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## The Rate and Variety Of Birth Defects In Newborns At Babylon Teaching Hospital For Maternity And Children For The Period (2022-2023)

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ARTICLE INFO	ABSTRACT	
<i>Article history:</i> Received 5 Dec. 2024 Revised 1 Jan 2025, Accepted 9 Jan 2025, Available online 15 Mar. 2025	Birth defects refer to structural changes that appear at birth and can affect any part parts of the body, such as the heart, brain, or feet and represent an important hea issue globally, impacting several millions newborns each year. Of these, 3.3 milli die when they reaching the five age, while 3.2 million experience somatic or mi disabilities. Understanding the various causes of these anomalies is crucial to enal effective healthcare management and intervention. The objective was to assess	
Keywords:	frequency and variety of birth defects in newborns in Hilla City from 2022 to 2023.A	
Birth defects	descriptive design to examine and describe the variables and their interrelationships	
Anomalies	related to birth defects in newborns in Hilla City from 2022 to 2023. In 2022 and	
Malformations	2023 years, a total of 5,034 newborns were delivered among them, 60 were detected	
Frequency	with birth defects, resulting in a rate of 1.19%. The predominant malformations	
Congenital	involved in 2022 were Multiple Congenital malformations (56.52%) and Down's	
	syndrome (13.04%) while in 2023 the predominant malformation involved Multiple	
	Congenital malformations (59.46%) followed by Congenital heart disease (18.92%).	
	An association was found between maternal age and birth defects, with a p-value of	
	<0.05. Additionally, newborn gender was also related to the occurrence of birth	
	defects, with a p-value of <0.05. The research suggests that advancing maternal age is	
	linked to an elevated risk of congenital abnormalities. Consistent prenatal care and	
	diagnostic testing are advised to support prevention, early detection, and, when	
	needed, planned termination.	

### 1. Introduction

Congenital defects are a major cause of perinatal and neonatal deaths globally, affecting both industrialized and low-income countries. These malformations arise from a range of factors, with approximately 40% of cases being idiopathic. There is evidence suggesting a higher prevalence in populations with consanguineous marriages(Sallout etal.,2008; Shetty etal.,2023).

The development of a fetus from a fertilized egg involves a series of intricate processes, any of which can potentially lead to defects or variations. For instance, the

formation of the upper extremities occurs between the fourth and sixth weeks of pregnancy, when the embryo is about an inch long and resembles a miniature baby. Estimates suggest that 1 in 20 infants may present with some form of imperfection (Mashuda etal.,2014; Silesh etal.,2021).

Some congenital abnormalities are inherited in a manner similar to traits such as hair and eye color. These genetic traits may skip generations and manifest only when both parents pass on a recessive gene to their child. Additionally, certain genetic conditions may arise de novo, meaning the baby is the first to exhibit the condition, though it may be passed on to future generations (AbouEl-Ella etal.,2018; Kurdi etal.,2019).

Furthermore, specific medications, such as chemotherapy agents and thalidomide, are known to cause birth defects. Understanding the causes of congenital abnormalities is crucial for improving prevention and treatment strategies(Togoo etal.,2012; Eltyeb etal.,2023).

Recreational drugs, tobacco, and alcohol are known to impact fetal development, but there is no specific evidence linking them to upper extremity abnormalities(Alshehri,2005). condition notable associated One with congenital deformities is congenital constriction bands, where threads of the amniotic membrane detach and wrap around the fingers or hands of the baby (Fida etal.,2007). The exact cause of these bands remains unknown. congenital Many abnormalities arise without a clear explanation, potentially resulting from disruptions in the complex developmental processes of the fetus. Researchers and geneticists are actively working to identify specific diagnoses and determine whether these conditions are hereditary(Sallout etal., 2015).

### 2. Methodology

A descriptive study design was employed to examine the variables and their relationships

#### 3. Results and discussion

The 5,034 newborns were delivered in 2022 and 2023. Of these, 60 were diagnosed with birth defects, resulting in rate of 1.19%.

The predominant malformation involved was Multiple Congenital malformations (56.52%) and Down's syndrome (13.04%) [Table 1]. in this research. This design was selected to fulfill the objective of assessing **the frequency and variety** of birth defects in newborns in Hilla City during the period from 2022 to 2023.

The study was conducted with the necessary permissions from the relevant administrative bodies and ethical committees to ensure compliance with ethical standards and regulations.

The study was carried out at the Babylon Teaching Hospital for Maternity and Children in Hilla City.

The data was collected from the hospital's statistics department for the years 2022 and 2023 where study utilized a random sampling method, involving 5,034 newborns. Out of these, 60 were diagnosed with congenital malformations. Data was statistically analyzed using the (SPSS) version 25 and Microsoft Excel (2016). Various statistical measures were applied to evaluate frequency, percentage and p value. All continuous quantitative variables and overall knowledge scores were tested for normality distribution and adhered to the normal statistical distribution.

# Table 1 Distribution of congenital malformations in 2022.

Malformations	Frequency	Percentage
Down's syndrome	3	13.04
Reproductive system	1	4.35
malformations		
Ichthyosis	2	8.7
Multiple Congenital	13	56.52
malformations		
Musculo-skeletal	1	4.35
system malformations		
Congenital heart disease	1	4.35
Imperforate anus	1	4.35
Abdominal	1	4.35
malformation		
Total	23	100.0

The predominant malformation involved was Multiple Congenital malformations (59.46%) followed by Congenital heart disease (18.92%) [Table 2].

Table 2 Distribution of congenital malformations in

Malformations	Frequency	Percentage	
Lungs malformation	1	2.7	
Multiple Congenital	22	59.46	
malformations			
Congenital heart	7	18.92	
disease			
Reproductive system	4	10.81	
malformations			
Hydrocephalus	1	2.7	
Skin and joints	2	5.41	
malformation			
Total	37	100.0	

There is a meaningful association between mothers' age and birth defects. Also, Newborn gender in relation with birth defects , p value was <0.0.5[Table 3]..

### Table 3 Association among birth defects and

mothers' age and gender of newborn for both years

2022 and 2023. Total number(60).

Variable	Group	Frequency (%)	Congenital anomalies P value
Maternal age	$\leq 20$	32(53.33)	0.027
	20-30	18(30.0)	
	31-40	8(13.33)	
	≥40	2(3.33)	
Newborn gender	Male	34(56.67)	0.041
	Female	26(43.33)	
Years	2022	23(38.33)	0.073
	2023	37(61.67)	

The and distribution occurrence of congenital abnormalities can change considerably over time and across various regions, influenced by a complex combination environmental, of genetic. socio-cultural, racial, and ethnic factors. In low-income countries, where progress has been made in addressing infections and nutritional deficiencies, birth defects have become a leading cause of perinatal death (Taksande et al.,2010).

In the current study, the occurrence of birth defects in newborns was found to be 1.19%. This figure is similar to the results of a study in India, which registered occurrence rates of 2.72% and 1.9%. However, a study conducted in Erbil City, Iraq, observed a higher prevalence of 3.5%. The differences in prevalence rates could be because of variations in study methods, population characteristics, or environmental influences (Othman ,2013).

An interesting feature of this study is the notable correlation between mothers' age and the incidence of birth defects. This result is consistent with similar researchs conducted in Morocco by (Elghanmi et al., 2020) and study of (Green et al., 2010) which also found a significant link between mother age and birth defects. In contrast, a study in India by (Sachdeva et al.2014) did not observe this association, possibly due to regional and contextual differences. Study conducted by ( Reefhuis and Honein, 2004) found that younger age of mother associated with several birth defects.Regarding gender distribution, this study observed a variation in the incidence of congenital anomalies between male and female newborns. Consistent with a study conducted in Iran by (Abdi-Rad et al., 2008)which found congenital anomalies to be more prevalent in female infants, our results also indicate a higher prevalence in females. However, a study in Kuwait by (Madi et al.2005) found a predominance of congenital anomalies in male babies. This discrepancy highlights the need for further investigation into the biological and environmental factors contributing to these gender differences.

In assessing the factors contributing to congenital anomalies, variables such as gender distribution, preterm births, birth weight, family history, and birth order were taken into account. Both hereditary and ecological influences are critical in conditions like congenital heart diseases and neural tube defects (McIntosh et al.,1995,Mashuda etal.,2014).The effect of consanguineous marriages on certain anomalies highlights the

requirement for genetic counseling and understanding initiatives for couples considering such marriages.

Furthermore, factors like family history, calcium intake, and pregnancy-related hypertension are important in relation to congenital anomalies. Health issues during pregnancy, use of medications, contraceptive methods, and supplementation with iron and folic acid also play a role in the risk of certain congenital defects. Recognizing these factors is vital for healthcare contributors to manage pregnancies properly and reduce the risks linked to congenital anomalies.

### 4. Conclusions

Our study revealed that whenever mother age decrease, the risk of congenital defects increases. Also, gender was in significant relation with congenital anomalies happening. Despite advancements in molecular diagnostics for congenital abnormalities, the initial diagnosis and decision to refer patients for further molecular analysis and costly genetic tests often depend on clinical and radiological evaluations. Given the little of other studies, additional research this area in is recommended.

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