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Study on effect of *Gelsemium sempervirens* potencies on central nervous system of experimental animals

P Bhuvanesh

Associate Professor, Department of Pharmacy, Sri Sairam Homoeopathy Medical College, Chennai, Tamil Nadu, India

Abstract

The work on clinical study and research work is to show the biological activity of *Gelsemium Sempervirens* on Central Nervous System using Actophotometre. One dose of *Gelsemium Sempervirens* 6, 200(0.1ml/kg), One dose of Nux vomica6x, 200 (0.1ml/kg) is administered orally to the Albino mice once in a day. The effect CNS locomotor activity is measured using the Actophotometre. The study that the action of drugs *Gelsemium Semperiverens* has signicant depressant activity on central nervous system, when compared with the Nux vomica.

Keywords: Gelsemium sempervirens, CNS activity, actophotometre, albino mice

Introduction

Gelsemium Sempervirens, parts of this plant contain the strychnine-related alkaloids Gelsemine Gelseminine. Which has a historical usage in the treatment of facial and other neuralgias, cardiac depressant and in spasmodic affections, but in this study we going to analyze the active principle of Gelsemium Sempervirens in central nervous system with Nux vomica, which has strychnine as alkaloid as using Actophotometre and Albino mice. The locomotor activity (horizontal activity) can be easily measured using an actophotometer which operates on photoelectric cells which are connected in circuit with a counter. When the beam of light falling on the photo cell is cut off by the animal, a count is recorded. An actophotometer could have either circular or square arena in which the animal moves. Both rats & mice may be used for testing in this equipment. The GABA receptors are a class of receptors that respond to the neurotransmitter gammaamino butyric acid (GABA), the chief inhibitory neurotransmitter in the vertebrate central nervous system. GABA is the chemical messenger that widely distributed in the brain. It reduce the activity of neuron to which it binds and also serve to control fear and anxiety experienced when neurons are overexcited. GABA receptors are probably the most common kind in the mammalian nervous system. It is estimated that close to 40% of the synapses in the human brain work with GABA and therefore have GABA receptors.

Aims and Objectives

To evaluate the effects of Drugs in Experimental Animals. To evaluate the efficacy of potencies of Nux vomica and *Gelsemium semperiverens* on Central nervous system of Experimental animals.

Material and Methods

1. Preparation of Drugs

The plant of *Gelsemium Sempervirens* was collected from Ooty under the supervision of Dr. D. Suresh Baburaj, Asst Director, medicinal plants survey and collection unit, CCRH

Ooty. The mother tincture of *Gelsemium Sempervirens* extracted as per the directions given in Homoeopathic Pharmacopoeia of India (Vol-1, 1971). The 6x potency of *Gelsemium Sempervirens* was prepared and 200C was brought from reputed firm used in this experiment.

The seed of Nux vomica plant is collected from was collected from Ooty under the supervision of Dr. D. Suresh Baburaj, Asst Director, medicinal plants survey and collection unit, CCRH Ooty. The mother tincture of *Nux vomica* extracted as per the directions given in Homoeopathic Pharmacopoeia of India (Vol-1, 1971). The 6x potency of *Nux vomica* was prepared and 200C was brought from reputed firm used in this experiment.

2. Experimental design

Male albino mice (swiss strain) were procured from Sri Venkateswara Enterprises, Bangalore and bred in animal house of Vinayaka Mission's College of Pharmacy, Salem. They were fed with commercial diet (Hindustan lever, Bangalore) and water adlibitum during the experiments. The pellet food containing 22.5% protein, 72.55% carbohydrate, 5% fat, and sufficient vitamins and minerals. The cages were placed in well ventilated place in the laboratory and were provided with 1.5 inches rice bran bedding, which was changed every day. The room temperature maintained at 25± 10c. The animals were selected randomly. The animals were divided into Four groups (I-IV), each comparising of six animals each weighing between 25- 30gms were selected.

Each group is given with 0.1ml/kg of *Gelsemium Sempervirens* 6,200 and Nux vomica 6,200 and standard drug chlorpromazine Hydrochloride (Dose: 3 mg/kg, ip; make a stock solution containing 0.3 mg/ml of the drug & inject 1 ml/100 g body wt of mice) and activity is tested using Actophoto meter. The instrument used for testing CNS activity, which operates photo electric cell.The beam of light is cut by the animal. The mice in each group is tested for prior to administration of drug for normal activity and after 30mins re-test each mouse for activity scores for 10mins after administration of corresponding drug in each

group. Note the difference in the activity, before & after is noted for each group.

Gamma-aminobutyric acid (GABA) is an amino acid which is the primary inhibitory neurotransmitter in the brain and a major inhibitory neurotransmitter in the spinal cord. It exerts its primary function in the synapse between neurons by binding to post-synaptic GABA receptors which modulate ion channels, hyperpolarizing the cell and inhibiting the transmission of an action potential.

Screening Methods for CNS Activity

The blind screening methodology, well known in psyconeuro pharmacology was utilized for central nervous system action investigations.

Drug Administration

1. Gelsemium Sempervirens

GROUP-1-Control group without any treatment

GROUP-11-Group treated with standard drugs (dose according to drugs used)

GROUP-111-Group treated with *Gelsemium Sempervirens* 6X (0.5ml/p.o) once in a day.

GROUP-1V-Group treated with *Gelsemium Sempervirens* 200C (0.5ml/p.o) once in a day.

2. NUX vomica

GROUP-1-Control group without any treatment

GROUP-11-Group treated with standard drugs (dose according to drugs used)

GROUP-111- Group treated with Nux vomica 6X (0.5ml/p.o) once in a day.

GROUP-1V- Group treated with Nux vomica 200C (0.5ml/p.o) once in a day.

The test with different drugs is made in two different occasions. The tests were made three hours after administration of drug.

Observation

- 1. The experimental animals used for the study, showed the loco motor activity at different levels and the findings are below.
- The mice with Control group showed the hyper active behavior.
- 3. The mice treated with Standard drug showed the decrease in loco motor activity.
- 4. The mice treated with Gelsemium semperiverns 6 has the hypo motility in action
 - The mice treated with Gelsemium semperiverns 200 also has decrease in loco motor action.
- 5. The mice treated with Nux vomica 6 has the moderate action of hypo motility

The mice treated with Nux vomica 200 has the slight hypo motility in action

The mice taken for the experimental studies are fed with appropriate food and water

Result

The loco motor activity before and after treatment is compared with the standard drug during the process of the experiment. The experimental results were tabulated below and the percentage has been estimated.

Table 1: Evaluation of CNS activity of Extracts of Nux vomica

C	Treatment	DOSE	Loco motor activity score for 10mins		Percentage change in
5.no			Before T/t	After T/t	activity
1.	Control	0.1ml	590.4 <u>+</u> 76.72	592.16 <u>+</u> 62.13	
2.	CPZ (Standard)	3mg/kg	590.6 <u>+</u> 94.79	88.33 <u>+</u> 28.20	85.04
3.	Nux vomica 6	0.1ml	553.33 <u>+</u> 64.47	253.0 <u>+</u> 93.44**	54.02
4.	Nux vomica 200	0.1ml	579.20 <u>+</u> 50.48	385.0 <u>+</u> 18.57*	33.52

All values are expressed as mean <u>+</u> S.E.M. p*<0.05 considered significant (n=6).Data will be analyzed by ANOVA followed by Dunnett's multiple comparison test

Data 1: Efficacy of nux vomica on cns activity

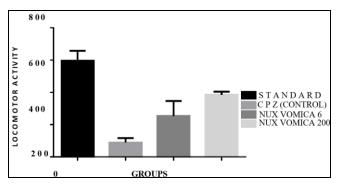


Fig 1

Table 2: Evaluation of CNS activity of Extracts of *Gelsemium Sempervirens*

S.no	Treatment	Dose	Loco motor activity score for 10mins		Percentage change in
			Before T/t	After T/t	activity
1.	Control	10ml/kg	342.20 <u>+</u> 15.30	340.30 <u>+</u> 5.8	
2.	CPZ (Standard)	3mg/kg	320 <u>+</u> 35.20	76.6 <u>+</u> 4.1	67.27
3.	Gels semp	0.1ml	337 <u>+</u> 80.08	168 <u>+</u> 9.3***	50.14
4.	Gels semp 200	0.1ml	513.33 <u>+</u> 40.46	169 <u>+</u> 8.17***	67.27

All values are expressed as mean \pm S.E.M. p*<0.05 considered significant (n=6).Data will be analyzed by ANOVA followed by Dunnett's multiple comparison test.

Data 2: Efficacy of gels semperiverns on cns activity

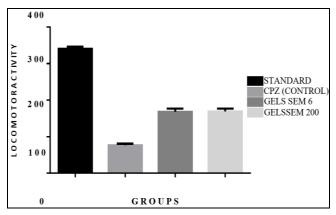


Fig 2

Stastical Analysis

The results were statically processed by Analysis of variance test ANOVA to compare the effects on treatment groups before and after treatment with control group Dunnett's *t* test were used to determine the effect of each treatment against the control group.

Discussion and Conclusion

Every drug has its action on the Human physiological system and here the drugs used for the study has its effects on the Central nervous system of experimental animals.

In Nux vomica has its action on different parts of the body but most probably on CNS, which causes paralysis and convulsions. In short it can be used to depress central nervous system activities. The work of the study also suggests that it can be used for the depressant activity.

Gelsemium semperiverens, As said earlier Barthlow says "I have demons trated that Gelsemium is a paralyzer of motility and sensibility, that sensibility is first affected in cold blooded animals (frog) and latter motility. The study shows that Gelsemium semperiverns has most significant activity on CNS depressant activity.

The study that the action of drugs Gelsemium Semperiverens has signicant activity on central nervous system, when compared with the Nux vomica.

The result of the study gives the opportunity for future researches.

References

- Amites Gangopadhyay, Jadupati Malakar et al. The Central Nervous System Activity of Barleria prionitis Linn. On the Locomotor Activity of swiss albino mice using Actophotometer, International Journal of Pharmaceutical & Biological Archives,2012:3(2):403-405.
- 2. Anitha K, Ranjith K *et al.* Anti-cataleptic activity of ethanol extract of *Ageratum conyzoides linn.*, International Journal of Innovative Drug Discovery,2012:2:48-54.
- 3. Arora Ankitkumar, Ashok M *et al.* Evaluation of Anxiolytic activity of aqueous and alcoholic extracts of leaves of *Crataegus oxycantha* in mice, International Journal of Pharma and Biomed Science, oct,2011:2(3):86-91.
- 4. Arun Kumar, Versha Parcha *et al.* Evaluation of CNS potential of *Hedychium spicatum*, Lambert Academic Publishing, 2012, 76.
- Ashish Dhir, Pattipati S Naidu et al. Modulatory Effect of Nimesulide on Apomorphine induced stereotypy and Mk-801 induced Hyperlocomotion, Annals of Neurosciences, 2005:12:1-5.
- Ashwaghosh Adhale, Subhash Yende, Ittadw AM et al. Pharmacological screening of Mimosa Pudica For its Alleged sedative hypnotic activity, International Journal Of Pharmacy & Technology, 2012:4:4451-4458.
- Ashish Dhir, Pattipati S Naidu et al. Modulatory Effect of Nimesulide on Apomorphine induced stereotypy and Mk-801 induced Hyperlocomotion, Annals of Neurosciences, 2005:12:1-5.
- 8. Avijit Chakraborty P, Amudha *et al.* Evaluation of anxiolytic activity of methanolic extract of *sapindus mukorossi gaertn*. In mice, International Journal of Pharma and Bio Sciences, 2010:1:1-8.

- 9. Balaji P, Thirumal M, Kumudhaveni B, Kishore G *et al.* Central nervous system depressant activity of *Barringtonia acutangula* (Linn.) Gaertn, Der Pharmacia Lettre, 2012:4(6):1786-1792.
- 10. Bhanumathy M, Shivaprasad *et al.* Anti-fatigue Activity of extracts of *Syzygium cumini* Leaves, International Journal for Pharmaceutical Research Scholars, 2013:2:1-1.
- 11. Chauhan Khushbu, Santwani Payal, *et al.* Evaluation of antidepressant activity of *Ailanthus Excelsa Roxb* using mice as experimental animal, Research journal of pharmacology and pharmacodynamics, 2011:3:102-104.
- 12. Choudhuri NM. A study on Materia medica, B.Jain publishers, Edt, 2001, 433-440, 732-740,
- 13. Deepika R Hemamalini *et al.* CNS Activity of the methanol extracts of *Solanum Pubescens* in experimental animal model, Journal of pharmacy and Biological sciences, 2013:5:48-51.
- 14. Dhayabaran D, Jeyaseeli Florance E *et al.* Anxiolytic and anticonvulsant activity of alcoholic extract of heart wood of *Cedrus deodara roxb*. in rodents, Journal of medicinal plants research, 4(14):1374-1381.
- 15. Farrington EA. Lectures on clinical Materia medica, B.Jain publishers edt, 2002, 187-196, 197-219.
- 16. Ghanshyam Yadav, Vipin Kumar Garg, Nishi Thakur *et al.* Loco motor activity of methanolic extract of *Saraca indica Bark*, Advances in Biological Research, 2013:7(1):01-03.
- 17. Habibur Rahman P, Muralidharan D *et al*. Evaluation of anxiolytic activity of ethanolic extracts from the leaves of *Trichosanthes cucumerina* L. in mice, Der Pharmacia Sinica, 2010, 86-94.
- 18. Homoeopathic pharmacopeia of India, Gelsemium Semperivens, Government of India, 1971:1:126
- 19. Homoeopathic pharmacopeia of India, Nux vomica, Government of India,1971:1:167
- Indurwade NH, Biyani KR et al. Evaluation of comparative and combined depressive effect of Brahmi, shankhpushpi and jatamansi in mice, Indian journal of medical science, 2000:5:339-341.
- 21. Joy Harris Hoskeri, Krishna Venkatarangaiah *et al.* CNS depressant activity of extracts from *Flaveria trinervia Spring C. Mohr*, Phytopharmacology,20111(4):100-107.
- 22. Kavita Gahlot, Abid M, Anupam Sharma *et al.* Pharmacological evaluation of *Gelsemium Sempervirens* roots for CNS depressant activity, International Journal of Pharmacy and Technology Research, 2011:3:693-697.
- 23. Kameshwaran Sugavanam, Suresh Velayutham *et al.* CNS depressent activity of different extracts of *Tecoma stans flowers*. Asian Journal of Traditional Medicines, 2012:7(1):39-42
- Kumar S, Madaan R, Sharma A. Pharmacological evaluation of bioactive principle *Turnera Aphrodisiaca*, Indian Journal of Pharma and Science, 2008:70(6):740-744.
- 25. Mahesh Radhakrishnan, Kumar, Baldev *et al.* Antidepressant-like activity of (4- phenylpiperazin-1-yl) (quinoxalin-2-yl) methanone (4a), a novel 5-HT3 receptor antagonist: An investigation in behaviour-based rodent models of depression, Indian Journal of Pharmacology,2012:44:560.

- 26. Maharudra S Rakh, Sanjay R Chaudhari, Evaluation of CNS depressant activity of *Momordica dioica roxbwilld fruit pulp*, International journal of pharmacy and pharmaceutical sciences, 2010:2:124-126.
- 27. Manjir Sarma, Kataki KT, Mani Senthil Kumar, *et al.* Neuro-psychopharmacological Profiling of Flunarizine: A Calcium Channel Blocker, International Journal of Pharm Tech Research, 2010:2:1703-1713,
- 28. Mayurr Bhurat. Effect of *Remusatia vivipara* (*roxb*) schott tubers on animal models of de pression International Journal of Current Pharmaceutical Research, 2011:3:90-92.
- 29. Nakur Singh, Sandeep *et al*. The study of analgesic and CNS stimulant activities of seeds of *Holarrhena antidysenterica* in laboratory animals, Journal of Pharmacology and Toxicology,2011:1(6):24-27.
- 30. Navanath MS, Naikwade NS, Mule SN. Analgesic antiinflammatory and CNS depressant activity of *Capparis decidua* Edgew flowers, Journal of Pharmacy Research, 2009:1-2:1307-1310.
- 31. Nimbal SK, Venkatrao N, Shivakumar Ladde *et al* Anxiolytic evaluation of *Benincasa Hispida* (*thunb*) *cogn.* fruit extracts, International Journal of Pharmacy and Pharmaceutical Science Research, 2011:1(3):93-97.
- 32. Pal A, Nayak S, Sahu PK *et al.* Piperine protects epilepsy associated depression: A study on role of monoamines, European Review for Medical and Pharmacological Sciences, 201115:288-1295.