

QUANTITATIVE LEVELS OF C- REACTIVE PROTEIN IN CEREBROSPINAL FLUID IN CHILDREN WITH BACTERIAL AND OTHER MENINGITIS AT PEDIATRICS

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Pediatrics

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ABSTRACT

BACKGROUND: C-reactive protein rises rapidly in the first 24-48 hours of occurrence of bacterial meningitis and in large incremental increases thereafter.

METHODS: All children (121) admitted during the period of study, with clinical suspicion of meningitis were clinically, biochemically, cytologically and bacteriologically investigated to clinch the diagnosis. Quantitative level of CSF was also sent for C-reactive protein assay by latex agglutination test by immuno-turbidometric method.

RESULTS: There were 121 cases of meningitis admitted in the hospital and only 10 cases were culture positive. There was significant difference of quantitative level of CSF CRP between bacterial and other than bacterial meningitis.

CONCLUSION: Quantitative estimation of CRP in CSF by IMMUNOTURBIDIMETRY is a useful test to differentiate pyogenic meningitis from tubercular meningitis, viral meningoencephalitis and other non-meningitis CNS disorders. It is an easy, quick to perform, sensitive, reliable and rapid diagnostic test for timely therapeutic intervention. The detection of CRP in CSF also helps in the choice of appropriate antibiotic and the duration of therapy.

KEYWORDS

C-reactive protein, CSF, Meningitis, viral meningoencephalitis

INTRODUCTION:

Meningitis is one of the most potentially serious infections occurring in infant and older children and important cause of morbidity and mortality in children worldwide. In the developing countries, acute bacterial meningitis (ABM) is one of the major cause of morbidity and mortality. Rapid and accurate diagnosis coupled with early appropriate therapy is the utmost importance in reducing morbidity and mortality of the patient. Biochemistry, Cytology, Gram stain, Culture sensitivity of Cerebrospinal fluid usually being done to diagnose and differentiate pyogenic from aseptic meningitis. In such circumstances the detection of C-reactive protein (CRP) in CSF appears to provide a new dimension to the diagnosis of meningitis.

In developing countries, with limited resources, manpower, and skill, we need to think about an easy and comprehensive test by which we can diagnose ABM. Routine use of detection of CSF CRP, could be a reliable and easy to do test and can be done for rapid diagnosis of meningitis.

OBJECTIVE:

GENERAL OBJECTIVE:

The study is aimed to estimate the level of CSF-CRP quantitatively in bacterial & other meningitis in among children admitted in Pediatric department of the IGIMS, Patna, Bihar.

METHODOLOGY:

Study design: The estimation of quantitative CSF-CRP among children of meningitis is a hospital based descriptive cross sectional study.

Place of study: All aspects of this study have been accomplished in the department of Pediatrics, of the Indira Gandhi Institute of Medical Sciences, Patna, Bihar.

Period of Study: The study was done in a period of 15 months from January 2019 to March 2020.

Sample Size: Total patients were 121.

Inclusion Criteria:

Infant and children of ages 1 day to 14 years diagnosed case of

meningitis, who required a CSF study to confirm or exclude the diagnosis.

Exclusion criteria:

Congenital anomalies of CNS, Non infectious conditions of CNS, Infections at the site of lumbar puncture.

Method of CSF-CRP was quantitatively detected by immuno-turbidimetry method by using Biosystems CRP kit.

121 patients of meningoencephalitis, between 1 day to 14 years of age were included in the study and were divided in 4 groups:- Group I- Pyogenic meningitis, Group II- Tubercular meningitis, Group III- Viral meningitis and Group IV- No CNS infection (Extra cranial infections).

RESULTS:

There were 53 cases of bacterial meningitis, 27 cases of viral meningoencephalitis, 13 cases of tubercular meningitis and 28 cases of control group. The male female ratio in the Acute Bacterial Meningitis, Tubercular Bacterial Meningitis, Viral Meningocephalitis and Control Group were 1.94:1, 5.5:1, 1.7:1 and 1.33:1 respectively. All groups were with male predominance.

Table no-1: Distribution according to sex & diagnosis of patients

Diagnosis	Sex		Total n (%)
	Male n (%)	Female n (%)	
Acute Bacterial Meningitis	35(66.03)	18(33.97)	53(100.00)
Tubercular Bacterial Meningitis	11(84.62)	2(15.38)	13(100.00)
Viral Meningocephalitis	17(62.97)	10(37.03)	27(100.00)
Control Group	16(57.15)	12(42.85)	28(100.00)
Total	79(65.29)	42(34.71)	121(100.00)

In present study, the total 53 cases of pyogenic meningitis, 18 (33.96%) were in the age group of <1 year, 12 (22.64%) were in the age group of 1-5 years and 23 (43.39%) were in the age group of 5-14 years. CSF-culture was positive only 10 cases of bacterial meningitis while all other cases of all groups were culture negative.

In our study, 47 (88.67%) cases out of 53 had quantitative level of CSF-CRP > 1000 ng/ml in bacterial meningitis patients and only 6 (11.32%) cases had <1000 ng/ml. 19 cases (35.84%) had >2000 ng/ml level of

CSF-CRP in this group. In tubercular meningitis patients 11 (84.61%) of 13 had <1000ng/ml quantitative level of CSF-CRP and only two had > 1000ng/ml CSF-CRP. In viral meningoencephalitis group CSF-CRP was >1000ng/ml in 5 cases of 27 and rest 22 cases had < 1000ng/ml quantitative level of CSF-CRP.

Table no-2 Distribution according to CSF CRP level & diagnosis of patients

CRP (ng/ml)	Diagnosis				Total n (%)
	ABM n(%)	TBM n(%)	VME n(%)	Control Group n (%)	
< 1000	6 (11.32)	11 (84.61)	22 (81.48)	25 (89.28)	64 (52.89)
1000-2000	28 (52.83)	2 (15.38)	5 (18.51)	3 (10.71)	38 (31.40)
> 2000	19 (35.84)	0 (0.00)	0 (0.00)	0 (0.00)	19 (15.70)
Total	53 (100.00)	13 (100.00)	27 (100.00)	28 (100.00)	121 (100.00)

The mean CSF CRP was 1862 ng/ml in bacterial meningitis while it was 597 ng/ml in control group and was 992 ng/ml and 816 ng/ml in viral meningoencephalitis & tubercular bacterial meningitis respectively. The difference in CSF CRP between bacterial and all other groups was highly significant.(table3)

Table no-3 Mean +SD of CSF CRP according to diagnosis of subjects

Diagnosis	No.	Mean +SD ng/ml
ABM	53	1862.20 +812.19
TBM	13	816.00 +361.55
VME	27	992.00 +462.92
Control Group	28	597.20 +301.38

Control Group v/s ABM	p <0 .001	Highly Significant	ABM v/s TBM	p <0.001	Highly Significant
Control Group v/s VME	p <0 .01	Significant	Control Group v/s TBM	p < 0.001	Highly Significant
ABM v/s VME	p <0 .01	Significant	TBM v/s VME	p > 0.05	Not Significant

DISCUSSION:

Bacterial meningitis is a potentially devastating illness. So prompt recognition and early appropriate treatment is essential. Males are more affected in pyogenic (66%) meningitis.

In the present study Gram's smear and CSF culture for pyogenic organism could detect only 24% and 20% cases respectively.

In our study IMMUNOTURBIDIMETRY method was used for quantitative analysis of CRP in CSF by using Biosystems CRP kit to detect CRP in CSF of suspected cases of meningitis. We studied the utility of CRP in the differential diagnosis of meningitis of varying etiology.

Our analysis showed the quantitative value of CSF CRP was significantly higher in bacterial meningitis group. The difference between pyogenic meningitis and control group ($p < 0.001$), between pyogenic meningitis and viral meningo encephalitis ($p < 0.001$) and between pyogenic meningitis and tubercular meningitis were highly significant ($p < 0.001$). The difference between tubercular meningitis and control group and between viral meningo-encephalitis and control group were significant ($p < 0.05$). While the difference between tubercular meningitis and viral meningo-encephalitis was not significant ($p > 0.05$). These findings were similar with other study findings like Donald PR et al, Gray BM et al, Hanson LO et al, Trienekens PH et al, Shimetani N et al, Belal M et al (3-8).

CONCLUSION:

Although culture of CSF is the gold standard for diagnosis of pyogenic meningitis for direct evidence of the organism, it has some limitation particularly in peripheral set up. Routine use of detection of CSF-C-reactive protein could be a reliable, rapid and easy to do test for the diagnosis of bacterial meningitis and differentiate from aseptic meningitis. It is not an alternative of examination of CSF biochemistry, cytology and culture.

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