Eucalyptus oil (EO) is an essential oil which has been used as a traditional remedy in upper respiratory tract infection. It contains approximately 90% cineole and is readily available worldwide in over-the-counter cough drops, liniments, toothpaste, mouthwashes, cold preparations, and hair lice remover. EO-induced adverse drug reaction is rare in both adults and children. The signs and symptoms of EO poisoning are CNS depression, hypotension, tachycardia, epigastric pain, nausea, vomiting, and contact dermatitis. Symptom onset is usually rapid and resolves within 24 h. We report the case series of four adult patients with EO-induced seizure in India, who inhaled EO for common cold and presented to the critical care with single first attack of generalized tonic-clonic seizures. On further evaluation, none of them had a family background of seizures/febrile seizures. EEG and brain MRI were found to be normal in all patients. All the patients were managed with anti-epileptic drugs and standard supportive care. All medical practitioners should be aware of the toxic effects of EO, a common OTC medication used in Indian households. Warning labels may be attached on EO comprised products.

**Keywords:** Eucalyptus oil, Poisoning, Seizures, OTC medication.

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**ABSTRACT**

Eucalyptus oil (EO) is an essential oil extracted from *Eucalyptus globulus*, a Myrtaceae plant family, native to Australia. EO has been used as a traditional remedy to upper respiratory tract infection, gastritis, and diabetes [4]. Chemical constituents of EO are 1,8-cineole and other compounds are monoterpenes (β-pinene, α-pinene, β-pinene, and p-cymene), alkaloids, eucalyptin, phenols, flavonoids 6,8-dimethylkaempferol-3,7-dimethyl ether, oleanolic acid (2,3-dihydroxy-urs-12-en-28-oic acid), 8-desmethyl-eucalyptin, tannins, terpenoid phenolaldehydes, 2,6-dihydroxy-3′-methyl-4′-methoxy-dihydrochalcone, and verbeneone, a monoterpene bicyclic ketone [5]. The pharmacological activities of EO include antimicrobial, antihypglycemic, antiinflammatory, anti-viral, and anti-inflammatory properties [5]. These volatile oils are found in many over-the-counter cough drops, liniments, toothpaste, mouthwashes, cold preparations, and hair lice remover and additionally used in transdermal drug delivery to enhance the penetration of minoxidil, zidovudine, and valsartan [6]. The public may not be aware of the potentially toxic effects of these oils and products if they are inhaled or ingested orally at higher concentrations. The signs and symptoms of EO poisoning are CNS depression, hypotension, tachycardia, epigastric pain, nausea, vomiting, and contact dermatitis. Symptom onset is usually rapid (within 5–10 min) and resolves within 24 h [7]. In this report, we present a case series of EO-induced seizures. Written informed consent was obtained from the patients for publication of this case report.

**DESCRIPTION OF CASES**

A 21-year-old male admitted to the emergency department with a single episode of generalized tonic-clonic seizure. He also had uprolling of the eyelid, one episode of vomiting, headache, and generalized weakness. The patient had a fever and nasal congestion with nasal block for a day. He sought to relieve his symptoms with home remedies by application of 6–7 drops of oil on his handkerchief. Following this, he kept inhaling the oil 3–4 times by covering his nose with the same handkerchief. Within a short while, he lost his balance and fell down. He had clonic movements for 5 min, postictal confusion and nausea on presentation to the hospital. This was described by his relative accompanying him. On investigation, the patient had no previous history of seizures and did not have febrile seizures in childhood. There was no family background of seizure attacks. He had never inhaled EO in the past. His brain MRI was normal and the EEG report was also normal. He was given injection fosphenytoin 150 mg IV route twice daily for 2 days along with other supportive treatment and was discharged after 2 days with an advice to continue anti-epileptic drugs for 1 month. During the follow-up for 6 months, no recurrence of seizure was observed. The patient was advised to stop the anti-epileptic drugs after a month. The causes of seizure were probably due to an increased cellular hyperexcitability induced by chemical compounds (cineole) in the EO.

**Case 2**

A 25-year-old male patient presented with one episode of a generalized tonic-clonic seizure. He had a lateral tongue bite with giddiness. The patient had inhaled the vapors of EO that was mixed in boiling water. He lost consciousness and had tonic posturing of all four limbs followed by clonic movements for 1 min. He had no previous history of a seizure attack. Family history revealed that his young brother had the same complaint 1 year ago. The treatment was started with intravenous levetiracetam 1000 mg (BD). On the next day, brain MRI and EEG were performed and reports are normal. The patient was continued with the same anti-epileptic drug for the next 2 days. He did not have any recurrent episodes of seizures. He was discharged on levetiracetam 500 mg twice daily for a month. After a month of follow-up, he was observed to be asymptomatic.
Case 3
A 31-year-old male patient presented with one episode of generalized tonic-clonic seizure, postictal drowsiness with nausea in the emergency department. The patient had steam inhalation with EO for 3 min before the seizure attack. He had nasal congestion for 2 days with headache. He had no history of seizure attack and had no family history. The treatment was started with intravenous levetiracetam 1000 mg (BD). Brain MRI and EEG reports revealed no abnormalities. The patient was discharged with medical advice on levetiracetam 500 mg twice daily for a month. After a month of follow-up, he was found to be asymptomatic.

Case 4
A 37-year-old male patient presented with an episode of generalized tonic-clonic seizures with generalized weakness. Within 5 min of steam inhalation with EO, he had a seizure attack and remained drowsy for 30 min. His brain MRI and EEG were normal. The patient received levetiracetam 1000 mg (BD) and was advised to continue anti-epileptic drugs for 2 months. The patient was not willing to continue the medication. He was discharged after 2 days with patient counseling.

DISCUSSION
EO is a volatile oil obtained by rectifying the leaves of E. globulus. Australian Aboriginals use it as a traditional medicine for the treatment of body pains, sinus congestion, fever, and cold [8,9]. Dennis Considen and John White, surgeons on the first fleet, distilled EO in 1788 [10]. The active constituents are monoterpenes 1, 8-cineole (eucalyptol), which comprises nearly 70%. It is readily available in many over-the-counter medications such as cough syrups, toothpastes, mouthwashes, and cold preparation slice remover and additionally used in transdermal drug delivery to enhance the penetration of minoxidil, zidovudine, and valsartan [6]. The reason behind the adverse effect of EO may be hydrocyanic acid which is highly toxic and may cause nausea, vomiting, drowsiness, seizures, coma, and at times death [11]. Such an adverse reaction induced by EO case was first found in Kerala on 1898 [12]. Similar case is reported in the U.K on 1911 [13]. In Toronto, Canada, they performed a retrospective analysis on EO poisoning over a period from December 1995 to March 1997. They found 232 cases of EO-induced adverse reaction which were reported [14]. Likewise, 42 cases of EO ingestion in children under 14 years of age were identified in a defined population between July 1, 1984, and June 30, 1991, in the Southeast, Queensland, by Webb and Pitt in December 1992 [15].

In 2011, New Zealand, Waldman explained 4-year-old girl with tonic-clonic attack due to external application of EO and more recently in 2019 Sitaraman and Rao posted a report of EO poisoning over a period from December 1995 to March 1997. They found 232 cases of EO-induced adverse reaction which were reported [14]. Likewise, 42 cases of EO ingestion in children under 14 years of age were identified in a defined population between July 1, 1984, and June 30, 1991, in the Southeast, Queensland, by Webb and Pitt in December 1992 [15]. In 2011, New Zealand, Waldman explained 4-year-old girl with tocic-clonic attack due to external application of EO and more recently in 2019 Sitaraman and Rao posted a report of EO poisoning over a period from December 1995 to March 1997. They found 232 cases of EO-induced adverse reaction which were reported [14]. Likewise, 42 cases of EO ingestion in children under 14 years of age were identified in a defined population between July 1, 1984, and June 30, 1991, in the Southeast, Queensland, by Webb and Pitt in December 1992 [15].

CONCLUSION
This case series describes the seizure inducing property of EO and the importance of disseminating this information among the health care workers so that the public can be dissuaded from their usage. Although EO-induced seizure has a low incidence rate, one should keep this as a potential cause of seizures in any patient presenting to the hospital with a single episode of seizures and a recent history of taking any over-the-counter medication for rhinitis.

CONFLICTS OF INTEREST
The author declared that there are no conflicts of interest related to this study.

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ETHICAL STATEMENT
Ethical approval was not applicable for case report in our institution.

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