Acute flaccid paralysis (AFP), as defined by Global Polio Eradication Initiative (GPEI), is sudden onset of weakness and floppiness in any part of the body in a child less than 15 years of age or when clinician suspects polio in any person of any age (excludes adults, spastic paralysis, old cases or cases with obvious causes like trauma). The term "flaccid" denotes the absence of spasticity or other signs of disordered central nervous system motor tracts such as hyperreflexia, clonus, or extensor plantar responses.

Amongst the various conditions leading to AFP, poliomyelitis has been an important differential. Since the Global Polio Eradication Initiative by WHO in 1988, Surveillance for AFP is an important public health activity in many countries and polio cases have decreased by over 99%. With WHO’s impressive progress, poliomyelitis is nearing its eradication in the world, other causes of AFP like Acute Encephalitis Syndrome (AES) also need to be monitored.

Guillain-Barre syndrome (GBS), transverse myelitis, traumatic paralysis, Vaccine Associated Paralytic Poliomyelitis (VAPP), West Nile virus and other enteroviruses, neurotropic viruses like Rabies, Varicella-zoster, and Japanese Encephalitis have gained importance. Similarly, critical illnesses like polyneuropathy, metabolic disorder like hypokalemic periodic paralysis, polymyositis, neuropathies and neuromuscular junction disorders may also present with AFP. WHO defines Acute Encephalitis Syndrome (AES) as a person of any age, in any geographical region, at any time of year with the acute onset of fever and a change in mental status (including symptoms such as confusion, disorientation, coma, or inability to talk) and/or new onset of seizures (excluding simple febrile seizures). Frequently, a child with AES may have AFP which also should be investigated and hence AES surveillance has now been integrated with AFP surveillance so that not a single case of acute flaccid paralysis goes unreported and thus helps in polio eradication campaign.

AFP can have varied presentation depending upon the underlying cause. Furthermore, the outcome depends on the severity at presentation and underlying cause. A detail evaluation of child’s clinical presentation and thereby proper and timely treatment directly affects the outcome. So, all the differentials of AFP need to be evaluated and AFP with AES should also be considered. Hence, this study was done to study the clinical features and outcome of AFP with AES in our part.

**MATERIALS AND METHODS**

It was a hospital-based prospective study done over one year (Jan 2011-Dec 2011) in the Department of Paediatrics & Adolescent Medicine, BPKIHS, Dharan. Any child of age ≤15 years presenting with AFP in any form alongwith AES as per WHO definition was included. Children of age >15 yrs , spastic paralysis, old cases or cases with obvious causes like trauma, obvious congenital anomalies or birth defects, AES without acute flaccid paralysis were excluded.

After taking informed consent from caregiver, all children meeting inclusion criteria were enrolled and detail history and clinical examination recorded. Relevant laboratory investigations were performed and stool samples sent for examination as per AFP surveillance guidelines. After instituting treatment, the patients were observed for any complications or sequelae. Regular follow-ups were done after discharge in Paediatrics OPD at 15 days/30 days/60 days along with the help of surveillance team. The stool sample reports were collected from the IPD field office and final classification made according to clinico-laboratory results and subsequent follow-up findings. The acquired data for all children were entered in SPSS Statistics software version 17 for analysis. Descriptive statistics were used and final inferences made.

**RESULTS**

Out of 3432 children admitted during the one-year study period (Jan 2011 to Dec 2011), 43 (1.25%) children had acute flaccid paralysis. The final classification showed Acute Encephalitis Syndrome (AES) as the most common cause in 23(53.5%) children.

Amongst the 23 children of AFP with AES, majority, ie 17 (74%) had high grade fever. Altered mental status was present in 20 (87%) children while seizures in 17 (74%) children. Other clinical features like headache and vomiting were present in 9 (39.1%) patients each while neck pain was found in 3 (13%).

On clinical examination, the mean GCS was 7.61±3.65. One or more signs of meningeal irritation were found in 13% children. The pattern of weakness in majority, i.e. 20 children of AES with AFP (87%), was quadriparesis while paraparesis, hemiparesis and monoparesis were found in 1 each. Cranial nerve involvement was seen in 2 (8.7%) children, having bilateral sixth nerve involvement. The deep tendon reflexes were absent or sluggish in 20 (87%). Sensory and bowel/bladder involvement were found in 2 (8.7%) and 8 (34.8%) respectively. Concurrent underlying paleness and undernutrition were detected in 8 (34.8%) and 16 (69.6%) children respectively. Although lumbar puncture for cerebrospinal fluid (CSF) analysis could not be done in 3 (13%) patients due to hemodynamic instability, 11 (47.9%) children had abnormal CSF findings with pyogenic meningitis in 2 (8.7%) and viral in rest. Subsequent serology for Japanese encephalitis was positive in 2 (8.7%) and malaria screening was positive in 1 (4.3%). The complications observed and treatment details of the AES children presenting with AFP are shown in table 1.

**DISCUSSION**

Acute flaccid paralysis is caused by various diseases, poliomyelitis being one of the most important causes. Nepal had re-emergence of polio
after four years in 2005 and few more imported cases were detected since then. Acute Encephalitis Syndrome surveillance has also been integrated with AFP surveillance now, so as not to miss any floppy child with AFP. This study was carried out to study the children presenting with AFP in our part and analyse the cases of AES amongst them. In our study, the majority 23 (53.5%) children having AES whereas GBS is the main cause of AFP in previous studies. Interestingly, sciatic nerve palsy accounted for majority (72.0%) of the AFP in a five-year review of cases by Hamzat et al. Tu YF et al found that three of 186 paediatric patients with acute bacterial meningitis presented with AFP due to myelopathy. In a study from Vietnam, 55% of patients identified with AFP were actually later diagnosed with JE. In a previous study of AES at BPKIHS, hypotonia was seen in 25% of herpetic encephalitis, 7.69% of non-JE encephalitis, 6.66% of meningitis and 5.26% of Japanese Encephalitis. Decreased power was found in 35% of the acute encephalopathies. Most of the children, ie 17 (74%) had high grade fever. Altered mental status was present in 20 (87%) children while seizures in 17 (84%) children. Similarly, fever was the most common symptom along with neck rigidity, convulsions, abnormal behavior and seizures in earlier studies. Fever was nearly universal (92%) in a study of AFP related to West Nile Virus by Moorman J et al. Rayamajhi A et al had found fever (>3 days) in 69%; altered sensorium in 50%; and seizures in 58% of acute febrile encephalopathy. The mean GCS in our study was 7.61±3.65. More than half (65.2%) of the children of AFP with AES had GCS in the range of three to eight. In a study of encephalitis by CR Kennedy et al, the modified GCS score fell below 11 in 52% and below six in 12% patients. Similar findings were also observed in a retrospective study of acute encephalitis syndrome at BPKIHS. In contrast the median GCS 13 (IQR 10 to 14) with a GCS of <8 in 10 (6.7%) children in a study from Papua New Guinea. The pattern of weakness in majority (87%) was quadriparesis and cranial nerve involvement was seen in two (8.7%) children, having bilateral sixth nerve involvement. Singh RR et al found neurological deficit in 9.34%- mostly 7th nerve. Amongst 11 (47.9%) children with abnormal CSF findings, there was pyogenic meningitis in two (8.7%) and viral meningoencephalitis in rest, of which two had positive JE serology. Similarly, viral encephalitis (46%) with JE (40%) was the most common cause in the study by Singh RR et al. In the study by Karmarkar SA et al, pyogenic meningitis was the most frequent diagnosis (33.8 %) while 37.3% were suspected as viral encephalitis- (35.1 %), mumps (10.5%), Japanese encephalitis (8.7%), and measles (7%) cases. Anga G et al confirmed S. pneumoniae and H. influenzae as major causes of febrile encephalopathy in children. A study of acute encephalopathy by Kumar et al showed pyogenic meningitis and JE to be responsible for 18% and 12% of cases, respectively. Khinchi R.et al found JE in 18% of AES cases with a case fatality of 16.6%. As most patients were referred after initial antibiotics, the yield of pyogenic meningitis could have decreased and not all cases of AES but only those with AFP were enrolled, the proportion of JE could have lowered. Also, there has been decrease in overall cases of JE over last few years after JE vaccination campaigns and inclusion of JE vaccine in EPI of Nepal. Although three (13%) children expired before collection of stool samples, AFP surveillance was carried out in the rest 20 children, amongst which no virus was isolated and thus it strengthened the

Table 1: Treatment details & complications in children of AFP with AES (n=23)

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>No. of patients (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>ICU requirement</td>
<td>14 (60.9)</td>
</tr>
<tr>
<td>Mechanical ventilation</td>
<td>11 (47.8)</td>
</tr>
<tr>
<td>Steroids</td>
<td>7 (30.4)</td>
</tr>
<tr>
<td>Osmotic diuretic</td>
<td>12 (52.1)</td>
</tr>
<tr>
<td>Inotropes</td>
<td>19 (82.6)</td>
</tr>
<tr>
<td>Complications</td>
<td></td>
</tr>
<tr>
<td>Aspiration pneumonitis</td>
<td>14 (60.9)</td>
</tr>
<tr>
<td>Respiratory failure</td>
<td>12 (52.1)</td>
</tr>
<tr>
<td>Autonomic dysfunction</td>
<td>8 (34.8)</td>
</tr>
<tr>
<td>Acute renal failure</td>
<td>4 (17.4)</td>
</tr>
<tr>
<td>Death by 60-day follow-up</td>
<td>11 (47.9)</td>
</tr>
<tr>
<td>Residual weakness</td>
<td>2 (8.7)</td>
</tr>
</tbody>
</table>
AFP surveillance. There are no published data of studies of AES with inclusion of stool examination as per AFP surveillance guidelines till now. Inclusion of AES in AFP surveillance has been initiated in Nepal since 2004 so as no case of AFP gets missed.

Out of 23 children of AFP with AES, 11 (47.9%) expired while two (8.7%) had residual paralysis. In contrast to this, Karmarkar SA et al found significant neurological sequelae in 33.33% of acute febrile encephalopathy syndrome. In a retrospective study by Rayamajhi A et al., viral encephalitis had poorer outcome than bacterial (31% versus 13%) and JE worse than AES of unknown viral aetiology (48% versus 24%).

CONCLUSION

AES is one of the common causes of AFP besides other causes. AFP with AES is commonly associated with quadriaparesis, low GCS, respiratory complications, neurological sequelae and a high mortality. Thus, this study stresses upon the importance of considering AES surveillance in all cases of AFP so as not a single case of AFP gets missed and thereby AFP with AES gets better diagnosis and management.

REFERENCES