

# Intractable hypocalcemic seizures due to hypoparathyroidism

Dear Editor,

In your recent issue, we read the case report “Intractable hypocalcemic seizures with neuropsychiatric symptoms- An under-diagnosed case” written by Kaeley *et al.*<sup>[1]</sup> This case report was very well summarized and highly informative. Hypocalcemia is one of the important causes of intractable seizures. The authors explained various acute and chronic manifestations of hypocalcemia along with felicitous take-home messages. Primary care physicians should be aware of hypocalcemic seizures whenever they encounter patients with refractory seizures.

Clinical presentation of hypoparathyroidism also varies depending on the duration of hypocalcemia. Muscular cramps, tetany, numbness, cardiac arrhythmias, altered behavior, and seizures are usually acute manifestations, and cataracts, basal ganglia calcifications, dilated cardiomyopathy, dementia, and cerebellar dysfunction are chronic manifestations.<sup>[2]</sup>

Seizures due to idiopathic hypoparathyroidism are often misdiagnosed as epilepsy. MRI Brain for this misdiagnosed epilepsy suggests intracranial calcification which increases suspicion of hypoparathyroidism. Idiopathic hypoparathyroidism needs to be kept as a differential diagnosis of intractable generalized, tonic-clonic seizures so that primary care physicians can prevent anonyms use of anti-epileptic drugs in this condition. Hypocalcemia with intracranial calcifications is a diagnostic clue for hypoparathyroidism. Early treatment with calcium and vitamin D helps to prevent the recurrence of seizures and other neurological complications.<sup>[3]</sup>

Intracranial calcification is an infrequent complication of hypoparathyroidism. Factors that predispose this intracranial calcification have not been completely explicated. Persistent hypocalcemia and hyperphosphatemia promote deposition of calcium-phosphorus complexes in brain parenchymal tissue. Increased expression of osteogenesis-related molecules like osteonectin/osteopontin in the caudate nucleus and grey matter could favor metastatic calcification in hypoparathyroidism.<sup>[4]</sup>

Autoimmune hypoparathyroidism is the second most common form of hypoparathyroidism in adults after postsurgical hypoparathyroidism. Auto-immune hypoparathyroidism and

genetic causes are important etiological factors that should be ruled out before labeling as idiopathic hypoparathyroidism. Anti-calcium sensing receptor (CaSR) antibodies and genetic testing for Glial Cell Missing-2 (GCM2) or CaSR genetic mutations can be done for further work-up.<sup>[5]</sup>

The non-availability of these genetic testing facilities in most of the Indian medical institutes and the economic constraints of patients are important limiting factors for the diagnosis of auto-immune hypoparathyroidism.

Primary care physicians and family physicians should keep a differential as hypocalcemic seizures whenever they face refractory seizures, not responding to antiepileptic drugs.

### Key points:

1. Hypocalcemia is an important cause of intractable seizures.
2. Auto-immune hypoparathyroidism should be ruled out before labelling as idiopathic hypoparathyroidism.
3. Primary care physicians should be aware about hypocalcemic seizures whenever they encounter the patients with intractable seizures.

### Financial support and sponsorship

Nil.

### Conflicts of interest

There are no conflicts of interest.

**Naresh K. Midha, Mahendra K. Garg**

*Department of Medicine, All India Institute of Medical Sciences, Jodhpur, Rajasthan, India*

**Address for correspondence:** Dr. Naresh K. Midha,  
Department of Medicine, All India Institute of Medical Sciences  
Jodhpur, Rajasthan - 342 005, India.  
E-mail: midha.naresh2005@gmail.com

### References

1. Kaeley N, Baid H, Chawang H, Vempalli N. Intractable hypocalcemic seizures with neuropsychiatric symptoms- An under-diagnosed case. *J Family Med Prim Care* 2021;10:2032-4.
2. Lopes MP, Kliemann BS, Bini IB, Kulchetscki R, Borsani V, Savi L, *et al.* Hypoparathyroidism and pseudohypoparathyroidism: Etiology, laboratory features and complications. *Arch Endocrinol Metab* 2016;60:532-6.
3. Letchinger R, Doyle K, Claassen J, Thakur KT. New-onset super-refractory status epilepticus: A case series of 26 patients. *Neurology* 2020;95:e2280-5.
4. Goswami R, Millo T, Mishra S, Das M, Kapoor M, Tomar N, *et al.* Expression of osteogenic molecules in the caudate

nucleus and gray matter and their potential relevance for basal ganglia calcification in hypoparathyroidism. J Clin Endocrinol Metab 2014;99:1741-8.

5. Midha NK, Garg MK, Kumar D, Meena DS, Bohra GK. Rapidly developing cataract in young adult patients: Always a matter for further evaluation. Cureus 2021;13:e17312. doi: 10.7759/cureus.17312.

This is an open access journal, and articles are distributed under the terms of the Creative Commons Attribution-NonCommercial-ShareAlike 4.0 License, which allows others to remix, tweak, and build upon the work non-commercially, as long as appropriate credit is given and the new creations are licensed under the identical terms.

**Received:** 01-06-2021

**Accepted:** 28-09-2021

**Published:** 27-12-2021

Access this article online	
<b>Quick Response Code:</b> 	<b>Website:</b> www.jfmpc.com
	<b>DOI:</b> 10.4103/jfmpc.jfmpc_1037_21

**How to cite this article:** Midha NK, Garg MK. Intractable hypocalcemic seizures due to hypoparathyroidism. J Family Med Prim Care 2021;10:4615-6.

© 2021 Journal of Family Medicine and Primary Care | Published by Wolters Kluwer - Medknow