

Incidence of Birth Defects at Birth among Babies Delivered at Maternity and Children Teaching Hospital in Ramadi

Fakhri Jamil Al-Dalla Ali 1, Najdat Shukur Mahmood 2, Bilal Kh. Al-Obaidi 3.

¹Ass.Prof. MBChB, MRCP (UK), Dept of pediatrics, Medical College, University of Anbar.

²Lecturer, MBChB, FICMS, Department of pediatrics, Medical College, University of Diyala.

³MBChB, DCH, Dept. of pediatrics, Maternity and Children Teaching Hospital in Ramadi.

Abstract

Background: Birth defects are an important contributor to infant morbidity and mortality among all racial/ethnic groups, and the 3rd leading cause of infant mortality after cancer and accidents in developed countries

Objective: To estimate the incidence of major and minor birth defects at birth and trying to identify some of possible associated factors.

Methods: Between the 1st of February/2009 and the end of October/2009, 1494 newborns were enrolled in this study in Al- Ramadi Maternity and Children Teaching Hospital was examined for birth defects. Full information regarding the mother, newborn, and some socio-demographic factors were recorded after interview of the mother, and full examination was done by the investigator.

Results: Forty-seven newborns (31.46/1000 total birth) had birth defects. The incidence of major anomalies was 16.73 /1000 total birth and minor 14.73/1000 total birth. Among total births, the most common system involved was genitourinary system (27.7%), followed by central nervous system (19.2%), musculoskeletal system (19.2%), oral anomalies (6.4%), skin (6.4%), and then the others. The incidence of birth defects was significantly higher in preterm infants, low birth weight infants, babies who were breech *in utero*, and in babies of mothers in both extreme age (below 20 and above 35 years), or had bleeding during pregnancy, urinary tract infection during pregnancy, and polyhydramnios or oligohydramnios.

Conclusion: The incidence of birth defects is alarmingly high in Al- Ramadi city. The catastrophically successive wars imposed on Iraq undoubtedly have played some role in this problem. Attention should be made for defects at birth in genitourinary system, central nervous system and musculoskeletal system, however defects of cardiovascular system, and internal gastrointestinal tract may be evident sometime after birth.

Keywords: Birth defects, Ramadi, West of Iraq.

Corresponding author: Fakhri Jamil Al-Dalla Ali E-Mail.anbarmedj@gmail.com

Introduction

Birth defects are defined as permanent change produced by an intrinsic abnormalities of development in a body structure during prenatal life¹, they are an important contributor to infant morbidity and mortality among all racial/ethnic groups², and the 3rd leading cause of infant

mortality after cancer and accidents in developed countries^{3,4}.

Birth defects accounted for 490.000 deaths world wide in 1997, the great majority of these deaths were in the 1st year of life⁵. Studies have shown that between 2% and 3% of all infants have major congenital anomalies identified at birth².

Birth defects are either single primary defect or multiple malformations⁶. Single primary defects could be a deformity, malformation, or destruction⁷.

Malformations either due to monogenic defects in 7.5% of patients; in 6%, they are caused by chromosomal anomalies; in 20%, they are the result of mutagenic defects; and in 6-7% they result from known environmental factors such as maternal diseases, infections, and teratogens. In the remaining 60%-70% of patients, malformations are due to unknown etiologies⁸.

Treatment and rehabilitation of children with birth defects is usually costly and complete recovery is usually impossible⁹, hence it is obligatory to find out causative and risk factors for birth defects and prevent them earlier¹⁰.

Currently no nationwide birth defects monitoring in Iraq. In the last two decades there was a feeling of an increase in the number of cases with birth defects in Iraq as a result of sequelae of environmental pollutions caused by the successive wars imposed on Iraq.

To our knowledge, there was no previous studies regarding birth defects in Ramadi, thus the aim of this study is a trial to identify the incidence of birth defects at birth among babies delivered at the Maternity and Children teaching hospital in Ramadi.

Patients and Methods

This is a cross-sectional study that was carried out at the Maternity and Children Teaching Hospital in Ramadi during the period from 1st of February/2009 to the end of October/2009. Because of the practical difficulties, two days per week selected for examination. The total number of all examined babies was 1494 (live and still birth).

History taking and physical examination were performed by one person (the

investigator). A structured form was used for data collection which consists of two parts; maternal characteristics included age, education, diseases during pregnancy, residence and consanguinity while neonatal characteristics included fetal presentation *in utero*, outcome of pregnancy, gestational age, sex and birth weight.

Full physical examination was done to all newborns (live and still birth).

X-rays, echocardiography and ultrasonography were done when necessary for some babies.

The defect was considered as major if potentially life threatening and/or if not corrected, might impair the child's development or well being while classed as minor when affected non vital organs, had little or no functional effect and doesn't cause distress in the neonatal period¹¹.

Gestational assessment of the newborn was done according to new expanded Ballard scoring system¹² taking in to consideration the last menstrual period and ultrasound done before delivery. Babies considered as preterm when gestational age less than 37 weeks¹³.

The weight of newborn was measured by using electronic scale measuring to the nearest 1 gram. The baby was considered to have low birth weight if the weight is equal or less than 2500 gram and normal weight if more than 2500 gram¹².

Distribution of birth defects was classified by the diagnostic standardization of congenital malformation from the *International Classification of Disease (ICD-10)* code¹⁴. Statistical analysis was done by using Chi Square test to compare differences between groups, P value less than 0.05 level is regarded as significant and highly significant when P value less than 0.01 level.

Results

The current study showed that 808(54.08 %) of the studied newborns were males while the others 686 (45.92 %) were females, 1478 babies (98.9%) were

Incidence of Birth Defects ...

Fakhri Jamil Al-Dalla Ali et al

live birth while still-births were 16(1.1%), normal birth weight were 1191(79.7%) and low birth weight were 303(20.3%), full term neonates were 1265(84.7%) while preterm were 229(15.3%).

The number of babies with birth defects at birth were 47 with incidence of 31.46/1000 total birth, twenty five of them had major birth defects (16.73/1000 total birth) while 22 had minor defects (14.73/1000 total birth) (Figure 1).

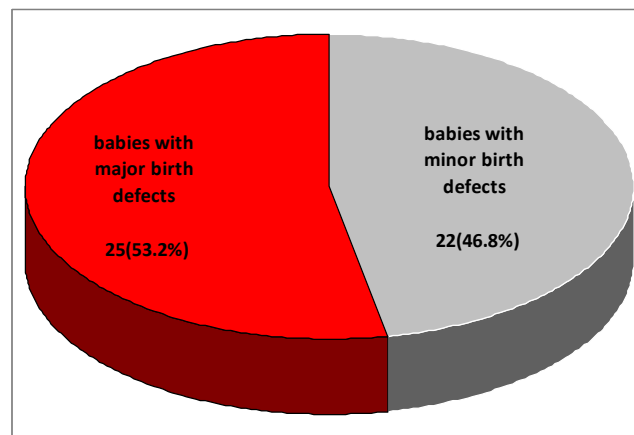
Single defect observed in 43 case (91.5%) while multiple defects in 4 cases (8.5%). Cases with multiple defects included one case with right hydrocele and black not hairy nevus on the abdomen, one case was having unclassified anomalies (single nostril, microphthalmia, small chest and polydactyly), and two cases with Down syndrome.

This study showed that the leading system involved was genitourinary system (27.7%), followed by central nervous system(19.2%), musculoskeletal system(19.2%), then oral anomalies(6.4%), skin(6.4%), and to less extent chromosomal disorders (4.3%), gastrointestinal

tract(4.3%), cardiovascular system(2.1%), eye(2.1%), nose anomalies (2.1%), and virtually unclassified anomalies(2.1%) as shown in (Table 1) .

The incidence of birth defects was statistically highly significant in low birth weight neonates, and preterm infants (P value < 0.01); however the association of birth defects was also significant but to less extent in babies whose mothers aged below 20 or above 35 years old, babies who were breech *in utero*, neonates of mothers who had bleeding, urinary tract infection during pregnancy and in babies of poly or oligohydramnic mothers (P value < 0.05). There was no significant association between birth defects and mother's education, maternal diabetes, maternal hypertension, residence, parental consanguinity, outcome of pregnancy, sex of the baby, (P value > 0.05).

Nevertheless the incidence of birth defects was higher in males, neonates whose parents had consanguineous marriage, and babies of mothers having hypertension, though this was statistically not significant (Table 2).



(Figure 1) Distribution of babies according to major and minor birth defects

Table 1: Distribution of birth defects according to the system involved

Birth Defect	No (%)	Incidence/1000 Total Birth	Birth Defect	No (%)	Incidence/1000 Total Birth
Genitourinary System	13(27.7)	8.7	Skin Anomalies	3(6.4)	2
Major			Minor		
Ambiguous genitalia	1	0.67	cutis aplasia	1	0.67
Bladder extrophy and epispadias	1	0.67	hemangioma on the right aspect of the neck	1	0.67
Minor			black not hairy nevus on the face	1	0.67
Unilateral undescended testes	6	4.01	Chromosomal Disorders	2(4.3)	1.34
Bilateral undescended testes	1	0.67	Major		
Hypospadias	2	1.34	down syndrome	2	1.34
Right hydrocele	1	1.34	Gastrointestinal Tract	2(4.3)	1.34
Right hydrocele and black not hairy nevus on the abdomen	1	1.34	Major		
Central Nervous System	9(19.2)	6.02	esophageal atresia	2	1.34
Major			Ear	2(4.3)	1.34
Anencephaly	2	1.34	Minor		
Hydrocephaly	3	2	bilateral microtia	1	0.67
Myelomeningocele	2	1.34	low set ear	1	0.67
Cranial meningocele	1	0.67	cardiovascular system	1(2.1)	0.67
Encephalocele and microcephaly	1	0.67	Major		
Musculoskeletal System	9(19.2)	6.02	left hypoplastic heart	1	0.67
Major			Eye	1(2.1)	0.67
Achondroplasia	1	0.67	Major		
Omphalocele	2	1.34	bilateral cataract	1	0.67
Minor			Nose	1(2.1)	0.67
Talipes varus	2	1.34	Major		
Talipes valgus	1	0.67	bilatertal co anal atresia	1	0.67
Bilateral polydactyly	1	0.67	unclassified		
Right hand syndactyly	1	0.67	in one baby(single nostril,	1(2.1)	0.67
Right hip dislocation	1	0.67	microphthalmia, small chest, polydactyly)	1	0.56
Oral anomalies	3(6.4)	2	Total	47(100)	31.46
Major					
Cleft lip and palate	2	1.34			
Cleft palate only	1	0.67			

Table 2: Distribution of birth defects by neonatal and maternal characteristics

Characters	Birth defects		Incidence /1000 total birth	P value	Characters	Birth defects		Incidence /1000 total birth	P value
	+ ve No(%) n= 47	- ve No(%) n=1447				+ve No(%) n=47	-ve No(%) n=1447		
Maternal characters					Liquor				
Age of mother(year)									
< 20	6(5)	114(95)	50	<0.05	Normal	40(2.8)	1372(97.2)	28.33	<0.05
20 – 35	33(2.6)	1232(97.4)	26.09		with	4(7.3)	51(92.7)	72.73	
> 35	8(7.3)	101(92.7)	73.39		polyhydramnios	3(11.1)	24(88.9)	111.1	
Mother education(year)					with oligohydramnios				
Illiterates	9(3.4)	255(96.6)	34.09	>0.05	Residence				>0.05
Primary	24(3)	788(97)	29.56		Urban	18(2.8)	635(97.2)	27.56	
Secondary	2(4.9)	39(95.1)	48.78		Rural	29(3.4)	812(96.6)	34.48	
Higher education	12(3.2)	365(96.8)	31.83						
Gestational diseases and characters					Consanguinity				
					Positive	34(3.8)	868(96.2)	37.69	>0.05
					Negative	13(2.2)	579(97.8)	21.95	
Diabetes mellitus					Neonatal characters				
With DM	2(4.1)	47(95.9)	40.82	>0.05					
Without DM	45(3.1)	1400(96.9)	31.14		Outcome of pregnancy				
Hypertension					Singleton	45(3.3)	1327(96.7)	32.79	>0.05
With hypertension	3(5.9)	48(94.1)	58.82	>0.05	Twin	2(2.1)	92(97.9)	21.27	
Without hypertension	44(3)	1399(97)	30.49		More than 2	0(0)	28(100)	0	
Bleeding					Gestational age (week)				
With bleeding	5(7.4)	63(92.6)	73.52	<0.05	< 37	13(5.7)	216(94.3)	56.77	<0.01
Without bleeding	42(2.9)	1384(97.1)	29.45		≥ 37	34(2.7)	1231(97.3)	26.88	
Urinary tract infection					Sex				
With UTI	17(5.8)	278(94.2)	57.62	<0.05	male	29(3.6)	779(96.4)	35.89	>0.05
Without UTI	30	1169(97.5)	25.02		female	18(2.6)	668(97.4)	26.23	
Presentation					Birth weight				
Cephalic	38(2.9)	1277(97.1)	28.89	<0.05	Normal birth weight	27(2.3)	1164(97.7)	22.67	<0.01
Breech	7(5.4)	122(94.6)	54.26		Low birth weight	20(6.6)	283(93.4)	66.08	
Transverse	2(4)	48(96)	40						

Discussion

This study showed that the incidence of birth defects at birth is 31.46/1000 total birth. This rate is higher than that found in many developed as well as some developing countries, while it appears similar to that observed in some neighboring countries⁽¹⁵⁻³⁰⁾ (Table 3). However the expected rate in our

study could be higher as some of the congenital malformations might not be evident at birth like cardiovascular, internal gastrointestinal tract, and renal anomalies. most of other studies made their observations on a follow-up basis over a period ranging from one week to several months.

Table 3: Incidence of birth defects in other regions of the World

Location	Incidence of BD* per 1000	
	Live birth	All birth
Al-Hasa / Saudi Arabia ¹⁵	-	33.4
Giza / Egypt ¹⁶	-	31.7
West of Saudi Arabia ¹⁷	-	29.46
Bahrain ¹⁸	27	-
Spain ¹⁹	20.2	-
Atlanta/USA ²⁰	20.2	-
Sweden ²¹	-	13
Britain ²²	-	12.9
Japan ²³	-	7.05
Tehran/Iran ²⁴	24.1	-
Oman ²⁵	-	24.6
UAE ²⁶	-	14.2
Singapore ²⁷	23.99	-
Maharashta/India ²⁸	-	12.2
Western area/China ²⁹	-	15.4
Korea ³⁰	-	18
this study(Al-Ramadi/Iraq)	-	31.45

* BD= Birth Defects.

The high rate of birth defects at birth in this study could be related to environmental pollutions resulted from the successive wars imposed on Iraq in the last 2 decades. The United States and NATO militaries used depleted uranium penetrator rounds in the 1991 Gulf war³¹, and the 2003 invasion of Iraq³².

The main problem of depleted uranium is that the aerosol produced during impact and combustion of

depleted uranium munitions can potentially contaminate wide areas around the impact site leading to possible inhalation by human beings³³.

Multiple studies using cultured cells and laboratory rodents suggest the possibility of leukemogenic, genetic, reproductive, and neurological effects from chronic exposure³⁴.

A 2005 epidemiology review conclude increased risk of birth defects in offspring of persons exposed to depleted uranium³⁵.

Medical personnel of Basrah hospital in southern of Iraq have reported an increase in the incidence of congenital malformation among babies born in the decade following the 1991 Gulf war³⁶. A medical survey in Fallujah city, Iraq reported an alarmingly high increase in birth defects following Fallujah battle in 2004³⁷.

In this study major birth defects were detected in 25 newborns (53.2%) with incidence of 16.73/1000 total birth while minor birth defects were found in 22 newborns (46.8 %) with incidence of 14.73/1000 total birth. This result is nearly similar to that in Al-Hasa/Saudi Arabia¹⁵ which reported 58% major defects and 42% minor defects. However, this result differs from that reported in the western of Saudi¹⁷ with 93.9% major and 6.1% minor. The current study result showed that single defects were found in 43 cases (91.5%) and multiple defects in 4 cases (8.5%), which is different from that reported in Tehran/Iran²⁴ (65.5% single, 34.5% multiple), and Asir/Saudi Arabia³⁸ (8.8% single, 91.2% multiple). Our results also showed that the most common systems involved respectively were genitourinary system (27.7%), central nervous system (19.2%), and musculoskeletal system (19.2%), while in the western of Saudi Arabia¹⁷ the most common systems involved in sequences were cardiovascular system (25.9%), genitourinary system (19.7%), and musculoskeletal system (15.7%).

In Atlanta/United State America²⁰ the incidence of congenital malformations according to systems involved per 1000 live birth were; cardiovascular system 8.8, genitourinary system 4.3, musculoskeletal system 2.3. In Singapore²⁷ the most common systems involved in sequences per 1000 live birth were; cardiovascular system 9.07, musculoskeletal system 4.98 , genitourinary system 3.12. The increment in the incidence of major birth defects, multiple birth defects and cardiovascular anomalies in these studies as compaired to this study is obviously related to the methodology, we depended mainly on physical examination at birth, while most of other studies were retrospective including all registered cases on follow-up bases. Most cases of cardiovascular system anomalies will not be evident at birth.

The incidence of neural tube defects in this study was 4.02/1000 total birth which was nearly similar to the incidence of Al-Ani study³⁹ (3.3/1000 birth), and in Afghanistan⁴⁰(3.0/1000 live birth), but more than that reported in other countries such as Saudi Arabia^{41,42} were ranged from (0.82 to 1.6/1000 total birth), Tehran/Iran²⁴ (1.7/1000 live birth), Bahrain⁴³ (1.5/1000 total birth), Atlanta/United State America²⁰ (0.73/1000 live birth), and Al-Ain/ United Arab Emirates⁴⁴ (1.3/1000 total birth). This difference may be related to the lack of use of folic acid one month before conception in our region. Furthermore prenatal diagnosis and termination of pregnancy in developed countries may contribute to further reduction.

Regarding the associated factors, there was highly significant association between birth defects and both preterm and low birth weight babies which is comparable to that found by others^{24, 25, 45}; in fact birth defects are a cause of prematurity and low birth weight rather than the reverse. Maternal age above 35 years and below 20 years was significantly associated with birth defects, similar finding was observed in Singapore²⁷. However most other studies showed increased risk only above 35 years old,^{25, 45, 46, 47}. Chromosomal defects are increased when maternal age above 35 years while non-chromosomal defects are increased in both extreme age²⁷. Results showed increase congenital anomalies in consanguineous marriages, though the association was not statistically significant, similar observation also seen in other studies^{17, 25, 36, 48, and 49}.

We conclude from this study that birth defects are alarmingly high as compared to many other countries. Probably this is related to the use of depleted uranium weapons by the united state and NATO militaries during the successive wars imposed on Iraq. Further studies are needed for registration of birth defects on follow up basis, and measurement of radiation in different parts of Iraq.

References

1. Jones KL. smith's Recognizable Pattern of Human Malformations, 4th ed. Pennsylvania (PA), WB Saunders, 1988:1-9.
2. Liu S, Joseph KS, Wen SW. Trends in fetal and infant deaths caused by congenital anomalies. *Semin Perinatal* 2002; 26: 268-76.
3. Behrman RE. The field of pediatrics. In: Behrman RE, Kliegman RM, eds. *Nelson textbook of pediatrics*, 14th Ed. Philadelphia, WB Saunders, 1992:1-5.
4. Forfar JO. Demography; vital statistics and the pattern of disease in childhood. In: Campbell AGM, McIntosh N, eds. *Forfar and Arneil's textbook of pediatrics*, 4th ed. London, Churchill Livingstone, 1992: 1-17.
5. World health organization. *World health report 1998*. Geneva: WHO, 1998; 43- 47.
6. Jones KL. Dysmorphology. In: Behrman RE, Kliegman RM, eds. *Nelson Textbook of Paediatrics*, 14th Ed. Philadelphia, WB Saunders, 1992: 473- 476.
7. Bishop IB, Witt KI, Sloane RA. Genetic toxicities of teratogen. *Mutation Research* 1997; 369:9- 43.
8. Anthony, Boris, Leslie G.B. Dysmorphology. In: Kliegman RM, Behrman RE, eds. *Nelson textbook of pediatrics*, 18th Ed. Philadelphia WB Saunders, 2007 :786-793.
9. Petrini J, Damus K, Johnston RB. Birth defects surveillance data from selected state. *Teratology* 1997; 56(1-2); 115-75.
10. Wald NJ, Smith GL, Denness JW. Serum screening For Down's syndrome bet. 8th and 14th Weeks of pregnancy B. J. obstet Gynecol 1996; 103: 407-12.
11. Christianson RE, Van den Berg BI, Milkovich L, Oechsli FW. Incidence of congenital anomalies among black and white live birth with long-term follow-up. *Am J Public Health* 1981; 71:1333-41.
12. Barbara J, Stoll, Ira A.C. The High Risk Infant. In: Kliegman RM, Behrman RE, eds. *Nelson textbook of pediatrics*, 18th Ed. Philadelphia WB Saunders, 2007: 698-711.
13. Altuncu E, Kavuncuoglu S, Ozemir P, Albayrak Z, Adruc A. The incidence of low birth weight in 5000 live born infants and the etiology of fetal risk factors. *Marmara Medical Journal* 2006; 19(2):46-51.
14. WHO. ICD-10. International Classification of Disease, 10th revision. Geneva: WHO; 2004.
15. Hassib N, Naji K. congenital Malformation; Are they more prevalent in populations with a high incidence of consanguineous marriage. *Annals of Saudi Medicine* 1997; 17(2): 254- 6.
16. Temtamy SA. Genetic epidemiological study of malformations at birth in Egypt. *Eastern Mediterranean health journal* 1998; 4(2): 252- 9.

Incidence of Birth Defects ...

Fakhri Jamil Al-Dalla Ali et al

17. Fida NM, Al-Amaa J, Nichols W, Alqabtani M. A prospective study of congenital malformations among live born neonates at University Hospital in Western Saudi Arabia . Saudi Med J 2007; 28(9):477- 84.
18. Al Arrayed SS. Epidemiology of congenital abnormalities in Bahrain. Eastern Mediterranean health journal 1995; 1(2): 248-52.
19. Martinez-Frias ML. Epidemiological aspect of mendelian Syndromes in a Spanish population sample. I. Autosomal dominant malformation syndromes. American journal of medical genetics 1991; 38: 622-5.
20. Metropolitan Atlanta Congenital defects program; US. Department of Health and Human Services, centers for disease control and prevention, national center on birth defects, developmental disabilities 2002. Atlanta: MACDP, 2002:1-17.
21. The Swedish Centre for Epidemiology. Registration of Congenital Malformations in the Swedish Health Registers 2004. Sweden: EPC;1-10 .
22. Rankin J, Pattenden S, Abram sky L, Boyd P, Jordan H, Stone D, et al. Prevalence of congenital anomalies in five British regions. Arch Dis Child Fetal Neonatal 2005; 90:374-379.
23. Sumiyoshi Y, Natsume N, Nimi T, Furukawa H, Toyota T, Obayashi N,et al . Research on malformation monitoring. Frequency of birth defects in 61859 newborns of Aichi, Gifu, and Mie prefecture. Kosei Rodo Kenkyu Jigyo (Kodomo Katei Sogo Kenkyu Jigyo) Hokousho Heisei, 2003;14: 333-7.
24. Tootoonchi P. Easily identifiable congenital anomalies Prevalence and risk factors. Acta Medica Iranica 2003; 41(1):15-9 .
25. Sawardekar KP. Profile of congenital malformations at Nizwa Hospital, Oman:10-years review. J Paediatr Child Health 2005; 41:323-330.
26. Mohamed S. National congenital abnormalities Registry in UAE. East Mediterr Health J 2006. Jul; 11(4):609-99.
27. Tan KH, Tan TYT, Tan J, Chew S K, Yeo G S. Birth defects in Singapore. Singapore Med J 2005; 46(10):545- 52 .
28. Datta V, Chaturvedi P. Congenital malformation in rural Maharashtra. Indian pediatrics 2000; 37: 998-1001.
29. Cheng N. A base-line survey on birth defects in Gansu Province , West China, Annals of tropical pediatrics 2003; 23: 25-9.
30. Jae-Hyug Y, Yon-Ju K, Jin-Hoon C, Moon-Young K, Jung-Yul H, Soon-Ha Y, et al . A multi-center study for birth defect monitoring systems in Korea. J Korean Med Sci 2004; 19:509-13.
31. Douglas Hamilton (25 January). "NATO: 50 Countries See No Depleted Uranium Illness". Reuters Health Information. Archived from the original, 2003-01-21. <http://web.archive.org/web/20030121211153/http://cancerpage.com/cancernews/cancernews2268.htm>
32. "Is an Armament Sickening U.S. Soldiers?". Associated Press. August 12, 2006. <http://www.commondreams.org/headlines06/0812-06.htm>. Retrieved 2006-11-01.
33. Mitsakou C, Eleftheriadis K, Housiadas C, Lazaridis M, Modeling of the dispersion of depleted uranium aerosol, 2003 Apr. Retrieved January 15, 2009.
34. Miller AC, McClain D. (2007 Jan–Mar). "A review of depleted uranium biological effects: in vitro and in vivo studies". *Rev Environ Health* **22** (1) : 75–89. PMID 17508699.
35. Hindin R. et al.. "Teratogenicity of depleted uranium aerosols: A review from an epidemiological perspective". *Environmental Health*, 2005; **4** (1): 17. doi:10.1186/1476-069X-4-17. PMC 1242351. PMID 16124873. <http://www.ehjournal.net/content/4/1/17>.
36. Al-Sadoon I, Hassan GG, Yacoub AA. Depleted Uranium and health of people in Basrah: epidemiological evidence. Incidence and pattern of congenital anomalies among births in Basrah during the period 1990-1998. Basrah (Iraq): Iraq Ministry of Higher Education and Scientific Research 2002; MJBU;1999;17,1-2. Available from URL:<http://idust.net/Docs/IQSRWrks/SelWks03.pdf>.
37. Busby C, Hamdan M, Ariabi E. Cancer, Infant Mortality and Birth Sex-Ratio in Fallujah, Iraq, 2005–2009, International Journal of Environmental Research and Public Health July, 2010, ISSN 1660-4601.
38. Asindi A. Asindi, Ibrahim Al Hifzi, Wagih A. Bassuni. Major Congenital malformations among Saudi infants in Asir Central Hospital. Annals of Saudi Medicine 1997; 17(2):250-253.

Incidence of Birth Defects ...

Fakhri Jamil Al-Dalla Ali *et al*

39. Al-Ani Z R, Al-Hiali SJ, Al-Mehimdi SM. Neural tube defects among neonates delivered in Al-Ramadi Maternity and Children's Hospital, western Iraq . Saudi Med J 2010; 31(2):163-9.
40. Singh M, Jawadi MH, Arya LS, Fatima. Congenital malformations at Birth among live birth infants in Afghanistan, a prospective study. Indian Journal of Pediatrics 1982; 49(3): 331-5.
41. Maclean MH. The frequency of spina bifida in parts of Saudi Arabia. Saudi Med J 1985; 6:69-73.
42. Thalji AA, Abu Osba YK, Hann RW, Shaman's J, Hamdan J. Incidence of neural tube defects in the eastern provinces of Saudi Arabia. J Kwt Assoc 1986; 20: 94-104.
43. Al-Arrayed S. Congenital anomalies in Bahrain. Med Bull 1987; 9: 70-2.
44. Al-Gazali LI, Sztrihai L, Dawodi A, Bakir M, Varghese M, Varady E , et al. pattern of central nervous system anomalies in a population with a high rate of consanguinity marriage. Clin Genet 1999; 55:95-102.
45. Patel ZM, Adhia RA. Birth defects surveillance study. Indian Journal of Pediatrics 2005; 72:489- 92.
46. Tomatir AG, Deminirhan H, Sorkun HC, Koksak A, Ozerdem F, Cilengir N . Major congenital anomalies: a five-year retrospective regional study in Turkey. Genetics and Molecular Research 2009; 8(1): 19-27.
47. Hegazy IS, AL-Beyari TG, AL-Amir AH, Qureshi NA, Abdelgadir MH. Congenital malformations in primary health care in Al-Qassim region. Ann Saudi Med 1995; 15:48-53.
48. Jaber L, Merlob P, Bu X, Rootter JT, Shohat M. Marked parental Consanguinity as a cause for increase major malformation in an Israeli Arab community. Ann J Med Genet 1992; 44:1-6.
49. Al-Gazali LI, Dawodu AH, Sabarinathan K. The profile of major congenital abnormalities in the United Arab Emirates (UAE) population. J Med Genet 1995; 32:7-13.