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Full Length Research Paper

In vitro evaluation of antiasthmatic activity of ethanol leaf extract of *Guiera senegalensis* j. F. GMEL (Combretaceae)

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Guiera senegalensis is a medicinal plant that is widely used in West Africa against many illnesses. In this work, the antiasthmatic properties of ethanol leaf extract of the *G. senegalensis* were evaluated by using various *in vitro* animal models. *In vitro* models like isolated guinea pig tracheal chain preparation, isolated guinea pig ileum and rabbit jejunum preparations were studied to evaluate possible mechanism by which extract shows relaxant activity. The preliminary phytochemical screening of the extract revealed the presence of alkaloids, anthraquinones, flavonoids, saponins and tannins. The study showed that the extract contained phytochemicals such as flavonoids, alkaloids, steroids, tannins and saponins. Isolated tracheal smooth muscles contractions induced by histamine was significantly (p<0.01) reduced by the extract, so also that induced by acetylcholine at concentrations of 4 and 8 mg/ml. The findings were similar to that of 0.4 µg/ml isoprenaline. Guinea pig and rabbit jejunum contractions induced by histamine and acetylcholine were significantly (p<0.01) reduced by the extract. The results of these studies indicated ethanol extract of *G. senegalensis* possess antiasthmatic activity.

Key words: Antiasthmatic activity, bronchoconstriction, Guiera senegalensis, guinea pig ileum.

INTRODUCTION

Asthma is a respiratory disease characterized by recurrent episodes of wheezing, chest tightness, cough and difficulty breathing brought about by bronchial constriction, inflammation, and excessive mucus secretion due to bronchial hyperesponsiveness (GINA, 2014). Although advancement in the treatment of asthma is on the increase, a great number of individuals have been affected by asthma. It has been estimated that about 300 million people in both developed and developing countries suffer asthma attacks worldwide (GINA, 2015). Global prevalence of asthma has been approximated to be 10% among children and 5% among adults population (Rathore et al., 2011). In Nigeria, asthmatic children has been reported to be between 5.1 and 14.3% (Musa and Aliyu, 2014) while estimate for adult Nigerians having asthma has been put to 10%

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Author(s) agree that this article remain permanently open access under the terms of the <u>Creative Commons Attribution</u> <u>License 4.0 International License</u> (Desalu et al., 2013). Prevalence of asthma has been estimated to increase by 59% worldwide in 2025 (Masoli et al., 2004). This can be attributed to increased industrialization and urbanization.

Although, pharmaceutical drugs are the major treatments given, a large number of patients worldwide (especially in Africa and Asia) still patronize traditional medicines as an alternative (WHO, 2013). It has been reported that 75 to 90% of the rural populations worldwide still depend on herbal preparations as their main source of medicines (Shettima et al., 2013). Research in traditional medicine is also increasing worldwide and has been encouraged by the World Health Organization because of the large number of patronizers of such kind of treatment for their healthcare (WHO 2013). Guiera senegalensis is used as a medicinal plant by many cultures and researches carried out on the medicinal uses of the plant turn out to give good results (Jigam et al., 2011). The plant has been used traditionally to treat various diseases including asthma and its use in treatment of pulmonary problems has been documented by Dénou et al., (2016) and Saraswathy et al. (2014). The plant could therefore be considered important in medical interventions and its potential as anti-asthmatic be exploited. This may generate data that can be used to complement pharmaceutical drugs for the benefit of humanity. This study therefore, aimed at evaluating the potential of ethanol extract of G.senegalensis (ELEGS) as anti-asthmatic.

MATERIALS AND METHODS

Drugs and chemical

The following drugs and chemicals were used in the experiment: Acetylcholine chloride (Sigma Chemicals, USA), histamine diphosphate (Sigma Chemicals, USA), atropine sulphate (Shandong Shenglu Pharmaceutical Co., Ltd.), isoprenaline hydrochloride (Sigma-Aldrid), mepyramine, sodium hydrogen carbonate (BDH Chemicals Ltd Poole, England), sodium chloride (Johnson and Solomon Ltd. London, England), D-glucose (BDH Chemicals Ltd. Poole, England), calcium chloride (BDH Chemicals Ltd. Poole, England), sulfuric acid, ferric chloride, acetic anhydride and ethanol.

Experimental animals

Adult guinea pigs (weighing 400 to 450 g) and rabbits (weighing 3.0 to 3.5 kg) were obtained from the animal house of the Department of Pharmacology and Therapeutics, Faculty of Pharmaceutical Sciences, Ahmadu Bello University, Zaria. The animals were kept in propylene cages under favourable conditions prior to the study. Water and feed were provided *ad libitum* according to the ethical guidelines of Ahmadu Bello University for the care and use of animals in research.

Plant

Fresh leaves of *G. senegalensis* were collected from Tofa Local Government Area in Kano State of Nigeria. Taxonomic identification

was done and voucher number (BUKHAN0032) was deposited at the herbarium of Biological Sciences Department, Bayero University, Kano.

Extraction

Leaves of the plant material were thoroughly washed with distilled water to remove dirt and soil and dried under the shade. The plant material was then pulverized into fine powder and the powder macerated in 70% ethanol for 48 h. The resultant extract was filtered and then through Whatman[®] Filter Paper No.1, 185 mm. The filtrate was concentrated in an oven at 40°C.

Preliminary phytochemicals screening

Phytochemicals screening of the ethanol leaf extract of *G. senegalensis* (ELEGS) were carried out to determine the presence of secondary metabolites such as alkaloids, steroids and flavonoids as described by Tiwari et al. (2011).

Isolated guinea pig trachea preparation

Guinea pig of either sex weighing 400 to 450 g were starved overnight but allowed free access to water. The animals were then sacrificed and the entire tracheae was dissected out and removed and cut into individual rings. Twelve rings were tied together and mounted in 25 ml organ bath containing Krebs physiological solution. The tissue was maintained at 37°C under tension of 1 g and constantly aerated with carbogen (Vogel and Vogel, 2002). The tissue was allowed to equilibrate during which the bath solution was replaced every 10 min. After the equilibrium period, the action of the extract alone was evaluated; three graded concentrations were selected and used for the study. Contraction was then induced by adding acetylcholine or histamine separately. When the contraction reached maximum (initial spasm), the concentration was recorded and after 5 min the standard drug (isoprenaline 0.4 µg/ml) was added (Kulkarni, 2005). Thereafter, the test extract (2, 4 and 8 mg/ml) was added serially in the presences of histamine and then acetylcholine. The tissues were observed for bronchodilation.

Isolated guinea pig ilea preparations

Guinea pig of either sex weighing 400 to 450 g were starved overnight but allowed free access to water. The animals were then sacrificed and then dissected. The intestine was removed and the distal portion was cut into pieces 2 to 3 cm long. Mesentery was removed and tyrode solution passed through the pieces of the tissue until the effluent was clear. The tissue was then mounted in 25 ml organ bath containing Tyrode physiological solution. The tissue was maintained at 37°C under tension of 1 g and constantly aerated (Vogel and Vogel, 2002). At the end of the equilibration period, increasing concentrations of histamine and the extract were added separately. The concentration of histamine that produced the submaximal response was then added. Five minutes after, the standard drug (mepramine 0.04 µg/ml) was added. The tissue responses were recorded. The tissue was then rinsed and allowed to equilibrate after which the test extracts (2, 4 and 8 mg/ml) was added serially in the presence of histamine. The tissues were observed for bronchodilation using microdynanometer.

In another set up of isolated guinea pig ileum preparation, effects of the acetylcholine and acetylcholine in presence of atropine, and graded concentrations of the extract were studied as described by Kulkarni (2005). Responses of the tissues were recorded.

Constituent	Inference
Alkaloids	+
Anthraquinones	+
Cardiac glycosides	-
Flavonoids	+
Phlabotanins	-
Saponins	+
Steroids	+
Tannins	+
Terpenoids	-

Table 1. Phytochemical constituents of ethanol leaf extract of *Guiera* senegalensis.

(+) present; (-) absent.

Isolated rabbit jejunum preparations

According to the method of Amos et al. (2000), overnight fasted rabbits (2.0 to 2.5 kg) were sacrificed and dissected. Segments of the rabbit jejunum of about 2 to 3 cm long were removed, washed with Tyrode solution and mounted in 25 ml organ bath and maintained at 37°C. The tissue was allowed to equilibrate and then, the effect of the various concentrations of the extract alone on the tissue was determined. After a washout period, various concentration that produced submaximal response determined. Using the concentration of acetylcholine that produced submaximal response, the procedure was repeated in the presences of atropine and then increasing concentrations of the extract. In another set up, the procedure was repeated using propranolol and isoprenaline in place of acetylcholine and atropine respectively.

Membrane stabilization assay

Bovine red blood cells (RBC) were obtained from slaughter house and made into 5% suspension. To 1 ml of the RBC suspension, 0.1 ml of 50 mM H_2O_2 was added and incubated at 37°C to initiate hemolysis. After 30 min, 4 ml distilled water was added to the mixture and centrifuged at 1000 rpm for 10 min and absorbance of the supernatant was then taken at 540 nm. This procedure was repeated in the presence of the various concentrations of the extract and percentage inhibition of hemolysis was calculated (Su et al., 2009).

Data analysis

The data obtained were expressed as Mean \pm standard error of mean (SEM) or percentages protection. Data were analyzed using one way analysis of variance (ANOVA), followed by Dunnett's posthoc tests. Values of p<0.05 were considered statistically significant. Results were presented in tables and charts.

RESULTS

Extraction

Maceration of the G. senegalensis leaf in 70% ethanol

yields 19.2% dark coloured extract.

Phytochemical screening

Preliminary qualitative phytochemical screening of ELEGS indicated the presence of alkaloids, flavonoids, saponins, steroids and tannins (Table 1). Similar study on the leaf extract conducted by Akuodor et al. (2013) also found these phytochemicals with the exception of anthraquinones.

Effects of ethanol leaf extract of *G. senegalensis* on isolated guinea pig trachea

The extract alone produced neither contraction nor relaxation. However, histamine (0.6 μ g/ml) induced-contraction was dose-dependently inhibited by the extract. The effect of the extract at 4 and 8 mg/ml was statistically significant (p<0.01) and was comparable to that of 0.4 μ g/ml isoprenaline (Figure 1). The extract also significantly (p<0.05) inhibited acetylcholine (0.4 μ g/ml) induced-contraction as compared to control, but lower than isoprenaline 0.4 μ g/ml (Figure 2).

Effect of ethanol leaf extract of *G. senegalensis* on isolated guinea pig ileum

Graded doses of the extract produced dose-dependent relaxation of isolated guinea pig ileum. Responses were all significant (p<0.05) compared to the baseline. Maximum relaxation was obtained at 3.2 mg/ml (Figure 3). On challenging the extract to histamine-induced contraction, significant and dose-dependent inhibition was observed. The extract at concentration of 1.6 mg/ml resulted to higher inhibition (p<0.001) than 0.2 μ g/ml mepyramine (Figure 4). Similar result was obtained on acetylcholine-induced contraction (Figure 5).

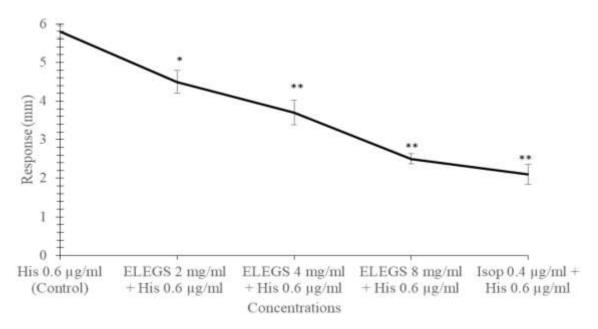


Figure 1. Effect of Ethanol Leaf Extract of *Guierasenegalensis* on Histamine Induced Contractions of Guinea Pig Trachea. Values represents mean \pm SEM (n=3). Data analysis was performed by ANOVA followed by Dunnett's post-hock test to determine significance (p<0.05)* (p<0.01)**. His: Histamine; Iso: Isoprenaline; ELEGS: Ethanol leaf extract of *Guiera senegalensis*.

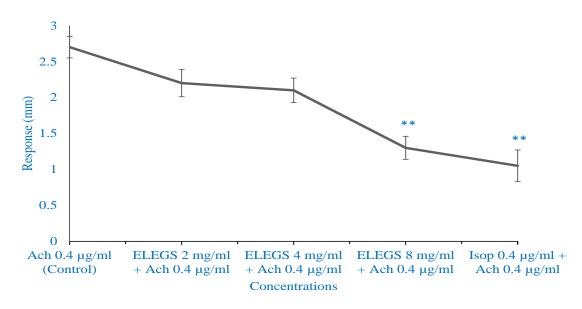
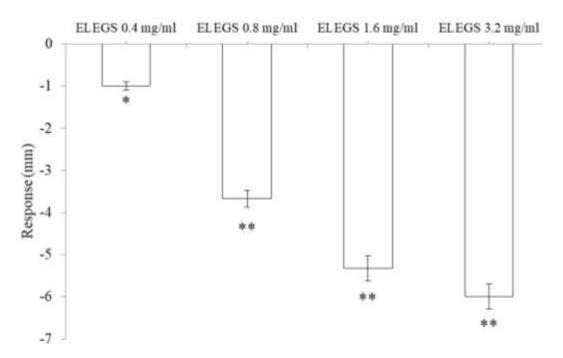


Figure 2. Effect of Ethanol Leaf Extract of *Guierasenegalensis*on Acetylcholine Induced Contraction of Guinea Pig Trachea. Values are expressed as mean \pm standard error of the mean (n=3). Statistical significances (p<0.05)* (p<0.01)** were determine by Dunnett's test after ANOVA. Ach: Acetylcholine; Iso + Isoprenaline; ELEGS: ethanol leaf extract of *Guiera senegalensis*.

Effect of ethanol leaf extract of *G. senegalensis* on rabbit jejunum

Dose-dependent inhibition of the spontaneous basal contractions of isolated rabbit jejunum was observed

when ELEGS was used alone. The contraction was almost abolished at 6.4 mg/ml (Figure 6). The contractile responses induced by acetylcholine (0.04 μ g/ml) were significantly (p<0.01) inhibited by graded doses of ELEGS in a dose-dependent manner. Atropine (0. 04



Concentration

Figure 3. Effect of ethanol leaf extract of *Guiera senegalensis* on isolated guinea pig ileum. Values are expressed as mean \pm standard error of mean (n = 3). Data analysis was performed by ANOVA followed by Dunnett's test to determine significance (p<0.05)*, (p<0.01)**. ELEGS: Ethanol leaf extract of *Guiera senegalensis*

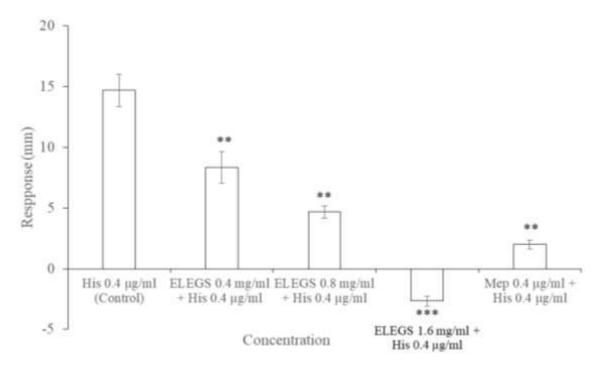


Figure 4. Effect of ethanol leaf extract of *Guiera senegalensis*on histamine induced contraction of guinea pig ileum. Values are expressed as mean \pm standard error of the mean (n=3). Statistical analysis was performed by ANOVA. Dunnet's post-hock was used to determine statistical significance (p<0.01)**, (p<0.001)***. His: Histamine; Mep: mepyramine; ELEGS: ethanol leaf extract of *Guiera senegalensis*.

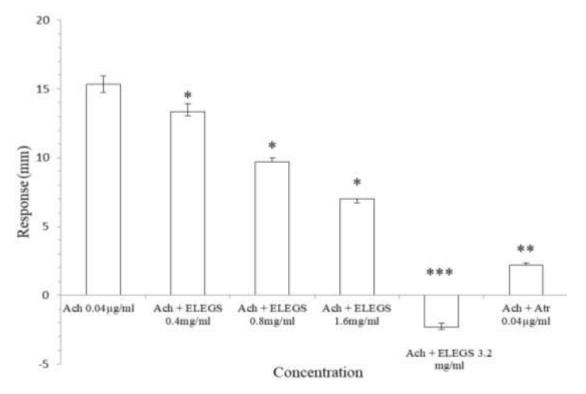


Figure 5. Effect of ethanol leaf extract of *Guiera senegalensis*on acetylcholine induced contraction of guinea pig ileum. Values are expressed as mean \pm standard error of the mean (n = 3). ANOVA was used for data analysis followed by Dunnett's post-hock to test for significance (p<0.05)* (p<0.01)** (p<0.001)***. Ach: Acetylcholine; Atr: atropine; ELEGS: ethanol leaf extract of *Guiera senegalensis*.

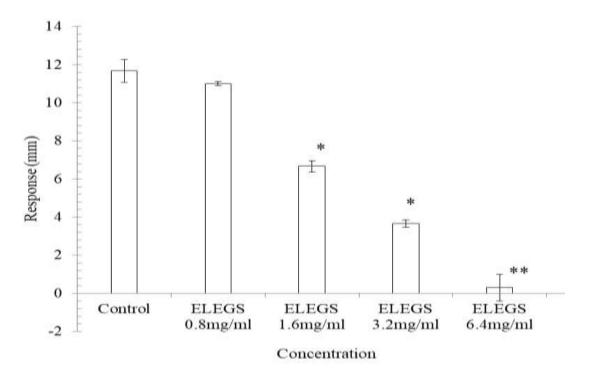


Figure 6. Effect of ethanol leaf extract of *Guiera senegalensis* on isolated rabbit jejunum. Values are expressed as mean \pm standard error of mean (n = 3). Data analysis was performed by ANOVA while significances (p<0.05)* (p<0.01)** were determined using Dunnett's post-hock test. Key: ELEGS: Ethanol leaf extract of *Guiera senegalensis*.

Treatment	Response (mm)
Ach 0.04 µg/ml (Control)	65.00 ± 4.5
Ach + ELEGS 0.8 mg/ml	47.33 ± 3.52**
Ach + ELEGS 1.6 mg/ml	38.33 ± 3.33**
Ach + ELEGS 3.2 mg/ml	30.66 ± 0.88**
Ach + ELEGS 6.4 mg/ml	11.33 ± 0.33***
Ach + Atropine 0.04 µg/ml	8.40 ± 0.70***

Table 2. Effects of ELEGS on acetylcholine induced contraction of rabbit jejunum.

Values are expressed as mean \pm standard error of mean (n=3). Data analysis was performed by ANOVA followed by Dunnett's test to obtain significance (p<0.01)**, (p<0.001)***. Ach: Acetylcholine; ELEGS: ethanol leaf extract of *Guiera senegalensis*.

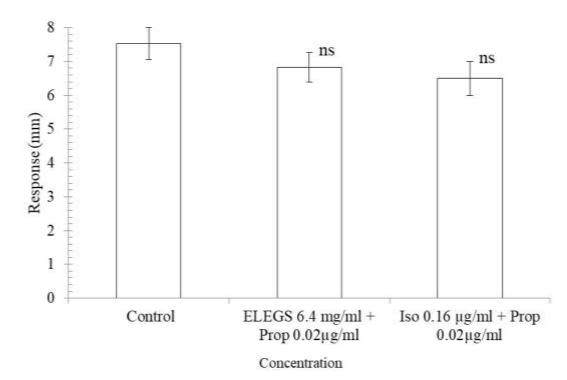


Figure 7. Effect of propranolol and ethanol leaf extract of *Guiera senegalensis* on rabbit jejunum. Values are expressed as mean \pm standard error of mean (n = 3). Data analysis was performed by ANOVA followed by dunnett's post-hock test. Iso: Isoprenaline; Prop: propranolol; ELEGS: ethanol leaf extract of *Guiera senegalensis*; ns: not significant (p>0.05).

 μ g/ml) also significantly (p<0.001) inhibited the contractile response induced by acetylcholine (Table 2).

Effect of ethanol leaf extract of *G. senegalensis* and propranolol on rabbit jejunum

Responses obtained from rabbit jejunum after addition of ELEGS (6.4 mg/ml) to baths pretreated with propranolol were not significantly different from the control. Similar effect was observed with 0.16 µg isoprenaline (Figure 7).

Membrane stabilization effect of ethanol leaf extract of *G. senegalensis*

The membrane stabilizing effect of ELEGS was assessed using erythrocytes hemolysis inhibition assay. Maximum protection was observed at ELEGS concentration 1 mg/ml compared to the control (Table 3).

DISCUSSION

Bronchial constriction in asthma is primarily mediated by

ELEGS (mg/ml)	Absorbance	% Inhibition
Control	0.3640 ± 0.032	-
0.1	0.3131 ± 0.013	13
0.2	$0.2630 \pm 0.022^*$	28
1.0	0.2247 ± 0.019**	38
2.0	0.3030 ± 0.014	16

 Table 3. Effect of ethanol leaf extract of Guiera senegalensis on erythrocytes membranes stability.

Values are expressed as mean \pm Standard Error of Mean (n=3). Data analysis was performed by ANOVA followed by Dunnetts post-hock test to determine the level of significance (p<0.05)*, (p<0.01)** existing between each value compared to the control. Absorbance was read at 540 nm. ELEGS: Ethanol leaf extract of *Guiera senegalensis*

histamine released from degranulated mast cells and by acetylcholine which cause contractions of tracheal smooth muscles and increased mucus secretion. Inflammation of the bronchi also sets due to lipid mediators from arachidonic acid pathway generated from phospholipids (Gosh, 1984). Clinical membrane interventions so far aim at maintaining normal airway muscles tone by the use of bronchodilators and antiinflammatory agents to reverse bronchoconstriction. Therefore, substance that can reverse bronchoconstrictions will possibly relieve symptoms of asthma. For this, the antihistaminic as well as the antimuscarinic activities of the plant extract were investigated using animals' isolated tissues. Smooth muscles are rich in many kinds of receptors with different localization specifications. Airway smooth muscles are rich in histamine, acetylcholine, and β_2 adrenergic receptors. Distribution of these receptors and their responses to agonists and antagonists is similar in both humans and guinea pigs. Therefore, guinea pigs are used in contrast to rats and mice.

In the present study, in vitro anti-asthmatic effect of the ethanol leaf extract of Guiera senegalensis (ELEGS) was evaluated. In isolated trachea guinea pig tracheal chain preparations, ELEGS was also found to possess antihistaminic and antimuscarinic effects. Although it did not exhibit any effect when used alone, the extract at concentrations of 8 mg/ml was able to significantly (p<0.01) inhibit contractions induced by both spasmogens (histamine and acetylcholine) on the tracheal chains. The action of the extract was found to be similar to that of the standard drug (isoprenaline).

Histamine and muscarinic receptors are the dominant receptors in guinea pig ileum and this makes it a good tissue for studies involving these spasmogens. ELEGS exhibited smooth muscle relaxing effect on guinea pigs ileal strips and was able to significantly inhibit contractions induced by histamine (p<0.01 to 0.001) and acetylcholine (p<0.05 to 0.01) in a dose-dependent manner. Atropine (antimuscarinic) was used against

acetylcholine while mepyramine (antihistamine) was used against histamine for comparison with the extract. These pharmacological antagonists significantly (p<0.01) inhibited the ileal induced contractions but to a lesser effect than 3.2 mg/ml of the extract. At the 3.2 mg/ml, ELEGS completely abolished the contractions-induced by both histamine and acetylcholine and resulted to relaxation of the tissues.

On isolated rabbit jejunum, the extract when given alone was able to significantly (p<0.05 to 0.01) reduced the basal spontaneous contractions as well as contractions induced by acetylcholine (p<0.01 to 0.001) in dose-dependent pattern. Similar result was observed with atropine. These indicate that ELEGS has a nonspecific spasmolytic activity on smooth muscles that may probably be due to β -adrenergic stimulatory (Martin et al., 1994), histamine H1 (Popa et al., 1984) and/or muscarinic inhibition (Linden et al., 1993). However, result obtained from rabbit jejunum after addition of ELEGS up to 6.4 mg/ml to baths pretreated with propranolol was not significantly different from the control. Similar effect was observed with 0.16 µg isoprenaline. Propranolol is a nonselective β adrenergic receptor blocker that binds to all β receptors and inhibits their activity. This suggests that, ELEGS might be acting on the same receptors resulting in smooth muscle relaxation by activation of β_2 adrenoceptors.

Allergic asthma is a chronic inflammatory process occurring due to exposure to allergen resulting in the activation of T-lymphocyte with subsequent release of inflammatory mediators. Immuno-modulating agents are useful in the treatment of asthma by inhibiting the antigen-antibody (AG-AB) reaction and thereby inhibiting the release of inflammatory mediators. Membrane stabilization assay was used to study the possible inhibition of lipid mediators of inflammation in asthma. Percent inhibition of erythrocytes hemolysis was calculated and the ELEGS at all concentrations used was able to protect the membranes of erythrocytes from possible damage. Maximum effective concentration was observed at 1 mg/ml.

Drugs effective in asthma are mostly steroidal in nature. ELEGS contained steroid, alkaloids, flavonoids, saponins, anthraquinones and tannins. These secondary metabolites have been known to possess various biological activities including antibacterial, antioxidant, anticancer, spasmolytic and anti-inflammatory effects (Parmar et al., 1999). Flavonoids have been known to inhibit the release of several mediators of inflammations such as prostaglandins, histamine, serotonin and bradykinins by inhibiting the biosynthetic pathways of inflammatory mediators (Macauder, 1986). Steroids are known for their maintenance of membrane integrity by inhibiting lipoxygenase and cyclo-oxygenase pathways and are used for their anti-inflammatory properties (Speroni et al., 2005). So antiasthmatic activity showed by G. senegalensis might be because of these chemical moieties.

CONFLICT OF INTERESTS

The authors have not declared any conflict of interest.

REFERENCES

- Akuodor GC, Essien AD, David-Oku E, Chilaka KC, Akpan JL, Ezeokpo B, Ezeonwumelu JOC (2013). Gastroprotective effect of the aqueous leaf extract of Guiera senegalensis in albino rats. Asian Pac. J. Trop. Med. 6(10):771-775.
- Amos S, Gammaniel K, Adamu M, Bukar B, Akah P, Wambebe C (2000). Pharmacological effects of aqueous extracts of *Chrysanthellum indicum* on gastrointestinal smooth muscles. J. Herbs Spices Med. Plants. 7(3):45-53.
- Dénou A, Togola A, Haïdara M, Sanogo R, Diallo D. and Koumaré M. (2016). Review on Phytochemistry and Pharmacological Aspects of Guiera Senegalensis J. F. Gmel (Combretaceae). Int. J. New Technol. Res. 2(3):30-32.
- Desalu OO, Onyedum CC, Adeoti AO, Ozoh OB, Fadare JO, Salawu FK, Danburam A, Fawibe AE, Adewole OO (2013). Unmet needs in asthma treatment in a resource-limited setting: findings from the survey of adult asthma patients and their physicians in Nigeria. Pan. Afr. Med. J. 16:20.
- Global Initiative for Asthma (GINA) (2014). The Global Asthma Report. P40. Available at: globalasthmareport.org
- Global Initiative for Asthma (GINA) (2015). Pocket guide for asthma management and prevention. P3. Available at: http://ginasthma.org/wp-

content/uploads/2016/01/GINA_Pocket_2015.pdf

- Gosh MN (1984). Fundamental of Experimental Pharmacology. 2nd edition. Calcutta: Scientic Book Agency.
- Jigam AA, Akanya HO, Dauda BE, Ogbadoyi EO (2011). Antiplasmodial, analgesic and anti-inflammatory effects of crude *Guiera senegalensis* Gmel (Combretaceae) leaf extracts in mice infected with Plasmodium berghei. J. Pharmacogn. Phytother. 3(10):150-154.
- Kulkarni SK (2005). Handbook of Experimental Pharmacology, 3rdedition. pp. 92-93.
- Linden A, Bergendal A, Ullman A, Skoogh BE, Lofdahl CG (1993). Salmetrol, formetrol, and salbutamol in the isolated guinea-pig trachea: Differences in maximum relaxant effect and potency but not in functional antagonism. Thorax 48(5):547-553.

- Macauder PJ (1986). Flavonoids affect acetylcholine, prostaglandin E2 and antigen-mediated smooth muscle contraction. Prog. Clin. Biol. Res. 213:489-492.
- Martin CA, Naline E, Bakdach H, Advenier C (1994). Beta 3adrenoceptor agonists, BRL 37344 and SR 58611A, do not induce relaxation of human, sheep and guinea-pig airway smooth muscle in vitro. Eur. Respir. J. 7(9):1610-1615.
- Masoli M, Fabian D, Holt S, Beasley R (2004). Global Initiative for Asthma: executive summary of GINA Dissemination Committee report. Allergy 59:469-478.
- Musa BM, Aliyu MD (2014). Asthma prevalence in Nigerian adolescents and adults: systematic review and meta-analysis. Afr. J. Respir. Med. 10(1):4
- Parmar VS, Jha A, Bisht KS, Taneja P, Singh SK, Kumar A, Jain R, Olsen CE (1999). Review Article Number 138: Constituents of the yew trees. Phytochemistry 50:1267-304.
- Popa VT, Somani P, Simon P, Simon V (1984). The effect of inhaled verapamil on resting bronchial tone and airway constriction by histamine and acetylcholine in normal and asthmatic subjects. Am. Rev. Respir. Dis. 130(6):1006-1013.
- Rathore C, Dutt KR, Sahu S, Deb L (2011). Antiasthmatic activity of the mEthanol extract of *Physalis angulata* Linn. J. Med. Plants Res. 5(22):5351-5355.
- Saraswathy GR, Sathiya R, Anbu J, Maheswari E (2014). Antitussive Medicinal Herbs - An Update Review. Int. J. Pharm. Sci. Drug Res. 6(1):12-19
- Shettima AY, Karumi Y, Sodipo OA, Usman H, Tijjani MA (2013). Acute and sub-acute study on ethylacetate root extract of *Guiera* senegalensis in rats. J. Pharm. Sci. Innov. 2(1):1-6.
- Speroni E, Cervellati R, Innocenti G, Costa S, Guerra MC, Dall S, Govonid P (2005). Anti-inflammatory, anti-nociceptive and antioxidant activities of *Balanites aegyptiaca*. J. Ethnopharmacol. 98(2):117-121.
- Su XY, Wang Z, Liu J (2009). *In vitro* and *in vivo* antioxidant activity of *Pinus koraiensis* seed extract containing phenolic compounds. Food Chem. 117:681-686.
- Tiwari P, Kumar B, Kaur M, Gurpreit K, Harleen K (2011). Phytochemical Screening and Extraction: Review. Int. Pharm. Sci. 1(1):98-106.
- Vogel GH, Vogel WH (2002). Drug Discovery and Evaluation. Pharmacological Assays. IInd edition, Germany, Springer Verlag Berlin Heideberg Newyork, pp. 419-425.
- World Health Organization (WHO) (2013). Traditional Medicine Strategy
2014-2023.Availableat:at:
 - http://www.who.int/medicines/publications/traditional/policy/en/