Neural Tube Defects in Native Fars Ethnicity in Northern Iran

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(Received 19 Feb 2010; accepted 2 Aug 2010)

Abstract

Background: Neural tube defects (NTD) are one of the leading causes of infant mortality worldwide. This study was designed to determine the prevalence of NTDs among native Fars ethnic groups during 1998-2005, and to identify maternal and demographic factors associated with NTDs.

Methods: We performed a descriptive cross-sectional hospital-based study in Dezyani Hospital, Gorgan, North of Iran, since January 1998 until December 2005. The design was based on a sample of 30,639 births of native Fars ethnic groups. Data were analyzed by using spss V13.5 software and were compared with the chi-square test.

Results: The prevalence of NTDs in Native Fars during the 8-year period was 25.4 per 10000 births (95% confidence interval: 20.1-31.8). The prevalence of NTDs was 20.6/10000 and 30.6/10000 in males and females respectively but this difference was not significant. The prevalence of spina bifida, anencephaly and encephalocele were 12.7, 11.4 and 1.3 per 10000 respectively. The rate of NTD was 48.9/10000 in newborns with mothers aged > 35 years. The highest rate of NTDs and spina bifida was in 2002. The highest and lowest rate of anencephaly was in 2005 and 2003 respectively. Twenty eight percent of the parents had consanguineous marriages. Degree relatedness 3, 4, 5 and 6 of consanguineous marriages were 12.8%, 9%, 3.8%, 2.5%, respectively. Also 47.5% of the parents resided in rural areas.

Conclusion: This investigation showed that the rate of NTDs in Native Fars was higher in Iran. In addition, this rate is higher than the Canada and Ukraine and lower than Chinese people.

Keywords: Neural tube defects, Ethnicity, Epidemiology, Iran

Introduction

Birth defects are one of the leading causes of infant mortality worldwide. Incomplete or incorrect closure of the neural tube during early embryologic development causes neural tube defects (1-3). Miscarriage, stillbirth and disability during lifetime were outcomes of NTDs (4). The etiologies of NTDs are considered complex, and in most cases, the causes of these conditions remain elusive (5). Multifactorial disturbances in embryonic neurulation have been identified as cause of NTDs (6, 7). NTDs are caused primarily by chromosomal abnormalities, single-gene disorders, and environmental agents (8). Exposure to methotrexate, aminopterin and valporic acid, maternal characteristics, racial, ethnic, geographical, nutritional, biological factors and low socioeconomic condition have been recognized as risk factors for developing NTDs (9-12). The prevalence of NTDs in different studies; varies from 1 case in 100 in some regions of China to about 1 case in 2000 in some Scandinavian countries. Overall, the prevalence is approximately 1 in 1000 births (12-15). Previous studies have suggested that there is a racial predilection for this condition. In previous US studies, the NTDs rate varied with ethnicity (16, 17), but potential confounders, including maternal weight (18) and the presence of diabetes mellitus, (19) were not controlled (20). The incidence was highest among the Malay population, compared with the Chinese population and other races (21). Gorgan is the capital city of Golestan Province

Gorgan is the capital city of Golestan Province in northern Iran, where different ethnicities such as native Fars, Turkman, and Sistani reside. The native Fars group is the predominant inhabitants of the region that included 45% of total population. Dezyani is a teaching hospital and a gynological referral center, which is the main site for about 80% of deliveries in Gorgan. This hospital is a referral hospital with an annual rate of more than 6000 deliveries, accounting for 20% of annual birth in Golestan Province of Iran and the largest portion of deliveries (80%) in the city.

The aim of this study was to determine the prevalence of NTDs among native Fars population and to identify maternal and demographic factors associated with NTDs in this area.

Materials and Methods

Data collection

We performed a descriptive cross-sectional hospital-based study and included all live and still-births newborns delivered in the Dezyani Teaching Hospital, Gorgan, from January 1998 until December 2005. This hospital is the largest referral hospital in the city with an annual rate of more than 6000 deliveries that accounts for 80% of deliveries in the city and 20% of annual births in Golestan Province. Patients are usually from moderate to low socioeconomic class families of various ethnic backgrounds.

In Golestan, the three main ethnic groups are Fars, Turkman, and Sistani. The region has a population of about 1.8 million and covers an area of about 20,460 square kilometers. NTDs were defined according to the International Classification of Diseases, Tenth Revision (ICD-10). NTDs were confirmed by a pediatrician (neonatologist). This study aims to estimate the prevalence and trends of congenital malformations in native Fars groups who had three previous generations in this area and was not Turkman. Sistani or other ethnicities and their correlation with maternal variables, and type of neural tube defect, associated malformations, prenatal diagnosis, type of consanguineous marriages and the other demographic information. The design was based on a sample of 30,639 postpartum women after admission for childbirth in maternity hospital in Gorgan, capital city of Golestan province that is a referral center for obstetrics and gynecologic problems. Data were collected through interviews with mothers in the immediate postpartum, as well as by consulting the patient records of both the mothers and newborn infants.

The data were analyzed using SPSS version 15 and STATA SE version 10 softwares and were compared with the chi-square and ANOVA test. Because of rarity of NTD, the 95% confidence interval for prevalence was estimated depends on binomial exact methods. A *P*-value of 0.05 or less was considered statistically significant. Crude and multivariate odds ratios (ORs), along with 95% confidence intervals (CIs), were derived using unconditional logistic regression analysis. All variables were included in the model a priori.

Results

Between 1998 and 2005 there were 30639 births in Native Fars in Dezyani teaching hospital, Gorgan, with 78 newborns and stillbirths recorded with NTDs. The prevalence of NTDs in Native Fars during the 8-year period was therefore 25.4 (95% confidence interval: 20.1-31.8) per 10000 births. There were 33 males and 45 females; the rate of NTD was 20.6/10000 and 30.6/10000 in males and females respectively (χ^2 = 3.0, P= 0.08) (Table 1).

Out of the 78 NTD cases in Native Fars, 39 spina bifida, 35 were anencephalic and 4 had encephalocele. The corresponding prevalence for spina bifida was 12.7/10000 births (10.02 and 14.4/10000 for males and females respectively) ($x^2 = 1.9$, P = 0.17), for anencephaly 11.4/10000 (8.7 and 14.3/10000 for males and females) ($x^2 = 2.1$, P = 0.15) and for encephalocele 1.3/10000 (1.87 and 0.68/10000 for males and females) (Fisher's Exact P value= 0.63).

According to mother's age; the highest rate of NTD was 48.9/10000 in newborns with mothers aged > 35 yr (Table 1). This study showed that 22(28%) of the parents with affected newborns had

consanguineous marriages. degree relatedness 3, 4, 5 and 6 of consanguineous marriages were 12.8%, 9%, 3.8%, 2.5% respectively. one mother was diabetic patient and during pregnancy treated with insulin.

Also 47.5% of the parents resided in rural areas and 52.5% in urban areas.

The rate of NTD and the rate of spina bifida, anencephaly and encephalocele for each year are shown in Fig. 1. The highest rate of NTD was in the year 2002 (40/10000). In addition, the highest rate of spina bifida and encephalocele was in the year 2002. The lowest rate of NTD was in 2000. The highest and lowest rate of anencephaly was in 2005 and 2003. The trend of NTDs, spina bifida, anencephaly and encephalocele during 1998-2005 is depicted in Fig. 1.

According to the ANOVA variant test and due to TUKY test we could observe the significant difference in the prevalence of NTD in 8 yr separately (Table 2).

Table 1: Prevalence of neural tube defects (per 10 000) by sex and mother's age, 1998-2005

Variable	Total No. of births			Anencephaly No/10000		Encephalocele No/10000		Total No/10000		x^2	P value
Sex male female	15962 14677	16 23	10.02 14.4	14 21	8.7 14.3	3	1.87 0.68	33 45	20.6 30.6	3.0	0.083
Mother's age (yr)											
15-19	3677	4	10.8	6	16.3	1	2.7	11	29.9	2.25	0.20
20-34	25737	31	12.04	27	10.4	3	1.1	61	23.7	3.25	0.20
≥ 35	1225	4	32.6	2	16.3	0	0	6	48.9		

 Table 2: Differences of NTD prevalence during 8 years of study (1998-2005)

Year	Years	P value	95%CI (lower bound)	95%CI (upper bound)
1998	2002	0.001	-29.4	-6.57
1999	2002	0.020	-24.4	-1.57
2000	2002	0.000	-32.7	-9.87
	2005	0.008	-26.1	-3.27
2001	2002	0.003	-27.8	-4.97
2002	1998	0.001	6.57	29.42
	1999	0.020	1.57	24.42
	2000	0.000	9.87	32.72
	2001	0.003	4.97	27.82
	2003	0.000	8.94	31.79
	2004	0.001	7.32	32.87
2002	2002	0.000	-31.7	-8.94
2003	2005	0.013	-25.1	-2.34
2004	2002	0.001	-32.8	-7.32
2004	2005	0.035	-26.2	-0.72
	2000	0.008	3.27	26.1
2005	2003	0.013	2.34	25.1
	2004	0.035	0.72	26.2

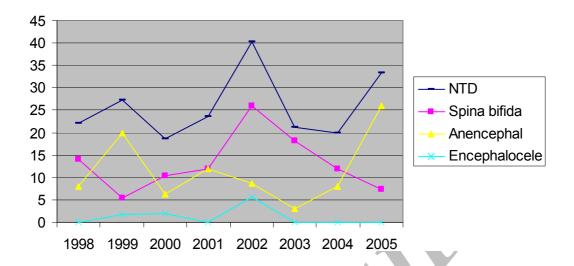


Fig. 1: Annual rates of neural tube defects and its classification (/10 000 births)

Discussion

The overall NTD rate in native Fars ethnicity for the period of 8 yr study period was found to be 25.4 /10 000 births, which this finding is nearly in agreement with previous report. Our previous report showed that the rate of NTDs in native Fars group during 1998-2000 was 23.5/10 000 births but in total population in Gorgan region the rate was 31/10 000 births. Furthermore the rate of NTDs in this study is more than reports from other parts of world with various races/ethnicities including Canada with 1.41/1000 (15) and 3.9/1,000 in 2002 (22), Cape Town 1.74-0.63/ 1,000, 20 yr period (23), United States of America (USA) 9.3 to 14.6/10 000 (12), South Africa 1.74/1000 (24), Southern Africa 000)1980-1984, (25) southern Nigeria 0.95/1000 (26), Northwestern Ukraine 2.1/1000 (2000-2002) (27), Germany 15.0/10 000 (28), the north of England 17.9/10 000 (29) and north of France 10.9/10 000 (30). It is also higher than the capital city of Iran "Tehran" where it was 17.6/10000 (31) (1969-78). The rate is lower than the rate of north-west of Iran (Hamadan) 50.1/10 000 (32) and Kordestan 55.0/10000 (33). This rate was also lower than the other countries such as in rural Transkei district of Umzimkulu which was 3.79 /1000(25), in China 6.0/1000 (34), in Shanxi province of China

199.38/10 000 (4), in Turkey 30.1/10 000 (10), in South Carolina 10.29/10 000 (1992-1996) (4, 35) and in Cameron County, Texas (USA) 27/10,000 (1990-1991) (36).

These variations in different studies could be explained by the influence of racial and social factors in various parts of the world, which are commonly explained as genetic disorders. Geographical, nutritional, socioeconomic, and biological factors could also be involved. Other reasons for these variations in birth defect prevalence are the type of sample (referral hospitals would be expected to have higher rates) and method of diagnosis. Spina bifida was the most common NTD in our study, which agrees with other studies (37, 38), followed by anencephaly and encephalocele. The rate of cystic spina bifida in our study population was 12.7/10 000, which is higher than 6.2/10 000 in France (39), 5.5/10 000 in Atlanta (40), 7.1/10 000 in Texas (12), 1.09/ 10 000 in Saudi Arabia (41), 3.8/10 000 in Tehran (capital of Islamic Republic of Iran) (31) and in Hamadan (north-west province) with 6.98/10 000 (32). The rate of anencephaly in our study was 11.4/10 000, which is higher than the other studies such as 6.0/10 000 in South America (42), 3.7/10 000 in Atlanta (40), 6.4/10 000 in Texas (12) and 8.0/10 000 in Tehran (31). However, the rate in our study was lower than in

Hamedan with 15.6/ 10 000, China with 87.0/10 000 and Turkey with 16.4/10 000 (32). The rate of encephalocele (1.3/10 000) that was nearly similar to studies in the USA (1.03/10 000) (12) and Atlanta (1.4/10 000) (40). However, encephaloceles were significantly more common among the offspring of Hispanic women (adjusted prevalence ratio: 1.91) 1999-2002 (5).

Previous studies reported that the rate and distributions of many of the birth defects such as NTDs related to the sex (4). Regarding sex differences, our results indicate that the rate of NTD was higher in females than males (male to female ratio= 0.73), as reported by other researchers (4, 12, 23, 25, 35, 43, 44). The male to female ratio was 0.66 for an encephaly and 0.69 for spina bifida, which is also comparable to other studies (4, 12, 23, 31, 35, 39, 41, 42, 45). For example, in the USA the ratio for all NTD was 0.62, for anencephaly 0.54 and for spina bifida 0.68 (12). Our research showed that the highest rate of affected newborns was in mothers aged \geq 35 years (48.9/10 000), 29.9/10 000 in mothers aged 15-19 yr and 23.7/10 000 aged 20-34 yr. Our study showed a U-shaped curve with higher rates in mothers aged under 19/20 years and over 35 yr (12, 23, 24, 39, 46, 47) which was in contrast with other studies (4, 5, 35). Thus, age is a complex risk factor in NTD and this issue needs more investigation. Some researches have shown that the rate of consanguineous marriage is high in NTD births (38, 41). In our study, 28% of parents with affected newborns had consanguineous marriage, although this rate is lower than in Saudi Arabia (89% of the spina bifida parents) (41) and higher than in South Africa (24). In addition, a report from North-west of Iran showed that 23% of parents with healthy infants had familial marriages (48) .The possibility that consanguinity could be a risk factor for NTD in a population requires further research. In this study, 47.5% and 52.5% of parents with affected newborns lived in rural and urban areas respectively. A greater prevalence of NTDs at birth has

been shown for rural areas compared with urban areas (49, 50). A report from China (1988–1991) indicated the prevalence of NTD in rural areas (44.3/10 000) was 3 times higher than urban areas (14.4/10 000) (51). It may be due to factors such as high population growth rates and socioeconomic factors. In our study the highest rate was seen in the year 2002 (40/ 10000), but in Quebec city of Canada, the average NTD prevalence decreased from 12.2/1,000 in 1993 to 3.9/1,000 in 2002 (22). In our study the relation between prevalence and year is seen in Fig. 1, the highest rate is seen in 1999, 2002 and 2005 which is due to immigration, socioeconomic, agriculture condition or nutritional factors (52).

According to our findings, interfamilial marriage may play a role in the NTD rate in this region of the Islamic Republic of Iran, although there could also be effects of environmental and nutritional factors. In this research, we could not study abortions and therefore our results may be underestimated.

In conclusion, the present study confirmed the previous reports of high prevalence of NTDs in this region (53). This approach has the advantage of capturing all essential information necessary for an accurate evaluation of NTDs prevalence in our region, and is applicable for other studies estimating the prevalence of birth defects. These findings will help establish a database for future studies, which will focus on multiplex causes and preventive factors to reduce the prevalence of NTD in this region.

Ethical considerations

Ethical issues (Including plagiarism, Informed Consent, misconduct, data fabrication and/or falsification, double publication and/or submission, redundancy, etc) have been completely observed by the author (s).

Acknowledgments

With special thanks to Dr Amir Hossein Noohi for his kindly thoughtful guides, we would like to thank Dr Arzo Mirfazeli and the personnel of Pediatric and Gynecology Wards of Dezyani Hospital specially Mrs Sedehi M, Nazhdaghy A, Asgaripour N, Naziri H, and Heidary R for their participation in this study. The authors declare that they have no conflicts of interest.

References

- 1. Northrup H, Volcik KA (2000). Spina bifida and other neural tube defects. *Curr Probl Pediatr*, 30: 313-32.
- 2. Frey L, Hauser WA (2003). Epidemiology of neural tube defects. *Epilepsia*, 3: 4-13.
- 3. Bol KA, Collins JS, Kirby RS (2006). Survival of infants with neural tube defects in the presence of folic acid fortification. *Pediatr*, 117(3): 803-813.
- 4. Gu X, Lin L, Zheng X, Zhang T, Song X, Wang J, et al. (2007). High Prevalence of NTDs in Shanxi Province. A Combined Epidemiological Approach. *Birth Defects Research*. (Part A), 79: 702–7.
- 5. Wen S, Ethen M, Langlois PH, Mitchell LE (2007). Prevalence of encephalocele in Texas, 1999-2002. *Am J Med Genet*, 15; 143(18): 2150-55.
- 6. Copp AJ (1990). The embryonic development of mammalian neural tube defects. *Progress In Neurobiology*, 35: 363-403.
- 7. Golalipour MJ, Mobasheri E, Vakili MA, Keshtkar AA (2007). Epidemiology of neural tube defects in northern Iran, 1998–2003. *East Mediterr Health J*, 13(3): 560-566.
- 8. Hall JG, Friedman JM, Kenna BA, Popkin J, Jawanda M, Arnold W (1988). Clinical, genetic, and epidemiological factors in neural tube defects. *Am J Hum Genet*, 43: 827-37.
- 9. Gucciardi E, Pietrusiak MA, Reynolds DL, Rouleau J (2002). Incidence of neural tube defects in Ontario, 1986–1999. *CMAJ*, 167(3):237–40.
- 10. Tuncbilek E, Boduroglo K, Alikasifoglu M (1999). Neural tube defects in Turkey: prevalence distribution and risk factors. *Turk J Pediatr*, 41(3):299–305.

- 11. Van der Put NMJ, Steegers-Theunissen RPM, Frosst P, Trijbels FJM, Eskes TKAB, van den Heuvel LP, Mariman ECM, den Heyer M, Rozen R, Blom HJ (1995). Mutated methylenetetrahydrofolate reductase as a risk factor for spina bifida. *Lancet*, 346:1070–1071.
- 12. Hendricks KA, Simpson JS, Larsen RD (1999). Neural tube defect along the Texas- Mexico border, 1993-1995. *Am J Epidemiol*, 149(12):1119–27.
- 13. Li Z, Ren A, Zhang L, Ye R, Li S, Zheng J, Hong S, Wang T, Li Z (2006). Extremely high prevalence of neural tube defects in a 4-county area in Shanxi Province, China. Birth defects research. Part A. *Clinical and Molecular Teratol*, 74(4): 237–40.
- 14. Jacobsen P (1996). Regional variations in neural tube defects and alfa-fetoprotein screening in Denmark 1983–88. *Acta Obstet Gynecol Scand*, 75(7):620–3.
- 15. DeWals P, Trochet C, Pinsonneault (L1992). Prevalence of neural tube defect in the province of Quebec. *Canad J Public Health*, 90(4): 237–79.
- Greene WB, Terry RC, DeMasi RA, Herrington RT (1991). Effect of race and gender on neurological level in myelomeningocele. *Dev Med Child Neurol*, 33: 110-7.
- 17. Feuchtbaum LB, Currier RJ, Riggle S, Roberson M, Lorey FW, Cunningham GC (1999). Neural tube defect prevalence in California (1990–94): eliciting patterns by type of defect and maternal race/ ethnicity. *Genet Test*, 3:265-72.
- 18. Werler MM, Louik C, Shapiro S, Mitchell AA (1996). Prepregnant weight in relation to risk of neural tube defects. *JAMA*, 275:1089-92.
- 19. McLeod L, Ray JG (2002). Prevention and detection of diabetic embryopathy. *Community Genet*, 5:33-9.
- 20. Ray JG, Vermeulen MJ, Meier Ch, Cole DEC, Wyatt PR (2004). Maternal ethnicity and risk of neural tube defects: a

- population-based study. *CMAJ*, 171(4): 343-45.
- 21. Tan KB, Tan SH, Tan KH, Yeo GS (2007). Anencephaly in Singapore: a ten-year series 1993-2002. *Singapore Med J*, 48(1): 12-5.
- 22. Tairou F, De Wals P, Bastide A (2006). Validity of death and stillbirth certificates and hospital discharge summaries for the identification of neural tube defects in Quebec City. *Chronic Dis Can*, 27(3):120-4.
- 23. Buccimazza SS, Molteno CD, Dunne TT, Viljoen DL(1994). Prevalence of neural tube defects in Cape Town, South Africa. *Teratology*, 50(3):194-9.
- 24. Buccimazza SS, Molteno CD, Dunne TT, Viljoen DL (1994). Prevalence of neural tube defects in Cape Town, South Africa. *Teratol*. 1994; 50: 194-9.
- 25. Ncayiyana DJ (1986). Neural tube defects among rural blacks in a Transkei district. A preliminary report and analysis. S Afr Med J, 69(10): 618-20.
- 26. Ugwu RO, Eneh AU, Oruamabo RS (2007). Neural tube defects in a university teaching hospital in southern Nigeria: trends and outcome. *Niger J Med*, 16(4):368-71.
- 27. Wladimir Werteleck (2006). Birth defects surveillance in Ukraine: a process. *J Appl Genet*, 47(2): 143–149.
- 28. Koch M, Fuhrmann W (1984). Epidemiology of neural tube defects in Germany. *Hum genet*, 68(2): 97-103.
- 29. Rankin J, Glinianaia S, Brown R, Renwick M (2000). The changing prevalence of neural tube defects: a population-based study in the north of England, 1984-96. *Paediatr Perinat Epidemiol*, 14(2):104-10.
- 30. Alembik Y, Dott B, Roth MP, Stoll C (1995). Prevalence of neural tube defects in northeastern France, 1979-1992 impact of prenatal diagnosis. *Ann Genet*, 38(1):49–53.
- 31. Farhud DD, Walizadeh GhR, Kamali MS (1986). Congenital malformations and ge-

- netic diseases in Iranian infants. *Hum genet*, 74:382–5.
- 32. Farhud DD, Hadavi V, Sadeghi H (2000). Epidemiology of neural tube defects in the world and Iran. *Iranian J Public Health*, 29(1-4):83–90.
- 33. Mohammadbegi R, Rahimi E (2002). Neural tube defects in newborns in Bacat hospital of Sannadaj in Kordestan province in Iran. *Kord Univer Med J*, 6(4): 36–40.
- 34. Moore CA, Li S, Li Z, Hong SX, Gu HQ, Berry RJ, Mulinare J, Erickson JD. (1997). Elevated rates of severe neural tube defects in a high prevalence area in northern China. *Am J Med Genet*, 73:113–8.
- 35. Stevenson RE, Allen WP, Shashidhar Pai G, Best R, Seaver LH, Dean J, et al. (2000). Decline in Prevalence of Neural Tube Defects in a High-Risk Region of the United States. *Pediatr*, 106(4): 677-83.
- 36. Missmer SA, Suarez L, Felkner M, Wang E, Merrill AH, Rothman KJ, et al. (2006). Exposure to Fumonisins and the Occurrence of Neural Tube Defects along the Texas–Mexico Border. *Environ Health Perspect*, 114(2): 237–241.
- 37. Harris JA, James L (1997). State-by-state cost of birth defects-1992. *Teratol* 56(1–2): 11–6.
- 38. Lynberg MC, Khoury MJ (1990). Contribution of birth defects to infant mortality among racial/ethnic minority groups, United States, 1983. *Morbidity and Mortality Weekly Report*, 39 (No. SS-3):1–12.
- 39. Strassburg MA, Greenland S, Portigal LD, Sever LE (1983). A population-based case control study of anencephalus and spina bifida in a low-risk area. *Dev Med Child Neurol*, 25(5):632–41.
- 40. Rowland CA, Correa A, Cragan JD, Alverson CJ (2006). Are encephaloceles neural tube defects? *Pediatrics*, 118(3): 916-23.
- 41. Murshid WR (2000). Spina bifida in Saudi Arabia: is consanguinity among the parents a risk factor? *Pediatr Neurosurg*, 32(1):10–2.

- 42. Castilla EE, Orioli IM (1985). Epidemiology of neural tube defects in South *America*. *Am J Med Genet*, 22(4): 695-702.
- 43. Stoll C, Dott B, Roth M. P, Alembik Y (1988). Aspects etiologiques et epidemiologiques des anomalies du tube neural. Etiologic and epidemiologic aspects of neural tube defects. *Archives françaises de pédiatrie*, 45(9):617–22.
- 44. Dolk H, De Wals P, Gillerot Y, Lechat MF, Ayme S, Cornel M, Cuschieri A, Garne E, Goujard J, Laurence KM (1991). Heterogeneity of neural tube defects in Europe: the significance of site of defect and presence of other major anomalies in relation to geographic differences in prevalence. *Teratol*, 44: 547-59.
- 45. Cragan JD, Roberts HE, Edmonds LD, Khoury MJ, Kirby RS, Shaw GM, Velie EM, Merz RD, Forrester MB, Williamson RA, Krishnamurti DS, Stevenson RE, Dean JH (1995). Surveillance for anencephaly and spina bifida and the impact of prenatal diagnosis-United States, 1985-1994. MMWR CDC Surveill Summ, 44 (4):1–13.
- 46. McDonnell RJ, Johnson Z, Delaney V., Dack P (1999). East Ireland 1980–1994: epidemiology of neural tube defects. *Epidemiol Community Health*, 53(12): 782–8.

- 47. Bound JP, Francis BJ, Harvey PW (1991), Neural tube defects, maternal cohorts, and age: a pointer to aetiology. *Arc Disease In Child*, 66: 1223-6.
- 48. Pourjafari H, Anvari N (2000). The survey of prevalence and different forms of consanguinity marriages in Hamedan (West of Iran). *J Hamed Univer Med Scienc*, 7(3):30–3.
- 49. Carter CO (1974). Clues to the aetiology of neural tube malformation. *Dev Med Child Neurol*. 16(suppl. 32):3–15.
- 50. Slattery ML, Janerich DT (1991). The epidemiology of neural tube defects. A review of dietary intake and related factors as etiologic agents. *Am J Epidemiol*, 133(6): 526-40.
- 51. Wu Y, Zeng M, Xu C, Liang J, Wang Y, Miao L, Xiao K. (1995). Analyses of the prevalences for neural tube defects and cleft lip and palate in China from 1988 to 1991. *Henan yi ke da xue xue bao*, 26(2):215–9.
- 52. Afshar M, Golalipour MJ, Farhud D (2006). Epidemiological aspects of neural tube defects in South East Iran. *Neurosci*, 11(4): 289-92.
- 53. Golalipour MJ, Vakili MA, Arya B (2003). Neural tube defects in newborns in the south-east of the Caspian sea border (Gorgan, Iran 1998-2000). *Med J Islam Repub Iran*, 16:199-203.