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Immune Response to Measles Vaccine in Primary School Students

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Abstract: This cross sectional study was performed to estimate the prevalence of serological evidence of immunity to Measles in primary school students (age 6-12) that have a history of twice vaccination against measles, on 9 and 15 month old and determining the appropriate age to re-vaccination. Multistage sampling was used to select students to participate in the present study. Immunization status and documentary evidence of immunization was recorded from the Personal Health Record. Measles antibodies were measured using enzyme-linked immunosorbent assay (ELISA). Sufficient blood for antibody testing was obtained from 1665 children, 975 (57%) were girls and 720 (43%) were boys, respectively and mean age was 9.17 ± 1.53 years. On the whole, 1198 subjects (72%) were seropositive and 467 (28%) were seronegative at all. Among girls and boys, 72 and 71%, were seropositive respectively. These differences were not statistically significant between the two sexes ($p = 0.404$). Antimeasles antibody titer decreased with increasing age from 6 to 10 years old, (76.5% in 6 years old group vs. 71% in 10 years old group) and then rise to 78% in 11-12 years old group. These differences were not statistically significant between age groups ($p = 0.775$). the immunity produced by the measles vaccine is not enough. Decrease of protective effect of measles vaccine, suggests the necessary of pre-school revaccination.

Key words: Measles, immunization, antibody, children

INTRODUCTION

Measles is one of the contagious diseases which is still identify as one of fatal diseases in childhood. The incidence rate of measles has obviously decreased after extensive vaccination program, but some limited outbreaks occur in older patients, because of antibody falling down (secondary vaccine failure) (Cutts *et al.*, 1991).

In spite of efficient vaccines and widespread vaccination programs, Measles Virus (MV) remains a major human pathogen. Worldwide, it still causes over half a million deaths per year and is a major cause of child mortality. Death is often caused by accompanying secondary infections, which are favoured by MV-induced immunosuppression. Vaccine coverage of over 94% is required to interrupt transmission of measles in a population (Anderson and May, 1985). In under 6 years old children, Vaccine coverage of over 93% reduce the transmission to a very low level (Hutchins *et al.*, 2004). Actual coverage is much lower, particularly in developing countries, where cost and lack of infrastructure are a hindrance. However, even in countries with adequate health systems, vaccine coverage is not high enough to prevent spread of

the highly contagious virus. In 2002 over 10,000 cases of measles were reported in Western Europe and many more may have gone unreported, as measles is not a notifiable disease in many countries (Anonymous, 2000).

Revaccination in the vaccines with low or undetectable neutralizing antibody titres can maintain a significant lymphoproliferative response 6 months after revaccination (Ward *et al.*, 1995). Since the half-life of human IgG is 1-2 months (Lee *et al.*, 2001; Caceres *et al.*, 2000), the long-term virus-specific antibody levels are maintained by antibody-secreting cells (Plasma cells) and memory B cells (Slifka and Ahmed, 1996). Thus, vaccines with a lower decay rate may keep their antibody-secreting cells living longer. Therefore, antibody titers in the 1 and 2-dose vaccines may not differ significantly in an intermediate period (≤ 6 years) post vaccination but the 2-dose vaccines may have a better cellular immune response and immune memory. This hypothesis may explain why the 2-dose vaccines have a lower risk of measles infection than the 1-dose vaccines who have received vaccination after 12 months of age to reduce the risk of primary vaccination failure in the USA and Canada where the boosting from wild virus is rare (Thomas *et al.*, 1999; DeSerres *et al.*, 1999).

Routine vaccination against measles for infants was introduced in Iran in 1967. Despite the reduction of measles's mortality and morbidity, we are still having local epidemics especially in ages of 15-20 years and this disease is still one of important health sector problems in Iran (Moradi *et al.*, 2001). Measles is a preventable disease and can be eradicated by increasing the vaccine coverage and promoting the motivation for vaccination, in accordance with the worldwide measles strategy (Nakayama *et al.*, 2003). The effectiveness of vaccination programs varies depends on age of subjects in vaccination time, coverage of vaccination programs, quality and specifications of vaccines and planning of immunization programs (Cvjetanovic *et al.*, 1998).

Community-based seroepidemiologic studies are performed to monitor the effectiveness of measles immunization programs and to estimate the decay rate of vaccine-induced measles IgG titers. This study was performed to estimate the prevalence of serological evidence of immunity to Measles in primary school students (age 6-12) that have a history of twice vaccination against measles, on 9 and 15 month old and determining the appropriate age to re-vaccination.

MATERIALS AND METHODS

The survey was performed on 6-12 years old primary school students from 19 educational sectors of Tehran (2003-2005). The sample size was determined considering an expected prevalence of measles virus antibodies of 70% with a worst acceptable error of 5% and a confidence interval of 95%. Multi stage sampling was used to select schools and then students from 19 educational sectors of Tehran; Overall, 1779 students were enrolled voluntarily and Informed consent was obtained from parents or legal guardians of students after informing them about the study.

Samples of 2 mL venous blood were taken from each subjects and the serum was separated and frozen at -20°C. Measles specific antibody titer (IgG) was estimated using MEASLES IgG ELISA DSL-05-10-MEG, DSL (Italy). Sensitivity = 96.4% and Specificity = 100%. Using this kit, IgG titers >1.1 IU mL⁻¹ were considered as positive. Amounts between 0.9 and 1.1 IU mL⁻¹ were considered as borderline and IgG titers less than 0.9 were considered as negative. Data analysis was carried out with spss 11.5 using chi square and Fisher tests.

RESULTS

Of the 1779 students selected, 114 were excluded from the study due to incomplete measles two dose vaccination, inadequate amount of blood, hemolysis of blood, or absence of a questionnaire. Thus 1665 subjects were enrolled, 945 (57%) of whom were girls and 720 (43%) were boys. mean age was 9.17±1.53 years.

Anti measles antibody titer was negative in 385 subjects, borderline in 82 subjects and positive in 1198 subjects. Borderline cases were also considered as seronegative because of their immunity status are not acceptable. Thus overall seronegative cases were 385+82 = 467 (= 28% of all cases) subjects and seropositive cases were 1198 (72% of all cases) (Table 1).

Three hundred and eighty five negative cases consist of 210 girls and 175 boys. Eighty two borderline cases consist of 51 girls and 31 boys. Also from 1198 positive cases, 684 cases were girls and 514 cases were boys. From 945 girls, 261 cases (28%) were seronegative and from 720 boys, 206 cases (29%) were seronegative. There was no statistically significant difference in seronegativity proportion with regard to gender (29%, in males and 27%, in females (p = 0.404).

The percentage of seronegative subjects in 6, 7, 8, 9, 10, 11 and 12 years groups were 23.5, 26, 29, 27, 29, 29 and 22, respectively. There were no statistically significant differences in seronegativity proportion with regard to age (p = 0.775) (Table 2).

The proportion of seronegativity was highest (31%) in the western educational sector of Tehran and in the northern, eastern and southern educational sectors were 28, 28 and 29, respectively. These differences were not statistically significant (p = 0.450) (Table 3).

From 1298 subjects that had a history of 2-dose measles vaccination in 9 and 15 months of age, 356 (27.5%) were seronegative and 942 (72.5%) were seropositive. From 376 subjects that received 2-dose vaccine with 1 dose of measles vaccine at 9 months of age and 1 dose of MMR after 15 months of age, 111 (30%) were seronegative and 256 (70%) were seropositive. There was no statistically significant difference in seronegativity proportion with regard to kind of vaccination (p = 0.296) (Table 4).

Table 1: Prevalence of antimeasles virus IgG by sex in Tehran

Serology	Girls		Boys		Total	
	No	%	No	%	No	%
Negative	220	23	175	24	395	24
Borderline	53	6	31	4.5	84	5
Positive	672	71	514	71.5	1186	71
Total	945	100	720	100	1665	100

Table 2: Prevalence of antimeasles virus antibodies by age group in Tehran

Serology age group	Negative		Borderline		Positive		Total	
	No	%	No	%	No	%	No	%
6	6	17.5	2	6	26	76.5	34	100
7	61	23.5	7	3	192	73.5	260	100
8	74	25	13	4	210	71	297	100
9	71	21	19	6	243	73	333	100
10	79	24	17	5	233	71	329	100
11	25	38.5	5	7.5	35	54	65	100

Table 3: Prevalence of antimeasles virus antibodies by geographical group in Tehran

Serology	Northern		Western		Eastern		Southern	
	No	%	No	%	No	%	No	%
Negative	102	20	103	26	77	24	103	24
Borderline	27	5	21	5	13	4	21	5
Positive	385	75	274	69	229	72	310	71
Total	514	100	398	100	319	100	434	100

Northern part consists of 1,3,4,7 and 8 educational sectors of Tehran, Western part consist of 2,5,6,9 and 18 educational sectors of Tehran, Eastern part consists of 12,13,14 and 15 educational sectors of Tehran, Southern part consists of 10,11,16,17 and 19 educational sectors of Tehran

Table 4: Prevalence of antimeasles virus antibodies by history of vaccination against rubella in Tehran

Serology	Routine vaccination		Measles vaccine+ MMR after 15 month		Total	
	No	%	No	%	No	%
Negative	356	27.5	111	30	467	28
Positive	942	72.5	256	67	1198	72
Total	1298	100	367	100	1665	100

DISCUSSION

In this study 1665 school age children (6-12 years old primary school students) from the 19 educational sectors of Tehran were enrolled, 945 (57%) of whom were girls and 720 (43%) were boys.

Anti measles antibody titer was positive in 1198 subjects, borderline in 82 subjects and negative in 385 subjects. Thus there were 467 (28%) of study subjects susceptible to disease. All of subjects have a history of twice vaccination against measles, on 9 and 15 month old. In a study that was performed in mashhad City in 2002 the measles IgG seroprevalence, reached almost 70%. They collected 172 students from 7 parts of Mashhad and tested for measles specific IgG antibodies by ELISA method. The age of these subjects was 14-19 years and divided in 6 groups. They found that 50 (29%) of the subjects were seronegative (Sarvghad, 2003). In another study that performed in less than 5 years old children, of them 668 subjects from western azarbaijan, 507 subjects from southern part of Tehran and 887 subjects from boushehr, showed that immunity ratio in western azarbaijan was 80%, in southern part of Tehran was 79% and in boushehr was 78% (Saidi *et al.*, 1984). The first reason of these differences is difference of age between these studies. The second reason is that answer to vaccine and antibody decay phenomenon are depend on several factors that maybe different in some areas.

Secondary vaccine failure in present study was 13-20% that increase with age; and in other studies has reported between 2 and 20% (Saidi *et al.*, 1984; Mathias, 1989; Zhonghua, 1987; Xiang and Chen, 1983; Krugman, 1983) and alter in different populations (Christenson and Bottiger, 1994; Markowitz, 1990; Olsha, 1994).

In present study from 945 girls that enrolled to study, 684 cases (72%) were seropositive and 261 cases (28%) were seronegative and from 720 boys that enrolled to study, 514 cases (71%) were seropositive and 206 cases (29%) were seronegative (Table 1). There was no statistically significant difference in seronegativity proportion with regard to gender 206 of 720, 29%, in males and 261 of 945, 28%, in females ($p = 0.404$). Our result is comparable with similar studies. In study that was performed in 1988, in 3 level of primary school, secondary, School and high school, girls were more susceptible to measles; but there was not a statistical significant difference between two sexes (Mokhtari Azad *et al.*, 1993). Males were significantly more seronegative than females ($p < 0.0001$) (Sarvghad, 2003).

In a study that was performed to evaluate the immune status to measles among 375 school children in south of Tehran, seronegativity was 20%. Similar to present study they did not detect a significant difference and also between two gender groups, but they detect a significant difference between age groups and lower age groups were more susceptible to measles. The reasons for this difference in the children in parts of Iran are unknown and need further study to clarify (Mokhtari Azad *et al.*, 1993).

The percentage of seronegative subjects in 6, 7, 8, 9, 10, 11, 12 years groups were 23.5, 26, 29, 27, 29, 29 and 22, respectively. It seems that the age group of 8-10 years old, are the most susceptible group to

measles; but there were no statistically significant differences in seronegativity proportion with regard to age ($p = 0.775$) (Table 2).

In Mashhad study, the most cases of seronegative subjects found in age 14-17. Thus, the correlation of the age and negative results in that study was significant ($p = 0.0003$) (Sarvghad, 2003). In comparison with present study, we do not detect a significant difference between our age groups probably because that the age of present groups were very close and so did not differ.

As regards vaccination coverage is higher than of 1988, all of present subjects have a history of twice vaccination against measles, but in previous study subjects had a history of one dose vaccination; may be because of absence of disease in community, the possibility of contact with virus and then increase in antibody titer has reduced and gradually in primary school students, immunity level due to vaccination has decreased (Mokhtari Azad *et al.*, 1993; Nkowane *et al.*, 1987). But in 1988, because of lower vaccination coverage, measles has existed in the community and causes relative increase in antibody titer. Although in present study, subjects had a history of twice vaccination against measles, on 9 and 15 month old, the maximum positive antibody titer was 76.5% in 6 years old group, which can be due to:

- Primary failure
- Secondary failure

Primary failure of vaccine: Vaccination program is appropriate if by use of it, after 12 month of age, more than 90 to 98% of susceptible individuals, become immune (Mokhtari Azad *et al.*, 1993), that show us between 2 to 10 percent of population would be susceptible (AAPC, 1990; Chen *et al.*, 1989; CDC, 1989).

Primary failure of vaccine has different causes include age of subjects in vaccination time, coverage of vaccination programs, vaccines quality, planning of immunization programs and specifications and type of vaccines (Cvjetanovic *et al.*, 1998). vaccination with the high-dose Edmonston-Zagreb or the AIK-C strain at 4-5 months was as good as vaccination with the AIK-C strain at 8-10 months and better than vaccination with the standard dose Schwarz strain at 8-10 months (Tidjani *et al.*, 1989; Morkowitz, 1990).

Method of prescription, maintenance quality of vaccines, Method and amount of injection, exposure of light and temperature, vaccine usage in accompany with immunoglobulin and existence of maternal antibodies (Shelton *et al.*, 1978; Wesley *et al.*, 1978) and nutritional status of individual (Wesley *et al.*, 1978), existence

of an acute disease in the time of vaccination (Halsey *et al.*, 1985; Krober *et al.*, 1991; Ndikuyeze *et al.*, 1988), vaccine usage in accompany with immunoglobulin (Krugman, 1971), race, environmental factors and hygiene status (Black *et al.*, 1986; Nieburg *et al.*, 1986) sex (Bromberg *et al.*, 1994) and immunity system status (Dai *et al.*, 1991; Markowitz *et al.*, 1992) are from Primary failure factors, that consist some of our susceptible individuals.

Secondary failure of vaccine: is the losing of vaccine induced immunity. Some of individuals in our study maybe lose their immunity to measles because of secondary vaccine failure.

In present study after the age of 10 years antibody titer rise and in 12 years old group, seropositivity receives to 78% and only 22% of individuals were susceptible to disease. In a similar study that performed at 1988 in southern part of Tehran, the results were similar to our study and the percentage of susceptible subjects in primary school students (9%) was higher than in guidance School students (4%) and in high school students (4%) (Mokhtari Azad *et al.*, 1993).

Increasing of antibody titer after the age of 10 years, in these studies is maybe due to contact with measles disease in the community and contact with wild virus (boosting phenomenon). In another study that performed in Ethiopia the seronegative proportion ($<100 \text{ IU L}^{-1}$) was 20% (95% CI: 16-25) in children 9-59 months old, declining to 9% (7-12) in 5-9 year olds and 5% (4-7) in 10-14 year olds (Enquselassie *et al.*, 2003). In other studies that have performed in china, secondary vaccine failure after the age of 8 years, receives to 13%. They haven't contact with wild virus (Xiang and Chen, 1983).

Antimeasles antibody titer in four geographical area of Tehran was different. The proportion of seronegativity was highest (31%) in the western educational sector of Tehran and in the northern, eastern and southern educational sectors were (28%), (28%) and (29%) respectively. These differences were not statistically significant ($p=0.450$) (Table 3). In other studies that have performed in less than 5 years old children, proportion of immunity have been 80% in azarbaijan, 79% in southern part of Tehran and 78% in boushehr (Saidi *et al.*, 1984). In another study that performed in southern part of Tehran in 1988, from 165 student, 120 had positive titer (73%) that is similar to our result (Mokhtari Azad *et al.*, 1993).

Comparing routine 2-dose measles vaccination in 9 and 15 months of age, with 2-dose vaccine with 1 dose of measles vaccine at 9 months of age and 1 dose of MMR after 15 months of age, showed that from 1298 subjects that had a history of 2-dose measles vaccination in 9 and

15 months of age, 356 (27.5%) were seronegative and 942 (72.5%) were seropositive and from 376 subjects that received 2-dose vaccine with 1 dose of measles vaccine at 9 months of age and 1 dose of MMR after 15 months of age, 111 (30%) were seronegative and 256 (70%) were seropositive. There was no statistically significant difference in seronegativity proportion with regard to kind of vaccination ($p = 0.296$) (Table 4).

According to these results and other studies that previously have performed in Iran, in 6-19 years old groups that have a history of one dose vaccination in 6 months of age, 80% of individuals had positive antibody titer. In comparing with studies that have performed in the United States and England, reported that 6-17 years after vaccination, more than 90% of individuals that had a history of measles vaccination in 1 year old, had positive antibody titer, indicate that vaccination before the age of one year, not only can induce primary vaccine failure, but also reduce vaccine stability. On the other hand serologic studies indicate that vaccine induced immunity is persistent and has been reported between 8 to 23 years (Nkowane *et al.*, 1987). These studies indicate that if we perform vaccination in appropriate time and reduce primary vaccine failure, effectiveness and persistence of vaccine will be increased.

Recent studies from Croatian population showed that vaccination coverage higher than the reported 90-94% should be attained if one is to expect measles elimination (Borcic *et al.*, 2003). The immunization campaign prevented 90-95% of predicted cases. The campaign was appropriately targeted at children aged 10 years and under (Hellenbrand *et al.*, 2003; Mansoor *et al.*, 1998).

CONCLUSION

In comparison of our study with previous 1988 study, susceptible age group decrease from 10-12 years to 8-10 years old, that maybe due to higher vaccination coverage and lesser virus contact in community; that indicate the necessity of revaccination in children under 10 years old. For availability, the best time is the time of school beginning (6 years of age).

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REFERENCES

- American Academy of Pediatrics Committee on Infectious Diseases (AAPC), 1990. Measles: Reassessment of the current immunization policy. *Pediatrics*, 85: 714.
- Anderson, R.M. and R.M. May, 1985. Age-related changes in the rate of disease transmission: Implications for the design of vaccination programmes. *J. Hyg.*, pp: 365-436.
- Anonymous, 1994. Expanded Programme on Immunization: Measles, WHO. *Weekly Epidemiol. Rec.* 1995, 70: 284-288.
- Anonymous, 2000. Measles surveillance in the European region, WHO, *EURO Measles Q.* (2003), pp: 1-4.
- Black, F.L., L.L. Berman, J.M. Borgono, R.A. Capper, A.A. Carvalho, C. Collins, O. Glover, Z. Hijazi, D.L. Jacobson and Y.L. Lee, 1986. Geographic variation in infant loss of maternal measles antibody and in prevalence of rubella antibody. *Am. J. Epidemiol.*, 124: 442-452.
- Black, F.L., 1989. Measles. In: Evans, A.S., 1989. *Viral Infection of Humans: Epidemiology and Control*. 3rd Edn., Plenum Medical Book Company, New York, pp: 451-465.
- Borcic, B., R. Mazuran and B. Kaic, 2003. Immunity to measles in the croatian population. *Eur. J. Epidemiol.*, 18: 1079-1083.
- Bromberg, K., B. Shah, M. Clark-Golden, H. Light, L. Marcellino, M. Rivera, P.W. Li, D. Erdman, J. Heath and W.J. Bellini, 1994. Maternal immunity to measles and infant immunity at less than twelve months of age relative to maternal place of birth. *J. Pediatr.*, 125: 579-581.
- Caceres, V.M., P.M. Strebel and R.W. Sutter, 2000. Factors determining prevalence of maternal antibody to measles virus throughout infancy: A review. *Clin. Infect. Dis.*, 31: 110-119.
- CDC, 1989. Centers for Disease Control. Measles prevention: Supplementary statement. *MMWR Morb Mortal Weekly Rep.*, 38: 11-14.
- Chen, R.T., G.M. Goldbaum, S.G. Wassilak, L.E. Markowitz and W.A. Orenstein, 1989. An explosive point-source measles outbreak in a highly vaccinated population. Modes of transmission and risk factors for disease. *Am. J. Epidemiol.*, 129: 173-182.

- Christenson, B. and M. Bottiger, 1994. Measles antibody: Comparison of long-term vaccination titres, early vaccination titres and naturally acquired immunity to and booster effects on the measles virus. *Vaccine*, 12: 129-133.
- Cutts, F.T., R.H. Henderson, C.J. Clements, R.T. Chen and P.A. Patriarca, 1991. Principles of measles control. *Bull World Health Organ*, 69: 1-7.
- Cvjetanovic, B., B. Grab and H. Dixon, 1998. Epidemiological models of poliomyelitis and measles and their application in the planning of immunization programmes. *Bul. WHO*, 60: 405-422.
- Dai, B., Z.H. Chen, Q.C. Liu, T. Wu, C.Y. Guo, X.Z. Wang, H.H. Fang and Y.Z. Xiang, 1991. Duration of immunity following immunization with live measles vaccine: 15 years of observation in Zhejiang Province, China. *Bull. World Health Organ.*, 69: 415-423.
- De Serres, G., J. Sciberras, M. Naus, N. Boulianne, B. Duval and L. Rochette, 1999. Protection after two doses of measles vaccine is independent of interval between doses. *J. Infect. Dis.*, 180: 187-190.
- Enquselassie, F., W. Ayele, A. Dejene, T. Messele, A. Abebe, F.T. Cutts and D.J. Nokes, 2003. Seroepidemiology of Measles in Addis Ababa, Ethiopia: Implications for control through vaccination. *Epidemiol. Infect.*, 130: 507-519.
- Halsey, N.A., R. Boulous, F. Mode, J. Andre, L. Bowman, R.G. Yaeger, S. Toureau, J. Rohde and C. Boulous, 1985. Response to measles vaccine in Haitian infants 6 to 12 months old. Influence of maternal antibodies, malnutrition and concurrent illnesses. *N. Eng. J. Med.*, 313: 544-549.
- Hellenbrand, W., A. Siedler, A. Tischer, C. Meyer, S. Reiter, G. Rasch, D. Teichmann, S. Santibanez, D. Altmann, H. Claus and M. Kramer, 2003. Progress toward measles elimination in Germany. *J. Infect. Dis.*, 187 Suppl 1: S208-216.
- Hutchins, S.S., A.L. Baughman, M. Orr, C. Haley and S. Hadler, 2004. Vaccination levels associated with lack of measles transmission among preschool-aged population in the united states. *J. infect. Dis.*, 189 Suppl: 108-115.
- Krober, M.S., C.E. Stracener and J.W. Bass, 1991. Decreased measles antibody response after measles-mumps-rubella vaccine in infants with colds. *JAMA.*, 265: 2095-2096.
- Krugman, S., 1983. Further-attenuated measles vaccine: Characteristics and use. *Rev. Infect. Dis.*, 5: 477-481.
- Krugman, S., 1971. Present status of measles and rubella immunization in the United States: A medical progress report. *J. Pediatr.*, 78: 1-16.
- Lee, M.S., P.M. Mendelman, C. Sangli, I. Cho, S.L. Mathie and M.J. August, 2001. Half-life of human parainfluenza virus type 3 (hPIV3) maternal antibody and cumulative proportion of hPIV3 infection in young infants. *J. Infect. Dis.*, 183: 1281-1284.
- Mansoor, O., T. Blakely, M. Baker, M. Tobias and A. Bloomfield, 1998. A measles epidemic controlled by immunisation. *N Z Med J.*, 111: 467-471.
- Markowitz, L.E., 1990. Preblud SR, Fine PE, Orenstein WA. Duration of live measles vaccine-induced immunity. *Pediatr. Infect. Dis. J.*, 9: 101-110.
- Markowitz, L.E., P. Albrecht, W.A. Orenstein, S.M. Lett, T.J. Pugliese and D. Farrell, 1992. Persistence of measles antibody after revaccination. *J. Infect. Dis.*, 166: 205-208.
- Mathias, R., 1989. The role of secondary vaccine failures in measles outbreak. *Am. J. Public Health*, 79: 475-478.
- Mokhtari Azad, T.M. Gharib, M. Mahmoudi, A. Moosavi, Z. Seadatmand and R. Nategh, 1993. Determination of immune status to Measles among school-children in south of Tehran Iranian *J. Public Health*, 4-1: 39-52.
- Moradi, A., M. Salehi and F. Rakhshani, 2001. Seroepidemiological study of measles among 25-60 months children in Iranshahr district tabib-e-shargh *J. zahedan Univ. Med. Sci. Health Serv.*, 3: 142-137.
- Morkowitz, L.E., 1990. Immunization of six-months old infants to different doses of Edmonston-Zagreb and Schwartz measles vaccine. *N. Eng. J. Med.*, 322: 580-587.
- Nakayama, T., J. Zhou and M. Fujino, 2003. Current status of measles in Japan. *J. Infect. Chemother.*, 9: 1-7.
- Ndikuyeze, A., A. Munoz, J. Stewart, L. Modlin, D. Heymann, K.L. Herrmann and B.F. Polk, 1988. Immunogenicity and safety of measles vaccine in ill African children. *Int. J. Epidemiol.*, 17: 448-455.
- Nieburg, P. and M.J. Dibley, 1986. Risk factors for fatal measles infections. *Int. J. Epidemiol.*, 15: 309-311.
- Nkowane, B.M., S.W. Bart, W.A. Orenstein and M. Baltier, 1987. Measles outbreak in a vaccinated school population: Epidemiology, chains of transmission and the role of vaccine failures. *Am. J. Public Health*, 77: 434-438.
- Olsha, M., 1994. Measles immunity in Israel young adults. *Isr. J. Med. Sci.*, 30: 596-599.
- Saidi, S., H. Malekafzali, M.A. Barzegar and M. Shafiyi, 1984. Effect of health system providers in measles immunization program. *J. Health in Iran*, 1: 1-10.

- Sarvghad, M.R., 2003. The serologic evaluation of Measles IgG antibody in vaccinated persons in 2002 in Mashhad. *Med. J. Mashhad Univ. Med. Sci.*, 81: 33-38.
- Shelton, J.D., J.E. Jacobson, W.A. Orenstein, K.F. Schulz and H.D. Jr. Donnell, 1978. Measles vaccine efficacy: Influence of age at vaccination vs. duration of time since vaccination. *Pediatrics*, 62: 961-964.
- Slifka, M.K. and R. Ahmed, 1996. Long-term humoral immunity against viruses: Revisiting the issue of plasma cell longevity. *Trends Microbiol.*, 4: 394-400.
- Thomas, A., D. Xu, K. Wooten, B. Morrow and S. Redd, 1999. Timing and effectiveness of requirements for a second dose of measles vaccine. *Pediatr. Infect. Dis. J.*, 18: 266-270.
- Tidjani, O., B. Grunitsky, N. Guerin, D. Levy-Bruhl, N. Lecam, C. Xuereff and K. Tatagan, 1989. Serological effects of Edmonston-Zagreb, Schwarz and AIK-C measles vaccine strains given at ages 4-5 or 8-10. *Lancet*, 2: 1357-1360.
- Ward, B.J., N. Boulianne, S. Ratnam, M.C. Guiot, M. Couillard and G. De Serres, 1995. Cellular immunity in measles vaccine failure: Demonstration of measles-specific lymphoproliferative responses despite limited serum antibody production after revaccination. *J. Infect. Dis.*, 172: 1591-1595.
- Wesley, A., H.M. Coovadia and L. Henderson, 1978. Immunological recovery after measles. *Clin. Exp. Immunol.*, 32: 540-544.
- Whittle, H.C., P. Aaby, B. Samb, H. Jensen, J. Bennett and F. Simondon, 1999. Effect of subclinical infection on maintaining immunity against measles in vaccinated children in West Africa. *Lancet*, 353: 98-102.
- Xiang, JZ. and Z.H. Chen, 1983. Measles vaccine in the People's Republic of China. *Rev. Infect. Dis.*, 5: 506-510.
- Zhonghua, 1987. measles vaccine study group. Epidemiologic examination of the duration of immunity of measles vaccine. *Zhonghua Yi Xue Za Zhi*, 67: 19-22.