

# GENUS CARALLUMA: PHYTOCHEMICAL DIVERSITY, THERAPEUTIC POTENTIAL, AND EMERGING OPPORTUNITIES FOR DRUG DISCOVERY

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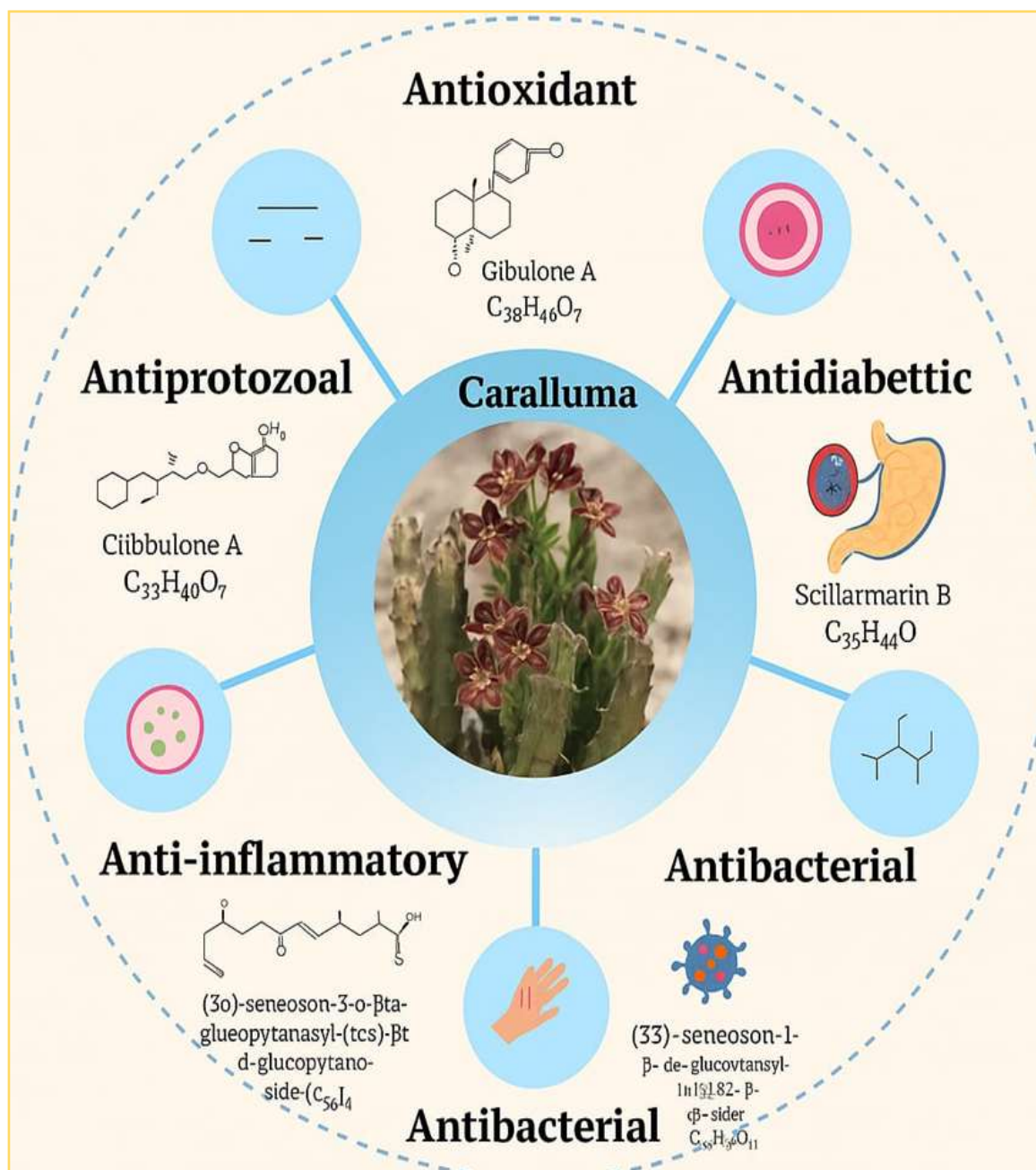
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## Abstract

The genus *Caralluma* has generated considerable attention due to its wide ethnomedicinal and phytochemical complexity. Still, lacking a comprehensive and update review that covers ethnomedicinal, phytochemical constituents and biological activities. This review aims to offer comprehensive and up-to-date information on genus *Caralluma* associating with its taxonomy, medicinal uses, phytochemical compound classes, anti-microbial properties and toxicology studies to lead future research. Web search engines such as, Science Direct, PubMed, Wiley Online Library, Google scholar and Springer Link were used to conduct an exhaustive examination of the literature. The genus *Caralluma* belonging from family (Apocynaceae) consists of succulent species that have long been used in traditional medicines across South Asia and the Arabian Peninsula. The results of this review presents a detailed synthesis of the available literature on the Distribution, medicinal uses, phytochemistry, antioxidant, antimicrobial and toxicity profiles of *Caralluma* species. The species of the *Caralluma* genus had been widely used to cure diabetes, inflammation, skin issues (freckles and pimples), hypertension, liver disorders, hepatitis B/C, cancer (oral, cervical, breast, lungs, colon, and hepatic cancer), rheumatism, blood disorders, leprosy, peptic ulcers, dysentery, gastric issues, constipation, kidney stones, cysts, cough, asthma, insomnia, urogenital infections, tuberculosis, weight management, malaria, appetite suppressing and microbial infections. It also covers the phytochemical compound classes, including pregnane glycosides, flavonoids, Saponins, gallic acid, rutins, alkaloids, and phenolic compound. These biological activities could be associated with the chemicals present in the species of *Caralluma*. This review highlights that Genus *Caralluma* had high medicinal values and also had antioxidant, toxicity, antimicrobial activities with its phytochemical constituents. It also demonstrated the urgent need for the future research on the drug discovery and ethnopharmacological studies of the genus *Caralluma*.

**Key words:** *Caralluma*, Phytochemical Diversity, Drug.

## GRAPHICAL ABSTRACT



## INTRODUCTION

There are 120 species in the xerophytic genus *Caralluma* R. Br., which belongs to the Apocynaceae family. The Arabic expression "qarh al-luhum," which indicates an abscess or a cut in the flesh, is where the word "*Caralluma*" originates. The arrangement of floral components is the only difference between the two taxa, thus *Caralluma* and *Boucerosia* are likewise regarded as synonyms (Stewart.,1972; abdel-sattar et al.,2025). The majority of *Caralluma* plant species are succulent perennial herbs with small, caducous leaves; some of these species have been found to be edible. (Naik and Krishnamurthy.,2012). A vast genus of the Asclepiadoideae subfamily (order: Ceropegieae), *Caralluma* R. Br. (syn. *Ceropegia* L.) has a variety of uses in traditional medicine (Karale and Karale .2017).

Many species of *Caralluma* contain large amounts of megastigmane glycosides, flavones, and PGs (Bader et al.,200; Braca et al.,2002; Abdel-sattar et al.,2007; abdel-sattar et al.,2025). Furthermore, some *Caralluma* extracts have shown antifungal, antibacterial, anti-eczemic, anticancer, antidiabetic, anti-inflammatory, and antimalarial properties (Waheed et al., 2011; Rauf et al., 2013; Anwar et al., 2022; Ansari et al., 2022; Amrati et al., 2021; abdel-sattar et al., 2025). In Pakistan, *Caralluma tuberculata* and *Caralluma edulis* were formerly classified as *Boucerosia edulis* and *Boucerosia tuberculata*, but they have since been recognized as *Caralluma* species (Rauf et al., 2013). It needs to be mentioned that *C. tuberculata* and *C. edulis* are currently recognized as *Caralluma* species, while they were

formerly classified as *B. tuberculata* and *B. edulis* in Pakistan. (Jayawardena et al., 2021; Zarei et al., 2020; Mounika et al., 2016)

*Caralluma* is a genus that is widely distributed over different regions. Asia and Africa (especially Pakistan, India, Afghanistan, Iran and Srilanka), These plants can be found in the Canary Islands, South Africa, Southeast Europe, and the Arabian Peninsula. This wide range illustrates the adaptability and resilience of *Caralluma* species in diverse environments (Gabali and Al-Gifri., 1990).

The genus *Caralluma* belongs to the class Magnoliopsida, subclass Asteridae, order Gentianales, and superorder Gentiananae (14). *Caralluma* was formerly a member of the Asclepiadaceae family, which is commonly referred to as milkweed. Asclepiadaceae should be categorized as a subfamily Asclepiadoideae within the Apocynaceae family (BENSUSAN.,1913) and tribe Ceropetegiaceae (Kamel et al., 2014), according to recent molecular and genetic research. With over 200 species, one of the main genera of the Asclepiadoideae subfamily is *Caralluma*. It is found in arid parts of tropical Asia, the southern Mediterranean, the near east and north, and central and eastern Africa (Gilbert.,1990; Albers and Meve.,2012).

Wight and Arnott divided the genus *Caralluma* into the new genera *Boucerosia* and *Hutchinia* in 1834 (Albers and Meve.,2012; Abdel-sattar et al.,2025). However, Brown regrouped all the genera into *Caralluma* in 1892, adding multiple identical succulents to the same genus, which increased dispute. Based on stem and flower morphology, Gilbert separated *Caralluma* into four subgenera once more in 1990: *Boucerosia*, *Caralluma*, *Desmidorchis*, and *Urmalcala*. Plowes (1995) later recognized 17 genera and roughly 70 species after further subdividing the genus according to different morphological principles (Albers and Meve.,2012; Abdel-sattar et al.,2025). *Caralluma* was classified into six subgenera by Meve and Liede in 2002: *Apteranthes*, *Boucerosia*, *Caralluma*, *Caudanthera*, *Desmidorchis*, and *Monolluma*. Taxonomists have made multiple attempts to categorize the various *Caralluma* taxa, but ambiguity persists. Apocynoideae, Asclepiadoideae, Periplocoideae, Rauvolfioideae, and Secamonoideae are the five subfamilies that make up the Apocynaceae family. One of the most important sources of C21 pregnane derivatives is Asclepiadoideae, formerly known as the Asclepiadaceae family (de Souza et al.,2024; Abdel-sattar et al.,2025).

The genus *Caralluma* is primarily abundant in pregnane glycosides, with several newly isolated including six novel pregnane glycosides (quadrangulosides A-F from *Caralluma quadrangular* (Ismail et al.,2023). *Caralluma quadrangula* produces flavone glycosides in addition to pregnane glycosides, which add to its range of bioactivity (Ismail et al.,2023). Glycosides, flavonoids, Alkaloids steroids, phenols, tannins and saponins, were found in several solvent extracts during the initial screening of *Caralluma stalagmifera* (veerabhadraiah et al.,2024). According to GC-MS analysis, the maximum concentration of flavonoids, alkaloids, phenolics and terpenoids was found in methanolic extracts of *Caralluma edulis* (Khan et al.,2022).

Hydroethanolic extracts of *Caralluma europaea* showed notable flavonoid and polyphenols concentrations as determined by HPLC, along with flavonoids like, myricetin, and hesperetin and quercetin (Amrati et al., 2020). It has been demonstrated that the stems of *Caralluma speciosa* contain a combination of fatty acids, hydrocarbons,  $\beta$ -sitosterol, megastigmane glycosides, flavones, and pregnane derivatives in both volatile and nonvolatile fractions. (Kiros et al.,2023).

High levels of phenolic and flavonoid compounds are found in *Caralluma tuberculata*; rutin and gallic acid have been found in methanol-based extracts, as well as, flavonoid glycosides, pregnane glycosides saponins, and triterpenes. (Baig et al., 2021; Mudrikah et al., 2021).

Through GC-MS analysis 30+ phytochemical compounds were reported in the *Caralluma indica*'s seed extract which also includes, esters and fatty acids. These compounds have antioxidant, antibacterial and cyto toxicity effects (Vidivu and Velavan, 2020). All over Mediterranean regions, Asia and Africa species of the genus *caralluma* are used in traditional medicines. Various species have been used as shortage foods, appetite suppressants, and treatments for a wide range of illnesses, such as gastrointestinal (dyspepsia, ulcers), metabolic (diabetes, obesity), febrile, inflammatory, and rheumatic conditions. (Adnan et al., 2014) In Yamen *Caralluma* is widely used for the treatment of gastrointestinal disorders, diabetes, obesity arthritis and inflammation (Abdel-sattar et al., 2025).

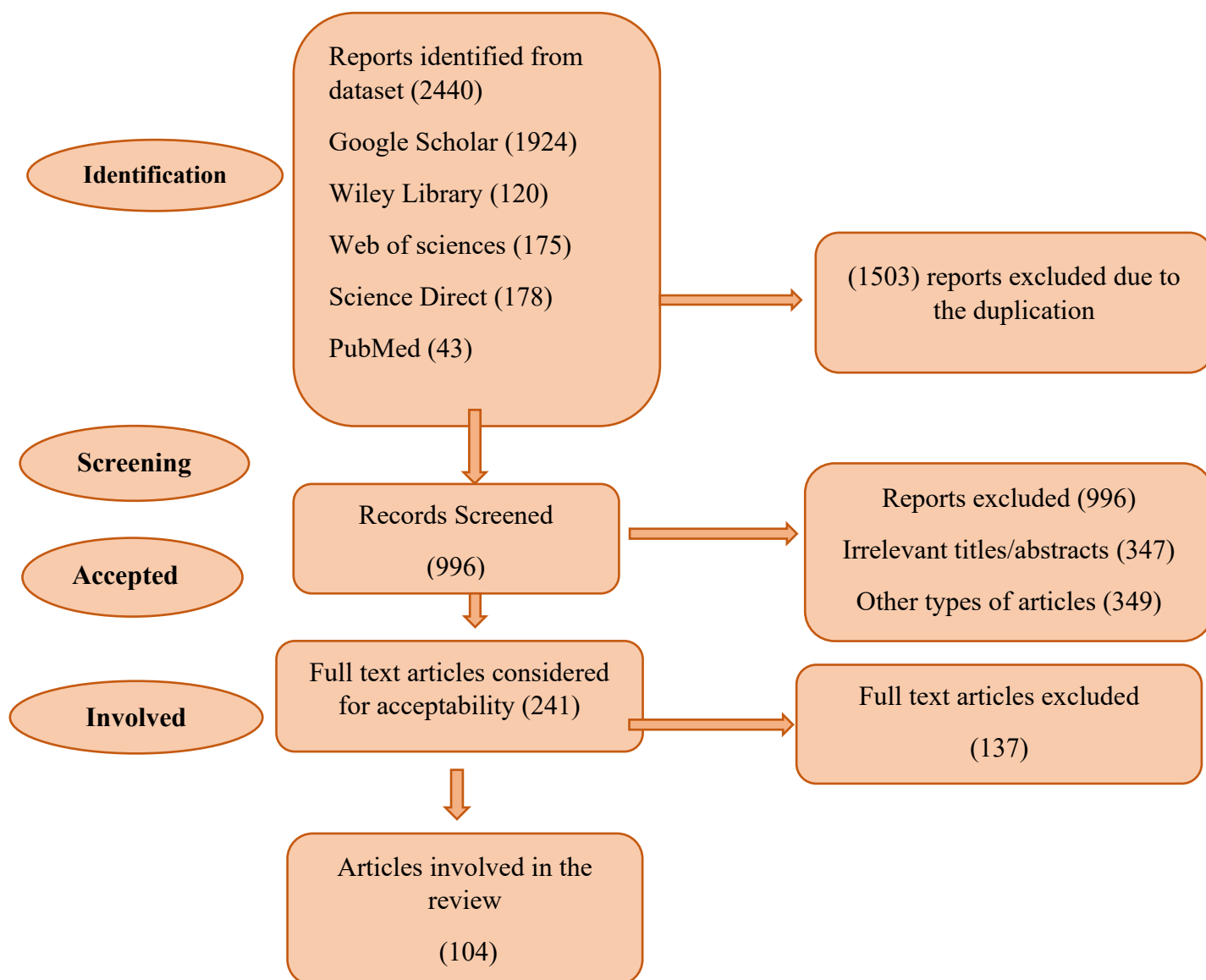
### 3.1. Review Methodology

An investigation was conducted on the available literature of genus *Caralluma* to evaluate the Phytochemical constituents, ethnomedicinal uses, antioxidant, anti-microbial properties and toxicity activities. A thorough search was conducted using Web of Science (<https://mjl.clarivate.com>), Science Direct, Wiley Online Library, Google Scholar, and Pub Med in accordance with the preferred reporting item PRISMA for Systematic Assessments and Meta-Analyses) recommendations. Important keyword searches include: genus *Caralluma*, species of the genus *Caralluma*, phytochemical contents, toxicological activities, antioxidant and antimicrobial properties, and phytochemicals of different *Caralluma* species. To ensure a thorough and accurate research, I employed a systematic filtering process to include and remove papers, as showed in Fig. 1. My first search screening yielded 2440 entries

from five sources: Google Scholar (n = 1924), PubMed (n = 43), Science direct (n = 178), Wiley online library (n=120) and Web Science (n = 175).

After removing 1503 duplicate items, I looked at 937 articles grounded on their abstracts and titles. During this initial screening step, 696 records were eliminated, including 347 publication categories deemed inappropriate (such as publications with unique elements) and 349 studies deemed unrelated to our emphasis on the genus *Caralluma* (such as studies of adjacent species). The remaining 241 full-text papers were carefully assessed using predetermined inclusion requirements that gave priority to research explicitly examining Genus *Caralluma* in terms of ethnomedicinal applications, phytochemical ingredients, toxicological activities, antioxidant, and anti-microbial capabilities. 137 studies were removed because they either (1) evaluated insufficient information for *Caralluma* species, (2) had insufficient data sources, or (3) were convention abstracts lacking full text availability or acceptable analytical information.

104 studies were included in this systematic review after a thorough assessment; and both the scientific and medicinal discriptions, acceptable Online resources such as Flora of Pakistan, The Plant List and Plants of the World Online were used to confirm the accurate species' name and author's name.



**Fig.1** Flow chart exhibited searched datasources for systematic literature review

## CHAPTER 4

### RESULTS

#### 4.1. Distribution and Conventional Uses of various *Caralluma* species

The collected data exhibited that *Caralluma* R. Br species have spread throughout the Asian continent, the Middle East, Africa, and portions of Europe. For instance, certain species, like *C. europea* (Guss.) N.E.Br., are indigenous to the Mediterranean and South Africa. *C. tuberculata* N.E.Br. are found in Saudi Arabia, Iran, the Republic of Nigeria, Egypt, Pakistan, India, and the United Arab Emirates. *C. retrospiciens* and *C. quadrangula* are primarily found in the East Africa and Arabian Peninsula. *C. edulis* (Edgew.) Benth. Ex Hook.f. and *C. umbelleta* Haw. are found in South Asia, Africa, India, and Pakistan. Differences by region demonstrate how well they adapt to semi-arid and drought-prone conditions. These herbs have long been used therapeutically to treat a variety of illnesses. Diabetes, stomach problems, liver problems, skin conditions, and blood purification are all treated with *C. tuberculata*. Kidney stones, respiratory conditions, obesity and cancer are all treated with *C. europea* (Guss.) N.E.Br.. A well-known remedy for snake bites, cancer, tuberculosis, and skin conditions is *C. quadrangula*, whereas *C. umbelleta* and *C. edulis* (Edgew.) Benth. Ex Hook.f. have been employed to treat intestinal diseases and stomach, inflammation, ulcers and diabetes. Certain species, such as *C. arabic* and *C. stalagmifera* are recommended for rheumatism, leprosy and wound healing, while, *C. retrospiciens*, *C. bhupendriana* and *C. indica* are especially mentioned due to their wound healing, anti-inflammatory, anti-cancer qualities.

#### 4.2. Preparation and Usage

According to the local traditions and the particular condition being treated, different *Caralluma* R. Br species are prepared and used in different ways. Most of the time, the plant's apical portions, roots or stems are used both as dried or fresh. *C. tuberculata* N.E. Br. is frequently taken fresh or in a decoction made from its stems to treat digestive issues, rheumatism and high sugar levels. Likewise, decoctions prepared from dried aerial portions of *C. edulis* (Edgew.) Benth. Ex Hook.f. are used to treat heart conditions, hyperglycemia and fever. The roots, stem and leaves of *C. europaea* (Guss.) N.E.Br. are frequently boiled to create a decoction that is used to treat respiratory conditions, digestive disorders and kidney stones.

*C. fimbriata* is crushed into powder or capsule form and used as an anti-diabetic and appetite suppressant medication in traditional Indian medicine. Usually, *C. adscendens* is made into decoctions or pests to treat diabetes and inflammation. Additionally, *C. umbelleta*'s stems and roots are brewed to treat metabolic issues. Additionally, *C. flava* is used in local medicine to treat fever, ulcers and infections using compositions prepared from the plant's sap or extracts. Although *C. russeliana* is used as aqueous or ethanolic stem extracts, particularly for the management for the enhancement of and kidney and liver functions, *C. sinaica* is historically prepared as ethanolic or methanolic extracts or decoctions to fight infections and digestive issues.

#### 4.3. Phytochemical compound classes of *Caralluma* R. Br species

The existence of a variety of pharmacologically significant chemicals was established by phytochemical screening of the chosen species. Alkaloids, terpenes, sterols,  $\alpha$  and  $\beta$  amyrin, lupeol, and pregnane glycosides were identified in *C. tuberculata*.  $\beta$ -pinene, verbenone, camphene, linalool, and other monoterpenes and sesquiterpenes are abundant in *C. europea* (Guss.) N.E.Br.. Pregnane glycosides, flavone glycosides, luteolin derivatives, triterpenes, and saponins are all present in *C. umbelleta*. Alkaloids, tannins, glycosides, carotenoids, and steroids are also characteristics of *C. edulis* (Edgew.) Benth. Ex Hook.f. Whereas *C. stalagmifera* is primarily composed of steroidal glycosides like lasianthoside A and B, *C. quadrangula* is rich in alkaloids, saponins, tannins, flavonoids, phenolics, and quercetin. Alkaloids, flavonoids, glycosides, terpenoids, and phenolphthaleins are found in *C. bhupendriana*, while, quinic acid, and hyperoside and apigenin-7-O-glucoside are found in *C. arabic*. Although *C. indica* is rich in Polyphenols, steroids, terpenoids, flavonoids, and saponins, *C. retrospiciens* demonstrated the existence of new pregnane glycosides.

#### 4.4. Phytochemical Analysis Units and Detection Techniques

Using standardized analytical procedures, the phytochemical profiles of *C. edulis*, *C. tuberculata*, *C. umbelleta* and *C. indica* were carried out to guarantee precise measurement of bioactive chemicals. The aluminum chloride colorimetric method was used to measure the total flavonoid content (TFC), which was then reported as milligrams of quercetin equivalents per gram of dry weight (mg QE/g DW). The total phenolic content (TPC) was measured using the Folin-Ciocalteu method, and the results were presented as micrograms or milligrams of gallic acid equivalents per gram of extract or dry weight ( $\mu\text{g GAE}/\text{mg}$  or  $\text{mg GAE}/\text{g DW}$ ). Alkaloids and tannins were estimated by both colorimetric and gravimetric methods, and their concentrations were expressed as  $\text{mg}/\text{g DW}$  or percentage weight by weight (% w/w). Saponins were analyzed in a similar way and reported either as  $\text{mg}/\text{g}$  or % w/w. To identify specific polyphenols such as rutin, quercetin, and gallic acid, advanced chromatographic techniques including HPLC and UPLC were applied. In addition, GC-MS analysis was carried out to determine the number of

chemical constituents and to identify fatty acids and volatile compounds. Together, these analytical approaches ensured accurate and reliable qualitative and quantitative evaluation of phytoconstituents in *Caralluma R. Br* species.

#### **4.5. Assessment of Toxicity and Safety**

Extracts from *C. tuberculata* N.E.Br. have been discovered to be non-toxic, with no negative effects on blood parameters, reproductive health and body weight. Because of its great safety margin and little cytotoxic effect, *C. europea* was likewise regarded as safe. Extracts from *C. umbelleta* showed non-toxic yet inhibiting properties, particularly when it came to pancreatic lipase. Extracts from *C. edulis* were also safe towards acute and long-term toxicity and non-antigenic. Extracts from *C. stalagmifera* proved to be efficient and tolerable against models of arthritis and inflammation, whereas *C. quadrangula* showed specific cytotoxicity against breast cancer cell lines. Butanol extracts had a minor impact on *C. arabic*, whereas methanol extracts demonstrated the strongest cytotoxicity, especially against breast cancer cells. According to reports, *C. retrospiciens* and *C. bhupendriana* are extremely protective and secure, particularly when it comes to medication delivery methods and DNA damage, correspondingly. Lastly, *C. indica* demonstrated substantial cytotoxic impacts on a cancer cell lines and variety of yeast models.

#### **4.6. Antioxidant activity**

Because of their high flavonoid and phenolic content, the majority of the species showed strong antioxidant activity. Extracts of *C. tuberculata* N.E.Br. made in hexane, methanol and chloroform shown exceptional antioxidant qualities. Flavonoids, phenolic substances and terpenoids in *C. europea* showed antioxidant activity. Although *C. edulis* extracts verified antioxidant effects from flavonoids and phenols. *C. umbelleta* additionally shown antioxidant activity because of triterpenoids and saponins. The primary antioxidant constituent in *C. quadrangula* was rutin, while glycosides and flavonoids shown action in *C. stalagmifera*. Polyphenolic acids, ascorbic acid and flavonoids in *C. arabic* had antioxidant properties. *C. bhupendriana*'s antioxidant chemicals showed a substantial prevention of UV-induced DNA damage. Flavonoids, tannins and alkaloids were present in *C. retrospiciens* extracts, which enhanced their antioxidant properties. Likewise, polyphenols and n-hexadecanoic acid in *C. indica* seeds offered defense against antioxidants.

#### **4.7. Antimicrobial and Cytotoxic Characteristics**

It was generally known that *Caralluma R. Br* species had antibacterial properties. Extracts from *C. tuberculata* N.E.Br. shown efficacy against *Salmonella typhi*, *E. coli*, *Shigella sonnei*, and *Bacillus subtilis*. *Candida albicans*, *Bacillus cereus*, *C. glabrata* and *Staphylococcus aureus*, were all significantly inhibited by *C. europea*. Extracts from *C. umbelleta* showed varying results against *P. aeruginosa*, *S. aureus*, and *E. coli*. *E. coli*, *P. vulgaris*, *P. aeruginosa*, and *K. pneumonia* were all shown to be inhibited by *C. edulis*. Significant antibacterial activity was demonstrated by *C. quadrangula* in both aqueous and organic extracts. Likewise, *C. stalagmifera* shown efficacy against *Proteus* species, *Pseudomonas* species, *Staphylococcus aureus*.

Ethanol extracts of *C. arabica* showed more potent antibacterial than antifungal properties. *C. retrospiciens* had both fungicidal and bacteriostatic properties, *C. bhupendriana* nanoparticles shown exceptional antibacterial ability. Lastly, *C. indica* extracts demonstrated broad-spectrum antibacterial efficacy against *Pseudomonas aeruginosa*, *Staphylococcus aureus*, *Salmonella typhi*, and oral pathogens.

S.No	Scientific Name	Distribution	Medicinal uses	Phytochemistry	Antitoxicity	Antioxidant	Antimicrobial	References
01.	C. Tuberculata	Pakistan (Balochistan, Nichara, Herboi, Gidar) India, United Arab Emirates, Saudi Arabia, Iran, Nigeria, South east of Egypt	Diabeties, Blood disorders, rheumatism, Hepatitis B and C, Leprosy, Peptic ulcers, Inflammation, Dysentery, digestive issues, Skin issues (pimples, freckles) purifier of the blood, Hypertention, Liver disorders, Gastric issues and Constipation	Terpenes ,Lupeol, $\alpha$ & $\beta$ Amyrin, $\alpha$ Amyrincinnamate, Pragnane Caratuberside, Sterols, Glucosides, Flavonoids, Luteolin-4-O-neohesperidosides, beta-cyaninan and alkaloids	Medicinally considered safe as in a research study ethyle acetate extractions was non toxic and had no negative affects on sperm health, body weight and on blood health	Methanol extracts, Chloroform friction and hexane friction demonstrated amazing anti- oxidant activity .	Bacillus subtilis, Bacillus streptococcus Eschrichiacoli, salmonella typhi and shigella sonnei, extracts of C.tuberculata were found effective against these microbes	(Duranni et al., 2009; Tareen et al., 2010; Shah et al., 2013 Rizwani, 1991; Venkatesh et al., 2003; Ahmed and Shaikh, 1989; Bensuzan, 2009; Collennate, 1991; Najam-us-saqib et al., 2013)
02.	C. Europea	Jordan, Morocco, Egypt, Libiya, Tunisia, Spain and Algeria	Coughs, Bronchospasm disorders, sleep disorders, Respiratory diseases, digestive issues, Urinary tract infections, Cancer, analgesic issues anti-parastic, anti-obesity, anti-atherosclerotic, kidney stones, cysts	Monoterpene hydrocarbons, sesquiterpenes hydrocarbons $\gamma$ -Terpinen/Linalool $\beta$ -Pinene, camphene, Thujopsene, $\alpha$ -Phellandrene $\beta$ -Eudesmol, $\alpha$ -Pinene, $\beta$ -Bisabolene, Tricyclene, verbonene and Linalool	Had minimum acute and cytotoxic affects and suggested a high safety margin	Terpenoids, flavonoids, and phenolic compounds	Candida albicans, Candida glabrata, Candida krusei, Candida parapsilosis, Micrococcus luteus, Bacillus cereus, Bacillus subtilis, and Staphylococcus aureus exhibited anti-microbial activity against these microbes	(Lahsissene et al., 2009; Dallahi et al., 2016; Mechchate et al., 2020; Meve and Heneidak., 2005; Zito et al., 2013; Ennacerie et al., 2017; Dra et al., 2018; Formisano et al., 2009; Zito et al., 2010; Ouassou et al., 2018; Issiki et al., 2017; Dra et al., 2018; Bourhia et al., 2020).
03.	C. umbellata	Pakistan, Africa, Spain, Middle East, Saudi Arabia, India and Middle East	Anti-inflammatory, Stomach disorders, Analgesic, anti-nociceptive, Anti ulcerogenic and intestinal disorders	pregnane glycosides, carumbelloside I to V, flavone glycoside luteolin-4'-O-neohesperidoside. megastigmane glycosides, bitter	Medicinally considered safe and the extract suggested inhibitroy affects against pancreatic lipase.	Sapanions, triterpenoids, falvone glycoside flavnoids, steroidal glycoside	Extracts from the root and stem of C.umbellata showed both effective and ineffective activity against P.aeruginosa, B.cereus and B.	(Abdel-sattar et al., 2007; Ahmed et al., 1993; Babu et al., 2014; Braca et al., 2002; Bader et al., 2003; Bellamakondi et al., 2014; Jyoti et al., 2015)

				principles saponins and triterpenes			subtillis, S.aureus P.vulgaris and E.Coli	
O4.	C.edulis	South Asia	cancer, tuberculosis, inflammation ,Skin Disorders, Diebeties Mellitus Malaria,Weight Management, obesity and appetite suppressing.	Alkaloids, Tannins, Glycosides, Carotenoids, Flavonoids Sterpenoids, and steroids.	Evaluated as safe and non- antigenic against acute and chornic toxic affects.	Phenols and flavonoids possessed antioxidant activity .	The plant extracts showed inhibitory effects preventing S. aureus, K., P. vulgaris, P. aeruginosa, E. coli and pneumoniaea	(Waheed et al., 2011; Adnan et al., 2014; Dutt et al., 2012; Mradu et al., 2012; Sofi et al., 2016; Elisha et al., 2017 and Ren et al., 2005; Mossa et al., 1995; Bukhari et al., 2007; Bukhari et al., 2016)
05	C.Quadrang ula	Arabian Peninnsula,India ,Srilanka,Iran,Afric a ,South ,Africa Canary Islands and Europe	High blood sugar, Sting from snakes and scorpions, tuberculosis, fever, skin disorders, cancer and inflammation	phenolics, flavonoids, tannins,quercitin alkaloids, and saponins	C.Quadrangula demonstrated favorable and active cytotoxic effects against MFC7 brest cancer cell lines.	Main antioxidant was Rutin(present in glycosidic form	Aqueous extracts and organic frictions of the C. quadrangula had been demonstrated as microbial	(Meve and Liede,2004; Leo et al., 2005; Abdallah et al., 2013; wua and Ng, 2007; Miliauskas et al., 2004; Perk et al., 2014; Enogieru et al., 2018; Farouk et al., 2016)
06	C.stalagmife ra	Southern India,(Andhara Pardesh,Tamil Nadu and Karnataka	Diabetes, leprosy, and rheumatism	Steroidal glycosides, carumbelloside III, lasianthoside A and B	Medicinally considered as safe as the Butanol and aqueous extracts showed significant effects as in a experiment upon koalin- associated arthritis within rats and carrageenin- mediated rat paw and also the extracts (in capsulated and and powdered form) showed effective results	glycosides, hydrocarbons, saponins and flavonoid exhibited antioxidant activity	crude extracts from the stem of the plant actively crude extracts from the stem of the plant actively proteus spp,Pseudomonas spp and staphylococcus aureus	(Reddy et al., 1996; (Parihar, 2016; Dutt et al., 2012; Aslam et al., 2019; Chandran et al., 2014; Packialakshmi and Naziya, 2014; Kunert et al., 2006)

					in reducing weight			
07	C.Arabica	Saudi Arabia ,Oman ,Yemen ,United Arab ,Emerites and Africa	Skin diseases and conditions (itchy skin,wounds cuts )	Epigenine-7-0-glucoside,Quinic acid,Naringenin, Hyperoside (quercitin-3-0-galactoside)	The methanol friction showed strongest cytotoxic effects while Butanol extract showed least friction against MCF-7 brest cancer cell line	polyphenolics compounds like phenolic acids flavonoids and ascorbic acid	Ethanollic extracts of C.Arabica showed stronger antibacterial effects while least antifungal effects evidenced by major inhabitant zones against funga strains compared to bacterial strains	(Abdel-sattar et al., 2022; williams et al., 2004; soobrattee et al., 2005; Khasawneh et al., 2014; Al-Mutaani et al., 2025)
08	C.bhupendriana Sarkaria	South India	Anti-inflammatory-Anti-cancer ,Anti-tumor,Anti-ulcer,Gastric-mucosa,	phlobatannins, phytosteroids, terpenoids, cardiac glycosides, alkaloids, quinones, phenols, tannins, carbohydrates,sap onins flavonoids	Methanol extracts of C.bhupenderiana remarkably inhibited UV-irradiated DNA damage therefore considered safe	The extracts from C.bhupenderiana sarkaria exhibited antioxidant activity and evaluate inhibitory effects against DNA damage .phenols and flavonoids	Nanoparticles prepared from C.sarkaria showed great performance against Anti-microbial activity	(Deepak et al., 1997; Ramesh et al., 1999; Zakaria et al., 2001; Al-Harbi et al., 1994; Zakaria et al., 2002; Umahmaheswari and Chatarjee., 2008; Nunes et al., 2012; Pachipala et al., 2022; Murugapandi et al., 2023)
09.	C.retrospicie ns	Eritrea, Ethiopia, Kenya, Saudi Arabia, Uganda and Yemen	Wounds healing	A novel-polyoxy pregnane glycosides (retrospinoids was successfully isolated from the aerial parts of C.retrospicie ns	The nanoparticles-based drug delivery mechanisms prepared from the plant reduced all drug toxicity	Alkaloids, flavonoids, tannins, steroids, and saponins	3,4-Altrosan showed fungicidal andbacteriostatic activites and the Benzonic acid identified from C.retrospicie ns demonstrated anti-bacterial effects.	Jadhav et al., 2014; Arokiyaraj et al., 2018; Eun-Soo et al., 2001; Bruyns et al., 2010; Patra et al., 2018; Wided et al., 2021; Alqahtani et al., 2022; Elsebai and Mohamed, 2015; Makeen et al., 2020)
10.	C.indica	Andhra pardesh ,karnataka and Tamil nadu	Cancer(oral curvical,brest lungs ,colon and hepatic cancers)	saponins, flavonoids, steroids, terpenoids,	C.indica showed remarkale cytotoxic activity on Oral (OECM-1)	Polyphenols and n-hexadecanoic acid. evaluated from C.indica seeds	Ethanollic extracts of C.indica plant demonstrated significant activities against	(Gnanashree et al.,2018; Sembiring et al., 2018; Periyamayagam et al., 2013; Koch et al., 2005; Chandra et al., 2023; Mostafa et al., 2018;

				polyphenols and tannis	cancerous cells lines and ethanolic seed extracts demonstrated significant cytotoxic action on yeast cells of <i>S. cerevisiae</i> .	exhibited anti-oxidant activity	<i>Pseudomonas aeruginosa</i> was significantly inhibited by ethanolic extracts of the <i>C. indica</i> plant. When it comes to treating oral infectious agents, <i>Staphylococcus aureus</i> , <i>Bacillus cereus</i> , <i>Escherichia coli</i> , <i>Salmonella typhi</i> , and the seeds show organic alternatives to synthetic microbials.	sanchez et al., 2016; Ramalingam et al., 2024; Kunert et al., 2006)
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Species	Area/Traditional uses	Preparation and mode of uses	References
C.tuberculata	reported from Pakistan's Khyber Pakhtunkhwa and Balochistan; frequently used to treat rheumatic pain, diabetes, and stomach distress	Fresh stems that are either eaten raw or made into a decoction	Baig et al., (2021);Haider et al., (2022)
c.edulis	used in Pakistan's Cholistan desert to treat heart conditions, high blood sugar and fever	Dry aerial components are used as decoction	Akram et al., (2023);Waris et al.,(2013);Ahmed et al., (2009)
c.europaea	Native to North Africa and some parts of Europe ,used to treat heart ailments ,kidney stones, infection, respiratory and digestive disorders.	Different plant parts (leaves, stems, roots used as decoction	Ouassau et al., (2021);Amrati et al., (2020)
C.fimbriata	used in Indian traditional medicine as an appetite suppressant, to control weight, and to treat diabetes.	Either as extracts in powder form or as capsules	Kuriyan et al., (2007)
C.adscendens	it was discovered in South India and is used to treat inflammatory diseases and diabetes.	Generally made as decoctions or pastes	Bhuvanewari and Manivannan.(2014)
C.umbellata	Reportedly from india,used to treat stomach disorders,diabetes .	Root and stem extracts or employed as Decoction	Bellamakondi et al., (2014); Babu et al., (2014)
C.flava NE Br. Int. J.	Detected from Arabian regions; utilized traditionally as anti-pyretic, anti-inflammatory, anti-ulcer, antidiabetic and anti-parasitic properties .	In local medicine ,plant extracts and sap are prepared	Raees .(2018)
C.sinaica	Native to Sinai regions and Egypt; utilized in traditional medicines for the treatment of infections, diabetes and digestive problems.	Traditionally prepared from the ethanolic/methanolic extracts or decoctions	Al-Massarani et al., (2012); Habibuddin et al., (2008)
C.russeliana	Derived from the arabian peninsula; stem extract has long been used to treat diabetes and related liver, lipid profile and kidney disorders.	Ethanolic/aqueous stem extracts are either ingested or used in experiments .	Zari and Al-Thebaiti.(2018)

Sr#	Phytochemical compound class	Species	Detection Method	Parts of the plant and Extracting medium	Reported data	References
1	Alkaloids	C.edulis	Colorimetric method	Entire plant ,Extracts of methanol	Alkaloids reported	Khan et al., (2022)
		C.indica	Qualitative and MS-GC tests	Stem extract	Alkaloids reported	Vadivu and Velavan.(2020)
		C.umbellata	Phytochemical qualitative method	Entire plant	Alkaloids reported	Bellamakondi et al., (2017)
		C.tuberculata	Colorimetric /Gravimetric method	Entire plant ,Fractions of methanol	Alkaloids reported	Baig et al., (2021)

2	<b>Tannins</b>	C.edulis	Colorimetric /Gravimetric assay	Leaves /extracts of methanol	Tannins reported positive	Khan et al., (2022)
		C.indica	Screening qualitative	Extract of stem	Tannins detected	Vadivu and Velavan. (2020)
		C.umbellata	Colorimetric test	Entire plant	Tannins present	Bellamakondi et al., (2017)
		C.tuberculata	Colorimetric /Gravimetric method	Hydroalcoholic extracts of the leaves	Tannins detected	Baig et al., (2021)
3	<b>Saponins</b>	C.edulis	Colorimetric technique	Methanolic extracts of the entire plant	Saponins present	Khan et al., (2022)
		C.indica	GC-MS assay	Extracts of the stem	Saponins present	Vadivu and Velavan.(2020)
		C.umbellata	Phytochemical techniques	Methanolic extracts of the entire plant	Saponins present	Bellamakondi et al., (2017)
		C.tuberculata	Gravimetric /colorimetric assay	Hydroalcoholic fractions of the entire plant		Baig et al., (2021)
4	<b>Total phenolic concentration (TPC)</b>	C.edulis	Folin-Ciocalteu technique	Apical parts,Acetone,ethanol,methanol.	Highest in methanol fractions	Khan et al., (2022)
		C.indica	Folin-Ciocalteu technique	Extracts of the stem	Phenolic compound present	Vadivu and Velavan.(2020)
		C.umbellata	Folin-Ciocalteu technique	Methanolic extracts of the entire plant	Great phenolic hepatoprotective levels	Bellamakondi et al., (2017)
		C.tuberculata	Folin-Ciocalteu technique	Water-acetone extracts extracts of entire plant	Rich phenolic content	Baig et al., (2021)
5	<b>Total flavonoids concentration</b>	C.edulis	Aluminium chloride technique	Methanolic extracts of the apical parts	Richest flavonoids content in methanol	Baig et al., (2021); Khan et al., (2022)
		C.indica	Aluminium chloride technique	Methanolic extract of the stem	Flavonoids detected (MS-GC) verified	Vadivu and Velavan. (2020)
		C.umbellata	Aluminium chloride technique	Methanolic extracts of the entire plant	Actively high flavonoid content detected	Bellamakondi et al., (2017)
		C.tuberculata	Aluminium chloride technique	Acetone/methanol extracts of the entire plant	Good levels of flavonoids detected	Baig et al., (2021); Khan et al., (2022)
6	<b>Specific polyphenols(Gallic acid, Quercetin, Rutin)</b>	C.edulis	UPLC/HPLC	Extracts of the methanol	Quercetin,gallic acid	Khan et al., (2022)
		C.indica	HPLC & MS-GC assay	Extracts of the stem	Derivatives of the fatty acid and gallic acid present	Vadivu and Velavan. (2020)
		C.umbellata	HPLC assay	Methanol extract	Gallic acid ,rutin detected	Bellamakondi et al., (2017)
		C.tuberculata	UPLC/HPLC	Extracts of the methanol	Gallic acid and rutin identified	Baig et al., (2021)

7	<b>Fatty acids /Volatile</b>	C.edulis	MS-GC technique	Extracts of the methanol	Fatty acid esters ,hexadecanoic detected	Iftikhar et al., (2022)
		C.indica	MS-GC technique	Extracts of the stem	n-heptadecanoic acid ,hexadecanoic acid detected	Vadivu and Velavan. (2020)
		C.umbellata	MS-GC technique	Methanolic extract	Sterols,phytol,hexadecanoic acid detected	Bellamakondi et al., (2017)
		C.tuberculata	MS-GC technique	Fractions of hexane/ethyl acetate	Hexadecanoic acid ,phytol identified	Khan et al., (2022)
8	<b>MS-GC over all compounds</b>	C.edulis	MS-GC technique	Extracts of methanol	32 compounds detected	Iftikhar et al., (2022)
		C.indica	MS-GC technique	Extracts of the stem	20 compounds detected	Vadivu and Velavan. (2020)
		C.umbellata	MS-GC technique	Extracts of methanol	30+ compounds identified	Bellamakondi et al., (2017)
		C.tuberculata	MS-GC technique	Extracts of methanol	30+ compounds identified	Khan et al., (2022)
9	<b>Various secondary metabolites(Terpenoids,phytosterols ,steroids)</b>	C.edulis	Screening phytochemicals tests	Methanol fractions of the entire plant	Terpenoids ,steroids,phytosterols identified	Khan et al., (2022)
		C.indica	Screening phytochemicals tests	Extracts of the stem	Phytosterols,fatty acid esters ,terpenoids identified	Vadivu and Velavan. (2020)
		C.umbellata	Screening phytochemicals tests	Methanol extracts of the entire plant	Glycosides,terpenoids ,steroids identified	Bellamakondi et al., (2017)
		C.tuberculata	Screening phytochemicals tests	Fractional extracts of the whole plant	Glycosides ,steroids ,terpenoids detected	Baig et al., (2021)

## CHAPTER 5

### DISCUSSION

***Caralluma tuberculata* N.E.Br.** is widely distributed in Pakistan (Balochistan, Nichara, Herboi, Gidar) India, United Arab Emirates Saudi Arabia, Iran, Nigeria, South east of Egypt (Duranni et al., 2009). The plant is used for several diseases including Diabetes, Blood disorders, rheumatism, Leprosy, (Venkatesh et al., 2003) Peptic ulcers, Inflammation (Ahmed and Shaikh, 1989) dysentery, hepatic disorder, skin issue (frackles and pimple), blood purification, gastric issue, hepatitis B and C, hypertension, stomach pain and constipation (Lawrence and Choudhary, 2004; Tareen et al., 2010; Shah et al., 2013). In contrast (Noreen, 2017) reported the use of *C. tuberculata* N.E.Br. for inflammation diabetes and obesity. The phytochemicals derived from *C. tuberculata* N.E.Br. including terpenes, Lupeol,  $\alpha$  &  $\beta$  Amyrin,  $\alpha$  Amyrincinnamate, Pragnane Caratuberside, Sterols, Glucosides, Flavonoids, Luteolin-4-O-neohesperidosides, beta-cyanin and alkaloids (Bensuzan, 2009). On the other hand, (Al-Rubaye et al., 2017) reported the phytochemicals of *C. tuberculata* such as Flavonoids terpinoids and other metabolites. The extracts of *C. tuberculata* N.E.Br. possessed anti-oxidant activity (Rauf, 2013). Methanol and chloroform fraction of *C. tuberculata* N.E.Br. showed great antioxidant activities. (Rizwani, 1991) examined that methanol fraction had highest antioxidant activity and it also possessed acute and chronic toxic activity. While in contrast to these studies (Ali et al., 2024) reported the phenolic chemicals of the plant that helped in strengthening the body's defence system against oxidative distractions. (Najam us saqib et al., 2013) reported in their research study that the extraction of ethyl acetate was nontoxic and had no negative effects on blood health, sperm health and body weight. And the extracts were also seemed to be active against all the Gram-positive and Gram-negative bacterial strains including *Staphylococcus Bacillus aureus*, *Bacillus subtilis* *Streptococcus viridens* and The Genera *Salmonella typhi*, *Shigella sonnei*, and *Escherichia coli* are examples of gram-negative bacteria (Collenette, 1999). Differing from this study (Baig et al., 2021) demonstrated that bioactive chemicals of *C. tuberculata* N.E.Br. had the capability to destroy the microbial cell wall.

***Caralluma europea* (Guss.) N.E.Br.** is distributed mainly in Jordan, Morocco, Egypt, Libya, Tunisia, Spain and Algeria (Meve and Heneidak, 2005; Zito et al., 2013). Adnan et al., (2014); Issiki et al., (2017) had similar results. The plant is used for various medicinal purposes coughs, asthma, insomnia, Cancer, Respiratory infections, digestive disorders, cysts, urogenital infections, Kidney stones, analgesic issue, anti-parasitic, anti-obesity, anti-atherosclerotic (Lahsissene et al., 2009; Dallahi et al., 2016; Mechchate et al., 2020). *C. europea* (Guss.) N.E.Br. contain various phytochemicals such as monoterpene hydrocarbons, sesquiterpenes hydrocarbons  $\beta$ -Eudesmol,  $\alpha$ -Pinene,  $\beta$ -Bisabolene, Tricyclene, verbonene, Linalool  $\gamma$ -Terpinen/Linalool  $\beta$ -Pinene, camphene, Thujopsene, and  $\alpha$ -Phellandrene (Ennacerie et al., 2017; Dra et al., 2018). While (Douhou et al., 2003) reported phytochemicals for example flavonoid, saponins alkaloids, terpenes, quinones steroids and tannins It had minimum acute and cytotoxic effects and suggested a high safety margin (Formisano et al., 2009; Zito et al., 2010). Ait Dara. (2019) reported that *C. europea* was safe in Up to 2000 kg/mg body weight of acute poisoning. Terpenoids, flavonoids, and phenolic compounds possessed antioxidant activity (Ouassou et al., 2018; Issiki et al., 2017). As compared to this study (Burits and Bucar, 2000) described the methanolic extracts of *C. europea* (Guss.) N.E.Br. evaluated antioxidant activity. Against numerous microbes like *Candida albicans*, *Candida glabrata*, *Candida krusei*, *Micrococcus luteus*, *Bacillus cereus*, *Bacillus subtilis*, and *Staphylococcus Aureus* *C. europea* exhibited anti-microbial activity (Dra et al., 2018; Bourhia et al., 2020). In contrast to this study (Amrati et al., 2021) reported that *C. europea* (Guss.) N.E.Br. fractions were highly active on gram positive bacteria and yeast.

***Caralluma umbellata* Haw.** is geographically distributed in Pakistan, Spain, Africa, Middle East, India Saudi Arabia Middle East and India (Jyoti et al., 2015). While (Shanmugam et al., 2013) reported the distribution of *C. umbellata* Haw. in India. This plant is used for Anti-inflammatory, stomach disorders, analgesic, anti-nociceptive, anti-ulcerogenic and intestinal disorders (Abdel-sattar et al., 2007). Differing from this study (Shanmugam et al., 2013) reported the uses of *C. umbellata* Haw. as vegetables. It contains various chemicals including pregnane glycosides, Flavone glycoside, luteolin-4'-O-neohesperidoside, carumbelloside I to V. megastigmane, glycosides, bitter principles, saponins and triterpenes (Ahmed et al., 1993) and (Shanmugam et al., 2013) had closely similar to this findings. Medicinally considered safe and the extract showed inhibitory effects against pancreatic lipase (Babu et al., 2014) On the other hand (Bellamakondi et al., 2014) revealed that extracts and fraction of the cytotoxicity test demonstrated safe concentration even at 500  $\mu$ g. Saponins, triterpenoids, flavone glycoside, flavonoids, steroidal glycosides exhibited antioxidant activity (Braca et al., 2002; Bader et al., 2003). Extracts from the root and stem of showed both effective and ineffective activity against *P. aeruginosa*, *B. cereus* and *B. subtilis*, *S. aureus*, *P. vulgaris* and *E. Coli*

(Bellamakondi et al., 2014). Similarly (Bennete, 2003) observed that Chloroform and acetone extracts showed antimicrobial activity against these bacteria, including *E. coli*, *B. subtilis*, *B. cereus*, and *S. aureus*.

**C. edulis (Edgew.) Benth. Ex Hook.f.** is distributed in South Asia (Malladi et al., 2018). The medicinal uses of the *C. edulis* are cancer, tuberculosis, inflammation, skin disorders, diabetes mellitus, malaria, weight management, obesity and appetite suppressings (Dutt et al., 2012; Waheed et al., 2011). Adnan et al., (2014) documented the uses of *Caralluma edulis* for Alzheimer's disease, leprosy, hypertension, rheumatism and obesity. The plant extracts showed inhibitory effects opposing *S. aureus*, *K. P. vulgaris*, *P. aeruginosa*, and *E. coli*, pneumoniae (Bukhari et al., 2016). Comparable to this study (Shailemo et al., 2016) demonstrated that various extracts of *C. edulis* showed best antimicrobial activity. Phenols and flavonoids showed antioxidant effects (Mossa et al., 1995; Bukhari et al., 2007). Ansari et al., (2022) highlighted that the extracts of the plant exhibited high antioxidant activity. Evaluated as safe and non-antigenic against acute and chronic toxic effects (Elisha et al., 2017; Ren et al., 2005). Similarly, Sanmugapriya and Venkataraman, (2006) revealed that plant was apparently safe in acute toxicity test. Chemicals present in the plant are alkaloids, tannins, glycosides, carotenoids, flavonoids, terpenoids, and steroids (Sofi et al., 2016; Mradu et al., 2012) whereas Ansari et al., (2022) recorded the phytochemicals in *Caralluma edulis* such as flavonoids, sterols and pregnane glycosides terpenoids.

**C. quadrangula (Forssk.) N.E.Br.** is distributed in Arabian Peninsula (Abdel-Sattar et al., 2017), India, Sri Lanka, Iran, Africa, South Africa, Canary Islands and Europe (Meve and Liede, 2004). In contrast to this study Guashash. (2006) documented that *C. quadrangula* (Forssk.) N.E.Br. is found in Saudi Arabia. Medicinal usage of the plant is diabetes, cancer, snake and scorpion bites, tuberculosis, fever, skin (Leo et al., 2005; Abdallah et al., 2013) while Guashash highlighted the uses of *C. edulis* (Edgew.) Benth. Ex Hook.f. for vitiligo, melasma, freckles and diabetes. *C. quadrangula* (Forssk.) N.E.Br. showed favorable and active cytotoxic effects against MFC7 breast cancer cell lines (Abdallah et al., 2013) while (Dutt et al., 2013) reported that this plant exhibited nontoxic effects against mammalian cells. Aqueous extracts and organic fractions of the *C. quadrangula* (Forssk.) N.E.Br. had been demonstrated as microbial characteristics (Farouk et al., 2016). Although (Al Sheikh et al., 2021) revealed that extracts of *C. quadrangula* like (Cq3, Cq2, Cq1) exhibited antibacterial activity compared to MDRAB AB5057 and MRSA USA300. Main antioxidant was Rutin present in glycosidic form (Perk et al., 2014; Enogieru et al., 2018). Chemicals of the *C. quadrangula* (Forssk.) N.E.Br. are phenolics, flavonoids, tannins, quercetin alkaloids, and saponins (Wua & Ng, 2007; Miliuskas et al., 2004). Compared to this study Abdallah et al., (2013); Al Sheikh et al., (2021), Ben Said et al., (2023) reported that various pregnane glycosides like flavonoid luteolin 4'-O-b-D-neo hesperidoside, ruscogenin, acylated boucerosides were evaluated from *C. quadrangula*.

**C. stalagmifera C.E.C.Fisch.** Parihar, (2016) documented the distribution of *C. stalagmifera* C.E.C.Fisch. in Southern India, (Andhra Pradesh, Tamil Nadu and Karnataka). Similarly, (Ramachandra et al., 2014) had similar findings (Kunert et al., 2006) reported the chemicals present in the plant are Carumbelloside III, lasianthosides A and B, and steroidal glycosides. While (Kunert et al., 2009) observed that *C. stalagmifera* is rich in alkaloids, flavonoids and glycosides. This plant is medicinally used for diabetes, leprosy, and rheumatism (Reddy et al., 1996). While Sreenivasacharyulu and Yogaratnakaram, (1939) revealed that mixing *C. stalagmifera* with black pepper is best for the treatment of migraine. Medicinally considered as safe as the butanol and aqueous extracts showed significant effects as in an experiment on rat paw caused by carrageenan and kaolin induced arthritis in rats and also the extracts (in capsulated and powder form) showed effective results in reducing weight (Dutt et al., 2012). In anti-microbial activity crude extracts from the stem of the plant actively inhibited the growth of *Escherichia coli*, *Bacillus subtilis* spp, *Pseudomonas* spp and *Staphylococcus aureus* (Packialakshmi and Naziya, 2014). Glycosides, hydrocarbons, and flavonoid saponins showed antioxidant activity (Aslam et al., 2019; Chandran et al., 2014). In contrast to these studies (Madhuri et al., 2011) highlighted the prominent antimicrobial, antioxidant activities of *C. stalagmifera* C.E.C.Fisch.

**C. arabica N.E.Br.** Abdel-sattar et al., (2022) revealed that *C. arabica* N.E.Br. is inhabited in Saudi Arabia, Oman, Yemen, United Arab Emirates and Africa and as compared to this study (Zakaria et al., 2001) reported that *C. arabica* N.E.Br. is found in United Arab Emirates. Abdel-sattar et al., (2022) reported that it is medicinally used for skin diseases and conditions (itchy skin, wounds cut) while (Zakaria et al., 2002) reported the use of *C. arabica* for anti-gastric ulcer. Ethanolic extracts of *C. arabica* N.E.Br. exhibited stronger antibacterial effects while least antifungal effects and as evidenced by major inhabitant zones against fungal strains compared to bacterial strains and as it was found that the antifungal extract of *Caralluma arabica* N.E.Br. showed resistant effects against *Fusarium oxysporum* (Al-Mutaani et al., 2025). Methanolic extracts of *C. arabica* included finite number of chemicals comprising of six flavonoids and only one phenolic compound. Epigenine-7-O-glucoside, Quinic acid, Naringenin, Hyperoside

(quercetin-3-O-galactoside) (Al-Mutaani et al., 2025). Opposed to this study (Abdel-Sattar et al., 2022) investigated that arabinoside One of the pregnane glycosides assessed is B. from the aerial parts of *C. arabica* N.E.Br. Antioxidant activity showed by polyphenolics compounds like phenolic acids, flavonoids and ascorbic acid (Williams et al., 2004; Soobrattee et al., 2005), As compared to this study (Khasawneh et al., 2014) revealed that alcoholic extracts of *C. arabica* exhibited antioxidant activity. The methanol friction showed strongest cytotoxic effects while butanol extract exhibited least friction against MCF-7 breast cancer cell line (Khasawneh et al., 2014).

**C. bhupinderana Sarkaria** is found in South India (Lawrence and Choudary, 2004) while (Ugraiah et al., 2011) reported that this plant is depleted by human activities and completely eaten by goats and sheep in Thirunelveli Tamil Nadu and Palayamkotai. It is used for anti-inflammatory, anti-cancer, anti-tumor, anti-ulcer, gastric-mucosa (Deepak et al., 1997; Ramesh et al., 1999; Zakaria et al., 2001). Alkaloids, quinones, phenols, tannins, polysaccharides, saponins, flavonoids, phlobatannins, phytosteroids, terpenoids, and cardiac glycosides (Al-Harbi et al., 1994; Zakaria et al., 2002) while (Pachipala et al., 2022) had similar findings. Methanol *C. bhupinderana* extracts remarkably inhibited UV-irradiated DNA damage therefore considered safe (Umahmaheswari and Chatarjee, 2008; Pachipala et al., 2022) had closely similar results. The *C. bhupinderana* Sarkaria extracts demonstrated action of antioxidants and assessed protective effects against damage to DNA. (Nunes et al., 2012) similarly examined that extracts of *C. sarkaria* had the capability of antioxidant for scavenging free radicals. Nanoparticles prepared from *C. sarkaria* showed great performance against anti-microbial activity (Murugapandi et al., 2023).

**C. retrospiciens (Ehrenb.) N.E.Br.** is mainly distributed in Kenya, Saudi Arabia, Eritrea, Ethiopia, Uganda and Yemen (Bruyns et al., 2010). While (Takhholm, 1974) reported that this plant was grown in south east of Egypt specially on the mountainous areas. The indigenous people of Saudi Arabia used this plant for wound healing. Makeen et al., (2020) reported the first authors that investigated the utilization of *Caralluma retrospiciens* (Ehrenb.) N.E.Br. in cancer treatment by the local people of Wadi-E-Damad and Al-Fayfa (Alallah et al., 2018). Methyl  $\beta$ -lilacinobioside was first isolated from *C. retrospiciens* (Ehrenb.) N.E.Br. (Alallah et al., 2018). While (Halaweish et al., 2004) reported the six polyoxypregnane glycosides evaluated from *C. retrospiciens*. Alkaloids, flavonoids, tannins, steroids, and saponins showed antioxidant effects (Alqahtani et al., 2022). Elsebai and Mohamed. (2015) investigated that A novel-polyoxy pregnane glycosides (retrospinoids) was successfully isolated from the aerial parts of *C. retrospiciens* (Ehrenb.) N.E.Br.. 3,4-Altrosan showed fungicidal and bacteriostatic activities (Jadhav et al., 2014) while the Benzonic acid identified from *C. retrospiciens* demonstrated anti-bacterial effects (Arokiyaraj et al., 2018; Eun-Soo et al., 2001). Compared to this study (Makeen et al., 2020) investigated that the exudate of *Caralluma retrospiciens* (Ehrenb.) N.E.Br. exhibited antimicrobial action against both gram-positive and gram-negative bacteria, with the antimicrobial activity against gram-negative bacteria being greater than that against gram-positive bacteria. All medication toxicity was decreased by the plant-based nanoparticle-based drug delivery systems. (patra et al., 2018; Wided et al., 2021). Opposed to this study (Halaweish et al., 2004) reported that the compounds isolated from *C. retrospiciens* (Ehrenb.) N.E.Br. exhibited cytotoxic effects against brain shrimp

**C. indica** (Wight and Arn.) N.E.Br. is numerously distributed in Tamil Nadu, Karnataka and Andhra Pradesh (Gnanashree et al., 2018). It is used for Cancer (oral curvical, breast lungs, colon and hepatic cancers) (Chandra et al., 2023). The ethanolic extracts of the seeds of *C. indica* (Wight and Arn.) N.E.Br. contained various categories of chemicals including saponins, flavonoids, steroids, terpenoids, polyphenols and tannins (Sembiring et al., 2018). Whereas (Kunert et al., 2006) investigated three steroidal glycosides and two novel bisdesmosidic glycosides from *C. indica*. Chemicals present in the extracts of *C. indica* (Wight and Arn.) N.E.Br. play a vital role in anti-oxidant activities (Shahinuzzaman et al., 2020). Polyphenols and n-hexadecanoic acid evaluated from *C. indica* (Wight and Arn.) N.E.Br. seeds exhibited anti-oxidant activity (Ramalingam et al., 2024). Ethanolic extracts of *C. indica* (Wight and Arn.) N.E.Br. plant demonstrated significant activities against *Pseudomonas* bacteria *Candida cereus* with *Staphylococcus aureus* *Searella typhi* with *Escherichia coli* (Mostafa et al., 2018). And the seeds exhibit natural alternatives to synthetic microbials in the handling of oral pathogens (Ramalingam et al., 2024). *C. indica* (Wight and Arn.) N.E.Br. exhibited remarkable cytotoxic activity on Oral OECM-1 (human oral carcinoma squamous cancerous cells lines) (Koch et al., 2005) and ethanolic seed extracts showed strong cytotoxic activity against *S. cerevisiae* yeast cells (Periyanyagam et al., 2013). Closely similar to these studies (Ramalingam et al., 2024) highlighted that *C. indica* (Wight and Arn.) N.E.Br. displayed strong antioxidant and antimicrobial activities, effectively targeting oral pathogens. It also showed moderate cytotoxic effects against oral cancer cells and yeast with limited selectivity.

## Medicinal uses of Genus *Caralluma* R. Br

*Caralluma tuberculata* N.E.Br. is reported in Pakistan's Balochistan province and Khyber pakhtunkhuwa and it is widely used for the treatment of diabetes, stomach pain and rheumatism, its fresh stems are used as decoctions (Haider et al., 2022; Baig et al., 2021) While (Akram et al., 2023; Ahmed et al., 2013; Waris et al., 2009) reported that *Caralluma edulis* (Edgew.) Benth. Ex Hook.f. is present in the Cholistan desert of Pakistan and is used for the treatment of diabetes, fever and heart problems, its dried apical parts were used as decoction. Similarly, *Caralluma fimbriata* is numerously utilized to cure diabetes and control weight in Indian local medicine, it is used as powder or in capsulated form (Kuriyan et al., 2007). Instantly *Caralluma europea* is present in the regions of Europe and North Africa and it is used to remove kidney stones, heart conditions, digestive and respiratory problem. Its different parts are used as decoctions (Amrati et al., 2020; Ouassau et al., 2021). *Caralluma adscendens* is reported in South India. It is widely used in the treatment of diabetes and inflammation, prepared as paste or decoction. (Bhuvanewari and Manivannan, 2014). Comperately (Bellamakondi et al., 2014; Babu et al., 2014) reported that *Caralluma umbellata* is located in India and it is used to treat stomach disease and diabetes. Its stem and root extracts are used as decoctions. *Caralluma flava* NE Br. Int.J. present in Arabian regions applied as anti-inflammatory, anti-ulcer, antidiabetic diseases and its extracts are prepared as sap (Raees, 2018). *Caralluma sinaica* is reported in Egypt and in Sinai areas. It is used in the treatment of infections, digestive disorders and diabetes. It is prepared as (Habibuddin et al., 2008; Al Massarani et al., 2012). While *Caralluma russeliana* is detected from Arabian regions and it is numerously used for the treatment of lipid profile, liver diseases, diabetes and for removing kidney stones. Stem extracts were ingested (Zari and Al-Thebaiti, 2018).

## PHYTOCHEMICAL COMPOUND CLASS OF GENUS CARALLUMA R. BR

### Flavonoids content:

The correlative screening of phytochemicals in *Caralluma R. Br* species exhibited remarkable differences in phenolic, flavonoids and secondary metabolite constituents among various parts of the plant and extracts solvent. Within the studied phytochemicals, total flavonoid compound (TFC) was identified to be distinguished in all four species. Among them, *C. umbellata* (Bellamakondi et al., 2017) and *C. tuberculata* (Baig et al., 2021; Khan et al., 2022) exhibited significant flavonoid content, while the extracts of *C. indica* confirmed the existence of flavonoids by means of MS-GC assay (Vadivu and Velavan, 2020). Likewise, *C. edulis* (Edgew.) Benth. Ex Hook.f. in methanol extract also demonstrated significant levels of flavonoids. (Baig et al., 2021; Khan et al., 2022).

### Phenolic content

In terms of total phenolic concentration, the Folin-Ciocalteu methods showed that *C. tuberculata* N.E.Br. and *Caralluma edulis* (Edgew.) Benth. Ex Hook.f. have prominently higher phenolic content as compare to other species, of which the most efficient solvent is the methanol (Baig et al., 2021; Khan et al., 2022). *C.umbellata* Haw. also demonstrated prominent levels of phenol which were which were related to hepatoprotective impacts (Bellamakondi et al., 2017). According to Vadivu and Velavan. (2020) *C. indica*'s secondary metabolites originated from phenolic contents of the plant.

### Other Phytochemical classes (Saponins, Alkaloids, Tannins, Steroids, Gallic acid, Rutin)

In addition to phenolics and flavonoids, both qualitative and quantitative tests identified that saponins, alkaloids and tannins were found among all the examined species of *Caralluma R. Br* of these, *C. tuberculata* N.E.Br. and *C. edulis* (Edgew.) Benth. Ex Hook.f. especially showed great levels of these metabolites in their methanolic and hydroalcoholic extracts (Baig et al., 2021; Khan et al., 2022). Likewise, GC-MS assays and colorimetric standard tests also identified the presence of these phytochemicals in the *C. umbellata* and *C. indica* (Bellamakondi et al., 2017; Vadivu and Velavan, 2020).

According to (Baig et al., 2021; Khan et al., 2022; Bellamakondi et al., 2017) the developed techniques of chromatography including (GC-MS and UPLC/HPLC identified the presence of Rutin, quercetin and gallic acid in *C. tuberculata*, *C. edulis* (Edgew.) Benth. Ex Hook.f. and *Caralluma umbellata*, whereas Vadivu and Velavan. (2020) reported phenolic acids and fatty acids derivatives in the *C. indica*. Chemical diversity is moreover elaborated by GC-MS techniques. Among them 30+ compounds reported in *C. umbellata* and *tuberculata* (Khan et al., 2022; Bellamakondi et al., 2017) and approximately 20-32 compounds detected in *C. edulis* (Edgew.) Benth. Ex Hook.f. and *C. indica* (Iftikhar et al., 2022; Vadivu and Velavan, 2020).

Across these aromatic compounds, phytol, fatty acid esters, hexadecanoic acid were detected among *C. tuberculata*, *C. edulis* (Edgew.) Benth. Ex Hook.f., *C. umbellata* and *C. indica* (Khan et al., 2022; Iftikhar et al., 2022; Bellamakondi et al., 2017; Vadivu and Velavan, 2020). Due to phytochemical screening assay all four species significantly showed reliable presence of phytosterols, terpenoids, glycosides and steroids. Particularly *C. edulis*

(Edgew.) Benth. Ex Hook.f. and *C. tuberculata* exhibited rich profiles of these metabolites, while *C. indica* and *C. umbellata* dominantly had fatty acid esters, terpenoids and phytosterols and glycosides. (Baig et al., 2021; Khan et al., 2022; Vadivu & Velavan, 2020; Bellamakondi et al., 2017).

## CONCLUSION

*Caralluma R. Br* is a desertic genus that belongs to the family Apocynaceae sub family Asclepiadoideae. It is classified under the order Ceropegieae suborder Stapeliinae. It contains approximately 120 species. This Genus has various phytochemical compounds that are linked to its medicinal, anti-oxidant, anti-microbial and toxicity properties. The species of the *Caralluma R. Br* genus are habited in Pakistan, Saudi Arabia, Egypt, Europe, India, Ethiopia, Yamen, Spain, Kenya, Srilanka and canar islands. They are variously used in traditional medicines. Including, cancer, Blood diseases, rhumatisim, inflammations, lipid profile, Dysentery, hepatitis, ulcers, hypertention, gastric, intestinal issues, malaria, TB (tuberculosis), Mellitus, obesity, appetite and weight management, Scorpions and snake bites. Alkaloids, glycosides, teropenoids, flavonoids, tannins, carotenoids, phenolics, quercetin, saponins, rutin, gallic acids, were found in various species of *Caralluma R. Br*. These chemicals were isolated by using different tecniques, including GC-MS, colorimetric-gravimetric, Screening and by folin-ciocalteu methods. It showed strong antimicrobial activity against most dangerous bacteria and fungi. The extracts from roots, leaves and stems were reported effectively towards *Bacillus* species and *Eschrichiacoli*. At moderate doses the species were generally found safe but at overdoses they may have mild toxicity. *Caralluma R. Br* is a genus of xerophytic regions and it have various biological activitiies that are associated with their phytochemical compounds.

## RECOMMENDATIONS

- More studies should be focused that how phytochemicals of the genus *Caralluma R. Br* worked in the body. For instanse the pathogenic bacteria and fungi should be controlled and prevent the body cells from injury.
- *Caralluma R. Br* species should be tested first in animal models and after then in humans to confirm the safety.
- Endangered species of Genus *Caralluma R. Br* should be conserved.
- After testing the *Caralluma R. Br* species in the laboratories herbal capsules should be made for humans.
- The indegenious knowledge from local people about the medicinal uses of genus *Caralluma R. Br* should be documented.

## REFERENCES

1. Stewart, R. R. (1972). An annotated catalogue of the vascular plants of West Pakistan and Kashmir.
2. Abdel-Sattar, O. E., Sabry, M. M., Shalabi, A. A., El-Halawany, A. M., Al-Hawshabi, O. S., Abdel-Sattar, E., ... & El-Shiekh, R. A. (2025). Genus *Caralluma R. Br*. in Yemen: A comprehensive review of taxonomy, ethnomedicine, phytochemistry, and biological activities. *Chemistry & Biodiversity*, e01112.
3. Naik, M. R., & Krishnamurthy, Y. L. (2012). Xerophyte *Caralluma stalagmiferavar. longipetala* (Asclepiadaceae): a new record to the flora of Karnataka, India. *Journal of Threatened Taxa*, 4(6), 2656-2659.
4. Karale, P. A., & Karale, M. A. (2017). A review on phytochemistry and pharmacological properties of milkweed family herbs (Asclepiadaceae). *Asian Journal of Pharmaceutical and Clinical Research*, 10(11), 27-34.
5. Braca, A., Bader, A., Morelli, I., Scarpato, R., Turchi, G., Pizza, C., & De Tommasi, N. (2002). New pregnane glycosides from *Caralluma negevensis*. *Tetrahedron*, 58(29), 5837-5848.
6. Braca, A., Bader, A., Morelli, I., Scarpato, R., Turchi, G., Pizza, C., & De Tommasi, N. (2002). New pregnane glycosides from *Caralluma negevensis*. *Tetrahedron*, 58(29), 5837-5848.
7. Bader, A., Braca, A., De Tommasi, N., & Morelli, I. (2003). Further constituents from *Caralluma negevensis*. *Phytochemistry*, 62(8), 1277-1281.
8. Abdallah, H. M., Osman, A. M. M., Almehdar, H., & Abdel-Sattar, E. (2013). Acylated pregnane glycosides from *Caralluma quadrangula*. *Phytochemistry*, 88, 54-60.
9. Abdel-Sattar, E. A., Abdallah, H. M., Khedr, A., Abdel-Naim, A. B., & Shehata, I. A. (2013). Antihyperglycemic activity of *Caralluma tuberculata* in streptozotocin-induced diabetic rats. *Food and chemical toxicology*, 59, 111-117.
10. Abdel-Sattar, E. A., Al-Hawshabi, O. S., Shalabi, A. A., El Halawany, A. M., & Meselhy, M. R. (2022). Arabincosides AD, pregnane glycosides isolated from *Caralluma arabica*. *Tetrahedron*, 119, 132858.
11. Abdel-Sattar, E., Ahmed, A. A., Hegazy, M. E. F., Farag, M. A., & Al-Yahya, M. A. A. (2007). Acylated pregnane glycosides from *Caralluma russeliana*. *Phytochemistry*, 68(10), 1459-1463.

12. Abdel-Sattar, E., EL-Maraghy, S. A., El-Dine, R. S., & Rizk, S. M. (2017). Antihyperglycemic activity of *Caralluma quadrangulata* streptozotocin-induced diabetic rats. *Bulletin of Faculty of Pharmacy, Cairo University*, 55(2), 269-272.
13. Abdel-Sattar, E., Harraz, F. M., Al-Ansari, S. M. A., El-Mekkawy, S., Ichino, C., Kiyohara, H., ... & Yamada, H. (2008). Acylated pregnane glycosides from *Caralluma tuberculata* and their antiparasitic activity. *Phytochemistry*, 69(11), 2180-2186.
14. Abdel-Sattar, E., Kutkat, O., El-Shiekh, R. A., El-Ashrey, M. K., & El Kerdawy, A. M. (2024). In Silico and in Vitro Screening of some pregnane glycosides isolated from certain *Caralluma R. Br.* species as SARS-COV-2 main protease inhibitors. *Chemistry & Biodiversity*, 21(4), e202301786.
15. Abdel-Sattar, E., Mehanna, E. T., El-Ghaiesh, S. H., Mohammad, H. M., Elgendy, H. A., & Zaitone, S. A. (2018). Pharmacological action of a pregnane glycoside, russelioside B, in dietary obese rats: impact on weight gain and energy expenditure. *Frontiers in pharmacology*, 9, 990.
16. Abdel-Sattar, O. E., Sabry, M. M., Shalabi, A. A., El-Halawany, A. M., Al-Hawshabi, O. S., Abdel-Sattar, E., ... & El-Shiekh, R. A. (2025). Genus *Caralluma R. Br.* in Yemen: A comprehensive review of taxonomy, ethnomedicine, phytochemistry, and biological activities. *Chemistry & Biodiversity*, e01112.
17. Abdel-Sattar, O. E., Sabry, M. M., Shalabi, A. A., El-Halawany, A. M., Al-Hawshabi, O. S., Abdel-Sattar, E., ... & El-Shiekh, R. A. (2025). Genus *Caralluma R. Br.* in Yemen: A comprehensive review of taxonomy, ethnomedicine, phytochemistry, and biological activities. *Chemistry & Biodiversity*, e01112.
18. Adnan, M., Jan, S., Mussarat, S., Tariq, A., Begum, S., Afroz, A., & Shinwari, Z. K. (2014). A review on ethnobotany, phytochemistry and pharmacology of plant genus *Caralluma R. Br.*. *Journal of Pharmacy and Pharmacology*, 66(10), 1351-1368.
19. Ahmad, M. M., & Shaikh, M. M. (1989). Improvement in glucose tolerance by *Caralluma tuberculata*, *Acacia nilotica* and *Papaver somniferum*. *Pakistan Journal of Zoology (Pakistan)*, 21(4).
20. Ahmad, M., Qureshi, R., Arshad, M., Khan, M. A., & Zafar, M. (2009). Traditional herbal remedies used for the treatment of diabetes from district Attock (Pakistan). *Pak J Bot*, 41(6), 2777-2782.
21. Ahmed, M. M., Qureshi, S., Al-Bekairi, A. M., Shah, A. H., Rao, R. M., & Qazi, N. S. (1993). Anti-inflammatory activity of *Caralluma tuberculata* alcoholic extract.
22. Ahmed, M. M., Qureshi, S., Al-Bekairi, A. M., Shah, A. H., Rao, R. M., & Qazi, N. S. (1993). Anti-inflammatory activity of *Caralluma tuberculata* alcoholic extract.
23. Ait Dra, L., Aghraz, A., Boualy, B., Oubaassine, S., Barakate, M., Markouk, M., & Larhsini, M. (2018). Chemical characterization and in vitro antimicrobial activity of *Caralluma europea* essential oil and its synergistic potential with conventional antibiotics.
24. Ait Dra, L., Sellami, S., Rais, H., Aziz, F., Aghraz, A., Bekkouche, K., ... & Larhsini, M. (2019). Antidiabetic potential of *Caralluma europea* against alloxan-induced diabetes in mice. *Saudi journal of biological sciences*, 26(6), 1171-1178.
25. Akram, A., Jamshed, A., Anwaar, M., Rasheed, H. M. F., Haider, S. I., Aslam, N., & Jabeen, Q. (2023). Evaluation of *Caralluma edulis* for its potential against obesity, atherosclerosis and hypertension. *Dose-Response*, 21(1), 15593258231152112.
26. Alallah, M. I., Alhemaïd, F., Bai, F., Mothana, R. A., Elshikh, M. S., Farah, M. A., ... & Al-Anazi, K. M. (2018). The binding proximity of methyl  $\beta$ -lilacinobioside isolated from *Caralluma retrospiciens* with topoisomerase II attributes apoptosis in breast cancer cell line. *Saudi Journal of Biological Sciences*, 25(8), 1826-1833.
27. Alamier, W. M., Hasan, N., Syed, I. S., Bakry, A. M., Ismail, K. S., Gedda, G., & Girma, W. M. (2023). Silver nanoparticles' biogenic synthesis using *Caralluma R. Br. subulata* aqueous extract and application for dye degradation and antimicrobials activities. *Catalysts*, 13(9), 1290.
28. Albaser, N. A. (2024). *CARALLUMA PENICILLATA (DEFLERS) NE BR.(ASCLEPIADACEAE): A MINI REVIEW ON ITS TRADITIONAL USES, PHYTOCHEMICAL COMPOSITIONS, AND PHARMACOLOGICAL PROPERTIES.*
29. Albaser, N. A., & AL-Kamarany, M. A. (2023). Investigating *Caralluma penicillata*'s Medicinal Properties: Effects on Haematological Parameters and Renal Functions in Cotton Pellet Induced Granuloma in Adult Guinea Pigs. *Al-Razi University Journal for Medical Sciences*, 7(2), 1-7.
30. Albers, F., & Meve, U. (Eds.). (2012). *Illustrated handbook of succulent plants: Asclepiadaceae*. Springer Science & Business Media.
31. Alharbi, M. M., Qureshi, S., Raza, M., Ahmed, M. M., Afzal, M., & Shah, A. H. (1994). Evaluation of *Caralluma tuberculata* pretreatment for the protection of rat gastric mucosa against toxic damage. *Toxicology and applied pharmacology*, 128(1), 1-8.

32. Ali, A., Mashwani, Z. U. R., Raja, N. I., Mohammad, S., Ahmad, M. S., & Luna-Arias, J. P. (2024). Antioxidant and Hypoglycemic Potential of Phytogetic Selenium Nanoparticle-and Light Regime-Mediated In Vitro *Caralluma tuberculata* Callus Culture Extract. *ACS omega*, 9(18), 20101-20118.
33. Ali, M. M., Al-Mokaddem, A. K., Abdel-Sattar, E., El-Shiekh, R. A., Farag, M. M., Aljuaydi, S. H., & Shaheed, I. B. (2024). Enhanced wound healing potential of arabinoside B isolated from *Caralluma arabica* in rat model; a possible dressing in veterinary practice. *BMC Veterinary Research*, 20(1), 282.
34. Almaqtari, M. A., & Mubarak, A. Y. (2024). Antioxidant and antimicrobial of three extracts of *Caralluma deflersiana* Laver. *Sana'a University Journal of Applied Sciences and Technology*, 2(2), 154-157.
35. Al-Massarani, S. M., Bertrand, S., Nievergelt, A., El-Shafae, A. M., Al-Howiriny, T. A., Al-Musayeb, N. M., ... & Wolfender, J. L. (2012). Acylated pregnane glycosides from *Caralluma R. Br. sinaica*. *Phytochemistry*, 79, 129-140.
36. Al-Mutaani, J., Zourgui, L., & Missaoui, N. (2025). Phytochemicals, bioactive compounds, and antimicrobial activities of *Ocimum basilicum*, *Teucrium polium*, *Cleome amblyocarpa*, and *Caralluma arabica* extracts: a comparative Omani study. *Cellular and Molecular Biology*, 71(3), 134-145.
37. Alodaini, H. A., Alarjani, K. M., Alkubaisi, N. A., Aziz, I. M., Alsayed, M. F., Almarfadi, O. M., ... & Aboul-Soud, M. A. (2025). *Caralluma tuberculata* (Chongan): Chemical Profiling, Antioxidant, Anticancer, Antibacterial and Antidiabetic Potential supported by In Silico Evidence. *ChemistrySelect*, 10(13), e202405946.
38. Alqahtani, S. N., Alkholy, S. O., & Ferreira, M. P. (2014). Antidiabetic and anticancer potential of native medicinal plants from Saudi Arabia. In *Polyphenols in human health and disease* (pp. 119-132). Academic Press.
39. Alqahtani, S. S., Makeen, H. A., Menachery, S. J., & Moni, S. S. (2020). Documentation of bioactive principles of the flower from *Caralluma retrospiciens* (Ehrenb) and in vitro antibacterial activity–Part B. *Arabian Journal of Chemistry*, 13(10), 7370-7377.
40. Al-Rubaye, A. F., Hameed, I. H., & Kadhim, M. J. (2017). A review: uses of gas chromatography-mass spectrometry (GC-MS) technique for analysis of bioactive natural compounds of some plants. *International Journal of Toxicological and Pharmacological Research*, 9(1), 81-85.
41. Alshehri, T., Alkhalifah, I., Alotaibi, A., Alsulaiman, A. F., Al Madani, A., Almutairi, B., & Balhaddad, A. A. (2025). The impact of *Caralluma R. Br. munbyana* extracts on *Streptococcus mutans* biofilm formation. *Frontiers in Dental Medicine*, 6, 1575161.
42. Amrati, F. E. Z., Bourhia, M., Saghrouchni, H., Slighoua, M., Grafov, A., Ullah, R., ... & Bousta, D. (2021). *Caralluma europea*(Guss.) NE Br.: Anti-inflammatory, antifungal, and antibacterial activities against nosocomial antibiotic-resistant microbes of chemically characterized fractions. *Molecules*, 26(3), 636.
43. Amrati, F. E. Z., Bourhia, M., Slighoua, M., Ibnemoussa, S., Bari, A., Ullah, R., ... & Bousta, D. (2020). Phytochemical study on antioxidant and antiproliferative activities of Moroccan *Caralluma europea* extract and its bioactive compound classes. *Evidence-Based Complementary and Alternative Medicine*, 2020(1), 8409718.
44. Amrati, F. E. Z., Elmadbouh, O. H. M., Chebaibi, M., Soufi, B., Conte, R., Slighoua, M., ... & Bousta, D. (2023). Evaluation of the toxicity of *Caralluma europea*(CE) extracts and their effects on apoptosis and chemoresistance in pancreatic cancer cells. *Journal of Biomolecular Structure and Dynamics*, 41(17), 8517-8534.
45. Amrati, F. E. Z., Lim, A., Slighoua, M., Chebaibi, M., Mssillou, I., Drioiche, A., ... & Bousta, D. (2025). Unraveling the hepatoprotective and anti-pancreatic cancer potential of *Caralluma europaea*: a comprehensive in vivo, in vitro and in silico evidence. *Drug and Chemical Toxicology*, 48(1), 120-135.
46. Ansari, B., Behl, T., Pirezada, A. S., & Khan, H. (2022). *Caralluma edulis*(Apocynaceae): A comprehensive review on its traditional uses, phytochemical profile and pharmacological effects. *Current Topics in Medicinal Chemistry*, 22(18), 1501-1514.
47. Anwar, R., Rabail, R., Rakha, A., Bryla, M., Roszko, M., Aadil, R. M., & Kieliszek, M. (2022). Delving the role of *Caralluma fimbriata*: an edible wild plant to mitigate the biomarkers of metabolic syndrome. *Oxidative medicine and cellular longevity*, 2022(1), 5720372.
48. Arif, A., Sultan, M. T., Nazir, F., Ahmad, K., Kashif, M., Ahmad, M. M., ... & Rocha, J. M. (2024). Exploring the therapeutic potential of *Caralluma fimbriata* for antioxidant and diabetes management: a 28-day rat model study. *Toxicology Research*, 13(4), tfae094.
49. Arokiyaraj, S., Bharanidharan, R., Agastian, P., & Shin, H. (2018). Chemical composition, antioxidant activity and antibacterial mechanism of action from *Marsilea minuta* leaf hexane: methanol extract. *Chemistry Central Journal*, 12(1), 105.
50. Aslam, I., Iqbal, J., Peerzada, S., Afridi, M. S., & Ishtiaq, S. (2019). Microscopic investigations and pharmacognostic techniques for the standardization of *Caralluma edulis*(Edgew.) Benth. ex Hook. f. *Microscopy research and technique*, 82(11), 1891-1902.

51. Babu, K. S., Malladi, S., Nadh, R. V., & Rambabu, S. S. (2014). Evaluation of in vitro antibacterial activity of *Caralluma umbelleta* Haw used in traditional medicine by Indian tribes. *Annual research & Review in Biology*, 4(6), 840.
52. Bader, A., Braca, A., De Tommasi, N., & Morelli, I. (2003). Further constituents from *Caralluma negevensis*. *Phytochemistry*, 62(8), 1277-1281.
53. Bader, A., Braca, A., De Tommasi, N., & Morelli, I. (2003). Further constituents from *Caralluma negevensis*. *Phytochemistry*, 62(8), 1277-1281.
54. Baig, M. W., Ahmed, M., Akhtar, N., Okla, M. K., Nasir, B., Haq, I. U., ... & AbdElgawad, H. (2021). *Caralluma tuberculata* NE Br manifests extraction medium reliant disparity in phytochemical and pharmacological analysis. *Molecules*, 26(24), 7530.
55. Baquar, S. R. (1989). Medicinal and poisonous plants of Pakistan. Printas.
56. Batool, S., Batool, S., Batool, T., Iram, F., Almas, T., Faizan, M., ... & Arif, H. (2024). Delving the Role of the Ameliorative Effects of *Caralluma tuberculata* NE Br.(Apocynaceae) on Diabetes and Its Effect on the Organs Weight of Alloxan-Induced Adult Male Mice. *Polish Journal of Environmental Studies*, 33(1).
57. Bellakhdar, J., Claisse, R., Fleurentin, J., & Younos, C. (1991). Repertory of standard herbal drugs in the Moroccan pharmacopoea. *Journal of ethnopharmacology*, 35(2), 123-143.
58. Bellamakondi, P. K., Godavarthi, A., & Ibrahim, M. (2014). Anti-hyperglycemic activity of *Caralluma umbelleta* Haw. *BioImpacts: BI*, 4(3), 113.
59. Bellamakondi, P. K., Godavarthi, A., & Ibrahim, M. (2017). *Caralluma umbelleta* Haw. protects liver against paracetamol toxicity and inhibits CYP2E1. *BioImpacts: BI*, 8(1), 23.
60. Ben Said, R., Ben Aissa, M. A., Rahali, S., Tangour, B., Kowalczyk, M., Oleszek, W., ... & Hamed, A. I. (2023). Fingerprinting of two an acylated polyoxypregnane glycosides from *Caralluma quadrangula*(Forssk.) NE Br. using UPLC-ESI-Q-TOF and computational study. *Natural Product Research*, 37(1), 136-140.
61. Benazzouz, B. (2009). Taxonomy and conservation status of Moroccan stapeliads (Apocynaceae-Asclepiadoideae-Ceropegieae-Stapeliinae).
62. Bennett, R. (2003). The 'direct costs' of livestock disease: the development of a system of models for the analysis of 30 endemic livestock diseases in Great Britain. *Journal of Agricultural Economics*, 54(1), 55-71.
63. BENSUSAN, K. (1913). Taxonomy and conservation status of moroccan stapeliads (Apocynaceae-Asclepiadoideae-Ceropegieae-Stapeliinae). *Bull. Misc. Inf.(Kew)*, 121(1913).
64. Bhuvanewari, S., & Manivannan, S. (2014). Anti-diabetic and anti-inflammatory activity of *Caralluma adscendens*var. *adscendens*.
65. Braca, A., Bader, A., Morelli, I., Scarpato, R., Turchi, G., Pizza, C., & De Tommasi, N. (2002). New pregnane glycosides from *Caralluma negevensis*. *Tetrahedron*, 58(29), 5837-5848.
66. Brenes, A., & Roura, E. (2010). Essential oils in poultry nutrition: Main effects and modes of action. *Animal feed science and technology*, 158(1-2), 1-14.
67. Bruyns, P. V. (1987). The genus *Caralluma* R. Br.own (Asclepiadaceae) in Israel. *Israel Journal of Plant Sciences*, 36(2), 73-86.
68. Bukhari, I. A., Gilani, A. H., Meo, S. A., & Saeed, A. (2016). Analgesic, anti-inflammatory and anti-platelet activities of *Buddleja crispa*. *BMC complementary and alternative medicine*, 16(1), 79.
69. Bukhari, I. A., Khan, R. A., Gilani, A. U. H., Shah, A. J., Hussain, J., & Ahmad, V. U. (2007). The analgesic, anti-inflammatory and calcium antagonist potential of *Tanacetum artemisioides*. *Archives of pharmacal research*, 30(3), 303-312.
70. Bukhari, S. A., Behramand, A. A. S. B., Shah, S. S., & Umm-e-Aiman, M. H. Z. Phytochemical Screening Of Different Fractions And Anti-Bacterial Activity Of *Caralluma tuberculata*.2024.
71. Burda, S., & Oleszek, W. (2001). Antioxidant and antiradical activities of flavonoids. *Journal of agricultural and food chemistry*, 49(6), 2774-2779.
72. Burt, S. (2004). Essential oils: their antibacterial properties and potential applications in foods—a review. *International journal of food microbiology*, 94(3), 223-253.
73. Chandra, S., Gahlot, M., Choudhary, A. N., Palai, S., de Almeida, R. S., de Vasconcelos, J. E. L., ... & Coutinho, H. D. M. (2023). Scientific evidences of anticancer potential of medicinal plants. *Food Chemistry Advances*, 2, 100239.
74. Chandran, R., Sajeesh, T., & Parimelazhagan, T. (2014). Total Phenolic Content, Anti-Radical property and HPLC profiles of *Caralluma* R. Br. *diffusa* (Wight) NE Br. *Journal of Biologically Active Products from Nature*, 4(3), 188-195.

75. Collenette S (1999). Wild Flowers of Saudi Arabia. National commission for wildlife conservation and development, Kingdom of Saudi Arabia, p. 52 Chemistry, 13(8), 6672-6681.
76. Dallahi, Y., El Aboudi, A., Aafi, A., & Belghazi, B. (2016). Inventory of medicinal plants in the site of biological and ecological interest of Kharouba (Central Plateau, Morocco). *Journal of Materials and Environmental Sciences*, 7(12), 3993–3999.
77. De Leo, M., De Tommasi, N., Sanogo, R., Autore, G., Marzocco, S., Pizza, C., ... & Braca, A. (2005). New pregnane glycosides from *Caralluma dalzielii*. *Steroids*, 70(9), 573-585.
78. de Souza, T. A., Lins, F. S. V., da Silva Lins, J., Alves, A. F., Cibulski, S. P., Brito, T. D. A. M., ... & Tavares, J. F. (2024). Asclepiadoideae subfamily (Apocynaceae): ethnopharmacology, biological activities and chemophenetics based on pregnane glycosides. *Phytochemistry Reviews*, 23(4), 1027-1063.
79. Deepak, D. (1996). *Progress in the chemistry of organic natural products*. Springer-Verlag.
80. Dohou, R., Yamni, K., Tahrouch, S., Hassani, L. I., Badoc, A., & Gmira, N. (2003). Screening phytochimique d'une endémique iberomarocaine, *Thymelaea lythroides*. *Bulletin-Société de Pharmacie de Bordeaux*, 142(1/4), 61-78.
81. Durrani, M. J., Manzoor, M., & Irfan, S. (2009). Folk uses of some plants of Quetta, Pakistan. *Pakistan Journal of Plant Sciences*, 15(1), 1–6.
82. Dutt, H. C., Singh, S., Avula, B., Khan, I. A., & Bedi, Y. S. (2012). Pharmacological review of *Caralluma R. Br...* with special reference to appetite suppression and anti-obesity. *Journal of medicinal food*, 15(2), 108-119.
83. Dutt, H. C., Singh, S., Avula, B., Khan, I. A., & Bedi, Y. S. (2012). Pharmacological review of *Caralluma R. Br...* with special reference to appetite suppression and anti-obesity. *Journal of medicinal food*, 15(2), 108-119.
84. El Khatabi, K., Alaqrbeh, M., Rehman, H. M., Ajana, M. A., Lakhlifi, T., & Bouachrine, M. (2024). Computational Investigation of Potent EGFR Inhibitors from Flavonoid-Based Phytochemical Constituents of *Caralluma europeaas* Pancreatic Cancer Agents. *Physical Chemistry Research*, 12(4), 963-974.
85. Elisha, I. L., Botha, F. S., McGaw, L. J., & Eloff, J. N. (2017). The antibacterial activity of extracts of nine plant species with good activity against *Escherichia coli* against five other bacteria and cytotoxicity of extracts. *BMC complementary and alternative medicine*, 17(1), 133.
86. Elsebai, M. F., & Mohamed, I. E. T. (2015). New pregnane glycoside derivative from *Caralluma retrospicens* (Ehrenb). *Natural Product Research*, 29(15), 1426-1431.
87. El-Shiekh, R. A., Hassan, M., Hashem, R. A., & Abdel-Sattar, E. (2021). Bioguided isolation of antibiofilm and antibacterial pregnane glycosides from *Caralluma quadrangula*: Disarming multidrug-resistant pathogens. *Antibiotics*, 10(7), 811.
88. El-Shiekh, R. A., Shalabi, A. A., Al-Hawshabi, O. S., Salkini, M. A., & Abdel-Sattar, E. (2023). Anticholinesterase and anti-inflammatory constituents from *Caralluma awdeliana*, a medicinal plant from Yemen. *Steroids*, 193, 109198.
89. Ennacerie, F.-Z., Rhazi Filali, F., & Rahou, A. (2017). Ethnobotanical study of medicinal plants used in traditional medicine in the province of Sidi Kacem, Morocco. *Asian Journal of Pharmaceutical and Clinical Research*, 10(1), 121–130.
90. Enogieru, A. B., Haylett, W., Hiss, D. C., Bardien, S., & Ekpo, O. E. (2018). Rutin as a potent antioxidant: Implications for neurodegenerative disorders. *Oxidative medicine and cellular longevity*, 2018(1), 6241017.
91. Farouk, A. E., Ahamed, N. T., AlZahrani, O., Alamer, K. H., Al-Sodany, Y., & Bahobail, A. A. (2016). Antimicrobial activity of *Caralluma quadrangula*(Forssk) NE Br latex from Al-ShafaTaif, Kingdom of Saudi Arabia. *Int J Curr Microbiol App Sci*, 5, 284-98.
92. Ferreira, C., Costa, S. S., Serrano, M., Oliveira, K., Trigueiro, G., Pomba, C., & Couto, I. (2021). Clonal lineages, antimicrobial resistance, and PVL carriage of *Staphylococcus aureus* associated to skin and soft-tissue infections from ambulatory patients in Portugal. *Antibiotics*, 10(4), 345.
93. Formisano, C., Senatore, F., Della Porta, G., Scognamiglio, M., Bruno, M., Maggio, A., ... & Sajeve, M. (2009). Headspace volatile composition of the flowers of *Caralluma europea*NE Br.(Apocynaceae). *Molecules*, 14(11), 4597-4613.
94. Gabali, S. A., & Al-Gifri, A. N. (1990). Flora of South Yemen—Angiospermae A provisional checklist. *Feddes Repertorium*, 101(7-8), 373-383.
95. Gilbert, M. G. (1990). A review of *Caralluma R. Br...* and its segregates. *Bradleya*, 1990(8), 1-32.
96. Gillani, B., Tariq, S., Shahzad, M. I., Fatima, T., Locatelli, M., Cai, X., ... & Ahmad, A. (2024). Phytochemical composition and therapeutic potential of *Caralluma edulisa cholistani* plant. *Journal of King Saud University-Science*, 36(11), 103519.

97. Gnanashree, G., Sirajudeen, P. M., & Sirajudeen, M. (2018). Determination of bioactive compounds in ethanolic extract of *Caralluma indica* using GC-MS technique. *J pharmacogn phytochem*, 7(6), 1675-7.
98. Gushash, A. S. (2006). *Plants in the Mountains of Sarat and Hejaz*. Sarawat Designer and Printers: Madinah, Saudi Arabia, 2, 15-19.
99. Habibuddin, M., Daghri, H. A., Humaira, T., Al Qahtani, M. S., & Hefzi, A. A. H. (2008). Antidiabetic effect of alcoholic extract of *Caralluma R. Br. sinaica L.* on streptozotocin-induced diabetic rabbits. *Journal of ethnopharmacology*, 117(2), 215-220.
100. Haider, S. I., Asif, A., Rasheed, H. M. F., Akram, A., & Jabeen, Q. (2022). *Caralluma tuberculata* exhibits analgesic and anti-arthritic potential by downregulating pro-inflammatory cytokines and attenuating oxidative stress. *Inflammopharmacology*, 30(2), 621-638.
101. Halaweish, F. T., Huntimer, E., & Khalil, A. T. (2004). Polyoxypregnane glycosides from *Caralluma retrospiciens*. *Phytochemical Analysis: An International Journal of Plant Chemical and Biochemical Techniques*, 15(3), 189-194.
102. Ibrahim, H. M., Saleem, H. A., Alhadi, F. A., Alhammadi, A. S., & Newton, L. E. (2024). Stem epidermal properties of four *Caralluma R. Br.* (Apocynaceae) species in Yemen and their taxonomic significance. *Phytologia Balcanica*, 30(3).
103. Iftikhar, N., Saleem, A., Akhtar, M. F., Abbas, G., Shah, S., Bibi, S., Ashraf, G. M., Alghamdi, B. S., & Abujamel, T. S. (2022). In vitro and in vivo anti-arthritic potential of *Caralluma tuberculata* N. E. Brown and its chemical characterization. *Molecules*, 27(19), 6323.
104. Ishaq, H., Rajendran, K., & Nisar, K. (2023). A comprehensive review of medicinal uses and phytochemicals isolated from *Caralluma tuberculata*. *Innovations in Agriculture*, 6, 01–11.
105. Iftikhar, N., Saleem, A., Akhtar, M. F., Abbas, G., Shah, S., Bibi, S., ... & Abujamel, T. S. (2022). In vitro and in vivo anti-arthritic potential of *Caralluma tuberculata* NE Brown. And its chemical characterization. *Molecules*, 27(19), 6323.
106. Ismail, G. A., Mostafa, M. E., Mubarak, A. Y., Dawidar, A. M., & Abdel-Mogib, M. (2024). Cytotoxic pregnane glycosides from *Monolluma quadrangula* (Forssk.). *Natural Product Research*, 1-10.
107. Issiki, Z., Moundir, C., Marnissi, F., Seddik, N., Benjelloun, N., Zaid, Y., & Oudghiri, M. (2017). Toxicological evaluation of the aqueous extract of *Caralluma europea* and its immunomodulatory and inflammatory activities. *Pharmacognosy research*, 9(4), 390.
108. Jamil, K., Khan, M. R., Jan, A., & Ali, G. M. (2023). Screening of phytochemical compounds and antimicrobial activity of *Caralluma tuberculata* by In-Silico study. *Sarhad Journal of Agriculture*, 39(4), 983-989.
109. Jaswanth, B. K., Reddy, P. V. B., & Kiranmai, C. (2025). In vitro Strategies for the Production of Bioactive Therapeutics from *Caralluma R. Br.* Species. *Asian Journal of Biological and Life Sciences*, 14(1), 1-7.
110. Jayaprakash, K., Manokari, M., Cokulraj, M., Dey, A., Faisal, M., Alatar, A. A., ... & Shekhawat, M. S. (2023). Improved organogenesis and micro-structural traits in micropropagated plantlets of *Caralluma umbellata* Haw. in response to Meta-Topolin. *Plant Cell, Tissue and Organ Culture (PCTOC)*, 153(1), 105-118.
111. Jayawardena, R., Francis, T. V., Abhayaratna, S., & Ranasinghe, P. (2021). The use of *Caralluma fimbriata* as an appetite suppressant and weight loss supplement: a systematic review and meta-analysis of clinical trials. *BMC complementary medicine and therapies*, 21(1), 279.
112. Jesudass, J. S., Sivaprakasam, B., Kulanthaivel, S. R., Muthukrishnan, A., Chinnaiyan, R., Ramasamy, R., ... & Balasubramani, R. (2025). Computational Identification of Bioactive Molecules from *Caralluma stalagmifera* L. as Potential VEGFR2 Inhibitors for Endometriosis Treatment. *Journal of Pharmaceutical Innovation*, 20(1), 1-21.
113. Kamel, E. A. R., Sharawy, S. M., & Karakish, E. A. K. (2014). Cytotaxonomical investigations of the tribes Asclepiadeae and Ceropegieae of the subfamily Asclepiadoideae-Apocynaceae. *Pak. J. Bot*, 46(4), 1351-1361.
114. Karale, P. A., & Karale, M. A. (2017). A review on phytochemistry and pharmacological properties of milkweed family herbs (Asclepiadaceae). *Asian Journal of Pharmaceutical and Clinical Research*, 10(11), 27-34.
115. Khan, M., Manzoor, Z., Rafiq, M., Munawar, S. H., Waqas, M. Y., Majeed, H., ... & Mojzych, M. (2022). Phytochemical screening, anti-inflammatory, and antidiabetic activities of different extracts from *Caralluma edulis* plant. *Molecules*, 27(16), 5346.
116. Kiros, T., Mohammed, S., Dekebo, A., & Melaku, Y. (2023). In Silico Pharmacokinetics Properties and In Vitro Bioactivities of Pregnane Derivatives and Other Compounds From Stems of *Caralluma R. Br. speciosa*. [Journal of Plant Research or relevant journal].
117. Koch, A., Tamez, P., Pezzuto, J., & Soejarto, D. (2005). Evaluation of plants used for antimalarial treatment by the Maasai of Kenya. *Journal of ethnopharmacology*, 101(1-3), 95-99.
118. Kumar, D. S., David Banji, D. B., & Harani, A. (2011). A medicinal plants survey for treatment of obesity.

119. Kumar, K. M. P., Murshida, U. C., Thomas, B., George, S., Balachandran, I., & Karuppusamy, S. (2014). Notes on *Caralluma adscendens* (Roxb.) Haw. var. *attenuata* (Wight) Grav. and Mayur. (Apocynaceae: Asclepiadoideae). *Journal of Threatened Taxa*, 6(9), 6282-6286.
120. Kunert, O., Rao, B. V. A., Babu, G. S., Padmavathi, M., Kumar, B. R., Alex, R. M., ... & Rao, A. V. N. A. (2006). Novel steroidal glycosides from two Indian *Caralluma* R. Br. species, *C. stalagmifera* and *C. indica*. *Helvetica chimica acta*, 89(2), 201-209.
121. Kuriyan, R., Raj, T., & Kurpad, A. V. (2007). Effect of *Caralluma fimbriata* extract on appetite, food intake and anthropometry in adult Indian men and women. *Appetite*, 48(3), 338-344.
122. Lahsissene, H., Kahouadji, A., Tijane, M., & Hseini, S. (2009). Catalogue des plantes médicinales utilisées dans la région de Zaër (Maroc Occidental). *Lejeunia, revue de botanique*.
123. Lawrence, R. M., & Choudhary, S. (2004, December). *Caralluma fimbriata* in the treatment of obesity. In 12th Annual World Congress of Anti-Aging Medicine.
124. Lone, A. B., Bhat, H. F., Kumar, S., Aït-Kaddour, A., Aadil, R. M., Hassoun, A., & Bhat, Z. F. (2025). Cricket protein-based film containing *Caralluma fimbriata* extract-based nanoparticles for preservation of cheddar cheese. *Ultrasonics Sonochemistry*, 112, 107167.
125. Makeen, H. A., Menachery, S. J., Moni, S. S., Alqahtani, S. S., ur Rehman, Z., Alam, M. S., ... & Albratty, M. (2020). Documentation of bioactive principles of the exudate gel (EG) from the stem of *Caralluma retrospiciens* (Ehrenb) and in vitro antibacterial activity—Part A. *Arabian Journal of Chemistry*, 13(8), 6672-6681.
126. Malladi, S., Ratnakaram, V. N., Babu, K. S., & Sreenivasulu, M. (2018). Pharmacological review of *Caralluma* R. Br.: a potential herbal genus. *Asian Journal of Pharmaceutics*, 12(4), S1146.
127. Manzoor, M., Ayesha, A., & Durrani, M. J. (2013). Uses of fruits, vegetables and herbs for the treatment of diabetes by the people of Quetta city. *Science Technology and Development*, 32.
128. Mechchate, H., Es-safi, I., Bari, A., Grafov, A., & Bousta, D. (2020). Ethnobotanical survey about the management of diabetes with medicinal plants used by diabetic patients in region of Fez Meknes, Morocco. *Journal of ethnobotany research and applications*.
129. Meng, Q. Q., Tong, S. Y., Peng, X. R., Zhao, Y. Q., Li, Z. H., Chen, H. P., & Liu, J. K. (2023). Nine new pregnane glycosides from the cultivated medicinal plant *Marsdenia tenacissima*. *Molecules*, 28(6), 2705.
130. Meve, U., & Heneidak, S. (2005). A morphological, karyological and chemical study of the *Apteranthes* (*Caralluma*) *europaea* complex. *Botanical Journal of the Linnean Society*, 149(4), 419-432.
131. Meve, U., & Liedtke, S. (2004). Subtribal classification in the Asclepiadeae (Apocynaceae). *Taxon*, 53(1), 61-76.
132. Miliauskas, G., Venskutonis, P. R., & Van Beek, T. A. (2004). Screening of radical scavenging activity of some medicinal and aromatic plant extracts. *Food Chemistry*, 85, 231-237.
133. Mohamed, O. G., Shalabi, A. A., El Halawany, A. M., Tripathi, A., & Abdel-Sattar, E. (2024). Hexagonosides AF: Pregnane glycosides isolated from *Caralluma hexagona*. *Phytochemistry*, 217, 113903.
134. Mossa, J. S., Rafatullah, S., Galal, A. M., & Al-Yahya, M. A. (1995). Pharmacological studies of *Rhus retinorrhoea*. *International Journal of Pharmacognosy*, 33(3), 242-246.
135. Mostafa, A. A., Al-Askar, A. A., Almaary, K. S., Dawoud, T. M., Sholkamy, E. N., & Bakri, M. M. (2018). Antimicrobial activity of some plant extracts against bacterial strains causing food poisoning diseases. *Saudi journal of biological sciences*, 25(2), 361-366.
136. Mounika, P., Jyothi, M. V., Ramalingam, P., Reddy, V. J., & Anusha, K. (2016). Anti tubercular and anthelmintic activities of aqueous methanolic extract of *Caralluma attenuata*. *International Journal of Pharmaceutical Sciences and Research*, 7(11), 4561.
137. Mradu, G., Saumyakanti, S., Sohini, M., & Arup, M. (2012). HPLC profiles of standard phenolic compounds present in medicinal plants. *International Journal of Pharmacognosy and Phytochemical Research*, 4(3), 162-167.
138. Mudrikah, Y., Bibi, Y., Tabassum, S., Zahara, K., & Bashir, T. (2021). Ethnomedicinal and pharmacological properties of *Caralluma tuberculata* N. E. Brown – A review. *Pure and Applied Biology*, 10(4), 503-510.
139. Murugapandi, M., Elanchezhian, S. S., Oh, T. H., Ramasundaram, S., & Muniyappan, N. (2023). Pharmacological applications of metal nanoparticles derived from *Caralluma sarkariae* species. *Process Biochemistry*, 132, 166-179.
140. Murugapandi, M., Elanchezhian, S. S., Oh, T. H., Ramasundaram, S., & Muniyappan, N. (2024). An evaluation of the pharmacological responses of metal nanoparticles derived from aqueous extract of *Caralluma adscendens* R. Br. var. *Bicolor*. *Journal of Molecular Liquids*, 401, 124618.
141. Mustafa, G., Arif, R., Atta, A., Sharif, S., & Jamil, A. (2017). Bioactive compounds from medicinal plants and their importance in drug discovery in Pakistan. *Matrix Sci. Pharma*, 1(1), 17-26.

142. Naik, M. R., & Krishnamurthy, Y. L. (2012). Xerophyte *Caralluma stalagmiferavar. longipetala* (Asclepiadaceae): a new record to the flora of Karnataka, India. *Journal of Threatened Taxa*, 4(6), 2656-2659.
143. Najam-us-Saqib, Q., Qayyum, R., & Waheed, A. (2013). Acute and chronic toxicity study of ethyl acetate fraction of *Caralluma tuberculata* in mice. *International Journal of Pharmacology and Clinical Sciences*, 2(4).
144. Navaneethan, R. D., NCJ, P. L., Ramaiah, M., Ravindran, R., Chinnathambi, A., Alharbi, S. A., ... & Mohamedibrahim, P. K. M. (2024). *Caralluma pauciflora* based Ag-NPs activate ROS-induced apoptosis through down-regulation of AKT, mTOR and p13K signaling in human Gastric Cancer (AGS) cells. *Nanotechnology*, 35(19), 195102.
145. Park, E. S., Moon, W. S., Song, M. J., Kim, M. N., Chung, K. H., & Yoon, J. S. (2001). Antimicrobial activity of phenol and benzoic acid derivatives. *International biodeterioration & biodegradation*, 47(4), 209-214.
146. Patra, J. K., Das, G., Fraceto, L. F., Campos, E. V. R., Rodriguez-Torres, M. D. P., Acosta-Torres, L. S., ... & Shin, H. S. (2018). Nano based drug delivery systems: recent developments and future prospects. *Journal of nanobiotechnology*, 16(1), 71.
147. Noor, N., Din, G. M. U., Nadeem, M., Qureshi, T. M., Khalid, W., Nadeem, M. A., ... & Aqlan, F. (2024). Development and Nutritional Evaluation of Ready-to-Drink Beverage Using the Choongan (*Caralluma tuberculata* L.) Extract. *Journal of Food Biochemistry*, 2024(1), 8831525.
148. Noreen, S. (2017). A mini review on a *Caralluma tuberculata* NE Br. uncommon and wild succulents but having exciting pharmacological attributes. *Pure and Applied Biology (PAB)*, 6(2), 748-761.
149. Noreen, S., Hussain, I., Tariq, M. I., Iqbal, S., Batool, F., Ghumman, S. A., ... & Kausar, T. (2018). Influence of Extraction Scheme on the Antioxidant Potential of *Caralluma tuberculata*. *Notulae Scientia Biologicae*, 10(3), 340-347.
150. Nunes, P. X., Silva, S. F., Guedes, R. J., & Almeida, S. (2012). Biological oxidations and antioxidant activity of natural products. *Phytochemicals as nutraceuticals-Global Approaches to Their Role in Nutrition and Health*, 278.
151. Odendaal, A. Y., Deshmukh, N. S., Marx, T. K., Schauss, A. G., Endres, J. R., & Clewell, A. E. (2013). Safety assessment of a hydroethanolic extract of *Caralluma fimbriata*. *International journal of toxicology*, 32(5), 385-394.
152. Ouassou, H., Bouhrim, M., Kharchoufa, L., Imtara, H., Daoudi, N., Benoutman, A., Bencheikh, N., Ouahhoud, S., Elbouzidi, A., Bnouham, M., & Benali, T. (2021). *Caralluma europea*(Guss) N.E.Br.: A review on ethnomedicinal uses, phytochemistry, pharmacological activities, and toxicology. *Journal of Ethnopharmacology*, 273, 113769.
153. Ouassou, H., Zahidi, T., Bouknana, S., Bouhrim, M., Mekhfi, H., Ziyat, A., ... & Bnouham, M. (2018). Inhibition of  $\alpha$ -glucosidase, intestinal glucose absorption, and antidiabetic properties by *Caralluma europaea*. *Evidence-Based Complementary and Alternative Medicine*, 2018(1), 9589472.
154. Mali, K. K., Kokate, M. D., Ghorpade, V. S., Dias, R. J., & Mahajan, N. S. (2025). Exploring the antiarthritic potential of *Caralluma fimbriata*: Phytochemical screening and preliminary observations in a rat model. *Journal of Ethnopharmacology*, 120417.
155. Pachipala, G., Vemula, R., Reddy, P. V. B., Kalita, P., & Chadipiralla, K. (2022). Phytochemical screening and in vitro evaluation of antioxidant and DNA inhibition activity of *Caralluma bhupenderiana* Sarkaria. *Biomedicine*, 42(4), 726-733.
156. Parihar S (2016) *Caralluma edulis*: an endemic, edible, medicinal and threatened plant species of Indian Thar Desert. *Biotech Today Int J Biol Sci* 6(1):37-40. *Appetite*, 48(3), 338-344.
157. Patra, J. K., Das, G., Fraceto, L. F., Campos, E. V. R., Rodriguez-Torres, M. D. P., Acosta-Torres, L. S., ... & Shin, H. S. (2018). Nano based drug delivery systems: recent developments and future prospects. *Journal of nanobiotechnology*, 16(1), 71.
158. Periyannayagam, K., Kasirajan, B., Karthikeyan, V., & Kumuda, R. I. T. (2013). *Vitis vinifera* L.(Vitaceae) leaves towards antimutagenic and anti-proliferative activity in anticancer drug discovery. *Innovare J. Health Sci*, 1(3), 32-35
159. Perk, A. A., Shatynska-Mytsyk, I., Gerçek, Y. C., Boztaş, K., Yazgan, M., Fayyaz, S., & Farooqi, A. A. (2014). Rutin mediated targeting of signaling machinery in cancer cells. *Cancer cell international*, 14(1), 124.
160. Poodineh, J., Khazaei Feizabad, A., & Nakhaee, A. (2015). Antioxidant activities of *Caralluma tuberculata* on streptozotocin-induced diabetic rats. *Drug Development Research*, 76(1), 40-47.
161. Qanwil, T., Malik, A., Mushtaq, A., Rehman, M. M. F. U., & Gohar, U. F. (2025). Hypolipidemic and Vasoprotective Potential of *Caralluma edulis*: A Histological and Biochemical Study. *Jordan Journal of Pharmaceutical Sciences*, 18(1), 21-35.
162. Raees, M. A. (2018). A phytopharmacological review on an Arabian medicinal plant: *Caralluma flava* NE Br. *Int. J. Phytomed*, 10, 148-152.

163. Ramachandra, M. N., Karuppusamy, S., & Krishnamurthy, Y. L. (2014). Micropropagation of *Caralluma stalagmiferavar. longipetala*: A rare succulent medicinal plant from Karnataka, India. *African Journal of Biotechnology*, 13(35).
164. Ramalingam, S. V., Bakthavatchalam, S., Ramachandran, K., Gnanarani Soloman, V., Ajmal, A. K., Al-Sadoon, M. K., & Vinayagam, R. (2024). Potential antimicrobial and cytotoxic activity of *Caralluma indicaseed* extract. *Antibiotics*, 13(12), 1193.
165. Ramesh, M., Rao, Y. N., Kumar, M. R., Mohan, G. K., Kumar, B. R., Rao, A. V. N., ... & Reddy, B. M. (1999). Flavone glycoside from three *Caralluma* R. Br. species. *Biochemical systematics and ecology*, 27(1), 85-86.
166. Rauf, A., Jan, M., Rehman, W., & Muhammad, N. (2013). Phytochemical, phytotoxic and antioxidant profile of *Caralluma tuberculata* NE Brown. *Wudpecker Journal of Pharmacy and Pharmacology*, 2(2), 21-25.
167. Reddy, B. M., Byahatti, V. V., Rao, A. A., & Ramesh, M. (1996). Anti-inflammatory activity of *Stapelia nobilis* and *Caralluma stalagmifera*.
168. Reddy, K. D., Rao, B. V., Babu, G. S., Kumar, B. R., Braca, A., Vassallo, A., De Tommasi, N., Rao, G. V., & Rao, A. V. (2011). Pregnane glycosides from *Caralluma adscendensvar. fimbriata*. *Fitoterapia*, 82(7), 1039–1043.
169. Rehman, R. U., Chaudhary, M. F., Khawar, K. M., Lu, G., Mannan, A., & Zia, M. (2014). In vitro propagation of *Caralluma tuberculata* and evaluation of antioxidant potential. *Biologia*, 69(3), 341-349.
170. Ren, D., Zuo, R., González Barrios, A. F., Bedzyk, L. A., Eldridge, G. R., Pasmore, M. E., & Wood, T. K. (2005). Differential gene expression for investigation of *Escherichia coli* biofilm inhibition by plant extract ursolic acid. *Applied and Environmental Microbiology*, 71(7), 4022-4034.
171. RIZWANI, G. H. (1991). Phytochemical and biological studies on medicinal herbs, *Caralluma tuberculata* and *Caralluma edulis* (Doctoral dissertation, UNIVERSITY OF KARACHI, KARACHI).
172. Sánchez, E., Rivas Morales, C., Castillo, S., Leos-Rivas, C., García-Becerra, L., & Ortiz Martínez, D. M. (2016). Antibacterial and antibiofilm activity of methanolic plant extracts against nosocomial microorganisms. *Evidence-Based Complementary and Alternative Medicine*, 2016(1), 1572697.
173. Sanmugapriya, E., & Venkataraman, S. (2006). Toxicological investigations on *Strychnos potatorum* Linn seeds in experimental animal models. *Journal of health science*, 52(4), 339-343.
174. Sembiring, E. N., Elya, B., & Sauriasari, R. (2018). Phytochemical screening, total flavonoid and total phenolic content and antioxidant activity of different parts of *Caesalpinia bonduc* (L.) Roxb. *Pharmacognosy journal*, 10(1).
175. Shah, A., Marwat, S. K., Gohar, F., Khan, A., Bhatti, K. H., Amin, M., ... & Zafar, M. (2013). Ethnobotanical study of medicinal plants of semi-tribal area of Makerwal & Gulla Khel (lying between Khyber Pakhtunkhwa and Punjab Provinces), Pakistan.
176. Shahinuzzaman, M., Yaakob, Z., Anuar, F. H., Akhtar, P., Kadir, N. H. A., Hasan, A. M., ... & Akhtaruzzaman, M. (2020). In vitro antioxidant activity of *Ficus carica* L. latex from 18 different cultivars. *Scientific reports*, 10(1), 10852.
177. Shailemo, D. H., Kwaambwa, H. M., Kandawa-Schulz, M., & Msagati, T. A. (2016). Antibacterial activity of *Moringa ovalifolia* and *Moringa oleifera* methanol, N-hexane and water seeds and bark extracts against pathogens that are implicated in water borne diseases. *Green and Sustainable Chemistry*, 6(2), 71-77.
178. Shanmugam, G., Ayyavu, M., Rao, D. M., Devarajan, T., & Subramaniam, G. (2013). Hepatoprotective effect of *Caralluma* R. Br. umbellate against acetaminophen induced oxidative stress and liver damage in rat. *Journal of Pharmacy Research*, 6(3), 342-345.
179. Si, Y., Sha, X. S., Shi, L. L., Wei, H. Y., Jin, Y. X., Ma, G. X., & Zhang, J. (2022). Review on pregnane glycosides and their biological activities. *Phytochemistry Letters*, 47, 1-17.
180. Sofi, F. R., Raju, C. V., Lakshmisha, I. P., Singh, R. R., & All India Coordinated Research Project on Post Harvest Technology. (2016). Antioxidant and antimicrobial properties of grape and papaya seed extracts and their application on the preservation of Indian mackerel (*Rastrelliger kanagartha*) during ice storage. *Journal of Food Science and Technology*, 53(1), 104-117.
181. Sreenivasacharyulu, M. (1939). *Yogarathnakaram* (Vol. 2, p. 678) [in Telugu]. Nellore, India: Swatantra Press.
182. Stewart, R. R. (1972). An annotated catalogue of the vascular plants of West Pakistan and Kashmir.
183. Tackholm, V., & Boulos, L. (1974). Students' flora of Egypt.
184. Tareen, R. B., Bibi, T., Khan, M. A., Ahmad, M., Zafar, M., & Hina, S. (2010). Indigenous knowledge of folk medicine by the women of Kalat and Khuzdar regions of Balochistan, Pakistan. *Pak J Bot*, 42(3), 1465-1485.
185. Thunuguntla, V. B. S. C., Gadanec, L. K., McGrath, C., Griggs, J. L., Sinnayah, P., Apostolopoulos, V., ... & Mathai, M. L. (2024). *Caralluma fimbriata* Extract Improves Vascular Dysfunction in Obese Mice Fed a High-Fat Diet. *Nutrients*, 16(24), 4296.

186. Ugraiah, A., Sreelatha, V. R., Reddy, P. K., Rajasekhar, K., Rani, S. S., Karuppusamy, S., & Pullaiah, T. (2011). In vitro shoot multiplication and conservation of *Caralluma bhupenderiana* Sarkaria-an endangered medicinal plant from South India. *African Journal of Biotechnology*, 10(46), 9328-9336.
187. Ugwah-Oguejiofor, C. J., Alkali, Y. I., Inuwa, A. M., Pender, G. C., & Chindo, B. A. (2024). Studies on neurobehavioural properties of *Caralluma dalzielii* NE Br. aqueous aerial parts extract in mice. *Journal of Ethnopharmacology*, 324, 117774.
188. Umamaheswari, M., & Chatterjee, T. K. (2008). In vitro antioxidant activities of the fractions of *Coccinia grandis* L. leaf extract. *African Journal of Traditional, Complementary and Alternative Medicines*, 5(1), 61-73.
189. Umamaheswari, M., & Chatterjee, T. K. (2008). In vitro antioxidant activities of the fractions of *Coccinia grandis* L. leaf extract. *African Journal of Traditional, Complementary and Alternative Medicines*, 5(1), 61-73.
190. Vadivu, R. S., & Velavan, S. (2020). PHYTOCHEMICAL CHARACTERIZATION OF *CARALLUMA INDICA* STEM EXTRACT USING GC MS TECHNIQUE. *Journal of Advanced Scientific Research*, 11(02), 213-216.
191. Veerabhadraiah, T., Sabitha Rani, A., Prabhakar, G., & Keerthi, M. (2024). Phytochemical analysis of *Caralluma stalagmifera* C.E.C. Fisch, an endemic and important medicinal plant. *Journal of Pharmacognosy and Phytochemistry*, 13(3), 291-293.
192. Venkatesh, S., Reddy, G. D., Reddy, B. M., Ramesh, M., & Rao, A. A. (2003). Antihyperglycemic activity of *Caralluma attenuata*. *Fitoterapia*, 74(3), 274-279.
193. Venkatesh, S., Reddy, G. D., Reddy, B. M., Ramesh, M., & Rao, A. A. (2003). Antihyperglycemic activity of *Caralluma R. Br. attenuata*. *Fitoterapia*, 74(3), 274-279.
194. Venkatesh, S., Reddy, G. D., Reddy, B. M., Ramesh, M., & Rao, A. A. (2003). Antihyperglycemic activity of *Caralluma R. Br. attenuata*. *Fitoterapia*, 74(3), 274-279.
195. Waheed, A., Barker, J., Barton, S. J., Khan, G. M., Najm-us-Saqib, Q., Hussain, M., ... & Carew, M. A. (2011). Novel acylated steroidal glycosides from *Caralluma tuberculata* induce caspase-dependent apoptosis in cancer cells. *Journal of ethnopharmacology*, 137(3), 1189-1196.
196. Wariss, H. M., Mukhtar, M., Anjum, S., Bhatti, G. R., Pirzada, S. A., & Alam, K. (2013). Floristic composition of the plants of the Cholistan Desert, Pakistan. *American Journal of Plant Sciences*, 4(12), 58-65.
197. Wided, N. M., Robert, D. A., & Brian, S. C. (2021). Safe nanoparticles: *International Journal of Molecular Sciences*, 22(1), 385.
198. Wu, S. J., & Ng, L. T. (2008). Antioxidant and free radical scavenging activities of wild bitter melon (*Momordica charantia* Linn. var. *abbreviata* Ser.) in Taiwan. *LWT-Food Science and Technology*, 41(2), 323-330.
199. Zaib, B., Ahmad, S., Yaqub, A., & Ambreen, F. (2018). Effect of gamma irradiation on antioxidant activity of phytochemicals in selected medicinal plants. *Int. J. Eng. Res*, 9(6), 1530-4.
200. Zakaria, M. N. M., Islam, M. W., Radhakrishnan, R., Chen, H. B., Kamil, M., Al-Gifri, A. N., ... & Al-Attas, A. (2001). Anti-nociceptive and anti-inflammatory properties of *Caralluma arabica*. *Journal of Ethnopharmacology*, 76(2), 155-158.
201. Zarei, Z., Razmjoue, D., & Karimi, J. (2020). Green synthesis of silver nanoparticles from *Caralluma tuberculata* extract and its antibacterial activity. *Journal of Inorganic and Organometallic Polymers and Materials*, 30(11), 4606-4614.
202. Zari, T. A., & Al-Thebaiti, M. A. (2018). Effects of *Caralluma R. Br. russeliana* stem extract on some physiological parameters in streptozotocin-induced diabetic male rats. *Diabetes, metabolic syndrome and obesity: targets and therapy*, 619-631.
203. Zito, P., Guarino, S., Peri, E., Sajeve, M., & Colazza, S. (2013). Electrophysiological and behavioural responses of the housefly to "sweet" volatiles of the flowers of *Caralluma europea*(Guss.) NE Br. *Arthropod-Plant Interactions*, 7(5), 485-489.
204. Zito, P., Sajeve, M., Bruno, M., Maggio, A., Rosselli, S., Formisano, C., & Senatore, F. (2010). Essential oil composition of stems and fruits of *Caralluma europea* NE Br.(Apocynaceae). *Molecules*, 15(2), 627-638.