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DEAFNESS IN HYPOTHYROIDISM, AN EVALUATION OF 47 PATIENTS.

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Summary

A prospective study of 47 patients with acquired hypothyroidism seen at Basrah General Hospital between March 1998 to March 1999. It comprises 17 males and 30 females with a ratio of 1.7:3. The evaluation concentrated on the hearing state, audiometric assessment and ECG changes. Out of 47 hypothyroid patients, 35 of them found to have either clinically and/or audiometrically impairment of hearing. Although sensorineural deafness was predominate (17 patients, about 48.6%), an un-expected high incidence of conductive hearing loss was detected (14 patients, 40%). The remaining 4 patients (about 11.4%) emphasized mixed deafness. The severity of deafness was ranged from mild to moderate. No one has suffered a sever deafness. Symmetrical-frequency hearing loss was predominate (17 patients, 48.5%).

Introduction

 \mathbf{T} hyroid hormone plays an important role in hearing development. Hereditary hypothyroidism is frequently associated with sensorineural hearing loss as identified in both animal models and human patients¹.

Correspondence to: Dr. Isam M. Al-Shuraidah Department of Surgery, College of Medicine University of Basrah, Basrah; IRAQ. Reports of hearing loss in hypothyroidism date from Bircher (1883) and in a major text on the thyroid, De Groot and Stanbury (1975) quoted auditory involvement in one-third of all cases^{2,3}.

Hypothyroidism is a cause of hearing impairment worldwide, probably accounting for at least 50000 hearing-impaired infants born each year⁴.

The problem is particularly prevalent in those parts of the world where endemic

goiter is common, as the use of iodized salt can prevent deafness due to hypothyroidism⁵.

The aim of this study is to find out the incidence, severity, and types of deafness in a consecutive series of 47 patients with acquired hypothyroidism.

Patients and Methods

A team of a physician, otolaryngologist and surgeon assessed all patients provisionally-diagnosed as acquired hypothyroidism seen at Basrah General Hospital for a mean period of 12 months (ranging from March 1998 to March 1999).

A total number of 47 patients included in this study. Their ages range from 18 to 75 years with a median age of 44 years. The evaluation included the history physical examination, hearing state, audiometric assessment and ECG changes.

Other possible causes of deafness whether systemic or local have been excluded such as aging-process (prespyacusis), noise trauma, infection, and others. So the inclusion criteria are deafness without cause and primary acquired hypothyroidism.

Results

Subjectively, out of 47 hypothyroid patients 24 of them complained of diminished hearing acuity and 8 patients has tinnitus. Audiometrically, 35 patients (74.4%) found to have a several different element of hearing loss, 12 patients have normal threshold of hearing.

Table I shows the age and sex distribution of the studied group. Females are found to be more affected than males. The male : female ratio was 1.7:3. The age group of 40-50 years was more prevalent.

The geographical distribution of our hypothyroid patients is illustrated in

Table II. The majority (27 patient, 57.4%) are from Basrah Center. The other important bulk (17 patients, 36.1%) are living in North of Basrah.

Among the 35 deaf hypothyroid patients evaluated clinically and audiometrically only 5 patients had unilateral deafness (about 14.3%) while the rest 30 patients had bilateral deafness (about 85.7%).

Sex					
Age	F	%	Μ	%	Total
(in years)					
1-10	0	0	0	0	0
11-20	2	6.7	0	0	2
21-30	7	23.3	3	17.7	10
31-40	7	23.3	4	23.6	11
41-50	7	23.3	7	41.3	14
51-60	6	60.0	0	0	6
61-70	1	3.4	2	11.6	3
> 70	0	0	1	5.8	1
Total	30	63.8	17	36.2	47

Table I. Age and sex distribution of patientswith hypothyroidism

Center of Basrah	North of Basrah	South of Basrah	East				
27	17	2	1				
Table II. Residency of patients withhypothyroidism.							

Types of deafness of the 35 patients are shown in Table III. Although sensorineural deafness was predominate (17 patients, about 48.6%), an un-expected high incidence of conductive hearing loss was detected (14 patients, 40%). The remaining 4 patients (about 11.4%) emphasized mixed deafness.

Type of deafness	Sensorine	Conduct	Mixed	Norma	Total
Side of deafness	ural	ive	—		
Unilateral	2	3	0	0	5
Bilateral	15	11	4	12	42
Total	17	14	4	12	47

Table III. Types and sides of deafness

60 decibels hearing loss).

Regarding the severity of deafness among the 35 deaf hypothyroid patients, 18 patients (51.5%) were within the range of mild deafness (10-30 decibels hearing loss), 17 patients (48.5%) had moderate deafness (40-60 decibels hearing loss). No one of those 35 patients showed a sever deafness (above

Table IV clarifies the different degrees of deafness severity and also correlates the severity of deafness with the frequency of hearing. Symmetricalfrequency hearing loss was predominate (17 patients, 48.5%). High-frequency loss was detected in 14 patients (40%), while only 4 patients (11.4%) suffered a low frequency loss.

Frequency	Low	High	symmetrica	Total	
Severity	W	ţh	trical		
Mild	4	6	8	18	
Moderate	0	8	9	17	
Sever	0	0	0	0	
Total	4	14	17	35	

Table IV. Severity of deafness in correlationwith frequency.

Figures (1,2,3) are examples of pure tone audiograms in different types of frequency loss regardless of the types of deafness.

Discussion

Meyerhoff (1979) concluded that the lower central auditory pathways were functioning and identified the cochlea as the site for the hearing loss associated with hypothyroidism. He was able to support this as the site of the lesion for the sersorineural hearing loss by morphological and biochemical findings⁶.

Ritter (1969) found a conductive loss in rats with myxoedematous infiltrates in the mucosa of the middle ear and austachian tube and Devos (1963) found changes in the spiral ganglion in rats, mice and hamsters⁷.

The deposition of mucopolysaccharides in the skin of the external auditory meatus with thickening reduction of the lumen size may explain conductive deafness in some hypothyroid patients.

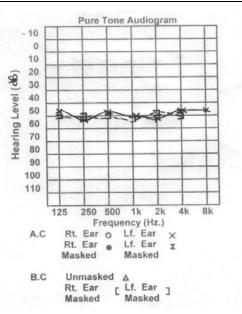
Van't Hoff and Stuart (1979) have reported an incidence of deafness of 85% in a series of 48 patients with myxoedema⁸ in comparison with an incidence of deafness of 74.4% in our study. Also, in their report (Van't Hoff and Stuart), there was no difference between the effect on the high or low frequencies and in some cases the loss was unilateral⁸, in contrast with this study where most of patients were suffered symmetrical hearing loss. (Table II). Van't Hoff and Stuart were in doubt that the deafness no was sensorineural⁸.

Parving and Lyngse (1983) in a series of 15 patients with confirmed myxoedema, median age 76 years demonstrated a bilateral symmetrical or nearly symmetrical sensorineural loss in all patients⁹.

We emphasize that a good number of our hypothyroid patients were suffering from conductive hearing loss.

Hall et al (1985) reported a prospective study undertaken to compare the auditory acuity in hypothyroid patients for a mean period of 7.5 months (range 2-24 months). Auditory thresholds were reduced over all frequencies but the difference being significant only at $2000-4000 \text{ Hz}^{10}$.

Anand et al (1989) carried out auditory investigations in 20 patients with hypothyroidism. Sixteen demonstrated a hearing loss: mild in five (31.25%), moderate in 11 (68.75%).



Audiometrically, 12 patients suffered sensorineural hearing loss, the rest (4 patients) with mixed deafness 11 .

Bhatia et al, 1977 reported Audiometricallyubt that a conductive hearing loss can occur in humans with myxoedema, but more commonly, any hearing loss is

sensorineural¹².

We concluded that among many different causes of deafness, acquired hypo-

Pure Tone Audiogram

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Figure 1. **Pure-tone** Audiogram Symmertical Left Sensorineural Deafness

thyroidism is prooved to be a common cause of deafness both its conductive and sensorineural type, ranging from mild to moderate in severity.

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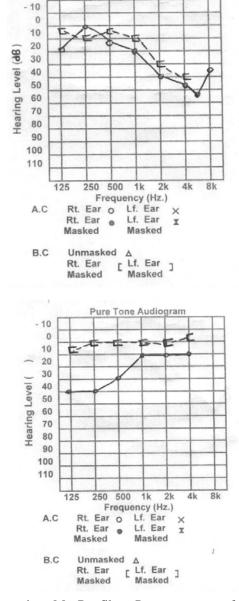
Figure 2. Pure-tone Adiogram victorian neonatal thyroid

High Frequency Right Segsorphogram Deafners

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Figure 3. Pure-tone Adiogram⁷⁹; 89: (suppl. 19).

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