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Review Article

Traditional Medicinal Plants Used for the Treatment of Diabetes

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Abstract

This review focuses on Indian Herbal drugs and plants used in the treatment of diabetes, especially in India. Diabetes is an important human ailment afflicting many from various walks of life in different countries. In India it is proving to be a major health problem, especially in the urban areas. Though there are various approaches to reduce the ill effects of diabetes and its secondary complications, herbal formulations are preferred due to lesser side effects and low cost. These are the some herbal drug used in the treatment of diabetes mellitus are *Emblca officinalis*, Gaertn Garlic (*Allium sativum*), Ginger (*Zingiber officinale*), *Tinospora cordifolia*, banana flower, *Abroma Augusta*, *Butea monosperma*, *Catharanthus roseus*, *Allium Cepa* (onion), *Azadirachta indica*, *Mangifera Indica* (Mango), *Aegle Marmelos* (bel), *Caesalpinia bonducella*, *Eugenia jambolana*, *Abelmoschus esculentus*, *Curcuma longa*, *Coriandrum Sativum*, *Berberis aristata*, *Helicteres isora* roots, *Murraya koenigii*, *Momordica charantia* etc.

1. INTRODUCTION

Diabetes mellitus (DM) is a metabolic disorder resulting from a defect in insulin secretion, insulin action, or both. Insulin deficiency in turn leads to chronic hyperglycemia with disturbances of carbohydrate, fat and protein metabolism¹.

Diabetes mellitus may be categorized into several types but the two major types are type 1 and type 2. On the basis of aetiology, the term type 1 and type 2 were widely used to describe IDDM (insulin dependent diabetes mellitus) and NIDDM (Non insulin dependent diabetes mellitus), respectively; other specific types of diabetes and gestational diabetes are given in the term juvenile-onset diabetes has sometimes been used for IDDM and maturity-onset for NIDDM².

1.1 Definition According to WHO

The term diabetes mellitus describes a metabolic disorder of multiple aetiology characterized by chronic hyperglycaemia with disturbances of carbohydrate, fat and protein metabolism resulting from defects in insulin secretion, insulin action, or both. The effects of diabetes mellitus include long-term damage, dysfunction and failure of various organs. Diabetes mellitus may present with characteristic symptoms such as thirst, polyuria, blurring of vision, and weight loss³.

1.2 Classification of Diabetes

Type 1 (IDDM) insulin dependent diabetes mellitus: β -cell destruction with little or no endogenous insulin secretory capacity Autoimmune Idiopathic.¹

Type 2 (NIDDM) Non insulin dependent diabetes mellitus ranges from relative insulin deficiency to disorders of insulin secretion and insulin resistance¹.

Type 3 (Gestational Diabetes): Mellitus refers to the onset or

initial recognition of glucose intolerance during pregnancy, usually in the second or third trimester. It occurs in about 4% of all pregnancies. Patients with GD have a 30% to 50% chance of developing DM, usually type 2 DM¹.

2. EPIDEMIOLOGY OF DIABETES

According to recent estimates, approximately 285 million people worldwide (6.6%) in the 20–79 year age group will have diabetes in 2010 and by 2030, 438 million people (7.8%) of the adult population, is expected to have diabetes. The largest increases will take place in the regions dominated by developing economies.^[4]

The WHO reports suggests that the prevalence of diabetes in adults worldwide would increase to 300 million in years 2025. It is the one of the main threats to human health in the 21st century and is the fifth leading cause of deaths in most developed countries. There are an estimated 86 million person with diabetes in Asia out of which around 25-30 millions are in India. High prevalence is increasing all over the world. By the 2025, more than 75% of individuals with diabetes will reside in developing countries⁵.

3. SOME MEDICINAL PLANT USED TO TREAT DIABETES MELLITUS

3.1 *Emblca officinalis* Gaertn

Commonly known as amla or Indian gooseberry is known for its medicinal and therapeutic properties from ancient time in India and considered as a wonder fruit for health conscious population. It is extensively found throughout India and some other Asian countries⁶.

3.1.1. Chemical Constituents

Tannins 30%, phyllembin (2.4%), phyllembic acid (6.3%), gallic acid (1.32%), ellagic acid in natural form and cytokine like substances identified as Zeatin, Z riboside, Z nucleotide etc⁷.

2.1.2. Mechanism of Action

Amla contains highest amount of Vitamin C (ascorbic acid), low and high molecular weight tannins 30%, phyllembin (2.4%), phyllembic acid (6.3%), gallic acid (1.32%), ellagic acid in natural form and

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cytokine like substances identified as Zeatin, Z riboside, Z nucleotide. Amla fruit ash contains chromium, 2.5; zinc, 4; and copper, 3 ppm. Presence of chromium is of therapeutic value in diabetes⁸. The fruit contains 482.14 units of superoxide dismutase/g fresh weight, and exhibited anti senescent activity. Chromium, a trace element possesses significant anti diabetic activity in various experimental models of diabetic mellitus. Chromium compounds also improved deranged lipid metabolism of both type 1 and type 2 diabetic rats. It has been reported that insulin derived with chromium is capable of reversing blood sugar, serum cholesterol and phospholipids levels to those of normal rats⁶.

3.1.3 Uses

The fruits are sour, astringent, bitter, acrid, sweet, cooling, anodyne, ophthalmic, carminative, digestive, stomachic, laxative, alterant, aphrodisiac, rejuvenative, diuretic, antipyretic and tonic. They are useful in vitiated conditions of tridosha, diabetes, cough, asthma, bronchitis,¹⁷ cephalalgia, ophthalmopathy, dyspepsia, colic, flatulence, hyperacidity, peptic ulcer, erysipelas, skin diseases, leprosy, haematogenesis, inflammations, anemia, emaciation, hepatopathy, jaundice, strangury, diarrhoea, dysentery, hemorrhages, leucorrhoea, menorrhagia, cardiac disorders, intermittent fevers and graying of hair⁸.

3.2 Garlic (*Allium sativum*)

Garlic (*Allium sativum*) is one of the most popular herbs Used worldwide to reduce various risk factors associated with cardiovascular diseases⁹.

3.2.1 Chemical constituents

Sulphur-containing compounds (e.g. S-allylcysteine sulphoxide) etc¹⁰.

3.2.2 Mechanism of Action

Most of the studies showed that garlic can reduce blood glucose levels in diabetic mice, rats and rabbits.⁹ Augusti and Sheela consistently showed that S-allyl cysteine sulphoxide, (allicin), a sulphur-containing amino acid in garlic (200 mg/kg body weight), had a potential to reduce the diabetic condition in rats almost to the same extent as did glibenclamide and insulin⁹.

3.2.3 Uses

Anticoagulant (anti-thrombotic), antioxidant, antibiotic, hypocholesterolaemic, hypoglycemic, as well as hypertensive activity^{9,10}.

3.3 Ginger (*Zingiber officinale*)

The *Zingiber officinale* is a hypoglycemic drug that has less or no side effects at all. Ginger exhibits hypoglycemic activity in both normal and diabetic rats. They further reported that ginger contains magnesium, calcium and phosphorus which play important roles in bone formation, curbing muscle spasm, depression, hypertension, convulsion, nausea, gastrointestinal disorders, paralysis, kidney damage and several other bio-functions necessary for keeping body in homeostatic condition.¹¹

3.3.1 Chemical Constituents

6-Gingerol, tannins, polyphenolic compounds, flavonoids and triterpenoids etc^{11,12}.

3.3.2 Mechanism of Action

It has been demonstrated that flavonoid compound act against diabetes mellitus either through their capacity to avoid glucose absorption or improve glucose tolerance. In vitro studies have shown that a soyabean extract containing the isoflavones genistein daidzein inhibits glucose absorption into the intestinal brush border membrane vesicles of rat or mice. So it has demonstrated that flavonoids can act per as insulin secretagogues or insulin mimetics¹².

3.3.3 Uses

Ginger contains magnesium, calcium and phosphorus which play important roles in bone formation, curbing muscle spasm, depression, hypertension, convulsion, nausea, gastrointestinal disorders, paralysis, kidney damage and several other bio-functions necessary for keeping body in homeostatic condition^{11,12}.

3.4 *Tinospora cordifolia* (TC)

Tinospora cordifolia (wild) Miers Ex Hook F. and Thoms.TC belong to the family Menispermaceae and is known as Gulancha in English, Guduchi in Sanskrit, and Giloya in hindi¹³.

3.4.1 Chemical Constituents

Flavonoids, tannins etc¹³.

3.4.2 Mechanism of Action

Mechanism of action of *Tinospora cordifolia* is same as mentioned under 3.3.2.

3.4.3 Uses

As tonic and vitalizer and as remedy for diabetes and other metabolic disorder^{12,13}.

3.5 Banana Flower (*Musa sapientum*)

Commonly known as 'banana' is widely used in Indian folk medicine for the treatment of diabetes mellitus. The available literature confirms that flavonoids are present in banana flowers. The chloroform, water and ethanol extract of *Musa sapientum* flowers were found to exhibit hypoglycemic activity in alloxan diabetic rat¹⁴.

3.5.1 Chemical Constituents

Bio flavonoids (e.g. epicatechin and galocatechin), are present in banana flower etc¹⁵.

3.5.2 Mechanism of Action

The chloroform, water and ethanol extract of *Musa sapientum* flowers were found to exhibit hypoglycemic activity in alloxan diabetic rat intraperitoneal administration of prunin produces a significant hypoglycemic effect in diabetic rats. Bio-flavonoids are well-known for their multi-directional biological activities including anti-diabetic efficacy¹⁴.

3.5.3 Use:

As an anti diabetes agent and as insulin secretagogues, insulin mimetics^{14,15}.

3.6 *Augusta abroma*

Abrinoma Augusta have hypoglycemic effect on alloxan induced diabetic rats. *Syzygium cumi* have alpha glycosidase inhibitory activity .In folk medicine *Syzygium cumini* were used in treating diabetes diarrhoea and ringworm And the leaves of *Syzygium cumini* are antibacterial and are used for strengthening the teeth and gums¹⁶.

3.6.1 Chemical Constituents

Syzygium cumini is the active constituents in *abroma augusta* etc¹⁷.

3.6.2 Mechanism of Action

Syzygium cumini have alpha glycosidase inhibitory activity. In folk medicine *Syzygium cumini* were used in treating diabetes diarrhea and ringworm And the leaves of *Syzygium cumini* are antibacterial and are used for strengthening the teeth and gums¹⁷.

3.6.3 Use

Treating diabetes, diarrhea, ringworm, antibacterial and used as strengthening the teeth and gums^{16,17}.

3.7 *Butea monosperma*

Butea monosperma belongs to family Fabaceae. In English name is bastard teak. Its vernacular name is parasa and it is commonly known as palaspara¹⁸.

3.7.1 Chemical Constituents

Butrin, butein, 7 trihydroxy flavones, palasonin etc^{18,19}.

3.7.2 Mechanism of Action

Because the presence of butrin, butein, 7 trihydroxy flavones *butea monosperma* shown antidiabetic activity¹⁹.

3.7.3 Use

It is use as antidiabetic agent²⁰.

3.8 *Catharanthus roseus*

There are a number of plants to control the blood glucose level one of them is *Nayantara* (*Catharanthus roseus*)²¹.

3.8.1 Chemical Constituents

Catharanthin, catharanthus, catharin etc^{21,22}.

3.8.2 Mechanism of Action

Due to the presence of catharanthin or catharin it show anti diabetes activity because catharanthin reduce blood glucose level in the albino wister rat and swiss albino mice²².

3.8.3 Use:

It is use as an anti-diarrheal, anti-diabetes agent²¹.

3.9 Allium Cepa (Onion)

Allium cepa belongs to the family Liliaceae and is probably native of south west Asia and is widely cultivated throughout the world,^[24]. *Allium cepa* (onion) dried powder shown anti hyperglycemic activity in diabetic rat. *Allium cepa* is also know to have antioxidant and hypolipidaemic activity²³.

3.9.1 Chemical Constituents

Sulfar containing amino acid s methyl cysteine sulphoxide etc.^[23, 24]

3.9.2 Mechanism of Action

Due to the presence of sulfur containing amino acid s methyl cysteine sulphoxide it control blood glucose as well as lipids in serum and tissues and normalized the activity of liver hexokinase glucose6 phosphate²⁴.

3.9.3 Use

It is use as anti diabetic, antioxidant, hypolipidaemic. Modern uses is to lower Blood pressure, antiseptic, hypoglycaemic and hypocholesterlemic properties^{23, 24}.

3.10 Azadirachta indica (Neem)

Hydro alcoholic extracts of this plant showed anti-hyperglycemic activity²⁵.

3.10.1 Chemical Constituents

Azadirachtin is the main active chemical constituents in the neem due to the presence of azadirachtin^{25,26}.

3.10.2 Mechanism of Action

Hydroalcoholic extracts of this plant showed anti-hyperglycemic activity in streptozotocin treated rats and this effect is because of increase in glucose uptake and glycogen deposition in isolated rat hemidiaphragm.^[26]

Use

Anti-bacterial, antimalarial, antifertility, hepatoprotective and antioxidant, anti diabetes²⁶.

3.11 Mangifera Indica (Mango)

The leaves of this plant are used as an antidiabetic agent in Nigerian folk medicine, although when aqueous extract given orally did not alter blood glucose level in either normoglycemic or streptozotocin induced diabetic rats^{27, 28}.

3.11.1 Chemical Constituents

Mangiferin, mangifera etc²⁷.

3.11.2 Mechanism of Action

Anti diabetic activity was seen when the extract and glucose were administered simultaneously and also when the extract was given to the rats 60 min before the glucose. The results indicate that aqueous extract of *Mangifera indica* possess hypoglycemic activity²⁸.

3.11.3 Use

Anti-diabetic agent²⁷.

3.12 Aegle Marmelos (Bel)

Administration of aqueous extract of aegle marmelos (bel) leaves improves digestion and reduces blood sugar and urea, serum cholesterol in alloxanized rats as compared to control.^[29]

3.12.1 Chemical Constituents

Flavonoids, tannins etc³⁰.

3.12.2 Mechanism of Action

Administration of aqueous extract of leaves improves digestion and reduces blood sugar and urea, serum cholesterol in alloxanized rats as compared to control. Along with exhibiting hypoglycemic activity, this extract also prevented peak rise in blood sugar at 1h in oral glucose tolerance test^{29,30}.

3.12.3 Use

As anti-oxidant and anti-diabetic agent³⁰.

3.13 Caesalpinia bonducella

Caesalpinia bonducella is widely distributed throughout the coastal region of India and used ethnically by the tribal people of India for controlling blood sugar. Both the aqueous and ethanolic extracts showed potent hypoglycemic activity in chronic type II diabetic models. These extracts also increased glycogenesis thereby increasing liver glycogen content. Two fractions BM 169 and BM 170 B could increase secretion of insulin from isolated islets.^[31]

3.13.1 Chemical Constituents

Flavonoids, is the active chemical constituents in the *Caesalpinia bonducella* etc^{31,32}.

3.13.2 Mechanism of Action

Due to the presence of flavonoid it shown anti diabetes activity by decreasing glucose absorption³².

3.13.3 Use

Antidiabetic as well as antihyperlipidemic³¹.

3.14 Eugenia jambolana (Indian gooseberry jamun)

In India decoction of kernels of *Eugenia jambolana* is used as household remedy for diabetes. This also forms a major constituent of many herbal formulations for diabetes³³.

3.14.1 Chemical Constituents

Tri-terpenoids, tannins, gallic acid, and oxalic acid were the chemical constituents detected in *Eugenia jambolana* seed etc³⁴.

3.14.2 Mechanism of Action

In India decoction of kernels of *Eugenia jambolana* is used as household remedy for diabetes. This also forms a major constituent of many herbal formulations for diabetes. Antihyperglycemic effect of aqueous and alcoholic extract as well as lyophilized powder shows reduction in blood glucose level. The extract of jamun pulp showed the hypoglycemic activity in streptozotocin induced diabetic mice within 30 min of administration while the seed of the same fruit required 24 h. The oral administration of the extract resulted in increase in serum insulin levels in diabetic rats.^[33,34]

3.13.3 Use

These extracts also inhibited insulinase activity from liver and kidney³³.

3.15 Abelmoschus esculentus

Abelmoschus esculentus Linn. is a plant of family Mallow (Malvaceae). It is Naturalized in all tropical countries and grown abundantly throughout India. The plant prefers acid, neutral and basic (alkaline) soils and can grow in very alkaline soil. Stem is erect, aerial, herbaceous or woody, usually solid, cylindrical and branched. Herbaceous portion of stem is covered with scaly hairs, woody part is fibrous. Leaves are Alternate and stipulate and its fruit is loculicidal Capsule³⁵.

3.15.1 Chemical Constituents

Carbohydrate, gums and mucilages, proteins, phytosterols, flavonoids, tannins, phenolic compounds and volatile oil etc³⁶.

3.15.2 Mechanism of Action

Due to the presence of flavonoids and phenolic compound it show anti diabetes activity by lowering blood glucose level^{35,36}.

3.15.3 Uses

Used traditionally as emollient, demulcent, diuretic, cooling, aphrodisiac, Antiseptic and in gonorrhoea³⁷.

3.16 *Curcuma longa*

Curcumin (diferuloylmethane) is a naturally occurring yellow pigment isolated from the rhizomes of the plant *Curcuma longa* (Linn) found in south Asia and is a potent antioxidant agent and free radical scavenger³⁸.

3.16.1 Chemical Constituents

Curcumin, bio flavonoid etc³⁸.

3.16.2 Mechanism of Action

Due to the presence of curcumin and flavonoid it show anti diabetes activity by lowering the blood glucose level³⁹.

3.16.3 Use

As anti-diabetic and antioxidants^{38,39}.

3.17 *Coriandrum Sativum*

Coriander is an important culinary herb. The fruits and the fresh leaves are widely used for flavouring food and the root can be cooked and eaten as a vegetable⁴⁰.

3.17.1 Chemical constituents:

Coriander, sataverin, tannins, flavonoid etc⁴¹.

3.17.2 Mechanism of Action

The present study was designed to investigate clinically the hypoglycemic effect of *Coriandrum sativum* in Type-2 diabetes mellitus⁴⁰. After assaying fasting plasma and urinary glucose, 10 patients of type-2 diabetes mellitus with no previous medication, 10 patients of type-2 diabetes mellitus taking oral hypoglycemic agents with history of inadequate control and six control subjects were given low (2.5 g tid) and high (4.5 g tid) doses of powdered part, aqueous extract and alcoholic extract of *Coriandrum sativum* for 14 days. On 15th day blood and urine samples for glucose were taken. Based on results obtained it was found that *Coriandrum sativum* has significant hypoglycemic activity in high dose and can be successfully combined with oral hypoglycemic agents in type-2 Diabetic patients whose diabetes is not controlled by these agents⁴¹.

3.17.3 Use

As anti-diabetic agent⁴⁰.

3.18. *Berberis aristata*

Berberis aristata DC, known as 'Daruharidra' in Ayurvedic system of medicine, is extensively used in various systems of indigenous medicine for treating a variety of ailments such as eye and ear diseases, rheumatism, jaundice, diabetes, stomach disorders, skin disease, malarial fever and as tonic etc⁴².

3.18.1 Chemical Constituents

Berberine, berbamine, aromoline, karachine, palmatine, oxyacanthine and oxyberberine etc⁴³.

3.18.2 Mechanism of Action

Due to the presence of berberine and aromoline it show anti-diabetic activity⁴².

3.18.3 Use

AS an antioxidant and anti-diabetic agent^{42,43}.

3.19. *Helicteres isora* root

The root of *helicteres isora* root used in the treatment of diabetes aqueous ethanol and butanol extracts had shown significant protection and lowered the blood glucose levels⁴⁴.

3.19.1 Chemical Constituents

Tannins, flavonoid, helicterin etc⁴⁵.

3.19.2 Mechanism of Action

It has been demonstrated that flavonoid compound act against diabetes mellitus either through their capacity to avoid glucose absorption or improve glucose tolerance.^[13]

3.19.3 Use

Used in the treatment of type2 diabetes.^[46]

3.20. *Murraya koenigii*

Leaves of *Murraya koenigii* (Rutaceae) are used traditionally in Indian Ayurvedic system to treat diabetes. The purpose of the study is to investigate the effect of mahanimbine carbazole alkaloid from *Murraya koenigii* leaves on blood glucose and serum lipid profiles on streptozotocin-induced diabetic rats.⁴⁷

3.20.1 Chemical Constituents

Mahanimbine, flavonoid, alkaloids^{48,49}.

3.20.2 Use

Used as anti-diabetic agent⁴⁸.

3.21 *Momordica charantia*

M. *Charantia* bitter Melon, Better gourd is a flowering vine in the family of cucurbitaceae⁵⁰. It is tropical plant that is widely cultivated in asia in India east Africa and south America. Used for the treatment of diabetes and used in cooking⁵¹.

3.21.1. Chemical Constituents

Saponins, glycosides, alkaloids, reducing sugar resins, fixed oil.^[52]

3.21.2. Uses

Antidiabetic, anticancer, anti inflammation, antiviral, antioxidants and cholesterol lowering effect.^[53]

4. CONCLUSION

The present study revealed that the administration of these plants *Emblica officinalis*, *Gaertn Garlic (Allium sativum)*, *Ginger (Zingiber officinale)*, *Tinospora cordifolia*, *banana flower*, *Abroma Augusta*, *Butea monosperma*, *Catharanthus roseus*, *Allium Cepa (onion)*, *Azadirachta indica*, *Mangifera Indica (Mango)*, *Aegle Marmelos (bel)*, *Caesalpinia bonducella*, *Eugenia jambolana*, *Abelmoschus esculentus*, *Curcuma longa*, *Coriandrum Sativum*, *Berberis aristata*, *Helicteres isora roots*, *Murrayc koenigii*, *Momordica charantia* etc. Extract showed antidiabetes effects in alloxan and streptozotocin (stz) induced diabetes rats, therefore it was concluded that these plant extract is helpful to lower glucose level in treatment of diabetes patients.

REFERENCES

1. Bastaki S "Review of Diabetes mellitus and its treatment" International journal of Diabetes and Metabolism, 2005, (13):111-134.
2. Riaz S "Review of Diabetes mellitus" Academic Journals, 2009, 4 (5): 367-373.
3. WHO Expert Committee on Diabetes Mellitus. Second Report. Geneva: WHO, 1980. Technical Report Series 2000, (15):646.
4. Ramachandran A, Das AK, et al "Current Status of Diabetes in India and need for Novel Therapeutic Agents Supplement to JAPI" 2010, (7):7-8.
5. World Health Organization. Diabetes Mellitus: Report of a WHO Study Group. Geneva: WHO Technical Report Series 727,1985,(9)
6. Tigar P R, Shah K, et al "Research journal of pharmaceutical biological and chemical sciences investigation into mechanism of action of Antidiabetic activity of *Emblica officinalis* on stz induce type 1 diabetic rat " 2010, (17):672.
7. Khurana SC, Gupta SK, et al "Study of pharmacodynamic properties of *emblica officinalis*" Ind. J. Physiol. Pharmacol,1970,(14) : 39
8. Satyavati G V, et al "Medicinal Plants of India", ICMR, (1):377.
9. Thomson M, amin-al Zainab M, Antidiabetic and Hypolipidaemic properties of garlic in streptozotocin induced diabetic rat, department of biological science faculty of science Kuwait University, Kuwait, 2007, (31) :108-115.
10. Ried K, Frank OR, et al "Effect of garlic on blood pressure" A systematic review and meta-analysis, 2008, (5):8-13.
11. Jafri.Ahmad S, Abbas S, et al, Institute of molecular biology, The University of Lahore. Hypoglycemic Effect of *Zingiber Officinale* in Alloxan induce diabetic rat. 2011, (5):160-162.

12. Brahmachari G, "Bio-flavonoids with promising antidiabetic potentials" 2001, (3):187-212.
13. Puranik N, Fakruddin KK, et al "Antidiabetic activity of *Tinospora Cord folia* in streptozotocin induced diabetic rat" 2010, (17) :265-270.
14. Ganugapati j, Balwda A, et al, Molecular docking studies of banana flower Flavonoids as insulin receptor tyrosine kinase activators as a cure for diabetes mellitus, March 17-2012, (10) :216-220.
15. Lima GPP, et al , Int J Food Sci Technol ,2008,(43):18-38.
16. Nahar L, Ripa Alam F, et al , Comparative study of antidiabetic effect of *Abroma agusta* and *Syzygium cumini* on alloxan induce diabetic rat. Lecture Department of pharmacy, south east university bamani, Dhaka 1231, 2010, (13):1268-1272.
17. Hussain M E M A, et al ,Preliminary studies on the hypoglycemic effect of *Abroma Augusta* in alloxan diabetic rats, Indian Journal of Clinical Biochemistry, 2001, (16):77-80.
18. Naeem f, Hassan khan S, Department of home economics division of education and extension university of agriculture Faisalabad Pakistan evaluation of hypoglycemic and hypolipidemic activity of *Butea monosperma* fruit in diabetic human subject. 2010, (21):189-197.
19. Mentreddy RS " Medicinal plant species with potential antidiabetic properties" J Sci of Food and Agric 2007, (87):743-750.
20. Mostofa M, Choudhury ME, et al , Department of pharmacology, Faculty of veterinary science, Bangladesh. 2007, (24):99-102.
21. Bahajiri SM, Mirza SA, et al, The effects of inorganic chromium and brewer's yeast supplementation on glucose tolerance, serum lipids and drug dosage in individuals with type-2 diabetes. Saudi Med, 2000, 21(9):831-837.
22. Mathew PT, Augusti KT "Hypoglycemic effects of onion, *Allium cepa* Linn. on diabetes mellitus- a preliminary report" Ind. J. Physiol. Pharmacol, 1975, (1):213-217.
23. Jevas C, Zougwu O, Physiology and biomedical research unit department of Zoology, university of Nigeria Nsukka, Enugu State. antidiabetic activity of *Allium cepa* Linn. on aqueous extracts on alloxan-induced diabetic rat, 5(7), 4 April 2011 (6):1134-1139.
24. Biswas K, Chattopadhyay I, et al "Biological activities and medicinal properties of neem (*Azadiracta indica*) Curr" Sci. 2002, (19) :1336-1345.
25. Banerjee RK, Ghosh KK "Antidiabetic activity of neem (*Azadiracta indica*), 2005, (21):12-15
26. Aderibigbe AO, Emudianghe TS, et al "Antihyperglycemic effect of *Mangifera indica* in rat" Phytother Res. 1999, (11):504-507.
27. Bhowmik A, Khan CA , antidiabetic activity of the stem barks and Leaves on non diabetic type1 and type2 diabetic rat models , Department of biochemistry and molecular biology jahangirnagar university savar Dhaka, department of pharmacology, 2009 , (13):110-114.
28. Karunanayake EH, Welihinda J, et al " Oral hypoglycemic activity of some medicinal plants of Sri Lanka" J. Ethnopharmacol, 1984, (9):223-231.
29. Sabu MC, Kuttan R, Antidiabetic activity of *Aegle Marmelos* and its relationship with antioxidant properties. Amala Cancer Research Center Amala Nagar, 2004, (19):81-88.
30. Kannur DM, Hukkeri VI, et al "Antidiabetic activity of *Caesalpinia bonducella* seed extracts in rats" Fitoterapia. In press, 2003, (21):17- 21.
31. Vats V, Grover JK, et al (2002). Evaluation of anti-hyperglycemic and hypoglycemic effect of *Caesalpinia bonducella* root bark in mice, 2006, (79):31-36.
32. Acherekar S, Kaklij GS, et al, Hypoglycemic activity of *Eugenia jambolana* and *ficus bengalensis*: mechanism of action , In vivo, 1991, (2):143-147.
33. Sridhar SB, Sheetal UD, et al , Preclinical evaluation of the antidiabetic effect of *Eugenia jambolana* seed powder in streptozotocin-diabetic rats, Department of pharmacology Kasturba medical college : Shastri Pharmaceutical , Mangalore India, 2005, (12):463-468.
34. Saha D, Jain B , et al , Phytochemical evaluation and characterization of various extract of *Abelmoschus Esculentas* Linn Fruit, School of pharmacy Lalkhadan Masturi Road , Bilaspur , 2011, (5) :183-185.
35. Ghosh R, Sharatchandra Kh, et al "Dept. of Pharmacology, GSL Medical College, laxmipuram" Indian Journal of Pharmacology, 2004, 36(4): 222 – 225.
36. Azza A, El Massy, Potential Therapeutic Effect of *Curcuma longa* on Streptozotocin Induced Diabetic rats, department of Zoology, faculty of science, Alexandria University, Egypt, 2012, (21) :91-98.
37. Babu PS, Srinivasan K, Influence of dietary curcumin and cholesterol on the progression of experimentally induced diabetes in albino rat. Mol. Cell. Biochem, 1995, 152: 13-21.
38. Waheed A, Miana GA, et al , Clinical investigation of hypoglycemic effect of *coriandrum sativum* on type two diabetes, department of pharmacology and therapeutics, Army Medical collage Rawalpindi, jan 2006, (9) :7-11.
39. Ahmed M, Ismail N, et al, Pharmacognostic profile of *Trigonella* seed and its hypoglycaemic activity. Natural Product Sciences, (1) :25-30.
40. Gupta JK, Mishra P, et al " Blood Glucose Lowering Potential of Stem Bark of *Berberis aristata* Dc in Alloxan-Induced Diabetic Rats" November 5, 2009, (9):21-24.
41. Rastogi RP, Mehrotra BN, Compendium of Indian Medicinal Plants(3), CDRI, Lucknow and PID CSIR, New Delhi; 199.2006 , (6) :11-15
42. Sama V, Reddy GD, et al, Department of pharmacognosy G. Pulla Reddy collage of pharmacy Mehdipatnam Hyderabad , Andhra Pradesh, 2011, (17) :105-108.
43. Singh SB, Singh AK, et al "Chemical Constituents of the leaves of *Helicteres isora*" Indian Journal of Pharmaceutical Sciences, 1987, (4):148-149.
44. Bean MF, Antoun M, et al "Cucurbitacin B and Isocucurbitacin B Cytotoxic components of *Helicteres isora*" Journal of Natural Product, 1985, (3): 500-503.
45. Dineshkumar B, Mitra A, et al "Antidiabetic effect of Mahanimbine from *Murraya koenigii* Leaves" International Journal of Phytomedicine, 2010, (7) :22-30
46. Kong YC, Ng KH, et al "Sources of the anti-implantation alkaloid yuehchukene in the genus *Murraya*" J Ethnopharmacol, 1986, (31), 15: 195-200.
47. Keasri AN, Kesari S, et al " Studies on the glycemic and lipidemic effect of *Murraya koenigii* in experimental animals" J Ethnopharmacol, 2007 , (2), 112: 305-311.
48. Abascal K, Yarnell E "Using bitter melon to treat diabetes" J Altern Complement Med, 2005 (1) : 179-184.
49. Lee SY, Eom SH, et al "Cucurbitane-type triterpenoids in *Momocharantia* Linn" J med Plnats Res, 2009, (13): 1264-1269.
50. Liu J, Chen J, et al, New cucurbitane triterpenoids and steroidal glycoside from *Momordica charantia*, Molecules, 2009, (14):4804-4813.
51. Budrat P, Shotipruk A "Extraction of phenolic compounds from fruits of bitter melon (*Momordica charantia*) with subcritical water extraction and antioxidant activities of these extracts" Chiang Mai J Sci, 2008, (13):123-130.
52. John JK, Simon P W, et al, Bitter gourd: Botany, horticulture, breeding. Horticulture Rev 2010, (37):101-141.
53. Wang C, John JK "Antidiabetic activity of methanolic extract of *Momordica Charantia* sci" 2009, (29) :131-141