Superimposed *Propionibacterium Acnes* Subdural Empyema in a Patient with Chronic Subdural Hematoma

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The authors present a case of subdural empyema in a macrocephalic patient. A 23-year-old male was admitted due to headache and fever. One month ago, he had mild head injury by his coworkers. Physical examination showed a macrocephaly and laboratory findings suggested purulent meningitis. Neuroimaging studies revealed a huge size of epidural space-occupying lesion. Under the impression of epidural abscess, operation was performed. Eventually, the lesion was located at subdural space and was proven to be subdural empyema. Later, histological examination of the specimen obtained by surgery demonstrated findings consistent with the capsule of the chronic subdural hematoma. Two weeks after operation, *Propionibacterium acnes* was isolated. The intravenous antibiotics were used for total of eight weeks under monitoring of the serum level of the C-reactive protein. Follow-up brain computed tomography (CT) scan showed the presence of significant amount of remaining subdural lesion. However, he has complained of minimal discomfort. It is suggested that the subdural empyema occurred with preexisting chronic subdural hematoma after head injury about one month prior to admission and it took a long time to treat *Propionibacterium acnes* subdural empyema with systemic antibiotics, at least over eight weeks.

**KEY WORDS**: Subdural empyema • Chronic subdural hematoma • *Propionibacterium acnes* • C-reactive protein.
mg/dL, respectively; the CSF protein concentrations was over the 300 mg/dL. The CSF Gram’s stain was negative, and culture remained sterile. A complete blood count revealed white blood cell count of 19,510 cells/mm with neutrophils 89%, a C-reactive protein (CRP) concentration of 205 mg/dL. Simple X-ray of skull showed macrocephaly and multiple calcifications at frontal area (Fig. 1). A precontrast computerized tomography (CT) scan demonstrated a lentiform-shaped high density lesion confined over left frontal convexity bordered with underlying cortex by a thick hyperdense calcified rim and another separated large left hemispheric isodense space-occupying lesion surrounded by hyperdense border with multiple calcifications (Fig. 2A, B). Under the impression of epidural abscess associated with pyogenic meningitis, he was started on intravenous antibiotics such as cefotaxime and gentamycin empirically and surgical decompression was planned. While waiting for operation, the patient’s level of consciousness deteriorated (Glasgow Coma Scale score of 6) and his left pupil was dilated and not reactive to light. The patient underwent urgent surgery via a left craniotomy. There were no abnormal findings in epidural space and the dura was very tight. A thick, hard, and yellowish capsule was identified during dural opening and this was firmly adhered to overlying dura. As soon as the capsule was incised and dissected, huge amount of purulent materials was gushed out. Also, inner capsule of the lesion was firmly adhered to underlying cortex and couldn’t be removed completely. A drain catheter was placed in the subdural space. Histological examination of the removed outer capsule revealed inflammatory changes with prominent neutrophil infiltration (Fig. 3A, B). Immediate postoperative CT scan showed preoperative midline shifting was improved but sizable amount of empyema still remained and occurrence of new contralateral large amount of EDH (Fig. 4A, B). Postoperative neurological examination revealed decrease in left pupil diameter and sluggish response to light but he was stuporous. Postoperative antibiotic therapy consisted with cefotaxime, vancomycin, amikacin, and clindamycin with injection of vancomycin into the subdural space. There were no bacteria found in the culture of the blood, urine, and sputum. There was no evidence of any other source of infection, including otorhinologic or gastrointestinal disease. Two weeks after operation, P. acnes was isolated from the purulent materials and the CRP concentration was 154 mg/dL. The antibiotic regimen was changed to vancomycin and imipenem for next six weeks. Thereafter, follow-up results of CRP were 98 mg/dL, 64 mg/dL, 27 mg/dL, 16 mg/dL, and <10 mg/dL, respectively third, fourth, fifth, sixth, and since seventh week. The patient was discharged four months after operation with both lower extremity weaknesses. One year after operation, follow-up brain CT scan showed large amount of subdural empyema still remained (Fig. 5), but he complained of minimal discomfort.

DISCUSSION

Our macrocephalic patient had a huge amount of subdural empyema surrounded by thick capsule. Although P. acnes was isolated from culture of
the purulent materials and it is reported that the clinical course of this infection was indolent, we thought that the amount of the subdural empyema was too much to be tolerable for a long time, and probably there had been another pathology within intracranial cavity. The chronic subdural hematoma must be differentiated from the causes of childhood macrocephaly. If undiagnosed and untreated, as a result of a gradual growth of the hematomas, it is possible to occur with atrophy in underlying cerebral hemisphere and to be accompanied by macrocephaly such as our patient. Also, it has been reported that calcification and ossification occur in 0.8-10% of chronic subdural hematoma patients. Postoperative histological examination of resected capsule revealed thick, fibrous, connective tissue membrane with chronic type inflammatory infiltrations and numerous necrotic amorphous fragments and degradation products of red blood cells within lumen under the membrane. It was suggested that these findings are indication of chronic nature of the disease. Therefore, it is presumed that the patient have had chronic subdural hematoma since childhood based on physical and radiological examination and histological findings despite lack of apparent clinical information. There is a possibility that head injury one month prior to admission may have been the provoking factor for occurrence of his subdural empyema on circumstance having had chronic subdural hematoma. However, portals of entry for microbes couldn’t be identified correctly despite thorough investigation. Postoperatively, he had been treated with empirical intravenous antibiotics such as cefotaxime, vancomycin, amikacin, and clindamycin until isolation of P. acnes at two weeks after operation. When P. acnes was isolated in our patient, at first we couldn’t believe that result because it had been known to be a benign commensal and being two-week delay in the culture study. However, given his huge amount of subdural empyema and relatively long history of present symptom opposing that most subdural empyemas are acute in onset, rapid in clinical course, it might be possible to consider P. acnes as the causative organism. Thereafter, antibiotics were changed to vancomycin plus imipenem for a next six weeks. At present, there is no guideline for optimal duration of systemic antibiotics management for P. acnes. However, most reports recommend the usage of systemic antibiotics over six weeks. However, the most important factor determining duration of antibiotics management must be the patient’s clinical condition. CRP is an acute phase protein synthesized by hepatocytes and levels rarely rise above 10 mg/dL in healthy subjects. We decided the value of CRP as the important parameter of clinical condition and followed every one week. And, scheduled brain CT scans were followed every two weeks. Six weeks after operation, the CRP value was declined to nearly normal level and neurological status was improved gradually despite the brain CT scan results that showed still considerable amount of subdural fluid with mixed density. Thus, there was a discrepancy between the patient’s clinical conditions and neuroimaging findings. One year after operation, follow-up brain CT scan revealed that there was still a significant amount of subdural fluid remained but the patient complained of minimal symptoms.

CONCLUSION

As in our case, it is possible that a huge amount of subdural empyema can be accumulated in subdural space with preexisting chronic subdural hematoma that may be a suitable media for P. acnes, especially after head trauma. Also, it takes a long time to treat P. acnes subdural empyema with systemic antibiotics, at least over six weeks, and because there is an apparent discrepancy between patient’s clinical course and neuroimaging findings, the most important parameters determining management duration are the patient’s neurological status and the laboratory value of the CRP rather than neuroimaging results.
References