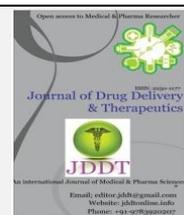


Available online on 15.02.2019 at <http://jddtonline.info>

Journal of Drug Delivery and Therapeutics

Open Access to Pharmaceutical and Medical Research

© 2011-18, publisher and licensee JDDT, This is an Open Access article which permits unrestricted non-commercial use, provided the original work is properly cited

Open  Access

Research Article

Effectual qualitative chemical evaluation of *Euphorbia neriifolia* Linn. by using fluorescence analysis

*¹Sharma Gaurav Kumar, ²Dhanawat Meenakshi¹Assistant Professor, Department of Pharmacy, Mewar University, NH-79, Gangrar, Chittorgarh -312901, Rajasthan, India. E-mail: garvsharma2050@gmail.com; Contact No. - +91-8290847107²Associate Professor, M. M. College of Pharmacy, Maharishi Markandeshwar (Deemed to be) University, Mullana, Ambala -133207, Haryana, India. E-mail: meenakshi.itbhu@gmail.com; Contact No. - +91-7726076396

ABSTRACT

Objective -Plants contain numerous biologically active compounds, many of these have been shown to exhibit therapeutic and antimicrobial properties and therefore they were in use as antimicrobial drugs in traditional medicines. *Euphorbia neriifolia* Linn. belongs to the family Euphorbiaceae. Assessment of medication implies affirmation of its personality and assurance of its quality and virtue and recognition of nature of debasement. Throughout the years the nature and level of assessment of rough medications has experienced an efficient changes. Hence, in the present study an attempt has been made for effectual qualitative evaluation of dried powdered leaves and stem of *Euphorbia neriifolia* Linn. by fluorescence analysis.

Design & Intervention- The dried powdered sample of *Euphorbia neriifolia* Linn. leaves and stem has been used to perform fluorescence analysis under the visible day light and UV light of short wavelength (254 nm) and long wavelength (365 nm) for their characteristic colour.

Main outcome measures- The ultra violet light creates fluorescence in numerous common items, which don't noticeably fluoresce in sunshine. In the event that the substances themselves are not fluorescent, they may frequently be changed over into fluorescent subordinates or disintegration items by applying diverse reagents. The hues created by these reagents speak to the nearness of dynamic constituents.

Results- The fluorescence character of any powdered character of any powdered medication is exceptionally particular and supportive distinctive highlights for the assurance of a medication.

Conclusion- The dried powder and herbal extracts of *Euphorbia neriifolia* Linn. leaves and stem showed varied fluorescence character which is an essential parameter for standardization of herbs.

Keywords: *Euphorbia neriifolia* Linn., Standardization, Fluorescence analysis, Evaluation, Euphorbiaceae.

Article Info: Received 14 Dec 2018; Review Completed 20 Jan 2019; Accepted 24 Jan 2019; Available online 15 Feb 2019



Cite this article as:

Sharma GK, Dhanawat M, Effectual qualitative chemical evaluation of *Euphorbia neriifolia* Linn. by using fluorescence analysis, Journal of Drug Delivery and Therapeutics. 2019; 9(1-s):44-47 <http://dx.doi.org/10.22270/jddt.v9i1-s.2248>

*Address for Correspondence:

Sharma Gaurav Kumar, Research Scholar & Assistant Professor, Department of Pharmacy, Mewar University, NH – 79, Gangrar, Chittorgarh, Rajasthan-312 901, India

1. INTRODUCTION

The restorative plants are of incredible enthusiasm to human wellbeing. The restorative plants (Rasayana) are the plants whose parts (leaves, seeds, stems, roots, natural products, foliage and so on.) removes, imbue, decoctions, powders have been broadly utilized in the Indian conventional (Ayurveda) arrangement of medication for the treatment of various sicknesses of people.¹

Plant based drugs have been a piece of customary social insurance in many parts of the world for a huge number of years.^{2, 3} Plants contain numerous biologically active compounds, many of these have been shown to exhibit therapeutic and antimicrobial properties and therefore they were in use as antimicrobial drugs in traditional medicines.

Plants utilized in conventional medication contain a huge range of substances that can be utilized to treat incessant and even irresistible ailments. As per a report of World Health Organization, over 80% of world's populaces rely upon conventional drug for their essential social insurance needs. Therapeutic properties of plants have likewise been researched in the light of later logical improvements all through the world, because of their strong pharmacological exercises, low poisonous quality and monetary suitability, when contrasted and engineered drugs.¹

Euphorbia neriifolia Linn. has a place with the family Euphorbiaceae develops lavishly around the dry, rough, sloping territories of North, Central and South India. *Euphorbia neriifolia* Linn. is worldwide dispersed in Baluchistan, Burma, India and Malaysian Islands. Inside

India, it is visit in rough ground all through Deccan Peninsula and Orissa. It is habitually cultivated for hedges in villages all over India. 4, 5 The scientific categorization of plant comprise of space: Eukaryota, kingdom: Plantae, sub-kingdom: Tracheobionta, division: Magnoliophyta, super-division: Spermatophyte, class: Magnoliopsida, sub-class: Rosidae, arrange: Euphorbiales, sort: Euphorbia, family: Euphorbiaceae and species: *neriifolia* Linn. ⁶

It is a herb loaded with spine, famously known as Sehund, Thohar and Milk Hedge. The leaves are thick succulent, 6 to 12 inches in length, ovular fit as a fiddle. Ayurveda portrays the plant as unpleasant, impactful, purgative, carminative, enhances hunger helpful in stomach inconveniences, bronchitis, tumors, loss of awareness, incoherence, leucoderma, heaps, aggravation, broadening of spleen, pallor, ulcers and fever. ^{7,8} The latex of *Euphorbia neriifolia* Linn. is a functioning element of numerous Ayurvedic definitions like Abhaya lavana, Avittoladi bhasma, Citrakadi taila, Jatyadi varti, Snuhidugdhadhi varti, Snuhi ghrta and Jalodarari ras. *Euphorbia neriifolia* Linn. has been customarily shown in Vatavyadhi, Gulma, Udara, Sula, Sotha, Arsas, Kusta and Medoroga. ^{9,10}

Assessment of medication implies affirmation of its character and assurance of its quality and virtue and recognition of nature of defilement. The assessment of an unrefined medication is important due to these fundamental reasons i) biochemical variety in the medications ii) deterioration because of treatment and capacity, and iii) substitution and contaminated, an after effect of thoughtlessness, numbness or misrepresentation.

Throughout the years the nature and level of assessment of unrefined medications has experienced a deliberate changes. At first, the unrefined medications were distinguished by examination just with the standard depiction accessible. Because of progression in the compound learning of rough medications, at present, assessment additionally incorporates strategy for evaluating dynamic constituents present in the unrefined medication, notwithstanding its morphological and infinitesimal examination. With the advent of separation techniques and instrumental analysis, it is possible to perform chemical evaluation of a crude drug, which could be both of qualitative and quantitative in nature. ¹¹

Hence, in the present study an attempt has been made for effectual qualitative evaluation of dried powdered leaves and stem of *Euphorbia neriifolia* Linn. by fluorescence analysis.

2. MATERIAL AND METHOD

2.1 Plant material- The leaves and stem of *Euphorbia neriifolia* Linn. were collected from the forest, a nearby place of Gangrar, Chittorgarh, Rajasthan. Their botanical identities were determined & authenticated by Scientist-in-Charge, Botanical Survey of India, Ministry of Environment and Forests, Govt. of India, Jodhpur- 342 008, Rajasthan, India. Several voucher specimen numbers were submitted to the authority for future references (Voucher Specimen Number-BSI/AZRC/I/12012/2018-19/326). The leaves and stem were dried under shade for 15 days, coarsely powdered and stored in air tight container for the further study.

2.2 Reagent and Chemicals- All reagents and chemicals used for fluorescence analysis and extraction were analytical grade obtained from SRL Chemical, Rankem, Otto, Himedia Pvt Ltd. India.

2.3 Extraction- Extraction of *Euphorbia neriifolia* Linn. leaves and stem carried out by Maceration and Soxhlation process with the use of various aqueous and non aqueous solvents. ¹¹

2.4 Fluorescence analysis- Take about 0.5gms of plant powder into clean and dried test tubes. To each tube 5ml of different organic solvents like distilled water, picric acid, glacial acetic acid, 1N HCl, 1N H₂SO₄, conc. HNO₃, ferric chloride (5%), iodine solution (5%), ammonia solution, 1N NaOH, potassium dichromate, HNO₃ + NH₃ solution, methanol, ethanol and toluene were added separately. At that point, every one of the cylinders was shaken and they were permitted to represent around 20-25 min. The solutions obtained were observed under the visible day light and UV light of short wavelength (254 nm) and UV light of long wavelength (365 nm) for their characteristic colour. ¹²⁻¹⁶

3. RESULTS

The behaviour of *Euphorbia neriifolia* Linn. leaves powder upon treatment with different chemical reagents showed Light green colour when powder was as such; Light green colour with distilled water; greenish yellow with picric acid; Light yellow with glacial acetic acid; Brownish yellow with 1N HCl; Magenta with 1N H₂SO₄; yellow with Conc. HNO₃; Green with ferric chloride (5%); Yellow with iodine solution (5%); Yellow green with ammonia solution; Orangish with 1N NaOH; Brown with potassium dichromate; Orangish with HNO₃ + NH₃ solution; Light Green with methanol, ethanol and toluene. Likewise the fluorescence characteristics of powdered leaves after treatment with different reagents emitted various colour radiations under ultraviolet light (Table-1). The fluorescence characteristics of different extracts of leaf was also studied under ordinary and UV light (365 nm), wherein the leaf extracts showed the visibility of varying colours which are tabulated in the Table-3.

The behaviour of *Euphorbia neriifolia* Linn. stem powder upon treatment with different chemical reagents showed Green colour when powder was as such; Green colour with distilled water; greenish yellow with picric acid; Yellow with glacial acetic acid; Brown with 1N HCl; Magenta with 1N H₂SO₄; yellow with Conc. HNO₃; Green with ferric chloride (5%); Yellow with iodine solution (5%); Yellow green with ammonia solution; Orangish brown with 1N NaOH; Dark brown with potassium dichromate; Orangish brown with HNO₃ + NH₃ solution; Green with methanol, ethanol and toluene. Likewise the fluorescence characteristics of powdered stem after treatment with different reagents emitted various colour radiations under ultraviolet light (Table-2). The fluorescence characteristics of different extracts of stem was also studied under ordinary and UV light (365 nm), wherein the stem extracts showed the visibility of varying colours which are tabulated in the Table-3.

S.N.	Treatments with leaves powder	UV Short (254 nm)	UV Long (365 nm)	Visible
1	Powder as such	Light yellow	Light yellow	Light green
2	Distilled water	Yellow green	Fluorescent green	Light green
3	Picric acid	Light green	Fluorescent green	Greenish yellow
4	Glacial acetic acid	Yellow	Light orange	Light yellow
5	1N HCl	Dark brown	Grayish	Brownish yellow
6	1N H ₂ SO ₄	Magenta	Magenta	Magenta
7	Conc. HNO ₃	Yellow	Dark brown	Yellow
8	Ferric Chloride (5%)	Dark brown	Fluorescent green	Green
9	Iodine solution (5%)	Greenish yellow	Brown	Yellow
10	Ammonia solution	Green	Fluorescent green	Yellow green
11	1N NaOH	Green	Fluorescent green	Orangish
12	Potassium dichromate	Dark green	Fluorescent green	Brown
13	HNO ₃ + NH ₃ solution	Yellow	Orangish	Orangish
14	Methanol	Light Green	Light Grey	Light Green
15	Ethanol	Light Green	Fluorescent green	Light Green
16	Toluene	Light Green	Yellow	Light Green

S.N.	Treatments with leaves powder	UV Short (254 nm)	UV Long (366 nm)	Visible
1	Powder as such	Yellow	Yellow	Green
2	Distilled water	Yellowish green	Green	Green
3	Picric acid	Mild green	Mild green	Greenish yellow
4	Glacial acetic acid	Yellow	Mild orange	Yellow
5	1N HCl	Brown	Dark brown	Brown
6	1N H ₂ SO ₄	Magenta	Magenta	Magenta
7	Conc. HNO ₃	Yellow	Dark Yellow	Yellow
8	Ferric Chloride (5%)	Dark brown	Mild green	Green
9	Iodine solution (5%)	Mild yellow	Dark brown	Yellow
10	Ammonia solution	Green	Mild green	Yellowish green
11	1N NaOH	Green	Mild green	Orangish brown
12	Potassium dichromate	Green	Mild green	Dark brown
13	HNO ₃ + NH ₃ solution	Yellow	Orangish	Orangish brown
14	Methanol	Green	Grey	Green
15	Ethanol	Green	Mild green	Green
16	Toluene	Green	Yellow	Green

S.N.	Solvent	Under Ordinary light	Under UV light (366 nm)
1	Pet-ether extract (L)	Yellowish green	Mild green
2	Benzene extract	Dark green	Mild blue
3	CHCl ₃ extract (L)	Light green	Light green
4	Ethyl Acetate extract (L)	Brownish green	Greenish brown
5	Ethanol extract (L)	Dark brown	Purplish Brown
6	Aqueous extract (L)	Dark Brown	Blackish brown
7	Aqueous extract (S)	Dark Brown	Blackish brown

(L)- Leave extract (S)- Stem extract

4. DISCUSSION

Fluorescence is a critical marvel shown by different concoction constituents present in plant material. A few constituents indicate fluorescence in the unmistakable range in sunshine. The ultra violet light delivers fluorescence in numerous characteristic items (for example alkaloids like berberine), which don't noticeably fluoresce in sunlight. On the off chance that the substances themselves are not fluorescent, they may frequently be changed over into fluorescent subsidiaries or decay items by applying

distinctive reagents, subsequently some rough medications are regularly surveyed subjectively thusly and it is an essential parameter of pharmacognostical assessment.¹⁷⁻¹⁹

The qualitative chemical evaluation of the drug was carried out with different concentrated mineral acids. The hues delivered by these reagents speak to the nearness of dynamic constituents. The fluorescence character of any powdered character of any powdered medication is exceptionally unmistakable and supportive distinctive highlights for the assurance of a medication. The

examination of powdered medication under bright light sets up the shade of the medication, all things considered and after treatment with various reagents.

5. CONCLUSION

In creating nations more than 80 percent of the populace depends on conventional medications, for the most part plant drugs, for their essential social insurance. The present study was conducted to evaluate the standardizing parameter for qualitative chemical evaluation of *Euphorbia neriifolia* Linn. with respect to fluorescence activity. The dried powder and herbal extracts of *Euphorbia neriifolia* Linn. leaves and stem showed varied fluorescence character which is an essential parameter for standardization of herbs.

ACKNOWLEDGMENT

On the occasion of presenting this article, It is my privilege to express my sincere thanks to my guide, mentor and supervisor Dr. Meenakshi Dhanawat, M. M. College of Pharmacy, Maharishi Markandeshwar (Deemed to be) University, Mullana, Ambala -133207, Haryana, India, Who has provided excellent guidance, valuable advices, and shared intelligent thoughts, criticisms and inculcated discipline. I am highly indebted to her for her valuable presence even in his busy schedule, which helped me to complete this work successfully. I extend my profound respect and heartfelt gratitude to my beloved Parents Late. Rajendra Kumar Sharma and Rajkumari. I also express my affection to my wife Deeksha and brother Kapil for their constant love, support, and encouragement throughout my life.

CONFLICTS OF INTEREST

We wish to confirm that there are no known conflicts of interest associated with this publication and there has been no significant financial support for this work that could have influenced its outcome.

REFERENCES

1. Vadlapudi V, Naidu KC. In vitro Bioautography of different Indian Medicinal plants. *Drug Invention Today* 2010; 2:53-56.
2. Chariandy CM, Seaforth CE, Phelps RH. Screening of medicinal plants from Trinidad and Tobago for antimicrobial and insecticidal properties. *J. Ethnopharmacol.*, 1999; 64:265-270.
3. Newman DJ, Cragg GM, Snader KM. The influence of natural products upon drug discovery. *Nat. Prod. Reports*, 2000; 17:215-234.
4. Anonymous. The wealth of India, a dictionary of Indian raw materials and industrial products (Raw materials), Vol. III (D-E). New Delhi: Central Institute of Medicinal and Aromatic Plants; 2003, p. 226-228.
5. Ved DK, Sureshchandra ST, Barve V, Srinivas V, Sangeetha S, Ravikumar K, et al. Plant details. Bengaluru: FRLHT's ENVIS Centre on Medicinal Plants; 2016. [Online]. Available from: http://envis.frlht.org/plant_details.php?disp_id=936&parname=0 [Accessed on 11th April 2017]
6. Anonymous. The plant list, version 1.1. 2013 [Online]. Available from: <http://www.theplantlist.org/tpl1.1/record/kew81079> [Accessed on 21st April 2017]
7. Nadkarni AK. *Indian Materia Medica*. Popular Prakashan, Bombay; 1976. p. 810-816.
8. Kirtikar KR, Basu BD. *Indian Medicinal Plants, II*, International Book Distributors, Dehradun 1996. p. 1581.
9. Chunekar KC. *Illustrated Dravyaguna Vijnana*. 2nd ed., vol. II. Varanasi: Chaukhambha Orientalia; 2005, p. 924-925.
10. Controller of Publications, Ministry of Health and Family Welfare, Department of Indian Systems of Medicine and Homoeopathy, Government of India. *The ayurvedic pharmacopoeia of India. Part-I. 1st ed., vol. I*. New Delhi: National Institute of Science Communication (CSIR); 2001, p. 100.
11. Kokate CK, Purohit AP, Gokhale SB. *Pharmacognosy*. Nirali Prakashan; 38th edition, Pune: 2007.
12. Huang CS, Yin MC, Chiu LC. Antihyperglycemic and antioxidative potential of *Psidium guajava* fruit in streptozotocin-induced diabetic rats. *Food Chem Toxicol.* 2011; 49:2189-2195.
13. Kokate CK, *Practical Pharmacognosy*, 1st edn, Vallabh Prakashan, New Delhi, 1986, 111.
14. Usha S, Pannine J and Sharma HP, Pharmacognostic studies on *Artemisia scoparia* Waldst and Kit, *Proc Indian Acad Sci (Plant science)*, 1984; 93:151-164.
15. Anonymous, Guidelines for the Assessment of Herbal Medicines, World Health Organization, Geneva, 1998, 8-9, 28-29, 30, 31-33.
16. Chase CR and Pratt RJ, Florescence of powdered vegetable drugs with particular reference to development of system of identification, *J Am Pharm Assoc*, 1949; 38:324-331.
17. Janchen D and Issaq HJ, Modern thin layer chromatography: advances and perspectives, *J Liquid Chromatogr*, 1988; 11:1941-1965.
18. Gupta MK, Sharma PK, Ansari SH, Lagarkha R. Pharmacognostical evaluation of *Grewia asiatica* fruits. *Int J Plant Science* 2006; 1(2):249-251.
19. Ansari SH. *Essentials of Pharmacognosy*. Birla Publications Pvt. Ltd 1st edition. New Delhi: 2006.