# Antimicrobial Activity of Secondary Metabolites in Medicinal Plants: An Update

Poonam Arora<sup>1</sup>\*, Madhukar Garg<sup>1</sup>, Tanvi Gera<sup>1</sup>, Lavish Vaid<sup>1</sup>, Parul Sood<sup>2</sup>, Loveleen Kaur<sup>1</sup>\*, Paranjeet Kaur<sup>1</sup>, Sanjeev Kumar Sahu<sup>3</sup>

<sup>1</sup>Chitkara College of Pharmacy, Chitkara University, Punjab-140401

<sup>2</sup>School of Pharmacy, Chitkara University, Baddi Himachal Pardesh

<sup>3</sup>School of Pharmaceutical Sciences, Lovely Professional University, Punjab, India

\*Corresponding author: poonam.arora@chitkara.edu.in, loveleen.kaur@chitkara.edu.in

Abstract. Plants have filled in as a significant wellspring of elements for conventional drugs for centuries. Verifiable records and present day ethno botanical field examines feature their significance in the conventional treatment of irresistible malady. Be that as it may, plants form just a minor level of present collection of FDA-endorsed antimicrobial medications. The present article gives an outline of active components of plants as hint for other wellspring of antimicrobial agents to be used in the battle against microscopic organisms. It additionally surveys the ethno botanical way to deal with sedate disclosure and talks about various inventive focuses for future medication revelation endeavours in this field. Without an uncertainty, antimicrobials are miracle tranquilizers. They have represented a very long time against different irresistible infections and spared a huge number of lives. The ongoing disappointment of antimicrobials because of the sensational rise of multidrug-safe microbes and fast spread of new diseases, be that as it may, prompts wellbeing associations and the pharmaceutical business worldwide to change their methodology and to expand improvement of antimicrobials against quickly rising anti-infection safe microorganisms. In spite of the fact that there is thinking about elective wellsprings of characteristic antimicrobial molecules from plants having different methods of activity, some of which have been utilized in regular medication for a considerable length of time and have appeared to have serious impacts contrasted with other antimicrobials. This examination portrays plant intensity as an elective hotspot for antimicrobial agents.

#### 1 Introduction

Medicinal plants are primarily considered for the variety of ingredients of plants used to cure chronic and infectious diseases in the pharmaceutical industry. Even before the human race was able to uncover the nature of bacteria, the theory was well known that such plants also possessed antimicrobial concepts that we now define. Traditional herbal medicines possess historical and cultural beliefs lading to their worldwide. In day today scenario herb refers to woody and non-woody part of the plant [1].

Rich origins of antimicrobial agents are medicinal plants. Medicinal plants are the main supplier of a variety of drug molecules according to the WHO and 80 per cent of the world banks on herbal medicine and usage of plant extracts or active ingredient for a large portion of the conventional therapies. However, a comparatively recent field is a research analysis for the identification of their antimicrobial active compounds [2].

Infectious conditions are normal, particularly skin and mucosal infections. Fungi and bacteria are a major category of such skin pathogens. The most severe inoculating disorders include dermal irritation, folliculitis, degradation of the skin, acne, dermatitis, Rosaceae etc. For higher skin care items, multi drug-resistant bacteria have been a significant source. The indiscriminate usage of synthetic antimicrobial medications widely used for combating infectious diseases has contributed to the growth of multiple drug resistance. Immune-impaired individuals are also observed to have difficult to treat infections. An enticing solution to multi-drug resistant bacteria is a novel compound with variation in modes of antibiotic action against microbes. Medicines currently in use for the prevention of infectious diseases often raise significant questions regarding the health of medicines. Many synthetic medications cause adverse reactions. Antimicrobial compounds from possible plants should be explored to mitigate

© The Authors, published by EDP Sciences. This is an open access article distributed under the terms of the Creative Commons Attribution License 4.0 (https://creativecommons.org/licenses/by/4.0/).

this issue. These herbal medications are less toxic; they have no and cost-effective side effects. These combat infectious diseases successfully, almost comparable to synthetic antimicrobials [3-5].

Antimicrobials dependent on plants represent a huge, untapped supply of medicines and it takes the hour to further explore plant antimicrobials. Plant-based antimicrobials have significant potential for therapy. The current one signifies such plants recognizing the enormous capacity of plants as sources of antimicrobial drugs [6].

## 2 Categories of antimicrobial molecules obtained from plants

Plants have an almost boundless capacity. This is discretionary metabolites, of which 12,000 were at all times excluded, an approximate amount under 10% of the number. These compounds are often used as plant defence components by microorganisms, creepy crawls and herbivores to avoid [7,8]. Many plants, for example, terpenoids, are responsible for pigmenting seeds, for example quinones among tannins. Plants may incorporate pleasant scenting exacerbates that are phenol or their subordinate oxygen inlaid.

## 2.1 Terpenoids and Basic Oils

Plants are the reservoir of biochemical and pharmaceuticals producing infinite biochemical mixtures. Human beings and animals use a tiny portion of earth's plants throughout their lives (1-10%) (250,000–500,000 species). The scent of the plant arises from the expected essential quinta or division of the simple material. The two oils are available metabolites, and in isoprene-based blends are remarkably enhanced. The overall compound form is C<sub>10</sub>H<sub>16</sub> and it is developed as diterpenes (C20, C30 and C40) and tetraterpenes (C5) and hemiterpenes (C15). If the mixes contain external ingredients, normally oxygen, they are terpenoids [9].

Terpenoids are incorporated in acetic acid derivative units and give their foundations as well with unsaturated fats. They are distinct from unsaturated fats, since they are wide-ranging and cyclic. The mill is commonly used as an anti-malarial product, methanol, camphor (monoterpenes) and farnesol (sesquiterpenoids) as well as arteremisin (sesquiterpenoids), which are the mill terpenoids. The World Health Association's control board in 1985 decided to establish this last drug as a cure for cerebral intestinal illness [10,11].

Microorganisms, pathogens and protozoa are active toward terpenes. In 1977, 60 per cent of the specific petroleum substances that have been attempting to date are specifically parasite inhibitors, although 30 per cent of the simple petroleum are restricted to bacteria. Terpenes' function is not well understood, but lipophile blends that damage the membrane are recognized. Mendoza et al. thus observed that the rise in methyl production to improve diterpenoids' hydrophilic behaviour has completely reduced their antimicrobial action [12,13].

The terpenoids in simple plant oils are found to be beneficial to manage *Listeria monocytogenes* Basil, an inexpensive home grown, is used as 125 ppm of chlorine in the sanitization of lettuce leaves. In several Mesoamerican cultures, Chile peppers are a staple that has been used all around the globe. In excess of a craving for seasonal food, their usage will represent. Many important supplements are contained in chiles, such as nutrients and vitamins. Capsaicin, a terpenoid portion, has a broad range of human organic ability, influences apprehensive cardiovascular and stomach-based structures and finds usage as a treatment of pain. The proof is paired with its function with antimicrobials. Cichewicz and Thorpe discovered the improvement in the growth of *Candida albicans* by capsaicin to different stages, however it is also bactericidal to *H. pylori*, causing damage gum mucosa. An antifungal large range is an extra hot-taste, aframodialditerpene with a cameroonian fragrance [14,15].

The dissolvable part of violet prairie clover produces a petalostemumol terpenoid that has had excellent results on *S. aureus* as did *C. albicans*. Batista *et al.* identified two diterpenes that slowly separated their popularity; they served well in the battle against mentioned strains added by *P. aeruginosa* and *V. cholerae*. The scientists reported that the terpenoid portions found in 10 rats were used in the diagnosis of gastric ulcer by Mali residents to use a tree bark called *Ptelopsissuberosa*, while substantial ulcers were caused by rats. This is not known if such symptoms were induced by antimicrobial behaviour or gastronic mucosal tolerance. We observed that terpenes prevented ulcers, which reduced the occurrence of current ulcers. Kadota et al. have discovered that *H. pylori* can be specifically inhibited by trichorabdal An, a Japanese diterpene produced [16, 17].

Since plants are highly susceptible to infections caused by Pathogens, to safeguard themselves, they produce Phytoalexins, which provide them with antimicrobial property.

#### 2.2 Flavones, flavonoids, and flavanols

Complexes that comprise one carbonyl set (instead two carbonyl quinones). In order to manufacture flavanol, a 3hydroxyl bunch is added. In comparison to the hydroxylated phenolic compounds, flavonoids exist as a C6-C3 unit attached to the good smelling chain. As plants are known to combine them with the microbial diseases, the fact that they are effective antimicrobials against a wide variety of microbes in vitro should not be shocking. Their activity is likely to be complex with extracellular and solvent proteins, as mentioned above for quinone because of their relation to bacterial cell divisors. Lipophilic flavonoids can also kill microorganisms gradually [18].

Catechins, the C3 class with the least diminished flavonoid composition, are capable of uncommon analysis. Such flavonoids have been studied absolutely in oolong green teas because of their origin. Earlier it was discovered that tea had antimicrobial influence due to a combination of catechin blends. The two blends bore in vitro *Vibrio cholerae*, *Streptococcus mutans*, *shigella* and other microscopic species. Catechins also repressed reclusive glucose transferases. The last operation was performed in normal rodent concentrates in vivo in view of the complexing activities for the above-referenced quinones [19].

There are few inhibitory properties in flavonoid mixtures. Flavone subordinates is used in more than one test as an inhibitory respiratory Syncytial (RSV) virus.. Kaul *et al.* contained quercetin, naringin, hesperetin, and catechin procedures and methods in *in vitro* as single-layer monolayers. Catechin inhibited infectiousness but did not intracellularly replicate RSV and HSV-1; and quercetin had a clear impact in decreasing diseases. The authors propose that minor auxiliary varieties in mixed plants are important to its capabilities and point out a further advantage to several subordinate plant individuals: their low level of noxious diseases day by day the standard Western diet contains roughly 1 g of flavonoids (mixed); the pharmacologically complex quantities are unlike [20].

It was also found that the menthol extract derived from *Ficus hispida*, which were later found to exhibit antimicrobial properties attributed due to presence of mentioned components [21].

### 2.3 Phenolics and polyphenols

These are fundamental bioactive phytochemicals which are less complicated consist of a single, substituted phenolic chain. Cinnamic acid and caffeic acids are the most prominent cause of oxidation in the mill from a large range of blends of phenylpropane. Estragon and thyme plants have caffeic corrosives, which neutralize pathogens, microorganisms and growths [22,23].

The two hydroxylated phenols are catechol and pyrogallol and are harmful for microbes. Catechol molecule has two –OH, and 3 pyrogallol classes. The sites and the amounts of hydroxyl phenol bunches were having relative risk to microbes, with a rise in hydroxylation contributing to their enhanced threat. However, some researchers note that the more highly oxidized phenol molecules are slowly becoming inhibited. The ingredients that were assumed to be responsible for phenolic poisonous qualities for smaller animals involve chemical deficiency by oxidized blends, often through reaction of sulfhydryl collections or somewhat ambiguous protein connections. Phenolic intensifying compounds that have a lower oxidation level on the C3 side chain that do not include oxygen molecule are categorised as essential oils and have been used as antimicrobials for several years. Eugenol is a major molecule contained in clove oil which is treated as bacteriostatic by the two parasites and microorganisms [24].

#### 2.4 Alkaloids

The morphine is the primary restoration of an alkaloid and was extracted from the *Papaver somniferum*. Antimicrobial effects have usually been identified to be diterpenoid alkaloids, which are typically extracted from Ranunculaceae plants or buttercups. Solamargin and other alkaloids may be priced for HIV exposure by Solanumkhasianum and HIV-related bowel infections. While alkaloids (counting species of *Giardia* and *Entamoeba* have proven to have a microbicidal function, their simple anti-diarrheal activity is possibly attributable to the influence of their transit periods in the small intestine. Berberine is a significant molecule in the alkaloid network [25].

#### 2.5 Tannins

'Tannin' is an illustrative generic term for the array of polymeric phenolic compounds ready for tanning or promoting gelatin, regarded as astringency from arrangement. The sub-atomic loads vary from 500 to 3000 and can be present in nearly all areas of the plant: bark, wood, seeds, food derived from the field and hydrolyzed and thick tannins are classified into two groups. Alternatively, tannins are formed by the polymerisation of the quinone plant.

Tannins may be provided by developed up flavan branches that have been transferred to woody tissue. Their mixture has been very much considered as it has been suggested to fix or forestall a number of diseases by taking the tannins containing refreshments, particularly green teas and red wines [26]. Ellagic acid which falls under the category of hydrolysable tannin of true tannin was also found to show the antimicrobial property [27].

#### 2.6 Different mixes

Many of the suspected phytosanitary compounds have been identified to be antimicrobial. The purpose of this study is to concentrate on evidence of toxic concoctions believed to be concerned. Nevertheless, the antimicrobial properties linked to polyamines, isothiocyanates, thiosulfinates and glucosides should be reported, in particular spermidin. Estevez-Braun *et al.* reported a polyacetylene C17 plant innate to the Canary islands, confined from *Bupleurum salicifolium.*<sup>28</sup> Aqueous extract of *Celastrus paniculatus* was found to be effective against bacterial and fungal strains owing to presence of phenolic components [29].

There has been considerable evidence of the effect on antimicrobials from cranberry grip. People were simply advised to drink their juice to retain a safe buffer from or just to handle urinary tract contaminations. In the mid 1990's, scientists saw the monosaccharide fructose that was severely forested by pathogenic adsorption in cranberry and blueberry squeezes. *E. coli* of urinary tract epithelial cells, and is as plain as mannose. Scientific work has proven that cranberry juice has positive effects. Numerous organic products contain fructose, however, and specialists are currently searching for a subsequent dynamic cranberry juice compound which adds the antimicrobial characteristics of this current juice [30].

# 3 Advantages and difficulties of plants as a wellspring of antimicrobial

In any case, the present amount of reported antibacterial plant drugs does not necessarily reflect the potential of plant characteristic items to be used in prospective treatments as antimicrobials. In some degree, the development of growing plant products such as antimicrobial drugs faces certain characteristic challenges:

1. Plant concentrate is synthesized inconceivable – even more so than parasites for example, since a single arrangement of a concentrate will produce multiple different concoction components. The classification of individual mixtures into the perfect antimicrobial bioactivity can be repetitive [31].

2. There are challenges in rediscovering identical blends from different sites, and careful dereplication from that in the disclosure process needs minimize the times and actions of identified radioactive elements.

3. Links to plant samples may be challenging to pursue action courses once in a while, in a global setting in particular. Plant spectrum guidelines provide for and send / import grants contrast where the research is performed. In addition, the exchange of equal access and profit-sharing understanding for these works is needed in compliance with the guidance and direction given by the United Countries Display for Natural Decent Variety and the Nagoya Convention [32].

4. Many therapies on plants operate by synergistic strategies. Synergism between mixes in an uncertain combination is a kind of difficulty because the cognitive science has not yet completely matured to accept multiple intermixes which function as one with specific natural goals [33]. It may be claimed again that, with the creation of an anti-infection barrier, the synergistic movement of certain plant concentrates will bring extraordinary open doors. It begs the issue of how monotherapy will withstand much more unnecessarily complicated strategies by rendering it more impossible for organisations to establish a multi-faceted attack defence [34].

# 4 Discussion

Advance strategies are important in order to fill the antimicrobial product pipeline during a time of increasingly growing resistance to infection. Clearly, in pursuit of new approaches, there are a range of innovative methods to explore. Plants are a rare and untapped source of bioactive mixtures, so ethnobotanical investigations into instruments should provide potential investigations so as to specify the most possible source. It is also important to check complicated plant concentrate and individual mixes for motion against elective bacterial goals, e.g., destruction and pathogenesis, as well as host organized objectives, ignoring tests to conduct excellent bacteriostatic and bactericidal acts.<sup>35</sup> In reference to quoted above the antimicrobials activity shown by the active components in the herbal mixtures, and a portion of the herbal active components is given in table 1.

S. No	Common name	Scientific name	Compound	Class	Structures	Referen ce
	name					u
1	Neem	Azadirachtaindica	β-sitosterol	Triterpenes		[36]
					βsitosterol	
			Quercetin	Flavonol glycosides	HO C C C C C C C C C C C C C C C C C C C	
2	Ashwagand ha	Withaniasomniferu m	Withaferin A	Lactone		[37]
3	Black pepper	Piper nigrum	Piperine	Alkaloid	Withafarin A	[38]
4	Buttercup	Ramunculusbulbos us	Protoaneminin	Lactone	CH <sub>2</sub>	
5	Clove	Syzgiumaromaticu m	Eugenol	Terpenoid	HO Eugenol	[39]

#### compounds having antimicrobial properties with structures of active ingredients

Table 1 Main classes

of

6	Coca	Erythroxylum coca	cocaine	Alkaloid	H <sup>3</sup> C	[40]
		5 5			Ň Lo-or	
					Cocaine	
7	Eucalyptus	Eucalyptus	1,8-Cineole	Monoterpene	CH3	[41]
		globulus	(Eucalyptol)	-		
					Ŭ,	
					N	
					H <sub>3</sub> C CH <sub>3</sub>	
					1,8-Cineole	
8	Garlic	Allium sativum	Allicin	Sulfoxide	Î au	
					H <sub>2</sub> C S S CH2	[42]
					Allicin	['2]
			Ajoene	Sulfatedterpen		
				oid		
					"2" S S CH2	
					ů H	
	-	-			Ajoene	
9	Ginseng	Panaxnotoginseng	Ginsenoside- Rg1	Saponin	H <sub>3</sub> C	[43]
			Kgi		OH <sub>H</sub> H <sub>3</sub> C	CH3
					H <sub>3</sub> C CH <sub>3</sub> H HO	
					HO HO OH	
					но он	
					Ginsenoside-Rg1	
10	Hemp	Cannabis sativa	β-Resercyclic	Organic acid	HO	
			acid		F	[44]
					ОН	[44]
					он	
					β-Resorcilic acid	
L	1				P 1 toool one dold	

11	Henna	Lawsoniainermis	Gallic acid	Phenolic	о он	[45]
					Y	
					но	
					Ьн	
12	Licorice	Glycyrrhizaglabra	Glabrol	Phenolic	Gallic acid	[46]
				alcohol	Н3С СН3	[]
					HO	
					U Contraction of the second se	
13	Olive oil	Oleaeuropae	Hexanal	Aldehyde	Glabrol	
					¥ У У	[47]
					Hexanal	
14	Onion	Allium cepa	Allicin	Sulfoxide	0	
					H <sub>2</sub> C	[48]
15	Peppermint	Menthapiperata	Menthol	Terpenoid	Allicin CH3	[49]
15	reppermit	Meninapiperaia	Wention	reipenoid		[49]
					Ц	
					нас сна	
					Menthol	
16	Rauwolfia	Rauwolfiaserpenti na	Reserpine	Alkaloid		[50]
					CHIO H	
					H I I I	6
						-OCH3
					Reserpine	fa
17	Senna	Cassia agustifolia	Rhein	Anthraquinone		[51]
					∥ ∣	
10			<b>TT</b>		Rhein он о он	
18	St. John's wort	Hypericumperforat um	Hypericin	Anthraquinone	OH O OH	
					н <sub>з</sub> с он	[52]
					H <sub>3</sub> C OH	
					ОН О ОН	
		·	•	·	-	·

					Hypericin	
19	Turmeric	Curcuma longa	curcumin	Terpenoids	"Q <sup>sli</sup> Q.	[53]
20	Apple	Malussylvestris	Phloretin	Flavonoid	Hjor <sup>d</sup> Curcumin OH Q	
20	Арре	manussyrvestris	rmoreun	derivative	но он	[54]
21	Betel pepper	Piper betel	Catechol's	Essential oils	OH	[55]
			Eugenols		Catechol's	
22	Green tea	Cameliasinensis	Catechin	flavonoid	Eugenol HO OH OH Catechin	[56]
23	Savory	Saturejamontana	Carvacrol	Terpenoid	CH <sub>3</sub> OH H <sub>3</sub> C CH <sub>3</sub> Carvacrol	[57]
24	Glory lily	Gloriosasuperba	Colchicine	Alkaloid		[58]
25	Fava bean	Viciafaba	Fabatin	Thionin	Colchicine Cl N H	[59]

					Fabatin	
26	Allspice	Pimentadioica	Eugenol	Essential oils	о <sup>_сн</sup> <sub>3</sub> [60]	
					HO	
					CH2	
					Eugenol	
27	Cashew	Anacardium	Salicylic acid	Polyphenols	H0 0 [61]	
		pulsatilla				
					но	
					Salicylic acid	
28	chamomile	Matricariachamo	Anthemic acid	Phenolic acid		
		milla			[62]	
					но снз	
					$\mathbf{v}$	
					CH3	
					chrysanthemic acid	
29	Chili	Capsicum annum	Capsaicin	Terpenoid	HO	
	peppers,					
	paprika				0	
20	G + 1 1				Capsaicin	
30	Gotu kola	Centellaasiatica	Asiatocoside	Terpenoid	[64]	
					O OH	
					HO HH HO O	
					"	
					но он	
					<sup>Õн</sup> Asiatocoside	
31	Hops	Humuluslupulus	Lupulone,	Phenolic acids	[65]	
			humulone			
					ОН	
					UH O .	
					Lupulone	
1						
					но	
					OH	

					Humulone	
32	Legume	Milletiathonningii	Alpinumisoflav one	Flavone		[66] ОН
33	Mountain tobacco	Arnica montana	Helanins	Lactones	Alpinumisoflavone	[67] O
34	Periwinkle	Vinca minor	Reserpine	Alkaloid	$\begin{array}{c} & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ &$	[68]
35	Tree bard	Podocarpusnagi	totarol	Flavonol	OH H Totarol	[69]

**Acknowledgements** The authors would like to thank the Chitkara College of Pharmacy, Chitkara University, Punjab, for providing institutional resources for data compilation.

## **Conflicts of Interest**

None

# **5** References

- Bourinbaiar AS, Lee-Huang S. The Activity of Plant-Derived Antiretroviral Proteins MAP30 and GAP31 against Herpes Simplex Virus Infectionin Vitro. Biochemical and biophysical research communications. 1996 Feb 27;219(3):923-9. doi: 10.1006/bbrc.1996.0334.
- 2. Anand U, Jacobo-Herrera N, Altemimi A, Lakhssassi N. A comprehensive review on medicinal plants as antimicrobial therapeutics: potential avenues of biocompatible drug discovery. Metabolites. 2019 Nov 1;9(11):258. DOI: 10.3390/metabo9110258.
- Kubo I, Muroi H, Kubo A. Naturally occurring antiacne agents. Journal of natural products. 1994 Jan;57(1):9-17. Doi: 10.1021/np50103a002.
- Ahmed AA, Mahmoud AA, Williams HJ, Scott AI, Reibenspies JH, Mabry TJ. New sesquiterpene αmethylene lactones from the Egyptian plant Jasonia candicans. Journal of natural products. 1993 Aug;56(8):1276-80. Doi: 10.1021/np50098a011.
- 5. Jones Jr SB, Luchsinger AE. Plant systematics. McGraw-Hill.; 1986 Dec. DOI: 10.4236/ahs.2015.44023.

- 6. Hunter MD, Hull LA. Variation in concentrations of phloridzin and phloretin in apple foliage. Phytochemistry. 1993 Nov 1;34(5):1251-4. DOI: 10.1016/0031-9422(91)80010-x.
- Lv Y, Tian T, Wang YJ, Huang JP, Huang SX. Advances in chemistry and bioactivity of the genus Erythroxylum. Natural Products and Bioprospecting. 2022 Dec;12(1):15. doi: 10.1007/s13659-022-00338z.
- Prakash A, Vadivel V, Banu SF, Nithyanand P, Lalitha C, Brindha P. Evaluation of antioxidant and antimicrobial properties of solvent extracts of agro-food by-products (cashew nut shell, coconut shell and groundnut hull). Agriculture and Natural Resources. 2018 Oct 1;52(5):451-9. https://li01.tcithaijo.org/index.php/anres/article/view/231843.
- 9. Islas JF, Acosta E, Zuca G, Delgado-Gallegos JL, Moreno-Treviño MG, Escalante B, Moreno-Cuevas JE. An overview of Neem (Azadirachta indica) and its potential impact on health. Journal of Functional Foods. 2020 Nov 1; 74:104171. https://doi.org/10.1016/j.jff.2020.104171.
- Ikram A, Khalid W, Saeed F, Arshad MS, Afzaal M, Arshad MU. Senna: As immunity boosting herb against COVID-19 and several other diseases. Journal of Herbal Medicine. 2023 Jan 10:100626. Doi: 10.1016/j.hermed.2023.100626.
- 11. Taylor RS, Edel F, Manandhar NP, Towers GH. Antimicrobial activities of southern Nepalese medicinal plants. Journal of ethnopharmacology. 1996 Feb 1;50(2):97-102. Doi: 10.1016/0378-8741(95)01335-0.
- Amoros M, Simõs CM, Girre L, Sauvager F, Cormier M. Synergistic effect of flavones and flavonols against herpes simplex virus type 1 in cell culture. Comparison with the antiviral activity of propolis. Journal of Natural Products. 1992 Dec;55(12):1732-40. Doi: 10.1021/np50090a003.
- 13. Anonymous. Commission on Dietary Supplement Labels issues final report. J Am Diet Assoc. 1998;98:270. Doi: 10.1016/S0002-8223(98)00064-9.
- 14. Borris RP. Natural products research: perspectives from a major pharmaceutical company. Journal of ethnopharmacology. 1996 Apr 1;51(1-3):29-38. Doi: 10.1016/0378-8741(95)01347-4.
- Islas JF, Acosta E, Zuca G, Delgado-Gallegos JL, Moreno-Treviño MG, Escalante B, Moreno-Cuevas JE. An overview of Neem (Azadirachta indica) and its potential impact on health. Journal of Functional Foods. 2020 Nov 1; 74:104171. https://doi.org/10.1016/j.jff.2020.104171.
- Mendoza L, Wilkens M, Urzua A. Antimicrobial study of the resinous exudates and of diterpenoids and flavonoids isolated from some Chilean Pseudognaphalium (Asteraceae). Journal of ethnopharmacology. 1997 Oct 1;58(2):85-8. Doi: 10.1016/s0378-8741(97)00084-6.
- 17. Ahmed AA, Mahmoud AA, Williams HJ, Scott AI, Reibenspies JH, Mabry TJ. New sesquiterpene αmethylene lactones from the Egyptian plant Jasonia candicans. Journal of natural products. 1993 Aug;56(8):1276-80. Doi: 10.1021/np50098a011.
- Behl T, Kumar C, Singh RK, Arora TK, Arora S. Traditional and Novel Herbal Drugs Emerging as Potent Novel Combinations for Managing Morbidities by Pharmacological and Mechanistic Studies. DOI:10.15415/jptrm.2018.61004.
- Amoros M, Simõs CM, Girre L, Sauvager F, Cormier M. Synergistic effect of flavones and flavonols against herpes simplex virus type 1 in cell culture. Comparison with the antiviral activity of propolis. Journal of Natural Products. 1992 Dec;55(12):1732-40. https://pharmajournals.stmjournals.in/index.php/RRJoPC/article/view/1117.
- Rana BK, Singh UP, Taneja V. Antifungal activity and kinetics of inhibition by essential oil isolated from leaves of Aegle marmelos. Journal of ethnopharmacology. 1997 Jun 1;57(1):29-34. Doi: 10.1016/s0378-8741(97)00044-5.
- Hotea I, Dragomirescu M, Berbecea A, Radulov I. Phytochemicals as Alternatives to Antibiotics in Animal Production. In Antibiotics and Probiotics in Animal Food-Impact and Regulation 2022 Sep 16. IntechOpen. Doi: 10.1186/s13567-018-0562-6.
- 22. Kubo A, Lunde CS, Kubo I. Antimicrobial activity of the olive oil flavor compounds. Journal of Agricultural and Food Chemistry. 1995 Jun;43(6):1629-33. https://doi.org/10.1021/jf00054a040.
- Hamburger M, Hostettmann K. 7. Bioactivity in plants: the link between phytochemistry and medicine. Phytochemistry. 1991 Jan 1;30(12):3864-74. Doi: 10.3389/fphar.2021.593856.
- 24. Ahmed AA, Mahmoud AA, Williams HJ, Scott AI, Reibenspies JH, Mabry TJ. New sesquiterpene αmethylene lactones from the Egyptian plant Jasonia candicans. Journal of natural products. 1993 Aug;56(8):1276-80. Doi: 10.1021/np50098a011.
- Pezzani R, Salehi B, Vitalini S, Iriti M, Zuñiga FA, Sharifi-Rad J, Martorell M, Martins N. Synergistic effects of plant derivatives and conventional chemotherapeutic agents: an update on the cancer perspective. Medicina. 2019 Apr 17;55(4):110. Doi: 10.3390/medicina55040110.

- Barre JT, Bowden BF, Coll JC, De Jesus J, Victoria E, Janairo GC, Ragasa CY. A bioactive triterpene from Lantana camara. Phytochemistry. 1997 May 1;45(2):321-4. Doi: 10.1016/s0031-9422(96)00805-9.
- 27. Taylor RS, Edel F, Manandhar NP, Towers GH. Antimicrobial activities of southern Nepalese medicinal plants. Journal of ethnopharmacology. 1996 Feb 1;50(2):97-102. Doi: 10.1016/0378-8741(95)01335-0.
- Amoros M, Sauvager F, Girre L, Cormier M. In vitro antiviral activity of propolis. Apidologie. 1992; 23:231–240. doi: 10.1155/2018/7092416.
- 29. Nagpal K, Garg M, Arora D, Dubey A, Grewal AS. An extensive review on phytochemistry and pharmacological activities of Indian medicinal plant *Celastrus paniculatus* Willd. Phytotherapy Research 2022 36(5),1930-51. DOI: 10.1002/ptr.7424
- Batista O, Duarte A, Nascimento J, Simões MF, de la Torre MC, Rodríguez B. Structure and antimicrobial activity of diterpenes from the roots of Plectranthus hereroensis. Journal of natural products. 1994 Jun;57(6):858-61. Doi: 10.1021/np50108a031.
- Ahmed AA, Mahmoud AA, Williams HJ, Scott AI, Reibenspies JH, Mabry TJ. New sesquiterpene αmethylene lactones from the Egyptian plant Jasonia candicans. Journal of natural products. 1993 Aug;56(8):1276-80. Doi: 10.1021/np50098a011.
- Tong Y, Song X, Zhang Y, Xu Y, Liu Q. Insight on structural modification, biological activity, structureactivity relationship of PPD-type ginsenoside derivatives. Fitoterapia. 2022 Apr 1; 158:105135. Doi: 10.1016/j.fitote.2022.105135
- Kriegeskorte N, Douglas PK. Cognitive computational neuroscience. Nature neuroscience. 2018 Sep;21(9):1148-60. Doi: 10.1038/s41593-018-0210-5.
- 34. Alam M, Bano N, Ahmad T, Sharangi AB, Upadhyay TK, Alraey Y, Alabdallah NM, Rauf MA, Saeed M. Synergistic Role of Plant Extracts and Essential Oils against Multidrug Resistance and Gram-Negative Bacterial Strains Producing Extended-Spectrum β-Lactamases. Antibiotics 2022, 11, 855. https://doi.org/10.3390/antibiotics11070855.
- Atta-ur-Rahman M I, Choudhary Diterpenoid and steroidal alkaloids. Nat Prod Rep. 1995; 12:361–379. Doi: 10.1039/np9951200361.
- 36. A. Alzohairy M. Therapeutics Role of *Azadirachta indica* (Neem) and Their Active Constituents in Diseases Prevention and Treatment. Evid Based Complement Alternat Med. 2016; 2016: 7382506. Doi: 10.1155/2016/7382506
- 37. Ooshima T, Minami T, Aono W, Izumitani A, Sobue S, Fujiwara T, Kawabata S, Hamada S. Oolong tea polyphenols inhibit experimental dental caries in SPF rats infected with mutatis streptococci. Caries research. 1993 Nov 20;27(2):124-9. Doi: 10.1159/000261529.
- Habtemariam S, Gray AI, Waterman PG. A new antibacterial sesquiterpene from Premna oligotricha. Journal of Natural products. 1993 Jan;56(1):140-3. Doi: 10.1021/np50091a022.
- 39. Hamburger M, Hostettmann K. 7. Bioactivity in plants: the link between phytochemistry and medicine. Phytochemistry. 1991 Jan 1;30(12):3864-74. Doi: 10.1016/j.phymed.2022.154196.
- Mishra LC, Singh BB, Dagenais S. Scientific basis for the therapeutic use of Withania somnifera (ashwagandha): a review. Alternative medicine review. 2000 Aug 1;5(4):334-46. https://pubmed.ncbi.nlm.nih.gov/10956379/.
- 41. Taylor RS, Edel F, Manandhar NP, Towers GH. Antimicrobial activities of southern Nepalese medicinal plants. Journal of ethnopharmacology. 1996 Feb 1;50(2):97-102. Doi: 10.1016/0378-8741(95)01335-0.
- 42. Perrett S, Whitfield PJ, Sanderson L, Bartlett A. The plant molluscicide Millettia thonningii (Leguminosae) as a topical antischistosomal agent. Journal of ethnopharmacology. 1995 Jun 23;47(1):49-54. Doi: 10.1016/0378-8741(95)01253-a.
- Verma S, Thakur D, Pandey CM, Kumar D. Recent Prospects of Carbonaceous Nanomaterials-Based Laccase Biosensor for Electrochemical Detection of Phenolic Compounds. Biosensors. 2023 Feb 22;13(3):305. Doi: 10.3390/bios13030305.
- 44. Scortichini M, Rossi MP. Preliminary in vitro evaluation of the antimicrobial activity of terpenes and terpenoids towards Erwinia amylovora (Burrill) Winslow et al. Journal of Applied Bacteriology. 1991 Aug;71(2):109-12. https://doi.org/10.3390/biom11040554.
- 45. Sher A. Antimicrobial activity of natural products from medicinal plants. Gomal Journal of medical sciences. 2009;7(1). https://doi.org/10.1016/j.sjbs.2017.02.004.
- 46. Pastorino G, Cornara L, Soares S, Rodrigues F, Oliveira M. Liquorice (Glycerrhiza glabra): A phtychemical and pharmacological review. Phytother Res. 2018 Dec; 32(12): 2323–2339. Doi: 10.1002/ptr.6178.

- 47. Kubo I, Muroi H, Himejima M. Combination effects of antifungal nagilactones against Candida albicans and two other fungi with phenylpropanoids. Journal of natural products. 1993 Feb;56(2):220-6. Doi: 10.1021/np50092a006.
- 48. Kubo I, Muroi H, Himejima M. Combination effects of antifungal nagilactones against Candida albicans and two other fungi with phenylpropanoids. Journal of natural products. 1993 Feb;56(2):220-6. Doi: 10.1021/np50092a006.
- Vishwakarma RA. Stereoselective synthesis of α-arteether from artemisinin. Journal of Natural Products. 1990 Jan;53(1):216-7. Doi: 10.1021/np50086a018.
- De Clercq E. Antiviral therapy for human immunodeficiency virus infections. ClinMicrobiol Rev. 1995; 8:200–239. Doi: 10.1128/cmr.8.2.200.
- 51. Jana S, Shekhawat GS. Critical review on medicinally potent plant species: Gloriosa superba. Fitoterapia. 2011 Apr 1;82(3):293-301. Doi: 10.1016/j.fitote.2010.11.008.
- Atta-ur-Rahman M I, Choudhary Diterpenoid and steroidal alkaloids. Nat Prod Rep. 1995; 12:361–379. Doi: 10.1039/np9951200361.
- Mendoza L, Wilkens M, Urzua A. Antimicrobial study of the resinous exudates and of diterpenoids and flavonoids isolated from some Chilean Pseudognaphalium (Asteraceae). Journal of ethnopharmacology. 1997 Oct 1;58(2):85-8. Doi: 10.1016/s0378-8741(97)00084-6.
- 54. Islas JF, Acosta E, Zuca G, Delgado-Gallegos JL, Moreno-Treviño MG, Escalante B, Moreno-Cuevas JE. An overview of Neem (Azadirachta indica) and its potential impact on health. Journal of Functional Foods. 2020 Nov 1; 74:104171. https://doi.org/10.1016/j.jff.2020.104171.
- 55. Habtemariam S, Gray AI, Waterman PG. A new antibacterial sesquiterpene from Premna oligotricha. Journal of Natural products. 1993 Jan;56(1):140-3. Doi: 10.1021/np50091a022.
- Vaou N, Stavropoulou E, Voidarou C, Tsigalou C, Bezirtzoglou E. Towards advances in medicinal plant antimicrobial activity: A review study on challenges and future perspectives. Microorganisms. 2021 Sep 27;9(10):2041. Doi: 10.3390/microorganisms9102041.
- 57. Ali M, Chaudhary N. Ficus hispida Linn.: A review of its pharmacognostic and ethnomedicinal properties. Pharmacognosy reviews. 2011 Jan;5(9):96. doi: 10.4103/0973-7847.79104.
- 58. Jones Jr SB, Luchsinger AE. Plant systematics. McGraw-Hill.; 1986 Dec. https://scirp.org/reference/referencespapers.aspx?referenceid=659808.
- 59. Estevez-Braun A, Estevez-Reyes R, Moujir LM, Ravelo AG, Gonzalez AG. Antibiotic activity and absolute configuration of 8S-heptadeca-2 (Z), 9 (Z)-diene-4, 6-diyne-1, 8-diol from Bupleurum salicifolium. Journal of natural products. 1994 Aug;57(8):1178-82. Doi: 10.1021/np50110a009.
- 60. Duke JA. CRC Handbook of Medicinal Herbs CRC Press. Inc., Boca Raton, FL, 1985.-677p. 1985.
- Rana BK, Singh UP, Taneja V. Antifungal activity and kinetics of inhibition by essential oil isolated from leaves of Aegle marmelos. Journal of ethnopharmacology. 1997 Jun 1;57(1):29-34. Doi: 10.1016/s0378-8741(97)00044-5.
- 62. Al-Dabbagh B, A. Elhaty I, Elhaw M, Murali C, Al Mansoori A, Awad B, Amin A. Antioxidant and anticancer activities of chamomile (Matricaria recutita L.). BMC Research Notes volume 12, Article number: 3 (2019). Doi: 10.1186/s13104-018-3960-y.
- Cichewicz RH, Thorpe PA. The antimicrobial properties of chile peppers (Capsicum species) and their uses in Mayan medicine. Journal of ethnopharmacology. 1996 Jun 1;52(2):61-70. Doi: 10.1016/0378-8741(96)01384-0
- 64. Prakash A, Vadivel V, Banu SF, Nithyanand P, Lalitha C, Brindha P. Evaluation of antioxidant and antimicrobial properties of solvent extracts of agro-food by-products (cashew nut shell, coconut shell and groundnut hull). Agriculture and Natural Resources. 2018 Oct 1;52(5):451-9. https://kukrdb.lib.ku.ac.th/journal/KJNS/search detail/result/410421.
- Kolenc Z, Langerholc T, Hostnik G, Ocvirk M, Štumpf S, Pintarič M, Košir IJ, Čerenak A, Garmut A, Bren U. Antimicrobial Properties of Different Hop (Humulus lupulus) Genotypes. Plants. 2022 Dec 26;12(1):120. Doi: 10.3390/plants12010120.
- Perrett S, Whitfield PJ, Sanderson L, Bartlett A. The plant molluscicide Millettia thonningii (Leguminosae) as a topical antischistosomal agent. Journal of ethnopharmacology. 1995 Jun 23;47(1):49-54. Doi: 10.1016/0378-8741(95)01253-a.
- Hotea I, Dragomirescu M, Berbecea A, Radulov I. Phytochemicals as Alternatives to Antibiotics in Animal Production. In Antibiotics and Probiotics in Animal Food-Impact and Regulation 2022 Sep 16. IntechOpen. Doi: 10.1186/s13567-018-0562-6.

- Deng Z, Sheng F, Yang SY, Liu Y, Zou L, Zhang LL. A comprehensive review on the medicinal usage of Podocarpus species: Phytochemistry and pharmacology. Journal of Ethnopharmacology. 2023 Mar 23:116401. Doi: 10.1016/j.jep.2023.116401.
- 23:116401. Doi: 10.1016/j.jep.2023.116401.
  69. Kubo I, Muroi H, Kubo A. Naturally occurring antiacne agents. Journal of natural products. 1994 Jan;57(1):9-17. Doi: 10.1021/np50103a002.