Thalassemia Center in Erbil City and two hundred healthy school( control group) children and adolescents were selected. A questionnaire was designed that composed of two parts to deal with physical growth pattern the data were collected by searchers to interviewed the participants in Hawler Thalassemic Center and secondary schools, for children and adolescents in Erbil city and the data were the analysis by using the (SPSS, 17),

**Results:** The results found that the range of age in the study sample was between 8-11 years old , which present in healthy school children and adolescents (control group) ( 46.5%), while in Thalassemic children( study group ) was ( 52.0%), the highest percentage of gender in both groups were male (65.5%, 58.0%) respectively revealed that there was significant differences incomparative physical growth pattern between Thalassemic and healthy children and adolescents

**Conclusion :** The study conclude that Thalassemic children (Case group) regarding the age they have high mean score than control group (healthy child) in (sitting and standing ) height, weight and nutritional status as BMI in all age groups.

**Recommendation**: The study recommended that Thalassemia was a important health problem in pediatric and plans need to set up to reduce the number of cases born by development of population screening programmer and awareness creating.

Key words: Thalassaemia, growth retardation, height, weight and growth pattern.

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#### **Introduction:**

Thalassemia is one of the most common human genetic diseases in the world and it cause many problems for patients, families and health care system <sup>(1)</sup> and it can't be prevented because they're inherited can passed on from parents to their children. Resulting anemia is usually severe with several health problems like enlarged spleen, bone deformities, fatigue and requires regular life-long transfusion, therapy and medical supervision.Iraq is one of the countries in which 6-10% of the populations have hemoglobinopathy of which thalassaemia is a major part<sup>(2)</sup>There are over 2,000 cases of thalassemia in the Kurdistan Region and Kirkuk.. In Kurdistan region a round 30000 people are carries of  $\beta$ -Thalassaemia. HawlerThalassaemia center serves about 493 regular registered patients, a daily attendance to follow up and to have blood transfusion of about 20-25 patients per day <sup>(3)</sup>. Thalassemia is widely distribution in Iraqi with high prevalence rate .growth retardation is one of complication in thalassemia. There are no studies available on the growth patterns in region . modern medical therapy allow the thalassemic children in Kurdistan Thalassemic children to grow normally in first decade of life but growth retardation was continues to be observed in significant proportion in those patients 8888

Growth retardation in thalassemic patients can occur as early as the first or second year of life but these abnormalities are more apparent after 6 to 8 years<sup>(4)</sup>. Regular blood transfusion followed by iron chelation therapy is just a supportive treatment for those patients, which is associated with serious complications. The beneficial effects of regular transfusions on growth and bone disorders in patients with thalassemia were first reported in 1965 <sup>(5)</sup>. Thalassemia is a chronic disease that presents a range of serious clinical challenges. The effects of thalassemia on physical health can lead to physical deformity, growth retardation, and delayed puberty<sup>(6)</sup> in Indian study reported that thalassemic patients are short, have low rate of growth and BMI which is related to low hemoglobin and high ferritin levels and sub-optimal iron chelation therapy. In developing countries, poor socio- economic background compounds the problem <sup>(7)</sup> .Additionally iron-deficiency anemia has a negative effect on physical growth. There was no studies of physical growth pattern of Thalassemic children in Erbil city

This study was aimed to:

- 1. Identify the physical growth pattern of Thalassemic patient( case group) and healthy school( control group) children and adolescents
- 2. Compare the physical growth pattern between Thalassemic patient and healthy children and adolescents (secondary schools ) with their age and gender .

# **Subjects and Methods**

**Study Design And Sample** A cross – sectional descriptive study were carried out at three public(governmental) secondary schools and Hawler Thalassemic Center in Erbil city during the period from1-2-2012 to 30-12-2012 ).This study was conducted in (50) Thalassemic children and adolescents (58.0%) males and (42.0%) females, receiving regular blood transfusions and subcutaneous desferrioxamine therapy and (200 healthy school) children and adolescents (65.5%) male and (34.5%) female from public secondary schools. To identfy the physical growth pattern between Thalassemic and healthy school children and adolescents with their age and gender. A questionnaire was designed that composed of two parts to deal with physical growth pattern,part I consist : demographic information of the study samples such as (age and gender), and part II consist physical growth measurement questions such as ( standing height , sitting height ,weight and nutritional status as body mass index).

Anthropometrical Methods The physical growth parameters of thethalassemic children and adolescents, weight, standing heigh and sitting height were measured by standard scales and standardized stadiometers. Body mass index (BMI) was calculated according to the formula weight in kg=height in m2. Weight, height and BMI for age and sex was calculation in boys and girls, in Hawler Thalassemic Center/ Erbil and secondary schools to measured height and weight in a private area which done by searchers.

**Procedure:** The searchers interviewed participants in Hawler Thalassemic Center and these schools in Erbil city.In school ,those agreeing to participate were instructed how to complete the questionnaire sheet by the interviewers. Participants self-completed the questionnaire sheet and returned it , and in Hawler Thalassemic Center/ Erbil the researcher complete the questionnaire sheet by the interviewers Thalassemic children .

**Statistical Methods:** All the analysis of the results were performed by using the (SPSS, 17), depended on the Body Mass Index (BMI) for age and sex for boys and girls.

**Ethical consideration**:For the purpose of this study a written official permission has been obtained from College of Nursing / Hawler Medical University ,Ministry of Education and secondary schools . These schools agreed to take part in the study within the schools, all students were invited to take part in this study , and from Ministry of Health / Director of Health , HawlerThalassemic Center in Erbil city

# **Results:**

Table (1): Demographic Characteristic of Healthy School Children (ControlGroup) and Thalassemic Children (Case Group).

Variable		Co	Control group N=200		Case group N=50		
		F	%	F	%		
Age	8-11 Years	93	46.5	26	52.0		
_	12-15 Years	83	41.5	17	34.0		
	16-19 Years	24	12.0	6	12.0		
	≥20	0	0.00	1	2.0		
Gender	Male	131	65.5	29	58.0		
	Female	69	34.5	21	42.0		
BMI	< 18	72	20.5	1	2.0		
	18 - 24.9	126	20.5	25	50.0		
	25 - 30	18	9.0	11	22.0		
	> 30	15	7.5	13	26.0		
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F=Frequency %= percentage BMI= Body mass index

This table found that the range of age in the sample was between 8-11 years old ,which present in healthy school children (control group) (46.5%), while in Thalassemic children(Case group) was (52.0%), the highest percentage of gender in both groups were male (65.5%, 58.0%) respectively and the majority of body mass index in both group were range 18 - 24.9 and present(63.0% 50.0%) respectively.

Table 2 :	Physical	Growth	Pattern	Of	Thalassemic	Children	And	Adolescent
(Case Grou	p) Relati	on With	Their Ag	e A	nd Gender .			

Age	Age 8-11 Years			12-15 Years		16-19	Years	≥20 Years	
Physical	M. S.		М.	S.		M. S.		M. S.	
Growth pattern		S. D.			S.D.		SD.		SD.
Standing height	119.4	7.87	129.18		7.97	135.83	12.34	169.00	0.00
Sitting height	57	8	61		4	67	7	69	0.00
Weight	25.12	7.38	29.88		6.49	33.17	7.03	61.00	0.00
BMI	25.12	7.38	29.88		6.49	33.17	7.03	61.00	0.00
Gender	Male				Female				
Physical Growth pattern		M. S.		S.D		М	I. S.	SD.	
Standing height	127.24			13.3	39	123.52		9.68	
Sitting height	59			9		61		5	
Weight	30.24		10.09		)9	25.90		5.99	
BMI	30.24		10.09		25.90		5.99		

BMI= Body mass index S.D.= Stander deviation . M. S.= Mean score

This table shows that in (Thalassemic children) was normal in height (sitting and standing) and abnormal weigh in age group (8-11) but abnormal in age (12-15) years old that means they have high mean score in this age group, while in age group (16-19 and  $\geq$ 20) years old they have growth retardation with highly significant mean score when compared with growth chart. Regarding BMI they have underweight in age groups (8-11and 12-15) years old but they have normal BMI in age groups (16-19 and  $\geq$ 20) years old but they have normal BMI in age groups (16-19 and  $\geq$ 20) years old .

Regarding the age of Thalassemic children shows that males patients reported a normal in height( sitting and standing) but in females reported less than normal percentile, and they have normal weight in both gender, But they reported normal BMI in male patients higher than female.

Age	age									
Physical	8-11	Years	12-15	Years	16-19	16-19 Years				
Growth pattern	M. S.	S. D.	M. S.	<b>S. D.</b>	M. S.	S. D.				
Standing height	138	14	160	17	174	7				
Sitting height	57.47	17.94	81.36	10.84	89.92	4.75				
Weight	34.23	10.87	54.72	15.18	69.67	16.10				
BMI	17.75	3.56	21.61	5.27	22.93	4.53				
Gender										
		Male Female								
Physical Growth pattern	M. S.	S. D.		M. S.	S. D.					
Standing height	anding height 158 20		138	138						
Sitting height	79.21	16.32	56.23		15.23					
Weight	/eight 53.16 18.98		35.26	35.26						
BMI	20.96	5.25	18.09		3.52					

Table 3:	Physical	<b>Growth Pattern</b>	Of healthy	Children	And Ad	olescents (	Control
Group)	Relation	With Their Age	And Gende	r.			

BMI= Body mass index S.D.= Stander deviation .M. S.= Mean score

This table shows that the height(sitting and standing) and weight in healthy children and adolescents in control group were normal in all age groups respectively, also they have normal BMI in all age groups. Regarding the height(sitting and standing) fond that the males have higher than female while the weight in male they have more than in female,

Variable		Age											
	8-11 Years				12-15 Years			16-19 Years					
Physical Growth pattern	Case		Cont	rol	Case Cor		Control	ontrol Case		Control			
	Mea n scor e	SD	Mea n score	e SD	Mea n score	SD	Mean score	SD	Mean score	SD	Mean score	SD	
Standing height	119. 4	7.87	138	8 14	129.2	7.97	160	17	135.8 3	12.34	174	7	
Sitting height	57	8	57.47	7 17.9 4	61	4	81.36	10.84	67	7	89.92	4.75	
Weight	25.1 2	7.38	34.23	3 10.8 7	29.88	6.49	54.72	15.18	33.17	7.03	69.67	16.1 0	
BMI	16.9 6	7.38	17.75	5 3.56	18	6.49	21.61	5.27	18.5	7.03	22.93	4.53	
Variable							Gender						
			Ν	<b>/Iale</b>					Fe	Female			
Physical	Case C			ontrol Case		Control		Control					
pattern	Mean score	SI	)	Mean score	5	SD	Mean score		SD	Mean score	SI	D	
Standing height	127.24	13.39		158	20 5		123.52	9.68		138	11		
Sitting height	59	9		79.21	16.3	2	61	5		56.23	15.23		
Weight	30.24	10.09		53.16	18.9	8	25.90	5.99		35.26	10.04		
BMI	18.78	10.09		20.96	5.25		16.98	5.99		18.09	3.52		

regarding the BMI they have normal BMI in both gender within normal

# Table 4: Comparative Physical Growth Pattern Between Thalathemic Children And Adolescents ( case Group) And healthy Children And Adolescents ( Control Group)

BMI= Body mass index S.D.= Stander deviation . S.= Mean score

Table( 4) shows that Thalassemic children regarding the age they have high mean score than control group (normal child) in (sitting and standing ) height, weight and BMI in all age groups. Regarding the gender the result revealed that the males and females (Thalassemic children) have lower mean score than male and females in control group (Non Thalassemic children) in, height(sitting and standing) and eight but females in (Thalassemic children) have high mean score than male in control group (Non Thalassemic children). while BMI shows high mean score in both gender (Thalassemic children) than in control group (Non Thalassemic children).

#### **Discussion :**

All age groups were compare with respect to age and gender with (Thalassemic children) control group and( non Thalassemic children) case group.

The present study revealed that the majority age of the Thalassemic children (case group,)and healthy children and adolesents (control group) range between 8-11 years old, the highest percentage of gender in both groups were males and the majority of BMI in both group were range 18 - 24.9 as show in(**Table 1**)

Results of this study show growth retardation in age groups (15 -19) years old of the sample of Thalassemic children and adolescents with highly significant mean score when compared with growth chart in age groups (15 -19) years old . Regarding BMI they have underweight in age groups (8-11and 12-15) years old but they have normal BMI in age groups (16-19 and  $\geq 20$ ) years old when compared with growth chart. ( **Table** 2). There are many factors are involved in the growth retardation of patients with Thalassemic, Such as chronic anemia, iron overload, and bone dysplasia receiving regular blood transfusions and subcutaneous desferrioxamine therapy <sup>(9)</sup> The observation of this study was not supported by a study who mentioned that the retardation can occur as early as the first or second year of life in major thalassemia but these abnormalities are more apparent after 6 to 8 years <sup>(10).</sup> The pattern of growth abnormalities in Thalassemic children, in which wasting is prominent early in life followed by stunting in the later years of childhood <sup>(11)</sup>.In (**Table 3**) shows that the height(sitting and standing) weight and BMI in (Non Thalassemic children) control group were normal in all age groups when compared with growth chart. Regarding the height( sitting and standing) they have more than 50<sup>th</sup> percentile in males, but in female they have normal percentile while the weight in male they have more than 50<sup>th</sup> percentile but in female they normal percentile (50<sup>th</sup> percentile), regarding the BMI they have normal BMI in both gender within 50th normal percentile<sup>(6)</sup>. Mean score of height, weight and BMI of Thalassemic children were compared with non Thalassemic children for age and sex(, Thalassemic children regarding the age they have high mean score than control group (normal child) in (sitting and

standing ) height , weight and BMI in all age groups . Regarding the gender the result revealed that the males and females (Thalassemic children) have lower mean score than male and females in control group (Non Thalassemic children) in height(sitting and standing) and weight but females in (Thalassemic children) have high mean score than male in control group (Non Thalassemic children).while BMI shows high mean score in both gender (Thalassemic children) than in control group (Non Thalassemic children) as present in (**Table 4**). The observation of this study was supported by a study who observed the growth retardation in all patient age between 12 to 20 years old (3).This finding of the present study was supported by study who found that a late start in chelation therapy has a negative, yet minor effect, on growth and bone maturation <sup>(11)</sup>.

**Conclusion :** The study conclude that (Thalassemic children ) regarding the age they have high mean score than control group (normal child) in (sitting and standing ) height, weight and BMI in all age groups.

**Recommendation:** the study recommended that Thalassemia was a important health problem in pediatric population and plans need to set up to reduce the number of cases born by development, population screening programme and awareness creating.

# **References :**

- 1. Najafi, , Rabari and Sabzevari ; The Effect of Family-Centered Empowerment Model on the Mothers' Knowledge and Attitudes about Thalassemia Disorder; Iranian Journal of Pediatric Hematology oncology Vol 1. No 3. 2006.
- Rasheed N. Ezzaddin and Ahmed S. Adnan, (2005). Effect of β- Thalassemia on Some Biochemical Parameters. Middle East Journal of Family Medicine [on online]. 7(2). Available from: <u>www.mejfm.com.</u> [Accessed on 21/1/2010].
- 3. Simone, Verrotti, Iughetti, Palumbo, Bartolomeo, and Rosato; Post-transplant complications ,Final height of thalassemic patients who nderwent bone marrow transplantation during childhood; Bone Marrow Transplantation (2001) 28, 201–205
- Moayeri, Oloomi ; Prevalence of Growth and Puberty Failure with Respect to Growth Hormone and Gonadotropins Secretionin Beta-Thalassemia Major; Arch Iranian Med 2006; 9 (4): 329 – 334
- 5. Yaish H M, (2009). Thalassemia. Available from: <u>www.emedicine.medscape.com</u>. [Accessed on 21/1/2010].
- 6. Sayani F., Warner M., Wu J., Wong-Rieger D., Humphreys K., and Odame I., (2009). Guidelines for the Clinical Care of Patients with Thalassemia in Canada. <u>www.thalassemia.ca</u>. [Accessed on 9/6/2010].
- 7. Saxena;Growth Retardation in Thalassemia Major Patients; Int J Hum Genet, 3(4): 237-246 (2003)
- C., and Aprili G., (2000). Safety and efficacy of subcutaneous bolus injection of deferoxamine in adult patients with iron overload. BLOOD [on online]. 95(9): 2776-2780. Available from: <u>www.bloodjournal.hematologylibrary.org</u>. [Accessed on 21/1/2010].

- 9. Novartis Pharmaceuticals Corporation, (2008). Desferal. Available from: www.pharma.us.novartis.com. [Accessed on 21/1/2010].
- Dhamcharee V., Romyanan O. and Ninlagarn T., Genetic Counseling FOR Thalassemia in Thailand: Problems and Solutions. Southeast Asian J Trop MED Public Health [on online]. 2001. 32 No. 2:414-417. Available from: www.tm.mahidol.ac.th. [Accessed on 21/1/2010].
- 11. Eshghi P., (2007). Combined deferiprone and desferrioxamine treatment in thalassemic patients. Iran J Med Sci [on online]. 32(1): 40-44. <u>Available from: www.ijms.sums.ac.ir</u>. [Accessed on 1/4/2010].