



Received on 16 March, 2017; received in revised form, 01 June, 2017; accepted, 25 June, 2017; published 01 November, 2017

ETHNOBOTANICAL AND ETHNOPHARMACOLOGICAL ACTIVITIES OF *ARTEMISIA NILAGIRICA*, *LYONIA OVALIFOLIA*, *SARCOCOCCA SALIGNA* AND *TARAXACUM OFFICINALE*

Neha Sahu and K. R. Arya *

Botany Division, CSIR-Central Drug Research Institute, 10 Jankipuram Extension, Sitapur Road, Lucknow- 226031, Uttar Pradesh, India.

Keywords:

Artemisia nilagirica,
Lyonia ovalifolia, *Sarcococca saligna*, *Taraxacum officinale*,
Ethnobotany, Uttarakhand

Correspondence to Author:

Dr. K. R. Arya


Ph.D.,
Principal Scientist
Botany Division, CSIR-Central Drug
Research Institute, 10 Jankipuram
Extension, Sitapur Road, Lucknow -
226031, Uttar Pradesh, India.

E-mail: kr_arya@cdri.res.in

ABSTRACT: Indian medicinal plants and traditional knowledge have been an integral part of our lifestyle since ages and now became an important source for natural product research (NPR) and development. Due to its knowing ability and applications, this knowledge has long been used as a thriving source for discovery of new drug molecules. This study was conducted under the routine drug development program of the Institute with the aim to explore the first-hand ethnobotanical information of 4 important plants *i.e.* *Artemisia nilagirica*, *Lyonia ovalifolia*, *Sarcococca saligna* and *Taraxacum officinale* and to correlate their possible applications in future research and developmental activities. The reported information was collected from 86 informants (villagers and herbal practitioners) of Nainital, Almora and Bageshwar districts of Uttarakhand Himalaya, during 2014-2016. The popularity of individual plant species for their ethnopharmacological uses was calculated as percent citation (PC) and cross verified through other informants of studied areas as well as the literature reported for their modern therapeutic applications.

INTRODUCTION: During last decades, ethnomedicinal plants are being reconsidered as one of the main botanical resources for searching novel, bioactive or their derivatives for modern therapeutics to combat different kinds of life-threatening diseases. Many of the known drugs and therapeutic leads have been derived from ethnomedicinal plants used in different traditional system of medicines¹⁻³. It is evident that about 55% of recent chemotherapeutic drugs for the treatment of cancer are derived from plants or based upon natural product research (NPR)⁴.

Indian sub-continent is well known for its cultural, social and traditional diversity with several ethnic groups comprised of more than 84.4 million people with age-old culture, traditions, languages, lifestyle and healthcare systems. The greater Himalaya of Uttarakhand, India (3300 - 4290 M) constitutes richest plant diversity with remarkable medicinal flora comprising about 4000 endemic species *i.e.* 50% of total Indian Himalayan diversity along with a vast ethnobotanical knowledge⁵. Due to their significant applications, this knowledge has long been used as a thriving source of bioactive compounds and recent ethnobotanical and ethnopharmacological studies proved folklore knowledge as a powerful tool for searching lead or new drug molecules for modern therapeutics⁶⁻¹⁰. Religious inspiration, unavoidable factors of inaccessibility and lack of medicinal facilities in remote hilly areas seems to be the main causative factors for the dependency on local herbal

<p>QUICK RESPONSE CODE</p> 	<p>DOI: 10.13040/IJPSR.0975-8232.8(11).4818-25</p> <p>Article can be accessed online on: www.ijpsr.com</p>
<p>DOI link: http://dx.doi.org/10.13040/IJPSR.0975-8232.8(11).4818-25</p>	

practitioners (Vaidyas) for their primary healthcare. Evidently, chemical and pharmacological investigations on these ethnomedicines have very often provided novel bioactive compounds for modern therapeutics¹¹⁻¹³. Literature reveals that *Artemisia nilagirica*, *Lyonia ovalifolia*, *Sarcococca saligna* and *Taraxacum officinale* are traditionally well-known plant species in Uttarakhand Himalaya. A systematic search towards validation of ethnomedicinal claims is extremely important and still pending. This paper describes the recent insights of ethnobotanical information collected from 86 informants including common villagers and herbal practitioners of Nainital, Almora and Bageshwar districts of Kumaon, Uttarakhand state during 2014-2016 along with their known pharmacological activities and status of abundance in studied areas. On the basis of these documented insights, the reported plant species are under investigation for their modern therapeutical applications.

MATERIALS AND METHODS: Study sites *i.e.* Nainital, Almora and Bageshwar districts of Kumaon region, Uttarakhand state were selected for the present study. These sites varied in altitude from 800 MSL (mean sea level) to 1098 MSL, geomorphologic characters, substrate and ecological conditions. During this study (March 2014 to November 2016) a total number of 86 informants (male and female) aged between 45-75 involving common villagers (VG) and recognized traditional herbal healers (HP) known as *Vaidyas* were interrogated (**Table 1**). In each locality, all the informants (VG and HP) were interviewed directly in Kumaoni and Hindi languages. Interviews were arranged by village level health workers familiar with local languages and the medicinal plants used for the treatment. Information provided by them was cross verified from other localities as well as through literature.

Plant species (**Fig. 1**) were identified with the local names and identified taxonomically according to the Flora of District Garhwal, North West Himalaya, India¹⁴. Herbarium specimens were housed in Departmental Herbarium CSIR-Central Drug Research Institute, Lucknow, India. The species were listed in alphabetical order by scientific name, local name of the region, family, voucher specimen number and current status of

abundance. The popularity of individual plant species being utilized *i.e.* percent of citation (PC) was calculated as per the following formula⁹.

$$\text{Percent Citation (PC)} = \frac{\text{Number of respondents mentioned the use of particular species}}{\text{Total number of respondents interviewed}} \times 100$$

TABLE 1: DEMOGRAPHIC PROFILE OF INFORMANTS INTERROGATED DURING STUDY IN DIFFERENT STUDY AREAS

Study locations	Traditional healers (HP)		Common villagers (VG)		Total
	Male	Female	Male	Female	
Nainital	5	3	8	6	22
Almora	6	4	10	8	28
Bageshwar	6	8	10	12	36

Enumeration:

***Artemisia nilagirica* (CB Clarke) Pampanini:** English name: Mugwort; Hindi: *Dona*; Sanskrit: *Damanak*; Local name (Kumaon): *Pati*; (Garhwal): *Kunja*. Family: Asteraceae is a perennial aromatic herb or under shrub commonly distributed in places from 1800-3500M altitude throughout Western Himalayas. Voucher specimen number KRA 24494 is submitted to CSIR-CDRI departmental herbarium, Lucknow.

Ethnobotanical uses: Plant is mostly considered sacred and young juvenile leaves are used as incense (*Dhoop*) during different local ceremonies and also to purify the surrounding environments as a bio-insecticide. Leave juice is also given to children to kill intestinal worms. A paste of mature leaves is used to treat wound and sore. Commercially, young leaves are dried and used as an ingredients of *Agarbati* (incense).

Ethnopharmacological activities: Plant species is reported as anthelmintics, antiseptic, expectorant, astringent, aromatic, anti-inflammatory, appetizer, digestive and diuretic¹⁵⁻¹⁶ and is one of the most popular genus in Chinese traditional medicine, frequently used for the treatment of malaria, hepatitis, cancer, inflammation and also as antifungal, antibacterial, antiviral remedy¹⁷⁻¹⁹. It is also used to treat cough, asthma, leprosy, skin disease and epilepsy²⁰⁻²². The paste of leaves is used as an ointment in fevers and headache and contain cooling properties while juice aid in earache and eye inflammation²³.

The major phytoconstituents reported from this plants are camphor, β -caryophyllene, thujone and isothujone²⁴, α -pinene, myrcene, sabinene, 1,8 cineole, linalool, linalyl acetate, isoborneol, borneol, bomyl acetate²⁵, α -thujone, β -thujone, β -pinene, sesquiterpene lactones, exiguaflavone A and B, maccianin, terpinen-4-ol, germacrene D, eugenol, perillaldehyde, cuminaldehyde²⁶, lavandulyl acetate, bornyl heptfluorobutanoate, 4-nitrobenzoic acid-4-methoxyphenyl ester²⁷. 2 compound belonging to sesquiterpene lactone and flavonoid class were isolated from ethanolic extract¹⁶. Also, caryophyllene oxide, borneol, alpha-humulene, trans-beta- guaiane, trans-sabinene hydrate, 2-hexane - 1 - ol 2-hexene-1-ol²⁸ are reported from this plant. Earlier Sati reported a major content of monoterpenoids (79.91%) and sesquiterpenoids (18.25%)²⁹.

***Lyonia ovalifolia* (Wallich) Drude:** Locally known as *Angyar* or *Aiyar*; family Ericaceae is a deciduous medium size tree about 12-16 M high. Commonly distributed in the slopes of mountain Himalaya associated with Oak and *Rhododendron* forest between the altitude of 1200-3000 M in (Uttarakhand Himalaya) India, Pakistan, China Myanmar and S.E Asia. Voucher specimen number KRA 24495 submitted to CSIR-CDRI departmental herbarium.

Ethnobotanical uses: Plant species is commonly used to prepare various agricultural instruments, fuel, timber and as an important source of timber for construction of traditional houses. Young leaves are poisonous to the cattle, but mature leaves are commonly used as fodder for domestic animals and preparation of organic manures (Cow dung). Moreover, a paste of young twigs and seeds is sometimes used to treat pimples, wounds, and boils³⁰⁻³¹. Leaves are mixed with food grains as a bio-insecticide to protect them during storage³²⁻³³.

Ethnopharmacological activity: Alkaloids are the major class of compounds reported in this plant³⁴. Various compounds such as Lyoniol-A³⁵, scorhodomollolides A and D, grayanane, lyoniol D³⁶, hexacosane, hexacosanol, sitosterol, taraxerol, lyonin A-C³⁷, lyoninide, lyoniresinol³⁸, ovafolinins A-E³⁹, lyonitoxin, quercetin 3-galactoside, Apigenin, luteolin, quercetin and epicatechin⁴⁰ have been isolated from this plant.

Grayanane diterpenoids having specialized carbon skeletons with highly oxygenated functionalities are exclusively known to Ericaceae family⁴¹ and is extremely toxic to mammals, especially to the heart and nervous systems⁴². Moreover, lyoniol A, an amyostatic compound³⁵ is also reported to produce respiratory depression, hypotension⁴³ and induce paralysis of nerve centers and motor nerve terminals. In addition, plant also reported to possess antimicrobial, antioxidant activities³⁴, cAMP regulation activity⁴⁴ and cause muscle tremors³⁵.

***Sarcococca saligna* (D. Don) Muell-Arg:** Locally known as Piruli, Geru, Tilya or Tiliara belongs to family Buxaceae, is an evergreen shrub about 1.5 M high. The abundance of this plant is not very common but found in patches between 1200-2800 M in Oak (*Quercus leucotrichophora*) and *Rhododendron* forest of Uttarakhand Himalaya and is also distributed in Kashmir to Nepal, Afghanistan, Myanmar and Sri Lanka¹⁴. Voucher specimen number KRA 24498 submitted to CSIR-CDRI departmental herbarium.

Ethnobotanical uses: The percent citation for ethnobotanical uses of this plant is recorded very low as compared to other plant species (**Fig. 1**). However, young leaves are used as fodder for cattle and preparation of organic manure. The mature stem is used as fuel. Very rarely the leaf paste is known to use for the treatment of mouth infection and as antipyretic⁴⁵. A decoction of leaf and young stem is given for treatment of diabetes.

Ethnopharmacological activities: The pharmacological action of this plant is reported as antibacterial, antitumor, antiulcer and acetylcholine esterase inhibitory⁴⁶ and hypotensive⁴⁶⁻⁴⁸, respiratory disorders, gastrointestinal, liver diseases, syphilis, fever, inflammation, rheumatism⁴⁹, ganglion-blocking⁵⁰, Alzheimers disease⁴⁸, antileishmanial and antiplasmodial⁵¹⁻⁵². The plant species is known to possess a major source of sarcosine, saracodine and pachyaximine-A⁵³⁻⁵⁴. Tertiary alkaloid salignine; salignarine-C; salignenamide-A, B, 2,4-diacetoxy epipachysamine-D⁴⁸, hydroxy epipachysamine D, salignenamide E, and salignen amide F and axillarine C, axillarine F, sarcorine, N-3-demethyl saracodine, saligcinnamide, salignenamide A, vaganine A, axillaridine A,

sarsalignone, and sarsalignenone are important compounds of this plant⁵⁵.

Taraxacum officinale Weber: English name: Dandelion or bitterwort, locally known as Dudhee, Kanphool and Kanphulya belongs to family Asteraceae is a perennial herb. Commonly distributed throughout Himalayan range up to 4000 M altitude. Density and frequency of occurrence are 0.13 plants/M² and 7.5% respectively⁵⁶. The rate of exploitation is quite higher than its natural regeneration which may cause a serious threat for its genetic erosion in the near future. Voucher specimen number KRA 24497 is submitted to CSIR-CDRI departmental herbarium, Lucknow.

Ethnobotanical uses: Leaves of this plant are advised to use as a vegetable to treat anemic patients. Aqueous extract of roots is given for the treatment of different kinds of urinary problems particularly urinary tract infection, increase the flow of urine to get relieved from pain during urination. It is also used to treat skin problems and

cure jaundice. The root extract of this plant is used to treat headache, migraine and liver problems.

Ethnopharmacological activities: In modern therapeutics, it is used to treat infections, swelling, water retention, breast cancer, gall bladder disease, pneumonia, hepatitis, hypolepidemia and oxidative stress etc⁵⁷⁻⁵⁸. In Chinese, Arabian and Native American traditional medicine, it is used to treat a variety of diseases including cancer^{57, 59-61}. Roots of this plant are considered as a major source of taraxacin, acrystallin, taraxacerin, taraxerol, taraxerone, arnidiol and faradiol etc⁶²⁻⁶⁴. Moreover, leaves of dandelion have been reported to be rich source of polyphenolic content of which hydroxyl cinnamic acid derivatives are predominant in both flowers and leaves⁶⁵⁻⁶⁷. Triterpenoids of plant origin have been reported to exhibit *in vitro* as well as *in vivo* cytotoxicity with low toxicity against mammary tumors⁶⁸⁻⁶⁹. This plant was also recommended in FDA list as safe nutraceuticals and has been approved by European council⁷³.

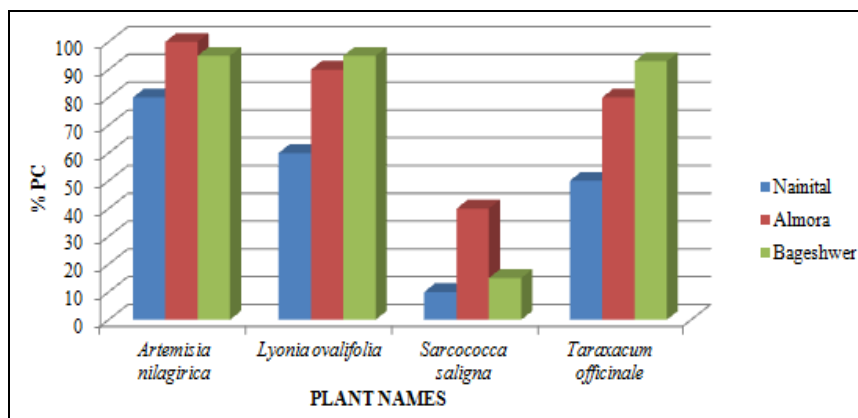


FIG. 1: CUMULATIVE PERCENT CITATION (PC) OF INDIVIDUAL PLANT SPECIES FOR DIFFERENT ETHNOBOTANICAL USES IN NAINITAL, ALMORA AND BAGESHWAR DISTRICTS OF KUMAON, UTTARAKHAND

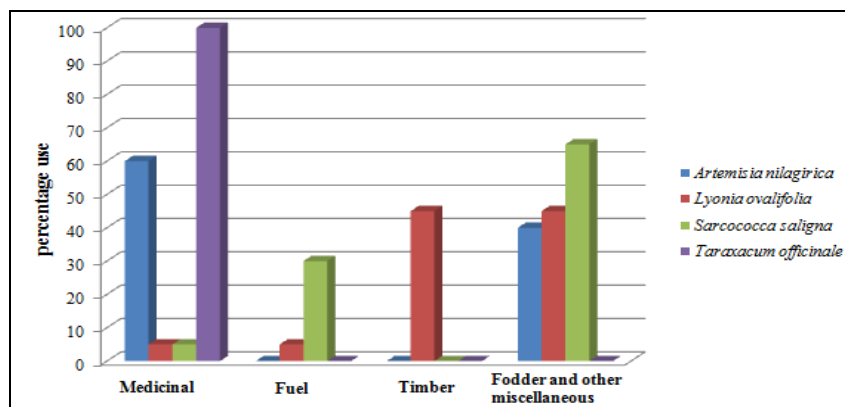


FIG. 2: PERCENT CITATION OF INDIVIDUAL PLANT SPECIES FOR DIFFERENT ETHNOBOTANICAL PURPOSES

DISCUSSION: Throughout history, nature and its resources have afforded a rich and important source of phytochemicals for different pharmaceutical applications in the field of medicine, cosmetics, pharmacy and biology. Of them, indigenous knowledge of medicinal plants and their uses have been considered as a unique source of raw materials and offer great opportunities for identification and isolation of novel pharmacological agents. These resources showed a strong relationship between ethnopharmacological use and medicinal properties of chemical compounds identified from them. Our recent past ethnobotanical and pharmacological studies also attested and proved its potential and utility to identify and isolate some valuable naturally occurring compounds for various modern therapeutical applications^{6-10, 12-13, 70-71}.

The higher frequency of percent citation popularity (80-100%) of *A. nilagirica*, *T. officinale* and *L. ovalifolia* (**Fig. 1**) in their day to day domestic needs and primary health care system reveals the importance of these plants in Uttarakhand Himalaya. Utility and percent citation of these ethnobotanical plants (**Fig. 1** and **2**) reflected a close relationship between geography, ecology, environment and depending upon the needs of local inhabitants which forced them to utilise these available natural resources on trial and error basis to meet out their day-to-day's needs as well as to combat their health related problems. The less popularity of *S. saligna* (5-40%), which is one of the richest sources of natural steroidal compounds and reported to use for the treatment of hypertension⁷² indicates the favourable environments to minimize these incidents in studies areas. But on other side, *L. ovalifolia* and *S. saligna* have a substantial utility (45-65%) for timber, fodder and other miscellaneous purposes (**Fig. 2**).

However, the cited chemical constituents of these plants showed tremendous scope for a number of pharmaceutical activities. Sore and wound healing properties of plants sometimes known to possess potential molecules for anti-cancer activities. This survey also provided some new insights on anti-diabetic and wound healing plants. These plants have been considered in our on-going research and developmental activities and their chemical and pharmacological investigations are under progress.

The study was also focused to get direct interaction with the user of these medicinal plants to know their importance or new uses, local names, current biodiversity status and kinds of problems associated with their availability, identification, adulteration etc. During survey and interrogation, some informants showed their non-cooperative nature to provide relevant information, because they had some bitter experiences from some of the previous surveyors like bio-piracy and sharing of benefits arises from their indigenous knowledge.

Interestingly, during survey of this study, it was also observed that some of the informants were quite confused with the identity of *T. officinale* and its closely related genus *Launaea*. Both the species belongs to the same family and also leaf and flowers of both the plants have quite similarities in their morphology and colors. This improper identification seems to be the strong reason of inadequate authentication and adulteration in their herbal products. For pharmaceutical points of view, the detail pharmacognostical markers need to workout to maintain the quality of both herbal products. Simultaneously, Rapid demand and over-exploitation of *T. officinale* roots from nature seems to be a serious threat to its existence at their natural habitats and may cause the rapid loss of its population density in the near future.

ACKNOWLEDGEMENT: One of the senior authors K.R. Arya (KRA) is thankful to Director, Central Drug Research Institute, Lucknow for encouragement and CSIR-Network project Bioprospection (BSC-0106) for financial support. Neha Sahu is grateful to University Grant Commission (UGC) New Delhi for providing Fellowship.

DECLARATION OF INTEREST: The authors reported no potential conflict of interest.

REFERENCES:

1. Newman DJ and Cragg GM: Advanced preclinical and clinical trials of natural products and related compounds from marine sources. *Current medicinal chemistry* 2004; 11(13): 1693-1713.
2. Butler MS: The Role of Natural Product Chemistry in Drug Discovery. *Journal of Natural Products* 2004; 67 (12): 2141-2153.
3. Modak M, Dixit P, Londhe J, Ghaskadbi S and Devasagayam TP: Indian herbs and herbal drugs used for

- the treatment of diabetes. *Journal of Clinical Biochemistry and Nutrition* 2007; 40(3): 163-173.
4. Newman DJ and Cragg GM: Natural products as sources of new drugs over the 30 years from 1981 to 2010. *Journal of Natural Products* 2012; 75(3): 311-335.
 5. Samant SS and Dhar U: Diversity, endemism and economic potential of wild edible plants of Indian Himalaya. *The International Journal of Sustainable Development & World Ecology* 1997; 4(3): 179-191.
 6. Arya KR and Agarwal SC: Folk therapy for eczema, bone fracture, boils, sores and gingivitis in Taragtal province of Uttaranchal. 2008.
 7. Arya KR, Sharma D and Kumar B: Validation and quality determination of an ethnobotanical lead for osteogenic activity isolated from *Ulmus wallichiana* Planch. *Journal of Scientific and Industrial Research* 2011; 70: 360-364.
 8. Sharan K, Siddiqui JA, Swarnkar G, Tyagi AM, Kumar A, Rawar P, Kumar M, Ngar GK, Arya KR, Manickavasagam L, Jain GK, Maurya R and Chattopadhyay N: Extraction and fraction from *Ulmus wallichiana* Planchon promotes peak bone achievement and have a nonestrogenic osteoprotective effects. *Menopause* 2010; 17: 393-402.
 9. Sharma C, Kumari T and Arya KR: Ethnopharmacological survey on bone healing plants with special reference to *Pholidota articulata* and *Coelogyne cristata* (Orchidaceae) used in folk tradition of Kumaon, Uttarakhand, India. *International Journal of Pharma Research And Health Sciences* 2014; 2(2): 185-190.
 10. Sharma C, Kumari T, Pant G, Bajpai V, Srivastava M, Mitra K, Kumar B, Arya KR: Plantlet formation via somatic embryogenesis and LC ESI Q-TOF MS determination of secondary metabolites in *Butea monosperma* (Lam.) Kuntze. *Acta Physiologiae Plantarum* 2015; 37(11): 239.
 11. Maurya R, Rawat P, Sharan K, Siddiqui JA, Swarnkar G, Mishra G, Manickavasagam L, Arya KR and Chattopadhyay N: Novel flavonol compounds, a bioactive extract/fraction from *Ulmus wallichiana* and its compounds for prevention for treatment of osteo-health related disorder. US Pat Application 2009 (110003).
 12. Sharan K, Mishra JS, Swarnkar G, Siddiqui JA, Khan K, Kumari R, Rawat P, Maurya R, Sanyal S and Chattopadhyay N: A novel quercetin analogue from a medicinal plant promotes peak bone mass achievement and bone healing after injury and exerts an anabolic effect on osteoporotic bone: the role of aryl hydrocarbon receptor as a mediator of osteogenic action. *Journal of Bone and Mineral Research* 2011; 26(9): 2096-3111.
 13. Sharma C, Dixit M, Singh R, Agrawal M, Mansoori MN, Kureel J, Singh D, Narender T and Arya KR: Potential osteogenic activity of ethanolic extract and oxoflavadin isolated from *Pholidota articulata* Lindley. *Journal of Ethnopharmacology* 2015; 170: 57-65.
 14. Gaur RD: Flora of the District Garhwal, North West Himalaya. Transmedia 1999.
 15. Kirtikar KP and Basu BD: Indian Medicinal Plant. 2nd edition Periodical expert, New Delhi 1975; 887: 4.
 16. Devmurari VP, Pandey S, Goyani MB and Jivani NP: Phytochemical screening of ethanolic extract of *Artemisia nilagirica*. *International Journal of Chemical Sciences* 2010; 8(4): 2099-2104.
 17. Simon JE, Charles D, Cebert E, Grant L, Janick J and Whipkey A: In *Artemisia annua* L.: A Promising Aromatic and Medicinal, *Advances in New Crops*, Oregon, Janick, 1990; 522-526.
 18. Kaul VK, Nigam SS and Dhar KL: Antimicrobial activities of the essential oils of *Artemisia absinthium* Linn, *Artemisia vestitia* Wall, and *Artemisia vulgaris* Linn. *Indian Journal of Pharmacy* 1976.
 19. Lin CC and Kan WS: Medicinal plants used for the treatment of hepatitis in Taiwan. *The American Journal of Chinese Medicine* 1990; 18: 35-43.
 20. Jack RA: A case of tuberous sclerosis treated with *Artemisia vulgaris*. *British Homoeopathic Journal* 1987; 76(4): 204-206.
 21. Ipsen H, Formgren H, Løswenstein H and Ingemann L: Immunochemical and biological characterization of a mugwort (*Artemisia vulgaris*) pollen extract. *Allergy* 1985; 40(4): 289-294.
 22. Chopra RN, Nayar SL and Chopra IR: Glossary of Indian Medicinal Plants.
 23. Jain SP and Puri HS: Ethnomedicinal plants of Jaunsar-bawar hills, Uttar Pradesh, India. *Journal of Ethnopharmacology* 1984; 12(2): 213-222.
 24. Näf-Müller R, Pickenhagen W and Willhalm B: New irregular monoterpenes in *Artemisia vulgaris*. *Helvetica Chimica Acta* 1981; 64(5): 1424-1430.
 25. Uniyal GC, Singh AK, Shah NC and Naqvi AA: Volatile constituents of *Artemisia nilagirica*. *Planta Medica* 1985; 51(05): 457-458.
 26. Shafi PM, Nambiar MG, Clery RA, Sarma YR and Veena SS: Composition and antifungal activity of the oil of *Artemisia nilagirica* (Clarke) Pamp. *Journal of Essential Oil Research* 2004; 16(4): 377-379.
 27. Badoni R, Semwal DK and Rawat U: Altitudinal variation in the volatile constituents of *Artemisia nilagirica*. *International Journal of Essential Oil Therapeutics* 2009; 3: 66-68.
 28. Haider F, Kumar N, Naqvi AA and Bagchi GD: Oil constituents of *Artemisia nilagirica* var. *septentrionalis* growing at different altitudes. *Natural Product Communications* 2010; 5(12): 1959-1960.
 29. Sati SC, Sati N, Ahluwalia V, Walia S and Sati OP: Chemical composition and antifungal activity of *Artemisia nilagirica* essential oil growing in northern hilly areas of India. *Natural Product Research* 2013; 27(1): 45-48.
 30. Joshi AR and Joshi K: Ethnomedicinal plants used against skin diseases in some villages of Kali Gandaki, Bagmati and Tadi Likhu watersheds of Nepal. *Ethnobotanical Leaflets* 2007; 2007(1): 27.
 31. Chauhan PS, Bisht S and Ahmed S: Traditional and Ethnobotanical uses of Medicinal Trees in District Tehri Garhwal (Western Himalayas). 2016.
 32. Shah NC and Joshi MC: An ethnobotanical study of the Kumaon region of India. *Economic Botany* 1971; 25(4): 414-422.
 33. Joshi AR and Joshi K: Insecticidal Plants of the Bagmati Watershed, Nepal: Ethnobotany and Traditional Uses. *Journal Bionotes* 2004; 6(2).
 34. Acharya SR: Antioxidant and Antimicrobial Properties of Leaves of *Lyonia ovalifolia* Wallich. *International Journal of Pharmaceutical & Biological Archive* 2015; 5(4).
 35. Fukuda H, Watanabe K and Ito T: Pharmacology of Lyoniol-A, an amyostatic component from *Lyonia ovalifolia* var. *elliptica* (Ericaceae). I. *Yakugakuzasshi: Journal of the Pharmaceutical Society of Japan* 1969; 89(3): 382-388.
 36. Sakakibara J, Ikai K and Yasue M: Studies on the constituents of *Lyonia ovalifolia* Drude var. *elliptica* Hand.-Mazz. XVIII. Structure of diterpenoid, lyoniol-D and correlation of lyoniol-A (lyoniatoxin) with grayanotoxin-I. *Yakugakuzasshi: Journal of the Pharmaceutical Society of Japan* 1974; 94(12): 1534-1540.

37. Yasue M, Kaiya T and Wada A: Studies on the constituents of *Lyonia ovalifolia* Sieb. Et Zucc. var. elliptica Hand.-Mazz. VI. On the triterpenoid and steroidal components of the leaves. Yakugakuzasshi: Journal of the Pharmaceutical Society of Japan 1967; 87(5): 581-584.
38. Yasue M, Kato T, Kishida T and Ota H: Isolation of Lyoniol-A,-B, and-C, the Toxic Principles of *Lyonia ovalifolia* var. elliptica. Chemical & Pharmaceutical Bulletin 1961; 9(2): 171.
39. Kashima K, Sano K, Yun YS, Ina H, Kunugi A and Inoue H: Ovafofinins A-E, five new lignans from *Lyonia ovalifolia*. Chemical and Pharmaceutical Bulletin. 2010; 58(2): 191-194.
40. Kato Y: Phenolic constituents of *Lyonia ovalifolia*. Phytochemistry 1973; 12(9): 2302.
41. Zhou SZ, Yao S, Tang C, Ke C, Li L, Lin G and Ye Y: Diterpenoids from the Flowers of *Rhododendron molle*. Journal of Natural Products 2014; 77(5): 1185-1192.
42. Wang LQ and Qin GW: Chemical and bioactive studies of diterpenoids from Ericaceae. Natural Products Research and Development 1997; 4: 82-90.
43. Ono H, Fukuda H and Kudo Y: Excitation by lyoniol-a of vagal afferent nerves and the reflex autonomic and somatic actions in rats. Journal of Pharmacobio-Dynamics 1981; 4(12): 940-946.
44. Wu ZY, Li HZ, Wang WG, Li HM, Chen R, Li RT and Luo HR: Lyonin A, a New 9, 10-Secograyanotoxin from *Lyonia ovalifolia*. Chemistry & Biodiversity 2011; 8(6): 1182-1187.
45. Kiamuddin MK and Hye HK: Pharmacological activity of an alkaloid from *Sarcococca saligna*. Pakistan Journal of Scientific and Industrial Research 1970.
46. Gilani AU, Ghayur MN, Khalid A and Choudhary MI: Presence of antispasmodic, antidiarrheal, antisecretory, calcium antagonist and acetylcholinesterase inhibitory steroidal alkaloids in *Sarcococca saligna*. Planta Medica 2005; 71(02): 120-125.
47. Kirtikar K. R and B. D. Basu: Indian medicinal plants, Basu, India: LM Allahabad. Blatter, E. Causis JR & Mhaskar KS 1933, 1: 77.
48. Atta-Ur-Rahman, Choudhary MI, Khan MR and Iqbal MZ: Three new steroidal amines from *Sarcococca saligna*. Natural Product Letters 1998; 11(2): 81-91.
49. Ghayur MN and Gilani AH: Studies on cardio-suppressant, vasodilator and tracheal relaxant effects of *Sarcococca saligna*. Archives of Pharmacal Research 2006; 29(11): 990-997.
50. Qiu M, Nie R and Li Z: Study on the chemical structures and bioactive screening of Pachysandra alkaloids. Acta Botanica Yunnanica 1993; 16(3): 296-300.
51. Devkota KP, Iqbal Choudhary M, Ranjit R, Samreen and Sewald N: Structure activity relationship studies on antileishmanial steroidal alkaloids from *Sarcococca hookeriana*. Natural Product Research 2007; 21(4): 292-297.
52. Devkota KP, Lenta BN, Choudhary MI, Naz Q, Fekam FB, Rosenthal PJ and Sewald N: Cholinesterase inhibiting and antiplasmodial steroidal alkaloids from *Sarcococca hookeriana*. Chemical and Pharmaceutical Bulletin 2007; 55(9): 1397-1401.
53. Naem I, Khan N and Choudhary MI: Alkaloids of *Sarcococca saligna*. Phytochemistry 1996; 43(4): 903-906.
54. Miana GA and Kiamuddin M: Alkaloids of *Sarcococca saligna* Muel: salignine. Pakistan Journal of Scientific and Industrial Research. 1969; 12: 161.
55. Atta-ur-Rahman ZU, Khalid A, Anjum S, Khan MR and Choudhary MI: Pregnane-type steroidal alkaloids of *Sarcococca saligna*: a new class of cholinesterases inhibitors. Helvetica Chimica Acta 2002; 85: 678-688.
56. Bhat JA, Kumar M and Bussmann RW: Ecological status and traditional knowledge of medicinal plants in Kedarnath Wildlife Sanctuary of Garhwal Himalaya, India. Journal of Ethnobiology and Ethnomedicine 2013; 9: 1.
57. Sigstedt SC, Hooten CJ, Callewaert MC, Jenkins AR, Romero AE, Pullin MJ, Kornienko A, Lowrey TK, Slambrouck SV and Steelant WF: Evaluation of aqueous extracts of *Taraxacum officinale* on growth and invasion of breast and prostate cancer cells. International Journal of Oncology 2008; 32(5): 1085-1090.
58. Choi UK, Lee OH, Yim JH, Cho CW, Rhee YK, Lim SI and Kim YC: Hypolipidemic and antioxidant effects of dandelion (*Taraxacum officinale*) root and leaf on cholesterol-fed rabbits. International Journal of Molecular Sciences 2010; 11(1): 67-78.
59. Bae TW, Park HR, Kwak YS, Lee HY and Ryu SB: *Agrobacterium tumefaciens*-mediated transformation of a medicinal plant *Taraxacum platycarpum*. Plant Cell, Tissue and Organ Culture 2005; 80(1): 51-57.
60. Schütz K, Carle R, Schieber A: *Taraxacum*—a review on its phytochemical and pharmacological profile. The Journal of Ethnopharmacology 2006; 107(3): 313-323.
61. Clare BA, Conroy RS and Spelman K: The diuretic effect in human subjects of an extract of *Taraxacum officinale* folium over a single day. The Journal of Alternative and Complementary Medicine 2009; 15(8): 929-934.
62. Burrows S and Simpson J: The triterpene alcohols of *Taraxacum* root. The triterpene group Part IV. Journal of the Chemical Society (Part II) Journal of the Chemical Society 1938; 141: 2042-2047.
63. Hansel R, Kartarahardja M, Huang J and Bohlmann F: Sesquiterpene lactone-beta-D- glucopyranoside and a new eudesmanolide from *Taraxacum officinale*. Phytochemistry 1980; 19: 857-861.
64. Akashi T, Furuno T, Takahashi T, Ayabe SI: Biosynthesis of triterpenoids in cultured cells, and regenerated and wild plant organs of *Taraxacum officinale*. Phytochemistry 1994; 36(2): 303-308.
65. Williams CA, Goldstone F and Greenham J: Flavonoids, cinnamic acids and coumarins from the different tissues and medicinal preparations of *Taraxacum officinale*. Phytochemistry 1996; 42: 121-127.
66. Budzianowski J: Coumarins, caffeoyl tartaric acids and their artifactual methyl esters from *Taraxacum officinale* leaves. Planta Medica 1997; 63(3): 288-289.
67. Wolbis M, Krolikowska M and Bednarek P: Polyphenolic compounds in *Taraxacum officinale*. Acta Poloniae Pharmaceutica 1993; 50: 153-158.
68. Bishayee A, Ahmed S, Brankov N and Perloff M: Triterpenoids as potential agents for the chemoprevention and therapy of breast cancer. Frontiers in Bioscience 2011; 16: 980-996.
69. Takasaki M, Konoshima T, Tokuda K, Masuda K, Arai Y, Shiojima K and Ageta H: Anti-carcinogenic activity of *Taraxacum* plant II. Biological and Pharmaceutical Bulletin 1999; 22(6): 606-610.
70. Sharma C, Mansoori MN, Dixit M, Shukla P, Kumari T, Bhandari SP, Narender T, Singh D and Arya KR.: Ethanolic extract of *Coelogyne cristata* Lindley (Orchidaceae) and its compound coelogin promote osteoprotective activity in ovariectomized estrogen deficient mice. Phytomedicine 2014; 21(12): 1702-1707.
71. Sharma C, Rajendar K, Kumari T and Arya KR: Indian traditional therapies and bio-prospecting: their role in drug development research. International Journal of

Pharmaceutical Sciences and Research 2014; 5(3): 730-741.

72. Ahmad B, Azam S and Bashir S: Biological screening of the aerial parts of the *Sarcococca saligna*. *Journal of Medicinal Plants Research* 2010; 4(22): 2404-3010.

73. Code of Federal Regulations Title 21. Access data.fda.gov. N.p., 2017. Web. 23 May 2017.

How to cite this article:

Sahu N and Arya KR: Ethnobotanical and ethnopharmacological activities of *Artemisia nilagirica*, *Lyonia ovalifolia*, *Sarcococca saligna* and *Taraxacum officinale*. *Int J Pharm Sci Res* 2017; 8(11): 4818-25. doi: 10.13040/IJPSR.0975-8232.8(11).4818-25.

All © 2013 are reserved by International Journal of Pharmaceutical Sciences and Research. This Journal licensed under a Creative Commons Attribution-NonCommercial-ShareAlike 3.0 Unported License.

This article can be downloaded to **ANDROID OS** based mobile. Scan QR Code using Code/Bar Scanner from your mobile. (Scanners are available on Google Playstore)