# PHYTOCHEMICALS AND BIOACTIVITIES OF *PIPER MAINGAYI* HK. F., *P. MAGNIBACCUM* C. DC. AND *P. CANINUM* BLUME SPECIES

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Specially to Husband and Ummar, For your unwavering support and energetic love. Both of you have been my greatest strength and thank you for always understand

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My siblings

The whole family

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#### **ABSTRACT**

The chemical compositions of the essential oils and phytochemicals of *Piper* maingayi Hk. F., P. magnibaccum C. DC. and P. caninum Blume were studied. The essential oils obtained by hydrodistillation from the fresh samples of P. maingayi (stem and fruit) and P. magnibaccum (stem and leaf) were analyzed by capillary gas chromatography (GC) (Kovats Indices) and gas chromatography-mass spectrometry (GC-MS). The stem and fruit oils of P. maingayi successfully afforded 34 and 18 components, respectively. The stem oil consisted of  $\beta$ -caryophyllene (26.2%) and  $\alpha$ cedrene (8.4%) as the major components, while the fruit oil was dominated by βcaryophyllene (39.6%) and  $\delta$ -cadinene (22.6%). The essential oils of leaf and stem of P. magnibaccum gave 25 and 34 constituents, respectively. Both the leaf and stem oils were rich with germacrene D (10.7-40.8%) and  $\beta$ -caryophyllene (8.5-19.7%). The phytochemical study was carried out on the dried samples using maceration technique with n-hexane, dichloromethane and methanol to acquire the crude extracts. Fractionation and purification of the crude extracts using various chromatographic techniques have resulted in the isolation of eighteen compounds belonging to eight classes of phytochemicals. Those classes of phytochemicals were identified spectroscopically as aporphine alkaloids, triterpenes, fatty acids and esters, phenolic, flavonoid, amide alkaloid and lignin. β-Sitosterol, oleic acid and cepharadione A were isolated from all the investigated species. Piperumbellactam A was isolated from P. maingayi and P. magnibaccum, linoleic acid was isolated from P. magnibaccum and P. caninum while methyl linolenate was isolated from P. caninum and P. maingayi. Six compounds were isolated exclusively from *P. maingayi* and elucidated as sesamin, butyl dodecanoate, isovanillic acid, cepharadione B, piperolactam A and one new unsaturated amide namely N-isobutyl-15-(18,19-methylenedioxyphenyl)-2E,4E,12Zpentadecatrienamide. Two compounds characterised as 24S-ethylcholesta-5,22,25trien-3 $\beta$ -ol and stigmast-3,6-dione were obtained from P. magnibaccum while four compounds, namely 24-methylenecycloartan-3-one, 5,7-dimethoxyflavone, cepharanone A and aristolactam AII were revealed from P. caninum. Screenings on antibacterial, antioxidant, anti-inflammatory and antityrosinase bioactivities of the selected crude extracts, essential oils and pure compounds were also investigated. The leaf essential oil of P. magnibaccum showed a moderate antibacterial activity with MIC value of 250 µg/mL against *Pseudomonas aeruginosa* compared to the other oils, while N-isobutyl-15-(18,19-methylenedioxyphenyl)-2E,4E,12Z-pentadecatrienamide showed MIC value of 250 µg/mL each on B. subtillis and P. aeruginosa. The essential oil of P. maingayi and P. magnibaccum stems depicted a significant activity in DPPH assay with SC<sub>50</sub> value of 14.9 and 17.5 µg/mL, respectively. Study on antiinflammatory activity was carried out using 15-LOX enzymatic assay. Amide of Nisobutyl-15- (18,19- methylenedioxyphenyl)- 2E,4E,12Z- pentadecatrienamide exhibited the strongest inhibition against 15-LOX at IC<sub>50</sub> 42.52 µM. The tyrosinase inhibition activity showed moderate activity (59.6%) for P. maingayi stem oil and ethyl acetate crude extract (69.2%) each at a concentration of 1 mg/mL.

#### **ABSTRAK**

Komposisi kimia minyak pati dan fitokimia spesies *Piper maingayi* Hk. F., P. magnibaccum C. DC. dan P. caninum Blume telah dikaji. Minyak pati yang diperoleh daripada penyulingan hidro sampel segar P. maingayi (batang dan buah) dan P. magnibaccum (batang dan daun) telah dianalisis menggunakan kromatografi gas (GC) kapilari (Indeks Kovat) dan kromatografi gas-spektrometri jisim (GC-MS). Minyak daripada batang dan buah P. maingayi masing-masing telah berjaya memberikan 34 dan 18 komponen. Minyak daripada batang terdiri daripada β-kariofilena (26.2%) dan α-kedrena (8.4%) sebagai komponen utama, manakala minyak daripada buah didominasi oleh β-kariofilena (39.6%) dan δ-kadinena (22.6%). Minyak pati daripada daun dan batang P. magnibaccum masing-masing memberikan 25 dan 34 sebatian. Kedua-dua minyak daripada daun dan batang didapati kaya dengan germakrena D (10.7-40.8%) dan β-kariofilena (8.5-19.7%). Kajian fitokimia telah dijalankan ke atas sampel kering menggunakan kaedah rendaman dengan *n*-heksana, diklorometana dan methanol untuk mendapatkan ekstrak mentah. Pemeringkatan dan penulenan ekstrak mentah menggunakan pelbagai teknik kromatografi telah menghasilkan lapan belas sebatian yang tergolong dalam lapan kelas fitokimia. Kelas fitokimia ini telah dikenalpasti secara spektroskopi sebagai alkaloid aforfina, triterpena, asid lemak dan ester, fenolik, flavonoid, alkaloid amida dan lignin. β-Sitosterol, asid oleik dan A telah diasingkan daripada kesemua spesies Piperumbellaktam A telah diasingkan daripada P. maingayi dan P. magnibaccum, asid linoleik telah diasingkan daripada P. magnibaccum dan P. caninum manakala metil linolinat telah diasingkan daripada P. caninum dan P. maingayi. Enam sebatian telah diasingkan secara eksklusif daripada P. maingayi dan telah dikenalpasti sebagai sesamin, butil dodekanoat, asid isovanilik, sefaradion B, piperolaktam A dan satu sebatian amida tak-tepu iaitu N-isobutil-15-(18,19-metilenadioksifenil)-2E,4E,12Zpentadekatrienamida. Dua sebatian yang dicirikan sebagai 24S-etilkolesta-5,22,25trien- $3\beta$ -ol dan stigmast-3,6-dion telah diperoleh daripada *P. magnibaccum* manakala empat sebatian iaitu 24-metilenasikloartan-3-on, 5,7-dimetoksiflavon, sefaranon A and aristolaktam AII telah dikenalpasti daripada P. caninum. Penyaringan bioaktiviti antibakteria, antioksidan, antiradang dan antitirosinasa terhadap ekstrak mentah, minyak pati dan sebatian tulen terpilih telah juga dikaji. Minyak pati daun P. magnibaccum menunjukkan aktiviti antibakteria yang sederhana dengan nilai MIC 250 µg/mL terhadap *Pseudomonas aeruginosa* berbanding dengan minyak pati yang lain, manakala N- isobutyl - 15 - (18,19 - metilenadioksifenil) - 2E, 4E, 12Z pentadekatrienamida menunjukkan nilai MIC 250 µg/mL setiap satu bagi B. subtillis dan P. aeruginosa. Minyak pati daripada batang P. maingayi dan P. magnibaccum menunjukkan aktiviti signifikan dalam cerakin DPPH masing-masing dengan nilai SC<sub>50</sub> 14.9 dan 17.5 µg/mL. Kajian aktiviti antiradang telah dijalankan dengan menggunakan cerakin enzim 15-LOX. Sebatian amida N-isobutil-15-(18,19metilenadioksifenil)-2E,4E,12Z-pentadekatrienamida menunjukkan perencatan yang paling kuat terhadap 15-LOX pada IC<sub>50</sub> 42.52 µM. Aktiviti perencatan tirosinasa menunjukkan aktiviti yang sederhana bagi minyak pati batang *P. maingayi* (59.6%) dan ekstrak mentah etil asetat (69.2%) setiap satu pada kepekatan 1 mg/mL.

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## LIST OF ABBREVIATIONS

<sup>13</sup>C - Carbon-13

1D - 1 Dimension

<sup>1</sup>H - Proton

2D - 2 Dimension
AA - Ascorbic Acid

Abs - Absorbance

Ac - Acetone

BaCl<sub>2</sub> - Barium chloride

BHT - Butylated hydroxytoluene

br - broad

*c* - Concentration

CC - Column Chromatography

CDCl<sub>3</sub> - Deuterated chloroform

CHCl<sub>3</sub> - Chloroform

CH<sub>2</sub>Cl<sub>2</sub> - Dichloromethane

cm - Centimeter

cm<sup>-1</sup> - Per centimeter

COSY - Correlation Spectroscopy

d - doublet

dd - doublet of doublets

DCM - Dichloromethane

DEPT - Distortionless Enhancement by Polarization Transfer

DMSO - Dimethyl sulfoxide

DPPH - 2,2-Diphenyl-1-picrylhydrazyl

EIMS - Electron Impact Mass Spectrometry

EtOAc - Diethyl ether

EtOAc - Ethyl acetate

GA - Gallic acid

GC - Gas Chromatography

GC-MS - Gas Chromatography-Mass Spectrometry

h - Hour(s)

n-Hex - Hexane

HMBC - Heteronuclear Multiple Bond Correlation

HMQC - Heteronuclear Multiple Quantum Coherence

Hz - Hertz

IC<sub>50</sub> - Inhibition Concentration at 50%

IR - Infrared

J - Coupling ConstantKBr - Potassium Bromide

Lit. - Literature m - multiplet

m.p - melting point

m/z - mass to charge ion

M<sup>+</sup> - Molecular ion

mg - Milligram

MIC - Minimum Inhibition Concentration

min - Minute(s)
mL - milliliter
mm - millimeter

MS - Mass Spectrometry

NA - Nutrient agar
NB - Nutrient broth

nm - nanometer

NMR - Nuclear Magnetic Resonance

NO - Nitric Oxide

PE - Petroleum ether ppm - parts per million  $R_f$  - Retention factor

rpm - Revolution per minute

s - singlet

SD - Standard Deviation

t - triplet

TLC - Thin Layer Chromatography

TMS - Tetramethylsilane

 $t_R$  - Retention time

UV - Ultraviolet

VLC - Vacuum Liquid Chromatography

 $\alpha$  - Alpha  $\beta$  - Beta

 $\delta$  - Chemical shift

 $\mu M$  - Micro molar

 $\mu m$  - Micrometer

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### **CHAPTER 1**

#### INTRODUCTION

### 1.1 Introduction

Isolation and characterization of pharmacologically active compounds from medicinal plants continuously been demanded today. In recent years, interest in traditional medicine has increased greatly among researchers and the general public [1]. The considerable interest for replacing synthetic drugs with natural sources from parts of plants has led to intensified exploration and research for variety of purposes to cure illness. Thousands of plants have been used traditionally to treat various diseases, thus, natural remedies have become popular, especially in the part of lower risk of adverse reaction. [2].

Drug discovery from medicinal plants has developed to include numerous fields of study and various approaches of analysis. Commonly, the procedure begins with collection and identification of potential plant(s) species by a plants expertise. Collection may involve species of known biological activity with interesting active compounds which have been used traditionally as natural remedies or may involve taxonomic collected randomly for a new study [3, 4]. Attentively, phytochemists will investigated the plants by preparing extracts, forming biological screening of the extracts using pharmacologically pertinent assays, and begins the process of isolation and characterization of the active compound(s) through various chromatographic methods [3].

Quinine (1), atropine (2), morphine (3) and codeine (4) are a few of novel drug entities isolated from plants that had been listed in WHO List of Essential Medicines and developed synthetically by pharmaceutical industry [5-7]. Quinine (1) was first isolated from *Cinchona* bark and used to prevent and treat malaria [5, 8], atropine (2) from family of Solanaceae was used as an intravenous drug during anaesthesia [5, 9], meanwhile morphine (3) and codeine (4) were isolated from latex of opium poppy, *Papaver somniferum* in which the former was devoted as analgesic to control chronic cancer pain [5, 10].

Literally, the practice of plants as natural medicine dates back to a very primitive period of known civilizations. The increasing interest in herbs is based on the beliefs that plants have a vast potential as a healing medicine [11]. In South Asian countries, they are frequently preferred for prophylactic and therapeutic uses [12]. This scenario has also reflected in Malaysia in view of the fact that the market demand for traditional herbs as health supplements or for medicinal purposes has increased gradually over the past years [13].

## 1.2 Medicinal Plants in Malaysia

Our Malaysia's rainforest, encompasses more than 2000 plants species which have been reported to possess various medicinal values. The traditional herbal plants and their parts are primary sources of products for the nutraceutical and pharmaceutical industries. They are used in preparations for various products ranging from traditional remedies to extracts with standardized contents of active constituents to chemically pure compounds used in drugs. Furthermore, herbal plants are also utilized in food, beverage, flavor and fragrance industries. Therefore, traditional herbal plants species have a good prospect not only for the traditional medicinal industries but also for country's pharmaceutical industry as a whole [14].

Several traditional plants in Malaysia are well known to possess medicinal values and largely consumable as an 'ulam', which is chewed alone or with other plants or food materials. The plants leaves, fruits, seeds, tuber and roots are enriched with nutrients [15]. **Table 1.1** shows few selected traditional plants in Malaysia which are consumed as 'ulam' and used as ingredients for traditional medicine. Plants from the genus *Piper* such as *Piper sarmentosum*, *P. betle* and *P. nigrum* are also categorize among the important medicinal plants used in various system of medicine in Malaysia [15, 16].

**Table 1.1**: Selected Traditional Medicinal Plants in Malaysia as '*Ulam*' [15, 16]

Local name	Botanical name
Cekur Manis	Sauropus androgynus
Daun Selom	Oenanthea javanica
Hempedu Bumi	Andrographis paniculata
Jarum Tujuh Bilah	Pereskia sacharosa
Kaduk	Piper sarmentosum
Kemangi	Ocimum americanum
Sirih	Piper betle
Mas Cotek	Ficus deltoidea
Pegaga	Centella asiatica
Tenggek Burung	Euodia redlevi

Plants of Piperaceae such as *P. betle* L. and *P. nigrum* Linn. are the most sought after medicinal plants among Malaysian. They are widely growing in the tropical humid climates and leaves of *P. betle*, with a strong pungent and aromatic flavor are largely used as a mouth freshener [17]. In previous studies, the *P. betle* leaves, roots and whole extracts of this glabrous climbing vine showed a very strong antimicrobial [18], anti-inflammatory [19], reduction of cholesterol level [20] and good antioxidant activities [21]. Meanwhile, *P. nigrum* is the primary source of spices worldwide [22].

Comparing with *P. sarmentosem* Roxb. which locally known as *kaduk*, this species is also shows a remarkable antioxidant activity [23], besides as potential anticancer [24], anti-inflammatory [25], antidiabetic [26] and protective effect against atherosclerosis [27]. Due to these pharmacological and nutraceutical prospectives, both *P. betle* and *P. sarmentosum* have been studied for their toxicology and drug exposures as prescription and recently improved into product formulations [28]. However, only these species were extensively studied for their oils, phytochemicals and bioactivities although ironically, there are a numerous species of *Piper* grown abundantly need to be discovered.

## 1.3 Piperaceae Family

The Piperaceae family is assigned in the order of Piperales and widely distributed in the topics and subtropics regions. The family has about five genera and over 1950 species [29]. *Manekia*, *Verhuellia*, *Zippelia*, *Piper* and *Peperomia* are the genera in Piperaceae plant taxonomy [30]. *Piper* and *Peperomia* contributed the most number of species in this family with the latter used as ornamental plants [31]. Commonly many species of *Piper* were used as spices, folk medicines and pests control agents [32, 33].

*Piper* as the largest genus in the family of this pantropical group are estimated to contains 2000 species dispersed widely in American and Asian tropic including India, Indonesian and Malaysian tropical rainforest [34]. Most species of *Piper* 

appeared to be restricted to altitudes ranging from 0 to 2500 m, and very few occurred above 3000 m which grow in wet and shaded places [34, 35]. This genus is usually erect or scandent herbs, shrubs or infrequently trees [36, 37]. The structure is rather uniform morphologically, with simple alternate leaves and joined stems with enlarged nodes and possessed aromatic or pungent smell. Many produce pearl bodies on the leaves or stems, but the most distinctive morphological feature is the production of inflorescences of tiny seeds packed into upright or pendant spikes [38]. **Table 1.2** tabulated few examples of common *Piper* species found in Malaysia with traditional uses [39-41]. Due to the endless traditional uses of *Piper* species, the search for chemical compositions and active constituents from different *Piper* species has been intensified in recent years as a source of natural products with potential bioactivity properties [42].

**Table 1.2**: Several Local *Piper* Species and their Traditional Uses [39-41]

Piper Species	Local Name	Traditional Uses
P. argyrites	Sireh rimau puteh	Masticatory as a stimulant to sweeten the
		breath
P. baccatum	Gadong hutan	Relief cough and treating venereal diseases
P. betle	Sireh China/	Masticatory as a stimulant to sweeten the
	Sireh Melayu	breath
	•	Relief cough and asthma
		To stimulate secretion of milk
		Treat vaginal odor and sagging breast
		externally
P. caninum	Sireh hantu	Treating hoarseness
P. cubeba	Kemungkus	Tonic and relief rheumatism
P. febrifugum	Akar sangkap	Treating fever
P. nigrum	Lada hitam/	Food seasoning
Ü	Lada putih	Tonic and 'jamu' drink during confinement
P. porphyrophyllum	Sireh rimau	Relief weakness and pains in bones
P. chaba	Sireh kadok	Treating hemorrhoids
P. retrofractum	Lada panjang	Food seasoning
v	1 0 0	Tonics for digestive/ intestinal disorder
		To relief muscular stiffness and
		inflammation
P. umbellatum	Segumbar urat	Poulticing and applied to wound

## 1.4 Problem Statements and Significant of Research

Piperaceae family has provided many past and present civilizations with a source of medicines and food spices. The well-known species as stated previously; *P. betle* Lin, *P. nigrum* Linn and *P. sarmentosum* Roxb had been brought up to the highest level of usage in perfumery and herbal products. However, there are still a wide numbers of *Piper* species from Malaysia that have not yet being explored scientifically. Among the *Piper* species that have not been investigated extensively are *P. maingayi* Hk. F., *P. magnibaccum* C. DC. and *P. caninum* Blume.

A study on the essential oil of *P. maingayi* leaf has been reported by Sirat *et al.*, [43] while another study on the chemical constituents of the oil from *P.caninum* was published in 2011 [44]. However, no study on the *P. magnibaccum* essential oil has been reported elsewhere. With regards on the phytochemicals investigation, only one study of phytochemicals from *P. magnibaccum* cultivated in Indonesia has been reported by Emrizal *et al.*, [45] and a short communication on the phytochemicals of *P. maingayi* has been published by Ahmad *et al.*, [46]. Thus far, in the aspect of biological activity, only one report on the anti-inflammatory activity of the phytochemicals of *P. magnibacum* [45] from Indonesia has been published but none on *P. maingayi*. Although *P. caninum* has been studied for its phytochemistry and biological activities, the species investigated was originally collected from Borneo [44, 47], not from the Peninsular of Malaysia.

Based on the above reports, there is an urgent need to explore the essential oil compositions of the other parts (stems and fruits) of *P. maingayi* as well as the essential oils of *P. magnibaccum*. Extensive studies on the phytochemicals of *P. maingayi*, *P. magnibaccum* and *P. caninum* originated from Peninsular Malaysia rainforest need to be carried out using modern technique in isolation of novel compounds from these species. Thus, comparison of the phytochemical profiles of the current findings with previous reports can also be compared. The biological activities of the oils, crudes and pure phytochemicals of *P. maingayi*, *P. magnibaccum* and *P. caninum* are similarly important to be investigated for the development of pharmaceutical and herbal formulation documentations.

## 1.5 Objectives of Research

The objectives of this study were divided into three parts. The first was to isolate and determine the chemical compositions of the essential oils of selected parts of *Piper* species (*P. maingayi* and *P. magnibaccum*) using GC and GC-MS techniques. The second part was to isolate, purify and elucidate phytochemicals from *P. maingayi*, *P. magnibaccum* and *P. caninum*. The third part was to screen the biological activities of the essential oils, extracts and pure isolated phytochemicals.

## 1.6 Scope of Study

The scope of this study was focused on the extraction of the essential oils from fresh stems and fruits of *P. maingayi*; stems and leaves of *P. magnibaccum* and *P. caninum* using hydrodistillation technique. The oil compositions were identified using GC and GC-MS instruments as well as Kovats indices.

The dried samples of each of the *Piper* species were extracted using cold extraction techniques with different polarity of solvents. The crude extracts were fractionated and purified by chromatographic techniques which include vacuum liquid chromatography (VLC), open gravity column chromatography (CC), versa flash chromatography and recycle-preparative high performance liquid chromatography (recycle-HPLC) to obtain the pure phytochemicals which were analysed spectroscopically by using IR, 1D NMR (<sup>1</sup>H and <sup>13</sup>C), 2D NMR (COSY, HMQC, HMBC, NOESY), MS and UV. Finally, the characterised phytochemicals, essential oils and crude extracts were subjected to biological activities which include antibacterial, antioxidant, antityrosinase and antiinflammatory activities. Antibacterial activity utilised minimum inhibition concentration (MIC) method. Antioxidant activity was determined by using total phenolic content, and DPPH free radical scavenging. As for the antiinflammatory and antityrosinase activities, the essential oils and phytochemicals were tested against 15-lipoxygenase (15-LOX) and mushroom tyrosinase enzymes *in vitro*.

#### REFERENCES

- 1. Fakim, A.G. (2006). Medicinal plants: Traditions of yesterday and drugs of tomorrow. *Molecular Aspects of Medicine*. 27: 1-93.
- 2. Natarajan, K.S., Narasimhan, M., Shanmugasunduraram, K.R. and Shanmugusundaram, E.R.B. (2006). Antioxidant activity of a salt-spice-herbal mixture against free radical induction. *Journal of Ethnopharmacology*. 105: 76-83.
- 3. Balunas, M.J. and Kinghorn A.D. (2005). Drug discovery from medicinal plants. *Life Sciences*. 78: 431-441.
- 4. Baker, D.D., Chu, M., Oza, U. and Rajgarhia, V. (2007). The value of natural products to future pharmaceutical discovery. *Natural Products Repository*. 24: 1225-1244.
- 5. Philipson, J.D. (2001). Phytochemistry and medicinal plants. *Phytochemistry*. 56: 237-243.
- 6. Rates, S.M.K. (2001). Plants as source of drugs. *Toxicon*. 39: 603-613.
- 7. WHO Model List of Essential Medicines. 19<sup>th</sup> Edition. Unpublished note, World Health Organization (WHO).
- 8. Taylor, R.B., Shakoor, O., Behrens, R.H., Everard, M., Low, A.S., Wangboonskul, J., Reid, R.G and Kolawole, J.A. (2001). Pharmacopoeial quality of drugs supplied by Nigerians pharmacies. *The Lancet*. 357: 1933-1936.
- 9. Moawad, H.E.S., Hefnawy, A.S.E. (2015). Spinal vs. general anesthesia for percutaneous nephrolithotomy: A prospective randomized trial. *Egyptian Journal of Anaesthesia*. 31: 71-75.
- 10. Walsh, T.D. (1984). Oral morphine in chronic cancer pain. *Pain*. 18: 1-11.
- 11. Kumar, S., Kamboj, J., Suman, and Sharma, S. (2011). Overview of various aspects of the health benefits of *Piper Longum* Linn. Fruit. *Journal of Acupuncture Meridian Study*. 4: 134-140.

- 12. Majumdar, B. and Islam, M.N. (2013). Effect of ethanolic extract of *Piper betle* Linn as an immunodulatory agent: A unique role of phytochemicals. *Journal of Biomedical Pharmacology Research*. 2: 42-47.
- 13. Bakhru, H.K. (2008). Herbs that Heal: Natural remedies for good health. Orient Paperbacks, India. p: 17-18.
- 14. Gerber, R. and Williams, M. (2002). Geography, Culture and Education. Kluwer Academic Publisher, The Netherlands. p: 62.
- 15. Zaifuddin, F.A.M., Hassan, N.M. and Othman, R. (2014). Quantification of pro-vitamin A activities and content in 22 selected '*Ulam*' species or Malaysian traditional vegetables. *International Journal of Pharmacology and Pharmaceuticals*. 6: 9-12.
- 16. Jamal, J.A. (2006). Malay Traditional Medicine: an overview of scientific and technological progress. *Technology Monitor*. Nov-Dec: 37-49.
- 17. Prabu, S.M., Muthmani, M. and Shagirtha, K. (2012). Protective effect of *Piper betle* leaf extract against cadmium-induced oxidative stress and hepatic dysfunction in rats. *Saudi Journal of Biological Sciences*. 19: 229-239.
- Sugumaran, M., Gandhi, S.M., Sankarnarayanan, K., Yokesh, M., Poornima, M. and Sree R.R. (2011). Chemical composition and antimicrobial activity of vellaikodi variety of *Piper betle* Linn leaf oil against dental pathogens. *International Journal of PharmTech Research*. 3: 2135-2139.
- 19. Chwan-Fwu, L., Tsong-Long, H., Chun-Chien, C., Huei-Yu, T. and Horng-Liang, L. (2013). A new hydroxychavicol dimer from the roots of *Piper betle*. *Molecules*. 18: 2563-2570.
- 20. Thirumalai, T., Tamilselvan, N. and David, E. (2014). Hypolipidemic activity of *Piper betle* in high fat diet induced hyperlipidemic rat. *Journal of Acute Diseases*. 12: 131-135.
- 21. Vadlapudi, V. and Kaladhar, D.S.V.G.K. (2012). Phytochemical evaluation and molecular characterization of some important medicinal plants. *Asian Pacific Journal of Tropical Diseases*. 11: S26-S32.
- 22. Ahmad, N., Fazal, H., Abbasi, B.H., Farooq, S., Ali, M. and Ali, M.K. (2012). Biological role of *Piper nigrum* L. (Black pepper): A review. *Asian Pacific Journal of Tropical Biomed*ical. S1: 1945-1953.

- 23. Hussain, K., Ismail, Z. and Ibrahim, P. (2009). Standardization and *in-vivo* antioxidant activity of ethanol extracts and fruit and leaf of *Piper sarmentosum*. *Planta Medica*. 76: 418-425.
- 24. Hussain, K., Ismail, Z., Sadikun, A., Ibrahim, P. (2009). Cytotoxicity evaluation and characterization of chloroform extract of leaf of *Piper sarmentosum* possessing antiangiogenic activity. *Pharmacology online*. 2: 379-391.
- 25. Seyyedan, A., Yahya F., Kamarolzaman, M.F.F., Suhaili, Z., Desa, M.N.M., Khairi, H.M., Somchit, M.N., Fatimah, C.A., The, L.K., Salleh, M.Z. and Zakaria, Z.A. (2013). Review on the ethnomedicinal, photochemical and pharmacological properties of *Piper sarmentosum*: scientific justification of its traditional use. *Humanitas Traditional. Medical.* 3: 19-23.
- 26. Damsud, T., Adisakwattana, S. and Phuwapraisirisan, P. (2013). Three new phenylpropanoyl amides from the leaves of *Piper sarmentosum* and their α-glucosidase inhibitory activites. *Phytochemistry Letters*. 6: 350-354.
- Amran, A.A., Zakaria, Z., Othman F., Das, S., Raj, S. and Nor-Anita, M.M.N. (2010). Aqueous extract of *Piper sarmentosum* decreases atherosclerotic lesions in high cholesteromic experimental rabbits. *Lipid Health Diseases*. 9: 44-50.
- 28. Hussain, K., Ismail, Ismail, Z., Sadikun, A. and Ibrahim, P. (2011). Bioactive markers based pharmacokinetic evaluation of extracts of a traditional medicinal plant, *Piper sarmentosum*. *Evidence-Based Complementary and Alternative Medicines*. 1-7.
- 29. Mabberley, D.J. (1997). *The Plant-book. A Portable Dictionary of the Higher Plants* (13<sup>th</sup> Edition). New York, USA: Cambridge University Press.
- 30. Samain, M.S., Mathieu, G. 1., Wanke, S., Neinhuis, C., Goetghebeur, P. (2008). *Verhuellia* revisited-unravelling its intricate taxonomic history and a new subfamilial classification of *Piperaceae*. *Taxonomy*. 57: 583-587.
- 31. Parra, J.E.P., Oscar J.P., Juliet A.D., Wilman A.C., Luis E. (2013). A new benzoic acid derivative isolated from *Piper cumanense* Kunth (Piperaceae). *Phytochemistry Letters*. 6: 590-592.
- 32. Arnason, J.T., Steven, R.S. and Ian, M.S. (2005). Natural products from plants as insecticides. *Phytochemistry and Pharmacognosy*. 5: 37-46.

- 33. Sengupta, S., Ray, A.B. (1987). Chemistry of *Piper* species: a review. *Fitoterapia*. 3: 147–166
- 34. Jaramillo, M.A. and Paul S.M. (2001). Phylogeny and patterns of floral diversity in the genus *Piper* (Piperaceae). *American Journal of Botany*. 88: 706-716.
- 35. Quijano-Abril, M.A., Callejas-Posada, R., Rafael, D. and Miranda, E. (2006). Areas of endemism and distribution patterns for Neotropical *Piper* species (Piperaceae). *Journal of Biogeography*. 33: 1266-1278.
- 36. Parmar, V.S., Jain, S.C., Bishit, K.S., Jain, R., Taneja, P., Jha, A., Tyagi, A.D., Prasad, A.K., Wengei, J., Olsen, C.E. and Boll, P.M. (1997). Phytochemistry of the genus *Piper*. *Phytochemistry*. 46: 597-673.
- 37. Meghwal, M. and Goswami, T.K. (2013). *Piper nigrum*: An update. *Phytotherapy Research*. 27: 1121-1130.
- 38. Gupta, M., Gupta, A. and Gupta, S. (2013). In vitro antimicrobial and phytochemical analysis of dichloromethane extracts of *Piper nigrum* (Black Pepper). *Oriental Journal Chemistry*. 29: 777-782.
- 39. Burkill, I.H., Birtwistle, W. and Foxworthy, F.W. (1993). Dictionary of the Economic Products of the Malay Peninsula. (Vol. II). Publisher Governments of Malaysia by the Ministry of Agriculture Malaysia. p: 1766-1784.
- 40. Ong, H.C., Zuki, R.M. and Milow. P. (2011). Traditional knowledge of medicinal plants among the Malay villagers in Kampung Mak Kemas, Terenggani, Malaysia. *Ethnopharmacology Medical*. 5: 175-185
- 41. Kumar, S., Kamboj, J. and Suman, S.S. (2011). Overview for Various Aspects of the Health Benefits of *Piper longum* Linn. Fruit. *Journal of Acupuncture and Meridian Studies*. 4: 134–140.
- 42. Péres, V.F.1., Moura, D.J., Sperotto, A.R., Damasceno, F.C., Caramão, E.B., Zini, C.A. and Saffi, J. (2009). Chemical composition and cytotoxic, mutagenic and genotoxic activities of the essential oil from *Piper gaudichaudianum* Kunth leaves. *Food Chemical and Toxicology*. 47: 2389-95.
- 43. Sirat, H.M., Thai, O.B. and Ahmad, F. (2010). Chemical composition of the essential oil of *Piper maingayi* Hk. F. *Journal of Essential Oil Research*. 33: 323-324.
- 44. Salleh, W.M.N.H.W, Ahmad, F., Yen, K.H. and Sirat, H.M. (2011). Chemical Compositions, Antioxidant and Antimicrobial Activities of Essential Oils of

- *Piper caninum* Blume. *International Journal of Molecule Sciences*. 12: 7720-7731
- 45. Emrizal, Ahmad, F., Sirat, H., Jamaluddin, F., Mustapha, N.M., Ali, R.M. and Arbain, D. (2008). Anti-inflammatory of *Piper magnibaccum* (Piperaceae). *Natural products Communications*. 3: 1719-1721.
- 46. Ahmad, F., Jamil, S., Ibrahim, A.Z. and Read, R.W. (1996). Alkenylamides from *Piper maingayi* Hk. F. *Pertanika Journal of Science & Technology*. 4: 167-171.
- 47. Salleh, W.M.N.H.W., Ahmad, F. and Yen, K.H. (2015). Chemical constituents from *Piper caninum* and antibacterial activity. *Journal of Applied Pharmaceutical Science*. 5: 20-25.
- 48. Oliveira, G.L., Cardoso, S.K., Júnior, C.R.L., Veira, T.M., Guimarăes, E.F., Figueiredo, L.S., Martins, E.R., Moreira, D.L. and Kaplan, M.A.C. (2013). Chemical Study and Larvicidal Activity against *Aedes aegypti* of Essential Oil of *Piper aduncum* L. (Piperaceae). *Annals Brazilian Academy of Sciences*. 21: 167-169.
- 49. Jeena, K., Liju, V.B., Umadevi, N.P. and Kuttan, R. (2014). Antioxidant, antiinflammatory and antinociceptive properties of black pepper essential oil (*Piper nigrum* Linn). *Journal of Essential Oil Bearing Plants*. 17: 1-12.
- Michel, T., Pierre, J.D., Modeste, L.S., Fombotioh, N., Arlette, V.W.N., Paul, H.A.Z., Menut, C. (2009). Comparative essential oils composition and insecticidal effect of different tissues of *Piper capense* L., *Piper guineense* Schum. et Thonn., *Piper nigrum* L. and *Piper umbellatum* L. grown in Cameroon Francois. *African Journal of Biotechnology*. 81: 424-431.
- 51. Menon, A.N., Padmakumari, K.P. and Jayalekshmy, A. (2003). Essential oil composition of four major cultivars of black pepper (*Piper nigrum L.*) III. *Journal of Essential Oil Research*. 15: 155-157.
- 52. Salleh, W.M.N.H.W, Ahmad, F., Yen, K.H. and Sirat, H.M. (2012). Chemical compositions, antioxidant and antimicrobial activity of essential oils of *Piper officinarum* (Piperaceae). *Natural Product Communications*. 7: 1659-1662.
- 53. Lesueur, D., Bighelli, A. and Casanova, J. (2009). Composition of the essential oil of *Piper bavinum* C.DC. from Vietnam. *Journal of Essential Oil Research*. 21: 113-115.

- 54. Andrade, E.H.A., Alves, C.N., Guimaraes, E.F., Carreira, L.M.M. and Maia, J.G.S. (2011). Variability in essential oil composition of *Piper dilatatum* L.C. Rich. *Biochemical Systematics and Ecology*. 39: 669-675.
- 55. Salleh, W.M.N.H.W, Ahmad, F., Khong, H.Y. (2014). Chemical composition of *Piper stylosum* Miq. and *Piper ribesioides* Wall. Essential oils, and their antioxidant, antimicrobial and tyrosinase inhibition activities. *Boletin Latinoamericano y del Caribe de Plantas Medicinales y Aromaticas*. 13: 488-497.
- Maia, J.G.S., Zohhbi, M.G., Andrade, E.H.A., Santos, A.S., Silva, M.H., Luz, A.I.R. and Bastos, C.N. (1998). Constituents of the essential oil of *Piper aduncum* L. growing wild in the Amazon region. *Flavour and Fragrance Journal*. 13: 269-272.
- 57. Guerrini. A., Sacchetti, G., Rossi, D., Paganetto, G., Muzzoli, M. Andreotti, E., Tagnolini, M., Maldonado, M.E. and Bruni, R. (2009). Bioactivities of *Piper aduncum* L. and *Piper obliquum* Ruiz & Pavon (Pipeaceae) essential oils from eastern Ecuador. *Environmental Toxicology and Pharmacology*. 27: 39-48.
- 58. Machicado, A.R., Zimermann, D. and Rebelo, R.A. (2002). *Piper mikaniaum* (Kunth) Steudel from Santa Catarina, Brazil- a new source of safrole. *Journal of Essential Oil Research*. 14: 361-363.
- Monzote, L., Garcia, M., Montalvo, A.M., Scull, R. and Miranda, M. (2010).
   Chemistry, cytotoxicity and antileshmanial activity of the essential oil from Piper auritum. Memórias do Instituto Oswaldo Cruz. 105: 168-173.
- 60. Morais, S.M., Facundo, V.A., Bertini, L.M., Cavalcanti, E.S.B., Junior, J.F.A., Ferreira, S.A., Brito, E.S. and Neto, M.A.S. (2007). Chemical composition and larvacidal activity of essential oils from *Piper* species. *Biochemical Systematic and Ecology*. 35: 670-675.
- 61. Genderen, M.H.P., Leclercq, P.A., Delgado, H.S., Kanjilal, P.B. and Singh, R.S. (1999). Compositional analysis of the leaf oils of *Piper callosum* Ruiz & Pav. From Peru and *Michelia Montana* Blume from Peru. *Spectroscopy*. 14: 51-59.
- 62. Moreira, D.L., Souza, P.O., Kalan, M.A.C., Pereira, N.A., Cardoso, G.L. and Guimaraes, E.F. (2000). Effect of leaf essential oil from *Piper solmsianum* C.DC. in mice behavior. *Analytical of Academic Brassiliere*. 73: 1.

- 63. Vargas, L., Perez-Alonso, M.J., Velasco-Negueruela, A., Pala-Paul, J., Vallejo, M.C.G. (2003). Leaf essential oil of *Piper barbatum* H.B.K. (Piperaceae) from Peru. *Journal of Essential Oil Research*. 15: 163-164.
- 64. Takahashi, S. (1969). The presence of the tumor inhibitor crotepoxide (futoxide) in *Piper futokadzura*. *Phytochemistry*. 8: 321-322.
- 65. Takahashi, S., Kurabayashi, M., Ogiso, A. and Mishima, H. (1969). The structure of futoamide: a constituent of *Piper futokadzura* SIEB et. ZUCC. *Phytochemistry*. 17: 1225-1228.
- 66. Chatterjee, A. and Dutta, C.P. (1967). Alkaloids of *Piper longum* Linn-I: Structure and synthesis of piperlongumine and piperlonguminine. *Tetrahedron*. 23: 1769-1781.
- 67. Tarannum, N., Mosaddika A., Rahmana, M.M., Muhammad I., Haque M.E. and Cho, S.K. (2012). Antimicrobial, antileishmanial and cytotoxic compounds from *Piper chaba*. *Natural Product Research*. 26: 979–986.
- 68. Muharini, R., Liu, Z., Lin, W., Prokssh, P. (2015). New amides from the fruits of *Piper retrofractum*. *Tetrahedron Letters*. 56: 2521-2525.
- 69. Jiang, Z., Liu, W., Huang, C. and Huang, X. (2013). New amide alkaloids from *Piper longum. Fitoterapia*. 84: 222-226.
- 70. Marques, J.V., Kitamura, R.O.S., Lago, J.H.G., Young, M.C.M., Guinmareas, E.F. and Kato, M.J. (2007). Antifungal amides from *Piper scutifolium* and *Piper hoffmanseggianum*. *Journal of Natural Products*. 70: 2036-2039.
- 71. Rukachaisirikul, T., Siriwattanakit, P., Sukcharoenphol, K., Wongvein, C., Ruttanaweang, P., Wongwattanavuch, P. and Suksamram, A. (2004). Chemical constituents and bioactivity of *Piper sarmentosum*. *Journal of Ethnopharmacology*. 93: 173-176.
- 72. Sunila, E.S and Kttan, G. (2004). Immunomodulatory and antitumor activity of *Piper longum* Linn. and piperine. *Journal of Ethopharmacology*. 90: 339-346.
- 73. Dewick, P.M. (1997). Medicinal Natural Products: A Biosynthetic Approach. John Wiley & Sons. Third edition. p:286-288.
- 74. Duh, C.Y., Chang, Y. and Wang, S.K. (1990). Cytotoxic pyridine alkaloids from *Piper aborescens*. *Pytochemistry*. 29: 2689-2691.

- 75. Facundo, V.A., Silveria, A.S.P. and Morais, S.M. (2005). Constituents of *Piper alatabaccum* Trel & Yuncker (Piperaceae). *Biochemical Systematics and Ecology*. 33: 753–756.
- 76. Lee, S.E., Park, B.S., Kim, M.K., Choi, W.S., Kim, H.T., Cho, K.Y., Lee, S.G., Lee, H.S. (2001). Fungicidal activity of pipernonaline, a piperidine alkaloid derived from long pepper, *Piper longum* L., against phytopathogenic fungi. *Crop Protection*. 20: 523-528.
- 77. Kaou, A.M., Mahiou-Leddet, V., Canlet, C., Debrauwer, L., Hutter, S., Azas, N. and Ollivier, E. (2010). New amide alkaloid from the aerial part of *Piper capense* L.f. (Piperaceae). *Fitoterapia*. 81: 632-635.
- 78. Lei, J., Burgess, E.J., Richardson, A.T.B., Hawkins, B.C., Baird, S.K., Smallfield, B.M., van Klink, J.W. and Perry, N.B. (2015). Cytotoxic amide from fruits of Kawakawa, *Micropiper excelsum*. Planta Medica. 81: 1161-1163.
- 79. Matsuda, H., Ninomiya, K., Morikawa, T., Yasuda, D., Yamaguchia, I. and Yoshikawa, M. (2008). Protective effects of amide constituents from the fruit of *Piper chaba* on D-galactosamine/TNF-a-induced cell death in mouse hepatocytes. *Biorganic & Medicinal Chemistry Letters*. 18: 2038-2042.
- 80. Morikawa, T., Matsuda, H., Yamaguchia, I., Pongpiriyadacha, Y., and Yoshikawa, M. (2004). New amide and gastroprotective constituents from the fruit of *Piper chaba*. *Natural Products Chemistry*. 70: 152-159.
- 81. Wang, L., Vieth, R., Landes, R.C., Suzuki, Y., and Philip D.W. (1993). Antiepileptic effect of antiepilepsirine in pentylenetetrazol and amygdala kindled rats. *Epilepsy Research*. 15: 1-5.
- 82. Li, S., Wang, C., Li, W., Koike, K., Nikaido, T. and Wang, M.W. (2007). Antidepressant-like effects of piperine and its derivative, antiepilepsirine. *Journal of Asian Natural Products Research*. 9: 421-430.
- 83. Rao, V.R.S., Suresh, G., Babu, K.S., Raju, S.S., Vardhan, M.V.P.S.V., Ramakrishna, S. and Rao, J.M. (2011). Novel dimeric amide alkaloids from *Piper chaba* Hunter: isolation, cytotoxic activity and their biomimetic activites. *Tetrahedron*. 67: 1885-1892.
- 84. Navickiene, H.M.D., Alecio, A.C., Kato, M.J., Bolzani, V.S., Young, M.C.M., Cavalheiro, A.J. and Furlan, M. (2000). Antifungal amides from *Piper hispidum* and *Piper tuberculatum*. *Phytochemistry*. 55: 621-626.

- 85. Speck, K. and Magauer, T. (2013). The chemistry of isoindole natural products. *Beilstein Journal of Organic Chemistry*. 9: 2048–2078.
- 86. Ee, G. C. L., Lim, C. M., Lim, C. K., Rahmani, M., Shaari, K. and Bong, C. F. J. (2009). Alkaloids from *Piper sarmentosum* and *Piper nigrum*. *Natural Product Research*. 23: 1416-1423
- 87. Zhang, K., Ni, W. and Chen, C. (1999). Chemical constituents of *Piper boehmeriaefolium var*. Tonkinense. *Tianran Chanwu Yanjiu Yu Kaifa*. 11: 44-47.
- 88. Desai, S.J, Prabhu, B.R. and Mulchandani, N.B. (1988). Aristolactams and 4,5-dioxoporphines from *Piper longum. Phytochemistry*. 27: 1511-1515.
- 89. Olsen, C.E., Tyagi, O.D., Boll, P.M., Hussaini, F.A., Parmar, V.S., Sharma, N.K., Taneja, P. and Jain, S.C. (1992). An aristolactams from *Piper acutisleginum* and revision of the structures of piperolactam B and D. *Phytochemistry*. 33: 518-520.
- 90. Tabopda, T.K., Ngoupayo, J., Liu, J., Mitiane-Offer, A.C., Tanoli, S.A.K., Khan, S.N., Ali, M.S., Ngadjui, B.T., Tsamo, E., Dubois, M.A.L. and Luu, B. (2008). Bioactive aristolactams from *Piper umbellatam*. *Phytochemistry*. 69: 1726-1731.
- 91. Cunha, W.R., Silva, M.L.A., Veneziani, R.C.S., Ambrósio, S.R. and Bastos, J.K. Lignans: Chemical and Biological Properties. Phytochemical- A Global Perspective of Their Role in Nutrition and Health, Dr. Venketeshwer Rao. Croatia. InTech. ISBN: 978-953-51-0296-0.
- 92. Bodiwala, H.S., Singh, G., Singh, R., Dey, C.S., Sharma, S.S., Bhutani, K.K., and Singh, I.P. (2007). Antileishmanial amides and lignans from *Piper cubeba* and *Piper retrofractum*. *Journal of Natural Medicines*. 61: 418-421.
- 93. Dhanya S. Rajalekshmi, D.S., Kabeer, F.A., Madhusoodhanan, A.R., Bahulayan, A.K., Prathapan, R., Prakasan, N., Varughese, S. and Nair, M.S. (2016). Anticancer activity studies of cubebin isolated from *Piper cubeba* and its synthetic derivatives. *Bioorganic & Medicinal Chemistry Letters*. 26: 1767–1771.
- 94. Niwa, A.M., Paula, N.A., Campos, D., Sartori, V.D., Maistro, E.L., Ribeiro, L.R. and Mantovani, M.S. (2016). Evaluation of lignan (-)-cubebin extracted from *Piper cubeba* on human colon adenocarcinoma cells (HT29). *Journal of Toxicology and Environmental Health*. 79: 92–100.

- 95. Danelutte, A.P., Costantin, M.B., Delgado, G.M., Braz-Filho, R. and Kato, M.J. (2015). Divergence of Secondary Metabolism in Cell Suspension Cultures and Differentiated Plants of *Piper cernuum* and *P. crassinervium. Journal of Brazilian Chemical Society*. 16: 1425-1430.
- 96. Kothapalli Haribabu, K., Ajitha, M. and Mallavadhani, U.M. (2014). Simultaneous Determination of Asaranin and Sesamin in *Piper chaba* Fruit by using HPTLC-MS Method: Effect of Different Extraction Methods on the Yield of Marker Compounds. *Journal of Analytical and Bioanalytical Techniques*. 5: 1127-1132.
- 97. Tuntiwachwuttikul, P., Phansa, P., Pootaeng-on, P. and Taylor, W.C. (2006). Chemical Constituents of the Roots of *Piper Sarmentosum*. *Chemical and Pharmaceutical Bulletin*. 54: 149-151.
- 98. Baldoqui, D.C., Bolzani, V.S. and Furlan, M. (2009). Flavones, lignans and terpene from *Piper umbellate* (Piperaceae). *Química Nova*. 32: 1107-1109.
- 99. Srivastava, S., Gupta, M.M., Prajapti, V., Tripathi, A.K. and Kumar, S. (2001). Sesamin a Potent Antifeedant Principle from *Piper mullesua*. *Phytotherapy Research*. 15: 70-72.
- 100. Chen, S., Huang, H., Cheng, M., Wu, C., Ishikawa, T., Peng, C., Chang, H., Wong, S. and Chen, I. (2013). Neolignans and phenylproanoids from the roots of *Piper Taiwanese* and their antiplatelet and antitubercular activities. *Phytochemistry*. 93: 203-209.
- 101. Chauret, D.C., Bernard, C.B., Arnason, J.T. and Durst, T. (1996). Insecticidal Neolignans from *Piper decurrens*. *Journal of Natural Products*. 59: 152-155.
- 102. Koroishi, A.M., Foss, S.R., Cortez, D.A.G., Nakamura, T.U., Nakamura, C.V. and Filho, B.P.D. (2008). In vitro antifungal activity of extracts of neolignans from *Piper regnellii* against dermatophytes. *Journal of Ethnopharmacology*. 117: 270-277.
- 103. Scodro, R.B.L., Espelho, S.C., Pires, C.T.A., Garcia, V.A.S., Filho, L.C., Cortez, L.E.R., Pilau, E.J., Ferracioli, K.R.C., Siquiera, V.C.D., Cardoso, R.F. and Cortez, D.A.G. (2015). A new benzoic acid derivative from *Piper diospyrifolium* and its *anti-Mycobacterium tuberculosis* activity. *Phytochemistry Letters*. 11: 18-23.

- 104. Malami, I., Gibbons, S. and Malkinson, J.P. (2014). Synthesis and antibacterial evaluation of 3-farnesyl-2-hydroxybenzoic acid from *Piper multiplinervium*. *Fitoterapia*. 93: 189-193.
- 105. Zarai, Z., Boujelbene, E., Salem, N.B., Gargouri, Y. and Sayari, A. (2013). Antioxidant and antimicrobial activities of various solvent extracts, piperine and piperic acid from *Piper nigrum. LWT-Food Science and Technology*. 50: 634-641.
- 106. Parra, J.E., Delgado, W.A. and Cuca, L.E. (2011). Cumanensic acid, a new chromene isolated from *Piper cf. cumanense* Kunth. (Piperaceae). *Phytochemistry Letters*. 4: 280-282.
- 107. Freitas, G.C., Kitamura, R.O.S., Lago, J.H.G., Young, M.C.M., Guimaraes, E.F. and Kato, M.J. (2009). Caldensinic acid, a prenylated benzoic acid from *Piper caldense*. *Phytochemistry Letters*. 2: 119-122.
- 108. Ruiz, C., Haddad, M., Alban, J., Bourdy, G., Reategui, R., Castillo, D., Sauvain, M., Deharo, E., Esteves, Y., Arevalo, J. and Rojas, R. (2010). Activity-guided isolation of antileishmanial compounds from *Piper hispidum*. *Phytochemical Letters*. 4: 363-366.
- 109. Costa, G.M., Endo, E.H., Cortez, D.A.G., Nakamura, T.U., Nakamura, C.V. and Dias, B.P. (2016). Antimicrobial effects of *Piper hispidum* extract, fractions and chalcones against *Candida albicans* and *Staphylococcus aureus*.

  Journal de Mycologie Médicale. 26: 217-226.
- 110. Lo'pez, A., Ming, D.S., and Towers, G.H.N. (2002). Antifungal Activity of Benzoic Acid Derivatives from *Piper lanceaefolium*. *Journal of Natural Products*. 65: 62-64.
- 111. Danulette, A.P., Lago, J.H.G., Young, M.C.M. and Kato, M.J. (2003). Antifungal flavanones and prenylated hydroquinones from *Piper crassinervium* Kunth. *Phytochemistry*. 64: 555-559.
- 112. Freitas, G.C., Batisya, J.M., Franchi, G.C., Nowii, A.E., Yamaguchi, L.F., Vichacagua, J.D., Favaro, D.C., Furlan, M., Guimaraes, E.F., Jeffrey, C.S and Kato, M.J. (2014). Cytotoxic non-aromatic B-ring flavanones from *Piper carniconnectivum* C.DC. *Phytochemistry*. 97: 81-87.
- 113. Facundo, V.A., Balico, L.J., Lima, D.K.S., Santos, A.R.S., Morais, S.M., da Silva, G.V. and Militao, J.S.L.T. (2012). Non-substituted B-ring flavonoids

- and an indole alkaloid from *Piper aleyreanum* (Piperaceae). *Biochemical and Systematic Ecology*. 45: 206-208.
- 114. Rajudin, E., Ahmad, F., Sirat, H.M., Arbain, D. and Aboul-Enein, H.Y. (2010). Chemical constituents from tiger's betel, *Piper porphyrophyllum* N.E.Br. (Fam. Piperaceae). *Natural Product Research*. 24: 387–390.
- 115. Salleh, W.M.N.H.W, Ahmad, F., Yen, K.H. and Sirat, H.M. (2012). Chemical compositions and antibacterial activity of the leaf and stem oils of *Piper Porphyrophyllum* (Lindl.) N.E. BR. *EXCLI Journal*. 11: 399-406.
- 116. Navickiene, H.M.D., Morandim, A.A., Alécio, A.C., Regasini, L.O., Bergamo, D.C.B., Telascrea, M., Cavalheiro, A.J., Lopes, M.N., Furlan, V.S.B.M., Marques, M.O.M., Young, M.C.M., Kato, M.J. (2006). Composition and antifungal activity of essential oils from *Piper aduncum*, *Piper arboreum* and *Piper tuberculatum*. *Química Nova*. 29: 467-470.
- 117. Silva, D.R., Endo, E.H., Filho, B.P.D., Nakamura, C.V., Svidzinski, T.I.E., Souza, A., Young, M.C.M., Nakamura, T.U., Cortez, D.A.G. (2009). Chemical Composition and Antimicrobial Properties of *Piper ovatum* Vahl. *Molecules*. 14: 1171-1182.
- 118. Perigo, C.V., Torres, R.B., Bernacci, L.C., Guimarães, E.F., Haber, L.L., Facanali, R., Vieira, M.A.R., Quecini, V. and Marques, M.O.M. (2016). The chemical composition and antibacterial activity of eleven *Piper* species from distinct rainforest areas in Southeastern Brazil. *Industrial Crops and Products*. 94: 528–539.
- 119. Silva, J.K.R., Silva, J.R.A., Nascimento, S.B., Luz, S.F.M., Meireles, E.N., Alves, C.N., Ramos, A.R. and Maia, J.G.S. (2014). Antifungal Activity and Computational Study of Constituents from *Piper divaricatum* Essential Oil against Fusarium Infection in Black Pepper. *Molecules*. 19: 17926-17942.
- 120. Silva, J.K.R., Pinto, L.C., Burbano, R.M.R., Montenegro, R.C., Guimaraes, E.F., Andrade, E.H.A. and Maia, J.G.S. (2014). Essential oils of Amazon *Piper* species and their cytotoxic, antifungal, antioxidant and ati-cholinesterase activities. *Industrial Crops and Products*. 58: 55-60.
- 121. Matasyoh, J.C., Wagara, I.N., Nakavuma, J.L. and Chepkorir, R. (2013). Chemical composition and antifungal activity of *Piper capensi* oil against mycotoxigenic *Aspergillus*, *Fusarium* and *Penicillium* species. *International Journal of Biological and Chemical Sciences*. 7: 1441-1451.

- 122. Oyedeji, O.A., Adeniyi, B.A. and Konig, W.A. (2005). Essential oil composition of *Piper guineense* and its antimicrobial activity. Another chemotype from Nigeria. *Phytotherapy Research*. 19: 362-364.
- 123. Santana, H.T., Trindade, F.T.T., Stabeli, R.G., Silva, A.A.E., Militao, J.S.L.T. and Facundo, V.A. (2015). Essential oils of leaves of *Piper* species display larvicidal activity against the dengue vector, *Aedes aegypti* (Diptera: Culicidae). *Revista Brasileira de Plantas Medicinais*. 17: 105-111.
- 124. Autran, E.S., Neves, I.A., da Silva, C.S.B., Santos, G.K.N., Camara, C.A.G. and Navarro, D.M.A.F. (2009). Chemical composition, oviposition deterrent and larvacidal activities against *Aedes aegypti* of essential oils from *Piper marginatum* Jacq. (Piperaceae). *Bioresource Technology*. 100: 2284-2288.
- 125. Santana, A.I., Vila, R., Cañigueral, S. and Gupta, M.P. (2016). Chemical Composition and Biological Activity of Essential Oils from Different Species of *Piper* from Panama. *Planta Medica*. 82: 986–991.
- 126. Bagheri, H., Manap, M.Y.A. and Solati, Z. (2014). Antioxidant activity of *Piper nigrum* L. essential oil extracted by supercritical CO<sub>2</sub> extraction and hydro-distillation. *Talanta*. 121: 220-228.
- 127. Mageed, M.A.A., Mansour, A.F., Massry, K.F.E., Ramadan, M.M. and Shaheen, M.S. (2011). The effect of microwave of essential oils of white and black pepper (*Piper nigrum* L.) and their antioxidant activity. *Journal of Essential Oil Bearing Plants*. 14: 214-223.
- 128. Oboh, G., Ademosun, A.O., Odubanjo, O.V. and Akinbola, I.A. (2013). Antioxidative properties and inhibition of key enzymes relevant to type-2 diabetis and hypertension by essential oils from black pepper. *Advance Pharmaceutical Sciences*, 47: 1-6.
- Carmo, D.F.M., Amaral, A.C.F., Machado, G.M.C., Leon, L.L., and Silva, J.R.
   (2012). Chemical and Biological Analyses of the Essential Oils and Main Constituents of *Piper Species*. *Molecules*. 17: 1819-1829.
- 130. Leal, S.M., Pino, N., Stashenko, E.E., Martínez, J.R. and Escobar, P. (2013). Antiprotozoal activity of essential oils derived from *Piper* spp. grown in Colombia. *The Journal of Essential Oil Research*. 25: 512–519.
- 131. Marques, A.M., Barreto, A.L.S., Curvelo, J.A.R., Romanos, M.T.V., Soares, R.M. and Kaplan, M.A.C. (2011). Antileshmanial activity of nerolidol-rich

- essential oil from *Piper claussenianum*. *Revista Brasileira de Farmacognosia*. 21: 908-914.
- 132. Avella, E. and Motta, J.R. (2010). Main constituents and cytotoxic activity of the essential oil of *Piper artanthe*. *Chemistry of Natural Compounds*. 46: 651-654.
- 133. Ramirez, J., Cartuche, L., Morocho, V., Aguilar, S. and Malagon, O. (2009). Antifungal activity of raw extract and flavanons isolated from *Piper ecuadorense* from Ecuador. *Brazilian Journal of Pharmacognosy*. 23: 370-373.
- 134. Scodro, R.B.L., Pires, C.T.A., Carraraa, V.S., Lemos, C.O.T., Filho, L.C., Souza, V.A., Corrêa, A.G., Siqueira, V.L.D., Lonardoni, M.V.C., Cardoso, R.F. and Cortez, D.A.G. (2013). Anti-tuberculosis neolignans from *Piper regnellii*. *Phytomedicine*. 20: 600- 604.
- 135. Robert, P. and Adams. (2007). Identification of Essential Oil Components By Gas Chromatography/Mass Spectrometry. 4th Edition. Allured Publishing Corporation.
- 136. Facundo, V.A., Polli, A.R., Rodrigues, R.V., Militao, J.S.L.T., Stabeli, R.G. and Cardoso, C.T. (2008). Fixed and volatile chemical constituents from stems and fruits of *Piper tuberculatum* Jacq. and from roots of *P. hispidum* H. Bk. *Acta Amazonia*. 38: 4-6.
- 137. Thanh, L., Dung, N.X., Bighelli, A., Casanova, J. and Leclecrq, P.A. (1997). Combination of capillary GC, GC-MS and <sup>13</sup>C-NMR for the characterization of the rhizome oil of *Piper betel* L. (Piperaceae) from Vietnam. *Spectroscopy* (*Armsterdam*). 13: 131-136.
- 138. Lognay, G.C., Bouxin, P., Marlier, M., Haubruge, E., Gaspar, C. and Rodriguez, A. (1996). Composition of the essential oil of *Piper acutifolium* Ruiz. And Pav. from Peru. *Journal of Essential Oil*. 8: 689-691.
- 139. Knothe, G. and Kener, J.A. (2004). Determination of the fatty acid profile from the <sup>1</sup>H-NMR spectroscopy. *European Journal of Lipid Science and Technology*. 106: 88-96.
- 140. Purcell, J.M., Morris, S.G. and Susi, H. (1966). Proton Magnetic Resonance Spectra of Unsaturated Fatty Acids. *Analytical Chemistry*. 38: 588-591.

- 141. Kanti, A., Sukara, E., Latifah, K., Sukarno, N. and Boundy-Mills, K. (2013). Indonesian oleaginous yeasts isolated from *Piper betle* and *P. nigrum*. *Mycosphere*. 4: 1015–1026.
- 142. Lakshmi, A., Menuka, A. and Damisha, R. (2006). *Piper betle*: a potential natural antioxidant. *International Journal of Food Science and Technology*. 41: 10-14.
- 143. Takaya, Y., Kondo, Y., Furukawa, T. and Niwa, M. (2003). Antioxidant Constituents of Radish Sprout (Kaiware-daikon), *Raphanus sativus. Journal of Agricultural and Food Chemistry*. 51: 8061-8066.
- 144. Val'eria, F.P., Jenifer, S.M.I., Melecchi, S., Fernanda, C.A., Rosangela de Assis, J., Migdalia, M.M. and Eniz, C. O. (2006). Comparison of soxhlet, ultrasound-assisted and pressurized liquid extraction of terpenes, fatty acids and Vitamin E from *Piper gaudichaudianum* Kunth. *Journal of Chromatography A.* 1105: 115-118.
- 145. Becker, E.D. (1962). Proton magnetic resonance studies relating to the stereochemistry of sesamin, asarinin and episarinin. *Tetrahedron letters*. 4: 157-163.
- 146. Hsieh, T., Lu, L. and Su, C. (2005). NMR spectroscopic, xray crystallographic and theoretical studies of molecular mechanics of natural products: farformolide B and sesamin. *Biophysical Chemistry*. 114:13-20.
- 147. Sawangjaroen, N., Sawangjaroen, K and Poonpanag, P. (2004). Effects of *Piper longum* fruit, *Piper sarmentosum* root and *Quercus infectoria* nut gall on *caecal ameobiasis* in mice. *Journal of Ethnopharmacology*. 91: 357-360.
- 148. Pierre, L.L. and Moses, M.N. (2015). Isolation and Characterisation of Stigmasterol and β-Sitosterol from *Odontonema Strictum* (Acanthaceae). *Journal of Innovations in Pharmaceuticals and Biological Sciences*. 2: 88-95.
- 149. Fatema, M.K., Chen, Z.Y. and Wei, G. (2015). β-sitosterol reduce cholesterol levels in high cholesterol diet fed zebrafish. *International Journal of Natural and Social Sciences*. 2: 53-65.
- Oliveira, A.P., Ferreira, J.G., Riboira, S., Andrade, P.B. and Valentão, P.
   (2016). Bioactive Natural Products from *Piper betle* L. Leaves and their α-Glucosidase Inhibitory Potential. *Records of Natural Products*. 10: 771-781.
- 151. Lee, J.M., Lee, D.G., Lee, K.H., Cho, S.H., Nam, K.W. and Lee, S. (2013). Isolation and identification of phytochemical constituents from the fruits of

- Acanthopanax senticosus. African Journal of Pharmacy and Pharmacology. 7: 294-301.
- 152. Wu, Q., Wang, S., Tu, G., Feng, Y. and Yang, J. (1997). Alkaloids from *Piper puberullum*. *Phytochemistry*. 44: 727-730.
- 153. Kumar, V., Prasad, A.K. and Parmar, V.S. (2003). Naturally Occuring Aristolactams, Aristolochic Acids and Dioxoaporphines and their Biological Activities. *Natural Products Repository*. 20: 563-583.
- 154. Diaz, A.M.P., Diaz, P.P. and Nathan, P.J. (1990). Dioxoaporphine Alkaloid and Flavone from *Piper manausense* YUNK. *Revista Colombiana de Quimica*. 19: 63-66.
- 155. Ee, G.C.L., Lim, S.K., Lim, C.M. and Dzulkefly, K. (2008). Alkaloids and Carboxylic Acids from *Piper nigrum*. *Asian Journal of Chemistry*. 20: 5931-5940.
- 156. Tsuruta, A.Y., Bomm, M.D., Lopes, M.N. and Lopes, L.M.X. (2002). Aristolactams and further constituents from *Aristolochia chamissonis*. *Eclética Química*. 27: 1-4.
- 157. Ahmad, F., Jamil, S. and Read, R.W. (1995). Isobutylamides from *Piper ridleyi*. *Phytochemistry*. 40: 1163-116.
- 158. Sucrow, W. and Polysou, P. (1971). Biosynthesis of 24-ethylcholesta-5,22,25-trien-3β-ol. *Tetrahedron Letters*. 21: 1883-1884.
- 159. Pandey, R., Verma, R.K., Singh, S.C. and Gupta, M.M. (2003). 4α-Methyl-24β-ethyl-5α-cholesta-14,25-dien-3β-ol and 24β-ethylcholesta-5,9(11),22E-trien-3β-ol sterols from *Clerodendrum inerme*. *Phytochemistry*. 63: 415–420.
- 160. Bolger, L.M., Rees, H.H., Gisalberti, E.L., Goad, L.J. and Goodwin, T.W. (1970). Biosynthesis of 24-Ethylcholesta-5,22,25-trien-3β-ol. A New Sterol from *Clerodendrum campbellii*. *Journal of Biochemistry*. 118: 197-200.
- 161. Gaspar, H., Brito, F.M.S., de la, Torre. M.C. and Rodriguez, B. (1996). Sterols from *Teucrium abutiloides* and *T. betonicum*. *Phytochemistry*. 43: 613-615.
- 162. Gaspar, E.M.M. and Neves, H.J.C.D. (1993). Steroidal Constituents from Mature Wheat Straw. *Phytochemistry*. 34: 523-527.
- 163. Wei, K., Li, W., Koike, K., Pei, Y., Chen, Y. and Nikaido, T. (2004). Complete 

  <sup>1</sup>H and <sup>13</sup>C NMR assignments of two phytosterols from roots of *Piper nigrum*. *Magnetic Resonances Chemistry*. 42: 355–359.

- 164. Barla, A., Irman, B., Kultur, S. and Oksuz, S. (2000). Secondary Metabolites from *Euphorbia helioscopia*. *Turkish Journal of Chemistry*. 30: 325-332.
- 165. Galloa, M.B.C., Cavalcanti, B.C., Barros, F.W.A., de Moraes, M.O., Costa-Lotufo, L.V., Pessoa, C., Bastos, J.K. and Pupo, M.T. (2010). Chemical Constituents of *Papulaspora immersa*, an Endophyte from *Smallanthus sonchifolius* (Asteraceae), and Their Cytotoxic Activity. *Chemistry and Biodiversity*. 7: 2941-2541.
- 166. Othman, S.N.A.M., Sarker, S.D., Talukdar, A.D., Ningthoujam, S.S., Khamis, S., and Basar, N. (2014). Chemical Constituents and Bacterial Activity of Phaleria macrocarpa (Scheff.) BOERL. International Journal of Pharmaceutical Sciences and Research. 5: 3157-3162.
- 167. Alves, J.S., de Castro, J.C.M., Freire, M.O., da-Cunha, E.V.L., Filho, J.M.B. and de Silva, M.S. (2000). Complete assignment of the <sup>1</sup>H and <sup>13</sup>C NMR spectra of four triterpenes of the ursane, artane, lupane and friedelane groups. *Magnetic Resonance in Chemistry*. 38: 201–206.
- 168. Ke, Z., Wei, N. and Changxiang, C. Tianran. (1999). Chemical constituents of *Piper boehmeriaefolium* var. Tonkinense. *Chanwu Yanjiu Yu Kaifa*. 11: 44-47.
- 169. Iqbal, E., lim, L.B.L., Salim, K.A., Faizi, S., Ahmed, A. and Mohamed, A.J. (2017). Isolation and Characterization of Aristolactam Alkaloids from the Stem Bark of *Goniothalamus velutinus* (Airy Shaw) and their Biological Activities. *Journal of King Saud University-Science*. 12: 132-138.
- 170. Rudolf, H., Anneliese, L., Arturo, G.P. (1975). Aporphine-type Alkaloids from *Piper auritum. Lloydia*. 38: 529-530.
- 171. Rudolf, H. and Anneliese, L. (1976). An Aporphine Alkaloid from *Piper sanctum. Phytochemistry*. 15: 1323-1329.
- 172. Jaggy, H. and Achenbach, H. (1992). Cepharadione A from *Piper methysticum*. *Planta Medica*. 58: 111-118.
- 173. Eliseo, A., Pedro, D., Aura, M.P.D. (1994). Constituents from *Piper divarcatum*. *Planta Medica*. 60: 195-121.
- 174. Boll, P.M., Prasad, A.K., Tyagi, O.D., Wengel, J., Olsen, C.E., Kumar, N., Bishit, K.S. and Parmar, V.S. (1996). Neolignans, cyclohexane and alkaloids from *Piper wightii. Recueil des Travaux Chimiques des Pays-Bas*. 115: 9-12.

- 175. Singh, S.K., Prasad, A.K., Olsen, C.E., Jha, A., Jain, S.C., Parmar, V.S. and Wenger, J. (2008). Neolignans and Alkaloids from *Piper argyrophylum*. *Phytochemistry*. 43: 1355-1360.
- 176. Pena, L.A., Avella, E. and Puentes, D.A.M. (2000). Prenylated benzoquinone and hydroquinone and other isolated constituents of *Piper bogotense* C.DC. *Revista Colombiana de Quimica*. 29: 25-37.
- 177. Chen, Y.C., Chen, J.J., Chang, Y.L., Teng, C.M., Lin, W.Y., Wu, C.C. and Chen, I.S. (2004). A New Aristolactam Alkaloid and Anti-Platelet Aggregation Constituents from *Piper taiwanense*. *Planta Medica*. 70: 174-177.
- 178. Ma, J., Jones, S.H., Marshall, R., Johnson, R.K. and Hecht, S.M. (2004). A DNA-Damaing oxoaporphine Alkaloid from *Piper caninum. Journal of Natural Products*. 67: 1162-1164.
- 179. Li, C.Y., Tsai, W.J., Damu, A.G., Lee, E.J., Wu, T.S., Dung, N.X., Thang, T.D. and Thanh, L. (2007). Isolation and Identification of Antiplatelet Aggregatory Principles from the Leaves of *Piper lolot. Journal of Agricultural and Food Chemistry*. 55: 9436-9442.
- 180. Tabopda, T.K., Mitaine, O.A.C., Miyamoto, T., tanaka, C., Ngadjui, B.T. Lacaille, D.M.A. (2012). Secondary Metabolites from Polar Fractions of *Piper umbellatum*. *Natural Product Communications*. 7: 595-596.
- 181. Rajeev, G. and Jain, S.C. (2014). Phytochemical Investigation Afforded a Novel Cycloartane Triterpenoid from *Piper thomsoni*. *Research Journal of Chemical Sciences*. 4: 7-11.
- 182. Desai, S.J., Chaturvedi, R.N., Badheka, L.P., Mulchandani, N.B. (1989). Aristolactams and 4,5-dioxoaporphines from Indian *Piper* species. *Indian Journal of Chemistry: Section B: Organic Chemistry Including Medicinal Chemistry*. 288: 775-777.
- 183. Chaves, M.C.D.O., Oliveira, A.H. and Santos, B.V.D.O. (2006). Aristolactams from *Piper marginatum* Jacq. (Piperaceae). *Biochemical Systematics and Ecology*. 34: 75-77.
- 184. Chen, Y.C., Liao, C.H. and Chen, I.S. (2007). Lignans, an Amide and Antiplatelet Activities from *Piper philippinum*. *Phytochemistry*. 68: 2101-2111.
- 185. Kim, K.H., Choi, J.W., Choi, S.U., Ha, S.K., Kim, S.Y., Park, H.J. and Lee, K.R. (2011). The Chemical Constituents of *Piper kadsura* and Their Cytotoxic

- and Anti-neuroinflammatory Activities. *Journal of Enzyme Inhibition and Medicinal Chemistry*. 26: 254-260.
- 186. Nobsathian, S., Tuchinda, P., Soorukram, D., Pohmakotr, M., Reutrakul, V., Yoosook, C., Kasisit, J., Napaswad, C. (2012). A new conjugated amide-dimer from the aerial parts of *Piper submultinerve*. *Natural Product Research*. 26: 1824-1830.
- 187. Lei, H.P., Chen, X.Q., Qiao, C.F., Liu, Y. and Zhao, J. (2014). Chemical Constituents from Twigs of *Piper hancei. Zhongyaocai*. 37: 69-71.
- 188. Mata, R., Morales, I., Perez, O., Cruz, I.R., Acevedo, L., Mendoza, I.E., Bye, R., Franzblau, S. and Timmermann, B. (2004). Antimycobacterial Compounds from *Piper sanctum. Journal of Natural Products*. 67: 1961-1968.
- 189. Tsai, I.L., Lee, F.P., Wu, C.C., Duh, C.Y., Ishikawa, T., Chen, J.J., Chen, Y.C., Seki, H. and Chen, I.S. (2005). New Cytotoxic Cyclobutanoid Amides, a New Furanoid Lignan and Anti-Platelet Aggregation Constituents from *Piper arborescens*. *Natural Product Chemistry*. 71: 535-542.
- 190. Ghosh, K. and Bhattacharya, T.K. (2005). Chemical constituents of *Piper betle* Linn. (Piperaceae) roots. *Molecules*. 10: 798-802.
- 191. Liang, Z., Junshan, Y. and Guangzhong, T. (2005). Study on chemical components of *Piper hancei. Zhongguo Yaoxue Zazhi*. 40: 184-185.
- 192. Anhua, W., Daonian, Z., Jinlan, R., Yaling, C., Chaomei, X., Guanghua, W. and Huibin, L. (2011). Study on the contents of aristololactams in *Piper wallichii* and their renal cytotoxicities. *Journal of Food and Drug Analysis*. 19: 349-354.
- 193. Salleh, W.M.N.H.W., Ahmad, F.; Yen, K.H. (2014). Antioxidant and anti-tyrosinase activities from *Piper officinarum* C.DC (Piperaceae). *Journal of Applied Pharmaceutical Science*. 4: 087-091.
- 194. Wu, D., Nair, M.G. and DeWitt, D.L. (2002). Novel Compounds from *Piper methysticum* Forst (Fava Kava) Roots and their Effect on Cyclooxygenase Enzyme. *Journal of Agricultural and Food Chemistry*. 50:701-706.
- 195. Mounyr, B., Moulay, S. and Saad, K. I. (2016). Methods for in vitro evaluating antimicrobial activity: A review. *Journal of Pharmaceutical Analysis*. 6: 71–79.

- 196. Thenmozhi, M. and Sivaraj, R. (2010). Phytochemical Analysis and Antimicrobial Activity of *Polyathia longifolia*. *International Journal of Pharmaceutical Biological Science*. 1: 1-7.
- 197. Yazdankhah, S.P., Sorum, H., Larsen, H.J. and Gogstad, G. (2001). Use of Magnetic Beads for Gram Staining of Bacteria in Aqueous Suspension. *Journal of Microbiology Methods*. 47: 369-371.
- 198. Chai, H., William, E.A. and Rickey, P.H. (2014). Synthetic Antimicrobial Peptides Exhibit Two Different Binding Mechanisms to the Lipopolysaccharides Isolated from *Pseudomonas aeruginosa* and *Klebsiella pneumonia*. *International Journal of Medicinal Chemistry*. 1-13.
- 199. Guilloteau, J.P., Mathieu, M., Giglione, C., Blanc, V., Dupuy, A., Chevrier, M., Gil, P., Famechon, A., Meinnel, T. and Miko, V. (2002). The Crystal Structures of Four Peptide Deformylases Bound to the Antibiotic Actinonin Reveal Two Distinct Types: A Platform for the Structure-based Design of Antibacterial Agents. *Journal of Molecular Biology*. 32: 951–962.
- 200. Ellof, J.A. (1998). Quick Microplate Method to Determine the Minimum Inhibitory Sensitive and Concentration of Plant Extracts for Bacteria. *Planta Medica*. 64: 711-713.
- 201. Hood, J.R., Wilkinson, J.M. and Cavanagh, H.M.A. (2003). Evaluation of Common Antibacterial Screening Methods Utilized in Essential Oil Research. *Journal of Essential Oil Research*. 15: 428-433.
- 202. Langfield, R.D., Scarano, F.J., Heitzman, M.E., Kondo, M., Hammond, G.B. and Neto, C.C. (2004). Use of a modified microplate bioassay method to investigate antibacterial activity in the *Peruvian* medicinal plant *Peperomia galioides*. *Journal of Ethnopharmacology*. 94: 279-281.
- 203. Randhawa, M.A. (2006). The Effect of Dimethyl Sulfoxide (DMSO) on the growth of Dermatophytes. *Japan Journal of Medical Mycology*. 47: 515-518.
- 204. Perumal, S., Pillai, S. Cai, L.W., Mahmud, R. and Ramanathan, S. (2012). Determination of Minimum Inhibitory Concentration of *Euphorbia hirta* (L.) Extracts by Tetrazolium Microplate Assay. *Journal of Natural Products*. 5: 68-76.
- 205. Stiefel, P., Schneider, J., Amberg, C., Weber. and Ren, Q. (2016). A simple and rapid method for optical visualization and quantification of bacteria on textiles. *Scientific Reports*. 6: 39635.

- 206. Sousa, R.M.F., Morais, S.A.L.D., Vieira, R.B.K., Napolitano, D.R., Guzman, V.B., Moraes, T.S., Cunha, L.C.S., Martins, C.H.G., Chang, R., Aquino, F.J.T.D., Nascimento, E.A.D., and Oliveira, A.D. (2015). Chemical composition, cytotoxic, and antibacterial activity of the essential oil from *Eugenia calycina* Cambess. leaves against oral bacteria. *Industrial Crops and Products*. 65: 71–78.
- 207. Cárdenas, J., Rojas, J., Rojas, L.F., Lucena, M. and Buitrago. (2012). Essential oil composition and antibacterial activity of *Monticalia greenmaniana* (Asteraceae). *Natural Products Communications*. 7:243-247.
- 208. Yang, S.W., Chan, T.M. and Terracciano, J. (2009). Caryophyllenes from a fungal culture of *Chrysosporium pilosum* (perpendicular). *Journal of Natural Products*, 72: 484–487.
- 209. Sousa, E.O., Barreto, F.S., Rodrigues, F.G., Camposa, A.R. and Costa, J.G.M. (2012). Chemical composition of the essential oils of *Lantana camara* L. and *Lantana montevidensis* Briq. and their synergistic antibiotic effects on aminoglycosides. *The Journal of Essential Oil Research*. 24: 447–452.
- 210. Rios, J.L. and Recio, M.C. (2005). Medicinal Plants and antimicrobial Activity. *Journal of Ethnopharmacology*. 100: 80-84.
- 211. Desbois, A.P., and Lawlor, K.C. (2013). Antibacterial Activity of Long-Chain Polyunsaturated Fatty Acids against *Propionibacterium acnes* and *Staphylococcus aureus*. *Marine Drugs*. 11: 4544-4557.
- 212. Parsons, J.B., Yao, J., Frank, M.W., Jackson, P. and Rock, C.O. (2012). Membrane Disruption by Antimicrobial Fatty Acids Releases Low-Molecular Weight Proteins from *Staphylococcus aureus*. *Journal of Bacteriology*. 194: 5294–5304.
- Desbois, A.P. and Smith, V.J. (2010). Antibacterial free fatty acids: activities, mechanisms of action and biotechnological potential. *Applied Microbiology Biotechnology*. 85:1629–1642.
- 214. Pala, F.S. and Gurkan, H. (2008). The Role of Free Radicals in Ethiopathogenesis of Diseases. *Advance Molecular Biology*. 1: 1-9.
- 215. Tegeli, V., Karpe, P. and Vikas, K. (2006). Importance of Free Radical and Antioxidant in Human Health. *International Journal of Pharmaceutical, Chemical and Biological Sciences*. 4: 1038-1050.

- 216. Alafiatayo, A.A., Ahmad, S. and Maziah, M. (2014). Total Antioxidant Capacity, Total Phenolic Compounds and the Effects of Solvent Concentration on Flavonoid content in *Curcuma longa* and *Curcuma xanthorhhiza* Rhizomes. *Medicinal Aromatic Plants*. 3: 2.
- 217. Princemail, J., Ricour, C., Defrraigne, J.O. and Petermans, J. (2014). Oxidative Stress, Antioxidants and the Ageing Process. *Reveue Medicale de Liege*. 69: 270-275.
- 218. Yu, K., Zhao, M., Yang, B., Zhao, Q. and Jiang, Y. (2007). Phenolics from hull of *Garcinia mangostana* Fruit and their Antioxidant Activities. *Food Chemistry*. 104: 176-181.
- 219. Qayum, M., Zua-ul-Haq, M., Kaleem, W.A., Ahmad,S., Calani, L., Mazzeo, T. and Pellergini, N. (2104). Antioxidant Potential of Impatients bicolour Royle and Zizypus oxyphylla Edgew. Pakistan Journal of Botanical. 46: 1725-1729.
- 220. Daniela, S., bubelova, S., Snyed, J., Erb-Weber, S. and Mleck, J. (2015). Total Phenolics, Flavonoids, Antioxidant Capacity, Crude Fibre and Digestibility in non-Traditional Wheat Flakes and Muesli. *Food Chemistry*. 174: 319-325.
- 221. Ayaz, M.M., Junaid, M., Ahmad, J., Ulah, F., Sadiq, A., Ahmad, S. and Imran, M. (2014). Phenolic Contents, Antioxidant and Anticholinesterase Potentials of Crude Extracts, Subsequent Fractions and Crude Saponins from *Polygonum hydropiper L. BMC Complementary Alternative Medicine*. 14: 145.
- 222. Tewtrakul, S., Sanan, S. and Kummee, S. (2008). Anti-allergic Activity of Compounds from *Kaempferia parviflora*. *Journal of Ethnopharmacology*. 116: 191-193.
- 223. Belsare, D.P., Pal, S.C., Kankate, R.S. and Vanjari, S.S. (2010). Evaluation of Antioxidant Activity of Chalcones and Flavanoids. *International Journal ChemTech Research*. 2: 1080-1089.
- 224. Thaipong, K., Boonprakaob, U., Crosby, K., Cisneros, L.Z. and Byrne, D.H. (2006). Comparison of ABTS, DPPH, FRAP and ORAC Assays for estimating antioxidant activity from guava fruit extracts. *Journal of Food Computational Analysis*. 19: 669-675.
- 225. Miller, H.E., Rigelhof, F., Marquat, L., Prakash, A. and Kanter, M. (2000). Antioxidant Content of Whole Grain Breakfast Cereals, Fruits and Vegetables. *Journal of American College Nutrition*. 19: 312-319.

- 226. Medzhitov, R. (2010). Inflammation 2010: New Adventures of an Old Flame. *Cell.* 140: 771-776.
- 227. Leyen, K.V. (2013). Lipoxygenase: An Emerging Target for Stroke Therapy. CNS & Neurological Disorders - Drug Targets. 12: 191-199.
- 228. Bisht, R., Bhattacharya, S., Jaliwala, Y.A. (2014). COX and LOX inhibitory potential of *Abroma augusta* and *Desmodium gangeticum*. *The Journal of Phytopharmacology*. 3: 168-175
- 229. Pelletier, J.M., Lajeunesse, D., Reboul, P. and Pelletier, J.P. (2003). Therapeutic role of dual inhibitors of 5-LOX and COX, selective and non-selective non-steroidal anti-inflammatory drugs. *Annals of Rheumatic Desease*. 62: 501-509.
- 230. Wisastra, R. and Dekker, F.J. (2014). Inflammation, Cancer and Oxidative Lipoxygenase Activity are Intimately Linked. *Cancers*. 6: 1500-1521.
- 231. Salmon, A.A., Dzoyem, J.P., Shai, L.J. and Eloff, J.N. (2015). The antiinflammatory and antioxidant activity of 25 plant species used traditionally to treat pain in southern African. *BMC Complementary and Alternative Medicine*. 15: 159-163.
- 232. Rackova, L., Oblozinsky, M., Kostalova, D., Kettmann, V. and Bezakova, L. (2007). Free radical scavenging activity and lipoxygenase inhibition of *Mahonia aquifolium* extract and isoquinoline alkaloids. *Journal of Inflammation*. 4: 15-19.
- 233. Oomah, B.D., Corbe, A.L. and Balasubramaniam, P. (2010). Antioxidant and Anti-inflammatory Activities of Bean (*Phaseolus vulgaris* L.) Hulls. *Journal of Agricultural Food Chemistry*. 58: 8225–8230.
- 234. Lee, J.H. and Kim, G.H. (2010). Evaluation of Antioxidant and Inhibitory Activities for Different Subclasses Flavonoids on Enzymes for Rheumatoid Arthritis. *Journal of Food Science*. 75: 1-7.
- 235. Kgun, S.E., Gyeong, Y.H.C and Cho, Y.S. (2012). Synthesis and Biological Evaluation of *N*-aryl- 4-aryl-1,3-Thiazole-2-Amine Derivatives as Direct 5-Lipoxygenase Inhibitors. *Chemical Biology Drug Deseases*. 80: 90–99.
- 236. Pontik, E. and Litina, D.H. (2007). Synthesis of Phenyl-substituted Amides with Antioxidant and Antiinflammatory activity as Novel Lipoxygenase Inhibitors. *Medicinal Chemistry*. 3: 175-186

- 237. Rani, P., Pal, D., Hegde, R.R. and Hashim, S.R. (2014). Anticancer, Anti-Inflammatory, and Analgesic Activities of Synthesized 2-(Substituted phenoxy) Acetamide Derivatives. *BioMed Research International*. 9:1-9.
- 238. Lee, Y.T., Hsieh, Y.L. Yeh, Y.Y.H. and Huang, C.Y. (2015). Synthesis of phenolic amide as evaluation antioxidative and anti-inflammatory *in vitro* and *in vivo*. *RSC Advance*. DOI: 10.1039/C5RA14137K.
- 239. Chawla, S., deLong, M.A., Visscher, M.O., Wickett, R.R., Manga, P. and Boissy, R.E. (2008). Mechanism of tyrosinase inhibition by deoxy-Arbutin and its second-generation derivatives. *British Journal of Dermatology*. 159: 1267-1274.
- 240. Iozumi, K., Hoganson, G.E., Pennella, R., Everett, M.A. and Fuller, B.B. (1993). Role of Tyrosinase as the Determinant of Pigmentation in Cultured Human Melanocytes. *Journal of Investigated Dermatology*. 100: 806-812.
- 241. Lin, J.W., Chiang, H.M., Lin, Y.C. and Wen, K.C. (2008). Natural Products with Skin Whitening Effects. *Journal of Food and Drug Analysis*. 16: 1-10.
- 242. Maghsoudi, S., Adibi, H., Hamzeh, M., Kooshka, M.R.A. Taviranic, M.R. and Khodarahmi, K. (2013). Kinetic of Mushroom Tyrosinase Inhibition by Benzaldehyde Derivatives. *Journal of Reports in Pharmaceutical Sciences*. 2: 131-139.
- 243. Palumbo, A., Ischia, M., Misuraca, G. and Prota, G. (1991). Mechanism of inhibition of melanogenesis by hydroquinone. *Biochimica and Biophysic Acta*. 1073: 85-90.
- 244. Seo, S.Y., Sharma, V.K., and Sharma, N. (2003). Mushroom Tyrosinase: Recent Prospects. *Journal of Agricultural Food and Chemistry*. 51: 2837-2853.
- 245. Zaidi, K.U., Ali, A.S., Ali, S.A. and Naaz, I. (2014). Microbial Tyrosinases: Promising Enzymes for Pharmaceutical, Food Bioprocessing, and Environmental Industry. *Biochemistry Research International*. 12: 1-16.
- Azami, F., Lemeski, E.T., Janlou, M.A.M. (2017). Kojic Acid Effect on the Inhibitory Potency of Tyrosinase. *Journal of Chemical Health Risks*. 7: 147-155.
- 247. Ching, L., Rowa, M. and Ho, J.C. (2009). The Antimicrobial Activity, Mosquito Larvicidal Activity, Antioxidant Property and Tyrosinase Inhibition of *Piper betle. Journal of the Chinese Chemical Society*. 56: 653-658.

- 248. Salleh, W.M.N.H.W.M., Hashim, N.A., Ahmad, F. and Yen, K.H. (2014). Anticholinesterase and Antityrosinase Activities of Ten *Piper Species* from Malaysia. *Advance Pharmaceutical Bulletin*. 4: 527-53.
- 249. Kashima, Y. and Miyazawa, M. (2012). Synthesis and Biological Evaluation of Bergenin Analogues as Mushroom Tyrosinase Inhibitors. Archives Pharmacal Research. 35: 1533-1541.
- 250. Bernard, P. and Berthon, J.Y. (2000). Resveratrol: an original mechanism on tyrosinase inhibition. *International Journal of Cosmetic Science*. 22: 219-226.
- 251. Kim, Y.M., Yun, J., Lee, C.K., Lee, H., Min, K.R. and Kim, Y. (2002). Oxyresveratrol and Hydroxystilbene Compound: Inhibitory Effect on Tyrosinase and Mechanism of Action. *The Journal of Biological Chemistry*. 277: 16340-16344.
- 252. Rao, G.V., Rao, K.S., Mukhopadhyay, T. and Madhavi, M.S.L. (2012). Alkamides and their biological activity from *Piper longum* Linn. *Journal of Pharmacy Research*. 5: 165-168.
- Chapin, K.C. and Laudrdale, T. Reagents, Stains and Media: Bacteriology. In Murray, P.R., Baron, E.J., Jorgensen, J.H., Pfaller, M.A. and Yolken, R.H. (2003). *Manual of Clinical Microbiology*. Washington DC:ASM Press. p. 358.
- 254. Kaikabo, A.A. and Eloff, J.N. (2011). Antibacterial Activity of Two Biflavonoids from Garcinia livingstonei Leaves against *Mycobacterium smegmatis*. *Journal of Ethnopharmacology*. 138: 253-255.
- 255. Pallant, J. *SPSS Survival Manual*. 3<sup>rd</sup> Edition. Australia: Open University Press.