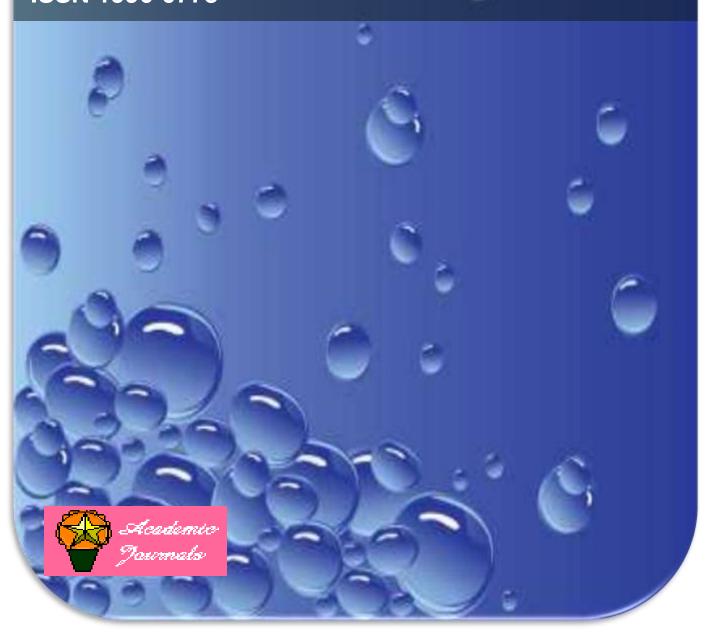


Volume 11 Number 1, January 2017 ISSN 1996-0778



ABOUT AJBR

The African Journal of Biochemistry Research (AJBR) (ISSN1996-0778) is published Monthly (one volume per year) by Academic Journals.

African Journal of Biochemistry Research (AJBR) provides rapid publication (monthly) of articles in all areas of Biochemistry such as Nutritional biochemistry, Analytical biochemistry, Clinical Biochemistry, Human and Plant Genetics, Molecular and Cell Biology, Enzymology, Toxicology, Plant Biochemistry, Biochemistry Education etc. The Journal welcomes the submission of manuscripts that meet the general criteria of significance and scientific excellence. Papers will be published shortly after acceptance. All articles are peer-reviewed.

Contact Us

Editorial Office: ajbr@academicjournals.org

Help Desk: helpdesk@academicjournals.org

Website: http://www.academicjournals.org/journal/AJBR

Submit manuscript online http://ms.academicjournals.me/

Editor

Prof. Johnson Lin

School of Biochemistry, Genetics, Microbiology and Plant Pathology University of KwaZulu-Natal (Westville) Private Bag X 54001, Durban Republic of South Africa

Associate Editors

Gregory Lloyd Blatch

Dept Biochemistry Microbilogy& Biotechnology Rhodes University Grahamstown 6140 South Africa

Dr. SerapYalin

Mersin University, Faculty of Pharmacy, Department of Biochemistry, YenisehirKampusu, Mezitli 33161 Mersin/Turkey

Dr. Om Prakash Gupta

Directorate of Wheat Research (ICAR) Post Box-158, A grasainMarg, Karnal-132001, Haryana, India

Editorial Board

Dr. Desouky A.M. Abd-El-Haleem

Biological Sciences Department, College of Arts and Sciences, Qatar University, Doha, Qatar

Dr. S.K. Trigun

Biochemistry and Molecular Biology Section, Banaras Hindu University Varanasi-221005, India

Dr. ImedGallouzi

McGill University, Biochemistry Department, 3655 Promenade Sir William OslerMontreal, Quebec, H3G 1Y6, Canada

Dr. Ashraf A Khalil

Protein Technology Lab, Mubarak City for Science, New Borg Elarab, Alexandria, Egypt.

Dr. Stanley Mukanganyama

Department of Biochemistry, University of Zimbabwe, Box MP 167, Mount Pleasant, Harare, Zimbabwe

Prof. Salah A. Sheweita

Taibah University, Faculty of Medicine, Department of Biochemistry, PO Box 30001, Madinah, Saudi Arabia

Dr Oluwafemi O Oguntibeju

Department of Clinical Biochemistry, School of Medicine, Spartan Health Sciences University, P.O. Box 324, Vieux Fort, St Lucia, West Indies

Dr. Robert L. Brown

USDA ARS, Southern Regional Research Center 1100 Robert E. Lee Blvd., New Orleans, LA 70124

Dr. Edward Eteshola

Biomedical Engineering Center Davis Heart and Lung Research Institute Ohio State University 473 W. 12th Avenue Columbus, OH 43210

G. Suresh Kumar

Senor Scientist and Head
Biophysical Chemistry Laboratory
Indian Institute of Chemical Biology
Council of Scientific and Industrial Research
Jadavpur,
Kolkata 700 032,
India

Xu Lι

Department of Biochemistry and Molecular Biology Colorado State University Fort Collins, CO 80523-1870 USA

Mohammed A.A Sarhan

Dept. Biological Sciences Faculty of Science King Khalid University Saudi Arabia

MehrdadBehmanesh

Department Of Genetics School Of Science P.O.Box 114-175 Tehran Iran Iran

Hans Verhagen

P.o Box 1 3720 Ba Bilthoven The Netherlands Netherlands

P.K.Sumodan

Post Graduate Department Of Zoology Government College Madappally India India

BalesengMoseki

University Of Botswana Botswana

Bhaskar C. Behera

Agharkar Research Institute Plant Science Division India India

Luiz Claudio Miletti

Universidade Do Estado De Santa Catarina Brasil

Oladipo Gabriel Sunday

University Of Port Harcourt Port Harcourt-Nigeria Nigeria

Basiouny Ahmed El-Gamal

Biochemistry Department Faculty Of Science Alexandria University Egypt

AminigoEbiokpo Rebecca

University Of Port Harcourt Portharcourt-Nigeria Nigeria

JiaZeng

Department Of Bioengineering Central South University Changsha Hunan 410083 P.R.China China

Adenike Kuku

ObafemiAwolowo University Department Of Biochemistry Nigeria

Elsayed Hafez

Genetic Engineering and Biotechnology Research Institute Egypt

Gabriella Castoria

Via L. De Crecchio 7 -80138 Naples Department Of General Pathology Italy

SalwaSeddik Abdel-Latif

21 Elbatal Ahmed Abdel Aziz Elmohandesien Giza Egypt

Erasto Vitus Mbugi

Muhimbili University Biochemistry Department School Of Medicine India

Mohamed Rholam

Université Paris7 - Denis-Diderot France

Hooi Ling Foo

Universiti Putra Malaysia Malaysia

JayanthRao

Biochemistry And Nutrition Cftri Mysore India

Maznah Ismail

Universiti Putra Malaysia

Svetlana Lutsenko

Oregon Health & Science University USA

Gabriel Ugwem

Rivers State University Of Science And Technology P.M.B. 5080 Port Harcourt Nigeria

HariChhatpar

Dept. Of Microbiology & Biotechnology Centre Faculty Of Science M.S.University Of Baroda Vadodara 390 002 Baroda India

MahiuddinAlamgir

The University Of New South Wales Sydney Nsw-2052 Australia

Sheeja Samuel Edwin

B.R Nahata College of Pharmacy & Research Centre India

William Cho

Room 1305 13/F Block R Department of Clinical Oncology Queen Elizabeth Hospital 30 Gascoigne Road Kowloon Hong Kong

Dr. SurainiAbd-Aziz

Universiti Putra Malaysia Malaysia

Dr. Mustafa NumanBucak

Lalahan Livestock Central Research Institute Lalahan Ankara Turkey

Alparslan Kadir Devrim

Department Of Biochemistry Faculty of Veterinary Medicine Kafkas University 36040 Kars Turkey

Vasudev R. Thakkar

Sardar Patel University Brd School of Biosciences Sardar Patel University Nagar

Prof. Emmanuel Anosike

Department Of Biochemistry University Of Port Harcourt Nigeria

Dr. Usama Beshay

New Bourg El-Arab City, Research Area Alexandria 21934 Egypt

Dr. Ramar Perumal Samy

Department of Anatomy Yong Loo Lin School of Medicine National University of Singapore Singapore

Dr. Shin-ichi ONO

Laboratory of Clinical Pharmacy College of Pharmacy, Nihon University Japan

Prof. Lawal Bilbis

Biochemistry Department UsmanuDanfodiyo University Sokoto Nigeria

Dr. Adriana G. Chicco

Department of Biochemistry University of Litoral, Santa Fe Argentina

Prof. Zia-Ur Rahman

Department Of Physiology and Pharmacology University Of Agriculture Falsalabad Pakistan

Dr. Oluwole Ariyo

Allen University USA

Prof. Francisco Torrens

Institut Universitari de Ciència Molecular Universitat de València Spain

Prof. Belkhodja Moulay

University of Senia Oran Algeria

Dr. Hossam M Ashour

Department of Microbiology and Immunology Faculty of Pharmacy, Cairo University Egypt

Dr. Fidelis Ocloo

Biotechnology and Nuclear Agriculture Research Institute/GAEC Ghana

Ass. Prof. Alfonso Baldi

Dept. Biochemistry, Sect. Pathology Second University of Naples, Italy

Dr. Anandh Babu Pon Velayutham

Department of Human Nutrition Foods and Exercise 253 Wallace Hall Virginia Tech Blacksburg VA 24061 USA

Dr. Tapan K. Chaudhuri

Department of Biochemical Engineering and Biotechnology Indian Institute of Technology Delhi, HauzKhas New Delhi-110016, India.

Dr. Rong Zhang

Shenyang Pharmaceutical University China

Ass. Prof. Tzong-Jih Cheng

Department of Bio-Industrial Mechatronics National Taiwan University Taiwan

Dr. Zuyong Xia

Department of Radiology, 1201 Welch Rd, Room P089, Stanford, CA 94301 USA

Dr. Pratap Kumar Das

Indian Institute of Chemical Biology India

Dr. Vasudeo Pandharinath Zambare

Advanced Enzyme Technologies Ltd India

Dr. A M Mujumdar

Agharkar Research Institute India

Prof. Christine Clayton

ZMBH ImNeuenheimer Feld 282 69120 Heidelberg Germany

Prof. Rekik Boul baba

ESA Mateur Département

Département des sciences et techniques de productions animales Tanzania

Dr. Farhad Mirzaei

National Dairy Research Institute, NDRI Karnal India

Dr. ROUABHI Rachid

Biology Department Tebessa University. Algeria

Prof. Vaclav Vetvicka

University of Louisville USA

Dr. Ramesh Putheti, Ph.D

Research scientist
Actavis Pharmaceuticals
10065 red run blvd,owings mills Blvd,Maryland.USA.21030

Prof. Dr. Mustafa NAZIROGLU

Head of Department of Biophysics Medical (TIP) Faculty, SuleymanDemirel University Cunur, TR-32260 Isparta TURKEY

Dr. José Luis Arias Mediano

GrupoInvestigaciónFarmaciaPráctica (CTS-205)
Dept. Farmacia y TecnologíaFarmacéutica
Facultad de Farmacia
Campus Universitario de Cartuja, s/n Universidad de
Granada
18071 Granada.

Ahmed Malki, PhD

Lecturer of Biochemistry and Molecular Biology Biochemistry Department Fcaulty Of Science Alexandria University Alexandria, Egypt

Dr. Alireza Seidavi (PhD)

Assistant Professor of Animal and Poultry Nutrition, Department of Animal Science, College of Agriculture, Islamic Azad University, Rasht Branch, Rasht, Iran

Amani S. Awaad

Professor of pharmacognosy, Chemistry Department Faculty of Sciences, King Saud University . Riyadh. KSA. P.O. Box 22452, Riyadh 11495. Saudi Arabia

Dr. Abdel-TawabMossa

Environmental Toxicology Research Unit (ETRU), Pesticide Chemistry Department, National Research Centre, Dokki, Egypt

Dr. Amal A. Mohamed

Plant Biochemistry Department, Agriculture Division - National Research Center, 31-El-Tahrir St., Dokki, Cairo – Egypt

Dr. Anabella Gaspar

Department of Biochemistry, University of Pretoria, South Africa

Dr. Anna Janecka

Department of Biomolecular Chemistry, Medical University of Lodz, Mazowiecka 6/8, 92-215 Lodz, Poland

Dr. Caser Abdel

Horticulture Department, Dohuk University, Iraq

Dr. David Sheehan

Dept Biochemistry, University College Cork, Ireland

Dr. Dayananda Chandrappa

Center for Bioenergy,
Department of Life and Physical
Sciences,
Cooperative Research,
Lincoln University,
Jefferson City,
USA

Dr. Elsayed Abdelaal

Special Graduate Faculty, University of Guelph, Onatrio, Canada

Dr. Etienne Marbaix

CELL Unit, de Duve Institute, UCL-75.41, 75 avenue Hippocrate, B-1200 Bruxelles, Belgium

Dr. Gary L. Firestone

Department of Molecular and Cell Biology, University of California, Berkeley, CA, 94720, USA

Dr. Henryk Zielinski

Institute of Animal Reproduction and Food Research, Polish Academy of Sciences, Poland

Dr. Irshad A. Nawchoo

Department of Botany, University of Kashmir, India

Dr. LuchaiButkhup

Department of Biotechnology, Faculty of Technology, Mahasarakham University, Mahasarakham 44000, Thailand

Dr. LuminitaVladescu

Department of Analytical Chemistry, Faculty of Chemistry, University of Bucharest, Romania

Dr. Mira Debnath

School of Biochemical Engineering, Institute of Technology - Banaras Hindu University, Varanasi, India

Dr. Nilesh S. Panchal

Department of Biosciences, Saurashtra University, Rajkot-360005, Gujarat. India

Dr. Rayappa A. Balikai

University of Agricultural Sciences, Dharwad, Karnataka- 580 005, India

Dr. SaadTayyab

Institute of Biological Sciences, University of Malaya, 50603 Kuala Lumpur, Malaysia

Dr. Shijun Fu

Institute of Health Sciences, Shanghai Institutes for Biological Sciences, Chinese Academy of Sciences and Shanghai Jiao Tong University School of Medicine, Shanghai, P. R. China

Dr. Shiming Zhang

Weis Center for Research, Geisinger Clinic, Danville, Pennsylvania, USA

Dr. Thomas Efferth

Department of Pharmaceutical Biology, Institute of Pharmacy and Biochemistry, University of Mainz, Heidelberg, 55128 Mainz, Germany

African Journal of Biochemistry Research

Table of Contents: Volume 11 Number 1 January 2017

ARTICLE

Effects of Xylopia aethiopica fruit extract on isolated rabbit ileum Okwari O.O, Obembe A.O, Jeje S.O, Honesty E.E and Osim E.E

1

academicJournals

Vol. 11(1), pp. 1-5, January 2017 DOI: 10.5897/AJBR2014.0775 Article Number: 0F1657462351 ISSN 1996-0778 Copyright © 2017 Author(s) retain the copyright of this article http://www.academicjournals.org/AJBR

African Journal of Biochemistry Research

Full Length Research Paper

Effects of Xylopia aethiopica fruit extract on isolated rabbit ileum

Okwari O.O¹, Obembe A.O², Jeje S.O^{1*}, Honesty E.E² and Osim E.E²

¹Department of Physiology, Cross River University of Technology, Okuku Campus, Nigeria. ²Department of Physiology, University of Calabar, Calabar, Nigeria.

Received 6 May, 2014; Accepted 29 March, 2016

In this study, pharmacological reactivity of rabbit ileum to crude extract of fruits of *Xylopia aethiopica* in the presence of other agonist and antagonist agents was evaluated. The contractility of the isolated rabbit ileum was recorded using kymograph. The effects of the crude extract of *X. aethiopica* on rabbit ileum were independently determined using a dose range of 10⁻⁸ to 10⁻⁴ mg/ml. However, to determine the effects of the extract on rabbit ileum in the presence of other agonist and antagonist a constant dose of 10⁻⁴ mg/ml of the extract was used. The result showed that the crude extract of *X. aethiopica*, nifedipine only and serotonin only produced a significant (p<0.001) dose dependent relaxation of the ileum muscle. Addition of the extract to nifedipine shifted the graph to the right. Administration of CaCl₂ alone produce a dose dependent contraction of the rabbit ileum with negative correlation coefficient (p<0.001). Co-administration of CaCl₂ with the extract shifted the CaCl₂ response curve to the left. Addition of extract to serotonin caused a significant (p<0.01) decreased in the relaxation of the rabbit ileum. In conclusion, this study suggests that *X. aethiopica* has smooth muscle relaxant effects on rabbit ileum and as such may reduce gastrointestinal motility.

Key word: Xylopia aethiopica, smooth muscle, lleum, motility, agonist.

INTRODUCTION

Xylopia aethiopica, has both nutritional and medicinal uses. It is a member of the custard apple family, Annonaceae, and is used as a spice in various traditional dishes of Western and Central Africa (Choumessi et al., 2012). It is one of the plants species man has discovered in the search for food and health care (Okwari et al., 2014).

Phytochemical analysis shows that the plant contain alkaloid (Ekong and Ogan, 1968), proteins, carbohydrate,

lipids, crude fibers in various concentrations, some minerals such as calcium, potassium and iodine (Berminas et al., 1999).

X. aethiopica has been applied in ethnomedicine in the treatment of cough, bronchitis, dysentery and female sterilization. It is believed to aid uterine contraction and is applied as an abortifacient agent (Iwu, 1993). Some of its investigated uses include antibacterial and antiseptic properties (Okwari et al., 2014).

*Corresponding author. Email: dhikrilat@yahoo.com

Author(s) agree that this article remains permanently open access under the terms of the <u>Creative Commons Attribution</u> <u>License 4.0 International License</u>

Nwafor and Kalio (2006) reported that aqueous extract of X. aethiopica caused a significant reduction in the gestation period, number of litters lost after seven days and weight of litters. In line with this, Ebong and Shode (2000) reported that the aqueous and petroleum ether extract of X. aethiopica possesses potent spasmolytic properties on the smooth muscles of the rat uterus. However, contrary to this claim, the study on the effect of aqueous and methanol extract of the fruits of the plant on gastrointestinal motility shows that, it has a smooth muscle relaxant effects in the gastrointestinal tract (Ebong et al., 1995). The contradiction in this report may be due to differences in the receptor activated by the extract of the plant in the uterine muscle and gastrointestinal tract. Hence, this study aims at evaluating the effect of aqueous extract of fruit of X. aethiopica on the motility of the rabbit intestine in the presence of other agonist and antagonist agents.

METHODOLOGY

Chemicals used

The chemical and drugs used for this study were obtained from sigma (United Kingdom) and prepared freshly by dissolving in distilled water.

Preparation of plant extract

Dried fruit of *X. aethiopica* were bought from Watt market in Calabar Cross River State. It was authenticated at Department of Botany University of Calabar. It was then grounded to a coarse powder using an engine grinder. The ground *X. aethiopica* was then subsequently weighed and 100 g of dried ground fruit was dissolved in 500 ml of deionized water (Ebong and Shode, 2000). The solution was left for 24 h before filtration with Whatman filter paper Number 1. The filtrate was then dried in an oven at about 40°C.

Preparation of isolated tissue

The animal for this study was fasted for 24 h prior to the beginning of the study to ensure complete emptying of the small intestine. The rabbit was then killed by cervical dislocation to ensure that gastrointestinal tract (GIT) reflexes were not inhibited. A midline incision was made on the abdomen along the *linea alba* to expose the small intestine. As soon as possible, a piece of the proximal ileum free from the mesentery near the duodenum was removed as it shows greater spontaneous activity as compared to the distal ileum. This part was then immersed in ice-cold saline and cut into small segments of about 3 to 5 cm long. The tissues were continually aerated with the aid of an aerator.

Experimental procedure

With the aid of threaded suture needle, a piece of thread was attached to each end of the tissue segment. At one end, the thread was made into a small loop and attached to the hook in the organ bath. The other piece of thread was attached to the writing lever of the kymograph with plasticine. The tissue was bathed with tyrode

solution (the solution composed of NaCl: 0.8%, KCl: 0.02%, NaHCO₃: 0.01%, NaHPO₄: 0.005%, CaCl₂: 0.02%, Glucose: 0.1%, Deionized water to make up 100 ml) and allowed to equilibrate for about 20 to 30 min. The bathing solution was flushed out at intervals of 10 min. The kymograph drum was set to a rotating velocity of 0.01 revolution/s with a speed of 0.1x0.5 mm/s. After equilibration period, the basal equilibrium response was obtained after which graded doses of the extract was added to the tissue and its result obtained. The tissue was flushed 3 to 4 times before the addition of other drugs. Each drug effect was considered alone using their graded doses and thereafter with 10⁻⁴ mg/ml of extract. The procedure was repeated five times using new ileal strips from different rabbit. The temperature of the organ bath was maintained at 37°C.

Statistical analysis

Results were expressed as Mean ± Standard Error of Mean (SEM). In all cases, the comparison of the different sets of data was done by using the unpaired student t-test. P-value of <0.05 was considered statistical significant. Correlation and regression analysis was done to show association.

RESULTS

The plant extract independently elicited dose-dependent relaxation of the rabbit ileum (correlation coefficient = 0.9990). The lowest concentration of the extract (10⁻⁴ mg/ml) produce 8.90±0.19% decrease in contraction of the smooth muscle. While the highest dose (1 mg/ml) produce 44.51±0.95% relaxation of the rabbit ileum (Figure 1). Nifedipine and serotonin also independently produce relaxation of the rabbit ileum (Figures 2 and 4). The correlation coefficient between dose and response was found to be significant at p<0.01 for both nifedipine and serotonin. Introduction of Cacl₂ alone evoked contraction of the smooth muscle in a dose dependent manner (p<0.01). The response of the smooth muscle to graded doses of nifedipine and CaCl₂ has a negative correlation coefficient (-0.9660 and -0.9766, respectively).

Addition of 10⁻⁴ mg/ml of the extract to nifedipine increased the relaxation of the smooth muscle with maximal response obtained at a dose of 10⁻⁴ mg (Figure 2). Co-administration of CaCl₂ with the 10⁻⁴ mg/ml of extract reduced the contractile response of the smooth muscle (Figure 3). Serotonin administration alone produces dose dependent increase in relaxation of the rabbit ileum. Addition of extract (10⁻⁴ mg/ml) to serotonin caused a significant decrease in the relaxation of the rabbit ileum (Figure 4).

DISCUSSION

The results obtained from this study showed that the extract of *X. aethiopica* produced a dose dependent relaxation of the rabbit ileum. This is in line with the report of Ebong et al. (1995), who reported that the aqueous and methanol extracts of the fruit of *X. aethiopica* has a

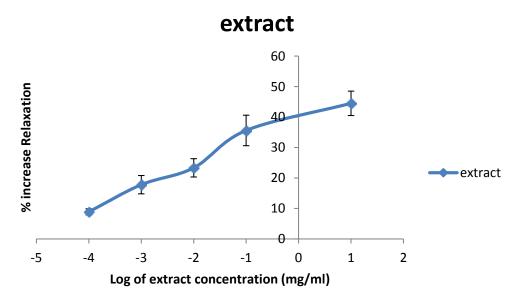


Figure 1. Effects of graded doses of the extract of Xylopia aethiopica on motility of the rabbit ileum.

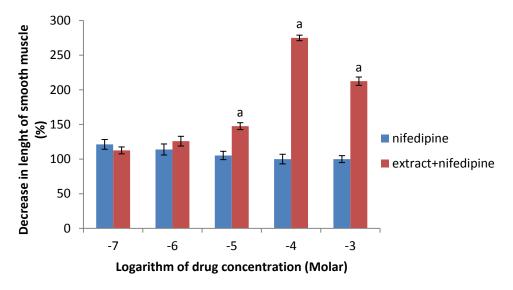


Figure 2. Effects of graded doses of Nifedipine + 10^{-4} mg/ml of *Xylopia aethiopica* on motility of rabbit ileum (a= p<0.05).

smooth muscle relaxant effect on gastrointestinal smooth muscle. The lowest concentration of the extract produced a decrease in contraction, but at high concentration, the extract produce strong relaxation. The smooth muscle relaxant effects of the extract may be due to the presence of alkaloids in the extract, since alkaloids have been previously implicated in the relaxation of smooth muscle (da Silva et al., 1999). Administration of nifedipine, a noncompetitive calcium channel blocker produced relaxation of the rabbit ileum. As the concentration of nifedipine

increased, the relaxation of the rabbit ileum decreased dose dependently. Addition of extract to graded doses of nifedipine increases the relaxation of smooth muscle of ileum. This may suggest that the extract and nifedipine are both acting through different receptor.

Administration of CaCl₂ produces contraction of the smooth muscle. This contractile response is reduced with increase concentration of CaCl₂. Addition of extract to graded doses of CaCl₂ further reduced the contractile response of the smooth muscle of ileum. It may be

4 Afr. J. Biochem. Res.

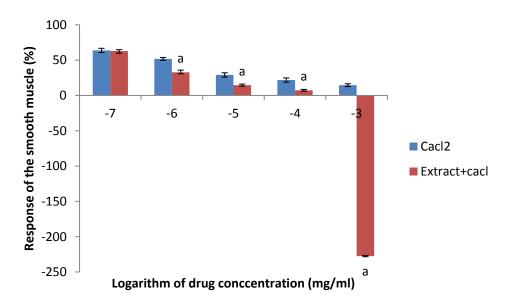


Figure 3. Effects of graded doses of $Cacl_2 + 10^{-4}$ mg/ml of *Xylopia aethiopica* on motility of rabbit ileum (a= p<0.05).

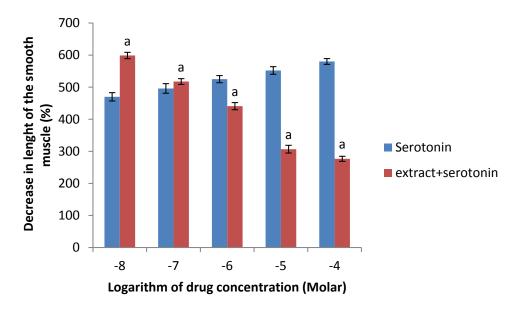


Figure 4. Effects of graded doses of Serotonin + 10⁻⁴ mg/ml of *Xylopia aethiopica* on motility of rabbit ileum (a= p<0.05).

inferred here that the extract potentiated the relaxant effect on the smooth muscle in the presence of CaCl₂. This may also indicate that both are acting independently to exert their effects.

Serotonin alone produced a dose dependent increase in relaxation of the rabbit ileum. Simultaneous administration of serotonin and extract produced a dose dependent decrease in relaxation of the rabbit ileum rather than the expected increase in relaxation by two

significant relaxants. This may suggest that the two substances may be using the same pathway and competing for receptor sites. Different types of serotonin receptors are expressed on different gastrointestinal cells, enteric nerves, smooth muscle and interstitial cells of cajal (Wouters et al., 2007). Expression and function of these receptors differ, some of them mediate relaxation, others only inhibition and activation of smooth muscle. This diversity makes it difficult to establish the clear

function of serotonin in the gastrointestinal system (Wouters et al., 2007). It is therefore possible that the extract of *X. aethiopica* employed this difference in serotonin receptor action to exert contractile effects in some smooth muscle such as uterus (Nwafor and Kalio, 2006), while exerting smooth muscle relaxant effects on rabbit ileum.

In conclusion, *X. aethiopica* has a smooth muscle relaxant effect on rabbit ileum and as such may reduce gastrointestinal motility.

Conflicts of interest

The authors have not declared any conflict of interest.

ACKNOWLEDGEMENT

The author acknowledged the technical staffs of Department of Physiology, University of Calabar.

REFERENCES

Barminas JT, James MK, Abubaka UM (1999). Chemical composition of seeds and oil of *Xylopia aethiopica* grown in Nigeria. Plant Foods Hum. Nutr. 53(3):193-198.

- Choumessi AT, Danel M, Chassaing S, Truchet I, Penlap VB, Pieme AC, Asonganyi T, Ducommun B, Valette A (2012). Characterization of the antiproliferative activity of *Xylopia aethiopica*. Cell Div. 7(1):8-15
- da Silva TMS, da Silva BA, Mukherjee R (1999). The monoterpene alkaloid canthyeyine from strychnos trinervis root and its spasmolytic properties. Phytomed. 6(3):169-176.
- Ebong OO, Warison BA, Orupabo I (1995). The gastrointestinal inhibitory actions of *Xylopia aethiopica* (Dunal) A. Rich (Annonaceae) in rats. West Afr. J. Pharmacol. Drug Res. 11(1):94-98.
- Ebong, Shode FO (2000). Inhibitory effects of aqueous and petroleum ether extracts of the fruit of *Xylopia aethiopica* (Dunal A. Rich (Annonaceae)) on rats uterine muscle. J. Med. Med. Sci. 2(1):45-51.
- Ekong DE, Ogan AU (1968). Chemistry of the constituent of *Xylopia aethiopica*. The structure of xylopic acid a diterpene acid. J. Chem. Soc. 54:311-312.
- Iwu MW, Duncan AR, Okunji CO (1993). New antimicrobial of plant origin. In Jankick J. (Eds). Perspective of new crops and uses AIIS press. Alexandria VA. pp. 457-462.
- Nwafor A, Kalio ID (2006). Physiological effects of *Xylopia aethiopica* on pregnancy in albino rats. J. Med. Pharmacol. Sci. 2(1):1-4.
- Okwari OO, Dasofunjo K, Obembe AO, Olatunji TL, Osim EE (2014). *Xylopia aethiopica* fruit extract lowers uric acid levels in wistar albino rats. Int. J. Pharm. Biol. Sci. 9(1):46-50.
- Wouters MM, Farrugia G, Schemann (2007): 5-HT receptors on interstitial cells of cajal smooth muscle and enteric nerves. J. Neurogastroenterol. Motil. 19(2):5-12.



Related Journals Published by Academic Journals

- International Journal of Plant Physiology and Biochemistry
- African Journal of Biotechnology
- Journal of Developmental Biology and Tissue Engineering

