

In Vitro Antidiabetic Activity of *Cardiospermum Halicacabum* leaves Extracts

Stalin¹

¹ Raos college of pharmacy

Received: 7 December 2012 Accepted: 5 January 2013 Published: 15 January 2013

Abstract

The present study was designed to investigate the glucose uptake of (antidiabetic activity) crude n-hexane, ethanol, methanol and aqueous leaf extracts of *Cardiospermum Halicacabum*. Methods: of *Cardiospermum Halicacabum* leaf extracts were subjected to inhibitory effect of glucose utilization using specific standard in vitro procedure. Results: results in four different leaf extracts revealed that, the methanol extract at a concentration of 50g plant extract/l was found to be more potent than other extracts with the lowest mean glucose concentration of 201 ± 1.69 mg/dl at the end of 27 hrs. Conclusions: The present findings suggest that, the methanolic extract showed a significant inhibitory effect on glucose diffusion in vitro thus validating the traditional claim of the plant.

Index terms— *cardiospermum halicacabum*, antidiabetic activity, glucose diffusion method.

1 Introduction

The plant *Cardiospermum halicacabum* Linn.

(Sapindaceae) is an annual or sometimes perennial climber, commonly found as a weed throughout India. The tender, young shoots are used as a vegetable, fodder, diuretic, stomachic, and a rubefacient [5,6]. It is used in rheumatism, lumbago, nervous diseases, and as a demulcent in orchitis and in dropsy. In Sri Lanka, it is used for the treatment of skeletal fractures. The juice of the herb is used to cure ear-ache and to reduce hardened tumours [7]. It exhibits significant analgesic, anti-inflammatory and vaso-depressant activity, which is transient in nature. In Authors [1]: Department of Pharmacology, Rao's college of pharmacy, Nellore. e-mail: stalinmpcol@gmail.com. In vitro studies have revealed its antispasmodic and curative actions confirming the use of the herb in Ayurvedic medicine [8]. The leaves of this plant mixed with castor oil are administered internally to treat rheumatism and to check lumbago [9]. The present investigation is directed to the exploration of the antidiabetic activity based on the study of the various extracts of *Cardiospermum halicacabum* which show inhibitory effect of glucose utilization and, are in use as hypoglycemic agent in traditional system of medicine.

2 II.

3 Materials and Methods

4 a) Plant material

The fresh plants of *Cardiospermum halicacabum* were collected from Nellore (Andhra Pradesh) and authenticated by Dr. P. Jayaraman, Ph.D., Director, Plant Anatomy Research Centre, Medicinal Plants Research Unit, Tambaram, Chennai-45. A portion of the sample was kept in the department museum for further reference (PARC/2010/579).

5 b) Preparation of extracts

The shade dried powdered form of leaves of *Cardiospermum halicacabum* was taken and subjected to successive extraction using n-hexane, Ethanol, and methanol by continuous percolation process in soxhlet apparatus. The aqueous extract was prepared by the maceration with double distilled water. Each extract was concentrated by distilling off the solvent and evaporated to dryness. The extracts were dissolved in 1% carboxy methyl cellulose (CMC) and used for the present study.

6 c) Effects of Various Extracts on In vitro Inhibitory

Glucose Diffusion A simple model system was used to evaluate the effects of *Cardiospermum halicacabum* leaf extracts on glucose movement in vitro. The model was adapted from a method described by Edwards et al. [10] which involved the use of a sealed dialysis tube into which 15ml of a solution of glucose and sodium chloride (0.15M) was introduced and the appearance of glucose in the external solution was measured. The model used in the present experiment consisted of a dialysis tube (6cmX15mm) into which 1ml of 50g/litre plant extract in 1% CMC and 1ml of 0.15M sodium chloride containing 0.22M D-glucose was added. The dialysis tube was sealed at each end placed in a 50ml centrifuge tube containing 45ml of 0.15M sodium chloride. The tubes were placed on an orbital shaker and kept at room B

Diabetes mellitus is a metabolic disorder characterized by a loss of glucose homeostasis with disturbances of carbohydrate, fat and protein metabolism resulting from defects in insulin secretion, insulin action, or both [1]. According to WHO, it is estimated that 3% of the world's population have diabetes and the prevalence is expected to double by the year 2025 to 6.3% [2]. Management of diabetes without any side effect is still a challenge to the medical community. The use of the drugs is restricted by their pharmacokinetic properties, secondary failure rates and accompanying side effects [3]. Thus searching for a new class of compounds is essential to overcome diabetic problems. There is continuous search for alternative drugs [4]. D temperature. The movement of glucose into the external solution was monitored at set time intervals.

7 d) Statistical Analysis

Data are expressed as mean \pm S.E.M. Statistical comparisons between groups were done by one way analysis of variance (ANOVA) followed by Tukey Kramer multiple comparison tests to analyze the differences. $p < 0.001$ were considered as significant.

8 III.

9 Results

10 a) Effect on Glucose Diffusion

With the distinctive traditional medical opinions and natural medicines mainly originated in herbs, traditional medicine offers good clinical opportunities and shows a bright future in the therapy of diabetes mellitus and its complications. The effect of *Cardiospermum halicacabum* leaves as anti-diabetic agents has been studied. All extracts showed varying effect on glucose utilization. These extracts caused a significant decrease in glucose concentration during the experiment.

The effects of *Cardiospermum halicacabum* leaves extracts on glucose diffusion inhibition were summarized in Table ??1. At the end of 27 hrs, glucose movement of control (without plant extract) in the external solution had reached a plateau with a mean glucose concentration above 300mg/dl (314 ± 2.89). It was evident from the table that the methanol and aqueous extracts were found to be potent inhibitors of glucose diffusion ($p < 0.001$) compared to control. The methanol extract was found to be more potent than other extracts showing the lowest mean glucose concentration of 201 ± 1.69 mg/dl at the end of 27 hrs (Table .1) IV.

11 Discussion

Diabetes mellitus is a debilitating and often life threatening disorder with increasing incidence throughout the world. There is a steady rise in the rate of incidence of Diabetes mellitus and estimated that 1 in 5 may be diabetic by 2025 [11]. Antihyperglycemic activities of most effective plants were in part explained by the ability of the phytoconstituents to increase glucose transport and metabolism in muscle and/or to stimulate insulin secretion [12]. In the present study, research has been carried out to evaluate the potential of various extracts to additionally retard the diffusion and movement of glucose in the intestinal tract [13].

A decoction of *Cardiospermum halicacabum* leaves is used worldwide for the treatment of various ailments including antidiabetic. The numerous polyphenolic compounds, triterpenoids and other chemical compounds present in the plant may account for the observed antidiabetic effects of the leaf extracts.

A Decoction of *Cardiospermum halicacabum* leaves was screened for hypoglycaemic activity on alloxan-induced diabetic rats. In both acute and sub-acute tests, the water extract, at an oral dose of 250 mg/kg, showed statistically significant hypoglycaemic activity [14]. The treatment with *Cardiospermum halicacabum* aqueous leaf extract (0.01-0.625 mg/mL) showed significant inhibition on LDL glycation in a dose-dependent manner. Tannins, flavonoids, apigenin, pinitol and luteolin, and other chemical compounds present in the plant are

speculated to account for the observed hypoglycaemic and hypotensive effects of the leaf extract. Values are expressed as mean \pm SEM of triplicate; Data were analysed using one way ANOVA followed by Tukey-Kramer multiple comparison test; ***P<0.001 compared to control.

12 Conclusion

The present study demonstrates the ability of various extracts of *Cardiospermum halicacabum* to inhibit glucose diffusion using an in vitro model of glucose absorption. In particular, methanol and aqueous extracts represent potential inhibitory of glucose diffusion supplements that may be useful for allowing flexibility in meal planning in type II diabetes. Further studies are required to elucidate whether in vitro effects represent therapeutic potential by limiting postprandial glucose absorptions and for improving glycemic control in type 2 diabetic subjects.

Extract	tube over 27hr incubation period				
	1h	3h	5h	24h	27h
Control(in the absence of extract)					

Figure 1: Table 1 :

¹© 2013 Global Journals Inc. (US)
²()BIn Vitro Antidiabetic Activity of *Cardiospermum Halicacabum* leaves Extracts

-
- [Gandhi and Sasikumar ()] 'Antidiabetic effect of MerremiaemarginataBurm. F. in streptozotocin induced diabetic rats'. Rajiv Gandhi , G Sasikumar , P . *Asian Pacific Journal of Tropical Biomedicine* 2012. 2 p.
- [Abdalla et al. ()] 'Antidiabetic Effects of Fenugreek (Trigonellafoenum -graecum) Seeds in the Domestic Rabbit (Oryctolagusuniculus)'. M Abdalla , Mariam Y Abdelatif , Mahmoud S Ibbrahim . *Res J of Medicinal Plant* 2012. 6 p. .
- [Reddy et al. ()] 'Antimicrobial screening of the plant extracts of Cardiospermum halicacabum L., Against selected microbes'. Thirupal Reddy , B , Ali Moulali , D Anjaneyulu , E Ramgopal , M , Hemanth Kumar , K Lokanatha , O Guruprasad , M Balaji , M . *Ethnobotanical Leaflets* 2010. 14 p. .
- [Ming-Hsinghuanga et al. ()] 'Antioxidant and anti-inflammatory properties of Cardiospermum halicacabum and its reference compounds ex vivo and in vivo'. Ming-Hsinghuanga , Shyh-Shyunhuangb , Bor-Senwangc , Ming-Jyhshueid Chieh-Hsiwud , Wen-Chi Houe , Guan-Jhonghuangb Shiang-Shioulinb . *J Ethnopharmacol* 2011. 133 p. .
- [Muthumani et al. ()] 'Chemical Investigation Of Tod-daliaAsiaticaLinn, AndCardiospermum Halicacabum Linn'. P Muthumani , R Meera , DeviP , Mohamed Sheik Arabath , S A Seshukumarkoduri , L V Sivaramanavarthi . *International Journal of Drug Formulation & Research* 2010. 1 (3) p. .
- [Annamalai et al. ()] 'Effect Of Drying Treatment On The Contents Of Antioxidants In Cardiospermum Halicacabum Linn'. A Annamalai , G Ponmaril , R Sathishkumar , P T Lakshmi . *International Journal of Pharma and Bio Sciences* 2011. 2 (1) p. .
- [Vishwakarma et al. ()] 'Evaluation of effect of aqueous extract of Enicostemmalittorale Blume. In streptozotocin induced type 1 diabetic rats'. S L Vishwakarma , S Rakesh , M Rajani , R K Goyal . *Indian J ExpBiol* 2010. 48 p. .
- [Priyadarshini et al. ()] 'Hypolipidaemic and Renoprotective study on the Ethanolic & Aqueous extracts of leaves of Ravenalamadagascariensis Sonn. onalloxan induced diabetic rats'. S S Priyadarshini , Vadivu , N Jayshreet . *International J Pharm Sci* 2010. 2 p. .
- [Palanuvej et al. ()] 'In Vitro Glucose Entrapment and Alpha-Glucosidase Inhibition of Mucilaginous Substances from Selected Thai Medicinal Plants'. C Palanuvej , S Hokputsa , T Tunsaringkarn , N Ruangrungsi . *Sci Pharm* 2009. 77 p. .
- [Gray and Flatt ()] 'Insulin-like and insulin-releasing actions of the traditional antidiabetic plant Sambucusnigra (elder)'. A M Gray , Abdel-Wahab Y H A Flatt , PR . *J Nutr* 2000. 130 p. .
- [Datta et al. ()] 'Phytochemical and biological evaluation of Cardiospermum halicacabum'. S Datta , A Ghosh , P Pal , M Das , P K Kar , Pharmacognostical . *Int J Pharm Sci Bio* 2010. 1 (1) p. .
- [Mittal et al. ()] 'Phytochemistry and pharmacological activeities of psidiumguajava: a review'. P Mittal , V Gupta , G Kaur , K G Ashish , Amarjeet Singh . *International J Phar Sci Res* 2010. 1 p. .
- [Syamsudin ()] 'Standardization of extract of Leucaenaleucocephala (lmk) De Wit seeds by ?glucosidase inhibitor'. Syamsudin . *International Journal of Phytomedicine* 2010. 2 p. .
- [Edwards et al. ()] 'Viscosity of food gums determined in vitro related to their hypoglycemic actions'. C A Edwards , N A Black Burn , L Cragne , P Daavidson , J Tomlin , K Sugden , I T Johnson , N W Read . *Am J CliNutr* 1987. 46 p. .