



STUDY OF INCIDENCE AND PREVALENCE OF RETINOPATHY OF PREMATUREITY IN TERTIARY HOSPITAL IN GARHWAL REGION.

Medical Science

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ABSTRACT

Purpose: To study the incidence and risk factors of retinopathy of prematurity (ROP).

Methods: A retrospective review of premature babies admitted in NICU over a period of one year, with a gestational age of 32 weeks or less at birth and a birth weight of 1500 g or less was made. Sick infants were included even if they were older and heavier. ROP screening was done in 200 cases from the fourth postnatal week and was followed up.

Results: The incidence of ROP was 19.5%. Most common maternal risk factor was pregnancy induced hypertension (17.9%). Low birth weight (LBW) and respiratory distress syndrome (RDS) (89.7%) were the most common risk factors in infants.

Conclusion: In our study the most common risk factors were LBW and RDS.

KEYWORDS

INTRODUCTION:

Retinopathy of prematurity (ROP), characterized by abnormal development of retinal vasculature, is an important and preventable cause of childhood blindness¹. It is estimated that ROP is the cause of 50,000 cases of childhood blindness in the world every year². According to the available data, there has been an increase in the incidence of ROP in developing countries, cause of which was considered to be higher survival rates in extremely premature babies³. A high concentration of supplemental oxygen therapy was previously thought to be the major risk factor for development of ROP^{4,5}. However, there are many reports that ROP has been observed in patients without oxygen therapy⁶. Low birth weight (BW) and low gestational age (GA) are well-known risk factors for ROP^{7,8}.

The criteria for ROP screening have been well determined in industrialized countries, but the available data suggests that blindness from ROP varies from region to region³. According to many studies performed in various parts of the world, it has been recognized that ROP screening in each region has its own characteristics and a single screening program cannot suit all regions⁹.

PURPOSE:

To study the incidence and risk factors of retinopathy of prematurity (ROP).

METHODS:

This is a retrospective cross sectional review which included premature babies who were referred for ROP screening, admitted in NICU over a period of one year, with a gestational age of 32 weeks or less at birth and a birth weight of 1500 g or less. Babies 32 weeks and 1500 grams were screened if the babies were sick as per paediatricians advice. ROP screening was done from the fourth postnatal week and within four weeks if the baby was ≤ 30 weeks and was followed up accordingly.

Data on birth weight, gestational age, postconceptional age was noted. Maternal risk factors like (pregnancy induced hypertension, gestational diabetes) and baby risk factors like (respiratory distress syndrome, sepsis) were also noted in the history. Babies in whom media opacity (not related to ROP) precluding fundus visualization disease, loss to follow-up were excluded from the study.

The pupils were dilated by application of tropicamide 0.5% and phenylephrine 1%, and the patients were examined by a single retina specialist using a binocular indirect ophthalmoscope with 20 D lens. A sterile lid speculum and a scleral depressor were routinely used.

ROP was classified according to the international classification of ROP and follow-up schedules were designed in accordance with suggestions of the American Academy of Pediatrics, American Academy of Ophthalmology and American Association for Pediatric Ophthalmology and Strabismus¹⁰. Treatment plans were based on recommendations of the Early Treatment of ROP cooperative group¹¹.

RESULTS:

During the period of 1 year, 200 babies were screened. Male babies screened were 110, and female babies were 90. Gestational age (GA) of babies ranged from 26 weeks-38 weeks (mean GA-33wks). Birth weight (BW) of the babies range from 1kg-3.5kg (mean BW-2.25kg). Incidence of ROP was 19.5% (39 babies) in our study. Different stages of ROP seen are as shown in the table 1. In the babies diagnosed to have ROP the gestational age ranged from 26 weeks- 37 weeks (with 30% being more than 32 weeks). Birth weight ranged from 1kg-2kg (with 25.6% being more than 1.5kg). Most common maternal risk factor for ROP was pregnancy induced hypertension in 7 patients (17.9%) followed by gestational diabetes in 2 cases (5.1%). The most common neonatal risk factors for ROP was respiratory distress syndrome in 35 babies (89.7%) followed by sepsis in 5 cases (12.8%).

TABLE 1:

ROP(stage)	NO. (%)
Stage 1	5 (12.8%)
Stage 2	16 (41%)
Stage 3	18 (46.2%)
Stage 4	0
Stage 5	0

DISCUSSION:

ROP is considered to be a preventable cause of childhood blindness especially if diagnosed and treated at an appropriate time. The incidence of ROP in premature infants in our study was 19.5%. In developed countries, the reported incidence of ROP ranges from 19.3% to 25%^(12,13). Likewise in India, the incidence of ROP depends on the region of study and ranges from 20% to 47.3%^(14,15). In the current study the incidence of ROP is almost same as other studies performed in India. As reported by Palmer, *et al.*⁽¹⁶⁾, incidence and severity of ROP was closely related to lower birth weight and lower postconceptional age, as was seen in our study.

There are varying screening criteria described by different authors. Vinekar, *et al.*⁽¹⁷⁾ suggested that the scenario in developing countries is quite different. Larger and gestationally 'older' infants are more likely to develop ROP compared to their counterparts in Western countries. Hence, the application of Western screening guidelines for developing countries has been questioned by Jalali, *et al.*⁽¹⁸⁾. As a higher cut off limit, they recommended screening babies born at <37 weeks gestation and/or birthweight <2000g in the presence of a high sickness score, in order to prevent missing any infant with threshold ROP.

So, we feel that all babies with birthweight less than 1500g and gestation ≤ 32 weeks should be routinely screened. Infants with birthweight between 1500-2000g and gestational age more than 32 weeks should be screened at the discretion of the neonatologist, depending on other risk factors during the course of stay in the NICU.

Many risk factors have been reported to predispose to the development of ROP. In our study the most common maternal risk factor noted was pregnancy induced hypertension seen in 17.9% followed by

gestational diabetes in 5.1%. Most common neonatal risk factors were respiratory distress syndrome in 89.7% followed by sepsis in 12.8%. Oxygen therapy was not found to influence the development of ROP in our study.

The higher rate of treatment required in our study is because our NICU is a tertiary referral center where sick babies are usually referred.

CONCLUSION:

ROP screening guidelines should vary according to the region. The most common risk factors were low birth weight, lower gestational age and respiratory distress syndrome and pregnancy induced hypertension.

REFERENCES

1. Yang CS, Chen SJ, Lee FL, Hsu WM, Liu JH. Retinopathy of prematurity: Screening, incidence and risk factors analysis. *Zhonghua Yi Xue Za Zhi (Taipei)* 2001;64:706–712.
2. Saugstad OD. Oxygen and retinopathy of prematurity. *J Perinatol.* 2006;26(Suppl 1):S46–50.
3. Gilbert C. Retinopathy of prematurity: A global perspective of the epidemics, population of babies at risk and implications for control. *Early Hum Dev.* 2008;84:77–82.
4. Saugstad OD. Is oxygen more toxic than currently believed? 2001;108:1203–1205.
5. Penn JS, Henry MM, Wall PT, Tolman BL. The range of PaO₂ variation determines the severity of oxygen-induced retinopathy in newborn rats. *Invest Ophthalmol Vis Sci.* 1995;36:2063–2070.
6. Lucey JF, Dangman B. A reexamination of the role of oxygen in retrolental fibroplasia. 1984;73:82–96.
7. Mousavi SZ, Karkhaneh R, Riazi-Esfahani M, Mansouri MR, Roohipoor R, Ghalichi L, et al. Retinopathy of prematurity in infants with late retinal examination. *J Ophthalmic Vis Res.* 2009;4:24–28.
8. Fortes Filho JB, Eckert GU, Procianny L, Barros CK, Procianny RS. Incidence and risk factors for retinopathy of prematurity in very low and in extremely low birth weight infants in a unit-based approach in southern Brazil. *Eye (Lond)* 2009;23:25–30.
9. Gilbert C, Fielder A, Gordillo L, Quinn G, Semiglia R, Visintin P, et al. Characteristics of infants with severe retinopathy of prematurity in countries with low, moderate, and high levels of development: Implications for screening programs. 2005;115:e518–525.
10. Section on Ophthalmology American Academy of Pediatrics; American Academy of Ophthalmology; American Association for Pediatric Ophthalmology and Strabismus. Screening examination of premature infants for retinopathy of prematurity. 2006;117:572–576.
11. Early Treatment for Retinopathy of Prematurity Cooperative Group. Revised indications for the treatment of retinopathy of prematurity: Results of the early treatment for retinopathy of prematurity randomized trial. *Arch Ophthalmol.* 2003;121:1684–1694.
12. Ho SF, Mathew MR, Wykes W, Lavy T, Marshall T. Retinopathy of prematurity: An optimum screening strategy. *J AAPOS.* 2005;9:584–588.
13. Larsson E, Holmström G. Screening for retinopathy of prematurity: Evaluation and modification of guidelines. *Br J Ophthalmol.* 2002;86:1399–1402.
14. Maheshwari R, Kumar H, Paul VK, Singh M, Deorari AK, Tiwari HK. Incidence and risk factors of retinopathy of prematurity in a tertiary care newborn unit in New Delhi. *Natl Med J India.* 1996;9:211–214.
15. Gupta VP, Dhaliwal U, Sharma R, Gupta P, Rohatgi J. Retinopathy of prematurity – Risk factors. *Indian J Pediatr.* 2004;71:887–892.
16. Palmer EA, Flynn JT, Hardy RJ, Phelps DL, Phillips CL, Schaffer DB, et al. Incidence and early course of retinopathy of prematurity. *Ophthalmology* 1991; 98: 1628-1640.
17. Vinekar A, Dogra M, Sangtam T, Narang A, Gupta A. Retinopathy of prematurity in Asian Indian babies weighing greater than 1250 grams at birth: ten year data from a tertiary care center in a developing country. *Indian J Ophthalmol* 2007; 55:331-336.
18. Jalali S, Anand R, Kumar H, Dogra MR, Azad RV, Gopal L. Programme planning and screening strategy in retinopathy of prematurity. *Indian J Ophthalmol* 2003; 51: 89-99.
19. Ng EY, Connolly BP, McNamara JA, Regillo CD, Vander JF, Tasman W. A comparison of laser photocoagulation with cryotherapy for threshold retinopathy of prematurity at 10 years. Part 1-Visual function and structural outcome. *Ophthalmology* 2002; 109: 928-934.
20. Connolly BP, Ng EY, McNamara JA, Regillo CD, Vander JF, Tasman W. A comparison of laser photocoagulation with cryotherapy for threshold retinopathy at 10 years. Part 2-Refractive outcome. *Ophthalmology* 2002; 109: 936-941.