Variations in antimalarial components of *Artemisia annua* Linn from three regions of Uganda

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**Abstract**

**Introduction:** Artemisia annua plant from the family Asteraceae is a powerful antimalarial plant introduced to Uganda around 2003. In addition to the artemisinin component, the plant also contains flavonoids which work in synergy to artemisinin against malaria parasites. The plant also contains aromatic oils which repel mosquitoes. In this paper we report the variations in antimalarial components of *A. annua* samples from the regions cultivating it in Uganda.

**Methods:** Artemisia annua samples were obtained from three regions that cultivated the plant at the time of this study. The samples were brought to laboratory, authenticated and processed. The levels of artemisinin, total flavonoids and aromatic components were quantified using high performance thin layer chromatography, ultra violet spectrophotometry and gas chromatography respectively.

**Results:** Artemisinin and total flavonoids levels were higher in samples obtained from high land areas (western and south western region) compared to that obtained from lowland regions (central) i.e 0.8% Vs 0.4% and 2.6% Vs 1.5% respectively. The aromatic oils (mosquito repellent components) were similar with camphor component being highest and levels ranging from 75.4% to 79.0%.

**Conclusion:** Our findings show that the active components in Artemisia annua cultivated and used in the Uganda vary with geographical regions and this calls for standardisation by source.

**Key words:** Variations, Antimalarial components, Artemisia annua, Uganda

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**Introduction**

Artemisia annua L. is a medicinal weed that belongs to the Asteracea family of plants. It is native to Asia especially China but also found in Europe and North America. In China, the leaves were used as infusion to cure malaria and also burnt to repel mosquitoes for over 3000 years¹. A. annua plant was introduced to Uganda around 2003 as the source of artemisinin needed by the pharmaceutical industries that manufacture artemisinin combination therapies (ACTs). Ugandan soils are reported to be suitable for cultivation of A.annua². Artemisinin the most potent antimalarial known or its derivative is not used singly in treatment of malaria but used in combination with other antimalarial drugs. The artemisinin combinations are currently WHO recommended treatments for uncomplicated malaria³. Apart from artemisinin, A. annua also contains other antimalarial compounds that work in synergy with artemisinin offering a natural combination therapy against malaria⁴,⁵,⁶,⁷.

In Uganda, the cultivation of the plant was initially restricted to farmers in highland areas mainly in western Uganda. However, the drop in the demand for artemisinin from Uganda by the industries that make its derivatives led to uncontrolled and wide spread local cultivation and use of the plant in Uganda. The plant is now cultivated and used by communities in Central, Eastern and Northern Uganda which are low land areas (less than 1200m above sea level) initially not recommended for the plant⁸,⁹. A recent study in Uganda reported that although Ugandan soils are suitable for...
cultivation of artemisinin, artemisinin levels vary with area of cultivation. The variations in levels of other antimalarial components remained unknown until this study. All plant components are known to vary according to geographical locations which in turn may lead to varying clinical outcomes in patients. The low land areas in Uganda have altitude of about 1200m, receive total rainfall of about 1000mm per annum and temperatures of up to 84°F. The high land areas have altitude of up to 2500m; receive rainfall of up to 1200mm per annum and temperatures of up to 73°F. In this paper, we report variations in the levels of artemisinin, total flavonoids and aromatic oils in samples from the three regions cultivating A. annua in Uganda.

Materials and methods

Material collection
Artemisia annua plant was collected from the Wagagai flower farm garden in Central Uganda in February 2009. The specimen was identified by a taxonomist at Natural Chemotherapeutics Research Institute (NCRI) herbarium, Ministry of Health, Uganda. A voucher specimen, NCJ 257 was deposited at the herbarium of NCRI for use as reference. Dry leaf powder of A. annua samples (1kg) was also obtained from Wagagai health clinic, where it has been used as tea for malaria prophylaxis among the farm workers since 2006. Other dry leaf powder samples (1kg each) were obtained from the two major commercial cultivators in Uganda located in high land areas i.e Kabale district in south western Uganda and Fortportal district in Western Uganda. The samples brought to the laboratory were stored in air tight opaque containers at room temperatures till the time of analysis.

Phytochemical screening of artemisia annua materials.
The A.annua dry leaf powders (100g) were extracted with soxhlet apparatus in a sequential manner using petroleum ether, followed by ethanol and then methanol solvent. While A.annua tea was prepared by adding boiling water to dry leaf powder to make 10g/L infusion following procedures previously described. Phytochemical ingredients in both leaf powders and tea extracts were determined using methods described in the manual of analysis of vegetable drugs as briefly described;

Polyuronides
To a test tube containing (10ml of formulation) was added drops of water, leading to formation of a thick precipitate. The precipitate obtained was placed on the filter paper and on staining with hematoxylin formed a blue precipitate for presence of polyuronides.

Reducing compounds
1ml of formulation was diluted with water (2ml) in test tube. Fehling’s solutions I (1ml) and Fehling’s solution II (1ml) were added and heated in a water bath at 90°C forming a brick-red precipitate.

Saponins
A diluted solution of the formulation (2ml) was placed in a test tube and shaken for 15 minutes. A soapy like column of about 2cm formed above liquid level.

Tannins
To the formulation (1ml) was added water (2ml) and 3 drops of ferric chloride. A blackish blue color formed.

Alkaloid salts
The formulation (15ml) was evaporated to dryness in an oven at 55°C and residue dissolved in 10% v/v Hydrochloric acid (10 mL). 10 % v/v ammonia solution (10ml) was added to precipitate the alkaloids and then extracted with ether (15ml). The ether portion was evaporated to dryness and hydrochloric acid (1.5ml) added. To 0.5ml of the acidic solution was added 2-3 drops of Mayer’s reagents forming opalescence precipitate.

To detect Steroid glycosides, Anthracenosides, coumariins and flavonosides, 25ml of the formulation was mixed in 10% v/v hydrochloric acid (15ml), refluxed for 30minutes, cooled and extracted with diethyl ether (36ml) in portions of 12ml each.

Steroid glycosides
To a residue obtained by evaporating to dryness ether extract (10 ml) was added acetic anhydride (0.50ml) and chloroform (0.50 ml) and transferred into a dry tube. Conc. Sulphuric acid (2 ml) was added by means of a pipette at the bottom of the tube forming reddish-brown ring at the contact zone of the two layers.

Anthracenosides
The ether extract (4 mL) was added to conc. Sulphuric acid (2 mL) and shaken with 25% v/v ammonia solution (2ml) forming cherished-red solution on the top layer.

Coumarin derivatives
To a residue obtained by evaporating ether extract (5
mL) was added hot water (2ml) to dissolve. 10%v/v ammonium solution (0.5 ml) was then added forming a blue fluorescence solution under UV.

**Flavonosides**
The residue obtained by evaporating ether extract (5ml) was heated in 50% methanol (2 mL). Metallic magnesium (0.5g) and conc. Hydrochloric acid (5drops) was added forming a red solution.

**Aromatic oil profile in artemisia annua materials.**
The aromatic oils were extracted by hydro-distillation of 500g of A. annua dry leaf powder. The distillate was cooled and fractionated with petroleumether (500ml) to extract the volatile oils. The oils in the ether fraction were analysed using Shimadzu GC-MS; model, C70374300170. The peak area of each component of the oil was identified through the in built database and content of each identified component computed based on peak areas as percentages.

**Determination of artemisinin content in the artemisia annua samples**
**Construction of artemisinin calibration curve**
Method previous described for quantification of artemisinin in bulk forms was adopted. Pure artemisinin (1mg) donated by African Laboratory for Natural Products (ALNAP) in Ethiopia was dissolved in ethanol to make 100ml stock solution. Concentrations of 200ng/ml, 300ng/ml, 500ng/ml and 1000ng/ml of pure artemisinin were applied onto Thin Layer Chromatographic (TLC) plates (Merck) by the automated High Performance Thin Layer Chromatography (HPTLC) applicator, the plate was developed in HPTLC chamber using solvent system, heptane- diethyl ether in the ratio1:1. The retardation factor (Rf) value for artemisinin was 0.5. After drying, the absorbances for each concentration was measured at 366 nm UV-wave length and artemisinin content obtained from the calibration curve shown. The concentration of artemisinin in A.annua tea was determined using the same procedures.

**Quantification of total flavonoids in artemisia annua powder**
**Construction of casticin calibration curve**
Casticin is one of the antimalarial and immunomodulatory flavonoids present in A.annua that works in synergy to artemisinin. The method previously described by Chen et al for determination of total flavonoids was adopted with slight modification. Casticin used as a standard was prepared by dissolving 14mg in 25mls of 70% ethanol to give a concentration of 0.56mg/ml. The standard (4 ml) was then pipetted into 10mls volumetric flask and 0.4mls of 5% NaNO₂ added, mixed and allowed to react for 6 minutes. A solution of 10% Al(NO₃)₃ (0.4ml) instead of AlCl₃ previously used by Chen et al was added, mixed and allowed to complex for 6 minutes before adding 4 mls of 4% NaOH to neutralise the acidity. Double distilled water was added to top to the 10mls mark giving a yellow green solution of the final concentration 0.224mg/ml. Of the standard prepared, 0.2mls, 0.5mls, 1.0mls, 2mls and 3mls were each placed into cuvetts and topped to 4ml using double distilled. Their absorbance was then measured at 510nm using UV-spectroscopy, model PERKIN ELMER LAMBDA 35 uv/vis computerised spectrophotometer double beam. The respective concentrations gave absorbance of 0.098, 0.2795, 0.4417, 0.9907 and 1.1556. A calibration curve was constructed using absorbance and concentration giving line of best fit in which R2 value= 0.991.

Ten (10 g) of dried A.annua of each sample was weighed into a separate conical glass and 100mls of 80% methanol then added, coked and allowed to stand for 4 h with constant shaking to extract flavonoids according method previously described by Edeoga et al. The extractives were filtered using whatmann filter paper (No.1) to obtain clear filtrates. Filtrate (4ml) was then pipetted into 10mls volumetric flask and 0.4mls of 5% NaNO₂ add-
ed, mixed and allowed to react for 6 minutes. A solution of 10% Al(NO$_3$)$_3$ (0.4ml) instead of AlCl$_3$ was added, mixed and allowed to complex for 6 minutes before adding 4 mls of 4% NaOH to neutralise the acidity. Double distilled water was added to top to the 10mls mark. For each sample, 2mls was placed into cuvette and topped to 4ml using double distilled, its absorbance measured at 510nm and total flavonoid concentration estimated from the calibration curve in reference to casticin. The concentrations were used to calculate the total flavonoids content in dry leaf samples as percentage of dry weight.

Quantification of total flavonoids in artemisia annua tea
The tea of A.annua from Wagagai farm and that prepared using powders from Kabale and Fortportal were filtered through what filter paper (no.1). Filtrate (4ml) was then pipetted into 10mls volumetric flask and 0.4mls of 5% NaNO$_2$ added, mixed and allowed to react for 6 minutes. A solution of 10% Al(NO$_3$)$_3$ (0.4ml) instead of AlCl$_3$ was added, mixed and allowed to complex for 6 minutes before adding 4 mls of 4% NaOH to neutralise the acidity. Double distilled water was added to top to the 10mls mark. For each sample, 2mls was placed into cuvet and topped to 4ml using double distilled water, its absorbance measured at 510nm and total flavonoid concentration estimated from the calibration curve with reference to casticin.

Results and discussion
Major phytochemical groups identified in artemisia annua samples
The qualitative phytochemical test results showed that A. annua from low land and high land areas had similar phytochemical groups including flavonoids and triterpenes (Table 1) the known antimalarial groups. Similarity in phytochemical constituents shows that the A.annua cultivated in the different regions of Uganda are of the same variety.

<table>
<thead>
<tr>
<th>Table 1: Phytochemical groups identified in Artemisia annua samples</th>
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<tr>
<td>Phytochemical group</td>
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<tr>
<td>Tannins</td>
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<tr>
<td>Reducing compounds</td>
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<td>Polyuronides</td>
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<td>Saponins</td>
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<tr>
<td>Alkaloid salts</td>
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<td>Anthracenosides</td>
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<tr>
<td>Coumarin derivatives</td>
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<tr>
<td>Steroid glycosides</td>
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<td>Flavonosides</td>
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Present (+), Present in abundance (++) , absent (-)

Aromatic oil constituents in artemisia annua L.
The major aromatic oil component in all three samples was camphor which constituted more than 75% of the oils. Camphor has been previously reported as the major aromatic oil component in A.annua cultivated in India$^{15}$ and in Iran$^{16}$. Unlike other aromatic components, camphor and borneol levels did not vary much between the regions (Figure 1). Camphor oil has been shown have mosquito repellent activity and provided up to 97.6% protection against Anopheles culicifacies$^{17}$.

Artemisinin and total flavonoid contents in powder and teas
Artemisinin content was highest in the sample from Fort portal in Western Uganda followed by that from Kabale in south western Uganda. The total flavonoids in samples from Wakiso in central was about half that from Fortportal and Kabale (Figure 4). These variations call for quality control and standardization of material for not only of artemisinin levels but also of total flavonoids. This is because flavonoids have been shown to play major role in use of A.annua for malaria treatment and prophylaxis$^{4,8,9}$.
The flavonoids content in Kabale and Fort Portal samples were similar to that previously reported in study on Brazilian A.annua cultivar which had 2.6% total flavonoid content\(^4\). The flavonoid contents in A.annua have been shown to vary with stage of growth, with highest amounts found during full bloom just like it is for artemisinin\(^1\)\(^8\). Although the time of harvesting the A.annua was not considered in this present study, farmers in Uganda generally harvest A.annua just before flowering which is the full bloom period. In lowland areas A.annua flowers much earlier than in highland areas perhaps due to higher temperatures and low rainfall compared to high land areas. The variations observed in this study could therefore be associated with the age of the plant at harvesting time.

**Figure 1:** Percentage of Aromatic oil constituents of *A. annua* by region.

**Figure 2:** Artemisinin calibration curve used for estimation of artemisinin content
Conclusion
This study reveals for the first time that the antimalarial components in A.annua vary with geographical area where the plant is cultivated. Standardization by source of A.annua raw materials used against malaria is vital for consistent clinical outcomes. A study on the variation of the components due to the age of the plant vis-à-vis geographical location is recommended to enable commercialization of the plant, and effective use by the communities.

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References