

Botanical Pharmacognostical Study of Al-Booda (*Striga hermonthica*) Distributed in Gezira State, Sudan

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ABSTRACT

Striga hermonthica is an invasive and destructive weed which is usually infects crops as well as pasture and wild grasses. It is widely distributed in many African countries including Sudan and it is traditionally been used as abortifacient, for pneumonia, fungal infections, as contraceptive, mosquito repellent and anti-diabetic. This study was aimed to identify morphology of whole plant and microscopic characteristics of the root, stem, and leaf and powder microscopy to support its botanical pharmacognostical characterization. Morphological and microscopic study was carried out using fresh and dry whole plant, the root, stem and leaves. The results of present macroscopic and histomicroscopic features of root stem and leaves of *S. hermonthica* and powder microscopic were reported which may provide useful information for the regulatory aspects of the quality control measures of the crude drugs.

Keywords: *Striga hermonthica*; Botanical pharmacognostical; Macroscopic; Microscopic

INTRODUCTION

Striga hermonthica (Del.) Benth. is a weed belongs to the family Orobanchaceae which is considered invasive and destructive and nearly hard to exterminate and usually infect crops as sorghum, maize, millet, rice, and sugar cane, as well as pasture and wild grasses [1,2]. It attaches itself to the roots of the host plant and diverts essential nutrients, which render the host stunted and decrease the yield of the crops more than 50% [3]. *S. hermonthica* widely distributed in many African countries and its commonly known as Witchweed and locally in Sudan it is called Al-Booda. It is the most distributed among *Striga* species in Africa throughout the Nile Delta and have the most damage to crops. It has great within-species variation which is expected as it is an obligate out-breeder [4]. Also, *S. hermonthica* was collected from different geographical regions exhibits variability (Figure 1) [5]. Floral variations may discourage cross pollinations between these strains and further enhance the differentiation of populations in *S. hermonthica* as evidenced in recent molecular studies in which populations of *S. hermonthica* collected from sorghum, millet, and maize showed genetic differences, the maize strains were more closely related to the sorghum-strain than to the millet-strain. Despite its effects on crops and agriculture, it is traditionally been used orally as an abortifacient, for pneumonia, fungal infections, as contraceptive, mosquito repellent [6], anti-cancer [7], and anti-diabetic agent in western Sudan [8].

Some pharmacological investigations of *S. hermonthica* extracts were showed an *in vitro* effect against *Trypanosoma congolense* and *Trypanosoma cruzi* [9], antimicrobial activities against *Staphylococcus aureus*, *Pseudomonas aeruginosa*, *Escherichia coli* and *Candida albicans*

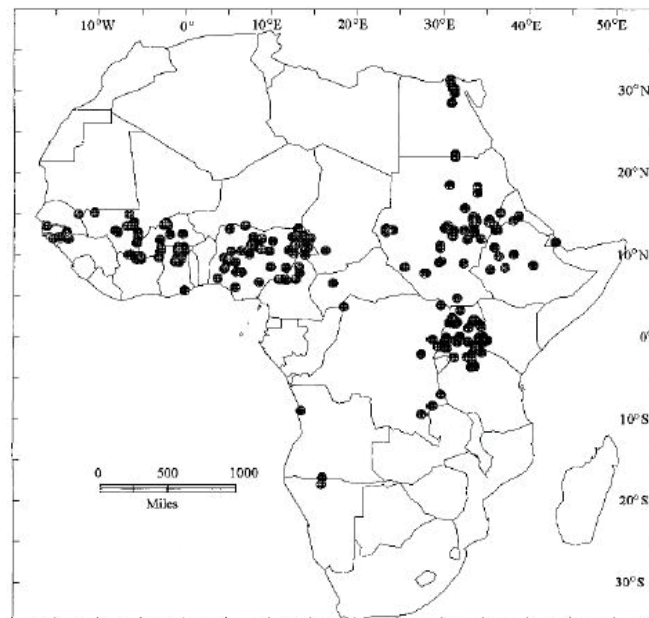


Figure 1: Geographical distribution map of *S. hermonthica* in Africa.

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[10]. The phytochemical studies of this plant showed the presence of terpenes, saponin, cardiac glycosides, alkaloids, anthracenocides, coumarins, tannins and flavonoids [11]. However, few pharmacognostical studies were carried out for evaluation of plant material of *S. hermonthica*. Therefore, this study was aimed to study the morphology of whole plant and microscopic characteristics of the root, stem, leaf and powder microscopy, in order to identify botanical-pharmacognostical markers, which may be an important contribution regarding the aspects of quality control measures of *S. hermonthica* and related similar species.

The taxonomical position of *S. hermonthica* [12]

Domain: Eukaryota

Kingdom: Plantae

Phylum: Spermatophyta

Subphylum: Angiospermae

Class: Dicotyledonae

Order: Scrophulariales

Family: Orobanchaceae

Genus: *Striga*

Species: *S. hermonthica*.

MATERIALS AND METHODS

S. hermonthica (whole plant) was collected from Shukaba province, South Gezira State, Sudan and identification and authentication were done at Department of Pharmacognosy University of Gezira. The used chemical materials and equipment include ethanol, distilled water, chloral hydrate 10%, safranin reagent 1%, glycerin, handheld magnifying lens (25x magnification), light microscope (OLYMPUS CH₂O_i, Tokyo, Japan) connected to 16.0-Megapixel digital camera, razors and dissection sets.

Microscopical studies were carried out for whole plant parts using the handheld magnifying lens. Fine hand sections of the lamina, stem, root, using very sharp razor were cut into very thin transverse sections, then mounted with chloral hydrate and heated using flame until bubbling occurs. User sections were stained with aqueous safranin and mounted in glycerin. Microphotographs of sections and powder analysis were made by using microscope and photos were collected too.

RESULTS AND DISCUSSION

Macroscopical study

S. hermonthica (Figures 2 and 3) is a herbaceous annual plant ranging in height of 30-100 cm, where the most infectious forms occurring in Sudan and Ethiopia [12,13]. Larger plants show more branching. Stems and leaves are covered with specific trichomes rendering the plant with a harsh texture. Leaves mostly opposite on the lower half of the stem but irregular above, narrowly lanceolate, or elliptic, 2-8 cm long, up to 1 cm wide. Inflorescence is a terminal spike of sessile flowers that branching from the axils of upper leaf. Flowers joined by bracts 1-2 cm long, up to 3 mm wide, with an edge of ciliate hairs. Calyx tubular reaching 1 cm long with 5 ribs and 5 teeth



Figure 2: Photo showing *S. hermonthica* in the wild.



Figure 3: Photo showing *S. hermonthica* (whole plant).

varying from 2-3 mm long. Flower asymmetrically campanulate, the tube 1-2 cm long, bent approximately halfway up in West African, Sudanese, and Ethiopian populations but usually well above halfway in East African populations [14]. Corolla lobes 4, one bi-lobed almost erect, the others spreading horizontally, up to 2 cm across, pink with some white markings within the throat. The stigma and stamens are concealed in the tube. The primary inflorescence spike may hold up to 100 flowers but only 6-10 are open at a time. The capsules are up to 1 cm long and each develops several hundred-minute seeds, approximately 0.3 mm long by 0.2 mm wide. The root system is delicate with little ability to uptake nutrients from the soil, but branches develop from lower nodes of the weed, ramifying and developing secondary haustoria and attachments on contact with other host roots [15], which facilitate the loss of water, carbon, nitrogen and nutrients from the host plant to *S. hermonthica* [16].

Microscopical study

Microscopic study of root: The root is elongated taproots, cylindrical, sharpened towards the ends with brownish-yellow color.

Transverse section (T.S.) of the young root (Figure 4) appeared circular in shape with tissue organization as outer piliferous layer, followed by the middle cortex and inner stele. Most tissues cells were empty and appeared reddish to brown in color, but some cells contained the deposition of colored secretary substances. Cortex was well marked and contains the important section of the root where parenchymatous cells were found to contain air cavities that facilitate the gaseous exchange [17]. Starch grains were in the cortical parenchymatous cells (Figure 5), where carbohydrate storage [18]. Calcium oxalate crystals (Figure 6) were found in the outer cortical parenchyma cells with various functions regulation of calcium levels in plant tissues and organs, protection against herbivory, detoxification of heavy metals and/or strengthening of the tissues [19]. The central part or stele were highly reduced and represented largely by xylem vessels and phloem (Figure 7).

Microscopic study of stem: T.S. of *S. hermonthica* stem was appeared with semi-regularly circular outline (Figure 8) with tissue organization as outer layer of epidermis, followed by the cortex and vascular bundle. Epidermis consisted of single cellular layer with thick cuticle. The Cortex was appeared in rows of cells contained chlorophyll and calcium oxalate crystals. Vascular bundles were mostly secluded with polygonal to ovoid shapes.

Microscopic study of leaf: T.S. of *S. hermonthica* leaf indicated that the lamina was differentiated into epidermis, mesophyll and

vascular tissues (Figure 9). Upper and lower epidermises were a single layer composed of tightly arranged rectangular cells with striated outer walls covered by cuticle to reduce epidermal water

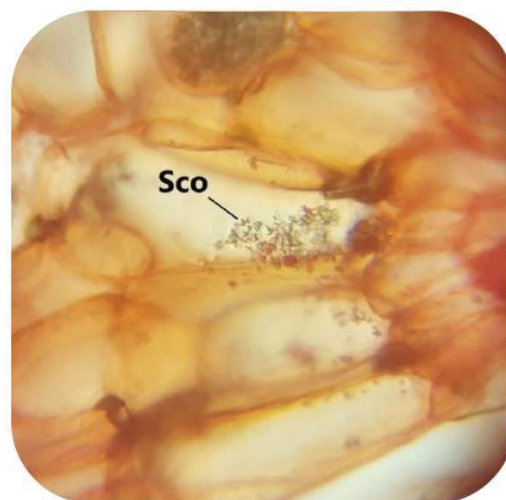


Figure 6: 40x T.S. of *S. hermonthica* Root showing Sco=Calcium oxalate crystals

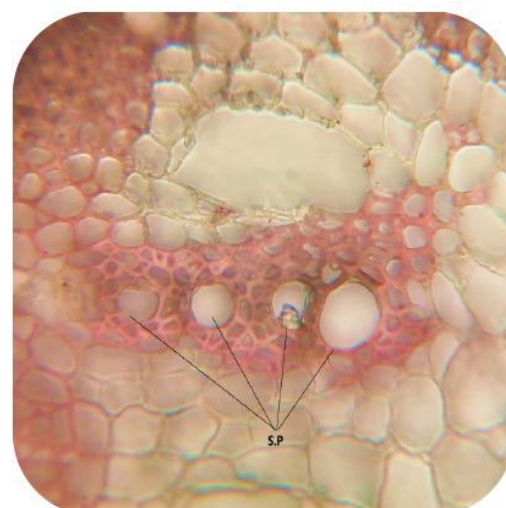


Figure 7: 40x T.S. of *S. hermonthica* Root showing S.P=phloem



Figure 4: 4x T. S. of *S. hermonthica* Root showing Sp=Phyllum, Sx=Xyllum, Spi=Pidth, Sc=Cortex tissue, Se=Epidermis

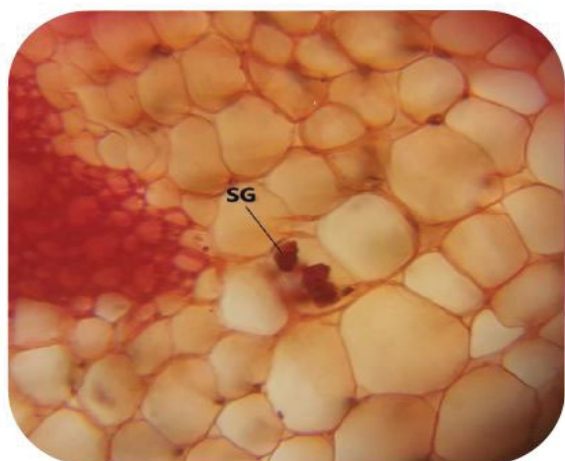


Figure 5: 40x T. S of *S. hermonthica* Root showing SG= starch grain

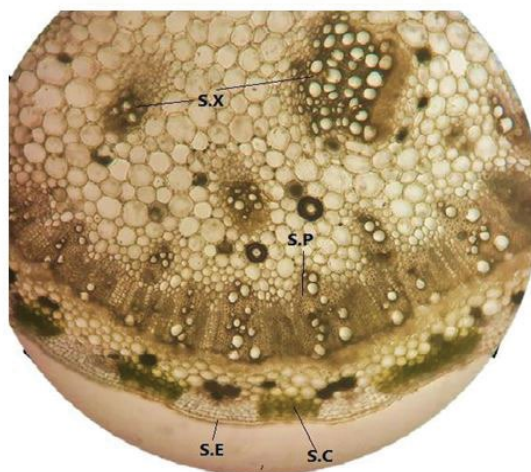


Figure 8: 10x Transverse section of *S. hermonthica* stem showing S.E.=Epidermis, S.P.=Primary Cortex, S.X.=Vascular bundle, S.C.=Schlrenchymatous cells.

loss [3] and presence of few trichomes. Below the both epidermis there was a layer of sclerenchymatous tissues. The mid rib area was appeared to have an acute depression on the adaxial side with broadly ovoid abaxial side. Midrib was composed of epidemics, sclerenchyma, collenchyma and vascular bundle (Figure 10). Lamina was differentiated upper elongated mesophyll having a compact palisade cells contain chloroplasts and lower spongy parenchymatous tissue (Figure 11) which allows efficient nutrient remobilization [20].

Microscopic study of powder: The shade dried *S. hermonthica* whole plant was milled into powder that subjected to microscopic examination which showed fragments of leaf epidermis with venations, flower stalk fragments, calcium oxalate crystals, sclerenchyma and parenchyma fragments and starch grains (Figure 12).

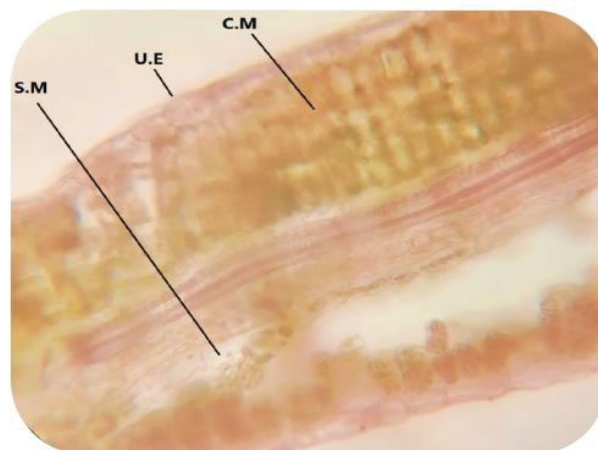


Figure 11: 10x Transverse section of *S. hermonthica* leaf showing U.E.=Upper epidermis, C.M.=Condensed mesophyll, S.M.=Spongy mesophyll.

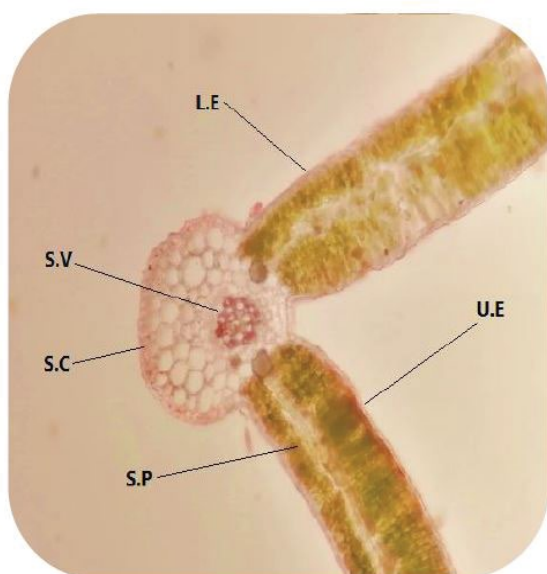


Figure 9: 10x Transverse section of *S. hermonthica* Leaf showing U.E.=Upper epidermis, S.P.=Spongy parenchymatous tissue, S.C.=Chlorenchymatous cells, S.V.=Vascular bundle.

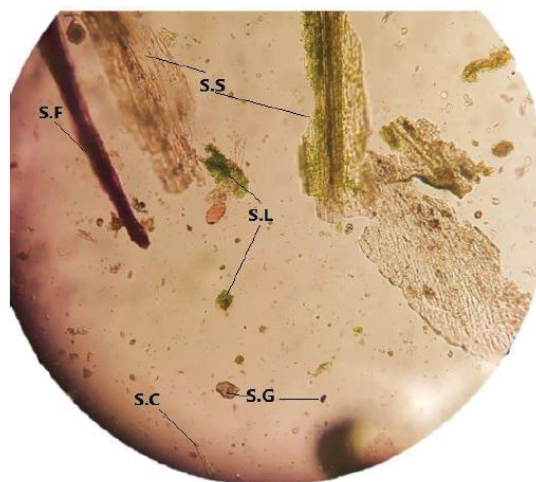


Figure 12: 10x Powdered drug of *S. hermonthica* whole plant showing different parts S.F.=Flower stalks, S.S.=Steam parts, S.L.=Leave laminaa, S.G=Grains, S.C.=Calcium oxalate crystals.

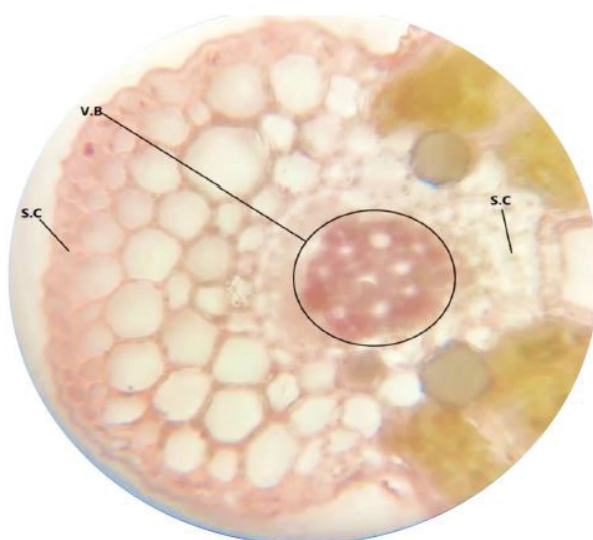


Figure 10: 10x Transverse section of *S. hermonthica* Leaf showing V.B.=Vascular bundle both Xylem and phloem, S.C.=Thickened collenchyma cells.

CONCLUSION

This study reported the morphological and anatomical features of *S. hermonthica* (Del.) Benth distributed in Sudan, including general macroscopical characters and microscopical study of root, stem, leaf and whole powdered plant materials. These findings could be remarkable anatomical markers useful for identifying this species that are useful for Botanical Pharmacognostic standards or markers for identification of crude drug materials which found scant for *S. hermonthica*.

DECLARATIONS OF COMPETING INTEREST

None.

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