



Therapeutic Properties and Nutritive Values of Some Fruit Bearing Medicinal Plants of Rajasthan State in India

Gupta GK¹, Tejas Joshi¹, Keshu Madhudiya¹, Arijit Chaudhuri^{2*}

Department of Agriculture, Madhav University, Sirohi- 307 026, Rajasthan, India

Department of Pharmacy, Manav Bharti University, Solan-173229, Himachal Pradesh, India

Article Information

Received 10 August 2017

Received in revised form 16 Oct 2017

Accepted 18 Oct 2017

Keywords:

Medicinal Plant,
Rajasthan,
Therapeutic Properties,
Nutritive

Corresponding Author:

E-mail : armanchaudhuri92@gmail.com

Mob.: +919459481842

Abstract

The medicinal plants imparts chief role in protecting our health from various disease. It is nature's gift to human being to live healthy life. Medicinal plants are believed to be much safer and proved as elixir in the treatment of various ailments. Medicinal plants used in Indian system of medicine from Rajasthan state have been surveyed and categorized systematically. The manuscript incorporated the therapeutic properties and nutritive values of medicinal plants of Rajasthan. The paper deals with 11 medicinal plants, thoroughly indexed along with their important traditional application for the cure of various ailments. This study also incorporates the ethno-botany and biological activities of these important plants.

1 Introduction

Folk medicines today play a key role in the developing countries due to a lack of or limited modern health service. From ancient times, plants have been a rich source of effective and safe medicines. Due to their safe, effective and inexpensive nature, indigenous remedies are popular among the people of both urban and rural areas in China and India. Information from ethnic groups about indigenous traditional medicines has played a vital role in the discovery of novel products from plants for use as chemotherapeutic agents. Herbal medicines have been the main source of primary health care in many nations. About 80% of the world's populations are still dependent on traditional medicines^{1,2}.

Traditions are dynamic entities of unchanging knowledge. Traditional medicine is in an evolutionary process as communities and individuals continue to discover new techniques that can transform practices. Ethnopharmacology and drug discovery using natural products remain important issues in the current target-rich, lead-poor scenario. Many modern drugs have their origin in ethnopharmacology. However, despite technologic advances, the drug discovery process is facing a major innovation deficit that is adversely affecting the pharmaceutical industry^{3,4}.

Rajasthan is the largest state in the Northeastern part of India, geographically it lies between 23°3' to 30°12'N longitude and 69°30' to 78°17'S latitude and is rich in diversity of medicinal plants¹. Medicinal uses of different plants have been recorded in numerous literatures standing from the age of Vedas. In Rajasthan also a lot of work has been done on ethnomedicinal plants used for various ailments by different tribal communities and researchers. The present paper represent the data of the therapeutic properties and nutritive values of fruit bearing medicinal plants occurring in the Rajasthan state which may be used in future as plant resources for modern system of medicine.

2 Therapeutic properties and nutritive values of medicinal plants of Rajasthan

Balanites aegyptiaca

Balanites aegyptiaca is a member of Zygophyllaceae family, is also known as 'Desert date' in English, one of the most common but neglected wild plant species of the dry land areas of Africa and South Asia⁷. It is a slow growing, small, not very spreading, multibranched, evergreen, spiny and medium size tree up to 10 m tall, or shrub⁸. This tree is native to Africa and parts of the Middle East. In India, it is particularly found in Rajasthan, Gujarat, Madhya Pradesh, and Deccan⁹. It can be found in

many kinds of habitat, tolerating a wide variety of soil types, from sand to heavy clay, and climatic moisture levels. This plant has been reported to have anthelmintic, insecticidal, antidiabetic, antimicrobial, antibacterial, antifungal, hepatoprotective, anticancerous, antiparasitic, anti-inflammatory, molluscicidal and antioxidant properties¹⁰⁻¹².

Balanites aegyptiaca is traditionally used in treatment of various ailments i.e. jaundice, intestinal worm infection, wounds, malaria, syphilis, epilepsy, dysentery, constipation, diarrhea, hemorrhoid, stomach aches, skin boils, leucoderma, malaria, wounds, colds, syphilis, liver and spleen disorders, asthma, snake bite¹³ and fever. The bark of the plant is useful in curing mental diseases, yellow fever, jaundice and syphilis and can also act as a fumigant to heal circumcision wounds. Fruit kernel has been found as a mild laxative, an antidote to arrow poison, and also acts as a vermifuge. Kernel oil helps in curing skin disease. The seeds are useful as ointments, to cure cough, colic pain and also have magicoreligious properties^{14,15}.

Balanites aegyptiaca contains saponin, furanocoumarin, and flavonoid namely quercetin 3-glucoside, quercetin-3-rutinoside; 3-glucoside, 3-rutinoside, 3-7-diglucoside and 3-rhamnogalactoside of isorhamnetin¹⁶. Fruit contains protein, sugars, organic acids, other constituents like 3-rutinoside and 3-rhamnogalactoside, diosgenin. kernel contains a xylopyranosyl derivative of above saponin present in mesocarp. Balanitoside (furostanol glycoside) and 6-methyldiosgenin, balanitin-3 (spirostanol glycoside) have been reported from fruits (mesocarp) of *B. aegyptiaca*. The kernels also contained oil and protein, oil contains mainly palmitic, stearic, oleic, and linoleic acids which were the main fatty acids. The leaves and fruit kernels of *B. aegyptiaca* L. were found to contain six diosgenin glucosides including di-, tri-, and tetraglucosides^{16,17}.

Roots is reported to contain steroidal saponin about 1% glycosides and major sapogenin is yamogenin other glycosides. Bark is reported to contain furanocoumarin bergapten and dihydrofuranocoumarin D- marmesin, two alkaloid namely, N-trans-feruloyltyramine and N-cis-feruloyltyramine, and three common metabolites, vanillic acid, syringic acid; and 3-hydroxy-1-(4-hydroxy-3-methoxyphenyl)-1-propanone^{18,19}.

Calligonum polygonoides

Calligonum polygonoides is a member of family Polygonaceae. It is a small leafless shrub, which has a reputation in folklore medicine as a stimulant and astringent²⁰. *Calligonum polygonoides* is locally known as Phog, Phogala or Phogaro²¹. Usually it is seen as a small glabrous, winter shedding, perennial shrub 3-4 feet high with whitish and fragile branches. Sometimes a small tree with 12-15 feet in height and a trunk with 2-3 feet in girth²². It is found from arid and semi-arid areas of Thar desert in India. It grows on sand dunes as a

psammophytic vegetation of Barmer, Bikaner, Churu, Jaisalmer, Jhunjhunu, Nagaur, Sikar and Shri Ganganagar.

Leaves and stems are chewed to wash teeth and to treat gummosis while young shoots infusion is used as tonic²³. Paste of Root is applied on the affected areas for the treatment of Prickly heat and scabies. Decoction is used for the treatment of sore gums, typhoid. Flowers buds are effective in sun stroke. Flowers are also used for the treatment of asthma, eczema, cough and cold. It is reported that juice of the plant is applied in eyes to remove poisonous effect of *Calotropis procera*²⁴.

Calligonum polygonoides possesses hypoglycemic, cytotoxic, antioxidant, antimicrobial, anti-cancer, antiulcer, anti-inflammatory, antifungal, and mosquitocidal activities²⁵⁻²⁷.

Findings show that flavonoids, alkaloids, tannins, steroids, phenols, carbohydrates and terpenoids are present in different parts of *C. polygonoides*²⁸. Chemical analysis of *Calligonum polygonoides* revealed the presence of catechin, dehydrocatechin A, kaempferol-3-O-rhamnopyranoside, quercitrin, β -sitosterol-3-O-glucoside, isoquercitrin, kaempferol-3-O-glucuronide, and mequilianin. Campesterol, stigmaterol, (3 β , 5 α , 24 S)-stigmastan-3-ol, and stigmast-4-en-3-one were isolated from the roots of the plant, whereas β -sitosterol, kaempferol, quercetin, taxfolin, gallic acid, and astragalin were isolated from leaves^{29,30}.

Citrullus colocynthis

Citrullus colocynthis is perennial herbs usually trailing belongs to family Cucurbitaceae, commonly known as Chitrapala or Bitter apple. It is found wild in the sandy lands of North West, the Punjab, Sind, and Central and southern India, and coromandal coast. Also found indigenous in Arabia, West Asia, and Tropical Africa and in the Mediterranean region. *Citrullus colocynthis* is tender climbing monoecious plant with 2-3 tendrils. Leaves are deeply 3-5 lobate and both, the male & female flowers are yellow. Fruits are globular, variegated, dark green with yellowish blotches. When ripe, it is filled with a dry spongy very bitter pulp^{31,32}.

Citrullus colocynthis shows mild stomachic, bitter tonic, anthelmintic, anti-cancer, antioxidant, antimicrobial, antidiabetic, analgesic, antipyretic, anti-inflammatory, carminative, diuretic and anthelmintic property^{33,34}.

Citrullus colocynthis is used generally in the cure of various diseases such as leprosy, gut disorders, diabetes, constipation, asthma, indigestion, colic, rheumatism, hypertension, gastroenteritis, dysentery, bronchitis, jaundice, joint pain, cancer and mastitis^{35,36}.

Citrullus colocynthis contain various bioactive compounds such as alkaloids, flavonoids, carbohydrates, glycosides, fatty acids and essential oils. Cucurbitacins (Cucurbitane type triterpen glycoside viz colocynthoside A & B) have been documented as

the major constituent of *Citrullus colocynthis* fruits³⁷. Seeds contain Fatty acid like Stearic, Myristic, Palmitic, oleic, Linoleic, Linolenic acid, Protein 8.25 % and rich content rich in lysine, leucin and sulfo amino acid like methionine, Vitamin B1, B2 and Niacin, Mineral like Ca, Mg, K, Mn, Fe, P and Zn. Aerial part and fruit contain flavonoid glycoside quercetin, Flavone- 3-glucoside viz iso-vitexin, iso-orientine and isoorientine -3-methyl ether³⁸.

Commiphora wightii

Guggulu consists of oleo-gum resin obtained as an exudate from the tapping of stem and branches of *Commiphora wightii* (Arnott) Bhandari; Family, Burseraceae. The plant is commonly known as guggal, gugar, and Indian bdellium tree and is found in arid areas of India, Bangladesh, and Pakistan. In India, it is found in Rajasthan, Gujarat, Assam, Madhya Pradesh, and Karnataka. It is a small, bushy tree with thorny branches and produces a yellowish gum resin (guggulu) in small ducts located throughout its bark. The trees are tapped by making an incision on the bark. The resin, which flows out, is allowed to harden before it is collected. The tree is tapped from November to January and the resin is collected through May to June³⁹.

Guggulu possesses hypolipidemic, anti-inflammatory, anti-arthritic, antifertility, Anti-atherosclerotic, astringent, anti-septic, anti-inflammatory, analgesic, wound healing, anti-obesity, anti-spasmodic activity⁴⁰.

In Indian traditional system of medicine, guggulu has been used for thousands of years in the treatment of arthritis, inflammation, stimulates libido, nervous diseases, bronchial congestion, cardiac and circulatory problems, weak digestion, wounds, abscess, foetid ear, fractures, gout, skin rashes, irregular menstruation, diarrhea, headache, mild nausea, liver toxicity, rheumatism, obesity, and disorders of lipids metabolism⁴¹.

Guggulu contains guggultetrols, monoterpenoids, sesquiterpenoids, diterpenoids, triterpenoids, steroids, long-chain aliphatic tetrols, aliphatic esters, ferulates, lignans, flavonoids, sugar, amino acids carbohydrates, and a variety of inorganic ions besides minor amounts of sesamin and other unidentified constituents. Guggul gum is a mixture of 61% resins, 29.3% gum, 6.1% water, 0.6% volatile oil and 3.2% foreign matter. The important biologically active principles of C21 or C27 steroids are: guggulsterol-I, guggulsterol-II, guggulsterol-III, guggulsterol-IV, guggulsterol-V, E-guggulsterone, Z-guggulsterone and related ketones⁴².

Cordia myxa

Cordia myxa belongs to family Boraginaceae, is also known as clammy-cherry, glueberry, Indian-cherry in English and Gondi in Hindi. *Cordia myxa* is Dioecious shrub or small tree up to 12 m tall, straight, forming a dense crown, with very prominent leaf

scars. *Cordia myxa* originated from the area stretching from the eastern Mediterranean region to eastern India, and was introduced long ago in tropical Africa, tropical Asia and Australia, and more recently also in the Americas^{43,44}.

Pharmacological studies revealed that *Cordia myxa* possessed analgesic, anti-inflammatory, immunomodulatory, antimicrobial, antiparasitic, insecticidal, cardiovascular, respiratory, gastrointestinal and protective effects^{45,46}.

Cordia myxa was eaten to suppress cough and for the treatment of respiratory infections and a sore throat, as it has demulcent properties. The pulp was also applied as an emollient to mature abscesses, to calm rheumatic pain and as an anthelmintic. Fruit pulp is applied on ringworm. leaves were applied to wounds and ulcers. A macerate of the leaves was taken to treat trypanosomiasis, and is externally applied as a lotion to tse-tse fly bites. Bark powder was used externally in the treatment of skin diseases. Bark juice together with coconut oil was taken to treat colic^{47, 48}.

The preliminary phytochemical screening carried out on *Cordia myxa* fruit extract revealed the presence of oil, glycosides, flavonoids, sterols, saponins, terpenoids, alkaloids, phenolic acids, coumarins, tannins, resins, gums and mucilage. The seeds of *Cordia myxa* was consisted of palmitic acid, stearic acid, oleic acid and linolenic acid. The flavonoids and phenolic derivative content of the five species of genus *Cordia* leaves (*C. francisci*, *C. martinicensis*, *C. myxa*, *C. serratifolia* and *C. ulmifolia*). Four flavonoid glycosides, robinin, rutin, datiscoside and hesperidin, one flavonoid aglycone, dihydrorobinetin, two phenolic derivatives, chlorogenic and caffeic acid⁴⁹.

Gymnema Sylvestre

Gymnema sylvestre belongs to family Asclepiadaceae, is also known as 'gurmar' or 'sugar destroyer' (If the leaves of the plant are chewed, the sense of taste for sweet and bitter substances is suppressed)⁵⁰. It is a woody, climbing traditional medicinal herb which has many therapeutic applications in Ayurvedic system of medicine. *Gymnema sylvestre* is a slow growing, perennial, medicinal woody climber found in southern part of China, Tropical Africa, Vietnam, Malaysia, and Srilanka and is widely available in Japan, Germany, USA, central and peninsular India (mostly in Rajasthan, Bihar, West Bengal)⁵¹. The bioactive compounds of plant have antidiabetic, atherosclerotic, antimicrobial, antiarthritic, antibiotic, hypolipidaemic, immunostimulatory, hepatoprotective, anti-hyperglycemic, antipyretic, diuretic, anti-inflammatory, wound healing and anticancer properties⁵²⁻⁵⁴.

Gymnema sylvestre is a traditional medicinal plant, with reported use as a remedy for diabetes mellitus, stomachic and diuretic problems. Its use has been indicated in adenopathy, cough⁵⁰, asthma, alexipharmic, anthelmintic, astringent, biliousness, bronchosis, cardiopathy, conjunctivosis, cornea,

dysuria, digestive, emetic, expectorant, fever, furunculosis, glycosuria, hemorrhoid, hepatosplenomegaly, inflammation, jaundice, leukoderma, rheumatismopacities, ophthalmia, and worm⁵⁵. The roots of *Gymnema sylvestre* has also been used in snake bite, boil, constipation, and water retention, epilepsy, pain, high cholesterol, IDDM, NIDDM and obesity⁵⁶.

The plant is a good source of a large number of bioactive substances. Its constituents include two resins, gymnemic acids, saponins, stigmasterol, quercitol; the amino acid derivative of betaine, choline and trim ethylamine and Gymnemagenin and gymnestogenin. Gymnemic acids, a group of triterpenoid saponins belonging to oleanane and dammarene classes. Oleanane saponins are gymnemic acids and gymnemasaponins, while dammarene saponins are gymnemasides. Gymnemic acids I-VI and gymnemic acids XV-XVIII were also isolated. Gymnemic acids VIII-XII have been elucidated as glucosideuronic acid derivatives of gymnemagenin^{50, 57}.

Jatropha curcas

Jatropha curcas a multipurpose, shrub or tree, drought resistant, perennial plant belongs to family 'Euphorbiaceae', is widely distributed in the wild or semicultivated areas in Central and South America, Africa, India and South East Asia⁵⁸. *Jatropha* grows almost anywhere except waterlogged lands, even on gravelly, sandy and saline soils. The tree has a straight trunk and grey or reddish bark, masked by large white patches. Various parts of the plant are of medicinal value, its bark contains tannin, the flowers attract bees and thus the plant has a honey production potential. Its wood and fruit can be used for numerous purposes including fuel. It is easy to establish and grows relatively quickly⁵⁹.

The bioactive compounds of *Jatropha curcas* have antioxidant, antidiabetic, atherosclerotic, antimicrobial, antiarthritic, antibiotic, hypolipidaemic, immunostimulatory, hepatoprotective, anti-hyperglycemic, antipyretic, antifungal, diuretic, anti-inflammatory, wound healing, Nickel toxicity, and anticancer properties^{60,61}.

It is traditionally used in arthritis, gout, jaundice & as contraceptives, fish poison, toothache, gum inflammation, gum bleeding, pyorrhea, dermatomucosal diseases, allergies, burns, cuts and wounds, inflammation, leprosy, leucoderma, scabies and small pox, HIV, tumor and wound healing⁶².

Jatropha curcas have shown the presence of various bioactive constituents such as Amyrin, sitosterol, taraxerol, cyclic triterpenes (stigmasterol, stigmast-5-en-3, 7 diol, stigmast-5-en-3,7 diol, cholest-5-en-3,7 diol, campesterol, sitosterol, 7-keto-sitosterol as well as the d-glucoside of sitosterol), Flavonoids (apigenin, vitexin, isovitexin). Leaves contain the dimer of a triterpene alcohol and two flavonoidal glycosides. Latex contain curcacycline A, a cyclic octapeptide curcain. Seeds contain

curcain, a lectin Phorbolsters Esterases and Lipase, saponins and a trypsin inhibitor. Roots contain sitosterol and its d-glucoside, marmesin, propacin, the curculathyrans A and B and the curcusones A–D, diterpenoids jatrophol and jatropholone A and B, the coumarin tomentin, the coumarinolignan jatrophin as well as taraxerol^{63, 64}.

Leptadenia pyrotechnica

Leptadenia pyrotechnica is a typical desert plant belongs to the Asclepiadaceae family, commonly known as Khimp, Kheep or Khip. It is leafless, erect, ascending, shrub up to 0.5 meter to 2.6 meter high with green stem and pale green alternating bushy branches with watery sap erect and evergreen shrub⁶⁵. *Leptadenia pyrotechnica* occurs throughout the state of Rajasthan and found in dry habitats particularly in desert zones. In India it is commonly found in Punjab, Banswara, Palod, Dungarpur, Kota and Western Uttar Pradesh. It is native to Mediterranean regions, semi-arid deserts of African and Asian countries⁶⁶.

Leptadenia pyrotechnica possess significant antioxidant, anti-inflammatory, antibacterial, anthelmintic, antilipoxygenase, cytotoxic, antifungal, anticancer, wound healing, antidiabetic, hepatoprotective, antitumour, hypolipidemic and anti atherosclerotic activity. The fiber of *Leptadenia pyrotechnica* is used as antihistaminic and expectorant^{67,68}.

It is traditionally used in fever, cough, kidney disorders, stones, urinary disease. Fresh juice of the plant is used for abortion. Plant sap is applied to eczema and other skin disease and is also given in diabetes. The latex or the leaf paste is applied over the thorn injury for thorn removal. Whole plant infusion is mixed with buttermilk and given for uterine prolapsed and stomach disorders in sariska region of Rajasthan. It is used to cure constipation and is considered good for health in Bikaner region of Rajasthan⁶⁹.

Leptadenia pyrotechnica have shown the presence of bioactive constituents such as steroidal glycoside, cardiac glycosides, cardenolides, alkaloids, flavonoids, triterpenes tannins, saponins and poloxypregnane derivatives⁷⁰.

Salvadora oleoides

Salvadora oleoides is a small, multipurpose tree commonly known as jhal, badapilu, pilu, vridhpilu and khakan, belongs to family Salvadoraceae. The tree is primarily sourced for its fruits known as desert grapes. This species is decreasing very rapidly due to over exploitation, indiscriminate collection, low rate of seed set, poor viability and inefficiency of propagation by vegetative means^{71, 72}. *Salvadora oleoides* grows on dry, saline and desert areas of Rajasthan, Haryana, Andhra Pradesh, West Bengal, Karnataka, Tamil Nadu, Punjab, Gujarat, and Madhya Pradesh. The tree species is known to tolerate a very dry environment with mean rainfall of less than 200 mm in Barmer,

Jalore, Jodhpur and Pali districts of Rajasthan. It grows well in the sand dunes of deserts to heavy soils, non-saline to highly saline soils and dry regions to marshy semi-arid and waterlogged areas^{73,74}.

Salvadora oleoides possesses anti-oxidant, anti-inflammatory, analgesic, anti-ulcer, anthelmintic, diuretic, hypoglycemic, hypolipidemic, antimicrobial, larvicidal, cytotoxicity activities.

The leaves were used as a cooling agent, blood purifier, low fever, laxative, piles, chest disease, relief of abdominal pain, expectorant,. Leaf juice can also be used for anemic patients. Fruits are sweet with cooling effect and employed in the treatment of rheumatism, low fever, piles, tumor, bronchitis, child birth and snake bites⁷⁵. Seed oil is widely used in commercial production of cosmetics, paints, varnish, lubricants and as an ointment base for the treatments of rheumatism (Goodman, 1992). The whole plant is used as cooling herb, wound healing herb and nerve tonic, in the treatment of various uterine and skin disorders by the local people of Kachchh region⁷⁶.

Various qualitative chemical tests revealed the presence of carbohydrates, alkaloids, steroids, glycosides, saponins, tannins, triterpenes, mucilage, fats and oils in the leaf and stem extracts⁷⁷. Fruits contain glucose, fructose, sucrose and are good source of calcium. The seeds have flavonoids like quercetin, rutin and its seed fat contains lauric, capric, malic acid, myristic, palmitic, oleic, linolenic acid, dibenzyl urea and proteins. Salvadorin, a new dimeric dihydroisocoumarin has been isolated from the chloroform fraction of *S. oleoides*. Its chemical structure was established as 8-benzyl-6-[6-(6-ethyl-7-methyl-5, 8-dihydro-2- naphthalenyl)-1-oxo-3, 4-dihydro-1H-isochromen- 8yl]-3,4-dihydro-1H-isochromen-1-one, through spectroscopic techniques and chemical analysis⁷⁸. A new compound, heptadecanoyl-2-methyl-heptanoate has been isolated from stem bark.

Ziziphus mauritiana

Ziziphus mauritiana is a tropical fruit tree belongs to the family 'Rhamnaceae', locally known as 'Ber'. It is a spiny, evergreen shrub or small tree up to 15 m high, spreading crown and many drooping branches. The fruit is of variable shape and size. When slightly unripe, this fruit is a bit juicy and has a pleasant aroma. The fruit's skin is smooth, glossy, thin but tight. Fruits are eaten in other forms, such as dried, candied, pickled, as juice, or as ber butter. It is the most commonly found in the tropical and sub-tropical regions. Originally native to India it is now widely naturalized in tropical region from Africa to Afghanistan and China, and also through Malaysia , Australia and in some pacific regions⁷⁹.

Jubosides (saponin) isolated from *Ziziphus* reported to have haemolytic, sedative, anxiolytic and sweetness inhibiting properties. Whereas, cyclopeptide alkaloids found to have

sedative, antimicrobial, hypoglycemic, antiplasmodial, anti-infectious, antidiabetic, diuretic, analgesic, anticonvulsant and anti-inflammatory activities⁸⁰.

Plant pacifies vitiated pitta, kapha, obesity, fever, burning sensations, cough, wound, skin disease, ulcers, stomatitis, diarrhea, sexual weakness, and general debility. The dried fruits are used as anodyne, anticancer, pectoral, refrigerant, sedative, stomachache, styptic and tonic. They are considered to purify the blood and aid digestion. The root is used in the treatment of dyspepsia. A decoction of the root has been used in the treatment of fevers^{81,82}.

Z mauritiana having tremendous medicinal properties, attributed by a diverse group of secondary metabolites such as alkaloids, flavonoids, terpenoids, saponin, pectin, triterpenoic acids and lipids.

It is a rich source of cyclopeptide alkaloids lupane and triterpenes. It have 14-membered ring cyclopeptides to be the largest subgroup of alkaloid obtained, whereas only one 13-membered macrocyclic alkaloid isolated from this plant. It also contain protein ,carotene and vitamin C. Ripe fruits contains 20 to 30% sugar, up to 2.5% protein and 12.8% carbohydrates⁸³.

3 Conclusion

In Rajasthan, several plants were used for maintaining the health and treatment of several ailments. These plants possess anticancer, nephroprotective, hepatoprotective, antimicrobial, antioxidant, anti-diabetic, radio-protective, anti-HIV, anti-hepatoprotective, contraceptive etc. The medicinal plants displayed in this article required immediate steps to be taken for their conservation and sustainable utilization. Thus, there is a great need for manufacturing newer herbal drugs from these medicinal plants.

4 Conflict of interests

The authors have no current conflict of interests in this work.

5 Author's contributions

GKG, TJ, KM and AC carried out literature review and draft the manuscript. All authors read and approved the final manuscript.

6 References

1. Rao MM, Meena AK. Folk herbal medicines used by the Meena community in Rajasthan. *Asian Journal of Traditional Medicines*. 2010; 5(1): 19-31.
2. Joshi P, Awasthi A. Life support plant species used in famine by the tribals of Aravalli. *Journal of Phytological Research*. 1991; 42(2): 193-196.
3. Katkar KV, Suthar AC, Chauhan VS. The chemistry, pharmacologic, and therapeutic applications of *Polyalthia longifolia*. *Pharmacogn Rev*. 2010; 4(7): 62–68.

4. Patwardhan, Vaidya ABD, Chorghade M. Ayurveda and natural products drug discovery. *Curr Sci.* 2004; 86: 789–799.
5. Sebastian MK, Bhandari MM. Medicinal plant lore of Udaipur district, Rajasthan. *Bull Med Ethnobot Res.* 1984;122-124.
6. Sebastian MK, Bhandari MM. Medicinal plant lore of Udaipur district, Rajasthan. *Bull Med Ethnobot Res.* 1988; 133-134.
7. Hall JB, Waljer DH. School of Agricultural and Forest Science. Banger: University of Wales; 1991. *Balanites aegyptiaca* Del. A monograph; pp. 1–12.
8. Anon. The useful plants of India. New Delhi: Publications and Information Council of Scientific and Industrial Research; 1986.
9. *Balanites aegyptiacus* (L.) *Delile*°. Germplasm Resources Information Network. United States Department of Agriculture. 2008.
10. Chothani DL, Vaghasiya HU. A review on *Balanites aegyptiaca* Del (desert date): phytochemical constituents, traditional uses, and pharmacological activity. *Pharmacogn Rev.* 2011; 5(9): 55–62.
11. Yadav JP, Panghal M. *Balanites aegyptiaca* (L.) Del. (Hingot): A review of its traditional uses, phytochemistry and pharmacological properties. *International Journal of Green Pharmacy.* 2010; 140-146.
12. Khare CP. Indian medicinal plants: An illustrated dictionary. Springer. 2007:77–8.
13. Ojo OO, Nadro MS, Tella IO. Protection of rats by extracts of some common Nigerian trees against acetaminophen-induced hepatotoxicity. *Afr J Biotechnol* 2006;5:755-60.
14. Hamid O, Wahab M, Hassan E. *Balanites aegyptiaca* extract for treatment of HIV/ AIDS and leukemia. International Publication Number WO 2001/49306 A1.
15. Bukar A, Danfillo IS, Adeleke OA, Ogunbodede EO. Traditional oral health practices among Kanuri women of Brono state Nigeria. *Odontostomatol Trop.* 2004;27:25-31.
16. Samuelsson G, Farah MH, Claeson P, Hagos M, Thulin M, Hedberg O, Warfa AM, Hassan AO, Elmi AH, Abdurahman AD. Inventory of plants used in traditional medicine in Somalia. I. Plants of the families Acanthaceae-Chenopodiaceae. *J Ethnopharmacol.* 1991; 35(1):25-63.
17. Chothani DL, Vaghasiya HU. A Review on *Balanites Aegyptiaca* Del (Desert Date) Phytochemical Constituents, Traditional Uses, And Pharmacological Activity. *Pharmacogn Rev* 2011; 5(9): 55–62.
18. Seida AA, Kinghorn GA, Cordell GA. Isolation of bergapten and marmesin from *Balanites aegyptiaca*. *Plant Medica.* 1981;43:92–3.
19. Sarker SD, Bartholomew B, Nash R.J. Alkaloids from *Balanites aegyptiaca*. *Fitoterapia.* 2000;71:328–30.
20. Nawash OS, Al-Horani AS. The most important medicinal plants in Wadi Araba desert in South West Jordan: A review article. *Adv Environ Biol.* 2011;5:418–25.
21. Bhandari MM. Flora of the Indian Desert (Scientific Publishers) Jodhpur. 1978; 331-332.
22. Parker RN. Forest Flora for the Punjab with Hazara and Delhi. The superintendent Government printing Punjab Lahor. 1918; 412-422.
23. Shah A, Marwat SK, Gohar F, Khan A, Bhatti KH, Amin M. Ethnobotanical study of medicinal plants of semi-tribal area of Makerwal & Gulla Khel (lying between Khyber Pakhtunkhwa and Punjab Provinces). *Pakistan Am J Plant Sci.* 2013;4:98–116.
24. Khan A, Khan RA, Ahmed M, Mushtaq N. In Vitro antioxidant antifungal and cytotoxic activity of methanolic extract of *Calligonum polygonoides*. *Bangladesh Journal of Pharmacology.* 2015;10(2): 316-320.
25. Liu XM, Zakaria MN, Islam MW, Radhakrishnan R, Ismail A, Chen HB. Anti-inflammatory and anti-ulcer activity of *Calligonum comosum* in rats. *Fitoterapia.* 2001;72:487–91.
26. Al-Abraham JS, Mohammed AE, Elobeid MM. Assessment of *in vitro* anti-fungal potential of ethanolic extract of *Calligonum comosum* against two fungal postharvest pathogens of fruits and vegetables in Saudi Arabia. *IJABPT.* 2014;5:90–4.
27. El-Hag E, Harraz F, Zaytoon A, Salama A. Evaluation of some wild herb extracts for control of mosquitoes. *J King Saud Univ.* 1996;8:135–45.
28. Samejo MQ, Memon S, Bhangar MI, Khan KM. Preliminary phytochemicals screening of *Calligonum polygonoides* Linn. *J Pharm Res.* 2011; 4: 4402-03.
29. Badria FA, Ameen M, Akl MR. Evaluation of cytotoxic compounds from *Calligonum comosum* L. growing in Egypt. *Z Naturforsch C.* 2007;62:656–60.

30. Ahmed H, Moawad A, Owis A, Abou Zid S, Ahmed O. Flavonoids of *Calligonum polygonoides* and their cytotoxicity. *Pharmaceutical Biology*. 2016
31. Ebrahimie M, Bahmani M, Shirzad H, Rafieian-Kopaei M, Saki K. A review study on the effect of Iranian herbal medicines on opioid withdrawal syndrome. *J Evid Based Complementary Altern Med*. 2015; 20(4):302-9.
32. Enioutina EY, Salis ER, Job KM, Gubarev MI, Krepkova LV, Sherwin CM. Herbal Medicines: Challenges in the modern world. Part 5. status and current directions of complementary and alternative herbal medicine worldwide. *Expert Rev Clin Pharmacol*. 2017;10(3):327-38.
33. Gurudeeban S, Ramanathan T. Antidiabetic effect of *Citrullus colocynthis* in alloxon-induced diabetic rats. *Inventi Rapid: Ethno Pharmacology*. 2010;1:112.
34. Marzouk B, Marzouk Z, Fenina N, Bouraoui A, Aouni M. Anti-inflammatory and analgesic activities of *Citrullus colocynthis* Schrad. immature fruit and seed organic extracts. *Eur Rev Med Pharmacol Sci*. 2011;15(6):665-72.
35. Abo K, Fred-Jaiyesimi A, Jaiyesimi A. Ethnobotanical studies of medicinal plants used in the management of diabetes mellitus in South Western Nigeria. *Journal of Ethnopharmacology*. 2008;115(1):67-71.
36. Sultan A, Khan F, Iqbal H, Khan M, Khan I. Evaluation of chemical analysis profile of *Citrullus colocynthis* growing in southern areas of Khyber Pukhtunkhwa Pakistan. *World Applied Sciences Journal*. 2010; 10(4):402-5.
37. Wasylkowa K, Van der Veen M. An archaeobotanical contribution to the history of watermelon, *Citrullus lanatus* (Thunb.) Matsum. & Nakai (syn. *C. vulgaris* Schrad.). *Vegetation History and Archaeobotany*. 2004;13(4):213-7.
38. Gurudeeban S, Satyavani K and Ramanathan T. Bitter Apple (*Citrullus colocynthis*): An overview of chemical composition and biomedical potentials. *Asian J Plant Sci*. 2010; 1: 1-8.
39. The Ayurvedic Pharmacopoeia of India (Formulations), Department of Indian Systems of Medicine and Homeopathy, Ministry of Health and Family Welfare, Government of India, New Delhi, India, 1st edition, 2007.
40. Ernest Small. Frankincense and Myrrh – imperilled divine symbols of religion's duty to conserve biodiversity. *Biodiversity*. 2017; 1–16.
41. Urizar NL, Moore DD. Gugulipid: a natural cholesterol-lowering agent. *Annual Review of Nutrition*. 2003; 23: 303–313.
42. Wickens GE. *Ecophysiology of Economic Plants in Arid and Semi-Arid Lands*. Berlin, New York: Springer Science & Business media. 2013; 343.
43. USDA, ARS, National Genetic Resources Program. Germplasm Resources Information Network- (GRIN). National Germplasm Resources Laboratory, Beltsville, Maryland. 2015.
44. Oudhia P. *Cordia myxa* L. Record from PROTA4U. Schmelzer, G.H. & Gurib-Fakim, A. (Editors). *Plant Resources of Tropical Africa*, Wageningen, Netherlands 2007.
45. Al-Snafi AE. Therapeutic properties of medicinal plants: a review of plants with anti-inflammatory, antipyretic and analgesic activity (part 1). *Int J of Pharmacy* 2015; 5(3): 125-147.
46. Ali Esmail, Al-Snafi. The Pharmacological and therapeutic importance of *Cordia myxa*- A review. *IOSR Journal Of Pharmacy*. 2016; 6(6): 47-57.
47. Rechinger KH. *Cordia*. In: Rechinger KH (ed). *Flora Iranica*. Graz: Akademische Durck-u Verlangantalt 1997; 48:6.
48. Al-Awadi FM, Srikumar TS, Anim JT and Khan I. Antiinflammatory effects of *Cordia myxa* fruit on experimentally induced colitis in rats. *Nutrition* 2001;17(5):391-396.
49. Tiwari RD, Srivastava KC, Shukla S, Bajpai RK. Chemical examination of the fixed oil from the seeds of *Cordia myxa*. *Planta Med*. 1967; 15(3):240-244.
50. Kapoor LD. *CRC Handbook of Ayurvedic Medicinal Plants*; CRC Press: Boca Raton, FL, 1990; 200-201.
51. Ye WC, Zhang Q, Liu X, Che C, Zhao S. Oleanane saponins from *Gymnema sylvestre*. *Phytochemistry*. 2000; 53: 893-899.
52. Kishore L, Kaur N, Singh R. Role of *Gymnema sylvestre* as Alternative Medicine. *J Homeop Ayurv Med*. 2015; 3:172.
53. Tiwari P, Mishra BN, Sangwan NS. Phytochemical and Pharmacological Properties of *Gymnema sylvestre*: An Important Medicinal Plant. *BioMed Research International*. 2014.
54. Thakur G, Sharma R, Sanodiya BS, Pandey M, Prasad GBKS, Bisen PS. *Gymnema sylvestre*: An Alternative Therapeutic Agent for Management of

- Diabetes. Journal of Applied Pharmaceutical Science. 2012; 2(12): 1-6.
55. Kirtikar KR, Basu BD (1st ed., 1918, 2nd ed., 1935 or 1938), Indian Medicinal Plants, 4 volumes text, 4 volumes illustrations, M/S Periodical Experts, New Delhi, 1975.
 56. Bone K. Clinical Applications of Ayurvedic and Chinese Herbs — Monographs for the Western Herbal Practitioner, Phytotherapy Press, Warwick, Australia, 1996.
 57. Porchezian E, Dobriyal RM. An overview on the advances of *Gymnema sylvestre*: chemistry, pharmacology and patents. Pharmazie. 2003; 58: 5-12.
 58. Cano-Asseleih LM. Chemical investigation of *Jatropha curcas* L. seeds. Ph.D. Thesis. University of London. UK, 1986.
 59. Gubitz GM, Mittelbach M, Trabi M. Biofuels and Industrial Products from *Jatropha curcas*, DBV, Graz 1997; 65–69.
 60. Balaji R, Rekha N, Deecaraman M, Manikandan L. Antimetastatic and antiproliferative activity of methanolic fraction of *Jatropha curcas* against B16F10 melanoma induced lung metastasis in C57BL/6 mice. African Journal of Pharmacy and Pharmacology 2009; 3(11):547-555.
 61. Sachdeva K, Singhal M, Srivastava B. A review on chemical and medicobiological applications of *Jatropha curcas*. IRJP. 2 (4) 2011 61-66.
 62. Duke JA. CRC Handbook of Medicinal Herbs. CRC Press, Boca Raton, FL, 1988; 253–254.
 63. Goonasekera MM, Gunawardana VK. Pregnancy terminating effect of *Jatropha curcas* in rats. Journal of Ethnopharmacology. 1995; 47(3):117-123.
 64. Kannappan N, Jaikumar S, Manavalan R, Muthu AK. Antiulcer activity of methanolic extract of *Jatropha curcas* linn on aspirin induced gastric lesions in wistar rats. Pharmacologyonline newsletters. 2008; 1:279-293.
 65. Qureshi R, Munazir M, Abul-Soad AA, Jatoi MA, Shabbir G. *In vitro* callus induction protocol for *Leptadenia pyrotechnica* using various explants. J. Med. Plants Res. 2012;6:379–382.
 66. Rastogi RP, Mehrotra BN. Compendium of Indian Medicinal Plant, First edition, (1980-1984), vol. 3, Central Drug Research Institute, Lucknow and National Institute of science communication and information Resources, New Delhi, 2007.
 67. Saleh IA, Gamal AEHS, Amani SA, Abd ER, Mohammed D. Anti-inflammatory activity, safety and protective effects of *Leptadenia pyrotechnica*, haloxylon salicornicum and ochradenus baccatus in ulcerative colitis. Phytopharmacology. 2012; 2: 58-71.
 68. Rastogi RP, Mehrotra BN. Compendium of Indian Medicinal Plant. First edition, vol.-4, 2008, (1985-1989), Central Drug Research Institute, Lucknow and National Institute of science communication and information Resources, New Delhi. 430p.
 69. Al-Fatimi M, Wurster M, Schroder G, Lindequist U. Antioxidant, antimicrobial and cytotoxic activities of selected medicinal plants from Yemen. Journal of Ethnopharmacology. 2007; 111: 657-666.
 70. Upadhyay P, Roy B, Kumar SA. Traditional use of medicinal plants among rural communities of churu district in the Thar desert. Indian Journal of Ethnopharmacology. 2007; (113): 387-399.
 71. Kaul RN. Need for afforestation in the arid zones of India, La-yaaran. 1963; 13.
 72. Khan AU. History of decline and present status of natural tropical thorn forest in Punjab. Biol Conserv. 1994; 67: 205-210.
 73. Singh MN, Mishra AK, Bhatnagar SP. *In vitro* production of plants from cotyledon explants of *Cucumis melo* L. and their successful transfer to field. Phytomorph. 1996; 46: 395-402.
 74. Chopra RN, Nayar AN. Glossary of Indian Medicinal Plants, 1956, Edn 3, Part 1, CSIR, New Delhi, pp. 219.
 75. Hussein G, Miyashiro H, Nakamura N, Hattori, Kakiuchi N, Shimotono K. Inhibitory effects of Sudanese medicinal plant extracts on hepatitis C virus (HCV) protease. Phytther Res. 2000;14: 510-516.
 76. Goodman SM, Ghafoor A. The ethnobotany of Southern Baluchistan, Pakistan with particular reference to medicinal plants, Fieldiana: Botany New Series. 1992; 31: 84.
 77. Arora M, Siddiqui AA, Paliwal S, Mishra R. Pharmacognostical and phytochemical investigation of *Salvadora oleoides* Decne. Int J Pharm Pharmac Sci. 2013; 5(2): 62-66.
 78. Mahmood T, Ahmed E and Malik A. Structure determination of Salvadorin, a novel dimeric

- dihydroisocoumarin from *Salvadora oleoides* by NMR spectroscopy, Magn Reson Chem. 2005; 43(8): 670-672.
79. Mishra T, Kullar N, Bhatia A. Anticancer potential of Aqueous ethanol seed extract of *Ziziphus mauritiana* against cancer cell lines and Ehrlich *Ascites Carcinoma*. Evidence Based Complementary And Alternative medicines 2011;:2011:11.
 80. Bhatia A, Mishra T. Free radical scavenging activity and inhibitory responses of *ziziphus mauritiana* seed extract on alcohol induced oxidative stress. An international forum for Evidence Based Practices. 2009;(1): 8.
 81. Dahiru D, Sini JM. John Africa L Antidiarrhoeal activity of *Ziziphus mauritiana* root extract in rodents. African Journal of Biotechnology 2006; 5:10.
 82. Dahiru D, Obidoa O. Evaluation of the Antioxidant Effects of *Z.mauritiana* Lam. Leaf Extracts Against Chronic Ethanol Induced Hepatotoxicity in Rat. Liver Afr J Tradit Complement Altern Med. 2007; 5(1): 39-45.
 83. Bhatia A, Mishra T. Hypoglycemic activity of *Ziziphus mauritiana* aqueous ethanol seed extract in alloxan induced diabetic mice. Pharmaceutical biology. 2010; 48,604.