Microbiological Study of Dacryocystitis in Paediatric Age Group

Prathiba¹, Aruna Sunder², Taruni³, V. Sudha Rani⁴

ABSTRACT

Introduction: Paediatric acute dacryocystitis (PAD) is a special subset with features that are unique and distinct from those of adults. There are few studies on paediatric dacryocystitis, hence the present study was undertaken to see the incidence of various microbial pathogens causing paediatric dacryocystitis, to identify and study their antibiogram.

Material and Methods: Retrospective study of 168 patients of pediatric age group, attending outpatient department of a tertiary care eye hospital, clinically diagnosed as dacrocystitits was done. Discharge from lachrymal punctum was collected by 2 sterile swabs, one was used for staining and another was inoculated onto Blood agar, Chocolate agar, Mac Conkey agar and Sabouraud's dextrose agar (SDA). The isolated organisms were identified using standard procedures. Sensitivity was tested by Kirby – Bauer's disc diffusion method.

Results: Out of the 168 samples collected,majority cases were between 1month to 3 years, dacryocystitis was common in right eye,72.6% were culture positive and 27.3% were negative. Staphylococcus epidermidis was the predominant organism. Fungal isolates were three, Aspergillus spp 2 and Fusarium spp 1.Majorityof S.epidermidiswere sensitive to Gatifloxacin, least sensitivity was to Ca, Staph.aureus were sensitive to Mo and Tb, Enterococci, Streptococci and Micrococciwere sensitive to all antibiotics except Ca. In case of Pseudomonas Oflaxacin and Gatifloxacin showed 100% sensitivity.

Conclusions: Paediatric dacryocystitis is a distinct entity with unique features of its own. In the era ofantibiotic resistance, microbiological work up of paediatric acute dacryocystitis is very useful forsubsequent treatment.

Keywords: Dacryocystitis, paediatric, antibiotic sensitivity.

INTRODUCTION

Dacryocystitis is caused by obstruction of nasolacrimal duct. It is due to malformation of tear duct, infection of eye, taruma or injury. Clinically patient presents with swelling over the inner aspect of the lower eyelid, redness and pain. There are 2 forms of Dacryocystitis, acute and chronic.² The acute form could be associated with severe morbidity and primarily related to the lacrimalsac abscess and spread of infection.^{2,3} There is a varied spectrum of its clinical presentations ranging from tenderness and erythemaof the overlying tissues to a frank lacrimal abscess.4 Untreated lacrimal abscess can progress to orbital cellulitis, superior ophthalmic vein thrombosis, and cavernoussinus thrombosis. 5-7 Acute dacryocystitis can present as a medical emergency with sudden pain, erythema and swelling, below the medial canthal tendon. Infection of lacrimal sac and perisac tissues can lead to epiphora.⁸ Clinically Paediatric acute dacryocystitis (PAD) presents as dacryocele in neonates. It can lead to complications like orbital cellulitis, orbital abscess, meningitis and loss of vision. 9-12 There are few studies on paediatric dacryocystitis, hence the present study was undertaken to see the incidence of various microbial pathogens causing paediatric dacryocystitis, to identify various bacterial isolates and study their antibiogram.

MATERIAL AND METHODS

A retrospective study of 168 patients of paediatric age group of either sex, attending outpatient department of a tertiary care eye hospital, clinically diagnosed as dacryocystitis by ophthalmologists was done after ethical board clearance.

Specimen Collection: After cleaning with normal saline swab, pressure was applied at medial epicanthic fold, the regurgitated pus or serosanguinous fluid was collected by sterile swab, two sterile cotton swabs moistened with physiological saline were used for collection of discharge from lachrymal punctum.

Specimen processing: One swab was spread on glass slide to prepare smear and stained by Grams stain. The second swab was used for inoculation ontoculture media like Blood agar, Chocolate agar, Mac Conkey agar and Sabauraud's dextrose agar (SDA). The inoculated media were incubated at 37°C for 24hrs to 48 hrs for aerobic cultures and SDA at room temperature for 3 weeks. The stained smears were screened for presence or absence of pus cells and bacteria, KOH mount for fungal elements. The isolated organisms were identified using standard procedures. Antibiotic sensitivity of organisms was tested by KirbyBauer's disc diffusion method on Muller hinton agar using the following antibiotics, 30Mcg-Chloramphenical (C), 30Mcg Ceflazidime (CA), 5Mcg Ciprofloxacin (CF), 5Mcg Oflaxacin (OF), 5Mcg Gatifloxacin (GF), 10Mcg Gentamycin (G), 5Mcg Moxifloxacin (MO), 10Mcg Tobramycin (TB).

RESULTS

Out of the 168 samples collected over a period of one year, 85 were in the age group of 1month -1 year, 36 were between 1-2 years, 23 were between 2-3 years and 25 were between 3-5 years. Out of 168 patients boys were 92, girls were 76, da-

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cryocystitis was common in right eye (93 patients) and in 75 patients left eye was affected. Majority cases were between 1month to 3years (Table - 1). Out of total 168 samples 122 (72.6%) were culture positive and 46 (27.3%) were culture negative. Bacterial organisms isolated were *Staphylococcus epidermidis* 86 (70.4%), *Staphylococcus aureus* 17 (13.9%), *Enterococcus* species 09 (7.3%), *Streptococcus* species 04 (3.2%), *Pseudomonas aeruginosa* 04 (3.2%), *Micrococci* 02 (1.6%). *Staphylococcus epidermidis* was the predominant organism. In Gram negative organisms it was only *Pseudomonas*. (Fig-1) Fungal isolates were three, *Aspergillus* spp 2 and *Fusarium* spp 1.

Sensitivity patterns: In case of *S.epidermidis*78 isolates were sensitive to Gatifloxacin, least sensitivity was to Ca, next CF, in case of *Staph.aureus* out of 17, Mo and Tb shown high sensitivity, CF least sensitive, for *Enterococci* sensitivity almost same for all antibiotics, *Streptococci* were almost sensitive to all antibiotics except Ca. In case of *Pseudomonas* Oflaxacin and Gatifloxacin showed good sensitivity of 100%. *Micrococci* were resistant only to Ca and sensitive to all. (Table 2)

DISCUSSION

In the present study out of 168 samples collected, male children were 92 and female children were 76, in 93 right eye was affected and in 75 left eye showed infection, right eye infection was common. In a study done by Mohammed Javed Ali et al, the female to male ratio was approximately 1.7:1, there was no preponderance of laterality.¹³ In a study done by 0. 0. Ffook *et al* StRoyal Infirmary, ¹⁴ of the total se-

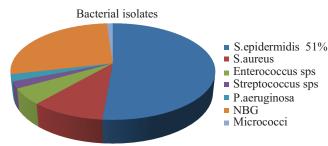


Figure-1: Showing different bacterial isolates.

Age	Number		
1month-1year	85		
1-2 years	36		
2-3 years	23		
3-4years	08		
4-5 years	07		
>5 years	09		
Table-1: Showing age wise distribution of cases.			

ries, 224 were girls and 213 were boys. This proportion is not very different from the normal ratio and is not considered significant. The right eyewas affected in 53 (47.3%) patients and the left eye in 59 (52.7%) patients. In a study done by Mohammed Javed Ali et al¹³, they have separately analyzed the 13 pediatric patients, there was no sex predilection with 7 malesand 6 females. The mean age at presentation was 30.6 months (range 22 days to 108 months). In our study majority of cases were in the age group ofbetween 1month to 36months. In a study done by Yared Assefa, Feleke Moges, Mengistu Endris, Banchamlak Zereayet al from the total of 51 dacryocystitis cases, bacterial origins were isolated among 31 (60.8%) cases. 15 In an interesting study of 47 children, Kuchar et al., observed that Gram positive bacteria were more frequently isolated in the samples obtained, S. pneumonia being the predominant microorganism in 36.4% of cases, followed by H. influenzae (19.6%).16 Mohammad Javed Ali, Swapna R Motukupally, Surbhi D Joshi and Milind N Naik observed in their study that the microbiological profile was not found to be different in the pediatric subset of their studygroup with S. aureus being the most common organism followed by S. pneumonia. 13 In our study Staphylococcus epidermidis was the predominant organism 86 (70.4%), followed by Staphylococcus aureus 17 (14.2%), and the only Gram negative organism isolated was Pseudomonas aeruginosa. We have isolated 3 fungi (3.4%). As forfungi, they have been reported to be present in 4% to 7% ofcases, the most commonly isolated genus being Candida, although Aspergillus and Mucor may also be found. 17 In a study done by Supriya Ghose, VM Mahajan¹⁸ fungal isolates were 12 (13.95%). -.5 were C. albicans and 5 were A. niger -. Our study showed 3 fungi, 2 Aspergillus and 1 Fusarium. Antibiotic sensitivity results showed that in casof S.epidermidis 78 isolates were sensitive to Gatifloxacin, least sensitivity was to Ca, next CF, in case of Staph.aureus out of 17, Mo and Tb shown high sensitivity, CF least sensitive, for Enterococci sensitivity almost same for all antibiotics, Streptococci were almost sensitive to all antibiotics except Ca. In case of *Pseudomonas* Oflaxacin and Gatifloxacin showed good sensitivity of 100%. Micrococci were resistant only to Ca and sensitive to all. In a study done by Yared Assefa, Feleke Moges, Mengistu Endris, Banchamlak Zereayet al the antimicrobial susceptibility tests revealedthat ceftriaxone (95.3%), erythromycin (84.2%), nalidic acid (87.1%), gentamycin (83.3%) were more effective than other antibiotics tested to all bacterial isolates.

CONCLUSIONS

Paediatric dacryocystitis is a distinct entity with unique features of its own. It is a serious infection that needs careful

C	CA	CF	OF	GF	G	MO	TB
74S,12R	31S,55R	48S,38R	57S,29R	78S,8R	69S,17R	71S,15R	72S,14R
12S,5R	8S,9R	3S,14R	6S,11R	8S,9R	9S,8R	13S,4R	13S,4R
8S,1R	6S,3R	7S,5R	8S,1R	8S,1R	7S,2R	8S,1R	7S,1R
4S,0R	1S,3R	4S,0R	4S,0R	4S,0R	4S,0R	4S,0R	4S,0R
2S,2R	2S,2R	2S,2R	4S,0R	4S,0R	3S,1R	3S,1R	3S,1R
2S,0R	0S,2R	2S,0R	2S,0R	2S,0R	2S,0R	2S,0R	2S,0R
	12S,5R 8S,1R 4S,0R 2S,2R	74S,12R 31S,55R 12S,5R 8S,9R 8S,1R 6S,3R 4S,0R 1S,3R 2S,2R 2S,2R	74S,12R 31S,55R 48S,38R 12S,5R 8S,9R 3S,14R 8S,1R 6S,3R 7S,5R 4S,0R 1S,3R 4S,0R 2S,2R 2S,2R 2S,2R	74S,12R 31S,55R 48S,38R 57S,29R 12S,5R 8S,9R 3S,14R 6S,11R 8S,1R 6S,3R 7S,5R 8S,1R 4S,0R 1S,3R 4S,0R 4S,0R 2S,2R 2S,2R 2S,2R 4S,0R	74S,12R 31S,55R 48S,38R 57S,29R 78S,8R 12S,5R 8S,9R 3S,14R 6S,11R 8S,9R 8S,1R 6S,3R 7S,5R 8S,1R 8S,1R 4S,0R 1S,3R 4S,0R 4S,0R 4S,0R 2S,2R 2S,2R 2S,2R 4S,0R 4S,0R	74S,12R 31S,55R 48S,38R 57S,29R 78S,8R 69S,17R 12S,5R 8S,9R 3S,14R 6S,11R 8S,9R 9S,8R 8S,1R 6S,3R 7S,5R 8S,1R 8S,1R 7S,2R 4S,0R 1S,3R 4S,0R 4S,0R 4S,0R 4S,0R 2S,2R 2S,2R 2S,2R 4S,0R 4S,0R 3S,1R	74S,12R 31S,55R 48S,38R 57S,29R 78S,8R 69S,17R 71S,15R 12S,5R 8S,9R 3S,14R 6S,11R 8S,9R 9S,8R 13S,4R 8S,1R 6S,3R 7S,5R 8S,1R 7S,2R 8S,1R 4S,0R 1S,3R 4S,0R 4S,0R 4S,0R 4S,0R 2S,2R 2S,2R 2S,2R 4S,0R 4S,0R 3S,1R 3S,1R

evaluation and immediate management. In the era ofantibiotic resistance, microbiological work up of paediatric acute dacryocystitis is very useful forsubsequent treatment.

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