Inhibitory Effects of Raw-Extract *Centella asiatica* (RECA) on Acetylcholinesterase, Inflammations, and Oxidative Stress Activities via In Vitro and In Vivo

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**Abstract:** *Centella asiatica* (*C. asiatica*) is one of the medicinal plants that has been reported to exert comprehensive neuroprotection in vitro and in vivo. In view of this, the present study was performed to investigate the effect of ethanolic extract of *C. asiatica*, designated as raw-extract of *C. asiatica* (RECA) in reducing the acetylcholinesterase (AChE), inflammations, and oxidative stress activities via both in vitro (SH-SY5Y and RAW 264.7 cells) and in vivo (Sprague Dawley rats). Quantitative high-performance liquid chromatography analysis reveals that RECA contains a significantly high proportion of glycosides than the aglycones with madecassoside as the highest component, followed by asiaticoside. Treatment of SH-SY5Y cells with RECA significantly reduced the AChE activity in a concentration-dependent manner with an IC$_{50}$ value of 31.09 ± 10.07 µg/mL. Furthermore, the anti-inflammatory and antioxidant effects of RECA were evaluated by lipopolysaccharides (LPS)-stimulated RAW 264.7 cells. Our results elucidated that treatment with RECA significantly suppressed the level of pro-inflammatory cytokine/mediators and oxidative stress released in a concentration-dependent manner. Interestingly, these patterns of inhibition were consistent as observed in the LPS-induced neuroinflammation Sprague Dawley rats’ model. The highest concentration used in the two models presented the most significant results. Herein, our findings strongly suggest that RECA may offer therapeutic potential for the treatment of Alzheimer’s disease through inhibiting the AChE, inflammation, and oxidative stress activities.

**Keywords:** Centella asiatica; anti-acetylcholinesterase; anti-inflammatory; antioxidant; lipopolysaccharides; in vitro; in vivo

1. Introduction

Alzheimer’s disease (AD), which is associated with progressive memory loss and impairment in cognitive and behavioral functions, is the most common neurodegenerative disease worldwide [1].
The etiopathogenesis of this disease is multifactorial and recent pieces of evidence have demonstrated that the deficits in central cholinergic neurotransmission in the brain, inflammatory injury, and induction of oxidative stress are also interrelated in the onset and progression of AD [2–4]. Suppressing the acetylcholinesterase activity in the brain is one of the most popular targets for AD treatment to ameliorate cognitive ability. This enzyme, when in excess, could further lead to depletion of the acetylcholine concentration that will eventually result in deficit neurotransmission within the synaptic region [5]. Consequently, most of the prescription drugs to manage the symptoms and halt the progression of AD are working as acetylcholinesterase inhibitors such as rivastigmine, donepezil, galantamine, and memantine [6,7].

Lipopolysaccharides (LPS), a pro-inflammatory bacterial mimetic derived from the outer membrane of gram-negative bacteria, is often used in the research field to promote neuroinflammation in vitro or in vivo. Once a model is stimulated by LPS, the levels of pro-inflammatory cytokines/mediators and oxidative stress would become significantly elevated. Prolonged elevation of this imbalanced inflammation and oxidative stress would eventually accelerate the manifestation and progression of AD [2,4,8]. Therefore, inhibition of the inflammatory response is crucial in restricting the development of this disease.

Currently, there is no highly effective medicine that can prevent, halt, or reverse the progressive course of AD [9,10]. Although there are well-established (semi-)synthetic drugs currently used for the management of AD, most of them have several adverse effects. Thus, public and scientific attention has turned toward phytopharmaceuticals as promising candidates for drug development owing to their wide therapeutic potential, superior safety, and consumer acceptability [11,12]. For instance, evodiamine (Evo), an extract of E. rutaecarpa Bentham, is widely used in traditional Chinese medicine as herbal medicine as it is believed to exhibit anti-AD, anti-inflammatory, antinociceptive, and thermoregulatory activities [13–15]. Meanwhile, a herbal medicine called ninjin’yoeito (NYT), a form of a traditional Japanese medicine (Kampo), has demonstrated both cognitive improvement and a reduction of AD-related depression to the patients that received a combination of donepezil supplemented with NYT [16].

C. asiatica (L.) Urban, or locally known as “pegaga” in Malaysia, is a psychoactive medicinal plant belonging to the Apiaceae family (formerly Umbelliferae) [17]. It is a small perennial plant that is commonly eaten raw as a salad (ulam) and used as a topical product, especially among the older generation in Malaysia [18]. Various uses are claimed for the plant, which has long been recognized by Ayurvedic and Chinese medical practitioners particularly for its rejuvenator and cognitive enhancer effects [19–21]. Moreover, it also has been shown to possess neurological, antioxidant, anti-inflammatory, and anticancer activities; yet, its mechanism of action is not clearly portrayed. The most prominent group of active compounds identified from this plant is pentacyclic triterpenoid saponins including asiaticoside, madecassoside, and their sapogenins (madecassic acid and asiatic acid), which are believed attributable to its wide spectrum of therapeutic actions [22,23]. A study done by Won et al. [24] reported that madecassic acid isolated from C. asiatica more potently suppressed the inflammatory mediators than madecassoside did via the downregulation of NF-kB activation in RAW 264.7 macrophage cells. This may point to the different affinities exhibited by madecassoside and madecassic acid for cellular membranes. Moreover, results from Park et al. [25] demonstrated that titrated C. asiatica (TECA), which comprises 29–30% madecassic acid, effectively inhibited LPS-induced inflammatory responses. However, the triterpene components in C. asiatica are not always the same due to the geographical origin, diverse environmental conditions, and different accessions [26]. The absence reports of adverse effects in any clinical studies further corroborate in the classification of this plant as a Class 1 herb (one that can safely be consumed when used appropriately) in the Botanical Safety Handbook [27].

Based on these rationales, this study is mainly devoted to finely evaluate the effect of ethanolic C. asiatica extract designated as raw-extract C. asiatica (RECA) in inhibiting the acetylcholinesterase (AChE) activity and suppressing inflammation and oxidative stress levels in vitro and in vivo. Results of this study will provide extra insights for using C. asiatica as an alternative for the treatment of AD.
2. Results

2.1. Quantitative High-Performance Liquid Chromatography (HPLC) Analysis

Figure 1A,B shows a chromatogram of RECA and the reference standards which are asiaticoside, madecassoside, asiatic acid, and madecassic acid. Four triterpenes in RECA, which are madecassoside, asiaticoside, madecassic acid, and asiatic acid were quantified by using the HPLC gradient method with an ultraviolet (UV) detector at 200 nm (Table 1). This extract contains a high proportion of glycosides than the aglycones, in which madecassoside is the highest component. Several researchers were able to detect these triterpenes within the wavelength range of 200–220 using a gradient system comprising water and acetonitrile [28–30].

Figure 1. Chromatograms of raw-extract of C. asiatica (RECA) (A) and standards mixture (B). (1): madecassoside; (2): asiaticoside; (3): madecassic acid; (4): asiatic acid.
was observed according to the modified US National Cancer Institute and Geran et al. [31] criteria, as follows: IC₅₀ ≤ 20 µg/mL = highly cytotoxic; IC₅₀: 21–200 µg/mL = moderately cytotoxic; IC₅₀: 201–500 µg/mL = weakly cytotoxic; and IC₅₀ > 501 µg/mL = non-cytotoxic. Since the concentration of RECA up to 1000 µg/mL did not exhibit cytotoxicity against both cell lines, these concentrations were selected for subsequent experiments.

2.2. Effect of RECA on Viability of SH-SY5Y and RAW 264.7 Cells

The cytotoxic effect of RECA at a concentration ranging from 3.91 to 1000 µg/mL was investigated on neuroblastoma cells (SH-SY5Y) and murine microglia cells (RAW 264.7) by using MTT assay. The effect of RECA on SH-SY5Y and RAW 264.7 cell lines was evaluated after 24 h and 48 h (Figure 2) of cell exposure to the extract. The cell viability percentage was calculated by measuring the resulting intracellular purple formazan formed from the reduction of MTT solution by the presence of mitochondrial dehydrogenase in viable cells. It was observed that cell viability on both cells increased with the increase in concentrations of RECA. However, the level of cell viability was slightly lowered at 48 h as compared to 24 h of exposure. RECA at the highest concentration of 1000 µg/mL showed maximal cell viability at both hours of exposure (*p ≤ 0.05 and ****p ≤ 0.0001). No significant degree of cytotoxicity was observed according to the modified US National Cancer Institute and Geran et al. [31] criteria, as follows: IC₅₀ ≤ 20 µg/mL = highly cytotoxic; IC₅₀: 21–200 µg/mL = moderately cytotoxic; IC₅₀: 201–500 µg/mL = weakly cytotoxic; and IC₅₀ > 501 µg/mL = non-cytotoxic. Since the concentration of RECA up to 1000 µg/mL did not exhibit cytotoxicity against both cell lines, these concentrations were selected for subsequent experiments.

Table 1. Triterpene concentration of RECA.

| Types of Triterpenes | Retention Time (min) | Concentration of Triterpenes in RECA ± SEM (mg/g) | % of Dry Plant |
|----------------------|----------------------|-----------------------------------------------|----------------|
| Madecassoside        | 9.077                | 231.44 ± 7.05                                 | 3.70 ± 0.11    |
| Asiaticoside         | 11.393               | 223.21 ± 5.63                                 | 3.57 ± 0.09    |
| Madecassic acid      | 22.207               | 118.95 ± 8.47                                 | 1.91 ± 0.14    |
| Asiatic acid         | 25.417               | 108.37 ± 2.25                                 | 1.73 ± 0.04    |

SEM = standard error of the mean.

Figure 2. Cytotoxicity effect of RECA against SH-SY5Y (A,B) and RAW 264.7 (C,D) cell lines after 24 h (A,C) and 48 h (B,D) of exposure using MTT assay at the concentration ranging from 3.91 to 1000 µg/mL. NC = negative control (untreated cells). Results are expressed as the mean ± SEM in triplicate. *p ≤ 0.05, **p ≤ 0.01, and ****p ≤ 0.0001 compared to NC denoted as 100%.
2.3. Inhibitory Effect of RECA on AChE Activity

To evaluate the inhibitory effect of RECA on AChE activity in vitro, SH-SY5Y cells were used in this study. The differentiated cells were exposed to RECA at a concentration ranging from 3.91 to 1000 µg/mL for 24 h. The enzyme activity was estimated by Ellman’s method and expressed where unit of enzyme catalyzes the production of 1 µmole of thiocholine per minute under the assay conditions (pH 7.5 and room temperature). Eserine at 0.16 µM was used as the standard AChE inhibitor in this study. Based on Figure 3, differentiated SH-SY5Y cells (RA) showed a significant increase (**** p ≤ 0.0001) in AChE activity as compared to undifferentiated cells. However, AChE activity in differentiated cells showed a significant decrease in a concentration-dependent manner when the cells were treated with RECA for 24 h. Interestingly, RECA at the highest concentration (1000 µg/mL) showed significant maximal inhibition (##### p ≤ 0.0001) on AChE activity as compared to RA alone. The IC50 values for RECA and eserine were found at values of 31.09 ± 10.07 µg/mL and 0.242 ± 0.227 µg/mL, respectively.

![Figure 3](image-url)

**Figure 3.** Effect of RECA on acetylcholinesterase activity after 24 h of exposure at the concentration ranging from 3.91 to 1000 µg/mL. NC = negative control (untreated cells) and RA = differentiated SH-SY5Y cells. Results are expressed as the mean ± SEM in triplicate. **** p ≤ 0.0001 compared NC, whereas #### p ≤ 0.01 and ##### p ≤ 0.0001 as compared to RA.

2.4. Inhibitory Effect of RECA on Anti-Inflammatory Activities

In order to elucidate the capability of RECA on suppressing the pro-inflammatory cytokines and mediators, initially, we examined the effect of RECA on inhibiting the nitrite level in LPS-induced RAW 264.7 cell line. The nitrite level in the cellular supernatants was assessed using Griess reagent. Dexamethasone at 5 µM was used as the reference drug. Nitrite (<2 µM) was present in unstimulated RAW 264.7 microglia cells; however, the level of nitrite was markedly increased (**** p ≤ 0.0001) after exposure to LPS for 24 h. Treatment with RECA at various concentrations ranging from 3.91 up to 1000 µg/mL significantly decreased (##### p ≤ 0.0001) the subsequent LPS-induced nitrite production in a concentration-dependent manner with the maximal inhibitory effect seen at a dosage of 1000 µg/mL. The present study also found the nitrite-suppressing effect of treatment with RECA at the highest dosage (1000 µg/mL) to be better than the reference drug, dexamethasone (Figure 4). RECA was found to exhibit the greatest maximal 50% inhibitory effect of nitrite with a value of 5.205 ± 3.13 µg/mL.
As RECA showed the greatest inhibition activity of nitrite production, we also examined its suppressive effect toward other pro-inflammatory cytokine/mediators, which are prostaglandin E\textsubscript{2} (PGE\textsubscript{2}) and tumor necrosis factor alpha (TNF-\textalpha{}). Dexamethasone at 5 µM and 6 µM were used as the reference drug for PGE\textsubscript{2} and TNF-\textalpha{} analyses, respectively. RAW 264.7 macrophage cells that were stimulated with 4 µg/mL LPS markedly (** p ≤ 0.001 and **** p ≤ 0.0001) showed an increment in PGE\textsubscript{2} and TNF-\textalpha{} expressions as compared to that generated by control. The result in Figure 5 showed a similar inhibition effect as demonstrated in Figure 4 as RECA inhibited the expression of PGE\textsubscript{2} and TNF-\textalpha{} in the macrophages stimulated with LPS in a concentration-dependent manner. Moreover, RECA at the highest concentration (1000 µg/mL) showed a remarkable inhibition effect (** p ≤ 0.001) as compared to untreated cells in both assays. RECA was found to exhibit a maximal 50% inhibitory effect of PGE\textsubscript{2} and TNF-\textalpha{} with values of 261.4 ± 14.2 µg/mL and 744 ± 14.38 µg/mL, respectively.

Figure 4. Effect of RECA on nitrite production. Result is expressed as the mean ± SEM in triplicate. NC = negative control (untreated cells), DEXA = dexamethasone and LPS = lipopolysaccharides. **** p ≤ 0.0001 compared to NC, whereas, #### p ≤ 0.0001 compared to LPS.

Figure 5. Cont.
Potential antioxidant properties. To this end, we first examined the effect of RECA on reactive oxygen species (ROS) production in RAW 264.7 macrophages stimulated with LPS. The ROS production was quantified using the stable nonpolar dye 2′,7′-dichlorodihydrofluorescein diacetate (H2DCF-DA), which readily diffuses into the cells and is enzymatically hydrolyzed by intracellular esterase to non-fluorescent DCF. A minimal ROS level (<40,000 RFU) was present in unstimulated cells; however, a noteworthy generation of ROS (**** p ≤ 0.0001) was observed once cells were stimulated with LPS (Figure 6). RECA was observed inhibiting the production of ROS in a concentration-dependent manner with RECA yielding the greatest suppression (##### p ≤ 0.0001) at the highest concentration (1000 µg/mL). RECA was found to exhibit greatest the maximal 50% inhibitory effect of ROS with a value of 20.15 ± 2.54 µg/mL.

2.5. Inhibitory Effect of RECA on Antioxidant Activities

Having observed the encouraging anti-inflammatory effects of RECA, we further explore its potential antioxidant properties. To this end, we first examined the effect of RECA on reactive oxygen species (ROS) production in RAW 264.7 macrophages stimulated with LPS. The ROS production was quantified using the stable nonpolar dye 2′,7′-dichlorofluorescein (DCF), which is rapidly oxidized to highly fluorescent DCF in the presence of ROS. α-tocopherol at 160 µM was used as the reference drug. A minimal ROS level (<40,000 RFU) was present in unstimulated cells; however, a noteworthy generation of ROS (**** p ≤ 0.0001) was observed once cells were stimulated with LPS (Figure 6). RECA was observed inhibiting the production of ROS in a concentration-dependent manner with RECA yielding the greatest suppression (##### p ≤ 0.0001) at the highest concentration (1000 µg/mL). RECA was found to exhibit greatest the maximal 50% inhibitory effect of ROS with a value of 20.15 ± 2.54 µg/mL.

Figure 5. Effect of RECA on (A) PGE2 and (B) TNF-α expressions. NC = negative control (untreated cells), DEXA = dexamethasone, and LPS = lipopolysaccharides. Results are expressed as the mean ± SEM in triplicate. *** p ≤ 0.001 and **** p ≤ 0.0001 compared to NC, whereas, # p ≤ 0.05, ## p ≤ 0.01, ### p ≤ 0.001, and #### p ≤ 0.0001 compared to LPS.

Figure 6. Effect of RECA on reactive oxygen species (ROS) production. NC = negative control (untreated cells), TOCO = α-tocopherol and LPS = lipopolysaccharides. Results are expressed as the mean ± SEM in triplicate. **** p ≤ 0.0001 compared to untreated cells (NC), whereas, ##### p ≤ 0.0001 compared to LPS.
Reduced glutathione (GSH) content in LPS-induced RAW 264.7 cell line was also quantified, as shown in Figure 7. α-tocopherol at 250 μM was used as the reference drug for this assay. Unstimulated RAW 264.7 microglia cells were found to exhibit a high level of GSH (≤1.46 μM); however, the GSH content was significantly (** p ≤ 0.00) depleted following stimulation with LPS. In general, incubation of RECA on stimulated cells prevented the LPS-induced depletion in GSH level in a dose-dependent manner up to the concentration of RECA at 125 μg/mL. Nonetheless, further incubation of RECA at a concentration of 250 μg/mL to 1000 μg/mL elicited suppression effect on the GSH content.

**Figure 7.** Effect of RECA on glutathione (GSH) content. NC = negative control (untreated cells), TOCO = α-tocopherol, and LPS = lipopolysaccharides. Results are expressed as the mean ± SEM in triplicate. ** p ≤ 0.01 as compared to NC.

2.6. Inhibitory Effect of RECA on AChE Activity In Vivo

The effect of increasing the concentration of RECA on AChE activity was further evaluated by using the LPS-induced Sprague Dawley rats’ model. Based on Figure 8, the LPS-induced group showed an elevation in AChE activity as compared to control. Treatment of rats with RECA after the LPS injection inhibited LPS-induced enhancement in AChE activity, in a dose-dependent manner. Moreover, it also found that the AChE suppressing effect of the RECA-treated group at a median dosage (300 mg/mL) was comparable to the reference drug group, rivastigmine.

**Figure 8.** Effect of RECA on acetylcholinesterase activity in vivo. LPS = lipopolysaccharides. Results are expressed as the mean ± SEM in triplicate.

2.7. Inhibitory Effect of RECA on Anti-Inflammatory Activities In Vivo

The effects of RECA on the PGE_2 and TNF-α levels are shown in Figure 9A,B. Rats induced with LPS alone exhibited a significant (**** p ≤ 0.0001) elevation in both levels of PGE_2 and TNF-α, as compared to the respective control group. The rivastigmine group, on the other hand, significantly (### p ≤ 0.001 and
### p ≤ 0.0001) suppressed the PGE₂ and TNF-α levels. Significant reductions (## p ≤ 0.01, ### p ≤ 0.001 and #### p ≤ 0.0001) in both mediators’ levels were also observed in RECA-treated groups at doses of 300 and 350 mg/kg; in contrast, the RECA-treated group at a dose of 250 mg/kg had no significant effect on the level of both mediators.

![Figure 8](image1.png)

**Figure 8.** Effect of RECA on acetylcholinesterase activity in vivo. LPS = lipopolysaccharides. Results are expressed as the mean ± SEM in triplicate.

#### 2.7. Inhibitory Effect of RECA on Anti-Inflammatory Activities In Vivo

The effects of RECA on the PGE₂ and TNF-α levels are shown in Figure 9A,B. Rats induced with LPS alone exhibited a significant (**** p ≤ 0.0001) elevation in both levels of PGE₂ and TNF-α, as compared to the respective control group. The rivastigmine group, on the other hand, significantly (### p ≤ 0.001 and #### p ≤ 0.0001) suppressed the PGE₂ and TNF-α levels. Significant reductions (## p ≤ 0.01, ### p ≤ 0.001 and #### p ≤ 0.0001) in both mediators' levels were also observed in RECA-treated groups at doses of 300 and 350 mg/kg; in contrast, the RECA-treated group at a dose of 250 mg/kg had no significant effect on the level of both mediators.

![Figure 9A](image2.png)

**Figure 9A.** Effect of RECA on PGE₂ expressions. Results are expressed as the mean ± SEM in triplicate. LPS = lipopolysaccharides. *** p ≤ 0.001 and **** p ≤ 0.0001 compared to control group, whereas, # p ≤ 0.05, ## p ≤ 0.01, ### p ≤ 0.001, and #### p ≤ 0.0001 compared to LPS-induced group.

![Figure 9B](image3.png)

**Figure 9B.** Effect of RECA on TNF-α expressions. Results are expressed as the mean ± SEM in triplicate. LPS = lipopolysaccharides. *** p ≤ 0.001 and **** p ≤ 0.0001 compared to control group, whereas, # p ≤ 0.05, ## p ≤ 0.01, ### p ≤ 0.001, and #### p ≤ 0.0001 compared to LPS-induced group.

#### 2.8. Inhibitory Effect of RECA on Antioxidant Activity In Vivo

In order to elucidate the anti-oxidant activity of RECA in the LPS-induced Sprague Dawley rats’ model, the reduced glutathione (GSH) level was quantified. As shown in Figure 10, the GSH level was depleted in the LPS-induced group. In contrast, the rivastigmine and RECA-treated groups at the highest dosage (350 mg/kg) significantly (# p ≤ 0.05) elevated the level of GSH as compared to the group that induced with LPS alone. The effect of the RECA-treated group at a dosage of 350 mg/kg was more pronounced than the standard drug rivastigmine.
with long and extensively branched neurites that connect to surrounding cells, as compared to Alzheimer’s disease (AD). Therefore, as a representative cell model to evaluate the inhibitory effect on AChE activity in vitro, differentiation of SH-SY5Y cells by the non-toxic and supraphysiologic concentration of all-trans-retinoic acid (ATRA) was used in this study.

In order to produce more neurons with a cholinergic phenotype and maximize basal AChE activity, SH-SY5Y cells are commonly induced to differentiate by the addition of retinoic acid (RA) [39,40]. RA has been shown to induce cellular differentiation through various mechanisms, in such a halting cell cycle progression out of G0/G1, enhancing expression of the cyclin-dependent kinase (CDK) inhibitors p21 and p27Kip1 and the anti-apoptotic proteins Bcl-2 and Bcl-xL, and increasing PI3K/AKT activity, which plays a role in neurite development and differentiation [41]. To address whether RA-differentiated SH-SY5Y cells are functionally mature in this present study, morphological changes of the cells were observed as in Figure 11, in which the differentiated cells manifested themselves with long and extensively branched neurites that connect to surrounding cells, as compared to undifferentiated cells [42]. The present study shows that treatment of differentiated cells with RECA for 24 h had significantly suppressed the elevated AChE activity with IC50 values of 57.47 ± 13.55 µg/mL.

3. Discussion

The present study reports the capability of ethanolic extract of C. asiatica (RECA) in inhibiting acetylcholinesterase (AChE) activity and its possible association in suppressing the elevated expression of pro-inflammatory cytokine/mediators and oxidative stress via in vitro and in vivo models of Alzheimer’s disease (AD).

Even though AD is manifested by the accumulation of neurofibrillary tangles and amyloid plaques, several pieces of evidence suggest the involvement of AChE in the pathogenesis of AD [3,32,33]. Severe loss of cholinergic neurons in brain potentially gives a great impact in all aspects of cognition and behavior in AD patients; hence, drugs targeting the cholinergic system represent the most popular option so far to treat AD patients [34,35]. The human neuroblastoma SH-SY5Y cell line is known to be a relevant cellular model for biochemical investigations on AD, with respect to molecular and neurochemical parameters [36,37]. However, this cell line is rapidly dividing and expresses immature neuronal markers by displaying low morphological and biochemical features of mature neurons [38]. Therefore, as a representative cell model to evaluate the inhibitory effect on AChE activity in vitro, differentiation of SH-SY5Y cells by the non-toxic and supraphysiologic concentration of all-trans-retinoic acid (ATRA) was used in this study.

Figure 10. Effect of RECA on GSH level. LPS = lipopolysaccharides. Results are expressed as the mean ± SEM in triplicate. * p ≤ 0.05 compared to LPS-induced group.
RECA’s inhibition of the AChE enzyme could boost the amount of neurotransmitter acetylcholine and thus enhance synaptic transmission in the AD brain. It is well known that the most significant modifications in AD patients are decreased levels of acetylcholine (ACh) in the brain’s cortex and hippocampus. Consequently, inhibition of AChE, in which the enzyme responsible for ACh hydrolysis in the cholinergic synapse interaction, may be a suitable target for AD therapy [43]. Results from Orhan et al. [44] demonstrated that ethanolic C. asiatica extract with 10.78% content of asiaticoside and madecassoside displayed close to 50% inhibition against AChE at 200 µg/mL. Therefore, the AChE inhibition observed in RECA could be attributed by the high proportion of triterpene saponins such as asiaticoside and madecassoside presented in the extract, as these bioactive components were reported to be responsible for its several beneficial pharmacological activities [45].

Neuroinflammation has been implicated to play a pivotal role in facilitating the development of several events in the pathological cascade of AD [46,47]. Microglia cells are the resident immune cells of the central nervous system (CNS) networks and can specifically interact with neurons in the brain once activated [48]. However, uncontrolled activation of microglia by pathological triggers, like neuronal death or protein aggregates, could result in the excessive release of various pro-inflammatory cytokines and neurotoxic agents. These will further lead to CNS damage and disease pathology [49]. On this basis, intervention against the activation process of microglia therefore presents a possible critical target in AD therapy.

In this study, the anti-inflammatory and antioxidant effects of RECA were evaluated against LPS-stimulated RAW 264.7 microglia cells. It is known that for the exposure of microglia cells to inflammatory stimuli, as in this study, LPS was preferred. It exacerbates a wide array of inflammatory mediators including pro-inflammatory cytokines, such as NO, PGE2, and TNF-α, which are deleterious to neurons and oligodendrocytes [50]. However, treatment of the activated cells with RECA significantly suppressed the elevated pro-inflammatory cytokine/mediators in a concentration-dependent manner, with the highest concentration (1000 µg/mL) showing remarkable inhibition in all assays. Among all, RECA displayed strong inhibition against nitrite and these inhibitory effects on inflammatory activities observed in RECA were not attributable to its cytotoxic effect, as assessed by MTT. This trend of inhibition clearly confirmed that madecassic acid plays an important role in the anti-inflammatory activity of C. asiatica as compared to their glycosides. Therefore, in the present study, RECA reversed the inflammatory activities moderately may be due to the smaller amount of madecassic acid presents in the extract.

Perpetual overactivation of microglia causes an overwhelming production of ROS that promotes oxidative stress and downregulates the level of antioxidants, such as reduced glutathione (GSH) [51]. GSH is the most abundant intracellular non-protein thiol in cells and plays a central role in maintaining cellular redox status. By increasing the GSH content, it would be expected to reduce the ROS level and antagonize apoptotic signals [52]. In support of previous studies, LPS exposure was shown to induce oxidative stress via accelerating the ROS level and, at the same time, GSH depletion [53–55]. However, treatment with RECA in the present study suppressed the oxidative changes stimulated by the LPS treatment, indicative of its role in inhibiting the production of pro-inflammatory cytokines and ROS.
by LPS. These inhibitory effects were associated with the reduction of ROS level and stimulation of GSH content, indicating the role of RECA in oxidative balance. A study done by Sasmita et al. [56] indicated that cells treated with the madecassoside at a maximum non-toxic dose (MNTD) were able to suppress the intracellular ROS generations by downregulating the gene and protein expression of pro-neuroinflammatory components. Therefore, the antioxidative properties observed in RECA could be attributed to the high proportion of madecassoside in the extract.

Generally, treatment with RECA in LPS-stimulated macrophages enhanced the GSH content in a concentration-dependent manner up to 125 µg/mL and GSH content at this concentration was observed to be better than the reference drug, α-tocopherol. In contrast, incubation of activated microglia cells with RECA at a further concentration from 250 µg/mL to 1000 µg/mL elicited suppression effect on the GSH content. Interestingly, this trend was found almost similar with the study done by Naidoo et al. [57], in which *C. asiatica* was observed to enhance the GSH content up to 0.2 mg/mL before it started to decrease at further concentrations. To the best of our knowledge, no previous literature has reported the effect of *C. asiatica* in attenuating GSH content in activated RAW 264.7 cells; hence, these findings warrant further investigation.

In order to ascertain whether the in vivo use of RECA can offer the same anti-acetylcholinesterase, anti-inflammatory, and antioxidative effects, we further examined the effect of RECA on the Sprague Dawley rats subjected to LPS injection. The establishment of an appropriate animal model for researching neuroinflammation associated with neurodegeneration in neurodegenerative diseases is very crucial. A large body of experimental evidence has demonstrated that LPS is one of the most suitable inducers for studying the mechanism of neuroinflammation and oxidative stress in animal models, which closely resembles AD [58–60]. A single injection of LPS intraperitonially caused neuroinflammation in rats via the TLR-4 signaling pathway, which further activates various intracellular molecules that affect the expression of a plethora of steady-state transcripts such as TNF-α, NO, and PGE₂ in the serum and brain homogenates [59,61]. In addition, the up-regulated of AChE activity upon the LPS injection in the present study was found in consonance with preceding reports [62–64]. This observation might reflect the deficit of cholinergic neurotransmission in the brain and further supports the impression that altered activity of AChE that has been found in patients with AD is attributed by the overexpression of pro- and anti-inflammatory cytokines [65]. Conversely, post-administration of RECA attenuated the AChE activity in a dose-dependent manner where the highest dosage showed a comparable AChE inhibition as rivastigmine. This finding was found in line with our previous work in which RECA showed a protective effect against LPS-pretreated rats [66]. Therefore, it is worth noting that RECA possesses both curative and protective properties as an AChE inhibitor.

In the current study, the depletion of GSH content in LPS-induced rats indicates that there was an overexpression of free radical generation and that the reduced GSH became depleted as it confers protection against oxidative stressors. According to an autopsy study, a magnitude of GSH depletion was found in the AD cingulated cortex and AD substantia innominate, and these unfavorable events were correlated with a decline in cognitive functions [67,68]. Nonetheless, post-administration of RECA has demonstrated a significant increment in GSH content and a reduction of TNF-α and PGE₂ levels in LPS-induced rats. Besides, it is also observed that GSH content at the highest dosage of RECA (350 mg/kg) was more prominent as compared to rivastigmine. Research has shown that intranasal GSH administration in patients with Parkinson’s disease (PD) elevates GSH levels in the brain [69], whereas clinical studies involving GSH have shown that the ingested GSH has important nutraceutical benefits for human health to improve oxidative stress and human protection [70–72]. In addition, the study of individuals with mild cognitive impairment (MCI) showed that the rate of GSH significantly elevated in both the anterior and posterior cingulate regions in patients with MCI as compared to the healthy control participants. The increase in GSH level was explained as an early compensatory or neuroprotective response [73]. Such findings offer important insight into the importance of an effective brain antioxidant system during longevity [74]. Indeed, we postulate that RECA may emerge its antioxidant property by reinforcing the GSH content, which not only serves
as an antioxidant agent but also plays as a protective role against inflammatory pathologies in the inflamed brain.

Through the comprehensive analysis of the data referred to above, we can presume that RECA has the capability in quashing the elevated level of AChE, inflammation, and oxidative stress activities.

4. Materials and Methods

4.1. Chemicals and Reagents

All solvents were HPLC grade, purchased from RCI Labscan (Bangkok, Thailand). The chemical standards of asiaticoside, madecassoside, asiatic acid, and madecassic acid, were purchased from Chemfaces (Wuhan, Hubei, China). Dulbecco’s modified Eagle’s medium (DMEM), Eagle’s minimum essential medium (MEM), fetal bovine serum (FBS), trypsin (0.25%, 1 mM EDTA), and penicillin streptomycin were obtained from Nacalai Tesque (Kyoto, Japan). Ham’s F-12K (Kaighn’s) Medium was obtained from Gibco, Thermo Fisher Scientific, Inc. (Waltham, MA, USA). Dimethyl sulfoxide, LPS from *Escherichia coli* (O26:B6), 3-[4,5-dimethylthiazol-2-yl]-2,5-diphenyltetrazolium bromide (MTT), 2,7-dichlorofluorescein diacetate (DCF-DA), Hanks Balanced Salt Solution, all-trans-retinoic acid, and phosphate-buffered saline (PBS) were obtained from Sigma-Aldrich (St. Louis, MO, USA). The triterpenoid standards, asiaticoside, madecassoside, asiatic acid, and madecassic acid were purchased from ChemFaces Natural Products Co., Ltd., China. AChE activity was measured by using QuantiChrom™ Acetylcholinesterase Assay kit (BioAssay Systems, Hayward, CA, USA). The Griess reagent kit for nitrite determination was purchased from Promega™ Griess Reagent System (Promega Biotech Co., Ltd., Madison, WI, USA). The enzyme-linked immunosorbent assay (ELISA) kit for TNF-α measurement was obtained from Elabscience (Rockford, IL, USA), and PGE₂ and GSH measurements were obtained from Cayman Chemical Company (Ann Arbor, MI, USA). All of the other reagents were of the highest commercially available grade.

4.2. Plant Material Preparation and Extraction

Raw materials were obtained from Herbagus Trading, Pulau Pinang, Malaysia. The whole plant was washed, cleaned, and oven-dried at 40 °C. The powdered plant material was extracted using a standard extraction protocol at the extraction facility of the Institute of Bioproducts Development (IBD), Universiti Teknologi Malaysia. The extraction was carried out by using 95% denaturated ethanol for 8 h at a temperature of 60 °C. The yield of crude extract with regard to the weight of dried plant material was about 16%. This extract was designated as raw-extract *C. asiatica* (RECA). Voucher specimen (CA-K017) was prepared and deposited in the Atta-ur-Rahman Institute for Natural Product Discovery, UiTM Puncak Alam (Selangor, Malaysia) for future reference.

4.3. Fingerprint Chromatogram Assessment

Chromatographic profiling of RECA and its standards was performed by using a Thermo Scientific™ UltiMate™ 3000 HPLC system coupled to a diode array detector (detector wavelength 200 nm), equipped with an ultra-high pressure pump and vial sampler, as described by previous literature with slight modifications [75]. Chromeleon chromatographic software was used for data collection and processing. Chromatographic separation was carried out on a reversed-phase Ascentis® C18 HPLC column (Supelco, USA) (15 cm x 4.6 mm i.d., 5 µm) with injection volume of 100 µL and a flow rate of 1.0 mL/min. The acquisition time for the analysis was 55 min using a mobile phase with ultra-pure water and HPLC grade acetonitrile, as shown in Table 2A,B, respectively. The concentration of sample for each analysis was 10 mg/mL dissolved in methanol-water (7:3) and filtered through a 0.22 µM nylon membrane syringe filter, prior to the injection. Standard solutions for the evaluation of linearity were prepared over a concentration range of 10, 50, 100, 500, and 1000 ppm in methanol:water (1:1). Quantification of marker compounds was done using calibration curves constructed from the
standards and the calibration curve criteria were accepted, with \( r^2 > 0.998 \) indicating an acceptable fit. The relative amount of the compound was expressed as milligram per gram of extract.

### Table 2. Gradient condition for HPLC.

| Time (min) | Pump A, Water (%) | Pump B, Acetonitrile (%) |
|------------|-------------------|-------------------------|
| 0          | 80                | 20                      |
| 15         | 65                | 35                      |
| 30         | 35                | 65                      |
| 35         | 20                | 80                      |
| 40         | 20                | 80                      |
| 45         | 80                | 20                      |
| 55         | 80                | 20                      |

### 4.4. Cell Culture

Human neuroblastoma cells, SH-SY5Y, and murine macrophage RAW 264.7 were obtained from the American Type Culture Collection (ATCC no. CRL-2266 and no. TIB-71™, Manassas, VA, USA). SH-SY5Y was cultured in the mixture of minimum essential medium (MEM) Eagle with Earle’s salt and F12-K medium in a 1:1 ratio, whereas RAW 264.7 was cultured in Dulbecco’s modified Eagle medium (DMEM). Both were supplemented with 10% heat-inactivated fetal bovine serum (FBS) and 1% penicillin/streptomycin in a humidified incubator maintained at 37 °C under 5% CO\(_2/95\%\) air. The media were changed every 2 days and subculture was performed when cells reached 80% confluence. Cell numbers were assessed using a hemocytometer. For cell viability assay, SH-SY5Y and RAW 264.7 cells were plated at density 10 \( \times \) 10\(^3\) cells/well and 30 \( \times \) 10\(^3\) cells/well in 96-well plates. Meanwhile, for assay to determine inhibition of AChE activity, SH-SY5Y cells were plated at a density of 5 \( \times \) 10\(^5\) cells/well in 6-well plates. For assays to determine inhibition activity of NO, TNF-\(\alpha\), PGE\(_2\), and ROS productions, RAW 264.7 cells were plated at density 30 \( \times \) 10\(^3\) cells/well in 96-well plates; whereas, to determine the production of GSH level, RAW 264.7 cells were plated at density 8 \( \times \) 10\(^5\) cells/well in 6-well plates.

### 4.5. Determination of Cell Viability Assay

The cytotoxic effects of RECA on the SH-SY5Y and RAW 264.7 cell lines were determined colorimetrically using the MTT method, which is based on the reduction of tetrazolium salt by mitochondrial dehydrogenase in viable cells. Serial two-fold dilutions of the extracts were added to the wells in triplicate with an increasing concentration (3.91–1000 \( \mu\)g/mL) and incubated for 24 h. A solution of the MTT reagent, 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyl tetrazolium bromide (5 mg/mL PBS) was added to both cells in each well and incubated for the next 3 h. Resulting formazan was solubilized with 100 \( \mu\)L dimethyl sulfoxide (DMSO). The plate was shaken for 15 min and absorbance determined at 540 nm using a SPECTROstar Nano plate reader (BMG Labtech Inc., Cary, NC, USA). Values were calculated in comparison to the control cells.

### 4.6. Determination of Anti-Acetylcholinesterase Activity

The AChE inhibitory activity was measured using Ellman’s method, as reported previously [66]. SH-SY5Y cells were induced to differentiate into neuronal cells by all-trans-retinoic acid (ATRA; 10 \( \mu\)M). After differentiation, culture media was removed, and cells incubated with various concentrations of RECA (3.91–1000 \( \mu\)g/mL) for the next 24 h. Eserine was used as the reference drug. Cell lysates were assayed into 96-well microplates for acetylcholinesterase activity as per manufacturer’s instructions. The intensity of the color change was measured at 412 nm of the 2nd and 10th min using the microplate reader.
4.7. Determination of Anti-Inflammatory Activities

4.7.1. Nitrite Production

Briefly, RAW 264.7 cells were plated and left incubated for 24 h. Dexamethasone was used as the standard drug. Cells were induced with LPS for 24 h prior to the treatment with RECA/dexamethasone at the indicated concentration for the next 24 h. The concentration of nitrite in the cultured media was measured at 540 nm. Nitrite was quantified by an external standard, using sodium nitrite to generate a standard curve.

4.7.2. Prostaglandin E2 (PGE2) and Tumor Necrosis Factor Alpha (TNF-α) Levels

In brief, RAW 264.7 cells were stimulated with LPS (4 µg/mL) 24 h prior to the treatment with several concentrations of RECA (3.91–1000 µg/mL) for the next 24 h. The PGE2 and TNF-α productions were quantified by measuring its cultured media at 412 nm.

4.8. Determination of Antioxidant Activities

4.8.1. Intracellular Reactive Oxygen Species (ROS) Production

Assay was performed as described in Section 4.7.1 with α-tocopherol was used as the standard drug. At the end of treatment, the medium was discarded, and the wells were gently washed thrice with Hanks Balanced Salt Solution (HBSS). Cells were stained with 20 µM H2DCF-DA and left incubated for 45 min at 37 °C in the dark. Then, cells were gently washed thrice with HBSS to remove excess dye solution. Finally, an additional 200 µL of HBSS was loaded to the cells and the fluorescence was detected in a spectrophotometer at an excitation wavelength of 485 nm and an emission wavelength of 520 nm.

4.8.2. Intracellular Reduced Glutathione (GSH) Level

In brief, RAW 264.7 cells were stimulated with LPS (4 µg/mL) 24 h prior to the treatment with several concentrations of RECA (3.91–1000 µg/mL) for the next 24 h. α-tocopherol was used as the standard drug. The GSH level was quantified by measuring the cell lysates at 450 nm as per manufacturer’s instructions.

4.9. Animal Study Design

Adult male Sprague Dawley rats, weighing approximately 200–250 g, were procured from the Laboratory Animal Facility and Management Faculty of Pharmacy (LAFAM), UiTM Puncak Alam, Selangor. Rats were housed in individually ventilated cages (IVC) cage and maintained under standard laboratory conditions that were automatically kept at 21–25 °C and relative humidity at 45%–65% with a controlled light–dark cycle. All rats had free access to standard laboratory food and tap water ad libitum. Rats were housed in groups and acclimatized for at least 7 days before using them for experiments. Experiments were carried out between 0800 h to 1800 h. The protocol was approved by the Committee on Animal Research and Ethics of the Universiti Teknologi MARA (UiTM) (Reference No: 600-FF (PS. 17/2/1); dated 28 August 2017). The laboratory animals were handled and managed in accordance to the Guide for the Care and Use of Laboratory Animal (National Research Council 1996). Thirty-six rats were randomly assigned into 6 groups (6 rats/group; Table 3) for biochemical analysis. Except for untreated control, all groups were injected intra-peritoneally (i.p) with LPS at a concentration of 250 µg/kg for first the 10 days, and subsequently from day 11 to day 21, RECA (250, 300 and 350 mg/kg) and rivastigmine (5 mg/kg) were administered through oral gavage. The experimental study design is shown in Figure 12. All the drugs (RECA, rivastigmine, and LPS) were freshly prepared before administration. The rats were sacrificed by cervical decapitation under light anesthesia for biochemical assessment on day 22.
The doses of RECA, rivastigmine, and LPS administered were selected based on previous work [66].

4.9.1. Vehicles

LPS was diluted in phosphate buffer saline (PBS). Rivastigmine (5 mg/kg) and RECA (250, 300, and 350 mg/kg) were suspended in distilled water for oral administration.

4.9.2. Biochemical Analyses

Immediately after decapitation, the whole brain was carefully removed from the skull, washed, and homogenized in ice-cold PBS to prepare 10% (w/v) homogenate by using a handheld rotor-stator homogenizer (TissueRuptor, QIAGEN, Germany). The brain homogenate was centrifuged at 3000 rpm at 4 °C for 15 min to remove cellular debris and resultant cloudy supernatant was collected and used for biochemical assessment such as quantification of AChE, anti-inflammatory, and antioxidant activities as per the manufacturer’s instructions.

4.9.3. Statistical Analysis

For statistical analysis, each of the experimental values was compared with its corresponding control. Results were expressed as the mean ± standard error of the mean (SEM). Mean differences among groups were evaluated by one-way analysis of variance (ANOVA) using the GraphPad Prism 6.0 software (SAS Institute, NC, USA). Post-hoc comparisons between groups were made using Tukey’s test. At least, at the level of \( p < 0.05 \), the results were considered as statistically significant.

5. Conclusions

In conclusion, current findings suggest that the raw-extract of \( C. \text{asiatica} \) (RECA) possesses capability in quashing the elevated level of acetylcholinesterase (AChE), inflammation, and oxidative stress activities in vitro and in vivo. The combination of findings from in vitro and animal studies in the present work are useful for current knowledge contribution and future research needs. Besides, it may also provide important insights that \( C. \text{asiatica} \) or compound derived from it can be considered as potential relevance in managing AD.
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References

1. Helzner, E.P.; Scarmeas, N.; Cosentino, S.; Tang, M.X.; Schupf, N.; Stern, Y. Survival in Alzheimer disease: A multiethnic, population-based study of incident cases. Neurology 2008, 71, 1489–1495. [CrossRef] [PubMed]
2. Luca, M.; Luca, A.; Calandra, C. The Role of Oxidative Damage in the Pathogenesis and Progression of Alzheimer’s Disease and Vascular Dementia. Oxid. Med. Cell. Longev. 2015, 2015, 1–8. [CrossRef] [PubMed]
3. Sanabria-Castro, A.; Alvarado-Echeverria, I.; Monge-Bonilla, C. Molecular pathogenesis of alzheimer’s disease: An update. Ann. Neurosci. 2017, 24, 46–54. [CrossRef] [PubMed]
4. Barone, E. Editorial (Thematic Issue: Oxidative Stress and Alzheimer Disease: Where Do We Stand?). Curr. Alzheimer Res. 2016, 13, 108–111. [CrossRef] [PubMed]
5. Overk, C.R.; Felder, C.C.; Tu, Y.; Schober, D.A.; Bales, K.R.; Wuu, J.; Mufson, E.J. Cortical M1 receptor concentration increases without a concomitant change in function in Alzheimer’s disease. J. Chem. Neuroanat. 2010, 40, 63–70. [CrossRef]
6. Pepeu, G.; Giovannini, M.G. Cholinesterase inhibitors and memory. Chem. Biol. Interact. 2010, 187, 403–408. [CrossRef]
7. Hung, S.Y.; Fu, W.M. Drug candidates in clinical trials for Alzheimer’s disease. J. Biomed. Sci. 2017, 47. [CrossRef]
8. Rosales-Corral, S.; Tan, D.-X.; Manchester, L.; Reiter, R.J. Diabetes and Alzheimer Disease, Two Overlapping Pathologies with the Same Background: Oxidative Stress. Oxid. Med. Cell. Longev. 2015, 2015, 1–18. [CrossRef] [PubMed]
9. Huang, G.J.; Huang, S.S.; Chiu, C.S.; Chen, H.J.; Hou, W.C.; Sheu, M.J.; Lin, Y.C.; Shie, P.H. Antinociceptive activities and the mechanisms of anti-inflammation of asiatic acid in mice. Evid. Based Complement. Altern. Med. 2011, 2011, 1. [CrossRef]
10. Osborn, G.G.; Saunders, A.V. Current treatments for patients with Alzheimer disease. J. Am. Osteopath. Assoc. 2010, 110, S16–S26.
11. Cooper, E.L.; Ma, M.J. Alzheimer Disease: Clues from traditional and complementary medicine. J. Tradit. Complement. Med. 2017, 7, 380–385. [CrossRef] [PubMed]
12. Li, J.W.; Vederas, J.C. Drug discovery and natural products: End of era or an endless frontier? Biomeditsinskaya Khimiya 2011, 57, 148. [CrossRef] [PubMed]
13. Heo, S.K.; Yun, H.J.; Yi, H.S.; Noh, E.K.; Park, S.D. Evodiamine and rutaecarpine inhibit migration by LIGHT via suppression of NADPH oxidase activation. J. Cell. Biochem. 2009, 107, 123–133. [CrossRef] [PubMed]
14. Tsai, T.H.; Lee, T.F.; Chen, C.F.; Wang, L.C.H. Thermoregulatory effects of alkaloids isolated from Wu-chu-yu in afebrile and febrile rats. Pharmacol. Biochem. Behav. 1995, 50, 293–298. [CrossRef]
15. Liu, A.-J.; Wang, S.-H.; Hou, S.-Y.; Lin, C.-J.; Chiu, W.-T.; Hsiao, S.-H.; Chen, T.-H.; Shih, C.-M. Evodiamine Induces Transient Receptor Potential Vanilloid-1-Mediated Protective Autophagy in U87-MG Astrocytes. Evid. Based Complement. Altern. Med. 2013, 2013, 1–9. [CrossRef]
16. Kudoh, C.; Arita, R.; Honda, M.; Kishi, T.; Komatsu, Y.; Asou, H.; Mimura, M. Effect of ninjin’yoeito, a Kampo (traditional Japanese) medicine, on cognitive impairment and depression in patients with Alzheimer’s disease: 2 years of observation. Psychogeriatrics 2016, 16, 85–92. [CrossRef]
17. Chanana, P.; Kumar, A. Possible Involvement of Nitric Oxide Modulatory Mechanisms in the Neuroprotective Effect of Centella asiatica Against Sleep Deprivation Induced Anxiety Like Behaviour, Oxidative Damage and Neuroinflammation. Phyther. Res. 2016, 30, 671–680. [CrossRef]
18. Hashim, P. Centella asiatica in food and beverage applications and its potential antioxidant and neuroprotective effect. Int. Food Res. J. 2011, 18, 1215.
19. Shinomol, G.K.; Muraleedhara, M.S.; Bharath, M. Exploring the Role of “Brahmi” (Bacopa monnieri and Centella asiatica) in Brain Function and Therapy. Recent Pat. Endocr. Metab. Immune Drug Discov. 2011, 5, 33–49.
20. Orhan, I.E. *Centella asiatica* (L.) Urban: From traditional medicine to modern medicine with neuroprotective potential. *Evid. Based Complement. Altern. Med.* **2012**, *2012*, 1–8. [CrossRef]

21. Singhal, A.; Bangar, O.; Naithani, V. Medicinal plants with a potential to treat Alzheimer and associated symptoms. *Int. J. Nutr. Pharmacol. Neurol. Dis.* **2012**, *2*, 84. [CrossRef]

22. Gohil, K.; Patel, J.; Gajjar, A. Pharmacological review on *Centella asiatica*: A potential herbal cure-all. *Indian J. Pharm. Sci.* **2010**, *72*, 546. [CrossRef] [PubMed]

23. James, J.; Dubery, I. Identification and quantification of triterpenoid centelloids in *Centella asiatica* (L.) Urban by densitometric TLC. *JPC. Planar Chromatogr. Mod. TLC* **2011**, *24*, 82–87. [CrossRef]

24. Won, J.H.; Shin, J.S.; Park, H.J.; Jung, H.J.; Koh, D.J.; Jo, B.G.; Lee, J.Y.; Yun, K.; Lee, K.T. Anti-inflammatory effects of madecassic acid via the suppression of NF-κB pathway in LPS-induced RAW 264.7 macrophage cells. *Planta Med.* **2010**, *76*, 251–257. [CrossRef]

25. Park, J.H.; Choi, J.Y.; Son, D.J.; Park, E.K.; Song, M.J.; Hellström, M.; Hong, J.T. Anti-inflammatory effect of titrated extract of *Centella asiatica* in phthalic anhydride-induced allergic dermatitis animal model. *Int. J. Mol. Sci.* **2017**, *18*, 738. [CrossRef]

26. Gray, N.E.; Alcazar Magana, A.; Lak, P.; Wright, K.M.; Quinn, J.; Stevens, J.F.; Maier, C.S.; Soumyanath, A. *Centella asiatica*: Phytochemistry and mechanisms of neuroprotection and cognitive enhancement. *Phytochem. Rev.* **2018**, *17*, 161–194. [CrossRef]

27. Edward, M.J. American Herbal Products Association’s Botanical Safety Handbook edited by M. McGuffin, C.; Hobbs, R. Upton, and A. Goldberg, CRC Press, Boca Raton, FL, 231 pages, 1997. $39.95. *J. Toxicol. Cutan. Ocul. Toxicol.* **2008**, *19*, 167–168. [CrossRef]

28. Randriamampionona, D.; Diallo, B.; Rakotoniriana, F.; Rabemanantsoa, C.; Cheuk, K.; Corbisier, A.M.; Mahillon, J.; Ratsimamanga, S.; El Jaziri, M. Comparative analysis of active constituents in *Centella asiatica* samples from Madagascar: Application for ex situ conservation and clonal propagation. *Fitoterapia* **2007**, *78*, 481–489. [CrossRef]

29. Alqahtani, A.; Tongkao-On, W.; Li, K.M.; Razmovski-Naumovski, V.; Chan, K.; Li, G.Q. Seasonal Variation of Triterpenes and Phenolic Compounds in Australian *Centella asiatica* (L.) Urb. *Phytochem. Anal.* **2015**, *26*, 436–443. [CrossRef]

30. Rafi, M.; Handayani, F.; Darusman, L.K.; Rohaeti, E.; Wahyu, Y.; Sulistiyani; Honda, K.; Putri, S.P. A combination of simultaneous quantification of four triterpenes and fingerprint analysis using HPLC for rapid identification of *Centella asiatica* from its related plants and classification based on cultivation ages. *Ind. Crops Prod.* **2018**, *122*, 93–97. [CrossRef]

31. Geran, R.I. Protocols for screening chemical agents and natural products against animal tumors and other biological systems. *Cancer Chemother. Rept.* **1972**, *3*, 17–27.

32. Micheau, J.; Marighetto, A. Acetylcholine and memory: A long, complex and chaotic but still living relationship. *Behav. Brain Res.* **2011**, *221*, 424–429. [CrossRef] [PubMed]

33. Li, S.-M.; Mo, M.-S.; Xu, P.-Y. Progress in mechanisms of acetylcholinesterase inhibitors and memantine for the treatment of Alzheimer’s disease. *Neuroimmunol Neuroinflammation* **2015**, *2*, 274.

34. Waldemar, G.; Hasslund, K.; Cowburn, R.F.; Bäckström, A.; Vasänge, M. The retinoic acid and brain-derived neurotrophic factor differentiated SH-SY5Y cell line as a model for Alzheimer’s disease-like tau phosphorylation. *Biochem. Biophys. Res. Commun.* **2004**, *319*, 993–1000. [CrossRef]

35. Pahlman, S.; Ruusala, A.I.; Abrahamsson, L.M.; Mattsson, M.E.; Esscher, T. Retinoic acid-induced differentiation of cultured human neuroblastoma cells: A comparison with phorbolester-induced differentiation. *Cell. Differ.* **1984**, *14*, 135–144. [CrossRef]

36. Paghans, S.P.; Ahmed, T.; Borwege, S.; Joyce, J.N. Terminally differentiated SH-SY5Y cells provide a model system for studying neuroprotective effects of dopamine agonists. *Neurotox. Res.* **2003**, *5*, 579–598. [CrossRef]
40. Korecka, J.A.; van Kesteren, R.E.; Blaas, E.; Spitzer, S.O.; Kamstra, J.H.; Smit, A.B.; Swaab, D.F.; Verhaagen, J.; Bossers, K. Phenotypic characterization of retinoic acid differentiated SH-SYSY cells by transcriptional profiling. *PLoS ONE* **2013**, *8*, e63862. [CrossRef]

41. Qiao, J.; Paul, P.; Lee, S.; Qiao, L.; Josifi, E.; Tiao, J.R.; Chung, D.H. PI3K/AKT and ERK regulate retinoic acid-induced neuroblastoma cellular differentiation. *Biochem. Biophys. Res. Commun.* **2012**, *424*, 421–426. [CrossRef]

42. Shipley, M.M.; Mangold, C.A.; Szpara, M.L. Differentiation of the SH-SYSY Human Neuroblastoma Cell Line. *J. Vis. Exp.* **2016**, *53193*. [CrossRef] [PubMed]

43. Saleem, H.; Ahmad, I.; Shahid, M.N.; Gill, M.S.A.; Nadeem, M.F.; Mahmood, W.; Rashid, I. In vitro acetylcholinesterase and butyrylcholinesterase inhibitory potentials of Jatropha gossypifolia plant extracts. *Acta Pol. Pharm. Drug Res.* **2016**, *73*, 419–423.

44. Orhan, I.E.; Atasu, E.; Senol, E.S.; Ozturk, N.; Demirci, B.; Das, K.; Sekeroglu, N. Comparative studies on Turkish and Indian *Centella asiatica* (L.) Urban (gotu kola) samples for their enzyme inhibitory and antioxidant effects and phytochemical characterization. *Ind. Crops Prod.* **2013**, *47*, 316–322. [CrossRef]

45. Niannuy, C.; Charoenchaitrakool, M.; Mayachiew, P.; Devahastin, S. Bioactive Compounds and Bioactivities of *Centella asiatica* (L.) Urban Prepared by Different Drying Methods and Conditions. *Dry. Technol.* **2013**, *31*, 2007–2015. [CrossRef]

46. Cuello, A.C. Early and Late CNS Inflammation in Alzheimer’s Disease: Two Extremes of a Continuum? *Trends Pharmacol. Sci.* **2017**, *38*, 956–966. [CrossRef]

47. Calsolaro, V.; Edison, P. Neuroinflammation in Alzheimer’s disease: Current evidence and future directions. *Alzheimer’s Dement.* **2016**, *12*, 719–732. [CrossRef]

48. Kettenmann, H.; Kirchhoff, F.; Verkhratsky, A. Microglia: New Roles for the Synaptic Stripper. *Neuron* **2013**, *77*, 10–18. [CrossRef]

49. Lyman, M.; Lloyd, D.G.; Ji, X.; Vizcaychipi, M.P.; Ma, D. Neuroinflammation: The role and consequences. *Neurosci. Res.* **2014**, *79*, 1–12. [CrossRef]

50. Dheen, S.T.; Kaur, C.; Ling, E.-A. Microglial Activation and its Implications in the Brain Diseases. *Curr. Med. Chem.* **2007**, *14*, 1189–1197. [CrossRef]

51. Park, J.; Min, J.S.; Kim, B.; Chae, U.B.; Yun, J.W.; Choi, M.S.; Kong, I.K.; Chang, K.T.; Lee, D.S. Mitochondrial ROS govern the LPS-induced pro-inflammatory response in microglia cells by regulating MAPK and NF-κB pathways. *Neurosci. Lett.* **2015**, *584*, 191–196. [CrossRef]

52. He, L.; He, T.; Farrar, S.; Ji, L.; Liu, T.; Ma, X. Antioxidants Maintain Cellular Redox Homeostasis by Elimination of Reactive Oxygen Species. *Cell. Physiol. Biochem.* **2017**, *44*, 532–553. [CrossRef] [PubMed]

53. Dilshara, M.G.; Lee, K.T.; Lee, C.M.; Choi, Y.H.; Lee, H.J.; Choi, I.W.; Kim, G.Y. New compound, 5-O-isoferuloyl-2-deoxy-D-ribofuranosyl-γ-lacton from Clematis mandshurica: Anti-inflammatory effects in lipopolysaccharide-stimulated BV2 microglial cells. *Int. Immunopharmacol.* **2015**, *24*, 14–23. [CrossRef]

54. Li, J.; Xu, X.; Cai, X.; Weng, Y.; Wang, Y.; Shen, Q.; Shi, X. Milk Fat Globule-Epidermal Growth Factor-Factor 8 Reverses Lipopolysaccharide-Induced Microglial Oxidative Stress. *Oxid. Med. Cell. Longev.* **2019**, 2019, 1–8. [CrossRef] [PubMed]

55. Park, E.; Chun, H.S. Melatonin Attenuates Manganese and Lipopolysaccharide-Induced Inflammatory Activation of BV2 Microglia. *Neurochem. Res.* **2017**, *42*, 656–666. [CrossRef]

56. Sasmita, A.O.; Ling, A.P.K.; Voon, K.G.L.; Koh, R.Y.; Wong, Y.P. Madecassoside activates anti-neuroinflammatory mechanisms by inhibiting lipopolysaccharide-induced microglial inflammation. *Int. J. Mol. Med.* **2018**, *41*, 3033–3040. [CrossRef]

57. Naidoo, D.B.; Chuturgoon, A.A.; Phulukdaree, A.; Guruprasad, K.P.; Satyamoorthy, K.; Sewram, V. Withania somnifera modulates cancer cachexia associated inflammatory cytokines and cell death in leukaemic THP-1 cells and peripheral blood mononuclear cells (PBMC’s). *BMC Complement. Altern. Med.* **2018**, *18*, 126. [CrossRef]

58. Yin, N.; Peng, Z.; Li, B.; Xia, J.; Wang, Z.; Yuan, J.; Fang, L.; Lu, X. Isoflurane attenuates lipopolysaccharide-induced acute lung injury by inhibiting ROS-mediated NLRP3 inflammasome activation. *Am. J. Transl. Res.* **2016**, *8*, 2033.

59. Batista, C.R.A.; Gomes, G.F.; Candelario-Jalil, E.; Fiebich, B.L.; de Oliveira, A.C.P. Lipopolysaccharide-Induced Neuroinflammation as a Bridge to Understand Neurodegeneration. *Int. J. Mol. Sci.* **2019**, *20*, 2293. [CrossRef]
60. Zakaria, R.; Wan Yaacob, W.M.H.; Othman, Z.; Long, I.; Ahmad, A.H.; Al-Rahbi, B. Lipopolysaccharide-Induced Memory Impairment in Rats: A Model of Alzheimer’s Disease. *Physiol. Res* 2017, 66, 553–565. [CrossRef] [PubMed]
61. Zhao, J.; Bi, W.; Xiao, S.; Lan, X.; Cheng, X.; Zhang, J.; Lu, D.; Wei, W.; Wang, Y.; Li, H.; et al. Neuroinflammation induced by lipopolysaccharide causes cognitive impairment in mice. *Sci. Rep.* 2019, 9, 5790. [CrossRef] [PubMed]
62. Nurul Huda, M.; Vasudevan, M.; Lim, S.; Sharmili, V.; Abu Bakar, A.M.; Kalavathy, R. Lactobacilli-fermented cow’s milk attenuated lipopolysaccharide-induced neuroinflammation and memory impairment in vitro and in vivo. *J. Dairy Res.* 2017, 84, 488–495.
63. Lykhmus, O.; Mishra, N.; Koval, L.; Kalashnyk, O.; Gergalova, G.; Uspekska, K.; Komisarenko, S.; Soreq, H.; Skok, M. Molecular Mechanisms Regulating LPS-Induced Inflammation in the Brain. *Front. Mol. Neurosci.* 2016, 9, 19. [CrossRef] [PubMed]
64. Lykhmus, O.; Voytenko, L.; Koval, L.; Mykhalskiy, S.; Kholin, V.; Peschana, K.; Zouridakis, M.; Tzartos, S.; Komisarenko, S.; Skok, M. α7 nicotinic acetylcholine receptor-specific antibody induces inflammation and amyloid β 42 accumulation in the mouse brain to impair memory. *PLoS ONE* 2015, 10, e0122706. [CrossRef] [PubMed]
65. Reale, M.; Iarlori, C.; Gambi, F.; Feliciani, C.; Salone, A.; Toma, L.; DeLuca, G.; Salvatore, M.; Conti, P.; Gambi, D. Treatment with an acetylcholinesterase inhibitor in Alzheimer patients modulates the expression and production of the pro-inflammatory and anti-inflammatory cytokines. *J. Neuroimmunol.* 2004, 148, 162–171. [CrossRef] [PubMed]
66. Hafiz, Z.Z.; Shamsuddin, N.; Mukhtar, S.M.; James, R.J.; Adenan, M.I. Anti-Oxidant Activities of Raw-Extract Centella Asiatica (RECA) on Lipopolysaccharide (LPS)-Induced Neuroinflammation Sprague Dawley Rats. *Int. J. Eng. Technol.* 2018, 7, 96–101. [CrossRef] [PubMed]
67. Cacciatore, I.; Baldassarre, L.; Fornasari, E.; Mollica, A.; Pinnen, F. Recent advances in the treatment of neurodegenerative diseases based on GSH delivery systems. *Oxid. Med. Cell. Longev.* 2012, 2012, 1–12. [CrossRef]
68. Ansari, M.A.; Scheff, S.W. Oxidative stress in the progression of alzheimer disease in the frontal cortex. *J. Neuropathol. Exp. Neurol.* 2010, 69, 155–167. [CrossRef]
69. Mischley, L.K.; Lau, R.C.; Shankland, E.G.; Wilbur, T.K.; Padowski, J.M. Phase Ib study of intranasal glutathione in Parkinson’s disease. *J. Parkinson’s Dis.* 2017, 7, 289–299. [CrossRef]
70. Richie, J.P.; Nichenametla, S.; Neidig, W.; Calcagnotto, A.; Haley, J.S.; Schell, T.D.; Muscat, J.E. Randomized controlled trial of oral glutathione supplementation on body stores of glutathione. *Eur. J. Nutr.* 2015, 54, 251–263. [CrossRef]
71. Kovacs-Nolan, J.; Rupa, P.; Matsui, T.; Tanaka, M.; Konishi, T.; Sauchi, Y.; Sato, K.; Ono, S.; Mine, Y. In vitro and ex vivo uptake of glutathione (GSH) across the intestinal epithelium and fate of oral GSH after in vivo supplementation. *J. Agric. Food Chem.* 2014, 62, 9499–9506. [CrossRef]
72. Allen, J.; Bradley, R.D. Effects of oral glutathione supplementation on systemic oxidative stress biomarkers in human volunteers. *J. Altern. Complement. Med.* 2011, 17, 827–833. [CrossRef]
73. Duffy, S.L.; Lagopoulos, J.; Hickie, I.B.; Diamond, K.; Graeber, M.B.; Lewis, S.J.G.; Naismith, S.L. Glutathione relates to neuropsychological functioning in mild cognitive impairment. *Alzheimer’s Dement.* 2014, 10, 67–75. [CrossRef] [PubMed]
74. Mandal, P.K.; Shukla, D.; Tripathi, M.; Ersland, L. Cognitive Improvement with Glutathione Supplement in Alzheimer’s Disease: A Way Forward. *J. Alzheimer’s Dis.* 2019, 1–5. [CrossRef] [PubMed]
75. Rafamantana, M.H.; Rozet, E.; Raelison, G.E.; Cheuk, K.; Ratsimamanga, S.U.; Hubert, P.; Quetin-Leclercq, J. An improved HPLC-UV method for the simultaneous quantification of triterpenic glycosides and aglycones in leaves of *Centella asiatica* (L.) Urb (APIACEAE). *J. Chromatogr. B Anal. Technol. Biomed. Life Sci.* 2009, 877, 2396–2402. [CrossRef] [PubMed]

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