Pharmacological Evaluation of Green Chilli in Alzheimer’s Disease

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ABSTRACT
Background: Green chilli pepper (Capsicum annum L. car frutescens) is popular as a spice used in many cuisines across the world. Capsaicin, as the pharmacologically active composition of chilli pepper, has been shown to possess antioxidant, local analgesic, and hypoglycemic effects according to many studies conducted on the red chilli pepper. However, there is no data associated with the role of green chilli peppers in Alzheimer’s disease (AD) caused by capsaicin, rather than red chilli peppers. Objectives: To evaluate the acetylcholinesterase (AChE) and beta-secretase enzyme inhibition of ethanolic extract of Capsicum frutescens (EECF) or known as green chilli in in-vitro manner. Materials and Methods: Extraction was carried out on the fruits of Capsicum frutescens with ethanol. Its ethanolic extract in different concentrations was applied as the analyte to test its anti-acetylcholinesterase activity and enzymatic inhibitory action on the beta-secretase. Donepezil with various concentrations was also prepared as the control set for comparison of effect with EECF. Both the analytes were tested using the QuantiChromTM and BACE-1 FRET assay kits, for evaluation of inhibition activity against acetylcholinesterase and BACE-1 respectively. Results: There is significant increase in the percentage of inhibition against acetylcholinesterase and BACE-1 when tested with increased concentration of EECF. Compared to inhibitory effects by EECF, donepezil exhibited higher values of IC50 against both the assayed key enzymes, though also demonstrated vital inhibitory actions in both assays. EECF also showed higher potency in controlling AD progression in terms of inhibition against acetylcholinesterase, shown by IC50 value for the acetylcholinesterase, if compared to IC50 value for BACE-1. Conclusion: Green chilli or EECF has potential to have inhibitory action on acetylcholinesterase and beta-secretase enzymes in patients with AD, but limited to optimum concentration or dose. Key words: Alzheimer’s disease, Green chilli, Betasecretase, Acetylcholinesterase, Capsaicin.

INTRODUCTION
A very common vegetable cum spice commodity that usually captures customers’ attention in the agricultural market, is chilli. Chilli belongs to the family of Solanaceae (nightshades or potato family) and is classified under the genus Capsicum L.[1] Probing further, the genus Capsicum is derived from the Greek word ‘Kapsimo’ which means ‘to bite’. Originating primarily from South and Central America, the demand for chilli is everlasting across the globe due to their attractive appearance, ability to maintain a constant sharp taste and the ease in methods employed in cultivation.[2] Records have shown the existence of more than 200 names commonly used to identify chilies. Some of the typical terms include chilli pepper, paprika (sweet varieties), christmas pepper (ornamental), bell pepper (hot varieties) and tabasco pepper.[1] Moreover, the genus Capsicum consists of approximately 22 wild species in which only five are referred as the domesticated species namely C. frutescens, C. chinense, C. baccatum, C. annum and C. pubescens.[1]

HISTORY
Chilli peppers are thought to have originated in the northern Amazon basin and so, by natural geographic spread, are indigenous throughout Central America, South America, the West Indies and the most southerly states of the U.S.A.[3] The Tepep or Chiltepin pepper (Capsicum annum var glabriusculum) is reputed to be the oldest variety in the world and is commonly called the “Mother Chilli”. The name chili was derived from the ancient Aztec dialect of Nahuatl which called the pepper “chilti”.[4] It grows wild in northern Mexico and up into Arizona and Texas where it is now the State Chilli.[5] It is particularly hard to domesticate but in the wild it grows best in seemingly impossible harsh habitats. In areas of extremely low rainfall, such as the Sonoran Desert, it can be found thriving in the partial shade provided by a Desert Oak or Mesquite. In these conditions, this supposedly annual plant has been known not only to survive but also to fruit for up to 20 years.[1,3] This is an interesting feature of most chilies, if they are in conditions they like,
they will not only thrive for several years, they will also be more prolific fruiters in their record, for the third and fourth years. Some 5000-6000 year ago, evidence has also been found for chilli cultivation in Ecuador from around the same period. Later the Aztecs were famous for their love of chilli and it featured heavily in their diet.[9] The ancient Inca and Aztec civilization used and worshipped the chilli. The Aztec used chilli literally in every meal and created the first chilli. The Incas worshipped the chilli as one of four mythical brothers who feature in the Incan creation.[10,11]

Chilli came into light globally when Christopher Columbus who discovered America in 1493 mistakenly thought chilli as a ‘black pepper’ and started the trading. The story began when Christopher Columbus encountered chillies on his first voyage to the Caribbean in 1492 and though he did not bring any back on that voyage [they were taken to Spain on his second voyage in 1495], he does write of “a ‘pepper’ that the natives called ‘Aji’ which was better is taste and nature than ordinary peppers”. [‘Aji’ is a Caribbean word for chillies] Wanting to prove that he had opened a new easyer route to the Indus, Columbus was keen to associate Aji with Asian “pepper”. [5]

Diego Alvarez Chanca, a physician on Columbus’s second voyage to the West Indies in 1493 brought the first chillies to Spain and first wrote about their medicinal qualities in 1494. From Europe chillies spread rapidly to India, China and Japan. In Europe they first were grown in the monastery gardens of Spain and Portugal as botanical curiosities but the monks experimented with their culinary potential and discovered that their pungency offered an inexpensive substitute for black pepper corns, which were so costly that they were used as legal currency in some countries. Within 50 years of its discovery the humble chillies pepper had spread across most of the then known world.[12,13]

From Spain, chilli spread to South Asian countries like India and China which spread further to Japan whereby soon chilli was accepted as an important source of spice in the local cuisines. At the initial stages, it was used only for ornamental purposes as many exotic plants then unknown but very quickly became a common food. It is assumed that the monks of the Royal Monastery of Santa Maria de Guadalua, in Estremadura, Spain were the first European to discover the taste of hot peppers and add them to their kitchen.[14] Originally cultivated in monasteries, seeds were then spread throughout Spain and Europe by traveller monks. [9] It was in 1600 that through trade routes from South America. Portuguese and Spanish traders introduced the Chilli pepper in Africa, India in Asia.[15,16]

The spread of Chilli peppers to Asia was most likely a natural consequence of its introduction to Portuguese traders, who aware of its trade value would have likely promoted its commerce in the Asian spice trade routes then dominated by Portuguese and Arab traders. Today Chillies are an integral part of South Asian and South East Asian cuisine.[17]

Chillies or hot peppers are small fruit-vegetable that commonly used as spice and flavour in multiethnic Malaysian cuisine. Capsicum annuum is the botanical name of chilli under the family of Solanaceae. In fact, capsicums appear to be of American origin and were introduced into India at a very early date, possibly by the Portuguese. At different ripening stages, fruits of Capsicum annuum exhibit varying colour, i.e., unripe gives green, mature gives yellow and over-ripe gives red. Asian Chillies, for e.g., Bird’s eye chilli, are oblong-conical in shape and the membranous dissepiment, where seeds are attached, separates fruit into two cells. The characteristic hot and pungent taste of chilli is attributed to the main active component – capsaicin (trans-8-methyl-N-vanillyl-non-6-enamide). Capsaicin is the pungent oleo-resin of the fruits of chilli, but its content is varying with varieties and stages of maturity. In 1876, Thresh extracted the compound with petroleum and treated the extract with aqueous alkali. Crystals of pungent compound, capsaicin was precipitated out by passing carbon dioxide through the alkaline liquid. Therefore, this implied that capsaicin is a phenolic compound.[6] Pure capsaicin when extracted out is water-insoluble, colourless, odourless and crystalline-to-waxy solid at room temperature.[6] Capsaicin, when given internally, act on vanilloid receptors, viz. vanilloid type 1 transient receptor potential receptor (TRPV1). This binding induces inflammation reaction and irritation, so it gives hotness and burning when ingested or applied topically. Meanwhile, it desensitizes sensory nerve endings to pain stimulation by depleting the neuropeptide, substance P from local C-type nerve fibres, thus making it a local analgesic for osteoarthritis, painful diabetic neuropathy and post-herpetic neuralgia.[7,8] In early days, capsaicin has already been used for cough and cold because its receptor was found as a sensor of airway irritation and initiator of the cough reflex. Due to its hotness characteristics, some creams or ointment containing capsaicin are intended to give counter-irritant effect, for e.g. for the relief of rheumatism, when applied externally, but it may produce severe irritation.[9]

Chilli pepper is well-known for their high Vitamin C contents (0.1-0.5%), thus having strong antioxidant capability. Red carotenoids such as capsanthin and capsorubin, thiamine and fixed oil (4-16%) are also present in C. annuum. The other capsaicinoids, for e.g. dihydrocapsaicin (22%), nort dihydrocapsaicin (7%), homocapsaicin (1%) and homoudihydrocapsaicin (1%) also account for the pungency and heat produced by chilli. They are esters of vanillyl amine with C8-C13 fatty acids.[10] Water-soluble constituents had been isolated in 1990 by Izumitani from C. annuum named capsianosides.

At over-mature stage, the brilliant red colour of C. annuum is due to capsanthin, a tetramerpenes compound, so chilli is sometimes used as colouring agent. Many things can happen as a double-edged sword, and chilli is not an exception. Contradictory data on the mutagenic and anticarcinogenic activities of capsaicin had been obtained from many studies, as a result, there is still a lot of uncertainty in whether the use of capsaicin is totally safe.[11] Eating too much chilli can also lead to constipation, heartburn, and even gastric cancer.[12,13] Chilli pepper had been studied on its insulitropic action which mainly through improving insulin resistance, but controversy on defective insulin secretion caused by capsaicin arised recently.[13,14]

A short flower stalk in an inflorescence or cluster of flowers. Pedicles hold individual flowers in place, but how they do that depends on the form of the inflorescence. The function of pedicels is to expose flowers to the sun and wind and put them in a position so their aroma and colour attracts pollinating insects more easily. An inflorescence grows from a main stem, also called a peduncle. The term “stem axis” is sometimes used instead of peduncle.[14,15]

**CHEMICAL CONSTITUENTS OF GREEN CHILLI**

Chillies are reported to contain an oleoresin (16%), ascorbic acid, protein (16% dry wt.), fat (6% mainly in seeds), carbohydrates (32%), carotenoids, volatile oil and Vitamin E. The oleoresin extracted by solvent yields capsaicinoids and some carotenoids.[16,17] The most important component of capsaicinoids is phenylamido capsaicin that counts for the pungency of the chilli. Other components include dehydrocapsaicin, nort dihydrocapsaicin, vanilly amides of caprylic, nonylc and decyclic acids, capsiamide and homocapsaicin.[18,19] Any variation in the chemical structures of the capsaicinoids, including the structure of the acyl moiety, affects the degree and the level of the pungency. The presence following secondary metabolites are useful in differentiating the red and green peppers.[17]

- Red chilli constituents: Capsanthin and capsorubin (carotenoids responsible for the red colour), β-carotene
• Green chilli constituents: Lutein, neoxanthin, violaxanthin, β-carotene, polyphenols
The composition of capsaicin and its other associated alkaloids in dried chilli peppers is much higher than the fresh picked ones due to the present water composition as well as water-soluble phytochemicals is minimized to little or none by evaporation. As compared to red chilli peppers, green chilli peppers are more relatively healthy due to higher water content. Owing to their lower capsaicin content, the green chilies are less likely to induce internal inflammation and indigestion instead of evoking peptic ulcer, if compared to red chilli consumption.[20,21]

MAJOR AILMENT OF CENTRAL NERVOUS SYSTEM- DEMENTIA
Our human nervous system consists of a compilation of the two systems, including peripheral nervous system and central nervous system. Normally, the central nervous system is linked to and coordinated with the peripheral nervous system consisting of numerous intertwining nerves. The brain is one of the key organs in our nervous system functioning as the main regulator of our memory, thoughts, feeling, and coordination of limb movements, besides the spinal cord. Once we are born to living, our human brain evolves from the tip from the longneural tube and the neural groove will bind at three to four weeks after conception process. [20] The basic units forming our brain are the neurons. There are around eighty-six billion neurons found in the human brain always assisting in transporting the messages to the main brain via energy trans-conversion of the electrical signals possessed by the stimuli.[21] Therefore, the complexity of the physiological functions and regulations within the nervous system brings up to millions of human beings with unpredictable thoughts and ideas. Recently, mental health is set as one of the global health priorities. In this case, several psychological and mental defects have been always taken into consideration of conducting clinical studies to explore further.

As the population age progresses, the normal activity of nervous system among the elderly would come to trends of declination as well as other metabolic functions of body systems. Due to determinant influences from the external environment and practice of the lifestyle surrounding the modern technology, health concerns especially regarding mental are recently not gaining attention from the public. However, the basic components of the nervous system will undergo degeneration and numerous problems will be aroused as a result of the destruction of the nervous system. One of the nervous ailments includes dementia. In fact, dementia is defined as constellation of symptoms or medical conditions which slows down the normal motor functions of the nervous system. The motor functions of central nervous system include the reduced in memory accumulation, thinking abilities, language production and stabilization of the microtubules. As tau is detached from the microtubule, increasing quantities of separately undissolved tau eventually formulate lots of tau aggregates.[22-24] If tau clearance is not successfully proceeded, this disintegrated tau aggregation will become insoluble and develop the neuronal pretangles accumulating into neurofibrillary tangles (NFT) by the phosphorylation expressed by one of the glycogen synthase kinase enzymes, GSK-3.[24] The presence of several NFTs induces neuronal toxicity in terms of interference with normal neuronal functions, impairment of motor and cognitive functions, and also worsening neuronal degeneration.[25] Hence, the recent studies had been conducted to formulate associated solutions against those two major pathologies mentioned above in case of all types of AD case events.

RELATIONSHIP OF ALZHEIMER’S DISEASE WITH PRESENT STUDY
The degeneration or functional loss of the body systems is one of the hallmarks determining the aging process due to the oxidative stress among the elderly populations. Therefore, the rapid destruction of the brain cells would also be one of the signals which resemble the oxidative stress occurring within the nervous system. Several studies had discovered that this AD is more associated with the diabetic risk and its hyperglycemic complications. From the previous studies, the capsaicin and its analogues were finally researched their hypoglycemic actions by stimulating the insulin secretion from the pancreas in the diabetic patients. However, the pharmacological actions of capsaicin or capsicum
from green chilli in AD has no relevant clinical data discovered out yet. Before the other chemicals had been identified out from the green chilli pepper sample extract which can deal with the accumulation of beta-amylloid and also lysis of ACh molecules, this can only merely be hypothesized that the composition of antioxidants like beta-carotene and ascorbic acid (Vit. C) would help combat the oxidative stress inducing AD, and anticholinesterase potency possessed by capsaicin.

MATERIALS AND METHODS

Materials

Fresh green chilli pepper (Capsicum annuum L. varfrutescens) was purchased from the local market, cotton wool, ethanol (Absolute alcohol), aluminium foil.

Apparatus

Measuring cylinder, knife, electric grinder, beakers, glass rod, weighing balance, retort stands, Soxhlet Apparatus, rotary evaporator, heating mantle, 500ml round-bottomed flask, water pipe, water bath (Tuff TEB-26), china dish, conical flask.

Methods

Preparation of green chilli powder

Fresh green chilli pepper (Capsicum annuum L. varfrutescens) was purchased from the local market. Its calyx and stalk were removed, leaving the fruit part to be cut into small pieces and dried with a soft tissue before drying under the shade for one to two weeks. The dried green chilli was finely ground into powder by using an electric grinder and preserved in an air-tight container at room temperature until the preparation of experimental diets.

 Extraction of capsaicin

Soxhlet extraction

Approximately 25g of powdered green chilli was weighed. A Soxhlet extraction tube was assembled into an apparatus with a condenser and round-bottomed boiling flask. Instead of using porous thimble, a small mass of cotton wool was placed at the bottom and top of the tube to sandwich the powdered bed added in between. Around 300ml of ethanol, along with porcelain chips was taken into the round-bottomed flask. The apparatus was placed on the heating mantle with retort stand and clamp. The cold tap water was circulated through the condenser. The solvent was made to boil to allow the siphoning cycle to take place. Therefore, extraction of capsaicin with ethanol at 70°C ± 5°C was completed within three days when the extracting solvent became clear.

Maceration

Another method for extraction of capsaicin that had been attempted was maceration to achieve larger quantity of extraction. 50g of powdered green chilli was weighed and placed in conical flask. Around 350ml of ethanol was mixed with the chilli powder. The flask was then stoppered or covered with aluminium foil to reduce the loss of solvents by evaporation. Frequent agitation was done by using the mechanical stirrer to allow homogenization of the final solution and saturation of the solvent. The whole solvent was allowed to stand at room temperature for five days with frequent agitation to allow penetration of extracting solvent into the cellular structure to dissolve the soluble compounds. After that period, the solvent was renewed again and left for another three days until the plant material was exhausted. The liquid was strained and marc was pressed. The liquid collected from first and second batch of maceration was filtered to obtain a clear extract.

 Rotary evaporation

After extraction, the solvent was eliminated by rotary evaporation under reduced pressure at temperature of around 65°C ± 3°C with approximately 90 rpm. The concentrated extract was obtained after few hours.

 Water bath

In order to safely achieve maximum concentration of the extract, water bath was used by adding the concentrated extract into the china dishes which were placed on the water bath at temperature of 45°C. After few days, the gummy residue from the extract was weighed and stored in the refrigerator to be used as experimental diets.

IN-VITRO ACETYLCHOLINESTERASE (ACHE) EVALUATION

The assay for measuring AChE activity was evaluated based on Ellman’s method. The principle of this method is based on enzyme hydrolyzes the substrate acetylthiocholine iodide (ATCI) into thiocholine and acetic acid. Then, thiocholine can react with 5,5-dithio-bis-(2-nitrobenzoic acid) (DTNB) and formation of yellow color. The intensity of the color formation is directly proportional to the activity of the enzyme. Briefly, 150 μl of 0.1 M sodium phosphate buffer (pH 8.0), 10 μl of the test compound, and 20 μl of the enzyme solution (0.1 units/mL) were added and incubated for 15 min at 25°C. After that, with mixed with 10 μl of DTNB (10 mM) and 10 μl of ATCI (14 mM) to initiate the reaction, incubate for 10 min. The color formation was measured at 410 nm wavelength. The controls contained the solvent to dissolve the compound instead of test compound.

The percentage inhibition for each test solution was then calculated using the following equation:

\[ \text{AChE Inhibition} = \left(1 - \frac{\text{Absorbance sample}}{\text{Absorbance control}}\right) \times 100 \]

IN-VITRO BETA-SECRETASE (BACE-1) ASSAY

Screening β-secretase inhibitors using purified enzyme

Compounds at concentration ranging from 0.1-100 μg/mL were assayed for BACE-1 inhibition using a fluorescence resonance energy transfer (FRET) assay. This assay uses baculovirus-expressed BACE-1 and peptide substrate based on Swedish mutation of APP. The mutation dramatically enhances the cleavage of APP by BACE-1 enzyme and this peptidic substrate becomes highly fluorescent upon enzymatic cleavage. A mixture of 10μl of test compound diluted in assay buffer, 10μl of BACE-1 substrate (Rh-EVNILDAEFK-quencher, in 50 nM ammonium bicarbonate) and 10μl of BACE-1 enzyme (1.0 U/mL) were incubated for 60 min at room temperature in a dark condition. After that, 10μl of BACE-1 stop buffer (2.5M sodium acetate) was added to the mixture. The fluorescence was read by using a spectrophotometer (TECAN) under the excitation at 545 nm and emission at 585 nm. The percentage of BACE-1 inhibition was measured using the equation below:

\[ \text{BACE-1 inhibition} = 1 - \frac{\text{Test Sample}}{\text{Positive Control}} \times 100 \]

STATISTICAL ANALYSIS

Data was presented as mean ± standard deviation (SD). All analyses were carried out intriplicates. GraphPad Prism 5 and Microsoft Excel 2007 were applied for the statistical and graphical evaluations.
RESULTS

In-vitro Acetylcholinesterase (AChE) Assay

The assay of acetylcholinesterase was based on an improved Ellman method in a 96 well plate reader using QuantiChrom assay kit (USA). One of the characteristic changes that occur in AD is increase in acetylcholinesterase (AChE) activity, the enzyme responsible for acetylcholine hydrolysis, from both cholinergic and non-cholinergic neurons of the brain. The results obtained from the ethanolic extract of Capsicum frutescens (EECF) against AChE enzyme inhibition activity and the percentage inhibition was evaluated and tabulated in Tables 1 and 2 respectively. IC50 value of EECF was found to be 0.056±0.030 (µM/mL). However, these compounds are shown to be highly effective as compared with control AChE inhibitor donepezil (0.096±0.030 µM/mL).

In-vitro Beta-secretase (BACE-1) Assay

The assay of β-secretase was based on fluorescence resonance energy transfer (FRET) method using Swedish mutation of amyloid precursor protein (APP). One of the characteristic changes that occur in AD is increase in β-secretase enzyme (BACE-1) activity, the enzyme responsible for β-Amloid (Aβ) formed by the continuously proteolytic processing of β-amyloid precursor protein (APP) by β- secretase and γ-secretase, plays a vital part in the pathogenesis of AD. The results obtained from the ethanolic extract of GCE against BACE-1 enzyme inhibition activity and the percentage inhibition was evaluated and tabulated in Tables 3 and 4. The IC50 value of EECF was found to be 0.074±0.006 mg/mL. However, these compounds are shown to be highly effective than control AChE inhibitor donepezil (0.096±0.030µM/mL).

Table 1: The AChE inhibition of ethanolic extract of CF and control.

<table>
<thead>
<tr>
<th>S. No</th>
<th>Concentration (mg/mL) of EECF</th>
<th>Percentage of inhibition (%)</th>
<th>Concentration (µM/mL) of Donepezil</th>
<th>Percentage of inhibition (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>0.25</td>
<td>25.75±0.18</td>
<td>0.1</td>
<td>68.38±0.14</td>
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<tr>
<td>2</td>
<td>0.5</td>
<td>61.44±0.33</td>
<td>1</td>
<td>91.29±0.18</td>
</tr>
<tr>
<td>3</td>
<td>1</td>
<td>66.46±0.22</td>
<td>10</td>
<td>94.09±0.19</td>
</tr>
<tr>
<td>4</td>
<td>2</td>
<td>76.30±0.30</td>
<td>100</td>
<td>96.13±0.17</td>
</tr>
<tr>
<td>IC50</td>
<td></td>
<td>0.056±0.005</td>
<td>IC50 (µM/mL)</td>
<td>0.096±0.030</td>
</tr>
</tbody>
</table>

*All the values are shown as mean ± SD (n = 3)*

Table 2: IC50 value of EECF and control (AChE inhibition).

<table>
<thead>
<tr>
<th>S. No</th>
<th>Name of the compound</th>
<th>IC50 (mg/mL)</th>
<th>IC50 (µM/mL)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>EECF</td>
<td>0.056±0.005</td>
<td>(mg/mL)</td>
</tr>
<tr>
<td>2</td>
<td>Donepezil</td>
<td>0.096±0.030</td>
<td>(µM/mL)</td>
</tr>
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</table>

Table 3: The BACE-1 inhibition of ethanolic extract of GC compared with control.

<table>
<thead>
<tr>
<th>S. No</th>
<th>Concentration (mg/mL) of EECF</th>
<th>Percentage of inhibition (%)</th>
<th>Concentration (µM/mL) of Donepezil</th>
<th>Percentage of inhibition (%)</th>
</tr>
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<tbody>
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<td>68.38±0.14</td>
</tr>
<tr>
<td>2</td>
<td>0.5</td>
<td>52.37±0.09</td>
<td>1</td>
<td>91.29±0.18</td>
</tr>
<tr>
<td>3</td>
<td>1</td>
<td>62.23±0.12</td>
<td>10</td>
<td>94.09±0.19</td>
</tr>
<tr>
<td>4</td>
<td>2</td>
<td>69.67±0.05</td>
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<td>96.13±0.17</td>
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<tr>
<td>IC50</td>
<td></td>
<td>0.074±0.006</td>
<td>IC50 (µM/mL)</td>
<td>0.096±0.030</td>
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</table>

*All the values are shown as mean ± SD (n = 3)*

DISCUSSION

Since Alzheimer’s disease (AD) is known to be listed in one of the ten global death-leading causes, the level of study workload related to this neurodegenerative disease is continuously increasing. It is merely to track the alternative way out to manage and alleviate the major dementia symptom of AD. Basically, the studies were aimed towards the two main pathological factors, including the APP processing and cholinesterase enzymes.[24] Undeniably, tau phosphorylation is also linked to the pathophysiological properties of this dementia ailment, which could occur as well with the APP processing. Previously, there were several studies regarding risk of AD and its correlation with the capsaicin administration in *in-vivo* manner.[30,36] However, the established data was merely limited to the utilization of red chilli peppers. Hence, the present study is designated for only the green chilli to estimate the inhibition potency of green chilliethanolic extracts against the AD risk in terms of enzymatic inhibition against acetylcholinesterase (AChE) and BACE-1, as one of the β-secretase enzymes.[27] According to the result data regarding the inhibition activity of EECF against AChE and BACE-1 collected throughout the present study *in-vitro* method, EECF is indicated to establish higher significance in lowering the occurrence of AD by terms of presenting higher percentage of inhibition against both the AChE and BACE-1, as compared with the data of using conventional medication for such study design.

Since the study was overall conducted in *in-vitro* manner, the enzyme assay kits which are used in the way of having well-plate were utilized, including AChE and BACE-1 assay kits. For the assay of AChE inhibition by capsaicin, one of the AChE assay kit products, i.e. QuantiChromTM assay kit was applied to determine and analyze the inhibition of AChE in quantitative manner using standard and sample analytes. BioAssay Systems QuantiChrom™ Acetylcholinesterase Assay is based on an improved Ellman’s method, which is limited for *in-vitro* studies. This method applies the principle based on the hydrolysis of thiocholine-derivative compounds, including ACh and butyrylcholine (BCh), by the enzyme cholinesterases (ChEs).[28,30] The final development of yellow colour is due to the production of the final product, i.e. 5-thio-2-nitrobenzoate, and the functionality of the ChEs could be indicated according to the intensity of yellow colour observed from the well-plate of the assay kit experimental setup.[30] The yellow colour exhibited by the final product from the reaction of thiocholine substrate and DTNB compound is due to the shifting of certain electrons from the ester linkage to one side of the nitrobenzoic acid within the DTNB while reacting to thiocholine substrate.[30] For the part of another estimation study of BACE-1 inhibition using capsaicin, the BACE-1 Fluorescence Resonance Energy Transfer (FRET) assay kit was utilized for this study experimental *in-vitro* setup.[30,36] It applies the principle of resonance energy transfer between the two-sensitizing light components; hence the darker environment is required to observe the changes. In both assay tests, the condition of assay medium should also be adjusted pH to its respective optimum acidity and basicity as the activity of AChE and BACE-1 is also dependent on the changes in the pH of the surrounding medium.[37,38] The evaluation of anti-Alzheimer properties through AChE and BACE-1 inhibition is also applied in one previous study regarding the baicalein.[38]
Compared to other anti-Alzheimer drugs including galantamine, donepezil is the medication present in the market available in the lowest price.[36] However, as for the medication used to manage the AD symptoms by opposing the BACE-1 functions, it seems to have limited market of this medication group type for AD patients. Due to benefits for all stages of AD (i.e. mild, moderate, and severe), donepezil is merely the selected available medicated application as the control purpose for the enzymatic inhibition of AChE as well as BACE-1, although the inhibition against any amylloid protein cleavage enzyme is not induced. As one of the AChE inhibitor medications, donepezil has been used for management or one of the drug regimens to control the progression of AD risks in either early or late onset. Generally, donepezil works its therapeutic functions by binding competitively to the cleavage site of AChE with acetylcholine (ACh) transmitters, being active substrate of AChE.[40] The unbound AChE within and outside the neurons degrades the intracellular and extracellular ACh which aids in memory function into separated choline and acetate molecules. By donepezil antagonizing actions, concentration of those ungraded ACh transmitters is under maintenance without any diminution or desensitization prior to undergo cholinergic transmission at the nerve terminals. For the control set (donepezil), the results obtained from the assay of its AChE and BACE-1 inhibition activity shows that the quantity of the inhibited AChE and BACE-1 exhibits a gradual increment with the consecutive increase in the concentration of the introduced donepezil.

As for the results attained from the present assay regarding the AChE inhibition activity, it could be assumed that the capsaicin from the EECG is effective in reducing the functionalities of AChE in-vitro, in terms of linear increase in the inhibition quantity of AChE with the consecutive increase in the EECG concentration. In case, the potent inhibitory action of capsaicin against AChE may be implicated from the results obtained. Capsaicin had been discussed in previous studies regarding its anti-inflammatory and pain soothing effects upon in-vitro local site application. This is related to its affirmed binding affinity to the TRPV1 receptor, being the typical capsaicin receptor, which channels the main afferent neurons that are responsible in dealing with the pain and heat stimuli from the external surrounding. Hence, it has exhibited its specific analgesic effect that it has been applied on the manufacture of the anti-inflammatory cream and ointment utilized for minimizing of pain at the local sites. In this current study, the mechanism of opposition against AChE activity was discovered to be similar as the pain mechanism mentioned above.[12,41,42]

Metabolism is known to be the essential process for all the chemicals and toxins to undergo after administration into the living system through various routes, including capsaicin itself. However, due to the complexities of its metabolism, so it is complicated to explore the mechanism of capsaicin after introduced into the system. In one previous study, Brock C. had investigated into and explained the metabolic pathway of capsaicin via various routes of administration, including in-vivo and in-vitro routes.[10] Prior to the explanation regarding the routes of capsaicin, its physical and chemical properties were revisited to understand clearly its mechanism of pharmacokinetic routes. As discussed above, capsaicin is chemically named as trans-8-methyl-N-vanillyl-6-nonenamide, and also grouped collectively as capsaicinoids as discussed above, capsaicin is chemically named as trans-8-methyl-N-vanillyl-6-nonenamide, and also grouped collectively as capsaicinoids as well with nordihydrocapsaicin compound. This active component is also possessed by most of the Capsicum sp., i.e. chili peppers, except sweet peppers or bell peppers. As the capsaicin proceeds unchanged beyond the gastric system, it would undergo biotransformation into its specific second metabolites. One of the second metabolites of capsaicin after being bio-transformed includes vanillin present in lesser quantity.[10] The other associated major metabolites of capsaicin include 16-hydroxycapsaicin, 17-hydroxycapsaicin, and 16, 17-dihydroxycapsaicin. In context to the study, it is suggested that merely little portion of capsaicin is found to be metabolized into two metabolites, i.e. vanillylamine and vanilinic acid.[43] Vanillylamine is produced as the metabolite of vanillin after enzymatic process of vanillin transferase, whereas vanillin acid being one of the plant metabolites, has also exhibited its antagonistic properties on the anionic site of AChE. Therefore, due to their presence, capsaicin within the chilli peppers is also known as one of the vanilloid compounds, which scores the highest SHU value based on the Scoville value for grading of spiciness.

ACh, as a major small-molecule transmitter in the cholinergic system, is mainly involved in the signaling of muscular contraction. Within most of the body systems, ACh transmitters are mainly involved in normal brain and muscle functions, in terms of memory and learning abilities and the control of muscle movements and limb coordination. Moreover, the presence of ACh also takes part in most of the metabolic functions including the activation of pain sensory receptors, vasodilation on the increase in temperature stimuli, regulation of body secretion including endocrine system.[43] Within cardiovascular and muscular systems, it also possesses its inhibitory actions at certain smooth muscles and at cardiac muscle. The excessive quantity of this cholinergic transmitters would likely to induce specific toxicities at the muscarinic and nicotinic receptors. In contrast, deficiency in ACh may be linked to prominent decline in cognitive function and also further impairment of memory function in elderly population. However, this hypothesis seems not yet to attain certain verification with strong evidence due to occurrence of the nervous symptoms related to insufficiency in ACh levels in certain brain areas has not been concluded yet and the phenomenon is also hardly determined.

Due to the lipophilic nature and its lower molecular weight, vanillin has greater potential to penetrate across the blood-brain barrier without difficulties. This has also been emphasized by one of the studies by Salau VF, et al.[14] Therefore, it could have diffused freely within the cholinergic neurons in the certain brain areas. In case, vanillin, which has been bio-transformed from capsaicin molecule itself, is suggested to possess anti-Alzheimer property in the way of AChE inhibition. One of the studies had discovered that one of the vanillin compounds exhibited greater potential in the inhibitory action against AChE in-vitro as compared to the cinnamaldehyde compounds besides having higher antioxidant properties during the analysis of azomethines derived from these two components.[44,45] On top of that, another causative factor contributing to AD is suggested to be the higher level of oxidative stress in the brain areas besides AChE working in the disease progression. Interestingly, certain phenolic compounds, including vanillin compound, are proclaimed to have exhibited their antioxidant potentials in AD. It is clearly demonstrated that vanillin derived from capsaicin has possessed affirmed antioxidant property besides exhibiting its neuroprotective effect in its role of inhibiting AChE in the control or management of AD risk progression.[46-48] For the part lacking, that is the dose-effect relationship of usage of vanillin in clinical trials, it is suggested to conduct related human studies regarding this arose issue.

It is known that the AD occurrence is also highly associated with the genetic factors, besides the aging factor, the practice of lifestyle habits which are mostly health-affecting and the previous acquired vascular diseases for especially the elderly population. One of the well-known hallmarks for this evoking of AD including the formation of senile plaques of Aβ, by the two enzymes including β- and γ-secretases.[23,49] In case, the presence and role of BACE-1 is the prominent risk factor in the occurrence of AD. From the comparison between potencies of BACE-1 inhibition by the EECG and donepezil control sets in the present study, EECF is interpreted to exhibit that its greater potency in declining BACE-1 functionalities which is represented by increase in the
percentage of BACE-1 inhibition correlated with the consecutive increase in the EECF concentration. Based on the results above, capsainc has also been suggested to have possible antagonizing effects in the accumulation of Aβ plaque production within the neurons, besides having possible its associated anti-dementia benefits in the way of inhibiting AChE.

In fact, Aβ peptide is one of the neuropeptides which has been emphasized in the role of boosting memory functions in normal healthy person, in the normal concentrations for the maintenance of physiological processes within the nervous system. These small-sized peptides are actually originated from the large molecular transmembrane integral protein surrounding the present neurons, known as the amyloid-precursor protein (APP) playing its role in memory and neurocognitive functions.\[90\] The presence of this peptide is normally concentrated at the nerve terminals as these APP molecules are mainly discovered within the synapses between two nerve cells present in the nervous system. As the concentration of the intracellular Aβ peptides increases uncontrollably through abnormal reproduction, this would lead to development of Aβ senile plaque due to insoluble characteristic of beta-amyloid peptides.\[23,49\] Their insoluble traits further cause more clefted Aβ peptides primarily by β-secretase become further saturated and eventually accumulated into irreversible plaques evoking undesired events in the nervous system of defects. Progressive accumulation of these Aβ plaques without monitoring will cause undeniable neuronal death due to loss of synaptic cleft functionalities. This process by BACE-1 was emphasized in the related theory known as “amyloid cascade hypothesis”.\[24\]

Similar to capsainc, this integral membrane protein also undergoes its vital metabolism or processing prior to its working of actual functions in the central nervous region. APP processing could take place in two opposite pathways, i.e. amyloidogenic pathway and non-amyloidogenic pathway.\[49\] Both the pathways will eventually result in formation of numerous soluble peptides which aid in normal neuronal functions in the brain region. Amyloidogenic pathway emphasizes the metabolism of APP by the functional working of beta-secretase (or mainly BACE-1), eventually result in production of soluble APPβ peptides.\[49,90\]

Compared to amyloidogenic pathway, non-amyloidogenic pathway occurs as the way of APP proteolysis by alpha-secretase enzyme into soluble APPsa peptides possessing neuroprotective properties.\[51\] As the age progresses, the frequency of non-amyloidogenic pathway becomes diminished due to the decline in functionalities of α-secretase enzyme in producing its complementary peptides, leading to enhance in the amyloidogenic pathway expression.\[51\] As prior to dementia problem in elderly, the excessive expression of amyloidogenic pathway would lead to neurotoxicity due to formation of senile amyloid plaques via aggregation and further oligomerization of beta-amyloid peptides.

The assumption of direct inhibition of BACE-1 by capsainc molecules itself attained limited verification with previous studies' evidence. There was also limited data regarding the molecular affinity of capsainc to the anionic property of aspartic residue within the structure of BACE-1. Interestingly, while the exact mechanism of capsainc upon AD progression is still unknown, one previous study suggested that this pungent capsainc works its inhibitory action upon BACE-1 out via the non-amyloidogenic processing of APP proteolysis which serves to reduce the APP-amyloidogenic processing and decrease the concentrations of Aβ peptides.\[52\] Within the non-amyloidogenic proteolysis of APP, one of the key alpha-secretases responsible in this process includes ADAM10.\[51\] In case, ADAM10 can work out its exact proteolytic functions only after it is matured or being cleaved from its prodomain part. Undeniably, prior to the AD pathogenesis, the quantity of ADAM10 declines and the amyloidogenic processing progresses without homeostatic modulation by the non-amyloidogenic pathway, and the production of Aβ is eventually beyond the Aa peptides. From the previous associated study, capsainc was suggested to have enhanced the rate of ADAM10 maturation prior to precluding of Aβ generation.\[30-52\]

Through upregulation of the peroxisome proliferator-activated receptor alpha (PPARα), capsainc administration could stimulate the expression of ADAM10 to produce more soluble alpha peptide molecules reducing AD risk.\[51\] However, one of the available report data showed opposition against the above statement, indicating amyloidogenic pathway could be enhanced by the capsainc.\[53,54\]

On top of that, our current study also suggested that precluding of Aβ production through BACE-1 inhibition is also linked to the activation of TRPV1 receptor, though the correlation between TRVP1 receptor stimulation and the changes in the BACE activity was opposed by one study.\[55,56\] Besides this pathological process of the amyloidogenic APP processing, the typical dementia and confusion symptoms of AD may also be induced by the uncontrolled boost in blood glucose levels. One of the medical studies reported that dementia problem or AD risk is also one of the complications lead by the progression of chronic or untreated type 2 diabetes mellitus (T2DM).\[57\] Several existed studies suggested the above correlation between T2DM and AD or its dementia risk.\[35,58\] The suggestion is based on the theory of potential destruction occurred with high value of blood glucose levels. The previous researchers also assumed that vascular dementia or destructive brain alterations are secondary to the possible vascular blockade in brain region by accumulation of glucose within the body system.\[57\]

However, research work is recently conducted based on the suggested theory of the linkage between these two parameters to clarify this above relationship clearly due to the complexity of the mechanism behind this glucose dyshomeostasis in inducing cognitive loss phenomenon. Besides, one previous study related to T2DM supported our assumption by emphasizing that insulin secretion could be improved via the activation TRVP1 expressed on CGRP-containing fibers, followed by the desensitization by capsainc introduction. The depletion in the concentration of calcitonin gene-related peptide (CGRP) is verified in the improvement in the insulin secretion.\[59\] In another in-vivo study, the inhibition of this vanilloid receptor which is Ca²⁺-ion-permeable, is also considered to be associated with decrease in amyloid-beta concentrations as well as in tau generation, as the excessive accumulation in the Ca²⁺ ion is conducted via permeation through TRVP1 receptors and other existed Ca²⁺ channels could lead to neuronal death as one of the hallmarks of AD pathophysiological phenomena.\[54\] However, another study demonstrated its controversial data findings, indicating that the destruction to the gamma-oscillations in the hippocampus region could be diminished by capsainc treatment via TRVP1 activation.\[41\] In case, there are verification limitations of the study information regarding the mechanism of TRVP1 stimulation found due to the conflict of the appeared data obtained from the previous associated reports.

Instead of giving health benefits to the spicy lovers, heavy consumption of capsainc also is discovered to induce adverse health events due to its irritating effect.\[40\] Capsainc found in most of the chilli peppers, including green chillies, is believed to cause a definite effect on normal digestion and further induce acute diarrhoea. As the TRVP1 receptor is stimulated through capsainc administration via daily diet, the burning sensation arises from inside the body and causes irritation to the gastrointestinal tract especially ileum from the small intestine.\[51\] In addition, the activation by capsainc is translated by the brain to release endorphins to cause the spicy-lovers feeling tipsy, in order to consume more peppery and spicy meals.\[62\] However, excessive consumption of capsainc or spicy food most likely to cause nausea, vomiting and abdominal pain due to stimulation of pain receptors overlying throughout the digestive tract, rather than providing the necessary antioxidant functions to
CONCLUSION

Overall, the data findings from this recent study suggest that EECF in the concentrations of 1 mg/mL and 2 mg/mL possess greater anti-Alzheimer effects, in the means of AChE and BACE-1 inhibition. However, the heavy or excessive consumption of capsaicin-based diet and the administration of high-dose capsaicin in various dosage forms is capable of inducing cognitive impairment in healthy population and even slimmer population. These adverse effects also most likely be induced towards the elderly including memory-declining AD patients. Previous, the results and findings are obtained from the studies regarding the red chilli peppers, but limited data regarding the green chillis peppers in animal and human studies could be found. Therefore, more animal studies related to the anti-Alzheimer effects of capsaicin from green chilli extracts should be carried out. Moreover, further associated work is required to unravel the mechanism of capsaicin within green chilli peppers in the declining phenomenon in the memory and cognitive functions in the elderly population, including healthy population at the age between 10 and 49.

CONFLICT OF INTEREST

The authors declare that there is no conflict of interest.

REFERENCES

SUMMARY

There is limited data related to pharmacological properties of green chilli peppers in the memory enhancing effects in AD, although many study reports regarding the effects of red chilli peppers in AD are found in either in-vivo or in-vitro manner. Due to difference in many study reports regarding the effects of red chilli peppers in AD are found in either in-vivo or in-vitro manner. Due to difference in these study reports, the present study was designed to evaluate the anti-Alzheimer effect through inhibition of AChE and BACE-1, and in order to test its potentials with red chilli peppers. The experiment setup was conducted in-vitro, using the application of AChE and BACE-1 assay kits. Besides, donepezil medication was prepared as the control setup to compare the anticholinesterase and BACE inhibition potential with the various concentrations of EECF in the in-vitro setup. As for the assay using BACE-1 kit, the experiment was carried out in a dark environment without any distraction of other light sources. The results were observed in the well-plate of both the kits and the absorbance of fluorescence in BACE-1 assay was measured by using spectrofluorometer. From the result data obtained, EECF have greater potential in associated control and management of AD, in terms of its anticholinesterase effects especially for elderly.
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GRAPHICAL ABSTRACT

EECF extract

Donepezil (Control set)

1. AChE Inhibition
2. 

EECF exhibited significant inhibitory actions towards AChE and 

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