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POSTNATAL RISK FACTORS IN PATIENTS WITH AUTISM SPECTRUM IN AL-NAJAF CITY

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ABSTRACT

Recently, autism spectrum disorder (ASD) has been considered as an increasing problem specially in Iraq. This study was designed to investigate the assessment of postnatal risk factors of ASD in Al-Najaf Province. The purpose of this study was to determine the main demographic and clinical postnatal risk factors of ASD among involved children and to find out the relationip between clinical risk factor and demographic factors. A cross-sectional descriptive approach was designed to meet the previously mentioned objectives of the current study. The period of the study is from Dec. 2018 to March. 2019. A Convenience sample of (40) children with autism spectrum was taken in this study. That the highest percentage of the women's subgroup are : women with ages between (6-8) years old which cotitute (47.5%), male patients (85%), those who live urban residents (87.5%), those who most of their fathers are employee (62.5%), those who most of their mothers are housewives (65%). The present study also revealed that the major clinical risk factors for autism are: patients with low birth weight (65%), jaundice (47.5%) and second hand smoking (2.5%). It was concluded that the most contributing risk factors in the occurrence of ASD are: low birth weight, jaundice, chronic diseases and respiratory infection.

1. INTRODUCTION

Autism is defined as severe psychiatric disorder of childhood marked by severe difficulties in communication and forming relatiohips with other people, in developing language, repetitive, and limited patter of behaviors and obsessive resistance to small changes in familiar surrounding (El-Baz et al., 2011).

Autism is a chronic disorder with an oet before the age of 3 years, characterized by the following three main sets of behavioral disturbances: social abnormalities, language abnormalities and stereotyped repetitive patter of behavior. It is considered one of the pervasive developmental disorders which represent a group of clinical syndromes that have two fundamental elements: developmental delays and developmental deviatio. The number of reported cases of autism increased dramatically in the 1990s and early 2000s. This increase is largely attributable to changes in diagnostic practices, referral patter, availability of services, age at diagnosis, and public awareness (Kidd, 2002).

Diagnosis typically comes from a complete patient history, physical and neurological evaluation. The possible causes of autism include perinatal factors as neonatal anemia, high incidence of respiratory distress syndrome and high incidence of medication usage during pregnancy in the mothers of autistic children, also maternal bleeding after the 1st trimester and meconium in the amniotic fluid. It was also found that autism has an important genetic component although how many genes may be involved remain unclear. The most frequently described are the structural and numerical abnormalities of sex chromosomes, anomalies of chromosome 15 and chromosome 17q21 (Arndt et al., 2005).

Environmental components are another important aspect of research in ASDs. Prior research suggests that parental characteristics, such as age and level of education, may be associated with a risk of autism. Parental age has been shown to be associated with many disorders, such as schizophrenia, childhood cancer and fetal death, however, results from studies of parental age and autism are incoistent. Studies focusing on single perinatal risk factor have reported a positive association for low birth weight (<2, 500 g), gestational age at birth of less than 37 weeks, and congenital malformatio. A gender stratification in one study indicated an increased risk of autism among boys, but not girls of low birth weight (<2, 500 g). Less than complete concordance in monozygotic twi reveals the necessary role of non-genetic factors in the etiology of autism (El-Baz et al., 2011).

Autism spectrum disorders (ASDs) are prevalent neurodevelopmental disorders. ASDs diagnoses are characterized by impairment in social interaction and communication, repetitive behaviors, abnormal movement patter, and seory dysfunction. A kid who has autism has trouble in linking words to their meaning, doesn't like changes in routines, and acts in unusual ways. There is increasing suspicion that autism doesn't have a single cause but it is a complex disorder with a triad of (social impairment, repetitive behavior, communication difficulties) that have distinct causes (Gockley et al., 2011).

METHODS

1. Design of the study

A cross-sectional descriptive approach was designed to meet the previously mentioned objectives of the current study. The period of the study is from Dec. 2018 to March. 2019. A Convenience sample of (40) children with autism spectrum was taken in this study.

2. The study itrument

The researchers have adopted the following tool to assess postnatal risk factors in patients with autism spectrum in Al-Najaf city. The final copy coists of the following parts: 1-Patient's socio-demographic data form. 2-Patient's clinical data form. 3. Statistical Analysis

The following statistical approaches are used in order to analyze the data of the study under application of the statistical package Mega stat (2005): two tailed Chi square-test was used test association between qualitative variables.

3. RESULTS

Table (3.1) shows statistical distribution of study sample (patients) by their socio-demographic data, it explai that the highest percentage of the women's subgroup are : women with ages between (6-8) years old which cotitute (47.5%), male patients (85%), those who live urban residents (87.5%), those who most of their fathers are employee (62.5%), those who most of their mothers are housewives (65%).

Table (3.1): Statistical distribution of study		group (patients) by their Socio-Demographi		phic Data.	
			Patie	nts group	

		Patients group		
Items	Sub-groups	Total = 40		
		Frequency	Percentage	
	3-5	3	7.5	
Age / Years	6-8	Patientsub-groupsTotalFrequencyTotal3-536-8199-1118Male34Female6Urban35Rural5Employee25Free Job15House wife26Employee14	47.5	
	9-11		45.0	
Condor	Male	18 34 6 35	85.0	
Gender	Female		15.0	
Dasidanau	Urban	35	87.5	
Residency	Rural	5	12.5	
Fathar's Job	Employee	25	62.5	
Famel \$ JOD	Free Job	25 15	37.5	
Mother's Joh	House wife	26	65.0	
Would S JOD	Employee	14	35.0	

Table (3.2): Statistical distribution of patients group by their Clinical Data.

		Patients group		
Items	Sub-groups	Total = 40		
		Frequency	Percentage	
Dirth Waight	Underweight	26	65.0	
Bitti weight	Normal	14	35.0	
Respiratory	Yes	8	20.0	
infection	No	32	80.0	
Joundias	Yes	19	47.5	
Jaundice	No	21	52.5	
A 11 amora	Yes	5	12.5	
Allergy	No	35	87.5	
Chronic Disease	Yes	11	27.5	
Chronic Disease	No	29	72.5	
Drug Taking	Yes	52.5	21	

	No	47.5	19
Auditory	Yes	3	7.5
Disease	No	37	92.5
Dland diagona	Yes	3	7.5
blood disease	No	37	92.5
Second Hand	Yes	1	2.5
Smoking	No	39	97.5

According to table (3.2), the statistical distribution of study sample (patients) by their clinical data explai the following risk factors : the major clinical risk factors for autism are : patients with low birth weight (65%), jaundice (47.5%), respiratory infectio (20%), allergy (12.5%), chronic disease (27.5%), auditory disease (7.5%), blood disease (7.5%) and second hand smoking (2.5%).

Table (3.3) shows that there is no significant association between clinical risk factors of patients with autisms and their demographic data, so that it refer to that there is no confounding factor that may interfere with the risk factors.

Table (3.3): Relatiohip b	etween clinical risk factors o	f patients with autisms and thei	r demographic data.
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Demographic Data	Age	Gender	Residence	Father's Job	Mother's Job
Clinical Data	Chi Square (P value)				
Dinth Waight	x2=1.76	x2=1.76	x2=1.24	x2=1.86	x2=0.77
Diftil weight	= 0.18	= 0.16	= 0.23	= 0.12	= 0.32
Respiratory infection	x2=1.86	x2=1.26	x2=1.66	x2=1.32	x2= 1.55
	= 0.14	= 0.15	= 0.26	= 0.46	= 0.26
Joundico	x2=1.34	x2=1.42	x2=1.54	x2=1.35	x2=1.66
Jaundice	= 0.26	= 0.13	= 0.36	= 0.26	= 0.26
Allergy	x2=1.71	x2=1.76	x2=1.55	x2=1.44	x2=1.34
	= 0.16	= 0.15	= 0.12	= 0.35	= 0.25
Chronic Disease	x2=2.76	x2=1.22	x2=1.52	x2=1.65	x2= 1.85
	= 0.06	= 0.18	= 0.14	= 0.17	= 0.35
Auditory Disease	x2=1.55	x2=1.22	x2= 1.42	×2= 1.65	x2= 1.85
	= 0.46	= 0.36	= 0.36	= 0.26	= 0.13
Blood disease	x2=1.76	x2=1.3	×2= 1.6	×2=0.96	x2=0.76
	= 0.16	= 0.2	= 0.26	= 0.46	= 0.56
Negative Smoking	x2=1.22	×2= 1.73	x2= 1.56	x2= 1.66	x2= 2.76
	= 0.16	= 0.16	= 0.16	= 0.16	= 0.16

DISCUSSION

According to table (3.1), the highest percentage age group is between (6-8) years, it mares about (47.5%) at the study sample, this may be attributed to delay in seeking medical care (help) by the family because of neglect or not observing the sings of autism.

The same table (3.1) shows that the a majority at children with autism are male, this result agrees with previous studies that recorded high prevalence at autism among male. A ratio of 4: 1 male to female sex bias has constantly been observe in autism spectrum disorder (ASD). Sex and gender provide unique angles for understanding causal mechanisms in atypical human developmental conditio and should be a central theme in the understanding to autism and its vast heterogeneity note that the term sex refers to the biological and physiological characteristics that define men and women, and 'gender' refer to 'the socially contracted roles, behaviors, activities, and attributes that a given society considers appropriate for men. It was also found that females with autism tend to be identified later than males; this result can be explained by the genetic differences between male & female that lead to difference in anatomy and physiology (Lai et al. 2015).

Researches also mentioned that white mater brain tissue in female is more efficient in communication than male: Nordahl et al. (2015) extended this by showing that there are sex/gender difference in the pattern of altered corpus callosum neuroanatomy in a longitudinal sample of preschooler with autism. They found male and female with autism, early in life in autism, possibly reflecting the key role of early biological factors giving rise to sexdifferential etiological mechanisms, such as prenatal steroids and associated regulatory mechanisms or early neuro-inflammatory mechanisms. Epidemiology and genetic studies suggest a female protective effect (FPE) may account for part of this bias; however, the mechanism of such protection is unknown. Quantitative assessment of ASD symptoms using the Social Responsiveness Scale (SRS) shows a bimodal distribution unique to females in multiplex families. This

leads to the hypothesis locus on chromosome X might mediate the FPE and produce the ASD sex bias. such a locus would represent a major therapeutic target and is likely to have been missed by conventional genome-wide association study (GWAS) analysis (Gockley et al., 2015).

For instance, increased inherited mutation occurring in generation previous and 'carried' by unaffected/undiagnosed females in the family that females-protective ubiquitous mechanism. For predication 2 to be observed, ubiquitous femaleprotective effects have to be overwhelmed by additional mechanisms. Elucidating different risk the mechanisms/factors contributing to FPE should be a focus of etiological investigation (Lai et al., 2015).

The current study showed that about (87.5%) at children are living in urban residence. This result can be explained by differences in life style between rural and urban areas most parents in urban are employee or have works outside their houses, so that their children may be left alone or with communication and emotional status that lead to development of autism some autism found an association between ASD and urbanicity (i.e. higher risk of autism in urban versus rural districts) has been documented (Lai et al., 2012).

According to table (3.2), the majority of patients with autism are classified as low birth weight (65%). this result agrees with the study conducted by Lampi et al. (2012) who recorded that low birth weight and small gestational age, this may be explained by the effect of low birth weight on brain growth. single perinatal risk factor have reported a positive association for low birth weight (2, 500 g), gestational age at birth of less than 37 weeks, and congenital malformation. A gender stratification in one study indicated an increased risk of autism among boys, but not girls of low birth weight (<2, 500 g). Premature infants with very low birth weight often need long hospitalization in the neonatal inteive care unit (NICU). Is also possible that the environment of the NICU adversely affecting emotional and social maturation may result in negative effects on child neurodevelopment. Another possibility is that prematurity ASD may share similar neurodevelopmental antecedents, including exposure to adverse prenatal factors. (Lampi et al., 2012).

The same table reveals that about (47.5%) of patient with autism had underwent from of jaundice. this come in agreement with the wore of Lozada et al. (2015) who revealed that children with ASD have a high percentage of diagnosis with jaundice during the neonatal period, this can be interpreted by the neurotoxicity of high bilirubin during neonatal period. That hyperbilirubinemia in the neonatal period is an important Their findings suggest factor to coider when studying causes of infantile autism. 55% of our patients presented with mild to severe mental retardation, 36% with below average

mentality. EL-Baz et al. (2011) found that children who develop ASD are more likely to have an admission with a diagnosis jaundice in the neonatal period and more likely to require treatment for this jaundice. Children who develop ASD are more likely to have an admission with a diagnosis jaundice in the neonatal period and more likely to require treatment for this jaundice. Conclusion, our study provides further evidence that neonatal unconjugated hyperbilirubinemia associated with the development of ASD. The estimates are consistent with prior literature and there is biologic plausibility. Further prospective studies are needed to clarify specific serum levels of bilirubin in combination with other neonatal risk factors that mediate the association of jaundice and ASD (Lozada et al., 2015). Regarding Allergy, only small percentage patient with ASD (12.5%) have allergic disorders. This can be supported by the study made by Hori et al. (2017) who found that about (14.7%) of ASD patient have asthmatic and allergic disorder. Concerning smoking, (second hand) smoking SHS., only (2.5%) of ASD patient have second hand smoking have smoking. This percentage is very close to that obtained by khalil et al (2018) who recorded by (2%) of patient with ASD had SHS exposure.

CONCLUSION

It was concluded that the most contributing risk factors in the occurrence of ASD are : low birth weight, jaundice, chronic diseases and respiratory infection.

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