The Dark Side of Seizure: Case Series of a T2 Dark-Through Pattern of Peri-ictal Diffusion Restriction

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Abstract

Analogous to T2 shine-through, T2 hypointensity can “dark-through” on diffusion weighted imaging (DWI) and mask diffusion restriction. In such cases, diffusion restriction is evident only by apparent diffusion coefficient (ADC) hypointensity, which is often subtle and easily missed without corresponding DWI hyperintensity. Because diffusion restriction may be present in the setting of seizure, avoiding this pitfall can aid in seizure diagnosis for patients without any other magnetic resonance imaging (MRI) findings. We present a case series of nine patients in the peri-ictal period with T2 dark-through on MRI to raise awareness of this finding and its potential clinical role.

Keywords

T2 hypointensity, Diffusion restriction, Seizure

Introduction

T2 dark-through describes diffusion restriction in the setting of T2 hypointensity. Just as T2 hyperintensity can “shine-through” on diffusion weighted imaging (DWI), T2 hypointensity can “dark-through” on DWI. DWI signal is influenced by multiple factors including T2 signal, which is demonstrated in the equation: $\text{DWI signal intensity} = kPDe^{-\frac{TE}{T2}}e^{-bADC}$, where $k$ is a constant, PD is the proton density, TE is the echo time, and $b$ is the b-value [1]. Understanding T2 dark-through is essential for detecting diffusion restriction of T2 hypointense processes where subtle apparent diffusion coefficient (ADC) hypointensity may be missed without corresponding DWI hyperintensity.

There have been reports of T2 dark-through in patients with seizure, hyperglycemia, Moma-Moya, Sturge-Weber, intracranial neoplasm, intracranial infection, and trauma [1-10]. We present a case series of nine patients with peri-ictal T2 dark-through, including the first documented cases in the setting of venous thrombosis and subdural hemorrhage.

Cases of Hyperglycemia

A 6-year-old male presented with 3 episodes of focal right-sided shaking and severe hyperglycemia (414 mg/dL). MRI showed left parietal T2 dark-through (Figure 1). Patient improved following the treatment of hyperglycemia.

A 52-year-old female presented with episodic confusion over the course...
of 6 months and severe hyperglycemia (413 mg/dL). MRI showed right parietal T2 dark-through (Figure 2). Electroencephalogram (EEG) showed right intermittent rhythmic delta activity and background slowing. Patient improved following the treatment of hyperglycemia.

A 56-year-old male presented with episodic left lower quadrant visual changes 50 times per day and severe hyperglycemia (451 mg/dL). Initial MRI showed right parietal T2 dark-through (Figure 3). Patient improved following the treatment of hyperglycemia. Findings were less prominent on 3-month follow-up MRI.

A 32-year-old male presented with generalized tonic-clonic convulsions lasting 1 minute and superior sagittal sinus thrombosis. MRI showed left parietal T2 dark-through, venous thrombosis, and leptomeningeal enhancement, the latter thought to be related to venous stasis (Figure 4). Patient was diagnosed with protein C deficiency and improved following anticoagulation.

Case of Hyperglycemia and Venous Thrombosis

A 62-year-old female presented with left upper extremity coordination difficulties over the course of 1 week with 2 episodes of shaking lasting 2-3 minutes, severe hyperglycemia (505 mg/dL), and right transverse sinus thrombosis. Initial MRI showed right parietal T2 dark-through, venous thrombosis, and leptomeningeal enhancement, the latter thought to be related to venous stasis (Figure 5). Patient improved following the treatment of hyperglycemia and anticoagulation. T2 dark-through was resolved on 1-month follow-up MRI.

Cases of Meningitis

A 9-year-old female presented with right upper extremity shaking and numbness in the setting of meningitis. Cerebrospinal fluid analysis showed an opening pressure
of 24 cm H2O, 93 WBC per uL (48% neutrophils, 45% lymphocytes), 26.5 mg/dL of protein, 60 mg/dL of glucose, and negative gram stain. MRI demonstrated left frontal T2 dark-through and leptomeningeal enhancement (Figure 6). Patient was lost to follow-up.

A 16-year-old male presented with left upper extremity and left facial twitching for 2 minutes in the setting of meningitis. MRI demonstrated right frontal T2 dark-through, sulcal FLAIR hyperintensity, and leptomeningeal enhancement (Figure 7). Patient was lost to follow-up.

Case of Sturge-Weber

A 2-year-old male with history of Sturge-Weber presented with seizure of unspecified semiology. MRI demonstrated right cerebral T2 dark-through and stigmata of Sturge-Weber including asymmetric smaller right cerebral hemisphere with pial angiomatosis (Figure 8). CT showed subcortical calcification only in the posterior right cerebral hemisphere, despite diffuse T2 dark-through throughout the right cerebral hemisphere. The patient underwent right hemispherectomy due to uncontrolled seizure.

Case of Subdural Hematoma

A 78-year-old female presented with one episode of right upper extremity tonic-clonic movement and intermittent right-sided weakness lasting 1-5 minutes followed by sleepiness in the setting of subdural hemorrhage. MRI showed left frontal T2 dark-through (Figure 9). The patient improved following hematoma evacuation and seizure management.

Discussion

T2 dark-through is an under-recognized pattern of diffusion restriction where T2 hypointensity "darks-through" on DWI and masks signal changes. When this happens, restricted diffusion can only be identified by ADC hypointensity, which may be subtle and easily missed. We present a case series of peri-ictal T2 dark-through, including the first documented cases in the setting of venous thrombosis and subdural hemorrhage, to raise awareness of this finding and its potential clinical role.

The T2 hypointensity component of T2 dark-through has been attributed to the susceptibility effects of paramagnetic substances like deoxyhemoglobin and free radicals [1]. These materials can accumulate from (1) oxygen demand exceeding oxygen supply despite compensatory increased blood flow, such as in the setting of seizure and super-imposed stressors, or (2) insufficient venous drainage, such as in Sturge-Weber, venous thrombosis, or mass effect from adjacent subdural hemorrhage. In the case of Sturge-Weber, axonal hypermyelination and other structural white matter abnormalities have also been proposed as mechanisms for T2 hypointensity [6].

The ADC hypointensity component of T2 dark-through may be related to a combination of hypoxia and excitatory neurotransmitters leading to intracellular edema. In the setting of hypoxia, sodium/potassium-ATPase pump failure results in intracellular shift of sodium and water [11]. Excitatory neurotransmitters released during seizure activate ion channel coupled receptors, which allow the influx of sodium and water and can activate the cell death cascade [12, 13]. Although the precise combination and role of these mechanisms are still unclear, it is believed that intracellular edema stemming from hypoxia and over-excitation in the peri-ictal state may be responsible for diffusion restriction.
Patients that received follow-up imaging did not have evidence of encephalomalacia. Plausible explanations for this include: (1) T2 dark-through is reversible if underlying seizure and stressors are treated promptly; (2) T2 dark-through may cause mild tissue injury evident only on histology; (3) our follow-up imaging did not allow sufficient time for the full extent of the T2 dark-through event to manifest. Although imaging was obtained within 24 hours of seizure, the precise time-course is unknown. It is possible many cases of T2 dark-through are resolved by the time patients are scanned. Prompt imaging in the acute setting and long-term follow-up imaging may increase the sensitivity for T2 dark-through and more reliably define its time course and reversibility.

Although there is a correlation between T2 dark-through and seizure, a relationship of causality is yet to be determined. Regardless, recognizing T2 dark-through may have important implications for seizure localization and managing seizures with potentially reversible underlying etiologies.

Our case series is limited by retrospective review of data, small sample size, and qualitative image analysis. Future prospective studies with a larger sample size and quantitative T2 dark-through analysis can determine the ADC and T2 thresholds for certain clinical outcomes, and better characterize the patient population who will exhibit T2 dark-through and benefit from a more comprehensive workup for comorbidities.

Conclusion

T2 dark-through is an under-recognized presentation of diffusion restriction occurring in the peri-ictal period. This finding should prompt the radiologist to look for superimposed stressors such as metabolic and vascular abnormalities. Although there is a correlation between T2 dark-through and seizure, a relationship of causality is yet to be determined. With further study, T2 dark-through may have a clinical role in identifying and managing seizure.

Conflict of Interest

The authors declare no conflict of interest.

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