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Seizure to Drug Induced Schizophrenia: A Rare Case of Keppra-Induced Psychosis

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Seizure to Drug Induced Schizophrenia: A Rare Case of Keppra-Induced Psychosis Shadi Shams^a BA, Riddhima Issar^a MS, Nardin El-Shammaa^{a,b} DO, Munaza Khan^{a,b} MD, ^aRowan School of Osteopathic Medicine, ^bVirtua Health-Our Lady of the Lourdes, NJ

Background

Levetiracetam is a broad-spectrum antiseizure medication and is approved as adjunctive thera to treat focal-onset seizures in children and add with epilepsy. Levetiracetam has a wide margin safety and patient-friendly pharmacokinetics th distinguish it from other currently available antiepileptic drugs.

Most common side effects are fatigue, somnole dizziness, and upper respiratory infection. Neuropsychiatric symptoms are reported. Psych symptoms, paranoid ideation, hallucinations, a behavioral problems (including aggressive beha agitation, anger, anxiety, apathy, confusion, depersonalization, depression, emotional labilit hostility, dyskinesia, irritability, neurosis, and personality disorder) may occur in adult and pediatric patients.

Amongst all adverse effects, the rate of psychos very low and ranges from less than 1% to 1.4%. retrospective study showed that this rate is high in older patients than in the younger population

Although the current literature provides cases levetiracetam-induced psychosis, the onset of psychotic symptoms is one-week post drug administration. Little to no data exists on the o of psychosis post levetiracetam induction.

Management of psychosis can be done with do reduction or discontinuation of the medication

References:

	Case Presentat
	A 67-year-old male with no psychiatric h
ару	the hospital for auditory hallucinations a
ults	ideations that a microchip was implante
n of	he was being tracked by the governmen
hat	voices telling him "he has brain cancer".
	nallucinations and paranoid ideations st
	idestions Upon further investigation th
ence	course and timeline were established
chec,	
hotic	Unwitnessed seizure O
ind	Levetiracetam initiated
avior,	30 days
	October 2021 No
ity,	
	The patient denied any recent changes t
	except starting levetiracetam 750mg.
	Diagnosis . Diagnostic workup included -
cic ic	count metabolic nanel thyroid function
	level urine analysis urine drug screen (
ther	tomography scan and magnetic resonan
on.	head, electroencephalography; which w
	prolonged video-EEG monitoring was no
of	paroxysmal activity, subclinical or clinica
	history, physical examination, diagnostic
	of symptoms post medication change, a
nset	levetiracetam-induced psychosis was ma
	Patient outcome: Levetiracetam was sw
JSe	with balangeridal 2 mg ance daily with a
Ι.	to twice daily. The nationt recovered over
	was discharged from the hospital

Chen B, Choi H, Hirsch LJ, et al. Psychiatric and behavioral side effects of antiepileptic drugs in adults with epilepsy. Epilepsy. Epilepsy. Epilepsy. Epilepsy. 2017;76:24-31. doi:10.1016/j.yebeh.2017.08.039 Patsalos PN. Pharmacokinetic profile of levetiracetam: toward ideal characteristics. *Pharmacol Ther*. 2000;85(2):77-85. doi:10.1016/s0163-7258(99)00052-2 Pinckaers FME, Boon ME, Majoie MHJM. Risk factors predisposing to psychotic symptoms during levetiracetam therapy: A retrospective study. Epilepsy Behav. 2019;100(Pt A):106344. doi:10.1016/j.yebeh.2019.05.039

on

nistory presented to and paranoid ed in his ear and that nt. He had heard The auditory tarted two weeks omicidal e following hospital



to his medication

a complete blood cell n tests, blood alcohol computed nce imaging of the vere unremarkable. A ormal with no l seizures. Based on tests, and the onset diagnosis of ade.

vitched to valproate ychosis was managed n eventual increase er next 48 hours and

Few days after stopping levetiracetam, the psychotic symptoms resolved which further proved that the symptoms were drug induced. Levetiracetam induced psychosis is mainly reported in patients who have predisposing factors, such as history of psychiatric illness or epilepsy, but our patient lacked both. Additionally, most reported cases of Levetiracetam induced psychosis occurred within a week of starting the medication, but in our case, the onset of psychosis was delayed to a month after medication induction. Furthermore, the current literature notes that levetiracetam induced psychosis, although rare, is seen in patients on concurrent use of other anti-epileptic medications.

Conclusion

Levetiracetam is found to be a safer option amongst all the available anti-epileptic medication. This profile may facilitate the clinical management of patients with epilepsy by providing a safer and less-complicated therapeutic strategy which is preferable option to prevent polypharmacy in elders. However, as shown in the case report, Levetiracetam-induced psychosis should be kept in mind regardless of the time of onset, predisposing factors, and the severity of the symptoms. When observed, physician should stop the medication and switch to other classes of anti-epileptic medications, such as valproic acid which could result in complete remission of side effects. This case highlights that it is essential for psychiatrists to consider drug induced psychosis even when the onset of symptoms is not acutely after drug administration.



Discussion