Todd’s paralysis (TP) is a neurological condition - a temporary weakness or paralysis and loss of sensation over part of the body experienced by individuals with epilepsy, in which a seizure is followed by a brief period of temporary paralysis. The paralysis may be partial or complete but usually occurs on just one side of the body. It can last from 30 minutes to 36 hours, with an average of 15 hours, before it resolves completely. Todd’s paralysis may also affect speech and vision. This transient focal neurological deficit seen after epileptic seizures was described by British neurologist Robert Todd, and as a result, the condition is often referred to as TP.1,2

The incidence of Todd paralysis following a seizure is variable and ranges between 0.6% and 13.4%.3 However, the clinical manifestations are heterogeneous and may include paresthesia, aphasia, hemianopsia, and impaired consciousness.4 An episode of TP can last anywhere from minutes to days, depending on the type of seizure, other accompanying structural damage, and any other signs that develop following the seizure.5

It is vital to distinguish Todd’s paralysis from stroke, which has a completely different treatment.6 However, this can be challenging when the paralysis is accompanied by imaging abnormalities such as diffusion restrictions seen in the acute stroke. This present case illustrated a 28-week pregnant female patient presented a Todd paralysis admitted as acute stroke following diffusion restriction seen in her brain MRI.

Case Presentation
A 27-year-old female patient who was 28 weeks pregnant was admitted to our facility during the postictal phase due to a focal-onset generalized tonic-clonic seizure she had 1 hour ago. According to the anamnesis, she initially experienced fainting in the
19th week of her pregnancy, which led to her first generalized tonic-clonic type seizure. At that time, Levetiracetam 500 BID was prescribed as a treatment. An EEG performed in the 19th week did not reveal any abnormal cerebral activity.

The patient’s neurological examination, conducted when she arrived at our emergency department, found that her left upper extremity was fully plegic at 0/5 (based on MRC); in the left lower extremity, she had 0/5 muscle strength in the proximal muscles and 2/5 muscle strength in the distal muscles. The plantar reflex was mute bilaterally (negative Babinski).

In the patient’s first MRI, acute diffusion restriction areas were observed in the right temporoparietal region, hypointense in ADC, and hyperintense in DWI (see Figure 1). After further examination, preeclampsia was excluded by the relevant department physicians. In order not to exceed the therapeutic window and detect a possible large vessel occlusion, non-contrast brain MR angiography and MR venography were performed. There was not any evidence of a major artery obstruction following the brain MR angiography. Furthermore, hypoplastic left transverse sinus was noted based on MRI venography results. Therefore, intravenous tissue plasminogen activator (TPA) or mechanical thrombectomy was not considered.

After the first examination of the patient, who was initially considered to have had an acute ischemic stroke, the neurological examination at the 45th minute showed that there was almost a complete recovery in the patient’s paresis. As a result, a control MRI was performed, and it was seen that the hyperintense areas of acute diffusion restriction in DWI and hypointensity in the ADC disappeared (see Figure 2). In line with these data, the patient’s muscle strength deficit was interpreted in favor of Todd’s paralysis, which is a complication of the postictal period.

**Figure 1** (A) initial DWI Shows hyperintensity in the right temporoparietal lobe consistent with diffusion restriction. (B) hypointensity in the right temporoparietal region in line with the hyperintensity in the DWI.

**Figure 2** (A-C) control MRI showing resolution of the diffusion restriction seen in the initial MRI after the disappearance of the patient’s weakness.
The levetiracetam dose was increased to 1000 mg BID. On an electroencephalogram performed during her hospitalization in the neurology unit, a spike-slow wave was observed in the right parietal region. The patient, whose seizures did not recur, was discharged in good condition after the fetal well-being examination was completed.

Discussion
Postictal paralysis is one of the wider spectrum of characteristics that accompany paroxysmal seizures; these symptoms are referred to as a variant of TP. Among the conditions that have been documented are postictal aphasia, hemianopsia, ideomotor apraxia, and retrograde non-memory impairment. The most important suggestion of TP is the presence of previous episodes of involuntary facial or limb contractions that may be associated with altered mental status. Therefore, anamnesis is paramount for identifying TP. Todd paralysis usually commences after a focal seizure and involves one or more limbs.7,8

The mechanism underlying the development of TP still remains uncertain. Numerous theories concerning its etiology have been proposed, including neuronal desensitization, neurotransmitter depletion, focal reduction of cerebral blood flow, active suppression, and extensive neuronal firing during seizures.9,10

Although the differential diagnosis of epilepsy and stroke is always easy to make, neurological deficits that develop after seizures should be specially considered in the differential diagnosis of stroke. TP should also be considered a rare etiological agent in cases presenting with acute-onset neurological deficits and accompanied by cytotoxic edema on DWI.5,11

At clinical presentation, the first symptom being a seizure, early complete recovery of the neurologic deficit, and some features of the lesion on DWI may be important clues in the differential diagnosis.5 In this case report, our pregnant patient’s initial imaging was suggestive of stroke due to diffusion restriction, but as the findings occurred following an epileptic seizure and the deficit in her neurological examination showed almost complete recovery 45 minutes later, the diagnosis shifted away from stroke. The control brain MRI, which was performed immediately after the disappearance of the neurologic deficit, demonstrated resolution of the diffusion restriction seen in the initial MRI. In addition, the brain MR angiography revealed normal findings, and the MR venogram did not show any occlusion of the cerebral veins and sinuses. Preeclampsia was ruled out by consultation with the corresponding physicians. An electroencephalogram demonstrated a spike-slow wave in the right parietal region. As a result, Todd’s palsy, which is attributed to transient global hemispheric hypoperfusion in the postictal period, was considered in the differential diagnosis. In this case, in which the beginning stroke was considered, the treatment was shifted to postictal paralysis. The dose of levetiracetam was increased to 1000 mg BID. The patient was discharged from the hospital after it was confirmed that the seizures were not recurring and that the fetal condition was stable.

Conclusion
We believe that our case illustrates the challenges in distinguishing between TP and stroke, which have entirely distinct etiologies and courses of treatment, and that it provides important clues that help the differentiation of the two conditions.

Ethical Approval
Institutional approval was not required to publish the case details.

Consent for Publication
Written informed consent was obtained from the patient to have the case details and any accompanying images published.

Author Contributions
All authors made a significant contribution to the work reported, whether that is in the conception, study design, execution, acquisition of data, analysis, and interpretation, or in all these areas; took part in drafting, revising, or critically reviewing the article; gave final approval of the version to be published; have agreed on the journal to which the article has been submitted; and agree to be accountable for all aspects of the work.
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