A Study of Clinical and CSF Characteristics in Cases of Acute Meningoencephalitis in Immunocompetent Adults in a Tertiary Care Hospital of Eastern India

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ABSTRACT

Meningoencephalitis is a very critical illness that is widespread and it remains a major cause of mortality and morbidity with neurological disabilities. It usually presents with varying degrees of symptoms of meningeal inflammation, posing difficulties in diagnosis and treatment. Study was conducted to Evaluate clinical severity and CSF (cerebrospinal fluid) findings at presentation. Encompassing the clinical status, CSF assays, along with revelations of recent trends of infection responsible for meningocencephalitis, this study also shows that early confirmation of clinical suspicion with judgement of severity & CSF study, is of great significance. Prompt diagnosis provides physicians with an opportunity to prevent undue mortality and morbidity. Here lies the relevance of this study.

Key Words: Acute Meningoencephalitis, Cerebrospinal fluid, MRI of brain

INTRODUCTION

The incidence of acute encephalitis in western countries is 7.4 cases per 100,000 population per year. In tropical countries, the incidence is 6.34 per 100,000 per year.¹ Herpes simplex encephalitis has an incidence of 2–4 per million population per year.² The common pathogens which are encountered in adult bacterial meningitis are Streptococcus pneumoniae (30–50%), Neisseria meningitidis (10–35%), Staphylococci (5–15%), other Streptococcus species, Haemophilus influenza (1–3%), Gram negative bacilli (1–10%) and Listeria monocytogenes.³ Prompt recognition, early diagnosis, efficient decision making followed by rapid institution of therapy plays a pivotal role in saving a large salvageable portion of the affected population and thus reducing mortality. Tuberculous meningitis (TBM) remains the most common presentation. In spite of advances in diagnostic technology and effective therapeutic options, it continues to pose significant management challenges. Despite anti-TB chemotherapy, 20-50% of the affected people die and many who survive have significant neurological deficits. The case fatality noted to be associated significantly with delay in diagnosis and treatment. Tuberculous meningitis (TBM) generally occur in course of a sub acute or chronic case but TBM may have an acute presentation. The duration of presenting symptoms may vary from 1 day to 9 months, although several cases may present with symptoms of less than 2 weeks duration. Diagnostic evaluation includes various microbiological, pathologic, molecular, and biochemical investigations & imaging modalities. Imaging helps in early diagnosis and helps in preventing morbidity and mortality.⁴
**Aims and objectives of these studies**
Evaluation of clinical severity and CSF (cerebrospinal fluid) findings at presentation in Acute Meningoencephalitis.

**MATERIALS AND METHODS**

a) Patients, who were admitted in between January 2013—August 2014 in the department of General Medicine of NRS Medical College and Hospital from rural and urban catchment area, were included in this study as simple random selection. 50 patients, aged >12 years were included in this prospective observational study with Fever and Signs of Meningitis (Nuchal rigidity, vomiting, and headache) or Signs of Meningoencephalitis: Meningeal signs with altered sensorium, focal neuro-deficits, and seizures. The following patients with Sepsis, Metabolic Encephalopathy, Dyselectrolytemia, Poisoning, Cerebrovascular Accident, Intracranial SOL, Neurocysticercosis, Enteric Fever with meningism, Vascular Aneurysms producing local compressive effect, Acute disseminated Encephalomyelitis (ADEM) and Cerebral Malaria were excluded from the study. Clinical Characteristics—Glasgow Coma Scale (GCS) Scoring (3 to 15) as a marker of clinical severity on admission. According to the score calculated on admission, patients divided into three groups—Gr. A (GCS 3–5), Gr. B (GCS 6–9), Gr. C (GCS 10–15). : The macroscopic appearance of the CSF recorded. A routine CSF total and differential count were done by a haemocytometer by standard methods. The CSF samples subjected to a cytospin by using Shandon cytospin MODEL 001/002. Gram stained of CSF was done and examined under microscope. ZN staining, Bacterial culture, TB culture (BACTEC) and Cryptococcal staining (India Ink Stain) were also done. Specific viral analysis was done by ELISA according to relevance, availability and feasibility. Final Diagnosis Based on Set Criteria and Segregation of cases according to etiological groups from CSF and MRI findings. Descriptive statistical methods were used, utilizing the SPSS software for data analysis.

**REVIEW OF LITERATURE**
Mortality of acute bacterial meningitis is 25%. Morbidity in the tem of neuro-deficit is also very high. Now are going through the advanced stage of antibiotic era & critical care facility. In spite of that we are facing increased mortality. The prognosis is worse with a delay in management. Increased morbidity and mortality is seen in both high and low-income group & countries. It is very tough job to select patients for timely antibiotic administration in emergency room.

Schutte CM and Van der Meyden CH in their study found good correlation between both the GCS and CSF-protein level at admission and the outcome of patients with meningitis was found. With the GCS value being a better prognostic indicator than high CSF protein levels, Syamal Modi and Amit Kumar Anand in their study found Streptococcus pneumonia was the most common pathogen which was isolated in 120 (60%) culture positive cases. Cell counts showed the predominance of neutrophils in all cases with ABM. In ABM protein is high & sugar is low. Gram staining is one of the cornerstones methods in diagnosing ABM in developing countries.

Mani R, Pradhan S et al in their study conducted in South India (NIMHANS) in 2007 observed that, as compared to Western studies, the relative incidence of meningitis caused by H. influenza, N. meningitides and Listeria is less in South-East Asia. On the contrary, gram negative bacilli such as Klebsiella pneumonia and Pseudomonas aeruginosa are increasingly being recognised as important pathogens of community-acquired as well as nosocomial meningitis especially among the elderly and in patients with chronic debilitating diseases like cirrhosis, diabetes and malignancies.

Steiner, H. Budka et al in their review of diagnosis and management recommendations in Viral encephalitis states that analysis of CSF for protein and glucose contents, cellular analysis and identification of the pathogen by biotechnology method like PCR polymerase chain reaction (PCR) and serology—Hospitalisation is mandatory with ICCU facility. Combinations of meningitis/encephalitis and myelitis/
radiculitis are associated with Epstein Barr Virus (EBV); myelitis with VZV, CMV, EBV, and HSV-2; and ventriculitis/encephalitis with VZV and CMV. Brainstem encephalitis due to HSV and VZV, and poly myeloradiculitis due to CMV are well documented. Recent large CSF PCR studies have shown that VZV, EBV, and CMV more frequently produce meningitis, encephalitis, or encephalopathy in immunocompetent hosts than was formerly realized.[12]

A study conducted by Rathore SK, Dwibedi B et al during April 2011 to July 2012. Blood and CSF samples of 526 AES cases are investigated by serology and/or PCR. Viral aetiology was identified in 91 (17.2%) cases. Herpes simplex virus (HSV; types I/ II) was most common (16.1%), followed by measles (2.6%), Japanese encephalitis virus (1.5%), dengue virus (0.57%), varicella zoster virus (0.38%) and enteroviruses (0.19%). Simultaneous infection of HSV-I and measles were observed in seven cases. This report provides the first evidence on viral aetiology of acute encephalitis syndrome viruses from eastern India showing dominance of HSV that will be useful in informing the public health system.[13] However, measurement of antibodies in single specimen of serum and CSF lacks sensitivity; they interpreted the positivity (positive for IgM and/or IgG antibody) in relation to other supportive clinical, electroencephalographic and neuro-radiological evidences. CSF-PCR is the diagnostic test of choice, which has sensitivity rate as high as 98-99% and specificity of 100%,.[14] HSE is the only form of sporadic encephalitis which has a specific antiviral therapy i.e. acyclovir,[15] F. Frantzidou, F. Kamaria et al found in their study that enterovirus was the most common cause of adult aseptic meningitis and together with HSV-1 the main causes of encephalitis.[16]

RESULTS AND ANALYSIS

GENDER DISTRIBUTION: (Figure 1 & Chart 1) All immune-competent patients aged 12 yrs or more were selected randomly over a period of 19 months for this observational study as per the clinical inclusion criteria. 18 cases (36%) were male and 32 cases (64%) were female. Patients ranging from ages 12 yrs to 77 yrs were selected and divided into age groups. Maximum number of patients (26%) belonged to 12-20 yrs age group.

Groups according to the Glasgow coma Scale: (See Figure 4 & Chart 2) The patients were clinically examined and classified into three groups according to the Glasgow coma scale. 15 cases (30%) belonged to group A (GCS 3 to 5), 22 cases (44%) Belonged to Gr. B (GCS 6 to 9) while 13 cases (26%) belonged to Gr. C (GCS 10 to 15) on admission, 30% patients had a very poor clinical condition (GCS 3 to 5) during admission as evident from the GCS category distribution while majority of patients belonged to group B. (See table 2 & Chart 2)

Etiological groups & gender distribution: (See Figure 2 & Chart 3) After obtaining the investigation results the cases were segregated into etiological groups and re classified according to the GCS groups and gender. Majority of cases had a viral aetiology (56%) followed by tuberculous (28%) and bacterial aetiology (16%). Pyogenic meningitis cases had a Male predilection (62.5%) while tuberculous and viral meningitis had female predilection in incidence, 57.14% and 75% respectively.

Distribution according to age groups: (See Figure 3 & Chart 5)

Majority of pyogenic meningitis cases (37.5%) belonged to 31-40 yrs age group while maximum number of tuberculous meningitis cases (35.71%) and viral Meningoencephalitis cases (28.57%) belonged to 12-20 yrs age group. Out of 50 cases 8 cases (16%) were diagnosed as pyogenic meningitis [5 males (62.5%) and 3 females (37.5%)]

Distribution of Patients according to GCS on admission: (See figure 4 & Chart 6) 1 case was in Gr A (GCS 3 to 5), 5 cases in Gr B (GCS 6 to 9) and 2 cases were in Gr C (GCS 10 to 15). So 12.5% cases presented in a clinically severe state (GCS 3 to 5) during admission. 14 cases (28%) were diagnosed as tuberculous meningitis [6 males (42.86%) and 8 females (57.14%)] 6 cases were in Gr A (GCS 3 to 5), 6 cases in Gr B (GCS 6 to 9) and 2 cases were in Gr C (GCS 10 to 15). 42.86% cases presented in a clinically poor condition (GCS 3 to 5) on admission. (See table 5 & Chart 5) 28 cases (56%) were diagnosed as viral meningoencephalitis. [7 males (25%) and 21 females (75%)] 8 cases were in Gr A (GCS 3 to 5), 11 cases in Gr B (GCS 6 to 9) and 9 cases were in Gr. C (GCS 10 to 15) 28.57% cases presented in a clinically poor state (GCS 3 to 5) on admission.

Classification according to gcs in different etiological groups: (See Figure 5)

After comparative analysis of three etiological groups and their GCS categories it was found that tuberculous meningitis cases (42.8%) presented in a more severe state followed by cases of viral meningoencephalitis (28.57%) and pyogenic meningitis (12.5%).

CSF Findings

Cytology: Cell Count & Cell Type: (See Figure 5 & Chart 7 & 8)

In cases that were later diagnosed as pyogenic meningitis, the cell counts ranged from 120 to 5900 cells/cumm with a mean count of 950.5± 3748.84 cells/cumm and neutrophil predominance. In cases with a final diagnosis of tuberculous meningitis, the cell counts ranged from 60 to 460 cells/cumm with a mean count of 225.86± 217.22 cells/cumm with a lymphocyte predomi-
nance. In cases categorized as viral meningoencephalitis, the cell counts ranged from 30 to 510 cells/μm with a mean count of 128.607±276.86 cells/μm and a predominantly lymphocytic picture. **Biochemistry & ADA levels:** (See Figure 6 & Chart 9) Pyogenic meningitis cases revealed a mean CSF glucose of 19±18.9 mg/dL, mean protein levels of 62.375±48.52 mg/dL, mean chloride levels of 112.88±12.48 meq/L with mean ADA level of 8.025±24.14. This picture was consistent with hypoglycorrachias expected in pyogenic meningitis. Tuberculous meningitis cases revealed a mean CSF glucose of 35.79±22.9 mg/dL, mean protein levels of 268.86±368.12 mg/dL, mean chloride levels of n112.714±12.54 meq/L with mean ADA level of 16.29±16.14 U/L. Protein levels were markedly increased with ADA levels also higher than normal range along with low glucose levels consistent with the diagnosis. Viral meningoencephalitis cases revealed a mean CSF glucose of 60.11±32.28 mg/dL, mean protein levels of 96.96±91.98 mg/dL, mean chloride levels of 113.45±14.32 meq/L with mean ADA level of 3.76±4.38 U/L. The protein levels were higher with normal glucose levels.

**CSF Culture, staining and viral PCR:** According to affordability of the patient and availability of appropriate facilities in Eastern India - CSF culture and staining, CSF viral antibody detection by ELISA (when suspected) were sent with the routine samples. Some cases were diagnosed accurately from CSF culture and ELISA that were corroborating with the clinical assumptions and radiological imaging findings.

**Pyogenic Meningitis etiologies:** (N=8): (See Figure 7 & Chart 10) CSF gram staining and culture revealed Streptococcus pneumoniae in 4 (50%) cases, Neisseria meningitides in 1 (12.5%) case, Staphylococcus aureus in 1 (12.5%) case and culture negative in 2 (25%) cases. This shows that majority of the culture positive cases were for Streptococcus pneumonia followed by equal incidence of Neisseria meningeitides and Staphylococcus aureus. 25 % cases that were culture negative had classical symptoms which resolved with empiric antibiotics.

**Tuberculous Meningitis:** (N=14) All the cases suspicious of tuberculous etiology were ZN smear negative. CSF culture was negative in 10 cases (71.4%) while culture could not be done in 4 cases (28.6%). Diagnosis had to be established on clinical features, CSF cytology, CSF biochemistry, brain imaging and therapeutic response to anti tubercular drugs.

**Viral Meningoencephalitis:** (N=28): (See Figure 8 & Chart 11) Patients with CSF cytology indicating a viral infection revealed the following viral etiologies in CSF Elisa (IgG). 10 cases (35.72%) were positive for Herpes simplex, 3 cases (10.72%) were positive for Varicella zoster, 4 cases (14.28%) were positive for Japanese B while 11 cases (39.28%) were negative or indeterminate.

**DISCUSSION**

**Pyogenic Meningitis** Acute bacterial meningitis is more common in resource-poor than resource-rich settings. Survival is dependent on rapid diagnosis and early treatment, both of which are difficult to achieve when laboratory support and antibiotics are scarce. Syamal Modi and Amit Kumar Anand in their study Phenotypic Characterization and Anti-biogram of CSF Isolates in Acute Bacterial Meningitis done in a tertiary care hospital Patna (India) found that 62.3% patients were males and 37.7% were females. The gender distribution and male preponderance in disease incidence was also marked (62.5 % males and 37.5% females) in our study done in Eastern India. In our study sample size was smaller in comparison to theirs. This male predilection reported in several previous studies. [17] Similar to the study by Marjolein J. Lucas, Matthijs C. Brouwer et al (2014),[18] patients using immunosuppressive drugs and those with asplenia, diabetes mellitus, alcoholism, or infection with immunodeficiency virus were considered immune compromised and excluded from our study. Schutte CM, van der Meyden CH in their study found patients with a Glasgow coma scale (GCS) value of > 12 had a good neurological outcome, while those with a GCS value of ≤ 8 had a poor outcome. They concluded that the GCS value was a better prognostic indicator than high CSF protein levels[8]. In our study we adapted GCS as the criteria for clinical severity. Group A (GCS 3 to 5) was considered as most severe clinical category. Pyogenic meningitis can be accurately and rapidly diagnosed by gram staining. Some studies have reported sensitivities of 60-90% and specificities of >97% of CSF gram staining in the diagnosis of ABM.[6] In our study: 50% cases were diagnosed as Streptococcus pneumoniae induced meningitis, followed by equal incidence of Neisseria meningitides and Staphylococcus aureus cases (12.5% each) on CSF gram staining and culture 25 percentage cases did not reveal any organism on CSF culture. Negative CSF cultures are estimated to occur in11%–20% of patients with bacterial meningitis.[19][20] However, the clinical presentation of patients with culture-positive bacterial meningitis and patients with culture-negative bacterial meningitis was reported to be similar. R Mani, S Pradhan et al in their study conducted in South India (NIMHANS), found that streptococcus pneumoniae was the most common etiological agent of community-acquired meningitis in all age groups. This
accounting for 238 (61.8%) cases in their study, reflecting
a similar trend reported in an earlier study from their insti-
tute (1978-1988). Most Indian studies have also reported
a high incidence of pneumococcal meningitis in our study,
also Streptococcus pneumonia was the most common
organism isolated from CSF culture (50% cases). Analysis
of the CSF is essential, and simple techniques can enhance
the yield of diagnostic microbiology. Penicillin-resistant and
chloramphenicol-resistant bacteria are a considerable threat
in resource-poor settings that go undetected if CSF and blood
cannot be cultured.

The future rests with the provision of effective conjugate
vaccines against S pneumoniae, Haemophilus influenzae,
and Neisseria meningitides to children in the poorest regions
of the World.

Viral Meningoencephalitis

Herpes simplex virus type 1 (HSV-1) is the most common
cause of sporadic encephalitis (Davison et al., 2003; Mailles
e et al., 2007). In some recent reports from Scandinavia and
central Europe, however, varicella zoster has been identified
as the most common viral agent responsible for encephalitis
(Cizman & Jazbec, 1993; Studahl et al., 1998; Koskiniemi
et al., 2001). A study named - Viral aetiology and clini-
co-epidemiological features of acute encephalitis syndrome
in eastern India conducted by Rathore SK, Dwibedi B et
al during April 2011 to July 2012. Blood and CSF samples
of 526 AES cases were investigated by serology and/
or PCR. Viral aetiology was identified in 91 (17.2%) cases.
Herpes simplex virus (HSV; types I or II) was most common
(16.1%). Simultaneous infection of HSV I and measles
was observed in seven cases. This report provides the first
evidence on viral aetiology of Acute encephalitis syndrome
viruses from eastern India showing dominance of HSV that
will be useful in providing the public health system. In our
study there were majority of HSV cases. In CSF Elisa (IgG).
10 cases (35.72%) were positive for Herpes simplex, 3 cases
(10.72%) were positive for Varicella zoster, 4 cases (14.28%)
were positive for Japanese B while 11 cases (39.28%) were
negative or indeterminate. So there is a similarity and etio-
logical preponderance of HSV in both western and eastern
India.

Panagariya A, Jain RS et al in a study conducted in North
West India with Herpes simplex encephalitis cases included
patients admitted with provisional diagnosis of an enceph-
alitic illness over a period of 30 months. Special investiga-
tions included CSF analysis, EEG, CT scan and MRI. Herpes
simplex virus (HSV) antibody estimation in CSF and blood
was done simultaneously using ELISA. 28 patients showed
electroencephalographic, serologic and/or neuro radiological
evidence of herpes simplex encephalitis. Exact incidence of
this disease (HSE) is the most difficult to estimate, because
only few patients with common cause of fatal sporadic acute
encephalitis with severe disease report to hospital whereas
mild and self limiting cases usually go unrecognised. In In-
dia, HSE appears to be under diagnosed; probably due to
lack of awareness and diagnostic facilities.CSF polymerase
chain reaction (PCR) and immuno-cytochemistry could not
be done in their study because of non-availability. CSF PCR
is not always available in laboratories in eastern part of In-
dia, due to which it could not be done in our study. Moreo-
ver affordability was also a point of concern in case of pa-
tients attending our hospital. Japanese encephalitis (JE) is
the leading cause of encephalitis in Southeast Asia, where
30,000--50,000 cases are recorded annually (Tsai, 1997).

The World Health Organisation estimated nearly 14,000
deaths due to JE in the year 2002. Of these, 8,500 occurred
in Southeast Asia, 3,000 in the western Pacific region and about
2,000 in the eastern Mediterranean region.

The incidence of neurologic complications associated with
varicella is estimated to be 1–3 per 10,000 cases. The cen-
tral nervous system (CNS) manifestations that occur most
frequently with varicella are cerebellar ataxia and encepha-
litis. The most serious CNS complication of varicella,
has an incidence of 1–2 episodes per 10,000 varicella cases,
with the highest incidence in adults and infants. The CSF
findings are usually abnormal with elevated opening pres-
sure, a mild-to-moderate lymphocytic pleocytosis (usually
100 cells/µL), mildly elevated protein (50–100 mg/dL),
and normal glucose levels. Dengue encephalopathy is a well-
recognized and common entity, the incidence ranging from
0.5 to 6.2 %. Dengue is not classically a neurotropic virus,
although there is recent evidence of direct neuronal injury.
Dengue encephalitis must be thought of in differentials of
encephalopathy, in patients with dengue. We found 3 such
cases in our study with feature of encephalopathy who were
diagnosed cases of IgM positive dengue with CSF analysis
indicating a viral etiology and a normal MRI. Dengue specif-
ic IgM antibody (ELISA) was negative in 2 and could not
be done in 1 case. In light of our knowledge regarding dengue
encephalopathy a negative CSF antibody cannot refute its
presence. So whether the encephalopathy was due to some
other viral pathogen or as consequence of dengue could not
be confirmed. All 3 patients survived without any residual
neuro deficit or CNS complications. Infection of the CNS
with the measles virus (MV) may result in 1) acute post in-
fec tious encephalitis, 2) acute progressive encephalitis, and
3) SSPE. Data about imaging findings in acute measles ence-
phalitis are sparse. T2WI may reveal cortical edema and
bilateral symmetric hyper intense lesions within the putamen
and caudate nuclei as well as within the centrum semiovale.

Sometimes patients also present bilateral thalamic lesions
and signal abnormalities within the corpus callosum. We
found 2 cases with a recent history of measles who subse-
quently developed features of encephalopathy. The disease
was self limiting and their MRI were normal.
We would sum up in accordance to I. Steiner, H. Budka et al and their review of diagnosis and management recommendations in Viral encephalitis. A holistic approach to diagnosis should be based on medical history, examination; followed by analysis of cerebrospinal fluid for protein and glucose contents, cellular analysis and identification of the pathogen by polymerase chain reaction (PCR) amplification (recommendation level A) and serology (recommendation level B). Lumbar puncture can follow neuroimaging when immediately available, but if this cannot be obtained at the shortest span of time it should be delayed only in the presence of strict contraindications. Brain biopsy should be reserved only for unusual and diagnostically difficult cases. All encephalitis cases must be hospitalized with an access to intensive care units. Supportive therapy is an important basis of management. Specific, evidence-based, anti-viral therapy, acyclovir, is available for herpes encephalitis (recommendation level A). Acyclovir might also be effective for varicella-zoster virus encephalitis, gancyclovir and foscarnet for cytomegalovirus encephalitis.

**Tuberculous Meningitis**

J Kalita, UK Misra in their study evaluated the clinical and radiological outcome of tuberculous meningitis (TBM) patients. In this study most of the patients were females who were anemic. MRI revealed hydrocephalus, exudates, infarction and multiple granuloma and the majority of the patients improved following ant tubercular therapy. In our study there was a female predilection (57.14%). Under nutrition and anaemia may be a result or risk factor for development of tuberculous meningitis. Adults with tuberculous meningitis (TBM) can often present with the classic meningitis symptoms of fever, headache and stiff neck along with focal neurological deficits, behavioural changes, and alterations in consciousness. The presence of active pulmonary tuberculosis on chest X-ray ranges from 30 to 50%. TBM may have an acute presentation. The duration of presenting symptoms may vary from 1 day to 9 months, although several cases may present with symptoms of less than 2 weeks duration. In our cases patients had a history of less than 2 weeks duration prior to admission. Cerebrovascular complications of tuberculous meningitis occur typically as multiple or bilateral lesions in the territories of the middle cerebral artery perforating vessels are termed as tuberculous vasculopathy. Vessel pathology appears to be a consequence of its immersion in the local inflammatory exudates. Infiltrative, proliferative and necrotising vessel pathologies have been described, leading to luminal thrombosis. There is some evidence that vasospasm may mediate strokes early in the course of the disease and proliferative intimal disease later strokes. In this study we encountered 28.57% cases with cerebral infarcts.

**SUMMARY**

This observational study was done in a tertiary care hospital of Eastern India. 50 consecutive hospital admitted patients (>12 yrs) fulfilling the inclusion criteria were randomly selected over a period of 19 months. A male preponderance was marked in Pyogenic meningitis cases while a female preponderance was noted in tuberculous meningitis and viral meningo encephalitis cases.

An increased propensity of disease occurrence was seen in age groups 31-40 yrs in pyogenic meningitis and younger age groups (12-30 yrs) were more affected in tuberculous and viral meningoencephalitis. Individual patients were assessed by clinical status on admission and divided into three GCS group. 15 cases (30%) belonged to group A (GCS 3 to 5), 22 cases (44%) belonged to group B (GCS 6 to 9) while 13 cases (26%) belonged to group C (GCS 10 to 15) on admission. 18 cases (36%) were male and 32 cases (64%) were female. Patient’s ages ranging from 12 yrs to 77 yrs were selected and divided into age groups. CSF study and MRI brain was done and etiologically the cases were reclassified. Out of 50 cases 8 cases (16%) were diagnosed as pyogenic meningitis [5 males (62.5%) and 3 females (37.5%)], 14 cases (28%) were diagnosed as tuberculous meningitis [6 males (42.86%) and 8 females (57.14%)] and 28 cases (56%) were diagnosed as viral meningoencephalitis [7 males (25%) and 21 females (75%)]. CSF study revealed neutrophilic picture with hypoglycorrhachia in bacterial meningitis with CSF culture studies revealing Streptococcus pneumoniae as the most commonly isolated pathogen (30% cases). Tuberculous meningitis revealed CSF lymphocytic pleocytosis with significantly increased protein and ADA levels. CSF ZN staining in all cases were negative, CSF culture for tuberculosis was negative in 71.4% cases and could not be done in 28.6% cases although the diagnosis was established on other parameters and imaging evidence.

The CSF samples of viral meningoencephalitis revealed lymphocytic picture and increased protein with majority of cases being positive for Herpes simplex antibody (in CSF by ELISA method) in 35.72%. 14.28% cases were positive for Japanese B antibody. 10.72% cases were diagnosed as Varicella zoster cases. Serological confirmation of diagnosis was not possible in other cases due to non-availability of the specific test (Viral PCR) or a negative serology in 39.28%.

**CONCLUSION**

In this Eastern India based study viral etiologies were more frequently detected (56%) followed by tubercular (28%) and pyogenic (16%) causes of disease. There was an overall female preponderance (64%) with maximum number of patients belonging to younger age groups. Male predilection...
in pyogenic meningitis (62.5%) and a female predilection (57.14%) in tuberculosis meningitis was noted, which was similar to several other studies done in other parts of the world. Most of the tuberculous meningitis cases (42.8%) were clinically more severe according to the GCS category during admission followed by viral meningoencephalitis (28.57%) and pyogenic meningitis (12.5%). CSF cytology revealed neutrophillic picture in pyogenic and lymphocytic pleocytosis in tuberculous and viral meningoencephalitis. CSF biochemistry revealed hypoglycorrhachia in pyogenic and tuberculous meningitis and high protein levels in viral and grossly high protein content in tuberculous meningitis cases. ADA levels were also high in tuberculous meningitis. CSF gram staining, culture and serology results showed Streptococcus pneumoniae as the most common pathogen causing pyogenic meningitis (50%) and Herpes simplex as most common viral pathogen (35.72%) causing meningoencephalitis. This is a potential area of research and rightfully demands attention in the near future. India being a resource poor nation, improvisation of the specific diagnostic modalities and implementation with regard to affordability should be prioritised. Especially in this part of the world, there is a dearth of multicentric prospective studies on meningoencephalitis. More studies should be conducted based on correlation of clinical aspects with brain imaging; prognostication and taking into account the long term outcomes.

Limitations of the study
Sample size is small (N= 50). This study was done over a certain catchment area, hence not a multicentric study. As the study was done in a tertiary care hospital, the mean values may not properly reflect the actual population mean. Due to small sample size and wide range of variation in the values of the CSF parameters, in certain cases the standard deviation exceeded the mean values. CSF PCR could not be done due to non-availability during that period. Empiric therapy had to be started in most of the cases prior to confirmation of diagnosis, for the sake of the patients. Some of the patients who fulfilled the inclusion criteria, had to be excluded due to economic constraints.

Conflict of interest- Authors did not have any conflict of interest.

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**Chart 6**: Clinical Group according to GCS on admission in Pyogenic Meningitis.

**Chart 7**: Mean CSF Total Cell.

**Chart 8**: CSF cell type in etiological group.

**Chart 9**: CSF BIOCHEMISTRY and ADA levels.

**Chart 10**: Organism isolated in CSF culture in pyogenic meningitis.

**Chart 11**: Viral meningoencephalitis according to CSF serology
### Table 1: Distribution According to Age Groups

| Age Groups (In Years) | Number of Patients (N=50) |
|-----------------------|--------------------------|
| 12 to 20              | 13                       |
| 21 to 30              | 12                       |
| 31 to 40              | 12                       |
| 41 to 50              | 07                       |
| 51 to 60              | 05                       |
| 61 to 70              | 0                        |
| >71                   | 01                       |

### Table 2: Gender Distribution in Etiological Groups and Percentages

| Etiology                   | Total Numbr | Males (N=18) | Females (N=32) | Percentage (%) |
|----------------------------|-------------|--------------|----------------|----------------|
| Pyogenic Meningitis        | 8           | 5            | 3              | Males- 62.5    |
| Tuberculous Meningitis     | 14          | 6            | 8              | Males- 42.86   |
| Viral Meningoencephalitis  | 28          | 7            | 21             | Males- 25      |

### Table 3: Distribution According to Age Groups

| Etiology                   | Age Groups of Patients (In Years) | Number of Patients | Mean Age | Percentage (%) |
|----------------------------|----------------------------------|--------------------|----------|----------------|
| Pyogenic Meningitis        | 12 to 20                         | 0                  | -        | -              |
|                            | 21 to 30                         | 1                  | 25       | 12.5           |
|                            | 31 to 40                         | 3                  | 37       | 37.5           |
|                            | 41 to 50                         | 2                  | 44       | 25             |
|                            | 51 to 60                         | 2                  | 52       | 25             |
|                            | 61 to 70                         | 0                  | -        | -              |
|                            | > 70                             | 0                  | -        | -              |
| Tuberculous Meningitis     | 12 to 20                         | 05                 | 16.8     | 35.71          |
|                            | 21 to 30                         | 04                 | 27       | 28.57          |
|                            | 31 to 40                         | 02                 | 38       | 14.28          |
|                            | 41 to 50                         | 02                 | 44.5     | 14.28          |
|                            | 51 to 60                         | 01                 | 53       | 07.1           |
|                            | 61 to 70                         | 0                  | -        | -              |
|                            | > 70                             | 0                  | -        | -              |
| Viral Meningoencephalitis  | 12 to 20                         | 08                 | 15.63    | 28.57          |
|                            | 21 to 30                         | 07                 | 25       | 25             |
|                            | 31 to 40                         | 07                 | 36       | 25             |
|                            | 41 to 50                         | 03                 | 44.33    | 10.71          |
|                            | 51 to 60                         | 02                 | 54       | 07.41          |
|                            | 61 to 70                         | 0                  | -        | -              |
|                            | > 70                             | 01                 | 77       | 03.57          |
### Table 4: Classification According to GCS in Different Etiological Groups

| Clinical groups (glasgow coma scale/ gcs) | Number of patients according to clinical severity on admission | Pyogenic Meningitis (n=8) | Tuberculous Meningitis (n=14) | Viral Meningoencephalitis (n=28) |
|------------------------------------------|---------------------------------------------------------------|--------------------------|-------------------------------|---------------------------------|
| Group - A(GCS: 3 to 5)                   | 15                                                           | 1                        | 6                             | 8                               |
| Group – B(GCS: 6 to 9)                   | 22                                                           | 5                        | 6                             | 11                              |
| Group – C(GCS: 10 to 15)                 | 13                                                           | 2                        | 2                             | 9                               |

### Table 5: CSF Cytology Comparison in Etiological Groups

| Aetiological group                  | CSF - total cell count (mean) | Neutrophils (mean) | Lymphocytes (mean) |
|-------------------------------------|------------------------------|--------------------|--------------------|
| Pyogenic meningitis                | 950.5 ± 3748.84              | 85.88 ± 25.18      | 14.13 ± 25.18      |
| Tuberculous meningitis             | 225.86 ± 217.22              | 13.14 ± 32.8       | 86.86 ± 32.8       |
| Viral meningoencephalitis          | 128.61 ± 276.86              | 14.93 ± 25.14      | 85.07 ± 25.14      |

### Table 6: CSF Biochemistry and ADA Levels in Different Etiological Groups

| Etiological Group                 | CSF Glucose (Mean) | CSF Protein (Mean) | CSF Chloride (Mean) | CSF ADA Levels Mean |
|-----------------------------------|--------------------|--------------------|---------------------|---------------------|
| Pyogenic Meningitis               | 19 ± 18.9          | 62.375 ± 48.52     | 112.88 ± 12.48      | 8.03 ± 24.14        |
| Tuberculous Meningitis            | 35.79 ± 22.9       | 268.86 ± 368.12    | 112.71 ± 12.54      | 16.29 ± 16.14       |
| Viral Meningoencephalitis         | 60.11 ± 32.28      | 96.96 ± 91.98      | 113.45 ± 14.32      | 3.76 ± 4.38         |

### Table 7: Organisms Isolated in CSF Culture of Pyogenic Meningitis Cases

| Sl. No. | Organisms isolated in CSF culture in Pyogenic Meningitis | Number of case (N=8) | Percentage (%) |
|---------|----------------------------------------------------------|----------------------|----------------|
| 1.      | Streptococcus pneumoniae                                  | 4                    | 50             |
| 2.      | Neisseria meningitidis                                    | 1                    | 12.5           |
| 3.      | Staphylococcus aureus                                     | 1                    | 12.5           |
| 4.      | Undetermined/Negative                                     | 2                    | 25             |

### Table 8: Confirmed viral Aetiology

| Etiology confirmed on CSF ELISA | Number of patients (N=28) | Percentage (%) |
|---------------------------------|----------------------------|----------------|
| Herpes Simplex                  | 10                         | 35.72          |
| Japanese B                      | 04                         | 14.28          |
| Varicella Zoster                | 03                         | 10.72          |
| Negative/Indeterminate          | 11                         | 39.28          |