

A pharmacological appraisal of medicinal plants with antidiabetic potential

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ABSTRACT

Diabetes mellitus is a complicated metabolic disorder that has gravely troubled the human health and quality of life. Conventional agents are being used to control diabetes along with lifestyle management. However, they are not entirely effective and no one has ever been reported to have fully recovered from diabetes. Numerous medicinal plants have been used for the management of diabetes mellitus in various traditional systems of medicine worldwide as they are a great source of biological constituents and many of them are known to be effective against diabetes. Medicinal plants with antihyperglycemic activities are being more desired, owing to lesser side-effects and low cost. This review focuses on the various plants that have been reported to be effective in diabetes. A record of various medicinal plants with their established antidiabetic and other health benefits has been reported. These include Allium sativa, Eugenia jambolana, Panax ginseng, Gymnema sylvestre, Momrodica charantia, Ocimum sanctum, Phyllanthus amarus, Pterocarpus marsupium, Trigonella foenum graecum and Tinospora cordifolia. All of them have shown a certain degree of antidiabetic activity by different mechanisms of action.

KEY WORDS: Antioxidant, diabetes mellitus, hypoglycemic, medicinal plants

iabetes mellitus is a global health crisis, which has been persistently affecting the humanity, irrespective of the socioeconomic profile and geographic location of the population. According to an estimate, one person is detected with diabetes every 5 s somewhere in the world, while someone dies of it every 10 s.^[1] Diabetes mellitus has attained a pandemic form. Hence, it is very important to control diabetes and its complications to alleviate the human suffering. Scientists are desperately trying to manage this crippling disorder. Because plants are of enormous medicinal importance, they are being extensively explored for their use against diabetes. Herbal drugs can be quite acceptable as these drugs are known to cause less adverse effects.^[2] They are quite popular in developing countries.^[3] The increased admiration of herbal medicines for diabetes may be due to the side-effects associated with

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the conventional antidiabetic drugs.^[4] The World Health Organization (WHO) has also substantiated the utilization of herbal remedies for the management of diabetes.^[5] Till date, numerous medicinal plants have been reported to be effective in diabetes, yet plenty of research is still needed to be done. This article focuses on the various plants that could be effective in the treatment of diabetes mellitus.

Prevalence of Diabetes Around the World

Diabetes is a metabolic disorder critically afflicting the population of both developed and developing countries. According to the Diabetes Atlas, the global prevalence of diabetes is estimated to be 4.6%, representing 151 million people, and is expected to go up to 333 million people by 2025. Recent reports have estimated an increase in these figures, with the global prevalence reaching up to 6.6%, representing 285 million people in 2010 and by 2030, it will rise up globally up to 7.8% (438 million people). Also, individual national prevalence rates from over 1% to almost 31% have been reported, severely affecting the developing countries and, more specifically, the lower socioeconomic groups. Diabetes is rapidly emerging as a major public health challenge. In 2010, diabetes will share 11.6% of the entire international healthcare expenses, much of which

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will comprise hospital admissions and medications. According to the International Diabetes Federation (IDF), the overall cost assessment for the global prevention and treatment of diabetes will run up to US\$490 billion by 2030.[1] The occurrence of diabetes is higher in men than in women, but more women are reported to be suffering from diabetes than men. A notable increase in the proportion of people suffering from diabetes with >65 years of age is also reported. [6] Studies have indicated a shocking rise in the prevalence of diabetes in India. [2] According to the WHO, India had 31.7 million diabetic subjects in the year 2000, and this number would increase up to 79.4 million by the year 2030.^[6] Currently, India has the highest number of diabetic patients, and India is being called the diabetic capital of the world.[7] Studies have shown a significant age-related prevalence in the urban population, largely among the people with sedentary life style.[8]

Pathophysiology of Diabetes Mellitus

Diabetes mellitus is divided into two main types: type I (insulin -dependent diabetes mellitus or IDDM) and type II (non -insulin -dependent diabetes mellitus or NIDDM). IDDM occurs due to insulin insufficiency because the body does not generate any insulin and patients entirely depend on an exogenous supply of insulin. IDDM is more pronounced in children and young adults. It causes severe damage to the pancreatic β-cells. It is categorized as autoimmune (immune mediated) diabetes (type 1A) or idiopathic diabetes with β -cell destruction (type 1B), although the precise description of the later is still unknown. [9] Patients suffering from NIDDM are unable to respond to insulin and can be treated with exercise, diet management and medication. Mostly, its onset is in adulthood, largely occurring in obese people over 40 years of age. NIDDM is the most widespread type. It indicates a condition with disturbed carbohydrate and fat metabolism. Hypertension, hyperlipidemia, hyperinsulinemia and atherosclerosis are often allied with diabetes. Both the types demonstrate some frequent symptoms like high blood sugar levels, unusual thirst, extreme hunger, frequent urination, extreme weakness, blurred vision etc. Although the pathophysiology of diabetes is not entirely understood, many studies indicate the participation of free radicals in the pathogenesis of diabetes^[10] and its complications.^[11-13] Free radicals are proficient enough of damaging cellular molecules, proteins, lipids and DNA, leading to alternation of cell functions. In fact, the abnormalities in lipids and proteins are one of the key reasons for the development of diabetic complications. During diabetes, free radicals oxidize the lipoproteins, and various irregularities of lipoprotein metabolism also occur in very low-density lipoprotein (VLDL), low-density lipoprotein (LDL) and high-density lipoprotein (HDL) in diabetes. [2] Different extracellular proteins are also modified into glycoprotein due to high blood glucose, which is associated with severe diabetic complications.^[14] Reactive oxygen species (ROS) are being reported to be formed in different tissues in diabetes^[15,16] by various sources such as the nonenzymatic glycosylation reaction, [17] the electron transport chain in mitochondria[18] and membrane -bound NADPH oxidase. [19,20] ROS are also involved in the progression of insulin resistance as well as pancreatic β-cell dysfunction.^[21] Also, advanced glycation end products (AGEs) are produced by non -enzymatic glycosylation of proteins, which tends to mount up on long-lived molecules in tissues creating abnormalities in cell and tissue functions.[22,23] AGEs also play a role in improved vascular permeability in both micro- and macro-vascular structures by sticking to specific macrophage receptors, which leads to free radical production and endothelial dysfunction. AGEs, produced on nucleic acids, may also lead to altered gene expression and mutation. In diabetes, oxidative stress coexists along with decrease in the antioxidant status, which can lead to the detrimental effects due to free radicals. [24] Vitamins C and E, the natural antioxidants, have been reported to decrease the oxidative stress in experimental diabetes. [25] Numerous plant products have been reported to have a significant antioxidant activity, which may be of some benefit in diabetes. [26,27]

Conventional Treatment

Diabetes is a multidimensional disorder and its management needs firm adherence to the prescribed treatment plan. The contemporary treatment of diabetes is focused on suppressing and controlling blood glucose to a normal level. The common agreement on management of type II diabetes is transformation in lifestyle along with appropriate diet and weight control. However, antidiabetic drugs are needed as these measures cannot provide satisfactory results. Antidiabetic drug therapy includes insulin injections and oral hypoglycemic drugs. These drugs act by various mechanisms to control the blood glucose level. However, many side-effects such as hypoglycemia, lactic acid intoxication and gastrointestinal upset, etc. have been reported in patients.^[28] Because the antidiabetic medication may sometimes involve prescribing more than one drug at the same time, which can augment the severity of these sideeffects, efforts are being made to find a suitable antidiabetic and antioxidant therapy.

Medicinal Plants with Antidiabetic Activity and other Beneficial Effects

There are various herbal antidiabetic remedies used in various traditional systems of medicine prevailing around the world, although only some of them have been scientifically assessed for their efficacy. A list of the various medicinal plants with their antidiabetic and associated useful effects is given in Table 1.

Aegle marmelos (Bengal quince*, Bel†): Family - Rutaceae

The leaf extract of Aegle marmelos (A. marmelos), when given to alloxanized rats, has shown to reduce the blood sugar, urea, liver glycogen and serum cholesterol and also improve digestion. Along with this, it has also checked the peak rise in blood sugar in the oral glucose tolerance test (OGTT). [150,174] Reduction in oxidative stress along with decreased blood glucose level has also been reported with the methanolic extract in alloxanized rats as apparent from the significant diminution in lipid peroxidation along with restored antioxidant enzyme

Table 1: Medicinal plants with antidiabetic and other medicinal properties

Plant name	Traditional/common name	Part used	Antidiabetic effect and other important effects	Reference
<i>Acacia arabica</i> (Leguminosae)	Guar gum*, Gum Arabic*	Seeds	Hypoglycemic, stimulate pancreatic β -cells for insulin release	[29]
Acorus calamus (Araceae)	Sweet flag* or Calamus*	Radix	Hypoglycemic, increased glucose consumption, decrease in triglyceride, free fatty acid, insulin sensitizer	[30,31]
Aloe vera (Liliaceae)	True or medicinal aloe*	Aloe gel, leaves	Hypoglycemic, hypolipidemic, antioxidant, increases glucose tolerance, stimulation of synthesis and/or release of insulin from pancreatic β -cells, enhanced insulin transport, anti-inflammatory	[32-37]
Anethum graveolens (Umbelliferae)	Dill*	Leaves	Hypoglycemic, antioxidant, decrease in serum insulin level	[38]
<i>Annona squamosa</i> (Annonaceae)	Sugar apple*	Leaves	Hypoglycemic, antihyperglycemic, antioxidant, increased plasma insulin level	[39-42]
Areca catechu (Palmae)	Betel nut*, supari†	Fruit	Hypoglycemic	[43]
Astragalus membranaceus (Leguminosae)	Yellow leader*	Root	Hypoglycemic, hypolipidemic, decrease in insulin resistance, potential insulin sensitizer, decreased expression of protein tyrosine phosphatase 1B (PTP1B)	[44-49]
Averrhoa bilimbi (Oxalidaceae)	Bilimbi [†] , cucumber tree* or tree sorrel*	Leaves	Hypoglycemic, hypolipidemic	[50, 51]
Azadirachta indica (Meliaceae)	Neem [†]	Leaves, seeds	Antihyperglycemic, antioxidant, antibacterial, antimalarial, antifertility, hepatoprotective	[52,53-56]
Baccharis trimera	-	Aerial parts	Hypoglycemic	[57]
(Asteraceae)		of plant		
Barleria lupulina (Acanthaceae)	-	Aerial parts	Antihyperglycemic	[58,59]
Bauhinia forficata (Fabaceae)	Pata de Vaca [†]	Leaves	Hypoglycemic, antioxidant effect, useful in preventing diabetic complications	[60,61]
Berberis aristata (Berberidaceae)	Barberries* or pepperidge bushes*	Root	Antihyperglycemic, antioxidant	[62]
Beta vulgaris (Chenopodiaceae)	Chukkander [†] , beetroot* or garden beet*	Root	Increase in glucose tolerance	[63]
Bixa orellana	Achiote [†] , aploppas [†]	Leaves	Concentration-independent inhibition of human pancreatic	[64]
(Bixaceae) Brassica juncea	Mustard greens*, indian	Leaves	α-amylase Hypoglycemic, antioxidant	[65,66]
(Brassicaceae)	mustard*, chinese mustard* and leaf mustard*		Hypogrycernic, antioxidant	[05,00]
Butea monosperma (Papilionaceae)	Palasa [†] , flame of the forest*	Bark, leaves, flower	Hypoglycemic, antihyperglycemic, antioxidant, thyroid inhibitory effect	[67-70]
Camellia sinensis (Theaceae)	Tea*	Leaves	Anti-hyperglycemic, antioxidant	[71,72]
Capparis spinosa (Capparaceae)	Caper bush*	Fruit	Antihyperglycemic, no change in basal insulin level	[73]
Carum carvi (Apiaceae)	Caraway*, meridian fennel*	Fruit	Antihyperglycemic, no change in basal insulin level	[73]
Casearia esculenta (Salicaceae)	-	Root	Antihyperglycemic, antihyperlipidemic, antioxidant	[74,75]
Cassia auriculata		Leaves, flowers	Antihyperglycemic and hypolipidemic activity, α -glucosidase	[76,77]
(Caesalpiniaceae) Cassia glauca	avaram senna* -	Leaves, bark	inhibitory activity Antihyperglycemic, antihyperlipidemic	[78]
(Caesalpiniaceae) Cinnamomum	Pseudocinnamomum* or	Leaves	Enhanced adinopectin secretion and activation of insulin-signaling	[79]
osmophloeum (Lauraceae) Cinnamomum verum	indigenous cinnamon* True cinnamon*, ceylon	Leaves	pathway Concentration-dependant inhibition of human pancreatic	[64]
(Lauraceae) Citrullus vulgaris	cinnamon* Water melon*	Peel of fruit	α-amylase Hypoglycemic, antihyperlipidemic, amelioration of thyroid	[80]
(Cucurbitaceae) Citrus sinensis	Sweet orange*	Peel of fruit	dysfunction, inhibit lipid peroxidation Antihyperglycemic, antiperoxidative, antithyroid, insulin	[81-84]
(Rutaceae)	-		stimulating property, hypolipidemic, cardioprotective	
Citrus paradisi (Rutaceae) Cochlosparmum vitifolium	Grapefruit*	Seeds	Hypoglycemic, hypolipidemic, decrease in cardiovascular risk factors	[85]
Cochlospermum vitifolium (Cochlospermaceae)		Bark	Hypoglycemic, vasorelaxant and hepatoprotective properties	[86]
Combretum micranthum (Combretaceae)	Kinkeliba [†]	Leaves	Antihyperglycemic	[87]
Coscinium fenestratum (Menispermaceae)	-	Stem	Antidiabetic , hypolipidemic	[88]
Costus igneus (Costaceae)	Fiery costus* or spiral flag*	Leaves	Hypoglycemia	[89]

Table 1 (Contd...)

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Plant name	Traditional/common name	Part used	Antidiabetic effect and other important effects	Reference
Costus speciosus (Costaceae)	Crape ginger*	Root	Anti-hyperglycemic, antihyperlipemic and antioxidative effects	[90]
<i>Curcuma longa</i> (Zingiberaceae)	Turmeric*	Rhizome	Concentration-dependant inhibition of human pancreatic α -amylase	[64]
Dendrobium chrysotoxum (Orchidaceae)	Golden-bow dendrobium* or fried-egg orchid*	Stem	Antioxidant and anti -hyperglycemic	[91]
Dioscorea polygonoides (Dioscoreaceae)	Jamaican bitter yam*	Root	Hypoglycemic, hypolipidemic	[92]
Emblica officinalis (Euphorbiaceae)	Amla†, dhatriphala†	Fruit	Hypoglycemic, antioxidant, decrease in lipid peroxidation, antibacterial, antiulcerogenic, hepatoprotective, gastroprotective	[93,94-96]
Ficus bengalensis	Bur [†] , banyan*	Aerial root,	Hypoglycemic, antioxidant, concentration dependant inhibition of	[64,97,98]
(Moraceae)	0 " "	stem bark	human pancreatic α-amylase	500 3033
Ficus carica (Moraceae)	Common fig*	Leaves	Hypoglycemic, hypolipidemic	[99-101]
	Cluster fig tree* or goolar [†]	Bark	Strong hypoglycemic effect	[102]
<i>Globularia alypum</i> (Plantaginaceae)	-	Leaves	Antihyperglycemic, no effect on insulin secretion	[103]
<i>Gymnema montanum</i> (Asclepiadaceae)	-	Leaves	Antihyperglycemic, antihyperlipidemic, decrease in lipid peroxides, inhibition of apoptotic pathway	[104,105]
Gynostemma pentaphyllum (Cucurbitaceae)	Jiaogulan [†]	Leaves	Alternation of activities of glucose metabolic enzymes	[106]
<i>Hedychium spicatum</i> (Zingiberaceae)	Spiked ginger lily*, sandharlika [†]	Rhizome	Antihyperglycemic, inhibition of intestinal $\alpha\text{-glucosidase}$ activity	[107]
Helichrysum plicatum (Asteraceae)	-	Flower	Antihyperglycemic, antioxidant	[108]
Helicteres isora (Sterculiaceae)	-	Root, fruit, bark	Hypoglycemic, antihyperlipidemic, insulin sensitizer	[109-111]
Hygrophila auriculata (Acanthaceae)	Marsh barbel*	Aerial parts of plant	Hypoglycemic, antioxidant, decreased lipid peroxidation	[112]
Hypoxis hemerocallidea (Hypoxidaceae)	African star grass*, african potato*		Anti-inflammatory, antineoplastic, antioxidant, antidiabetic and anti-infective properties, stimulate insulin release	[113-117]
Indigofera mysorensis (Fabaceae)	-	Whole plant	Antihyperglycemic, potential insulin sensitizer	[118]
Juniperus chinensis (Cupressaceae)	Chinese juniper*	Fruit	Antihyperglycemic, hypolipidemic	[119]
Lagerstroemia speciosa (Lythraceae)	Giant crape-myrtle*, queen's crape-myrtle*, banabá plant for philippines*, or pride of India*		lpha-glucosidase inhibitor	[120]
<i>Linum usitatisumum</i> (Linaceae)	Flax* (also known as common flax* or linseed*)	Seeds	Inhibition of pancreatic amylase activity	[121]
Liriope spicata (Ruscaceae) Matricaria chamomilla	Creeping lilyturf* German chamomile*	Tuberous roots Aerial part	Hypoglycemic and hypolipidemic Antihyperglycemic, antioxidant, protection of β -cells	[122,123] [124]
(Asteraceae)		of plant	, , , , , , , , , , , , , , , , , , , ,	
<i>Momordica cymabalaria</i> (Cucurbitaceae)	Kadavanchi [†]	Fruit	Hypoglycemic, hypolipidemic	[125]
Morus alba (Moraceae)	Mulberry*, white mulberry*	Leaves	Inhibition of α -glucosidase, sucrose, maltase, inhibit porcine pancreatic α -amylase activity and reduced starch hydrolysis	[121,126]
Morus indica (Moraceae)	-	Leaves	Hypoglycemia	[89]
<i>Murraya koenigii</i> (Rutaceae)	Curry patta [†] , curry tree*	Leaves	Antihyperglycemic, enhances glycogenesis and reduce glycosylated Hb, gluconeogenesis, urea, uric acid, creatinine and glycogenolysis, concentration-independent inhibition of human	[64,127,128]
			pancreatic α -amylase	
Musa sapietum (Musaceae)		Flower	Antihyperglycemic, antioxidant	[129-131]
Olea europaea (Oleaceae)	Olive tree*	Leaves	Hypoglycemic, antioxidant, antihypertensive, anti-inflammatory, antiatherogenic, hypocholesterolemic	[132]
<i>Paeonia lactiflora</i> (Paeoniaceae)	Chinese peony*	Roots	Inhibition of PTP1B	[133]
Paspalum scrobiculatum (Poaceae)	or kodra millet*	Grains	Antihyperglycemic, increase in serum insulin level, hypolipidemic	[134]
<i>Persea americana</i> (Lauraceae)	Avocado* or alligator pear*	Leaves	Hypoglycemic, hypocholesterolemic	[135]
<i>Phaseolus vulgaris</i> (Fabaceae)	Hulga [†] , white kidney bean*, common bean*	Seeds	Hypoglycemic, hypolipidemic, inhibition of α -amylase activity, antioxidant, alternation in level of insulin receptor and GLU -4 mRNA in skeletal muscle, enhanced production of adinopectin, decreased TNF- α level	[136,137- 140]

Table 1 (Contd...)

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Plant name	Traditional/common name	Part used	Antidiabetic effect and other important effects	Reference
Pistacia atlantica (Anacardiaceae)	Mt. Atlas mastic tree* and as the Persian turpentine tree*	-	In vitro inhibition of $\alpha\text{-amylase}$ and $\alpha\text{-glucosidase}$ activity, acute postprandial antihyperglycemic	[141]
Polygonatum odoratum (Ruscaceae)	Angular solomon's -seal* or scented solomon's -seal*	Roots	Insulin sensitizers	[142]
Punica granatum (Lythraceae)	Anar [†]	Flowers, peel of fruit	Antioxidant, anti-hyperglycemic effect, inhibit TNF- α -stimulated free fatty acid release and attenuated TNF- α inhibition of adinopectin secretion, prevent diabetes via binding with PPAR - γ receptor, production of NO, inhibition of lipid peroxidation, inhibition of α -glucosidase activity	[81,136, 143-146]
Raphanus sativus (Brassicaceae)	Japanese radish*	Root, sprout	Hypoglycemic, improved lipid metabolism	[147-149]
Retama raetam (Fabaceae)	-	Fruit	Antihyperglycemic, inhibition of intestinal glucose absorption, stimulate pancreatic insulin release	[150,151]
Rhodiola sachalinensis (Crassulaceae)	-	Root	Hypoglycemic, hypolipidemic, antioxidant, increase in serum insulin level	[152,153]
Rubus fructicosis (Rosaceae)	-	-	Hypoglycemic, antihyperglycemic, no effect on insulin secretion	[103]
Sclerocarya birrea (Anacardiaceae)	Murula [†]	Stem bark	Hypoglycemic, increased plasma insulin level, prevented body weight loss, hypocholesterolemic, renoprotective, cardioprotective, antidiarrhoeal, antidiabetic, anti-inflammatory, antimicrobial, antiplasmodial, antihypertensive, anticonvulsant, antinociceptive and antioxidant properties	[154-157]
Sutherlandia frutescens (Fabaceae)	Cancer bush*, balloon pea*	Shoot	Hypoglycemic, anti -inflammatory, restores insulin levels and reduces intestinal glucose uptake	[158,159]
Swietenia macrophylla (Meliaceae)	Big leaf* Mahogany*, big-leaf mahogany*	Seeds	Hypoglycemic, hypolipidemic	[160]
Swietenia mahagoni (Meliaceae)	West Indian mahogany*	Bark	Antidiabetic, antioxidant	[161]
Syagrus romanzoffiana (Arecaceae)	Queen palm* or cocos palm*	Seed	Inhibition of $\alpha-glucosidase, reduce post-prandial blood glucose level$	[162]
Syzygium alternifolium (Myrtaceae)	Shahajire [†]	Seeds	Hypoglycemic and antihyperglycemic	[163]
Syzygium cumini (Myrtaceae)	Jambul [†]	Seeds	Concentration-dependant inhibition of human pancreatic $\alpha\text{-amylase}$	[64]
Terminalia catappa (Combretaceae)	-	Fruit	Hypoglycemia	[164]
Tridax procumbens (Asteraceae)	Dhaman grass*	Whole plant	Antihyperglycemic, hypoglycemic, correction in body weight	[165]
Vaccinium angustifolium (Ericaceae)	Canadian low bush blueberry*	Root, stem, leaf, fruit	Enhanced glucose transport, increase in glucose stimulated insulin release, reduced apoptosis, enhanced β -cells proliferation	[166]
Vaccinium bracteatum (Ericaceae)	-	Leaves	Potential hypoglycemic effect, body weight recovery	[167]
Vaccinium vitis -idaea (Ericaceae)	-	Berry	Potent antiglycating action, prevent insulin resistance and other metabolic diseases	[168,169]
Vernonia amygdalina (Asteraceae)	Bitter leaf*	-	Hypoglycemic and antihyperglycemic	[170,171]
Withania somnifera (Solanaceae)	Ashvagandha [†] , winter cherry*	Root	Hypoglycemic, diuretic and hypocholesterolemic, antioxidant, suppresses metformin-induced hypothyroidism	[172,173]

^{*}English common name, †Regional name

levels. [175] Treatment of streptozotocin-diabetic rats with the leaf extract of A. *marmelos* demonstrated superior functional state of pancreatic β -cells as it facilitates in the regeneration of damaged pancreas. [176] Further investigation has revealed the presence of various important chemical constituents such as scopoletin and umbelliferone, which provides additional therapeutic benefits (in hyperthyroidism and collagen-mediated diabetic nephropathy). [177,178]

Allium cepa (Onion*): Family - Amaryllidaceae

The ether extract of Allium cepa (A. cepa) has shown

antihyperglycemic activity in diabetic rabbits. [179] Also, a 1 ml solution (0.4 g A. cepa/rat) when given to streptozotocin -diabetic rats causes an augmentation in fasting serum diabetic HDL levels, demonstrating alleviation of hyperglycemia along with considerable antioxidant activity. [180] Moreover, when diabetic patients were administered a single oral dose of onion juice, it markedly controlled the post-prandial glucose levels. [181] A. cepa is also reported to have a hypolipidemic effect. Administration of S-methyl cysteine sulfoxide (SCMS), a sulfur-containing amino acid from onion, significantly controlled the levels of blood glucose as well as lipids and

also normalized the activities of glucose 6 -phophatase and 3 -hydroxy -3 -methylglutaryl coenzyme -A (HMG Co -A) reductase in alloxanized rats. [182] A new compound, (S(S) R(C)) -S -(3 -pentenyl) -L -cysteine sulfoxide, which is obtained from the seed extract of A. cepa var. tropeana has also been reported to contain antioxidant properties. [183]

Allium sativa (Garlic*): Family - Amaryllidaceae

When the aqueous extract of garlic [Figure 1] is given orally to sucrose-fed rabbits, it considerably improved hepatic glycogen and free amino acid content, reduced fasting blood glucose and triglyceride levels in serum.[184] Also, garlic extract administered to streptozotocin-diabetic rats not only decreased the blood glucose level but also inhibited the lipid peroxidation and inhibited the superoxide formation. This study has also recommended its long-term use in preventing diabetic complications. However, the extrapolation of these results to humans needs further research. [52] Most recent findings have also proposed that aged garlic extract inhibits the generation of glycation -derived free radicals and ACEs in vitro. S -allyl cysteine, [Figure 1] a chief ingredient of aged garlic, is a potent antioxidant that can inhibit AGEs synthesis and, thus, deserves more attention. [185] Evidence also proposes that the antioxidative, anti-inflammatory and antiglycative properties of garlic are accountable for its role in preventing diabetes and its complications. [186] Allicin, [Figure 1] a sulfur -containing compound isolated from garlic, is the reason for its pungent odor and also has considerable hypoglycemic action, [187] which is thought to be due to the augmented hepatic metabolism, insulin release and/or insulin-sparing effect. [188] Researchers have also reported that allicin contains considerable antioxidant^[189] and antimalarial activities.^[190] S -allyl cysteine sulfoxide (SACS), the precursor of allicin and garlic oils, is reported to control lipid peroxidation better than glibenclamide and insulin and also stimulated in vitro insulin release from the β-cells isolated from normal rats. [191] Ajoene, [Figure 1] obtained from garlic, has been reported to show antithrombotic, anti-tumor, antifungal and antiparasitic properties, [192] thus making garlic a useful plant.

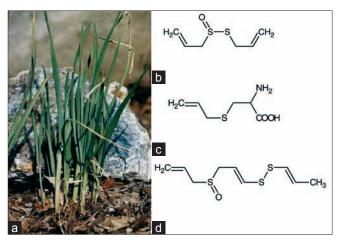


Figure 1: Garlic and its antidiabetic constitutents. (a) Allium sativa (garlic), (b) allicin, (c) S-allyl cysteine, (d) ajoene

Caesalpinia bonducella (Gray Nicker*): Family - Fabaceae

The aqueous and ethanolic extracts of *Caesalpinia bonducella* (*C. bonducella*) demonstrated potent hypoglycemic activity in chronic type II diabetes. These extracts also improved glycogenesis, thereby escalating the liver glycogen content. [193] Aqueous and 50% ethanolic extracts of *C. bonducella* seeds have also shown antihyperglycemic and hypolipidemic activities in streptozotocin -diabetic rats. [194] Oral administration of seed extracts to alloxanized rats not only produced considerable lowering of the blood urea nitrogen levels but it also lowered the elevated cholesterol as well as LDL levels, confirming that the drug also has the potential to show antidiabetic as well as antihyperlipidemic effects. This antihyperglycemic action may be due to obstruction in absorption of glucose. [195]

Capparis deciduas (Kerda[†], Kair[†], Karir[†]): Family - Capparaceae

Hypoglycemic effect was reported in alloxanized rats fed with the fruit powder of the *Capparis decidua* (*C. decidua*) plant. The powdered extract also significantly decreased the alloxan -induced lipid peroxidation in erythrocytes, kidney and heart. *C. decidua* is also reported to alter antioxidant enzyme levels and decreases oxidative stress. [196] The antidiabetic potential may be due to the alkaloids present in it. When the alkaloid-rich fraction of *C. decidua* plant was given to streptozotocin -diabetic rats, it demonstrated promising results, establishing its claim for further purification and categorization of the individual alkaloids. [197]

Coccinia indica (Ivy Gourd*, Little Gourd*): Family - Cucurbitaceae

The combined extract of Coccinia indica (C. indica) with Musa paradisiaca has shown a defensive effect against diabetes through β-cell regeneration in streptozotocin -diabetic albino rats. [198] Also, an aqueous -methanol extract of C. indica along with Musa paradisiaca, Tamarindus indica and Eugenia jambolana provided considerable defense against testicular dysfunction along with lowering of blood glucose. [199] In a study on diabetic patients, dried extracts of C. indica, when administered, helped to restore the normal activities of lipoprotein lipase, glucose -6 -phosphatase and lactate dehydrogenase, [200] while in a double -blind, placebo -controlled, randomized trial on type 2 diabetic patients, the alcoholic extract of the herb significantly decreased the fasting and postprandial blood glucose levels in the experimental group as compared with the placebo group. However, no considerable alterations in the serum lipid levels were observed.[201]

Eriobotrya japonica (Loquat[†]): Family - Rosaceae

Eriobotrya japonica (E. japonica) has been reported to show a distinct hypoglycemic action in normal and alloxanized rabbits and mice. [202-204] Studies have revealed that 300 mg/kg of the leaf extract in streptozotocin-diabetic mice induced significant decrease in plasma glucose concentration, glycosylated serum

protein, total cholesterol, triglycerides and oxidative stress. [205] The leaf extracts of *E. japonica* are also known to inhibit 11 β-hydroxy steroid dehydrogenase (HSD) type 1, preferentially over 11β-HSD2, which might add to the antidiabetic effect of the plant. [206] Further, studies demonstrated that sesquiterpene glycosides and polyhydroxylated triterpenoids are the active ingredients of *E. japonica* accountable for controlling diabetes mellitus. [207] Also, the total triperpene acid fraction from the leaves demonstrated a high anti-diabetic potential beside hypolipidemic and antioxidant profile in alloxan and streptozotocin-diabetic mice. [208]

Eugenia jambolana (Jambul†): Family - Myrtaceae

Decoction of Eugenia jambolana (E. jambolana) seed kernel is used as a domestic preparation for diabetes and it also forms a key ingredient of several antidiabetic herbal formulations. The pulp extract of Eugenia demonstrated hypoglycemic activity promptly than the seed extract in streptozotocindiabetic mice. Increased serum insulin levels and inhibition of insulinase activity from liver and kidney was observed on oral administration of the extract in diabetic rats. [209] Ethanolic extract of Eugenia seed kernel also established its antioxidant potential along with hypoglycemic effect in streptozotocindiabetic rats. [210] Combination treatment of lower dose of glimepiride together with ethanolic Eugenia seed extract showed potent hypoglycemic as well as antihyperglycemic activities without stern hypoglycemia in normal rats, concluding its possible use in considerable dose reduction of standard drugs.^[211] However, examination of the Brazilian Eugenia fruit has uncovered no beneficial effect of the plant extract in diabetic rats. [212]

Ginseng species: - Family - Araliaceae

Ginseng root has been used since long because of its medicinal properties. Medicinally important ginseng species include Panax ginseng (Asian ginseng*) [Figure 2] and Panax quinquefolius (American ginseng*). The main chemical constituents of the entire ginseng species are ginsenosides, polysaccharides, peptides, polyacetylenic alcohol and fatty acids, with ginsenosides being the most important pharmacological constituent. [213] Ginsenosides have been the object of countless researches as they are believed to be the key principles behind the efficacy and potential effects of ginseng. Therefore, it is also imperative to assess its worth using various analytical techniques.[214] Animal study data indicate that both Asian ginseng^[215,216] and American ginseng^[217,218] have a prominent hypoglycemic effect, which may be due to ginsenoside Rb -2 [Figure 2] and, more specifically, because of panaxans I, J, K and L in type 1 diabetic models. [219-222] American ginseng has also been reported to inhibit the tumor necrosis factor -alfa (TNF $-\alpha$)-stimulated free fatty acid release and also attenuated the TNFα -inhibition of adiponectin secretion. [136] Hypoglycemic activity of ginseng has also been established by clinical studies.[223,224] However, a few adverse effects of ginseng have been observed, most common of which are nervousness and excitation, which are diminished with continued use or dosage

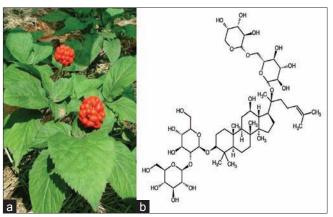


Figure 2: Ginseng and its antidiabetic constituent. (a) Panax ginseng (Asian ginseng), (b) Ginsenoside Rb2

reduction. Massive overdose of drug can lead to ginseng abuse syndrome. [225] The suggested dosage is 1 - 3 g of the plant's crude root daily or 200 - 600 mg of standardized extract. [226]

Gymnema sylvestre (Gymnema*, Australian cow plant*): Family - Asclepiadaceae

Gymnema sylvestre (G. sylvestre) has been used in the treatment of diabetes since ages. Assessment of the alcoholic extract of G. sylvestre on insulin secretion from the islets of langerhans and several pancreatic β-cell lines of rats has uncovered that the extract stimulated insulin release from several β-cells and islets, due to increased cell permeability.[227] The ethanolic extract of Gymnema displayed hypoglycemic and antihyperglycemic activity in rats when given alone or in combination with any standard drug, and could be used for dose reduction of the standard drug.[211] Recently, an active compound, dihydroxy gymnemic triacetate, has been isolated from its acetone extract. When it was given orally to streptozotocin-diabetic rats, it produced considerable hypoglycemia along with hypolipidemic effects. [228] In a study on type 2 diabetic subjects, the G. sylvestre extract was given daily to 22 patients along with oral hypoglycemic drugs, which improved blood sugar control. Five patients were able to retain blood sugar control with Gymnema extract without the help of oral medication. [229] One of its side-effects may be reduction/loss of taste sensation. The recommended dosage of G. sylvestre extract is 400 - 600 mg/day.[230]

Mangifera indica (Mango*): Family - Anacardiaceae

Although no change in blood glucose level was reported when the aqueous mango extract was given to streptozotocindiabetic rats, antidiabetic activity was observed when the aqueous extract was given before/along with glucose, which could be due to decreased intestinal glucose absorption. [231] Analysis of the peel extract of *Mangifera indica* (*M. indica*) [Figure 3] in rats revealed their potential to ameliorate diet-induced change in serum lipids, thyroid dysfunctions and hyperglycemia, which could be due to polyphenols and ascorbic acid present in the test peel extract. [80] Lupeol,

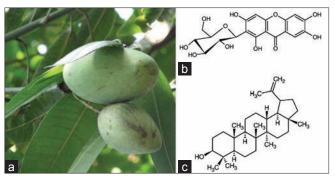


Figure 3: Mango and its antidiabetic constituents. (a) Mangifera indica (mango), (b) mangiferin, (c) lupeol

[Figure 3] a triterpene found in mango, is acknowledged to display antioxidant, antilithiatic and antidiabetic effects and is also established to be valuable in combating oxidative stress-induced cellular injury of mouse liver by modulating cell -growth regulators.^[232] Also, mangiferin, [Figure 3] a xanthone glucoside, isolated from mango leaves possesses considerable antidiabetic, antihyperlipidemic and antiatherogenic properties as obvious from lowering of fasting glucose level, decrease in total cholesterol, triglycerides and LDL-cholesterol along with elevation of the HDL-cholesterol level and diminution of atherogenic index in diabetic rats.^[233]

Momordica charantia (Bitter gourd*, Karela†): Family - Cucurbitaceae

It is commonly used as an antidiabetic agent. Extracts from various parts of the plant have shown hypoglycemic activity in different animal models. Ethanolic extracts of Momordica charantia (M. charantia) showed hypoglycemic and antihyperglycemic effect in normal and streptozotocin-diabetic rats, which could be due to inhibition of glucose -6 -phosphatase and stimulation of hepatic glucose - 6 -phosphate dehydrogenase activity. [234] Studies concerning mechanism of bitter gourd in alloxanized rats suggested its potential antidiabetic, antihyperlipidemic and other valuable effects in amelioration of diabetic complications. Furthermore, it restores the altered histological architecture of the islets of Langerhans.^[235] The latest finding on high -fat -fed rats treated with bitter gourd extract showed better insulin sensitivity, glucose tolerance and insulin signaling. Detection of the possible mechanism of these effects may unlock novel therapeutic targets for the management of obesity/dyslipidemia -induced insulin resistance. [236] Aqueous seed extract of Momordica also provides notable protection against lipid peroxidation in streptozotocindiabetic rats due to its antioxidant activity. [237,238] Antidiabetic screening of Momordica on differentiating 3T3 -L1 adipocytes has shown inhibition of TNF- α -stimulated free fatty acid release and attenuation of TNF-α -induced inhibition of adiponectin secretion.^[136] Animal studies have suggested many parallels between the actions of metformin and bitter gourd. [239] Recent extensive screening has identified triterpenoids to be the hypoglycemic components present in Momordica that may be responsible for insulin resistance and in activation of AMP -activated protein kinase. [240] Polypeptide p, isolated from the fruit, seeds and tissues of *M. charantia* showed significant hypoglycemic effect in human.^[241] Because diet is one of the approaches in the management of diabetes mellitus, there is scope for exploiting the antidiabetic potency of Momordica to the maximum extent.^[242] However, there are insufficient evidences to recommend it for type 2 diabetes mellitus due to a lack of significant data on morbidity, expected expenditure, etc.^[243] Traditionally, it is used to cure numerous disorders. A number of studies have authenticated its use in diabetes and its complications as antibacterial, antiviral, anticancer, abortifacient, etc. However, only a few reports showing positive results on its clinical use are available.^[244]

Ocimum sanctum (Holy basil*, Tulsi†): Family -Lamiaceae

Ocimum sanctum (O. sanctum) is known for its therapeutic benefits since ancient times. The aqueous extract of O. sanctum leaves demonstrated considerable decrease in blood sugar level in both normal and alloxanized rats. [245] Marked fall in fasting blood glucose along with decrease in uronic acid, total cholesterol, triglyceride and total lipid level point toward its health-benefitting effects. [246] Administration of the ethanolic leaf extract has been reported to lessen the plasma glucose level along with increase in the renal glycogen content, while skeletal muscle and hepatic glycogen levels are decreased in streptozotocin-diabetic rats. [247] This plant is also known to possess antimicrobial, adaptogenic, hepatoprotective, anti-inflammatory, anti-carcinogenic, neuroprotective, cardio-protective, mosquito repellent and numerous other therapeutic activities. [52,248]

Phyllanthus amarus (Stonebreaker*, Seed-Under-Leaf*): Family - Phyllanthaceae

Traditionally, *Phyllanthus amarus* (*P. amarus*) is used in the treatment of diabetes. The methanolic extract of *P. amarus* was found to reduce the blood sugar in alloxanized diabetic rats along with potential antioxidant activity.^[249] Complete inhibition of TNF-α -stimulated free fatty acid release and attenuation of TNF-α induced inhibition of adiponectin secretion was observed while screening its antidiabetic effects on 3T3 -L1 adipocytes.^[136] *P. amarus* is a potential diuretic and hypotensive drug for humans.^[250] Interestingly, other plants of the Phyllanthus species also possess medicinal properties.^[93,251,253]

Psidium guajava (Common guava*): Family - Myrtaceae

Various studies revealed that various parts of *Psidium guajava* have antidiabetic property. The aqueous guava leaf extract showed hypoglycemic activity in alloxanized and streptozotocin-diabetic rats, which is attributed to the various tannins, flavonoids and other chemical constituents of the plant.^[254,255] In a different study, anti -hyperglycemic activity of the ethanol extract of stem bark of the plant was assessed on normal, alloxanized and normal glucose-loaded rats, and the results showed that the extract displayed considerable

hypoglycemic activity in all except normal and glucose-loaded rats. $^{[256]}$ Antidiabetic property was also observed when the butanol-soluble fraction of the leaves was given to Lepr db/ Lepr db mice. $^{[257]}$ The leaf extract of guava was shown to inhibit the α -glucosidase activity in the small intestine of diabetic mice. $^{[258]}$ Considerable hypolipidemic activity has also been reported when the aqueous extract of the raw fruit peel was given to streptozotocin-diabetic rats, besides hypoglycemic and antidiabetic activity. $^{[259]}$

Pterocarpus marsupium (Indian kino tree*): Family - Fabaceae

Pterostilbene, obtained from *Pterocarpus marsupium* (*P. marsupium*) wood, showed hypoglycemic activity in dogs. [260] Flavonoids obtained from *P. marsupium* have been reported to cause pancreatic β -cell regranulation. [261] The flavanoid fraction has also showed antihyperlipidemic activity. [262] Also, the aqueous extract of the latex of *P. marsupium* was found to possess marked α -glucosidase inhibitory activity. [76] One of the active principles of Pterocarpus, (-) epicatechin, has insulinogenic action. It stimulate oxygen uptake in fat cells and tissue slices of different organs, and also enhances the glycogen content of the rat diaphragm in a dose-dependent manner. [263]

Trigonella foenum graecum (Fenugreek*): Family - Fabaceae

Fenugreek seeds are frequently used as a constituent of spices. Oral administration of the plant extract produced lowering of the blood glucose levels in both normal as well as diabetic rats. [264] Administration of fenugreek seeds also enhanced glucose metabolism and stabilized creatinine kinase activity in heart, skeletal muscle and liver of diabetic rats. [265] Fenugreek plant has also been reported to show antioxidant activity. [266,267] Further, daily oral treatment of its steroids to diabetic rats demonstrated a significant decrease of blood glucose level and a substantial enhancement of the area of insulin -immunoreactive β -cells along with considerable reduction in sperm shape abnormality and improved sperm count. [268] 4-hydroxyisoleucine, isolated from fenugreek seeds, exhibits marked potential as an anti -diabetic agent by suppressing progression of type II diabetes in the db/db mice model, as apparent from improvement of insulin sensitivity and glucose uptake in peripheral tissue. [269] When fenugreek oil is given to alloxanized rats, notable reduction in renal toxicity besides improved hematological status and antidiabetic effect has also been accounted, which could be due to the immunomodulatory activity and insulin stimulation action of fenugreek. [270] Fenugreek seeds are also known to possess a hypolipidemic effect^[271] and also offer antilithogenic potential due to its encouraging effect on cholesterol metabolism.^[272] Its seed has also been shown to be effective in the prevention of retinopathy and other diabetic complications when used alone or in combination with sodium orthovandate. [273,274] The seed extract also considerably repressed the 7, 12 -dimethylbenz (α) anthracene (DMBA) -induced mammary hyperplasia and reduced its incidence. [275] It has shown no genotoxic effect, and has a wide safety margin. Also, adding the fenugreek seed extract to foodstuffs for diabetic patients is predicted to be safe. [276]

Tinospora cordifolia (Guduchi[†]): Family - Menispermiaceae

Tinospora cordifolia (T. cordifolia) is extensively employed for treating diabetes mellitus in the traditional system of medicine. [277-279] Oral administration of the root extract of T. cordifolia led to a considerable decrease in blood and urine glucose and in lipids in alloxanized rats and also prevented reduction in body weight. [280] Also, the aqueous root extract of the plant resulted in considerable decrease in blood glucose and brain lipids in alloxanized rats. Although the extract showed significant antihyperglycemic effect, its effect was comparable only to 1 unit/kg of insulin. [281] It is also accounted that the daily administration of the aqueous or alcoholic extract of T. cordifolia reduces the blood glucose level and enhances glucose tolerance in rodents. [282]

Stevia rebaudina (Sweet Leaf*, Sugar leaf*): Family - Asteraceae

Stevioside, [Figure 4] a natural sweetener obtained from the plant Stevia rebaudiana (S. rebaudiana), [Figure 4] has been used in the treatment of diabetes since a long time. It stimulates secretion of insulin via a direct action on the pancreatic cells, and is thought to have significant antidiabetic potential. [283,284] However, a comparative study between S. rebaudiana and stevioside has revealed that the hypoglycemia due to S. rebaudiana leaves was partially mediated by inhibition of hepatic gluconeogenesis and did not involve stevioside and peroxisome proliferator -activated receptor -gamma (PPAR -γ) receptors activation. [285] Rebaudioside, [Figure 4] a new diterpene glycoside, also possesses insulinotropic effects and may provide support in the treatment in type 2 diabetes mellitus.[286] A study of stevioside on insulin -sensitive lean (Fa/-) and insulin -resistant obese (Fa/Fa) Zucker rats revealed the skeletal muscle glucose transport system to be one of its possible sites of action. [287] Stevioside is also reported to exercise antihyperglycemic, insulinotropic, blood pressure lowering and glucagonostatic effects in type 2 diabetic Goto - kakizaki

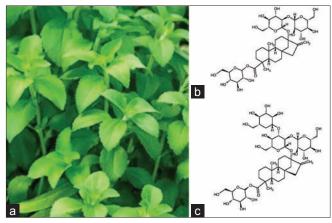


Figure 4: Stevia and its antidiabetic constituents. (a) Stevia rebaudiana (sweet leaf), (b) stevioside, (c) rebaudioside

rats, proving its worth in the treatment of type 2 diabetes and metabolic syndrome. [284,288] Stevioside and steviol may have a potential as an antihyperglycemic agent. While investigating their effects on insulin release from normal mouse islets and the β -cell line INS -1, it came to appear that the insulinotropic effects of stevioside and steviol were significantly reliant on the prevailing glucose concentration, and that they also stimulate insulin secretion by acting on pancreatic β -cells. [289]

Conclusion

The number of people suffering from diabetes mellitus has been increasing dramatically over the past few decades, and this demands special attention towards its management. The few conventional therapies available are either expensive or often related with adverse effects; therefore, various traditional therapies with antihyperglycemic effect are increasingly sought by patients. Medicinal plants provide better alternatives as they are generally less-toxic and affordable; yet, their safety and efficacy needs more evaluation by controlled clinical studies. Although herbs are less likely to have drawbacks of the conventional drugs used for diabetes, potential herb-drug interactions should be kept in mind for those receiving conventional antidiabetic medications. Taking all these details into account, further research is required to validate the antidiabetic effects of medicinal plants.

References

- Colagiuri R. Diabetes: A pandemic, a development issue or both? Expert Rev Cardiovasc Ther 2010;8:305-9.
- Modak M, Dixit P, Londhe J, Ghaskadbi S, Devasagayam TP. Indian herbs and herbal drugs used for the treatment of diabetes. J Clin Biochem Nutr 2007;40:163-73.
- Ali H, Houghton PJ, Soumyanath A. α-amylase inhibitory activity
 of some Malaysian plants used to treat diabetes; with particular
 reference to *Phyllanthus amarus*. J Ethnopharmacol 2006;107:449-55.
- Marles R, Farnsworth NR. Plants as sources of antidiabetic agents. In: Wagner H, Farmsworth NR, editors. Economic and Medicinal Plant Research. UK: Academic Press Ltd; 1994. p. 149-87.
- Bailey CJ, Day C. Traditional plant medicines as treatments for diabetes. Diabetes Care 1989;12:553-64.
- Wild S, Roglic G, Green A, Sicree R, King H. Global prevalence of diabetes: Estimates for the year 2000 and projections for 2030. Diabetes Care 2004;27:1047-53.
- Mohan V, Sandeep S, Deepa R, Shah B, Varghese C. Epidemiology of type 2 diabetes: Indian scenario. Indian J Med Res 2007;125:217-30.
- Gupta A, Gupta R, Sarna M, Rastogi S, Gupta VP, Kothari K. Prevalence of diabetes, impaired fasting glucose and insulin resistance syndrome in an urban Indian population. Diabetes Res Clin Pract 2003;61:69-76.
- Alberti KG, Zimmet PZ. Definition, diagnosis and classification of diabetes mellitus and its complications. Part 1: Diagnosis and classification of diabetes mellitus provisional report of a WHO consultation. Diabet Med 1998;15:539-53.
- Mattecci E, Giampietro O. Oxidative stress in families of type I diabetic patient. Diabetes Care 2000;23:1182-6.
- 11. Oberlay LW. Free radicals and diabetes. Free Radic Biol Med 1988:5:113-24.
- Baynes JW, Thorpe SR, Suzanne R. The role of oxidative stress in diabetic complications. Curr Opin Endocrinol Diabetes Obes 1996;3:277-84.
- Lipinski B. Pathophysiology of oxidative stress in diabetes mellitus. J Diabetes Complications 2001;15:203-10.
- Glugliano D, Ceriello A, Paolisso G. Oxidative stress and diabetic vascular complications. Diabetes Care 1996;19:257-67.
- 15. Baynes JW, Thorpe SR. Role of oxidative stress in diabetic

- complications: A new perspective on an old paradigm. Diabetes 1999;48:1-9.
- Dandona P, Thusu K, Cook S, Snyder B, Makowski J, Armstrong D, et al. Oxidative damage to DNA in diabetes mellitus. Lancet 1996;347:444-5.
- Sakurai T, Tsuchiya S. Superoxide production from nonenzymatically glycated protein. FEBS Lett 1988;236:406-10.
- Brownlee M. Biochemistry and molecular cell biology of diabetic complications. Nature 2001;414:813-20.
- Harrison D, Griendling KK, Landmesser U, Hornig B, Drexler H. Role of oxidative stress in atherosclerosis. Am J Cardiol 2003;91:7A-11.
- Mohazzab KM, Kaminski PM, Wolin MS. NADH oxidoreductase is a major source of superoxide anion in bovine coronary artery endothelium. Am J Physiol 1994;266:H2568-72.
- Evans JL, Goldfine ID, Maddux BA, Grodsky GM. Oxidative stress and stress-activated signalling pathways: A unifying hypothesis of type 2 diabetes. Endocr Rev 2002;23:599-622.
- 22. Brownlee M. Advanced protein glycosylation in diabetes in diabetes and ageing. Annu Rev Med 1996;46:223-34.
- Elgawish A, Glomb M, Friendlander M, Monnier VM. Involvement of hydrogen peroxide in collagen cross-linking by high glucose in vitro and in vivo. J Biol Chem 1999;271:12964-71.
- Collier A, Wilson R, Bradley H, Thomson JA, Small M. Free radical activity is type 2 diabetes. Diabet Med 1990;7:27-30.
- Garg MC, Bansal DD. Protective antioxidant effect of vitamins C and E in streptozotocin induced diabetic rats. Indian J Exp Biol 2000;38:101-4.
- Anjali P, Manoj KM. Same comments on diabetes and herbal therapy. Anc Sci Life 1995;15:27-9.
- Jain SR, Sharma SN. Hypoglycemic drugs of Indian indigenous origin. Planta Med 1967;15:439-42.
- Li WL, Zheng HC, Bukuru J, De Kimpeb N. Natural medicines used in the traditional Chinese medical system for therapy of diabetes mellitus. J Ethnopharmacol 2004;92:1-21.
- Wadood A, Wadood N, Shah SA. Effects of Acacia arabica and Caralluma edulis on blood glucose levels on normal and alloxan diabetic rabbits. J Pak Med Assoc 1989;39:208-12.
- Wu HS, Zhu DF, Zhou CX, Feng CR, Lou YJ, Yang B, et al. Insulin sensitizing activity of ethyl acetate fraction of Acorus calamus L. in vitro and in vivo. J Ethnopharmacol 2009;123:288-92.
- 31. Wu HS, Li YY, Weng LJ, Zhou CX, He QJ, Lou YJ. A fraction of *Acorus calamus* L. extract devoid of β-asarone enhances adipocyte differentiation in 3T3-L1 cells. Phytother Res 2007;21:562-4.
- 32. Al-awadi FM, Gumaa KA. Studies on the activity of individual plants of an antidiabetic plant mixture. Acta Diabetol Lat 1987;24:37-41.
- 33. Ajabnoor MA. Effect of aloes on blood glucose levels in normal and alloxan diabetic mice. J Ethnopharmacol 1990;28:215-25.
- Davis RH, Maro NP. Aloe vera and gibberellins, anti-inflammatory activity in diabetes. J Am Podiatr Med Assoc 1989;79:24-6.
- Chen W, Lu Z, Viljoen A, Hamman J. Intestinal drug transport enhancement by Aloe vera. Planta Med 2009;75:587-95.
- Rajasekaran S, Ravi K, Sivagnanam K, Subramanian S. Beneficial effects of *Aloe vera* leaf gel extract on lipid profile status in rats with streptozotocin diabetes. Clin Exp Pharmacol Physiol 2006;33:232-7.
- Rajasekaran S, Sivagnanam K, Subramanian S. Modulatory effects of *Aloe vera* leaf gel extract on oxidative stress in rats treated with streptozotocin. J Pharm Pharmacol 2005;57:241-6.
- 38. Panda S. The effect of *Anethum graveolens* L. (dill) on corticosteroid induced diabetes mellitus: Involvement of thyroid hormones. Phytother Res 2008;22:1695-7.
- Kaleem M, Asif M, Ahmed QU, Bano B. Antidiabetic and antioxidant activity of *Annona squamosa* extract in streptozotocin-induced diabetic rats. Singapore Med J 2006;47:670-5.
- Gupta RK, Kesari AN, Murthy PS, Chandra R, Tandon V, Watal G. Hypoglycaemic and antidiabetic effect of ethanolic extract of leaves of *Annona squamosa* L. in experimental animals. J Ethanopharmacol 2005;99:75-81.
- Gupta RK, Kesari AN, Watal G, Murthy PS, Chandra R, Tandon V. Nutritional and hypoglycaemic effect of fruit pulp of *Annona* squamosa in normal healthy and alloxan-induced diabetic rabbits. Ann Nutr Metab 2005:49:407-13.
- Panda S, Kar A. Antidiabetic and antioxidative effects of *Annona* squamosa leaves are possibly mediated through quercetin-3-Oqlucoside. Biofactors 2007;31:201-10.

- 43. Chempakam B. Hypoglycaemic activity of arecoline in betel nut *Areca catechu* L. Indian J Exp Biol 1993;31:474-5.
- Liu M, Wu K, Mao X, Wu Y, Ouyang J. Astragalus polysaccharide improves insulin sensitivity in KKAy mice: Regulation of PKB/GLUT4 signaling in skeletal muscle. J Ethnopharmacol 2010;127:32-7.
- Wang N, Zhang D, Mao X, Zou F, Jin H, Ouyang J. Astragalus polysaccharides decreased the expression of PTP1B through relieving ER stress induced activation of ATF6 in a rat model of type 2 diabetes. Mol Cell Endocrinol 2009;307:89-98.
- Mao XQ, Yu F, Wang N, Wu Y, Zou F, Wu K, et al. Hypoglycemic effect of polysaccharide enriched extract of Astragalus membranaceus in diet induced insulin resistant C57BL/6J mice and its potential mechanism. Phytomedicine 2009;16:416-25.
- Mao XQ, Wu Y, Wu K, Liu M, Zhang JF, Zou F, et al. Astragalus polysaccharide reduces hepatic endoplasmic reticulum stress and restores glucose homeostasis in a diabetic KKAy mouse model. Acta Pharmacol Sin 2007:28:1947-56.
- Shen P, Liu MH, Ng TY, Chan YH, Yong EL. Differential effects of isoflavones, from *Astragalus membranaceus* and *Pueraria thomsonii*, on the activation of PPAR-α, PPAR-γ, and adipocyte differentiation in vitro. J Nutr 2006;136:899-905.
- Wu Y, Ou-Yang JP, Wu K, Wang Y, Zhou YF, Wen CY. Hypoglycemic effect of Astragalus polysaccharide and its effect on PTP1B. Acta Pharmacol Sin 2005;26:345-52.
- Tan BK, Tan CH, Pushparaj PN. Anti-diabetic activity of the semi-purified fractions of *Averrhoa bilimbi* in high fat diet fedstreptozotocin-induced diabetic rats. Life Sci 2005;76:2827-39.
- 51. Pushparaj PN, Tan BK, Tan CH. The mechanism of hypoglycemic action of the semi-purified fractions of *Averrhoa bilimbi* in streptozotocin-diabetic rats. Life Sci 2001;70:535-47.
- Chandra A, Mahdi AA, Singh RK, Mahdi F, Chander R. Effect of Indian herbal hypoglycemic agents on antioxidant capacity and trace elements content in diabetic rats. J Med Food 2008;11:506-12.
- Chattopadhyay RR, Chattopadhyay RN, Nandy AK, Poddar G, Maitra SK. Preliminary report on antihyperglycemic effect of fraction of fraction of fresh leaves of *Azadirachta indica* (Beng neem). Bull Calcutta Sch Trop Med 1987b;35:29-35.
- Chattopadhyay RR, Chattopadhyay RN, Nandy AK, Poddar G, Maitra SK. The effect of fresh leaves of *Azadiracta indica* on glucose uptake and glycogen content in the isolated rat hemidiaphragm. Bull Calcutta Sch Trop Med 1987b;35:8-12.
- Biswas K, Chattopadhyay I, Banerjee RK, Bandyopadhyay U. Biological activities and medicinal properties of neem (*Azadiracta indica*). Curr Sci 2002;82:1336-45.
- Roop JK, Dhaliwal PK, Guraya SS. Extracts of Azadirachta indica and Melia azedarach seeds inhibit folliculogenesis in albino rats. Braz J Med Biol Res 2005;38:943-7.
- Oliveira AC, Endringer DC, Amorim LA, das Graças L Brandão M, Coelho MM. Effect of the extracts and fractions of *Baccharis trimera* and *Syzygium cumini* on glycaemia of diabetic and non-diabetic mice. J Ethnopharmacol 2005;102:465-9.
- Suba V, Murugesan T, Arunachalam G, Mandal SC, Saha BP. Antidiabetic potential of *Barleria lupulina* extract in rats. Phytomedicine 2004:11:202-5
- Suba V, Murugesan T, Rao RB, Ghosh L, Pal M, Mandal SC, Saha BP. Antidiabetic potential of *Barleria lupulina* extract in rats. Fitoterapia 2004:75:1-4.
- Khalil NM, Pepato MT, Brunetti IL. Free radical scavenging profile and myeloperoxidase inhibition of extracts from antidiabetic plants: Bauhinia forficata and Cissus sicyoides. Biol Res 2008;41:165-71.
- De Sousa E, Zanatta L, Seifriz I, Creczynski-Pasa TB, Pizzolatti MG, Szpoganicz B, Silva FR. Hypoglycemic effect and antioxidant potential of kaempferol-3,7-O-(α)-dirhamnoside from *Bauhinia forficata* leaves. J Nat Prod 2004;67:829-32.
- 62. Singh J, Kakkar P. Antihyperglycemic and antioxidant effect of *Berberis aristata* root extract and its role in regulating carbohydrate metabolism in diabetic rats. J Ethnopharmacol 2009;123:22-6.
- Yoshikawa M, Murakami T, Kadoya M, Matsuda H, Muraoka O, Yamahara J, et al. Medicinal stuff. III sugar beet. Hypoglycaemic oleanolic acid oligosaccharides, betavulgarosides I, II, III and IV, from the root of Beta vulgaris L. Chem Pharm Bull (Tokyo) 1996;44:1212-7.
- Ponnusamy S, Ravindran R, Zinjarde S, Bhargava S, Ravi Kumar A. Evaluation of traditional Indian antidiabetic medicinal plants for human pancreatic amylase inhibitory effect in vitro. Evid Based

- Complement Alternat Med 2011;2011. pii: 515647.
- 65. Kim HY, Yokozawa T, Cho EJ, Cheigh HS, Choi JS, Chung HY. *In vitro* and *in vivo* antioxidant effects of mustard leaf (*Brassica juncea*). Phytother Res 2003;17:465-71.
- Yokozawa T, Kim HY, Cho EJ, Yamabi N, Choi JS. Protective effects of mustard leaf (*Brassica juncea*) against diabetic oxidative stress. J Nutr Sci Vitaminol (Tokyo) 2003;49:87-93.
- 67. Somani R, Kasture S, Singhai AK. Antidiabetic potential of *Butea monosperma* in rats. Fitoterapia 2006;77:86-90.
- Panda S, Jafri M, Kar A, Meheta BK. Thyroid inhibitory, antiperoxidative and hypoglycemic effects of stigmasterol isolated from *Butea* monosperma. Fitoterapia 2009;80:123-6.
- Sharma N, Garg V. Antidiabetic and antioxidant potential of ethanolic extract of *Butea monosperma* leaves in alloxan-induced diabetic mice. Indian J Biochem Biophys 2009;46:99-105.
- Sharmna N, Garg V. Antihyperglycemic and antioxidative potential of hydroalcoholic extract of *Butea monosperma* Lam flowers in alloxaninduced diabetic mice. Indian J Exp Biol 2009;47:571-6.
- Gomes A, Vedasiromoni JR, Das M, Sharma RM, Ganguly DK. Antihyperglycemic effect of black tea (*Camellia sinensis*) in rats. J Ethanopharmacol 1995;45:223-6.
- Devasagayam TP, Kamat JP, Mohan H, Kesavan PC. Caffeine as an antioxidant: Inhibition of lipid peroxidation induced by reactive oxygen species. Biochim Biophys Acta 1996;1282:63-70.
- Eddouks M, Lemhadri A, Michel JB. Caraway and caper: Potential anti-hyperglycaemic plants in diabetic rats. J Ethnopharmacol 2004;94:143-8.
- Chandramohan G, Al-Numair KS, Sridevi M, Pugalendi KV. Antihyperlipidemic activity of 3-hydroxymethyl xylitol, a novel antidiabetic compound isolated from Casearia esculenta (Roxb.) root in streptozotocin-diabetic rats. J Biochem Mol Toxicol 2010;24:95-101.
- Prakasam A, Sethupathy S, Pugalendi KV. Effect of Casearia esculenta root extract on blood glucose and plasma antioxidant status in streptozotocin diabetic rats. Pol J Pharmacol 2003;55:43-9.
- Abesundara KJ, Matsui T, Matsumoto K. α-glucosidase inhibitory activity of some Sri Lanka plant extracts, one of which, Cassia auriculata, exerts a strong antihyperglycemic effect in rats comparable to the therapeutic drug acarbose. J Agric Food Chem 2004:52:2541-5.
- Gupta S, Sharma SB, Bansal SK, Prabhu KM. Antihyperglycemic and hypolipidemic activity of aqueous extract of *Cassia auriculata* L. leaves in experimental diabetes. J Ethnopharmacol 2009;123:499-503.
- Salahuddin M, Jalalpure SS, Gadge NB. Antidiabetic activity of aqueous bark extract of *Cassia glauca* in streptozotocin-induced diabetic rats. Can J Physiol Pharmacol 2010;88:153-60.
- Lee MJ, Rao YK, Chen K, Lee YC, Tzeng YM. Effect of flavonol glycosides from Cinnamomum osmophloeum leaves on adiponectin secretion and phosphorylation of insulin receptor β in 3T3-L1 adipocytes. J Ethnopharmacol 2009;126:79-85.
- 80. Parmar HS, Kar A. Possible amelioration of atherogenic diet induced dyslipidemia, hypothyroidism and hyperglycemia by the peel extracts of *Mangifera indica*, *Cucumis melo* and *Citrullus vulgaris* fruits in rats. Biofactors 2008;33:13-24.
- 81. Parmar HS, Kar A. Antidiabetic potential of *Citrus sinensis* and *Punica granatum* peel extracts in alloxan treated male mice. Biofactors 2007;31:17-24.
- 82. Parmar HS, Kar A. Medicinal values of fruit peels from *Citrus sinensis*, *Punica granatum*, and *Musa paradisiaca* with respect to alterations in tissue lipid peroxidation and serum concentration of glucose, insulin, and thyroid hormones. J Med Food 2008;11:376-81.
- 83. Parmar HS, Kar A. Antiperoxidative, antithyroidal, antihyperglycemic and cardioprotective role of *Citrus sinensis* peel extract in male mice. Phytother Res 2008;22:791-5.
- 84. Chau CF, Huang YL, Lee MH. *In vitro* hypoglycemic effects of different insoluble fibre-rich fractions prepared from the peel of *Citrus sinensis* L. cv. Liucheng. J Agric Food Chem 2003;51:6623-6.
- 85. Adeneye AA. Methanol seed extract of *Citrus paradisi* Macfad lowers blood glucose, lipids and cardiovascular disease risk indices in normal Wistar rats. Nig Q J Hosp Med 2008;18:16-20.
- 86. Sánchez-Salgado JC, Ortiz-Andrade RR, Aguirre-Crespo F, Vergara-Galicia J, León-Rivera I, Montes S, et al. Hypoglycemic, vasorelaxant and hepatoprotective effects of *Cochlospermum vitifolium* (Willd.) Sprengel: A potential agent for the treatment of metabolic syndrome. J Ethnopharmacol 2007;109:400-5.

- Chika A, Bello SO. Antihyperglycaemic activity of aqueous leaf extract of *Combretum micranthum* (Combretaceae) in normal and alloxaninduced diabetic rats. J Ethnopharmacol 2010;129:34-7.
- Shirwaikar A, Rajendran K, Punitha IS. Antidiabetic activity of alcoholic stem extract of *Coscinium fenestratum* in streptozotocin-nicotinamide induced type 2 diabetic rats. J Ethnopharmacol 2005;97:369-74.
- Devi VD, Urooj A. Hypoglycemic potential of *Morus indica*. L and *Costus igneus*. Nak.-a preliminary study. Indian J Exp Biol 2008;46:614-6.
- Bavarva JH, Narasimhacharya AV. Antihyperglycemic and hypolipidemic effects of *Costus speciosus* in alloxan induced diabetic rats. Phytother Res 2008;22:620-6.
- Zhao Y, Son YO, Kim SS, Jang YS, Lee JC. Antioxidant and antihyperglycemic activity of polysaccharide isolated from *Dendrobium* chrysotoxum Lindl. J Biochem Mol Biol 2007;40:670-7.
- Omoruyi FO. Jamaican bitter yam sapogenin: Potential mechanisms of action in diabetes. Plant Foods Hum Nutr 2008;63:135-40.
- 93. Krishnaveni M, Mirunalini S. Therapeutic potential of *Phyllanthus emblica* (Amla): The ayurvedic wonder. J Basic Clin Physiol Pharmacol 2010;21:93-105.
- 94. Bhattacharya A, Chatterjee A, Ghosal S, Bhattacharya SK. Antioxidant activity of active tannoid principles of *Emblica officinalis* (Amla). Indian J Exp Biol 1999;37:676-80.
- Kumar KC, Muller K. Medicinal plants from Nepal, II. Evaluation as inhibitors of lipid peroxidation in biological membranes. J Ethanopharmacol 1999;64:135-9.
- Devasagayam TP, Subramanian M, Singh BB, Ramanathan R, Das NP. Protection of plasmid pBR322 DNA by flavonoids against singlestrand breaks induced by singlet molecular oxygen. J Photochem Photobiol B 1995;30:97-103.
- Augusti KT, Daniel RS, Cherian S, Sheela CG, Nair CR. Effect of leucoperalgonin derivative from *Ficus benagalnesis* Linn. on diabetic dogs. Indian J Med Res 1994;99:82-6.
- Singh RK, Mehta S, Jaiswal D, Rai PK, Watal G. Antidiabetic effect of *Ficus bengalensis* aerial roots in experimental animals. J Ethnopharmacol 2009;123:110-4.
- Campillo JE, Perez C, Ramiro JM, Torres MD. Hypoglycemic activity
 of an aqueous extract from *Ficus carica* in streptozotocin diabetic
 rats (Abstract). Diabetologia 1991;34 Suppl 2:A-181.
- Torres MD, Dominguez E, Romero A, Campillo JE, Perez C. Hypoglycemic and hypolipidemic activity of an aqueous extract from Ficus carica in streptozotocin diabetic rats (Abstract). Diabetologia 1993;36 Suppl 1:A-181.
- Perez C, Dominguez E, Ramiro A, Campillo J, Torres MD. A study on the glycemic balance in streptozotocin diabetic rats treated with an aqueous extract of *Ficus carica* leaves. Phytother Res 1996;10:82-6.
- 102. Ahmed F, Urooj A. *In vitro* studies on the hypoglycemic potential of *Ficus racemosa* stem bark. J Sci Food Agric 2010;90:397-401.
- Jouad H, Maghrani M, Eddouks M. Hypoglycaemic effect of Rubus fructicosis L. and Globularia alypum L. in normal and streptozotocininduced diabetic rats. J Ethnopharmacol 2002;81:351-6.
- 104. Ramkumar KM, Lee AS, Krishnamurthi K, Devi SS, Chakrabarti T, Kang KP, et al. Gymnema montanum H. protects against alloxan-induced oxidative stress and apoptosis in pancreatic β-cells. Cell Physiol Biochem 2009;24:429-40.
- 105. Ananthan R, Latha M, Ramkumar KM, Pari L, Baskar C, Narmatha Bai V. Effect of *Gymnema montanum* leaves on serum and tissue lipids in alloxan diabetic rats. Exp Diabesity Res 2003;4:183-9.
- 106. Yeo J, Kang YJ, Jeon SM, Jung UJ, Lee MK, Song H, et al. Potential hypoglycemic effect of an ethanol extract of Gynostemma pentaphyllum in C57BL/KsJ-db/db mice. J Med Food 2008;11:709-16.
- 107. Prabhakar Reddy P, Tiwari AK, Ranga Rao R, Madhusudhana K, Rama Subba Rao V, Ali AZ, et al. New Labdane diterpenes as intestinal α-glucosidase inhibitor from antihyperglycemic extract of Hedychium spicatum (Ham. Ex Smith) rhizomes. Bioorg Med Chem Lett 2009;19:2562-5.
- 108. Aslan M, Deliorman Orhan D, Orhan N, Sezik E, Yesilada E. In vivo antidiabetic and antioxidant potential of Helichrysum plicatum ssp. Plicatum capitulums in streptozotocin-induced-diabetic rats. J Ethnopharmacol 2007;109:54-9.
- 109. Boopathy Raja A, Elanchezhiyan C, Sethupathy S. Antihyperlipidemic activity of *Helicteres isora* fruit extract on streptozotocin induced diabetic male Wistar rats. Eur Rev Med Pharmacol Sci 2010;14:191-6.
- 110. Kumar G, Banu GS, Murugesan AG, Pandian MR. Hypoglycaemic

- effect of *Helicteres isora* bark extract in rats. J Ethnopharmacol 2006;107:304-7.
- 111. Chakrabarti R, Vikramadithyan RK, Mullangi R, Sharma VM, Jagadheshan H, Rao YN, et al. Antidiabetic and hypolipidemic activity of *Helicteres isora* in animal models. J Ethnopharmacol 2002;81:343-9.
- 112. Vijayakumar M, Govindarajan R, Rao GM, Rao ChV, Shirwaikar A, Mehrotra S, et al. Action of Hygrophila auriculata against streptozotocin-induced oxidative stress. J Ethnopharmacol 2006:104:356-61.
- 113. Erasto P, Adebola PO, Grierson DS, Afolayan AJ. An ethnobotanical study of plants used for the treatment of diabetes in Eastern Cape Province, South Africa. Afr J Biotechnol 2005;4:1458-60.
- 114. Owira PM, Ojewole JA. 'African potato' (*Hypoxis hemerocallidea corm*): A plant-medicine for modern and 21st century diseases of mankind? A review. Phytother Res 2009;23:147-52.
- 115. Zibula SM, Ojewole JA. Hypoglycaemic effects of *Hypoxis hemerocallidea corm* 'African Potato' methanolic extract in rats. Med J Islam World Acad Sci 2000;13:75-8.
- 116. Ojewole JA. Antinociceptive, anti-inflammatory and antidiabetic properties of *Hypoxis hemerocallidea* Fisch. and C.A. Mey. (Hypoxidaceae) corm ['African Potato'] aqueous extract in mice and rats. J Ethnopharmacol 2006; 103:126-34.
- 117. Mahomed IM, Ojewole JA. Hypoglycemic effect of *Hypoxis hemerocallidea* corm (African potato) aqueous extract in rats. Methods Find Exp Clin Pharmacol 2003;25:617-23.
- 118. Chakrabarti R, Damarla RK, Mullangi R, Sharma VM, Vikramadithyan RK, Rajagopalan R. Insulin sensitizing property of *Indigofera* mysorensis extract. J Ethnopharmacol 2006;105:102-6.
- 119. Ju JB, Kim JS, Choi CW, Lee HK, Oh TK, Kim SC. Comparison between ethanolic and aqueous extracts from Chinese juniper berries for hypoglycemic and hypolipidemic effects in alloxan-induced diabetic rats. J Ethnopharmacol 2008;115:110-5.
- 120. Hou W, Li Y, Zhang Q, Wei X, Peng A, Chen L, et al. Triterpene acids isolated from $Lagerstroemia\ speciosa\$ leaves as α -glucosidase inhibitors. Phytother Res 2009;23:614-8.
- 121. P S, Zinjarde SS, Bhargava SY, Ravikumar A. Potent α -amylase inhibitory activity of Indian Ayurvedic medicinal plants. BMC Complement Altern Med 2011;11:5.
- 122. Chen X, Liu Y, Bai X, Wen L, Fang J, Ye M, et al. Hypoglycemic polysaccharides from the tuberous root of *Liriope spicata*. J Nat Prod 2009:72:1988-92.
- 123. Chen X, Bai X, Liu Y, Tian L, Zhou J, Zhou Q, et al. Anti-diabetic effects of water extract and crude polysaccharides from tuberous root of *Liriope spicata* var. prolifera in mice. J Ethnopharmacol 2009:122:205-9.
- 124. Cemek M, Kağa S, Simşek N, Büyükokuroğlu ME, Konuk M.. Antihyperglycemic and antioxidative potential of *Matricaria chamomilla* L. in streptozotocin-induced diabetic rats. J Nat Med 2008;62:284-93.
- 125. Rao BK, Kessavulu MM, Giri R, Appa Rao C. Antidiabetic and hypolipidemic effects of *Momordica cymbalaria* Hook fruit powder in alloxan-diabetic rats. J Ethanopharmacol 1999;67:103-9.
- Hansawasdi C, Kawabata J. α-glucosidase inhibitory effect of mulberry (Morus alba) leaves on Caco-2. Fitoterapia 2006;77:568-73.
- Khan BA, Abraham A, Leelamma S. Hypoglycaemic action of *Murraya koenigii* (curry leaf) and *Brassica juncea* (mustard): Mechanism of action. Indian J Biochem Biophys 1995;32:106-8.
- 128. Arulselvan P, Senthilkumar GP, Sathish Kumar D, Subramanian S. Antidiabetic effect of *Murraya koenigii* leaves on streptozotocin induced diabetic rats. Pharmazie 2006;61:874-7.
- 129. Dhanabal SP, Sureshkumar M, Ramanathan M, Suresh B. Hypoglycaemic effect of ethanolic extract of *Musa sapientum* on alloxan induced diabetes mellitus in rats and its relation with antioxidant potential. J Herb Pharmacother 2005;5:7-19.
- 130. Pari L, Umamaheswari J. Antihyperglycemic activity of *Musa sapientum* flowers: Effect on lipid peroxidation in alloxan-induced diabetic rats. Phytother Res 2000;14:136-8.
- 131. Pari L, Maheswari JU. Hypoglycaemic effect of *Musa sapientum* L. in alloxan-induced rats. J Ethanopharmacol 1999;68:321-5.
- 132. El SN, Karakaya S. Olive tree (*Olea europaea*) leaves: Potential beneficial effects on human health. Nutr Rev 2009;67:632-8.
- 133. Baumgartner RR, Steinmann D, Heiss EH, Atanasov AG, Ganzera M, Stuppner H, et al. Bioactivity-guided isolation of 1,2,3,4,6-Penta-

- O-galloyl-D-glucopyranose from *Paeonia lactiflora* roots as a PTP1B inhibitor. J Nat Prod 2010;73:1578-81.
- 134. Jain S, Bhatia G, Barik R, Kumar P, Jain A, Dixit VK. Antidiabetic activity of *Paspalum scrobiculatum* Linn. in alloxan induced diabetic rats. J Ethnopharmacol 2010;127:325-8.
- 135. Brai BI, Odetola AA, Agomo PU. Hypoglycemic and hypocholesterolemic potential of *Persea americana* leaf extracts. J Med Food 2007;10:356-60.
- Babish JG, Pacioretty LM, Bland JS, Minich DM, Hu J, Tripp ML. Antidiabetic screening of commercial botanical products in 3T3-L1 adipocytes and db/db mice. J Med Food 2010;13:535-47.
- 137. Tormo MA, Gil-Exojo I, Romero de Tejada A, Campillo JE. Hypoglycaemic and anorexigenic activities of an α-amylase inhibitor from white kidney beans (*Phaseolus vulgaris*) in wistar rats. Br J Nutr 2004:92:785-90.
- Pari L, Venkateswaran S. Protective role of *Phaseolus vulgaris* on changes in the fatty acid composition in experimental diabetes. J Med Food 2004;7:204-9.
- 139. Knott RM, Grant G, Bardocz S, Pusztai A, De Carvalho AF, Hesketh JE. Alternations in the level of insulin receptor and GLU-4 mRNA in skeletal muscle from rats fed a kidney bean (*Phaseolus vulgaris*) diet. Int J Biochem 1992;24:897-902.
- 140. Okada Y, Okada M, Sagesaka Y. Screening of dried plant seed extracts for adiponectin production activity and tumor necrosis factor-α inhibitory activity on 3T3-L1 adipocytes. Plant Foods Hum Nutr 2010:65:225-32.
- 141. Kasabri V, Afifi FU, Hamdan I. In vitro and in vivo acute antihyperglycemic effects of five selected indigenous plants from Jordan used in traditional medicine. J Ethnopharmacol 2011;133:888-96.
- Zhang H, Yang F, Qi J, Song XC, Hu ZF, Zhu DN, et al. Homoisoflavonoids from the fibrous roots of *Polygonatum odoratum* with glucose uptakestimulatory activity in 3T3-L1 adipocytes. J Nat Prod 2010;73:548-52.
- 143. Jafri MA, Aslam M, Javed K, Singh S. Effect of *Punica granatum* Linn. (Flowers) on blood glucose level in normal and alloxan induced diabetic rats. J Ethnopharmacol 2000;70:309-14.
- 144. Katz SR, Newman RA, Lansky EP. *Punica granatum*: Heuristic treatment for diabetes mellitus. J Med Food 2007;10:213-7.
- 145. Huang TH, Peng G, Kota BP, Li GQ, Yamahara J, Roufogalis BD, et al. Anti-diabetic action of Punica granatum flower extract: Activation of PPAR-γ and identification of an active component. Toxicol Appl Pharmacol 2005;207:160-9.
- 146. Li Y, Wen S, Kota BP, Peng G, Li GQ, Yamahara J, et al. Punica granatum flower extract, a potent α-glucosidase inhibitor, improves postprandial hyperglycemia in Zucker diabetic fatty rats. J Ethnopharmacol 2005;99:239-44.
- 147. Shukla S, Chatterji S, Mehta S, Rai PK, Singh RK, Yadav DK, et al. Antidiabetic effect of Raphanus sativus root juice. Pharm Biol 2011;49:32-7.
- 148. Taniguchi H, Muroi R, Kobayashi-Hattori K, Uda Y, Oishi Y, Takita T. Differing effects of water-soluble and fat-soluble extracts from Japanese radish (*Raphanus sativus*) sprouts on carbohydrate and lipid metabolism in normal and streptozotocin-induced diabetic rats. J Nutr Sci Vitaminol (Tokyo) 2007;53:261-6.
- 149. Taniguchi H, Kobayashi-Hattori K, Tenmyo C, Kamei T, Uda Y, Sugita-Konishi Y, et al. Effect of Japanese radish (Raphanus sativus) sprout (Kaiware-daikon) on carbohydrate and lipid metabolisms in normal and streptozotocin-induced diabetic rats. Phytother Res 2006;20:274-8.
- Karunanayake EH, Welihinda J, Sirimame SR, Sinnadorai G. Oral hypoglycemic activity of some medicinal plants of Sri lanka. J Ethnopharmacol 1984;11:223-31.
- 151. Algandaby MM, Alghamdi HA, Ashour OM, Abdel-Naim AB, Ghareib SA, Abdel-Sattar EA, et al. Mechanisms of the antihyperglycemic activity of Retama raetam in streptozotocin-induced diabetic rats. Food Chem Toxicol 2010;48:2448-53.
- Cheng XJ, Di L, Wu Y. Hyperglycemic effect of polysaccharides from Rhodiola sachalinensis A. China. J Chin Materia Med 1993;18:557-9.
- 153. Gao D, Li Q, Liu Z, Feng J, Li J, Han Z, et al. Antidiabetic potential of Rhodiola sachalinensis root extract in streptozotocin-induced diabetic rats. Methods Find Exp Clin Pharmacol 2009;31:375-81.
- 154. Ojewole JA. Hypoglycemic effect of Sclerocarya birrea [(A. Rich.) Hochst.] [Anacardiaceae] stem-bark aqueous extract in rats. Phytomedicine 2003;10:675-81.
- 155. Dimo T, Rakotonirina SV, Tan PV, Azay J, Dongo E, Kamtchouing

- P, et al. Effect of Sclerocarya birrea (Anacardiaceae) stem bark methylene chloride/methanol extract on streptozotocin-diabetic rats. J Ethnopharmacol 2007;110:434-8.
- 156. Gondwe M, Kamadyaapa DR, Tufts M, Chuturgoon AA, Musabayane CT. Sclerocarya birrea [(A. Rich.) Hochst.] [Anacardiaceae] stem-bark ethanolic extract (SBE) modulates blood glucose, glomerular filtration rate (GFR) and mean arterial blood pressure (MAP) of STZ-induced diabetic rats. Phytomedicine 2008;15:699-709.
- 157. Ojewole JA, Mawoza T, Chiwororo WD, Owira PM. Sclerocarya birrea (A. Rich) Hochst. ['Marula'] (Anacardiaceae): A review of its phytochemistry, pharmacology and toxicology and its ethnomedicinal uses. Phytother Res 2010;24:633-9.
- 158. Chadwick WA, Roux S, Van de Venter M, Louw J, Oelofsen W. Anti-diabetic effects of *Sutherlandia frutescencs* in Wistar rats fed a diabetogenic diet. J Ethnopharmacol 2007;109:121-7.
- Ojewole JA. Analgesic, anti-inflammatory and hypoglycemic effects of Sutherlandia frutescens R. BR. (variety Incana E. MEY.) [Fabaceae] shoot aqueous extract. Methods Find Exp Clin Pharmacol 2004;26:409-16.
- Dewanjee S, Maiti A, Das AK, Mandal SC, Dey SP. Swietenine: A potential oral hypoglycemic from Swietenia macrophylla seed. Fitoterapia 2009;80:249-51.
- Panda SP, Haldar PK, Bera S, Adhikary S, Kandar CC. Antidiabetic and antioxidant activity of Swietenia mahagoni in streptozotocin-induced diabetic rats. Pharm Biol 2010;48:974-9.
- Lam SH, Chen JM, Kang CJ, Chen CH, Lee SS. α-Glucosidase inhibitors from the seeds of Syagrus romanzoffiana. Phytochemistry 2008:69:1173-8.
- 163. Rao BK, Rao CH. Hypoglycaemic and antihyperglycemic activity of Syzygium alternifolium (Wt.) Walp. seed extracts in normal and diabetic rats. Phytomedicine 2001;8:88-93.
- Nagappa AN, Thakurdesai PA, Venkat Rao N, Singh J. Antidiabetic activity of *Terminalia catappa* Linn fruits. J Ethnopharmacol 2003;88:45-50.
- Pareek H, Sharma S, Khajja BS, Jain K, Jain GC. Evaluation of hypoglycemic and anti-hyperglycemic potential of *Tridax procumbens* (Linn.). BMC Complement Altern Med 2009;9:48.
- 166. Martineau LC, Couture A, Spoor D, Benhaddou-Andaloussi A, Harris C, Meddah B, et al. Anti-diabetic properties of the Canadian lowbush blueberry Vaccinium angustifolium Ait. Phytomedicine 2006;13:612-23.
- 167. Wang L, Zhang XT, Zhang HY, Yao HY, Zhang H. Effect of Vaccinium bracteatum Thunb. leaves extract on blood glucose and plasma lipid levels in streptozotocin-induced diabetic mice. J Ethnopharmacol 2010;130:465-9.
- 168. Eid HM, Martineau LC, Saleem A, Muhammad A, Vallerand D, Benhaddou-Andaloussi A, et al. Stimulation of AMP-activated protein kinase and enhancement of basal glucose uptake in muscle cells by quercetin and quercetin glycosides, active principles of the antidiabetic medicinal plant Vaccinium vitis-idaea. Mol Nutr Food Res 2010;54:991-1003.
- 169. Beaulieu LP, Harris CS, Saleem A, Cuerrier A, Haddad PS, Martineau LC, et al. Inhibitory effect of the Cree traditional medicine wiishichimanaanh (Vaccinium vitis-idaea) on advanced glycation end product formation: Identification of active principles. Phytother Res 2010;24:741-7.
- 170. Gyang SS, Nyam DD, Sokomba EN. Hypoglycaemic activity of Vernonia amygdalina (chloroform extract) in normoglycemic and alloxan-induced hyperglycemic rats. J Pharm Bioresour 2004;1:61-6.
- 171. Osinubi AA. Effects of *Vernonia amygdalina* and chlorpropamide on blood glucose. Med J Islam World Acad oSci 2008;16:115-9.
- Adallu B, Radhika B. Hypoglycaemic, diuretic and hypocholesterolemic effect of Winter cherry (Withania somnifera, Dunal) root. Indian J Exp Biol 2000:38:607-9.
- 173. Jatwa R, Kar A. Amelioration of metformin-induced hypothyroidism by Withania somnifera and Bauhinia purpurea extracts in Type 2 diabetic mice. Phytother Res 2009;23:1140-5.
- 174. Ponnachan PT, Paulose CS, Panikkar KR. Effect of leaf extract of Aegle marmelos in diabetic rats. Indian J Exp Biol 1993;31:345-7.
- Sabu MC, Kuttan R. Antidiabetic activity of Aegle marmelos and its relationship with its antioxidant properties. Indian J Physiol Pharmacol 2004;48:81-8.
- 176. Das AV, Padayatti PS, Paulose CS. Effect of leaf extract of Aegle marmelos (L.) Correa ex Roxb. on histological and ultra structural

- changes in tissues of streptozotocin induced diabetic rats. Indian J Exp Biol 1996;34:341-5.
- 177. Panda S, Kar A. Evaluation of the antithyroid, antioxidative and antihyperglycemic activity of scopoletin from *Aegle marmelos* leaves in hyperthyroid rats. Phytother Res 2006;20:1103-5.
- 178. Ramesh B, Pugalendi KV. Effect of umbelliferone on tail tendon collagen and haemostatic function in streptozotocin-diabetic rats. Basics Clin Pharmacol Toxicol 2007;101:73-7.
- 179. Roman-Ramos R, Flores-Saenz JL, Alaricon-Aguilar FJ. Antihyperglycemic effect of some edible plants. J Ethnopharmacol 1995;48:25-32.
- 180. Campos KE, Diniz YS, Cataneo AC, Faine LA, Alves MJ, Novelli EL. Hypoglycaemic and antioxidant effects of onion, Allium cepa: Dietary onion addition, antioxidant activity and hypoglycemic effects on diabetic rats. Int J Food Sci Nutr 2003;54:241-6.
- Mathew PT, Augusti KT. Hypoglycaemic effects of onion, Allium cepa Linn. on diabetes mellitus- a preliminary report. Indian J Physiol Pharmacol 1975;19:213-7.
- 182. Kumari K, Mathew BC, Augusti KT. Antidiabetic and hypolipidemic effects of S-methyl cysteine sulfoxide, isolated from *Allium cepa* Linn. Indian J Biochem Biophys 1995;32:49-54.
- Dini I, Tenore GC, Dini A. S-alkenyl cysteine sulfoxide and its antioxidant properties from Allium cepa var. tropeana (red onion) seeds. J Nat Prod 2008;71:2036-7.
- Zacharias NT, Sebastian KL, Philip B, Augusti KT. Hypoglycaemic and hypolipidemic effects of garlic in sucrose fed rabbits. Indian J Physiol Pharmacol 1980;24:151-4.
- Ahmad MS, Ahmed N. Antiglycation properties of aged garlic extract: Possible role in prevention of diabetic complications. J Nutr 2006;136 (Suppl 3):796S-799S.
- 186. Liu CT, Sheen LY, Lii CK. Does garlic have a role as an antidiabetic agent? Mol Nutr Food Res 2007;51:1353-64.
- Sheela CG, Augusti KT. Antidiabetic effects of S-allyl cysteine sulfoxide isolated from garlic *Allium sativum* Linn. Indian J Exp Biol 1992;30:523-6.
- Bever BO, Zahnd GR. Plants with oral hypoglycemic action. Q J Crude Drug Res 1979;17:139-49.
- Okada Y, Tanaka K, Sato E, Okajima H. Kinetic and mechanistic studies of allicin as an antioxidant. Org Biomol Chem 2006;4:4113-7.
- Coppi A, Cabinian M, Mirelman D, Sinnis P. Antimalarial activity of allicin, a biologically active compound from garlic cloves. Antimicrob Agents Chemother 2006;50:1731-7.
- Augusti KT, Shella CG. Antiperoxide effect of S-allyl cysteine sulfoxide, an insulin secretagouge in diabetic rats. Experientia 1996;52:115-20.
- Ledezma E, Apitz-Castro R. Ajoene the main active compound of garlic (Allium sativum): A new antifungal agent. Rev Iberoam Micol 2006;23:75-80.
- 193. Chakrabarti S, Biswas TK, Rokeya B, Ali L, Mosihuzzaman M, Nahar N, et al. Advanced studies on the hypoglycemic effect of Caesalpinia bonducella F. in type 1 and 2 diabetes in Long Evans rats. J Ethanopharmacol 2003;84:41-6.
- 194. Sharma SR, Dwiedi SK, Swarup D. Hypoglycaemic, antihyperglycemic and hypolipidemic activities of *Caesalpinia bonducella* seeds in rats. J Ethanopharmacol 1997;58:39-44.
- 195. Kannur DM, Hukkeri VI, Akki KS. Antidiabetic activity of *Caesalpinia bonducella* seed extracts in rats. Fitoterapia 2006;77:546-9.
- 196. Yadav P, Sarkar S, Bhatnagar D. Lipid peroxidation and enzymes in erythrocytes and tissues in aged diabetic rats. Indian J Exp Biol 1997;35:389-92.
- Sharma B, Salunke R, Balomajumder C, Daniel S, Roy P. Anti-diabetic potential of alkaloid rich fraction from *Capparis decidua* on diabetic mice. J Ethnopharmacol 2010:127:457-62.
- 198. Mallick C, De D, Ghosh D. Correction of protein metabolic disorders by composite extract of *Musa paradisiaca* and *Coccinia indica* in streptozotocin-induced diabetic albino rat: An approach through the pancreas. Pancreas 2009;38:322-9.
- 199. Mallick C, Mandal S, Barik B, Bhattacharya A, Ghosh D. Protection of testicular dysfunctions by MTEC, a formulated herbal drug, in streptozotocin induced diabetic rat. Biol Pharm Bull 2007;30:84-90.
- Kamble SM, Kamlakar PL, Vaidya S, Bambole VD. Influence of Coccinia indica on certain enzymes in glycolytic and lipolytic pathway in human diabetes. Indian J Med Sci 1998;52:143-6.
- 201. Kuriyan R, Rajendran R, Bantwal G, Kurpad AV. Effect of supplementation of *Coccinia cordifolia* extract on newly detected

- diabetic patients. Diabetes Care 2008;31:216-20.
- 202. El-Hossary GA, Fathy MM, Kassem HA, Kandil ZA, Abdel Latif HA. Phytochemical and biological investigations of *Eriobotrya japonica* L. growing in Egypt. Bull Fac Pharm 2000;38:129-42.
- Noreen, W, Wadood A, Hidayat HK, Wahid SA. Effect of *Eriobotrya japonica* on blood glucose levels of normal and alloxan-diabetic rabbits. Planta Med 1988;54:196-9.
- Roman-Ramos R, Flores-Saenz JL, Partida-Hernandez G, Lara-Lemus A, Alarcon-Aguilar F. Experimental study of the hypoglycemic effect of some antidiabetic plants. Arch Invest Med (Mex) 1991;22:87-93.
- Lü H, Chen J, Li WL, Ren BR, Wu JL, Zhang HQ. Hypoglycemic effect of the total flavanoid fraction from folium Eriobotryae. Phytomedicine 2009;16:967-71.
- 206. Gumy C, Thurnbichler C, Aubry EM, Balazs Z, Pfisterer P, Baumgartner L, et al. Inhibition of 11 β -hydroxysteroid dehydrogenase type 1 by plant extracts used as traditional antidiabetic medicines. Fitoterapia 2009:80:200-5.
- De Tommasi N, De Simone F, Cirino G, Cicala C, Pizza C. Hypoglycemic effects of sesquiterpene glycosides and polyhydroxylated triterpenoids of *Eriobotrya japonica*. Planta Med 1991;57:414-6.
- 208. Lü H, Chen J, Li WL, Ren BR, Wu JL, Kang HY, *et al.* Hypoglycemic and hypolipidemic effects of the total triterpene acid fraction from Folium Eriobotryae. J Ethnopharmacol 2009;122:486-91.
- 209. Acherekar S, Kakliji GS, Pote MS, Kelkar SM. Hypoglycemic activity of *Eugenia jambolana* and *Ficus bengalensis*: Mechanism of action. *In vivo* 1991;5:143-7.
- Ravi K, Ramachandran B, Subramanian S. Protective effect of Eugenia jambolana seed kernel on tissue antioxidants in streptozotocininduced diabetic rats. Biol Pharm Bull 2004;27:1212-7.
- 211. Yadav M, Lavania A, Tomar R, Prasad GB, Jain S, Yadav H. Complementary and comparative study on hypoglycemic and antihyperglycemic activity of various extracts of *Eugenia jambolana* seed, *Momordica charantia* fruits, *Gymnema sylvestre*, and Trigonella foenum graecum seeds in rats. Appl Biochem Biotechnol 2010;160:2388-400.
- Pepato MT, Mori DM, Baviera AM, Harami JB, Vendramini RC, Brunetti IL. Fruit of the jambolana tree (*Eugenia jambolana* Lam.) and experimental diabetes. J Ethnopharmacol 2005;96:43-8.
- 213. Attele AS, Wu JA, Yuan CS. Ginseng pharmacology: Multiple constituents and multiple actions. Biochem Pharmacol 1999;58:1685-93.
- 214. Christensen LP. Ginsenosides chemistry, biosynthesis, analysis, and potential health effects. Adv Food Nutr Res 2009;55:1-99.
- Liu CX, Xiao PG. Recent advances on ginseng research in China. J Ethnopharmacol 1992;36:27-38.
- 216. Ohnishi Y, Takagi S, Miura T, Usami M, Kako M, Ishihara E, et al. Effect of ginseng radix on GLUT2 protein content in mouse liver in normal and epinephrine-induced hyperglycemic mice. Biol Pharm Bull 1996;19:1238-40.
- Oshima Y, Sato K, Hikino H. Isolation and hypoglycemic activity of quinquefolans A, B, and C, glycans of Panax quinquefolium roots. J Nat Prod 1987;50:188-90.
- 218. Martinez B, Staba EJ. The physiological effects of Aralia, Panax and Eleutherococcus on exercised rats. Jpn J Pharmacol 1984;35:79-85.
- 219. Konno C, Sugiyama K, Kano M, Takahashi M, Hikino H. Isolation and hypoglycemic activity of panaxans A, B, C, D, and E, glycans of Panax ginseng roots. Planta Med 1984;50:434-6.
- 220. Konno C, Murakami M, Oshima Y, Hikino H. Isolation and hypoglycemic activity of panaxans Q, R, S, T, and U, glycans of Panax ginseng roots. J Ethnopharmacol 1985;14:69-74.
- Yokozawa T, Kobayashi T, Oura H, Kawashima Y. Studies on the mechanism of the hypoglycemic activity of ginsenoside-Rb2 in streptozotocin-diabetic rats. Chem Pharm Bull (Tokyo) 1985;33:869-72.
- 222. Oshima Y, Konno C, Hikino H. Isolation and hypoglycemic activity of panaxans I, J, K and L, glycans of Panax ginseng roots. J Ethnopharmacol 1985;14:255-9.
- Sotaniemi EA, Haapakoski E, Rautio A. Ginseng therapy in noninsulin-dependent diabetic patients. Diabetes Care 1995;18:1373-5.
- 224. Vuksan V, Sievenpiper JL, Koo VY, Francis T, Beljan-Zdravkovic U, Xu Z, et al. American ginseng (*Panax quinquefolius* L) reduces postprandial glycemia in nondiabetic subjects and subjects with type 2 diabetes mellitus. Arch Intern Med 2000;160:1009-13.
- 225. Punnonen R, Lukola A. Oestrogen-like effect of ginseng. Br Med J 1980;281:1110.

- Schulz V, Hansel R, Tyler VE. Rational phytotherapy. Agents that Increase Resistance to Diseases. New York, NY: Springer- Verlag; 1998: p. 269-72.
- Persaud SJ, Al-Majed H, Raman A, Jones PM. Gymnema sylvestre stimulates insulin release in vitro by increased membrane permeability. J Endocrinol 1999;163:207-12.
- Daisy P, Eliza J, Mohamed Farook KA. A novel dihydroxy gymnemic triacetate isolated from *Gymnema sylvestre* possessing normoglycemic and hypolipidemic activity on STZ-induced diabetic rats. J Ethnopharmacol 2009;126:339-44.
- 229. Baskaran K, Kizar Ahamath B, Radha Shanmugasundaram K, Shanmugasundaram ER. Antidiabetic effect of a leaf extracts from *Gymnema sylvestre* in non-insulin-dependent diabetes mellitus patients. J Ethnopharmacol 1990;30:295-300.
- Mozersky RP. Herbal products and supplemental nutrients used in the management of diabetes. J Am Osteopath Assoc 1999;99(12 Suppl):S4-9.
- 231. Aderibigbe AO, Emudianughe TS, Lawal BA. Antihyperglycemic effect of *Manaifera indica* in rat. Phytother Res 1999;13:504-7.
- Prasad S, Kalra N, Shukla Y. Hepatoprotective effects of lupeol and mango pulp extract of carcinogen induced alteration in Swiss albino mice. Mol Nutr Food Res 2007;51:352-9.
- Muruganandan S, Srinivasan K, Gupta S, Gupta PK, Lal J. Effect of mangiferin on hyperglycemia and atherogenicity in streptozotocin diabetic rats. J Ethnopharmacol 2005;97:497-501.
- 234. Shabib BA, Khan LA, Rahman R. Hypoglycemic activity of *Coccinia indica* and *Momordica charantia* in diabetic rats: Depression of the hepatic gluconeogenic enzymes glucose-6-phosphatase and fructose-1, 6-biphosphatase and elevation of liver and red-cell shunt enzyme glucose-6-phosphate dehydrogenase. Biochem J 1993;292:267-270.
- 235. Fernandes NP, Lagishetty CV, Panda VS, Naik SR. An experimental evaluation of the antidiabetic and antilipidemic properties of a standardized *Momordica charantia* fruit extract. BMC Complement Altern Med 2007;7:29.
- 236. Sridhar MG, Vinayagamoorthi R, Arul Suyambunathan V, Bobby Z, Selvaraj N. Bitter gourd (*Momordica charantia*) improves insulin sensitivity by increasing skeletal muscle insulin-stimulated IRS-1 tyrosine phosphorylation in high-fat-fed rats. Br J Nutr 2008;99:806-12.
- Sathishsekar D, Subramanian S. Antioxidant properties of *Momordica Charantia* (bitter gourd) seeds on Streptozotocin induced diabetic rats. Asia Pac J Clin Nutr 2005;14:153-8.
- 238. Sitasawad SL, Shewade Y, Bhonde R. Role of bittergourd fruit juice in STZ-induced diabetic state *in vivo* and *in vitro*. J Ethnopharmacol 2000;73:71-9.
- 239. McCarty MF. Does bitter melon contain an activator of AMP-activated kinase? Med Hypotheses 2004;63:340-3.
- 240. Cheng HL, Huang HK, Chang CI, Tsai CP, Chou CH. A cell-based screening identifies compounds from the stem of *Momordica* charantia that overcome insulin resistance and activate AMP-activated protein kinase. J Agric Food Chem 2008;56:6835-43.
- 241. Khanna P, Jain SC, Panagariya A, Dixit VP. Hypoglycemic activity of polypeptide-p from a plant source. J Nat Prod 1981;44:648-55.
- Platel K, Srinivasan K. Plant foods in the management of diabetes mellitus: Vegetables as potential hypoglycemic agents. Nahrung 1997;41:68-74.
- 243. Ooi CP, Yassin Z, Hamid TA. *Momordica charantia* for type 2 diabetes mellitus. Cochrane Database Syst Rev 2010;(2):CD007845.
- Grover JK, Yadav SP. Pharmacological actions and potential uses of Momordica charantia: A review. J Ethnopharmacol 2004;93:123-32.
- 245. Vats V, Grover JK, Rathi SS. Evaluation of antihyperglycemic and hypoglycemic effect of *Trigonella foenum graecum* Linn, *Ocimum sanctum* Linn and *Pterocarpus marsupium* Linn in normal and alloxanized diabetic rats. J Ethanopharmacol 2002;79:95-100.
- 246. Rai V, Iyer U, Mani UV. Effect of tulsi (*Ocimum sanctum*) leaf powder supplementation on blood sugar levels, serum lipids and tissue lipid in diabetic rats. Plant foods Hum Nutr 1997;50:9-16.
- 247. Vats V, Yadav SP, Grover JK. Ethanolic extract of *Ocimum sanctum* leaves partially attenuates streptozotocin induced alteration in glycogen content and carbohydrate metabolism in rats. J Ethanopharmacol 2004;90:155-60.
- Mondal S, Mirdha BR, Mahapatra SC. The science behind sacredness of Tulsi (*Ocimum sanctum* Linn.). Indian J Physiol Pharmacol 2009;53:291-306.

- 249. Raphael KR, Sabu MC, Kuttan R. Hypoglycemic effect of methanol extract of *Phyllanthus amarus* Schum and Thonn on alloxan induced diabetes mellitus in rats and its relation with antioxidant potential. Indian J Exp Biol 2002;40:905-9.
- 250. Srividya N, Periwal S. Diuretic, hypotensive and hypoglycemic effect of *Phyllanthus amarus*. Indian J Exp Biol 1995;33:861-4.
- 251. Shabeer J, Srivastava RS, Singh SK. Antidiabetic and antioxidant effect of various fractions of *Phyllanthus simplex* in alloxan diabetic rats. J Ethnopharmacol. 2009;124:34-8.
- 252. Sharma P, Parmar J, Verma P, Sharma P, Goyal PK. Anti-tumor activity of *Phyllanthus niruri* (a medicinal plant) on chemical-induced skin carcinogenesis in mice. Asian Pac J Cancer Prev 2009;10:1089-94.
- 253. Kumar S, Kumar D, Deshmukh RR, Lokhande PD, More SN, Rangari VD. Antidiabetic potential of *Phyllanthus reticulatus* in alloxan-induced diabetic mice. Fitoterapia 2008;79:21-3.
- Mukhtar HM, Ansari SH, Ali M, Naved T, Bhat ZA. Effect of water extract of *Psidium guajava* on alloxan induced diabetic rats. Pharmazie 2004:59:734-5.
- Ojewole JA. Hypoglycemic and hypotensive effects of *Psidium guajava* Linn. (Myrtaceae) leaf aqueous extract. Methods Find Exp Clin Pharmacol 2005;27:689-95.
- 256. Mukhtar HM, Ansari SH, Bhat ZA, Naved T, Singh P. Antidiabetic activity of an ethanol extract obtained from the stem bark of *Psidium guajava* (Myrtaceae). Pharmazie 2006;61:725-7.
- Oh WK, Lee CH, Lee MS, Bae EY, Sohn CB, Oh H, et al. Antidiabetic effects of extracts from *Psidium guajava*. J Ethnopharmacol 2005;96: 411-5.
- 258. Wang B, Liu HC, Hong JR, Li HG, Huang CY. Effect of *Psidium guajava* leaf extract on α-glucosidase activity in small intestine of diabetic mouse. Sichuan Da Xue Xue Bao Yi Xue Ban. 2007;38:298-301.
- 259. Rai PK, Mehta S, Watal G. Hypolipidemic and hepatoprotective effects of *Psidium guajava* raw fruit peel in experimental diabetes. Indian J Med Res 2010;131:820-4.
- 260. Haranath PS, Ranganthrao K, Anjaneyulu CR, Ramnathan JD. Studies on the hypoglycemic and pharmacological actions of some stilbenes. Indian J Med Sci 1958;12:85-9.
- Chakravarty BK, Gupta S, Gambhir SS, Gode KD. Pancreatic β-cell regeneration. A novel antidiabetic mechanism of *Pterocarpus* marsupium Roxb. Indian J Pharmacol 1980:12:123-7.
- 262. Jahromi MA, Ray AB. Antihyperlipidemic effect of flavonoids from *Pterocarpus marsupium*. J Nat Prod 1993;56:989-94.
- 263. Ahmad F, Khalid P, Khan MM, Rastogi AK, Kidwai JR. Insulin like activity in (-) epicatechin. Acta diabetol Lat 1989;26:291-300.
- 264. Khosla P, Gupta DD, Nagpal RK. Effect of *Trigonella foenum* graceum (fenugreek) on blood glucose in normal and diabetic rats. Indian J Physiol Pharmacol 1995;39:173-4.
- Gupta D, Raju J, Baquer NZ. Modulation of some gluconeogenic enzyme activities in diabetic rat liver and kidney: Effect of antidiabetic compounds. Indian J Exp Biol 1999;37:196-9.
- 266. Ravikumar P, Anuradha CV. Effect of fenugreek seeds on blood lipid peroxidation and antioxidants in diabetic rats. Phytother Res 1999;13:197-201.
- Dixit P, Ghaaskadbi, S, Mohan H, Devasagyam TP. Antioxidant properties of germinated fenugreek seeds. Phytother Res 2005:19:977-83
- 268. Hamden K, Jaouadi B, Carreau S, Aouidet A, El-Fazaa S, Gharbi N, et al. Potential protective effect on key steroidogenesis and metabolic enzymes and sperm abnormalities by fenugreek steroids in testis and epididymis of surviving diabetic rats. Arch Physiol Biochem 2010;116:146-55.
- 269. Singh AB, Tamarkar AK, Narender T, Srivastava AK. Antihyperglycemic effect of an unusual amino acid (4-hydroxyisoleucine) in C57BL/KsJ-db/db mice. Nat Prod Res 2010;24:258-65.
- Hamden K, Masmoudi H, Carreau S, Elfeki A. Immunomodulatory, β-cell, and neuroprotective actions of fenugreek oil from alloxaninduced diabetes. Immunopharmacol Immunotoxicol 2010;32:437-45.
- 271. Vijayakumar MV, Pandey V, Mishra GC, Bhat MK. Hypolipidemic effect of fenugreek seeds is mediated through inhibition of fat accumulation and up regulation of LDL receptor. Obesity (Silver Spring) 2010;18:667-74.
- Reddy RL, Srinivasan K. Fenugreek seeds reduce atherogenic dietinduced cholesterol gallstone formation in experimental mice. Can J Physiol Pharmacol 2009;87:933-43.
- 273. Preet A, Siddiqui MR, Taha A, Badhai J, Hussain ME, Yadava PK, et al.

- Long-term effect of *Trigonella foenum* graecum and its combination with sodium orthovanadate in preventing histopathological and biochemical abnormalities in diabetic rat ocular tissues. Mol Cell Biochem 2006;289:137-47.
- 274. Preet A, Gupta BL, Siddiqui MR, Yadava PK, Baquer NZ. Restoration of ultra structural and biochemical changes in alloxan-induced diabetic rat sciatic nerve on treatment with Na3VO4 and Trigonella-a promising antidiabetic agent. Mol Cell Biochem 2005;278:21-31.
- 275. Amin A, Alkaabi A, Al-Falasi S, Daoud SA. Chemopreventive activities of *Trigonella foenum* graecum (Fenugreek) against breast cancer. Cell Biol Int 2005;29:687-94.
- 276. Flammang AM, Cifone MA, Erexson GL, Stankowski LF Jr. Genotoxicity testing of a fenugreek extract. Food Chem Toxicol 2004;42:1769-75.
- Stanely Mainzen Prince P, Menon VP. Antioxidant action of *Tinospora cordifolia* root extract in alloxan diabetic rats. Phytother Res 2001;15:213-8.
- 278. Price PS, Menon VP. Antioxidant activity of *Tinospora cordifolia* roots in experimental diabetes. J Ethnopharmacol 1999;65:277-81.
- 279. Mathew S, Kuttan G. Antioxidant activity of *Tinospora cordifolia* and its usefulness in the amelioration of cyclophosphamide-induced toxicity. J Exp Clin Cancer Res 1997;16:407-11.
- Stanely Mainzen Prince P, Menon VP. Antioxidant action of *Tinospora cordiflora* roots in chemical induced diabetes in rats. Phytother Res 2003:17:410-3.
- 281. Dhaliwal KS. Method and composition for treatment of diabetes. US patent 5886029, 1999.
- Gupta SS, Verma SC, Garg VP, Rai M. Antidiabetic effect of *Tinospora cordifolia* I. Effect on fasting blood sugar level, glucose tolerance and adrenaline induced hyperglycemia. Indian J Med Res 1967;55:733-45.

- 283. White JR Jr, Kramer J, Campbell RK, Bernstein R. Oral use of a topical preparation containing an extract of *Stevia rebaudiana* and the chrysanthemum flower in the management of hyperglycemia. Diabetes Care 1994;17:940.
- 284. Jeppesen PB, Gregersen S, Alstrup KK, Hermansen K. Stevioside induces antihyperglycemic, insulinotropic and glucagonostatic effects *in vivo*: Studies in the diabetic Goto-Kakizaki (GK) rats. Phytomedicine 2002;9:9-14.
- 285. Ferreira EB, de Assis Rocha Neves F, da Costa MA, do Prado WA, de Araújo Funari Ferri L, Bazotte RB. Comparative effects of *Stevia rebaudiana* leaves and stevioside on glycaemia and hepatic gluconeogenesis. Planta Med 2006;72:691-6.
- 286. Abudula R, Jeppesen PB, Rolfsen SE, Xiao J, Hermansen K. Rebaudioside A potently stimulates insulin secretion from isolated mouse islets: Studies on the dose-, glucose-, and calcium-dependency. Metabolism 2004;53:1378-81.
- Lailerd N, Saengsirisuwan V, Sloniger JA, Toskulkao C, Henriksen EJ. Effects of stevioside on glucose transport activity in insulin-sensitive and insulin-resistant rat skeletal muscle. Metabolism 2004;53:101-7.
- 288. Jeppesen PB, Gregersen S, Rolfsen SE, Jepsen M, Colombo M, Agger A, *et al.* Antihyperglycemic and blood pressure-reducing effects of stevioside in the diabetic Goto-Kakizaki rat. Metabolism 2003:52:372-8
- 289. Jeppesen PB, Gregersen S, Poulsen CR, Hermansen K. Stevioside acts directly on pancreatic β-cells to secrete insulin: actions independent of cyclic adenosine monophosphate and adenosine triphosphatesensitive K+-channel activity. Metabolism 2000;49:208-14.

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