Seizures are the second most common neurologic complication after liver transplantation and may be caused by metabolic abnormalities, electrolyte imbalance, infection, and immunosuppressant toxicity. A 61-year-old male patient underwent liver transplantation due to hepatitis B virus-related liver cirrhosis with portal systemic encephalopathy. The immediate postoperative course of the patient was uncomplicated. However, on postoperative day (POD) 6, weakness developed in both lower extremities. No abnormal findings were detected on a brain computed tomography (CT) scan on POD 8, but a generalized tonic clonic seizure developed which was difficult to control even with multiple antiepileptic drugs. A follow-up brain CT scan on POD 15 showed a 2.7 cm sized acute intracranial hemorrhage (ICH) in the left parietal lobe. The patient's mental status improved after 2 months and he was able to communicate through eye blinking or head shaking. Our case reports an acute ICH that manifested into a refractory seizure in a patient who underwent a liver transplant.

**Key Words:** Encephalopathy, Intracranial hemorrhages, Liver transplantation, Seizures.
Seizure in a liver transplant patient

mosed vessels, tacrolimus and prednisolone for immunosuppression, and ampicillin/sulbactam, fluconazole, and nystatin for prophylaxis against infection.

The patient showed deteriorated mental status and decreased motor power on both upper extremities (grade III) and lower extremities (grade I) on postoperative day (POD) 6. Lower extremity motor power improved (from grade I to III) on POD 7. A neurologic examination conducted by a neurologist. The Glasgow Coma Score was 12 (4/2/6), and the pupil reflex was prompt. The patient developed GTCS on POD 8, with a deviated eye ball (3 o’clock direction) for 2 min, which stopped after administering lorazepam. Levetiracetam (2,000 mg/day) and pregabalin (150 mg/day) were administered to control the seizure. However, seizures with abnormal mouth chewing movements tended to be triggered when the patient’s position was changed or during pharyngeal suction. A brain computed tomography (CT) scan (Fig. 1A) failed to show any evidence of stroke. Over the next 6 days (PODs 9–15), the pregabalin dose was increased

![POD 9](image1.png) ![POD 15](image2.png) ![POD 16](image3.png)

**Fig. 1.** CT findings are demonstrated. (A) There is no evidence of acute intracranial hemorrhage or infarct. And there is no abnormal enhancing lesions or other focal lesions in the brain parenchyma. (B) Acute intracranial hemorrhage (ICH) sized 2.7 cm in left parietal lobe showed high attenuation. (C) The size of high attenuated ICH was slightly decreased. POD: postoperative day.

![Ictal EEG](image4.png) ![Interictal EEG](image5.png)

**Fig. 2.** (A) Ictal electroencephalography (EEG) showed left hemisphere dominant waves. (B) Interictal EEG showed slow waves with low voltage over the left lead, especially parietal lobe.
to 300 mg/day, and fosphenytoin (400 mg/day) was added. A subsequent brain CT (POD 15) scan revealed a 2.7 cm sized acute ICH in the left parietal lobe (Fig. 1B). Electroencephalography (EEG) showed left hemisphere dominant waves during the ictal state and slow waves with low voltage over the left lead, particularly the parietal lobe, during the interictal state (Fig. 2). Aspirin, which is prescribed routinely after liver transplantation, was stopped. A follow-up brain CT scan taken 2 days later showed that the existing ICH decreased slightly in size compared to that on previous CT scan, and there was no evidence of a new lesion (Fig. 1C). After increasing the dose of antiepileptic drugs (AED), adding another AED (400 mg/day lacosamide), and administering thiopental (100 mg) for breakthrough seizures, the frequency of seizures decreased slowly, and the EEG improved over time.

Brain magnetic resonance imaging (MRI) was performed 3 months after liver transplantation, and the ICH had decreased by 50% (Fig. 3). The patient was discharged 2 weeks after the MRI and was able to blink his eyes and shake his head.

### Discussion

As the definitive cure for end-stage liver disease, liver transplantation has become continuously more frequent over the past few decades. Postoperative complications and mortality have decreased with the improvements in surgical techniques and postoperative management, and survivors can enjoy an excellent quality of life [7,8]. However, the risk of complications remains high and critical complications can result in failure of the transplanted liver. Thus, caution and preventive measures are required.

Neurologic complications after liver transplantation are relatively rare with a reported rate of 1.5–4.7%. The most common neurologic complication is encephalopathy followed by seizure and ICH [1,3,9]. However, the mortality rate in patients with ICH after liver transplantation is > 65% [2,10,11]. Therefore early detection and adequate management are critical.

The risk of bleeding in liver transplant recipients is relatively high for several reasons. First, the transplanted liver becomes fully functional and regains adequate coagulation status after some time. Second, anti-platelet therapy is routinely used to prevent vascular thrombosis. Blood products including coagulation factors are avoided for the same reason and to evaluate recovery of the transplanted liver. Third, relative thrombocytopenia is tolerated to avoid thrombotic complications, which can be devastating. Considering our patient's coagulation profile (Table 1), the cause of ICH may have been the anti-platelet medication and postoperative coagulopathy due to post-transplantation liver dysfunction.

However, the interactions between AEDs and immunosuppressants and a possible change in activity of the AEDs due to altered serum protein levels have made newer AEDs preferred for use in liver transplant recipients. Moreover, our patient was drowsy before the liver transplantation. The laboratory findings of our patient, including the coagulation profile (Table 1) arterial blood gases, and electrolyte, glucose (Table 2), and hemoglobin levels were within normal ranges, and the initial brain CT scan after the first seizure showed no structural abnormalities. There was slight hyponatremia but the initial sodium level was slightly low, and no dramatic changes in sodium level occurred. Thus, we presumed that an electrolyte imbalance was probably not the

### Table 1. Changes in Coagulation Profile

|                | Post-reperfusion | POD 0 | POD 1 | POD 3 | POD 6 | POD 8 | POD 9 | POD 15 |
|----------------|------------------|-------|-------|-------|-------|-------|-------|--------|
| PT (INR)       | 2.06             | 1.43  | 1.33  | 1.06  | 0.98  | 0.96  | 0.97  | 1.15   |
| aPTT (sec)     | 64.9             | 45.3  | 39.9  | 32.0  | 25.9  | 24.8  | 24.2  | 32.7   |
| Platelet (×10^3/ul) | 52          | 74    | 30    | 57    | 47    | 45    | 67    | 137    |

POD: postoperative day.
Seizure in a liver transplant patient

main cause for the seizures.

Because ICH was low on the differential diagnosis list, aspirin (100 mg/day) was continued for 6 days after the seizure, which may have accelerated the progression of ICH. A better patient outcome may have been anticipated had we detected the ICH sooner and adjusted or stopped the antplatelet agent earlier.

The initial negative brain CT scan can be explained by the timing of the scan. Considering the final size of the hemorrhage and its slow rate of progression, the initial hemorrhage may have been too small to detect by the initial brain CT scan. Accuracy of the first head CT scan within 24 hours of an acute is about 70%, with 61.9% sensitivity and 71.4% specificity [12]. MRI has advantages of enabling diagnosis of an ischemic lesion and not exposing the patient to ionizing radiation [13,14]. Mullins et al. reported the superiority of diffusion-weighted MRI compared with that of CT in patients suffering an acute (within 24 hours) stroke with unknown onset of neurologic symptoms, particularly when performed within 6 hours. We could not determine the exact onset time of the neurologic symptoms because the mental status of our patient was not fully recovered; therefore, diffusion-weighted MRI may have been a better choice for the initial evaluation. However the long duration of MRI limits practical use, particularly in patients with strong suspicion of bleeding.

The EEG findings suggested that the focus of the seizure was the left parietal lobe, which correlated with the ICH location. The CT scan showed no evidence of high intracranial pressure (such as midline shift or loss of sulci), so the degree hemorrhaging did not meet the indication for surgical treatment. However, interactions between AEDs and immunosuppressants used for liver transplantation and the altered serum protein levels that can affect the activity of AEDs makes newer AEDs preferred for liver transplant recipients [4,6]. Levetiracetam, pregabalin, and lacosamide were selected because of their minimal drug interactions, and fosphenytoin was added for repetitive seizure control in this patient.

In our case, effective communication was impaired because of the underlying hepatic encephalopathy and the long duration mechanical ventilation. Although ICH is rare, considering that liver transplant recipients have high risk of bleeding, the presence of ICH should be considered when neurologic symptoms develop or when mental recovery is slow.

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