

Ictus emeticus presenting as an unusual seizure type in chromosome 22q11.2 deletion syndrome

Pi-Lien Hung¹, Li-Tung Huang¹, Shang-Yeong Kwan²,
Kai-Ping Chang³, Hsin-Hung Chen⁴, Yi-Yen Lee³,
Huang-Chuen Fan⁵, Chien Chen²

¹ Department of Pediatrics, Kaohsiung Chang Gung Memorial Hospital and Chang Gung University College of Medicine, Kaohsiung

² Department of Neurology, Taipei Veterans General Hospital and Department of Neurology, School of Medicine, National Yang-Ming University, Taipei

³ Department of Pediatrics, Taipei Veterans General Hospital, Taipei

⁴ Division of Pediatric Neurosurgery, Neurological Institute, Taipei Veterans General Hospital, Taipei

⁵ Department of Pediatrics, Tungs' Taichung Metro Harbor Hospital, Wuchi, Taichung, Taiwan

Received October 10, 2016; Accepted February 02, 2017

Ictus emeticus presenting as an unusual seizure type in chromosome 22q11.2 deletion syndrome

- Bilateral polymicrogyria is an indicative feature of CNS malformation in 22q11.2DS, which may precipitate seizures.
- In addition to hypocalcaemic seizures, ictus emeticus can be the peculiar seizure type in patients with 22q11.2DS who present with cyclic vomiting.

- The ictal EEG in our patient showed a slowing delta wave over the left temporal area associated with loss of ability to speak, which indicated the seizure originated from the language-dominant hemisphere.