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Glucose lowering effects and *in vitro* α -amylase and α -glucosidase inhibitory potentials of aqueous extract of *Adansonia digitata* (baobab) seed

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pharmaceuticals



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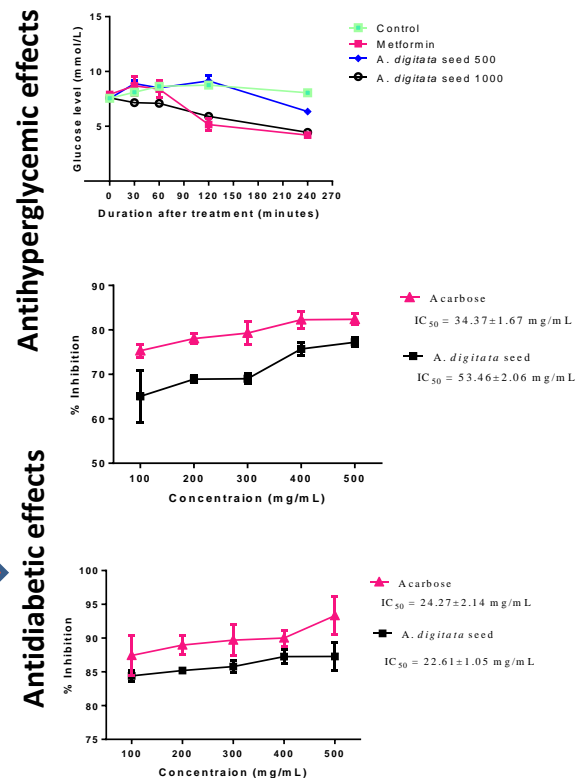
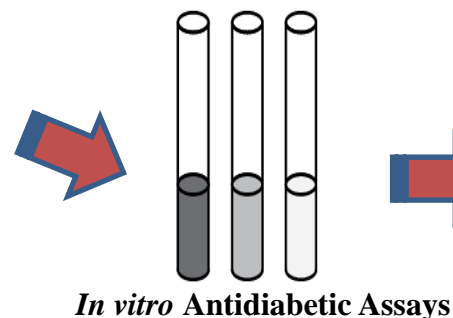
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Glucose lowering effects and *in vitro* α -amylase and α -glucosidase inhibitory potentials of aqueous extract of *Adansonia digitata* (baobab) seed

Graphical Abstract



Adansonia digitata
(baobab) seed



Abstract:

Adansonia digitata L. (*Malvaceae*, Baobab) is a medicinal tree with antimicrobial, antiviral, anti-inflammatory, and antioxidant properties. The leaves, fruit pulp, stem bark, and roots have been extensively studied. The aim of this study was to evaluate the glucose-lowering and *in vitro* antidiabetic potentials of the aqueous extract of *A. digitata* seed. The aqueous extract of *A. digitata* seed was prepared by dissolving 50g of powder seed in 500 mL of distilled water for 24 hours, filtered using Whatman filter paper, and concentrated using a rotary evaporator at 40°C. Following an oral administration of glucose (2 g/kg body weight), distilled water, metformin (14.2 mg/kg body weight), and *A. digitata* seed extracts at 500 and 1000 mg/kg body weight, respectively. Results show that the untreated mice had an average of 11.09% increase in plasma glucose concentration, while metformin, aqueous seed extract of *A. digitata* had an average of 17.05%, 0.99%, and 19.21% decrease in plasma glucose concentration respectively. The aqueous seed extract of *A. digitata* inhibited α amylase in a concentration-dependent manner with an IC_{50} of 24.27 ± 2.14 mg/mL compared with acarbose with IC_{50} of 22.61 ± 1.05 mg/mL. However, the α -glucosidase inhibitory activities of the extract (IC_{50} 34.37 ± 1.67 mg/mL) were significantly lower compared to acarbose (IC_{50} 53.46 ± 2.06). The study concludes that aqueous seed extract of *A. digitata* possesses glucose-lowering properties, *in vitro* α -amylase and α -glucosidase inhibitory potentials. Further studies will require a bioguided fractionation of aqueous seed extract of *A. digitata*, to identify its phytochemical constituents using fingerprint chromatography among other techniques.

Keywords: *Adansonia digitata*; Antihyperglycemia; *In vitro*; Seed; α -Amylase; α -Glucosidase

Introduction

Adansonia digitata L. (*Malvaceae*; Baobab) is a medicinal plant wide distribution throughout the sub-Saharan Africa and Western Madagascar areas. Their many important use include medicinal, beverages, and as food (Lamien-meda *et al.*, 2008).

Diabetes mellitus is a common and prevalent disease with estimated 1.5 million deaths in 2019 and reported 48% of death cases occurring before 70 years (WHO, 2021). Diabetes is a chronic metabolic disorder of fats, proteins, and carbohydrate metabolism, which are linked to insulin functions; blood insulin level or insensitivity of target organs or both (Ghosh *et al.*, 2004). Diabetes are associated with oxidative stress (Tijjani *et al.*, 2020a) and hyperlipidemia (Tijjani *et al.*, 2020b).

The seed of *A. digitata* (baobab) are used traditionally for the management of diabetes and its complications, among other derived benefits. The present study was designed to evaluate the glucose lowering properties, and *in vitro* antidiabetic potentials of the aqueous extract of *A. digitata* seed.

Materials and methods

Chemicals and reagents

α – amylase, α – glucosidase, metformin, *p*-Nitrophenyl- α -glucopyranoside (PNPG) and 3, 5-dinitrosalicylic acid (DNSA) were purchased from Sigma-Aldrich, Germany. All other reagents used were of analytical grade.

Experimental animals

Sixteen (16) albino mice weighing between 20 ± 2 g were purchased from the animal unit, University of Jos, Plateau state and acclimatized for 14 days in a plastic cage of 4 mice each, given standard mouse chow and water *ad libitum*. The animal care procedure was approved by the ethical review committee, University of Jos, with registration number UJ/FPS/F17-00379.

Assays

Antihyperglycemic assay was determined according to the method of Du Vigneaud and Karr (1925), while α -amylase and α -glucosidase inhibition assays were determined according to Andrade-Cetto *et al.* (2008) and Kuppusamy *et al.* (2011).

Results and discussion

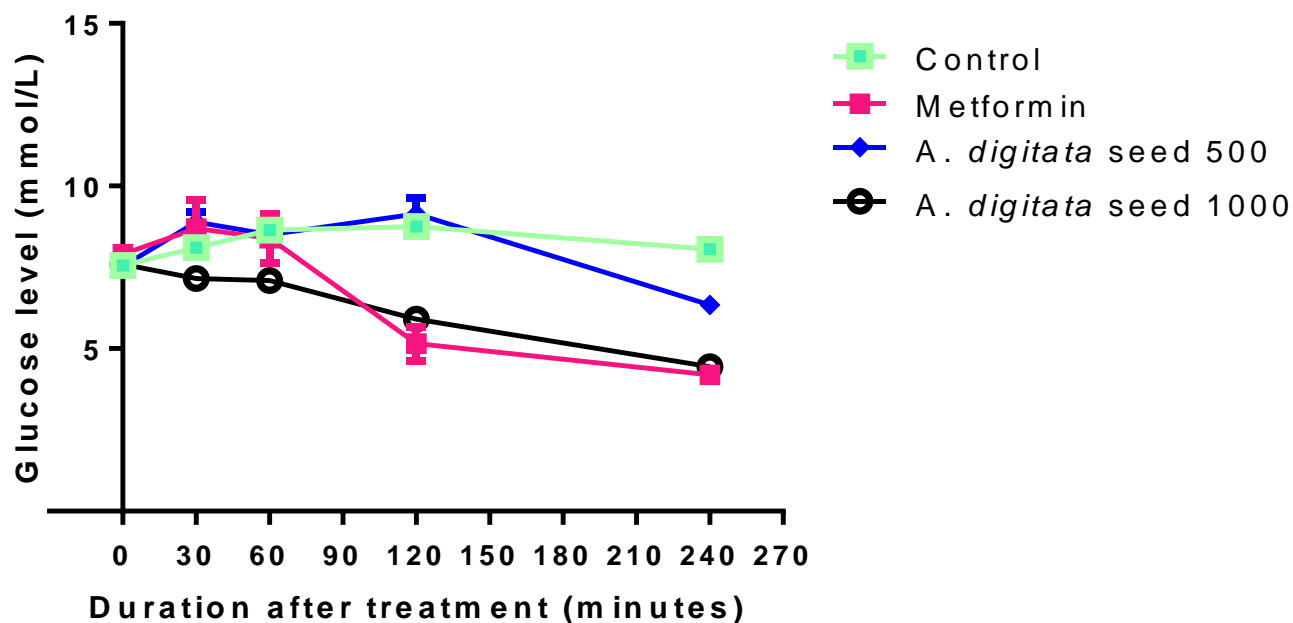


Figure 1: Antihyperglycemic effects of aqueous seed extract of *A. digitata* in experimental mice. Values are mean \pm SD, n=4, Dosages are in mg/kg body weight

Results and discussion

Table 1: Percentage decrease in blood glucose levels following treatments with aqueous seed extract of *A. digitata* in experimental mice

| Groups (mg/kg bwt) | Effects on glucose level (%) | | | | | Average |
|-------------------------|------------------------------|---------------|---------------|----------------|----------------|---------|
| | 0 minutes | 30 Minutes | 60 Minutes | 120 minutes | 240 Minutes | |
| Control | 7.55 | 8.10 (↑7.28) | 8.65 (↑14.56) | 8.75 (↑15.89) | 8.05 (↑6.62) | ↑11.09 |
| Metformin | 7.90 | 8.70 (↑10.59) | 8.40 (↑6.62) | 5.15 (↓36.42) | 4.20 (↓49.01) | ↓17.05 |
| A. digitata 500 | 8.30 | 8.90 (↑7.94) | 8.50 (↑2.64) | 9.15 (↑11.25) | 6.35 (↓25.82) | ↓0.99 |
| A. digitata 1000 | 7.60 | 7.15 (↓5.96) | 7.10 (↓6.62) | 5.90 (↓22.51) | 4.45 (↓41.72) | ↓19.21 |

Values are mean, n=4, ↑ = increase, ↓ = decrease, mg/kg bwt = mg/kg body weight

Results and discussion

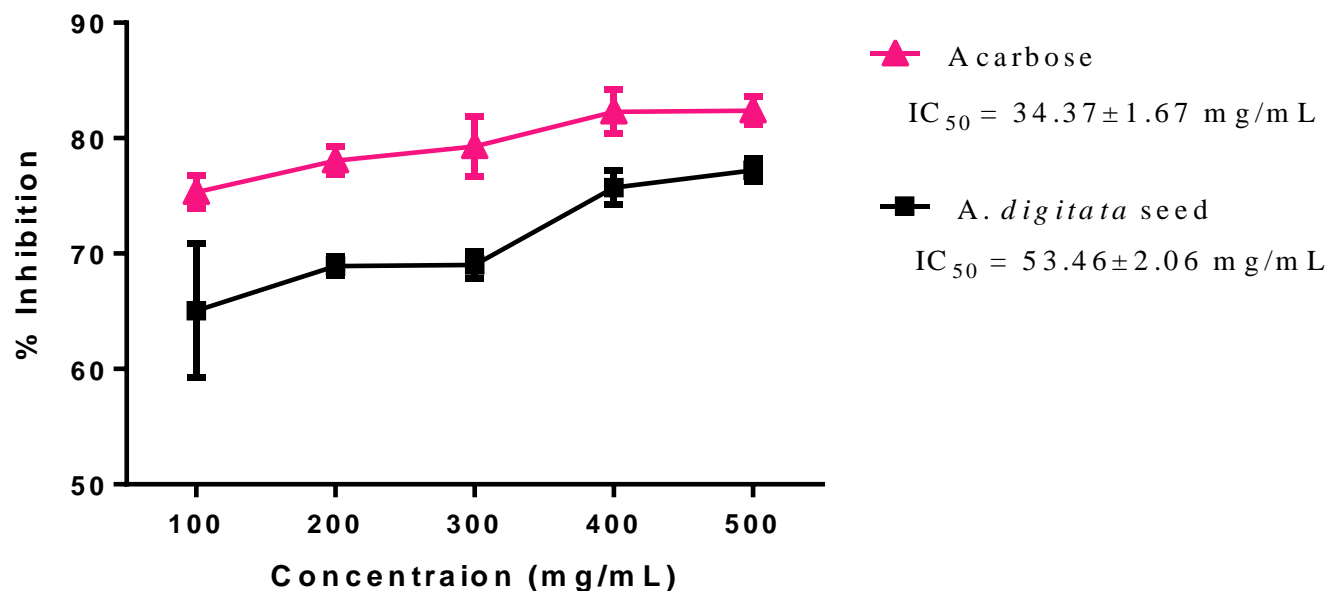


Figure 2: Inhibition of α -glucosidase by aqueous extract of *A. digitata* seed. Values are mean \pm SD of triplicate determination

Results and discussion

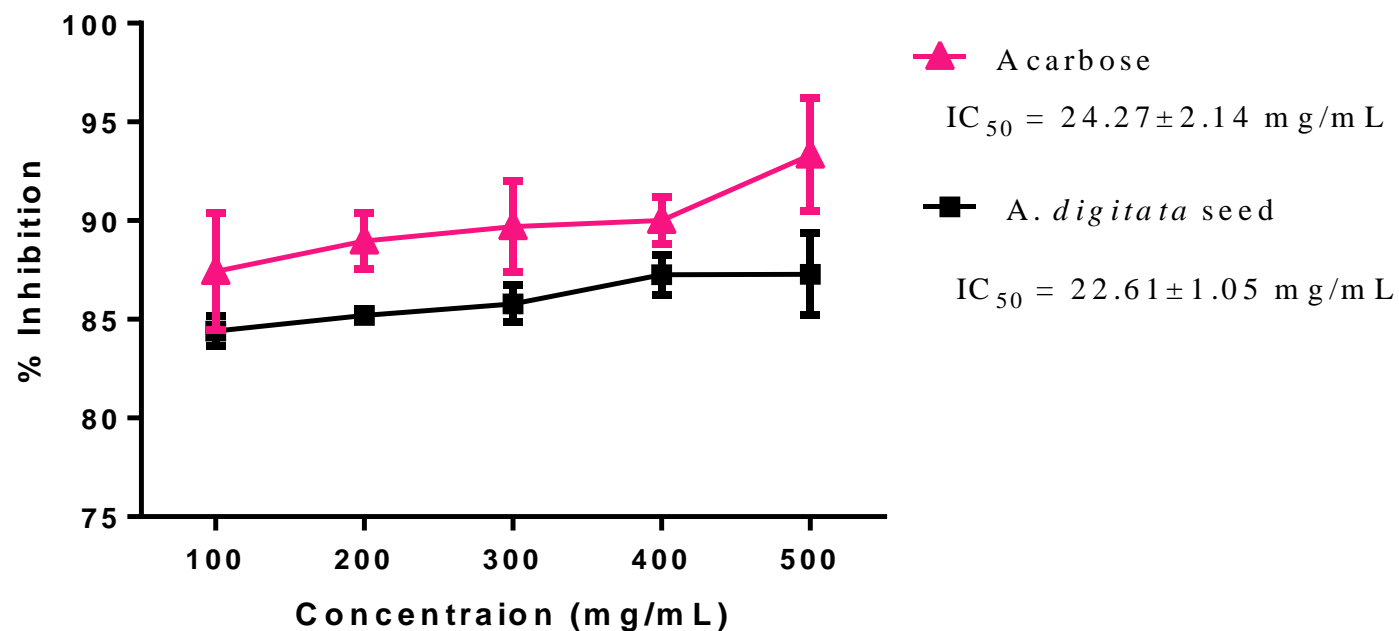


Figure 3: Inhibition of α -amylase by aqueous extract of *A. digitata* seed. Values are mean \pm SD of triplicate determination

Conclusions

The study conclude that aqueous extracts of *A. digitata* seed possess both *in vitro* α -amylase and α -glucosidase inhibitory potential and possess glucose lowering effects at 500 and 1000 mg/kg body weight. Thus, providing scientific support for the usage of *A. digitata* seed in the management of diabetes mellitus. Further studies will required a bioguided fractionation of aqueous seed extract of *A. digitata*, to identify its phytochemical constituents using fingerprint chromatography among other techniques.

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