**Prevalence Of Positive Antinuclear Antibodies In Thalassemic Patients In Babylon Thalassemic Center**

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**Abstract**

A prospective study was conducted on (104) persons: classified as 74 patients with B-thalassemia major.,30 patients as control group (healthy people ) for the period of May 2008 to February 2009.Their age range from (1-30) years who were attending the thalassemic center at Babylon Maternity and children hospital .By using the kit of Anti nuclear antibodies (ANA). They were studied, as the level of ANAtiter had been estimated, and the results analyzed by using the statistical methods. ( 10.8%) show positivity of ANAtiter at level of (55 U/ml or more) for thalassemic patients and (89.2% ) show negative result below (40 U/ml) for those patients ,while all control group were negative. The positive titers where studied according to deferoxamine treatment (Iron chelating drug) where 87.5% for patients receiving deferoxamine , and 12.5% for patients not receiving this drug.Also the positive titers according to the age of thalassemic patients were analyzed, and the results correlated with the age, where most of positive titers for patients older than ten years (represent 75% of positive titers) ,in compare to (12.5%) from the positive titers for patients their age five to ten years and a similar percent (12.5%) for patients under five years. Finally (62.5%) of positive titers for patient who were spleenectomised, whereas (37.5% )for non spleenectomised patient.

**الخلاصة**

تم أجراء هذه الدراسة على المرضى المصابين بفقر الدم البحري (فقر الم البحري الكبير من نوع B-thalassemia.major في مركز فقر الدم الوراثي في مستشفى بابل للولادة والأطفال للفترة من شهر ايار 2008 ولغاية شباط 2009 وباستخدام العدة التشخيصية المختبرية حيث تم دراسة 104 شخص ,30 شخص منهم كمجموعة سيطرة ,74 شخص مصابين بفقر الدم البحري ,أظهرت نتائج الدراسة 10,8 %نتيجة موجبة للأضداد (عند مستوى(55 وحدة لكل مل ) 89.2 %))نتيجة سالبة(بمستوى اقل من 40 وحدة لكل مل للمرضى المصابين بفقر الدم البحري بينما كانت النتيجة سالبة لمجموعة السيطرة .ثم بعد ذلك تم دراسة النتائج الموجبة حسب استلامهم لعلاج الديفروكسامين وحسب العمر وكذلك العلاقة مع رفع الطحال من عدمة وتبين الاتي حيث وجد أن 87,5%من المرضى الموجبة يستلمون الديفروكسامين و 12,5%للمرضى الذين لا يأخذون هذا العلاج و75%للمرضى الذين تزيد أعمارهم عن عشرة سنة و12,5%للمرضى الذين اعمارهم بين الخامسة والعاشرة ومثلها 12,5 للمرضى الذين تقل اعمارهم عن خمسة سنوات واخيرا تبين ان 62,5%من النتائج الموجبة للمرضى رافعي الطحال و37,5%للمرضى غير رافعي الطحال .

**Introduction**

Thalassemia is actually a group of inherited disease of the blood that affected a person's ability to produce heamoglobulin ,resulting in anemia (Aqarwal etal.,1992) .Heamoglobulin is a protein in red blood cells that carries as oxygen and nutrients to cells in the body (Naney andGray,1997). The two main types of thalassemia are called alpha and beta depending on which part of an oxygen carrying protein in the red cells is lacking (Behrman,2004). The B –thalassemia syndrome is treated by blood transfusion and chelating therapy as Deferoxamine (Desferal) .Deferoxamine is given subcutaneously over 10-12 hr, for 5-6 days a week. (Cunningham *et al*.,2004)

Most autoimmune diseases occurring in humans have a multifactorial and polygenic basis;

Various combinations of genetic, humeral, metabolic and environmental components such as chemicals, toxins, infections (viral, microbial) and also some drugs which serve as either the accelerating, promoting, precipitating or triggering factors determine an individual's disease susceptibility while tissue or organ damage most likely results from an aberrant pathogenic autoimmune response to self antigens (Roitt,1994;Solomon *et al.,*2002). Different studies have shown that in B-

thalassemia patients treated with iron chelators , various auto antibodies are often found, because of these iron chelators or possible development of an autoimmune phenomenon and also show clinical manifestations of arthralgi as, joint pains or joint swellings(Roitt,1994).

[Antinuclear antibodies](http://arthritis.about.com/cs/diagnostic/g/ana.htm) are a unique group of auto antibodies that have the ability to attack structures in the nucleus of cells. The nucleus of a cell contains genetic material referred to as DNA (deoxyribonucleic acid). (Solomon *et al.,* 2002; Turkantoni *et al.,*2008) .

ANAs are found in patients who have various [autoimmune diseases](http://arthritis.about.com/od/diseasesandconditions/g/autoimmune.htm), but not only autoimmune diseases. ANAs can be found also in patients with infections, [cancer](http://cancer.about.com/), [lung diseases](http://lungdiseases.about.com/), gastrointestinal diseases, hormonal diseases, blood diseases, [skin diseases](http://dermatology.about.com/), and in people with a family history of rheumatic disease or drugs like (deferoxamine). (Tan,1982; RaymondandYung,1994 )

The antinuclear antibodies test (ANA) is ordered to help screen for autoimmune disorders such as systemic lupus erythromatous (SLE).Or it is ordered when the doctor suspected that the patients may have developed an additional autoimmune disease (Solomon etal.,2002).

**Aims of the Study**

1- Estimation of prevalence of positive antinuclear antibodies in thalassemic patients in Babylon thalassemic center and compared with control group.

2- The correlation of ANA with certain variables including Age, Deferoxamine intake and spleenectomy .

**Patients and methods**

This study started from May 2008 to the February 2009; include one hundred and four persons (104), thirty (30)

Persons as control group, seventy-four (74) as thalasseimic patients their aged range from 1-30 years by

Using two Kits of ELISA for ANA test (IMTEC-ANAScreen,ELISA for the Quantitative

Determination of Antinuclear Antibodies (Ig(GAM)), the following data have been collected

-age

- deferoxamine intake.

-Spleenectomy or not

Sample collection s: Three ml of blood was collected from each person by vein puncture, serum sample were separated and kept at -20C until the time of processing.

Statistical analysis**:The percent** was used to compare different results during this study .

**Result**

**Fig (1) prevalence of positive ANA titer in thalassemic patients**


**Fig (2) Distribution of positive ANA according to age**

> 10 YEARS

< 5 YEARS

**Fig(3 ) Comparisim of positive ANA titers between patients received deferoxamine and patients not received it.**

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**Fig(4) Comparisim of positive ANA titers between spleenectomised patients and other not.**

**Discussion**

Deferoxamine is a drug used in the treatment of thalassemia but this drugs have many side effects one of these effects when used for long period cause many autoimmune diseases(Karager and Basel, 2003) .Patients with thalassemia major on iron chelators such as deferoxamine shows change in the autoimmune profile suggestive of some human immune alteration. This study shows positive ANA titers in (10.8%) of thalessemic patients and a negative results in(89.2%) in Fig (1) that it is different from other study like (Aqarwal *et al.,*1992) that’s shows a positive ANA titer in 14% of thalassemic patients from thirty –eight patients, our result may be due to irregular treating with this drug because it's not always available in the center and the poor compliance because the deferoxamine gives subcutaneously for five nights per week and it is painful ,also there's shortening in the infusion pumps that’s essential to infuse the drug over the night.

 Fig.(2) that shows an increase in percent of positive ANA with the age of patients which is similar with (Kapadia *et al* ;1980Grady *et al.,*1995; Naneyand Gray, 1997), that studied when the patients used treatment (iron chelator) in older age causes elevated in titer of ANA titer ,also the older ages patients had larger doses of deferoxamine for longer period.

Fig (3) shows acomparisim between the patients used deferoxamine and other not ,where most of positive ANA titers (87.5%) for patients receiving the treatment and only (12.5% ) of positive titers for patients not received this drug , this finding correlated with (Aqarwal etal.,1997;Karager and Basel,2003;Turkantoni etal.,2008) that thalasemic patients depended on deferoxamine for long period show changes on the autoimmune profile.

In the Fig (4) shows that patients who had spleenectomy are liable to the autoimmune disordered and that correlated with othe studies like (Kapadia *et al.,* 1980; Aqqrwal *et al.,*1997; Naney and Gray,1997) who shows that thalassemic patients with spleenectomy susceptible to autoimmune disordered more than other patients who are not spleenectomy.also the spleenectomised patients are older age and use deferoxamine for longer period, in this study 62.5% of positive ANA for spleenectomised patients and 37.5% for other patients .

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