

CRP, an acute phase reactant: Role in Differentiating Bacterial and Non-Bacterial CNS Infections

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Abstract

C reactive protein is marker of acute inflammation. Now quantitative measurement of CRP is possible. Its quantitative measurement is better indicator of inflammation than qualitative one.

C reactive Protein is known acute phase reactant since 1930¹. Its level increases with any kind of inflammation. Meningitis still remains very common reason for morbidity and mortality in developing countries like India. Although cerebrospinal fluid examination and Culture remains gold standard for diagnosis of meningitis but facilities are not easily available especially in emergency hours. There are very few studies available in literature to evaluate role of C reactive proteins in Cerebrospinal fluid to differentiate Non bacterial and bacterial CNS Infections^{2, 3, 4}. In developing countries patient usually reach to tertiary care centre after receiving various antibiotics. In this situation culture report is either negative or it takes few days before showing any result. Often CSF examination is not possible or show picture of partially treated meningitis. We give empirical antibiotics therapy in this situation.

Bejagavi et al⁶ in his study acute phase reactant Adenosine deaminase and CRP in Cerebrospinal fluid. ADA has a sensitivity and specificity of 73.9% and 92.6% respectively for Tubercular meningitis while CSF C- reactive Proteins have a sensitivity and specificity of 83.3 % and 100% respectively for Pyogenic meningitis.

Other studies^{7, 8, 9, 10, 11} have similar results. Sensitivity and specificity of CSF C reactive Protein was high for for Pyogenic meningitis. It also important tool to differentiate Viral and partially treated meningitis with Pyogenic meningitis. Measurement of other acute phase reactant with CSF C reactive Protein will further increases sensitivity and specificity of diagnostic tool.

Patel et al⁵. In his study he used CSF C reactive Protein as a bedside tool to differentiate between Bacterial and Non bacterial Meningitis. Results were very encouraging while awaiting result of CSF examination. Sensitivity and specificity was 83.3 and 87.5% respectively for Pyogenic meningitis. Similarly Positive predictive value was 75% and Negative predictive value was 92% for Bacterial meningitis. In tubercular meningitis sensitivity and specificity was 20% and 60.5% respectively. In viral meningitis CSF C reactive protein was negative in all cases. Similar results were seen with Mala et al². In her study sensitivity and specificity was 96.87 % and 74.73% respectively. For Tubercular meningitis sensitivity and specificity was 10% and 55.38%.

This study⁵ has some limitations. They have used a Qualitative CRP kit which shows either positive or negative results. CSF C reactive proteins by measuring quantitative methods could have been better. This study included a small sample size. Thus while measuring CSF CRP as screening methods we need further study with larger sample size to show its efficacy. Similarly a larger control group should be taken for better analysis. Measurement of virological markers in CSF could have been done to diagnose viral meningitis.

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