NOTE

Anticonvulsant Potential of Essential Oil of *Vetiveriazizanioides* Linn. Nash

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In light of the traditional claim of south Indian tribes to use root and stem juice of *Vetiveria zizanioides* (Linn.) Nash in the treatment of epilepsy, an investigation was carried out to evaluate the anticonvulsant activity of essential oil obtained from the fresh roots and stem of *V. zizanioides*. The anticonvulsant activity was screened in maximal electroshock induced seizure (MES) and subcutaneous pentylenetetrazole induced seizure models in mice. The neurotoxicity was assessed using the rotarod method. Oil obtained from roots of *V. zizanioides* showed 100 % protection in maximal electroshock induced seizures. But, no protection was observed in subcutaneous pentylenetetrazole induced seizures. Oil obtained from stems of *V. zizanioides* was inactive in both maximal electroshock and subcutaneous pentylenetetrazole induced seizures. No neurotoxicity was observed at the highest administered dose. Thus, the present study has demonstrated that the essential oil present in roots of *V. zizanioides* has anticonvulsant properties.

Key Words: Anticonvulsant activity, *Vetiveriazizanioides* Linn. Nash.

Vetiveria zizanioides (Linn.) Nash (Poaceae), popularly known as khus grass, has been known in India since ancient times. Khus grass grows wild in many states, namely Haryana, Uttar Pradesh, Rajasthan, Gujarat, Bihar, Orissa, Madhya Pradesh and throughout south India. Two species of *Vetiveria* are found in India, of which *V. zizanioides* is the common source of the well known oil of vetiver, which is used in medicine and in perfumery¹. The grass has been reported to have antimycobacterial², antibacterial³, antipyretic⁴, antioxidant⁵, larvicidal, repellent and insecticidal properties⁶. Various tribes use different parts of the grass for many of their ailments such as mouth ulcer, fever, boil, epilepsy, burn, snakebite, scorpion sting, rheumatism, fever, headache, etc. The Santhal tribes of Bihar and West Bengal use the paste of fresh roots for burn, snakebite and scorpion sting and a decoction of the roots as a tonic for weakness. The Lodhas tribes of West Bengal use the root paste for headache, rheumatism and sprain and a stem decoction for urinary tract infection; the Mandla and Bastar tribes of Madhya Pradesh use the leaf juice as anthelmintic; the tribes of the Varanasi district inhale the root vapour for malarial fever. The root ash is given to patients for acidity by the Oraon tribe. Likewise, there are many different applications of the plant for different ailments among different ethnic tribes. South Indian tribes use leaf paste of khus grass in rheumatism and sprain and root and stem juice

in boil, burn, epilepsy, scorpion sting, snakebite and mouth ulcer^{7,8}.

Thus, there are traditional evidences for the use of root and stem juice of *V. zizanioides* in epilepsy, but this activity has not yet been established experimentally. Essential oil is reported to be present in the roots of *V. zizanioides*⁹. As essential oils present in plants are shown to be responsible for the anticonvulsant activity of the plant¹⁰. An attempt was made to evaluate the anticonvulsant potential of essential oil obtained from fresh roots and stems of *V. zizanioides*. The anticonvulsant activity was screened in maximal electroshock induced seizure (MES) and subcutaneous pentylenetetrazole (scPTZ) induced seizure models in mice. The neurotoxicity was assessed using the rotarod method.

The roots and stem of *V. zizanioides* were collected from the local market of Muzaffarnagar and authenticated in the Department of Pharmacognosy, S. D. College of Pharmacy & Vocational Studies, Muzaffarnagar, India. A voucher specimen (PK-112) has been deposited in the herbarium.

Preparation of extract: The fresh roots and stems were grounded in a blender. The grounded material was subjected to hydrodistillation using a Clevenger apparatus for 4 h for isolation of oil (Oil of roots of *V. zizanioides*: ROVZ: 2.0 % v/w; Oil of stem of *V. zizanioides*: SOVZ: 0.8 % v/w). The essential oil was diluted with sesame oil to obtain the desired

TABLE 1
ANTICONVULSANT AND NEUROTOXIC ACTIVITY OF OIL OF ROOTS OF
V. zizanioides (ROVZ); OIL OF STEM OF *V. zizanioides* (SOVZ)

Compound	Intraperitoneal injection in mice*					
	MES screen (h)		scPTZ screen (h)		Neurotoxicity screen (h)	
	0.5	4.0	0.5	4.0	0.5	4.0
ROVZ	-	300**	-	-	-	-
SOVZ	-	-	-	-	-	-
Phenytoin	30	30	-	-	100	100
Sodium valproate	-	-	-	300	-	-

*Dose of 30, 100 and 300 mg/kg were administered. The figure in the table indicates the minimum dose whereby bioactivity was demonstrated in half or more mice. The dash (–) indicates an absence of activity at maximum dose administered (300 mg/kg). **In the MES screen ROVZ showed protection (1/1, 4 h) at a dose of 300 mg/kg, but SOVZ showed no protection in MES and scPTZ screen

doses and was immediately administered intraperitoneally (i.p.) to mice.

Anticonvulsant and neurotoxicity evaluation: The animal study was carried out according to the protocols approved by the Institutional Animal Ethical Committee, S.D. College of Pharmacy and Vocational Studies, Muzaffarnagar (Reg No. 876/AC/05/CPCSEA). Male albino mice (CF-1 strain, 18-25 g) were used as experimental animals. The animals were housed in metabolic cages and allowed free access to food and water. In the preliminary screening each compound was administered as an i.p. injection at three dose levels (30, 100 and 300 mg/kg) and anticonvulsant and neurotoxic effects were assessed at 0.5 h and 4 h intervals after administration. Anticonvulsant efficacy was measured by maximal electroshock induced seizure and subcutaneous pentylenetetrazole tests.

Essential oils are present in the roots and stem of *V. zizanioides*. The essential oil in the roots (ROVZ) is thick and dark brown in colour. The oil distilled from the stem (SOVZ) is inferior in quantity and quality from that obtained from roots.

ROVZ and SOVZ were evaluated for anticonvulsant activity in maximal electroshock induced seizure (MES) and subcutaneous pentylenetetrazole induced seizure models in mice using doses of 30, 100, 300 mg/kg. The observation was carried out at two different time intervals (0.5 and 4 h). The pharmacological test results are shown in Table-1. ROVZ showed 100 % protection (1/1, 4.0 h) at a dose of 300 mg/kg in maximal electroshock induced seizure test, indicating the

compounds ability to prevent seizure spread. But, ROVZ showed no protection in subcutaneous pentylenetetrazole induced seizures. SOVZ is inactive in both maximal electroshock induced seizure and subcutaneous pentylenetetrazole induced seizures. Both ROVZ and SOVZ exhibited no neurotoxicity at the highest administered dose (300 mg/kg) in rotorod method. Thus, the present study has demonstrated that the essential oil present in roots of *V. zizanioides* has anticonvulsant properties. Advanced anticonvulsant screening and mechanistic studies are required to further explore the anticonvulsant activity of roots of *V. zizanioides*.

REFERENCES

1. R.R. Rao and M.R. Suseela, Proceedings of Second International Conference on Vetiver. Office of the Royal Development Projects Board, Bangkok, pp. 444-448 (2000).
2. J.A. Hammond, D. Fielding and S.C. Bishop, *Vet. Res. Com.*, **21**, 213 (1997).
3. R.D. Vidyarthi, A Text Book of Zoology, S. Chand & Co., New Delhi, edn. 14 (1967).
4. O. Ajaiyeoba, P.A. Onocha and O.T. Olarenwaju, *Pharm. Biol.*, **39**, 217 (2001).
5. Y.M. Shivkumar and V.L. Kumar, *Pharma Biol.*, **41**, 263 (2003).
6. A.A. Gbolade and A.A. Adeyemi, *Fitoterapia*, **79**, 200 (2008).
7. S.K. Jain, Dictionary of Indian Folk Medicine and Ethnobotany. Deep Publ., New Delhi, India (1991).
8. K.K. Singh and J.K. Maheshwari, *J. Econ. Tax. Bot.*, **4**, 829 (1983).
9. N.I. Bhuiyan, J.U. Chowdhury and J. Begum, *Bangladesh J. Bot.*, **37**, 213 (2008).
10. R.N. Almeida, M.F. Agra, F.N.S. Maior and D.P. DeSousa, *Molecules*, **16**, 2726 (2011).