

- 5 Shah RS, Rajalakshmi R. Vitamin A status of the newborn in relation to gestational age, body weight, and maternal nutritional status. *Am J Clin Nutr* 1984;**40**:794–800.
- 6 Ganguly C, Mukherjee KL. Relationship between maternal serum vitamin A and vitamin A status of the corresponding fetuses. *J Trop Pediatr* 1986;**32**:287–9.
- 7 Saha K, Garg M. Studies on colostrum nutrient and immunologic factors. *Nutr Foundation India Bull* 1991;**12**:6.
- 8 DeSole G, Belay Y, Zegeye B. Vitamin A deficiency in southern Ethiopia. *Am J Clin Nutr* 1987;**45**:780–4.
- 9 Hellen Keller International and Institute of Public Health. Bangladesh nutritional blindness study: Risk factors. Dhaka: Helen Keller International Report, 1986.
- 10 Anderson RK, Robinson WD, Golvo J, Payne GC. Nutritional status during pregnancy and after delivery of a group of women in Mexico city. *J Am Diet Assoc* 1946;**22**:588–93.
- 11 Arroyare G. Nutrition in pregnancy studies in Central America and Panama. *Arch Hatinorum Nutr* 1976;**26**:129–57.
- 12 Rebecca JS, Mohammad Haki, Miller KW, Kethleen MS, Dawiesah S, Jean Pierre Hatricht, Michael JD. High dose vitamin A supplementation of breast-feeding Indonesian mothers: Effects on vitamin A status of mother and infant. *Am J Nutr* 1992;**123**:666–75.
- 13 Sommer A. Field guide to the detection and control of xerophthalmia. Geneva: World Health Organization, 1982.
- 14 Tarwojo I, Sommer A, Soegiharto T, Susanto D, Muhilahl H. Dietary practices and xerophthalmia among Indonesian children. *Am J Clin Nutr* 1982;**35**:574–81.
- 15 Sampathkumar V, Abel R. Xerophthalmia in rural South Indian children. *Indian Paediatr* 1993;**30**:246–8.
- 16 Gopalan C. Combating vitamin deficiency through dietary improvement. *Nutr Foundation India Bull* 1992; Special publication services 6:12.

Increasing toxoplasma seropositivity in women with bad obstetric history and in newborns

P. SHARMA, I. GUPTA, N. K. GANGULY,
R. C. MAHAJAN, N. MALLA

ABSTRACT

Background. Most infants with congenital *Toxoplasma gondii* infection have no symptoms at birth but few may develop retinal diseases or neurological abnormalities later in life. The presence of significant titres of antitoxoplasma antibodies in women in the reproductive age group indirectly indicates that *Toxoplasma gondii* is the cause of such congenital abnormalities and also sporadic abortions in some women.

Methods. We did a retrospective analysis of antitoxoplasma antibodies detected by indirect haemagglutination assay, in women with bad obstetrical history and in newborns clinically suspected of congenital toxoplasmosis during 1981–91.

Results. A significant increase in seropositivity in women and newborns was seen during 1989–91 as compared to 1981–88. More seropositive patients were recorded between April–June and October–December. However, no significant correlation could be observed between rising incidence of seropositivity and the seasonal distribution or age of women.

Conclusion. Epidemiological studies are required to ascertain the reason for the increasing trend of toxoplasma seropositivity and to suggest appropriate control strategies as it is possible to prevent congenital infection.

Natl Med J India 1997;**10**:65–6

INTRODUCTION

Toxoplasma gondii (*T. gondii*), an obligate intracellular proto-

Postgraduate Institute of Medical Education and Research,
Chandigarh 160012, India

P. SHARMA, R. C. MAHAJAN, N. MALLA
Department of Parasitology

I. GUPTA Department of Gynaecology and Obstetrics
N. K. GANGULY Department of Experimental Medicine
and Biotechnology

Correspondence to N. MALLA

© The National Medical Journal of India 1997

zoon that is ubiquitous in nature, is an important cause of infection and disease in humans and domestic animals. Human infection results from ingestion of tissue cysts or oocysts and the parasite has been shown to invade the placenta and the foetus resulting in abortion, stillbirth or congenital infection.^{1–8} The prevalence of toxoplasmosis in women with a bad obstetrical history (BOH) is known to be significantly higher than in those without it.^{3,8,9} Toxoplasma infection may be acute or chronic, symptomatic or asymptomatic. Acute infection poses the greatest hazard to the immunodeficient patient and the foetus *in utero*. The infant infected *in utero* with or without signs of infection at birth may develop serious sequelae such as impaired vision, neurological disorders and sensorineural hearing loss. The diagnosis is made on the basis of presence of specific antibodies as methods for isolation of the organism are less sensitive.

We did a prospective study during 1990–91 to determine the seropositivity rate of women with a BOH and of newborns clinically suspected to have toxoplasmosis. These data were compared with a retrospective analysis of the data available for 1981–89 in similar groups.

PATIENTS AND METHODS

A total of 7222 patients (5916 women with BOH and 1306 newborns, clinically suspected to have toxoplasmosis) who attended the Obstetrics and Gynaecology Department of the Nehru Hospital, Postgraduate Institute of Medical Education and Research, Chandigarh, between 1981 and 1991 were studied. Both the percentage seropositivity and the number of seropositive cases seen each year were taken into account to determine the trends of toxoplasmosis. Five ml of clotted blood was obtained from each patient and the serum was stored at –20 °C till used.

Preparation of antigen

The RH strain of *T. gondii* being maintained in Swiss albino mice was used for preparing the antigen.¹⁰ Tachyzoites were harvested from the peritoneal cavity of Swiss albino mice, infected 3 days earlier, and filtered on 3 µm polycarbonate membrane (Mucleopore, CA). These were then washed twice with 10 mM PBS (pH 7.2). The pellet was resuspended in distilled water at a concentration of 6×10⁸ tachyzoites/ml. The parasites were disrupted by sonication (×1 minute) in an MSE ultrasonic disintegrator (SONIPREP) and then centrifuged at 10 000 g for 60 minutes. The supernatant used as soluble antigen was stored at –20 °C in an aliquot till used.

Indirect haemagglutination assay (IHA)

All the serum samples were analysed for specific antitoxoplasma

antibodies by indirect haemagglutination assay as described by Lewis and Kessel.¹¹ The optimal dilution of every batch of antigen used was determined by conventional block titrations done against standard positive and negative control serum samples. The patterns of agglutination were read after 2 hours of incubation at room temperature followed by overnight incubation at 4 °C. Known positive and negative serum controls were tested simultaneously with every batch. Serum samples with a titre of more than 1:128 were considered positive.⁸ The results were analysed by the Student's t-test.

RESULTS

A steadily rising trend in percentage seropositivity and total number of seropositive cases (Table I) was observed during 1989–91, there being a three-fold increase in 1989, four-fold in 1990 and seven-fold in 1991. This rise in seropositivity was statistically significant ($p < 0.001$) and was seen both in women with BOH and in newborns suspected to have toxoplasmosis. The mean age of women with BOH was 25.5 years (range 18–37 years) and the majority (49.6%) were in the 23–27 years age group (Fig. 1). A bimodal peak of seasonal distribution of cases was observed with more seropositive patients during April–June and October–December (Fig. 2). There was no significant correlation of the rise in seropositivity with either the age of women with BOH or seasonal distribution of cases.

DISCUSSION

The presence of antibodies to *T. gondii* suggests that the person has either been exposed to the infection in the past or has been infected recently.^{2,3} It has been hypothesized that when these organisms are present in the genital tract especially near the implantation site, abortion, stillbirth or congenital infection may occur.^{3,4} The presence of significant titres of antitoxoplasma antibodies are considered an indirect evidence of the organism being the cause of BOH in women of the reproductive age group. Isolation of the parasite from the endometrium, placenta or products of conception is usually difficult.^{3,5,12–14}

Our data indicate a rising trend in seropositivity to *T. gondii* in women with BOH and newborns who were clinically suspected to have toxoplasmosis. The reason for this rising trend is not clear. Chandigarh is a city of white collar people and the living conditions are good and hygienic. There are only a few slums with stray animals. There has been a change in the eating habits of people, with meat being eaten more often. This might be one of the contributing factors. Ours being a referral hospital, patients from different socio-economic strata, religions and faith attend it from practically the entire northern region of the country. This report indicates that there is a need to investigate the epidemiological

TABLE I. Toxoplasma seropositivity from 1981 to 1991

Year	n	Total seropositive (%)	Women with BOH	Neonates
1981	563	13 (0.02)	12	1
1982	450	11 (0.02)	10	1
1983	553	11 (0.02)	11	0
1984	545	11 (0.02)	9	2
1985	525	11 (0.02)	9	2
1986	534	13 (0.02)	9	4
1987	704	12 (0.02)	9	3
1988	829	9 (0.01)	8	1
1989	800	49 (0.06)	42	7
1990	860	68 (0.08)	55	13
1991	859	127 (0.15)	107	20

BOH bad obstetric history

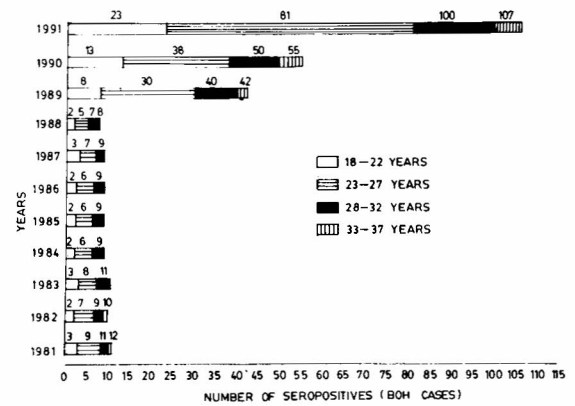


Fig 1. Seropositive women with bad obstetrical history (BOH) during 1981–91, classified by age

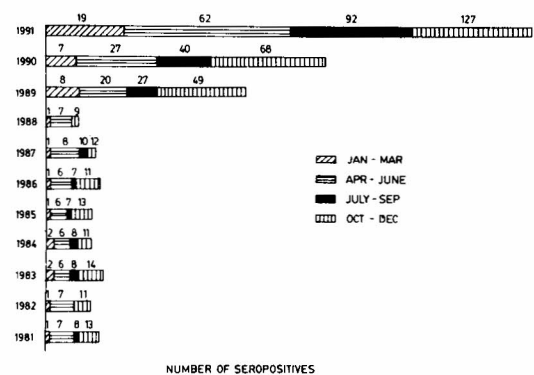


Fig 2. Seasonal distribution of seropositive cases during 1981–91

factors causing this rising trend in toxoplasma seropositivity in this region. With an increasing number of acquired immunodeficiency syndrome (AIDS) patients, this infection may produce acute life-threatening conditions such as encephalitis, myocarditis or pneumonitis.

REFERENCES

- Remington JS, Mcleod R, Desmonts G. Toxoplasmosis. In: Remington JS, Klein JO (eds). *Infectious diseases of the fetus and newborn infant*. Philadelphia: WB Saunders, 1995:140–267.
- Frenkel JK. Toxoplasmosis. *Paediatr Clin North Am* 1985;32:917–32.
- Remington JS, Desmonts G. Toxoplasmosis. In: Remington JS, Klein JO (eds). *Infectious diseases of the fetus and newborn infant*. Philadelphia: WB Saunders, 1990:89–195.
- Sabin AB, Eichenwald H, Feldman HA, Jacobs L. Present status of clinical manifestations of toxoplasmosis in man: Indications and provisions for routine serologic diagnosis. *JAMA* 1952;150:1063–7.
- Remington JS, Newell JW, Cavanaugh E. Spontaneous abortion and chronic toxoplasmosis: Report of a case with isolation of the parasite. *Obstet Gynaecol* 1964;24:25–31.
- Hingorani V, Prakash O, Chowdhry P, Kamalam TS. Toxoplasmosis—Abortions and stillbirths. *Indian J Med Res* 1970;58:967–74.
- Desmonts G, Couvreur J. Toxoplasmosis in pregnancy and its transmission to the fetus. *Bull NY Acad Med* 1974;50:146–59.
- Mahajan RC, Gupta I, Chhabra MB, Gupta AN, Devi PK, Ganguly NK. Toxoplasmosis—Its role in abortion. *Indian J Med Res* 1976;64:797–800.
- Kimball AC, Kean BH, Fuchs F. The role of toxoplasmosis in abortion. *Am J Obstet Gynecol* 1971;111:219–26.
- Mahajan RC, Chitkara NL, Jolly JG. Serological survey of toxoplasma antibodies in Chandigarh area (Northern India). *Indian J Med Res* 1974;62:1–6.
- Lewis WP, Kessel J. Haemagglutination test in the diagnosis of toxoplasmosis. *Arch Ophthalmol* 1961;66:471–6.
- Mahajan RC, Chhabra MB, Ganguly NK, Singh RP. Fluorescent antibody test in immunodiagnosis of toxoplasmosis. *Indian J Med Res* 1977;66:29.
- Chhabra MB, Mahajan RC, Mahajan MK. Isolation of *Toxoplasma gondii* from suspected human cases. *Indian J Med Res* 1979;69:746–51.
- Thokar MA, Malla N, Watal C. Serological study of patients clinically suspected to have toxoplasmosis in Kashmir. *Indian J Med Res* 1988;88:29–34.