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PHYTOCHEMICAL EVALUATION OF INDIGOFERA ASTRAGALINA

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ABSTRACT

The word diabetes is Greek for a draw off, referring to the ejection of a more quantity of urine; and mellitus is Latin used for sugar. Consequently diabetes mellitus means the passage of huge amounts of sweet urine. This is derived from the information that excess glucose in the blood spills over into the urine, absorbing fluids with it. Based on the exhaustive literature survey, the objective of the present work focussed on the search of herb to be proved as more suitable among the reported herbs/ herbal preparations/claimed herbs, towards the control of blood glucose level in hyperglycemic condition. In the present investigation herbs; *Indigofera astragalina* were considered based on their literature information. In general, there is very little biological knowledge on the specific modes of action in the treatment of diabetes, but most of the plants have been found to contain substances like phenols, glycosides, alkaloids, terpenoids, flavonoids etc., that are normally concerned as having antidiabetic effect.

Keywords: Antidiabetic activity, alkaloids, Flavonoid, Phenol and Indigofera Astragalina.

INTRODUCTION

Diabetes is a costly disease for the health care sector, at communal and at individual level. Expenditure of diabetes care is extremely high. The cost of concern increases a lot of folds when complications occur or when access to hospital, operation or insulin treatment is needed. A study by the authors has shown that the yearly median expenses by patients on diabetes care are Rs 10,000 in city and Rs 6,300 in rustic areas [1]. A low-income person spends nearly 25–35% of their yearly income on diabetes concern. Due to the high financial burden on the patients as well as their families, people are likely to neglect health care causing severe morbidities and early death.

Insulin biosynthesis occurs in rough endoplasmic reticulum from a single-chain precursor, preproinsulin, with a molecular weight of 11,500 with containing 109 amino acids. This molecule consists of proinsulin in addition a hydrophobic extension of 23 amino acids (preregion) on the N terminus of proinsulin. Proinsulin is claved to form equimolar amounts of C-peptide and insulin. Insulin is a protein composed of fifty one amino acids in two chains (A and B chains), linked by two disulfide bonds. Insulin is synthesized and accumulated in the β -cells of the islets of langerhans, which are situated in the pancreas. The normal pancreas contains 200 U insulin, and a basal amount of insulin is secreted always at a rate of roughly 0.5 to 1.0 U/h.

Diabetes mellitus emerges in two diversities, everyone with its individual cause: diabetes mellitus type I (previously known as juvenile onset diabetes), caused by lack of the pancreatic hormone insulin (whose principal function is to encourage the entry of glucose into cells); and diabetes mellitus type II (previously known as maturity onset diabetes), in which insulin is accessible but cannot be

appropriately utilized (The expert committee on the diagnosis and classification of diabetes

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mellitus, 2002). The third group consists of other less common types of diabetes that are caused or associated with certain specific conditions and/or syndromes. The very last group comprises diabetes diagnosed during pregnancy, called gestational diabetes (GDM) [3]. Long before the birth of orthodox Western medicine, medicinal herbs were applied to treat a wide range of disease categories [3]. Due to emphasis on scientism and other complicated reasons, Western medicine now prevails over "traditional" forms of medicine including herbal medicine systems. The use of a medicinal herb, alone or in combination with other herbs, can be thought of as a type of combination therapy because of the complexity of the phytochemicals and bioactivities in the plant. Thus, a single antidiabetic herb [4] with thousands of phytochemicals may have multiple benefits by targeting several metabolic pathways and essentially "killing several birds with one stone."

MATERIALS AND METHODS

Collection and Preparation of Plant Extract:

Drug discovery from medicinal plants has evolved to include numerous fields of inquiry and various methods of analysis. The process typically begins with a botanist, ethno botanist, ethnopharmacologist, or plant ecologist [5] who identifies the plant of interest. Collection may involve species with known biological activity for which active compound(s) have not been isolated (e.g., traditionally used herbal remedies) or may involve taxon collected randomly for a large screening program. On the basis of intensive literature survey; *I. astragalina* was selected for present study [6].

Ash Values:

The residues remaining after incineration is the ash content of the fruits powder. Ash values

are helpful in determining the quality and purity of crude drug, especially in the powdered form [7]. It usually represents the inorganic salts naturally occurring in the drug and adhering to it, but it may also include inorganic matter added for the purpose of adulteration.

Determination of total ash value:

Accurately weighed about 3 gm of air dried powdered drug was taken in a tared silica crucible and incinerated by gradually increasing the temperature to make it dull red hot until free from carbon. Cooled and weighed, repeated for constant value. Then the percentage of total ash was calculated with reference to the air dried drug [8].

Extraction and fractionation:

The extraction yield of the extracts from plant species is vastly depends on the solvent polarity, which find out both qualitatively and quantitatively the extracted compounds. Ethanol and water are the commonly used solvent for the extraction because of their low toxicity and high extraction yield with the advantage of modulating the polarity of the solvent by using mixtures at different ratios [9-11] The plant materials (1 kg) were initially defatted with petroleum ether and then extracted with alcohol and water using a Soxhlet apparatus. The yield of the plant extracts ethanol (95%) and aqueous measured about 20 g each after evaporating the solvent using water bath. The standard extracts obtained from Indigofera astragalin were then stored in a refrigerator at 4°C [12].

Phytochemical screening

Qualitative examination of phytoconstituents:

Extract has been tested for phytoconstituents present in extract as per procedure,

Preliminary phytochemical study of the I. extracts/fractions [13]. astragalin The medicinal plants are useful for healing as well as for curing of human diseases because of the presence of phytochemical constituents. The preliminary phytochemical screening was carried out to assess the gualitative chemical composition of crude extracts and fractions from I. astragalin by using precipitation and coloration reaction to identify the major natural chemical groups [14-18] General reactions in this analysis revealed the presence or absence of these compounds in the crude extracts and fractions tested.

CONCLUSION

The present investigations concluded that the

ethanolic and aqueous extracts of aerial parts of I. astragalin endowed with potential antidiabetic activity which could be attributed by their possible multiple effects on both pancreatic and extra-pancreatic site by influencing either the metabolism and/or absorption of glucose, which in turn also influence the lipid metabolism. Conversely the extracts and fractions of both plants exert very good potentials to scavenge toxic free radicals along with the inhibition of the liver lipid peroxidation products and activation of the enzymatic antioxidant defense mechanism in diabetic rats that might be due to the presence of high levels of sterols, phenolics, alkaloids and flavonoids, which may be responsible for the supporting properties of the extracts and fractions for their hypoglycaemic and antidiabetic activity. Stigmasterol, a plant sterol isolated from chloroform fraction of I. astragalin.

Chemical constituents	Chemical Test	Extracts/Fractions				
		Ethanol extract	Aqueous extracts	Chloroform fraction	Aqueous fraction	
Alkaloids	Mayer's	+	+	+	+	
	Dragendorff's	+	+	+	+	
	Wagner's	-	-	-	-	
	Hager's	+	+	+	+	
saponin	Foam	-	+	-	-	
	Haemolytic	-	-	-	-	
Phenolic compounds and Tannins	Ferric Chloride	+	+	+	+	
	Gelatin	-	-	-		
	Lead acetate test	+	+	+	+	
Proteins	Million's	+	-	+	-	
	Biuret	+	+	+	-	
	Xanthoprotein	-	-	-	-	

Table-1: Phytochemical screening of extracts and fractions of *I. astragalin*

Flavonoids	Ferric Chloride	+	+	-	+
	Shinoda	-	-	-	-
	Lead Acetate	+	+	+	+
Glycoside	Baljet's	-	-	-	-
	Legal's	-	-	-	-
	Borntrager's	-	-	-	-
	Killer killani	-	-	-	-
Fixed oil	Spot	-	-	-	-
Carbohydrate	Molisch's	-	-	-	-
	Fehling's	+	+	+	+
	Benedict's	-	-	-	-
	Barfoed's	+	+	+	+
	Cobalt-chloride	-	-	-	
Gums and mucilage	Swelling Index	-	-	-	-
Amino Acids	Ninhydrin	-	-	-	-
	Tyrosin	-	-	-	-
	Tryptophan	-	-	-	-
Sterols and triterpenes	Liebermann- Burchard's	+	+	+	-
	Salkowski's	+	-	+	-

Key (+) = Presence, (-) = Absent

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