

Review Article

# POTENTIAL OF NATURAL SUBSTANCE USAGE IN SOUTHEAST ASIA FOR MEMORY ENHANCEMENT: A REVIEW

Xinyi Zhu\*, Shun Kuroki\*, Jessebel V. Gadot\*\*, Aya Wada\*#

\*Department of Natural Product Chemistry and Biological Research, Delightex Pte. Ltd, Singapore, Singapore;

\*\*Centralized Analytical Testing Laboratory-Integrated Research Centers, University of Antique, Sibalom, Philippines

## Abstract

The use of natural substance-based supplements and treatments for mental wellness is increasingly gaining attention. Southeast Asia, with its rich heritage of medicinal practices and cultural reliance on natural remedies, presents a unique opportunity to explore such interventions. Delightex is actively collaborating with research partners in Southeast Asia to investigate natural substances that may enhance mental well-being and create enriching experiences. Memory, defined as the capacity to record, retain and recall sensory stimuli, events and information, is a fundamental aspect of mental health. Memory loss and Alzheimer's Disease (AD) are significant and growing concerns worldwide, particularly due to aging populations. Nootropics are generally well tolerated and typically mild. However, occasional complications can still occur. Hence, it is important to explore more natural alternatives for memory enhancement or treatment of memory loss. In this review, following an initial comprehensive literature search on mental well-being, we focused on memory improvement, identified and summarized 57 natural substances from 31 families with potential memory-enhancing effects. This review highlights their traditional use in Southeast Asia and examines the scientific evidence supporting their efficacy in enhancing memory and potential as nootropics alternatives. *ASEAN Journal of Psychiatry, Vol. 25 (7) September, 2024; 1-18.*

**Keywords:** Memory Enhancement; Memory Improvement; Memory Booster; Alzheimer's Disease (AD); Nootropics; Natural Substance; Southeast Asia

## Introduction

Memory encompasses the storage of information, the mental representation of experiences and the mental processes involved in learning, storing and retrieving information [1]. Human memory can be categorized into three essential types: Sensory memory, short-term memory and long-term memory [2]. Information processing begins in sensory memory, where it is held for a brief moment. It then transitions to short-term memory, which serves as the subject's working memory and may eventually be consolidated into long-term memory [2].

Memory loss, a central and highly feared consequence of aging, is also the signal symptom of dementia [3]. Memory issues, such as poor

recall, low retention, difficulty concentrating and weak analytical skills and are prevalent in today's world [4]. In some cases, these issues may escalate into more serious conditions, such as Alzheimer's Disease (AD) or schizophrenia [4,5]. Memory loss is often one of the initial symptoms of AD reported by patients with and their caretakers [6]. As of 2019, the global prevalence of AD was estimated at 57.4 million cases and this number could rise to 152.8 million by 2050 [7]. The increasing prevalence of AD, coupled with its strong impact on individuals and families, highlights the critical need for effective treatments and preventive strategies for AD and memory loss.

Nootropics, commonly referred to as smart drugs, encompass a diverse class of pharmacological agents that enhance cognitive

functions, including thinking, learning and memory, particularly in conditions where these capabilities are impaired [8]. Nootropics work by enhancing the brain's glucose and oxygen supply, exhibiting antihypoxic properties and protecting brain tissue from neurotoxicity [8-10]. Classical Nootropic Compounds include Deanol Dimethylaminoethanol (DMAE), Meclofenoxate, Nicergoline, Piracetam and Pyritinol [8].

Additionally, some substances exhibit simultaneous nootropic, hemorrheological and vasodilatory effects. Examples of these include vinpocetine, naftidrofuryl and dihydroergotoxine [8]. While there are benefit of nootropic drugs and nootropics are generally well tolerated and typically mild, there are still concerns over side effects, addictiveness and potential abuse of drugs [11,12]. Meanwhile, options to explore natural substances based on the traditional use have been regained interest as an alternative to those modern medicines. For example, traditional systems such as Ayurveda, have identified several herbs and plants with memory benefits [13,14]. Several species of plants have been selected for testing as nootropic agents because of their use in traditional medicine. Notable examples include *Ginkgo biloba*, *Panax ginseng*, *Paullinia cupana* and *Rhodiola rosea* [8].

Southeast Asia, with its rich biodiversity and traditional medicinal practices, offers a plethora of natural substances that could potentially enhance memory. However, there is a lack of comprehensive literature reviews summarizing the use of natural substances for memory enhancement in Southeast Asian regions. This review explores natural substances from this region, summarizing their taxonomic classification, the parts used, traditional applications and the scientific evidence supporting their potential for memory enhancement and their role as alternatives to nootropics. The bioactive compounds of highlighted substances are also discussed.

## Methodology

Through collaborations with Southeast Asian research partners, Delightex is exploring natural substances that may enhance mental well-being, in this review, we have refereed to works done by Delightex's research partners from Philippines (the University of Santo Tomas, the University of Antique) and Delightex's Vietnam research

team. The criteria of natural substance search using key words such as 'natural substances,' 'Southeast Asian countries' and 'medicinal plants' using databases and other online sources. Based on the broad search on mental wellness, we further narrowed specific natural substances with potential memory-enhancing effects, utilizing terms such as 'memory enhancement,' 'memory boosting', 'memory impairment,' 'memory loss' and 'nootropics.' This review discusses selected plants traditionally used or supported by scientific evidence for memory enhancement in Southeast Asian countries.

## Results and Discussion

Following comprehensive reviews of mental well-being enhancing natural substances conducted across Southeast Asia by Delightex's research partners, including the University of Santo Tomas, the University of Antique and Delightex's research team in Vietnam, we further focused on identifying natural substances with memory-enhancing properties. This investigation led to the identification of 57 natural substances (Table 1). The table provides detailed information for each substance, including its botanical family, scientific name, utilized part, traditional applications (if available) and supporting scientific evidence for its memory improvement effects.

In this review, the substances have been categorized and analyzed by botanical family to offer a more comprehensive understanding of their potential roles in promoting mental well-being. Figure 1 depicts the distribution of memory enhancing natural substances in Southeast Asia by families. 57 memory enhancing natural substances are from 31 families. Among them, 15.8% natural substances are from the Lamiaceae family, followed by the Fabaceae family (14.0%) and the Malvaceae family (5.3%).

### Plant families

The Lamiaceae family, also known as the mint or deadnettle family, comprises dicotyledonous flowering plants distributed globally. The enlarged Lamiaceae contains about 236 genera and about 6900 to 7200 species [15]. The Lamiaceae family is known for its high essential oil content and is also rich in polyphenolic compounds and terpenoids [16]. This review has listed 8 substances within this family (Table 1).

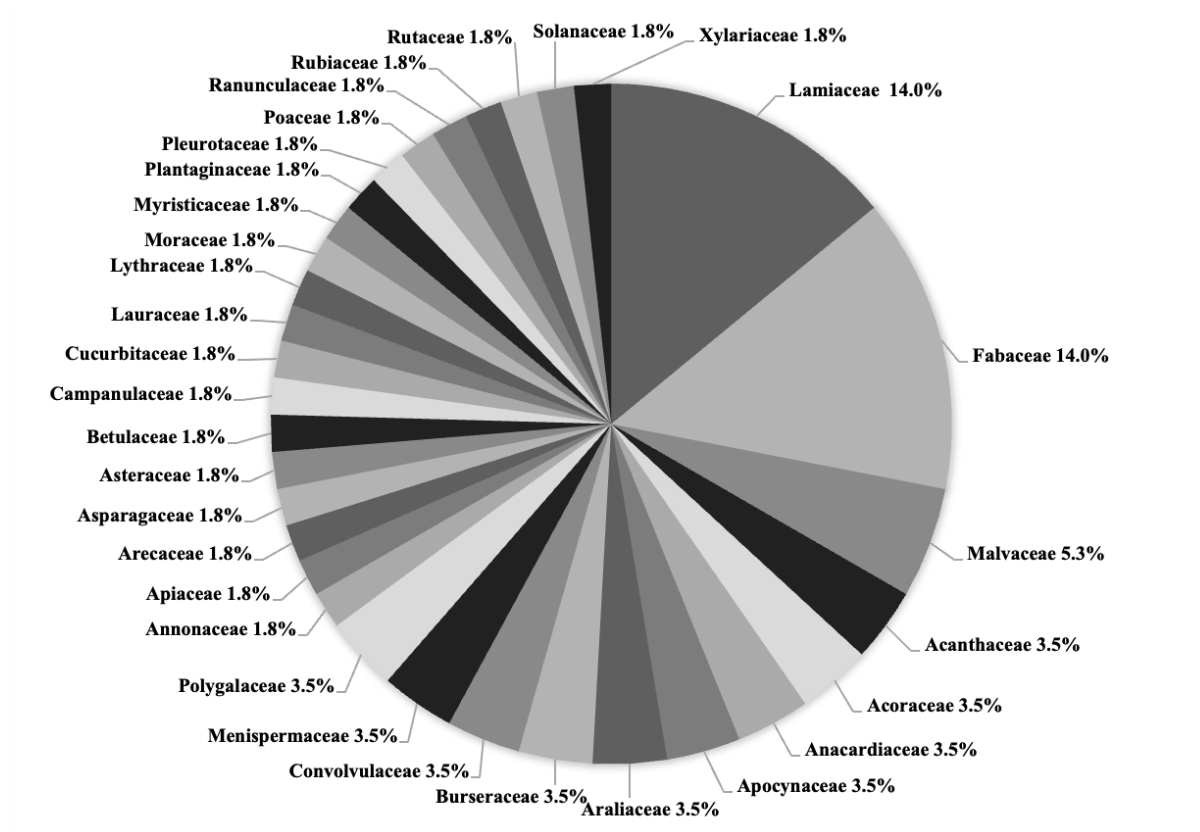


Figure 1. Pie chart categorizing memory-enhancing natural substances from Southeast Asia by family names.

*Hyptis suaveolens*, commonly known as pokok kemangi in Malaysia, amotan, suobkabayo, or loko-loko in the Philippines and kara or maeng lak kha in Thailand [17]. It is found in southeast countries such as Cambodia, Thailand, Indonesia, Malaysia, Vietnam and Philippines [18]. *H. suaveolens* is a rich source of medicinally significant phytochemicals, including essential oils, tannins, saponins, phenols, flavonoids, terpenoids, alkaloids and sterols. Main components identified in *H. suaveolens* essential oil include 1,8-Cineole (Eucalyptol), Sabinene,  $\beta$ -Caryophyllene and  $\beta$ -elemene [19]. These compounds belong to the terpene classes of monoterpene hydrocarbons and sesquiterpene hydrocarbons [19]. Traditionally, plant extracts of *H. suaveolens* were used as a memory aid [20]. In India, it was referred to as Chan or Wilaiti Tulsi, and the morning corn soup known as bate was believed to aid memory [21].

*Salvia rosmarinus*, commonly known as rosemary, is an evergreen, bushy shrub that thrives along the Mediterranean coast and in sub-Himalayan regions [22]. It is called dumero, romero and rosmiro in Philippines [23]. In Philippines, a leaf infusion of rosemary is utilized as eyewash for

mild catarrhal conjunctivitis, as vapor baths for the treatment of rheumatism, paralysis and early-stage catarrhs and for bathing postpartum women [23]. The main bioactive compounds in rosemary are triterpenes, phenolic diterpenes and phenolic acids including rosmarinic acid, carnosic acid, rosmanol, carnosol, ursolic acid and betulinic acid [24,25]. In traditional medicine, rosemary is believed to fortify the brain and refresh the memory [23]. The rosemary extracts have clinical effects on mood, learning, memory, pain, anxiety and sleep. Rosemary (500 mg administered twice daily for one month) has been shown to enhance both prospective and retrospective memory in a study involving 68 university students [26]. Additionally, Rosemary powder (750 mg), a dose comparable to typical culinary use, has demonstrated positive effects on memory speed and the efficiency of retrieving information from both episodic and working memory in 28 older adults (mean age: 75 years) [27]. Their finding suggests that rosemary may be a valuable memory enhancer.

*Vitex negundo*, commonly known as Chinese chaste tree, is native to tropical Eastern and

Southern Africa and Asia [28]. In Southeast Asia, it is indigenous to countries such as Bhutan, Cambodia, Indonesia, Malaysia, Myanmar, Philippines, Thailand and Vietnam [28]. Kanwal et al. reported that treatment with *V. negundo* aqueous extracts (300 mg/kg) for 5 days reduced scopolamine-induced amnesia in rats by enhancing memory and learning through antioxidant effects and decreasing AChE activity [29]. The administration of hydroalcoholic extract of *V. negundo* leaves at doses of 250 mg/kg and 500 mg/kg over 8 days significantly enhanced learning and memory in mice [30]. This improvement may be attributed to AChE inhibition, antioxidant activity and/or enhanced cholinergic transmission and the nootropic activity could be associated with the presence of compounds such as flavonoids, triterpenoids, phenolic acids and lignans in the extract [30].

**Family Fabaceae:** The Fabaceae family, commonly known as the legume, bean, or pea family, is the third largest family in the plant kingdom, comprising approximately 19,500 species, which constitute about 7% of all flowering plants [31]. This review has listed and summarized 7 substances within this family (Table 1).

*Arachis hypogaea*, peanut is one of the most consumed oil seeds worldwide. Peanuts have spread to other parts of the world from South America [32]. Peanuts are extensively utilized in the culinary traditions of Southeast Asia, particularly in countries such as Malaysia, Vietnam and Indonesia, where they are commonly processed into spicy sauces. In the Philippines, peanut is a key ingredient in the traditional dish kare-kare and fried shelled peanuts are widely consumed as an affordable snack throughout the Philippines. Regular peanut and peanut butter consumption may improve memory function and stress response in healthy young adults. These effects appear to be linked to the intake of peanut polyphenols [33]. In this 6-month randomized controlled trial, participants consuming 25 g/day of skin-roasted peanuts or consuming 32 g/day of peanut butter have shown improved immediate memory after the intervention [33].

*Clitoria ternatea*, known as butterfly pea and blue pea in various countries is originated from tropical Asia countries and it is widely available in the Asian regions [34,35]. As a traditional Ayurveda medicine as a brain tonic, memory and nootropic herb, butterfly pea flower plays an important role in nervine medicine as well as improving

brain system and boosting memory [35]. Chronic oral administration of butterfly pea root extract for 28 days at doses of 200 mg/kg and 300 mg/kg effectively restored memory impairments in rats induced by Chronic Cerebral Hypoperfusion (CCH) [36].

*Pueraria candollei* var. *mirifica* is endemic to Thailand and is widely distributed in the country's deciduous forests, particularly at altitudes of 300 to 800 meters. It is called kwao khrua khaw (various spellings) in the Thai language [37]. It is most commonly and abundantly found in the northern, western and northeastern regions of Thailand [38,39]. The main bioactive phytochemicals of *P. mirifica* are phytoestrogens including isoflavonoids, coumestans and chromenes [40]. Traditionally, pounding the roots of *P. mirifica* and mixing them with cow's milk is believed to enhance memory [41]. *P. mirifica* extracts have demonstrated beneficial effects on cognitive deficits associated with menopause or estrogen depletion, primarily through its phytoestrogenic and antioxidant properties [42]. In the same study, it was found that PM extracts at a daily dose of 25 mg/kg exerted anti-dementia effects that were almost equivalent to the effects elicited by 17 $\beta$ -estradiol at 1  $\mu$ g/kg [42].

**Family Malvaceae:** The Malvaceae, commonly known as the mallow family, is a group of dicotyledonous flowering plants comprising approximately 244 genera and around 4,225 species. These species are distributed across regions ranging from tropical to temperate climates [43]. This review has explored 3 substances within this family (Table 1).

*Abelmoschus manihot* is in South-East Asia it is cultivated particularly in the eastern parts of Indonesia and in Papua New Guinea [44]. In North Sulawesi, Indonesia, this plant is known as "gedi" and its leaves are a key ingredient in preparing porridge, a distinctive gourmet dish in North Sulawesi cuisine [45]. More than 128 phytochemical compounds have been extracted and identified from the flowers, seeds, stems and leaves of *A. manihot* [46]. These compounds primarily include flavonoids, amino acids, nucleosides, polysaccharides, organic acids, steroids and volatile oils [46]. The hydroalcoholic extract of *A. manihot* flowers has been shown to alleviate learning and memory impairments caused by sleep deprivation in mice [47]. This effect may be attributed to its antioxidant capacity



and enhanced BDNF/TrkB/GluR1 levels in the hippocampal memory [47].

*Thespesia populnea* commonly known as ‘Indian tulip tree. The species is native to Australia, China and India but can also be found on rocky coasts such as in Malaysia and other countries in SEA including Cambodia, Indonesia, Philippines and Thailand [48]. *T. populnea* contains glycosides such as quercetin, gossypol,  $\beta$ -sitosterol and sesquiterpene, lupenone and lupeol [49,50]. Thespesenone and dehydrooxoperezinone-6-methyl ether have been isolated from the red hardwood, while alanine, arginine, methionine and tryptophan have been identified in the seeds [51,52]. The bark of *T. populnea* shows potential as a memory-enhancing agent. Oral administration of ethanol bark extract of *T. populnea* at doses of 200 mg/kg and 400 mg/kg for 7 days significantly improved memory in both young and aged mice [53].

**Other families:** *Areca catechu*, belongs to the Arecaceae family, a species of palm native to the Philippines, is primarily grown for its areca nuts [54,55]. It is known by various common names in English, including betelnut palm, arecanut, betel palm, betel-nut, supari palm, pinang palm [56]. It is extensively cultivated in several Southeast Asian countries, including Malaysia, Indonesia, Cambodia, Laos, Myanmar, Thailand and Vietnam [57]. *A. catechu* contains alkaloids, polyphenols, polysaccharides, triterpenes, steroids, fatty acids and other components [58]. These compounds demonstrate a variety of bioactive functions, such as antibacterial, deworming, antiviral, antioxidant, anti-inflammatory and anti-tumour effects [58]. Extracts of arecanut have been reported to improve memory. Oral administration

of methanol extracts of arecanut at a dose of 500 mg/kg for 21 days significantly improved memory and learning in rats [59]. Arecoline, the major alkaloid of arecanut, has been reported to enhance memory in both laboratory animals and humans, as summarized by Bhat et al [60]. For example, in humans, a 4 mg injection of arecoline significantly enhanced serial learning and reversed impaired learning behaviour induced by scopolamine, a cholinergic antagonist [61]. Additionally, a 4 mg dose of arecoline significantly improved picture recognition in Alzheimer’s patients [62].

*Bacopa monnieri* also known as waterhyssop, thyme-leafed gratiola, bacopa and brahmi, is a perennial, creeping herb from the Plantaginaceae family. Bacopa is a medicinal herb native to South and Southeast Asia [63]. Ayurvedic medicine classifies Bacopa as a ‘medhya rasayana’ which refers to a category of herbs thought to enhance mental health, boost memory and intellect and support rejuvenation and longevity [64,65]. Preliminary clinical studies have suggested that Bacopa has the potential in ameliorating cognitive disorders, as well as prophylactic reduction of oxidative damage, neurotransmitter modulation and cognitive enhancement in healthy individuals [66]. Additionally, across six human randomized controlled trials were all conducted over 12 weeks have reported that 300 mg to 450 mg bacopa extract per day have improved performance in the domain of memory free recall [67]. *B. monnieri* is rich in a variety of distinct bioactive phytoconstituents, each offering a range of benefits. Among them, Bacopaside XI, bacopaside I and bacopasaponin C, administered at a concentration of 50 mg/kg, demonstrated nootropic effects and reduced memory impairment in mice [68,69].

**Table 1. List of natural substances in Southeast Asia countries and their usage for memory enhancement.**

Family	Scientific name	Part	Scientific evidence and traditional knowledge (if available) on memory enhancement	References
Acanthaceae	<i>Peristrophe bicalyculata</i>	Leaf	The methanolic leaf extract improves memory performance in a rat model of type 2 diabetes mellitus	[70]
	<i>Rhinacanthus nasutus</i>	Leaf, root	The ethanol extracts from the leaves and roots have demonstrated a dose-dependent ability to reduce neuron cell death induced by both glutamate and amyloid- $\beta$ exposure	[71]

Acoraceae	<i>Acorus calamus</i>	Root, Rhizome	<p>In Ayurvedic medicine, rhizomes are used for treatment of epilepsy, schizophrenia and memory disorders</p> <p>The memory impairment in groups that received aqueous fraction at 600 mg/kg dose, was less than control</p> <p>Alpha-asarone, as a major component of the <i>A. calamus</i> ameliorates memory deficit in mice</p>	[72-74]
	<i>Acorus gramineus</i>	Rhizome	<p>Across 34 studies involving 1,431 animals, extracts or active components was found to significantly enhance learning and memory function</p>	[75]
Anacardiaceae	<i>Rhus verniciflua</i>	Bark	<p>Active component fesetin reduced memory deficits induced by scopolamine through activation of the CREB–BDNF pathway</p>	[76]
	<i>Spondias mombin</i>	Leaf	<p>Oral administration of aqueous extracts at 400mg/kg for 28 days improved learning and memory capabilities in rats</p>	[77]
Annonaceae	<i>Annona squamosa</i>	Leaf	<p>The leaf extract and its isolated constituent-anonaine demonstrated memory boosting and memory regaining effects in rats</p>	[78]
Apiaceae	<i>Centella asiatica</i>	N/A	<p>An Ayurvedic herb used to enhance memory and nerve function</p> <p>Water extracts (200 mg/kg/day, 2 weeks) attenuated amyloid-<math>\beta</math>-associated behavioral abnormalities in the Tg2576 mouse, a murine model of AD</p>	[79]
Apocynaceae	<i>Catharanthus roseus</i>	Leaf, stem and root	<p>Aqueous extract of leaf, stem and root have been shown to effectively inhibit AchE <i>in-vitro</i> and active compound serpentine displayed a strong activity against Acetylcholinesterase (AchE) <i>in-vitro</i>, suggesting a potential as a therapeutic agent for AD</p>	[80]
	<i>Tabernaemontana divaricata</i>	Root	<p>The ethanolic root extract acts as a reversible Acetylcholinesterase Inhibitor (AChE-I) and may offer potential as a novel therapeutic agent for AD</p>	[81]

Araliaceae	<i>Eleutherococcus senticosus</i>	Leaf	Oral administration of the water extract for 17 days significantly enhanced object recognition memory pure compounds ciwujianoside C3, eleutheroside M, ciwujianoside B were administered orally for 17 days to normal mice and significantly enhanced object recognition memory	[82]
	<i>Panax ginseng</i>	Root	Oral administration of fermented cultured wild ginseng root extract (HLJG0701-β) (250 mg/kg, 8 weeks) resulted in memory improvement in mice	[83]
Arecaceae	<i>Areca catechu</i>	Fruit, seed, nut	Refer to the text above	
Asparagaceae	<i>Asparagus racemosus Willd</i>	Root	Administration of extract (200 mg/kg, 7 days) has shown a promising memory-enhancing effect in both young and aged mice	[84]
Asteraceae	<i>Achillea biebersteinii</i>	Aerial Part (stem, leaf and flower)	Inhalation of essential oil (1%, 3%) for 21 days significantly increased spontaneous alternation percentage, and decreased the working memory errors and reference memory errors in rats	[85]
Betulaceae	<i>Betula platyphylla</i>	Bark	Oral treatment using bark (BPB-316) significantly attenuated amyloid-β-induced memory impairment in mice	[86]
Burseraceae	<i>Commiphora caudata</i>	Leaf	Ethanollic leaves extract (200 mg/kg, 400 mg/kg, 27 days) enhanced learning and memory activity: Transfer Latency (TL), Time Taken To Reach Reward Chamber (TRC) and Swim Latency (SL) in comparison to scopolamine treated rats	[87]
	<i>Garuga pinnata</i>	Bark	Contains bioactive phytoconstituents, including alkaloids, tannins, phenols, and flavonoids, which possess free radical scavenging properties and memory-enhancing capabilities	[88]
Campanulaceae	<i>Platycodon grandiflorus</i>	Root	Administration of root to mice resulted in increased spontaneous alternation in the Y-maze test and promoted synaptogenesis in the hippocampus	[89]
Convolvulaceae	<i>Convolvulus pluricaulis</i>	Whole plant	Administration of extract (200 mg/kg, 7 days) has shown a promising memory-enhancing effect in both young and aged mice	[84]
	<i>Ipomoea batatas</i>	Tuber	Anthocyanins extracted from purple sweet potato demonstrate memory-enhancing effects, potentially linked to their antioxidant properties	[90]

Cucurbitaceae	<i>Cucurbita maxima</i>	Seed	Ethanollic seed extract (50 mg/kg, 100 mg/kg and 200 mg/kg) exhibited anxiolytic and antidepressant effects with memory improvement in mice	[91]
Fabaceae	<i>Mimosa pudica</i>	Leaf	Behavioral effects on learning and memory were augmented by leaf ethyl acetate extracts (200 mg/kg and 400 mg/kg)	[92]
	<i>Albizia adianthifolia</i>	Leaf	Aqueous extract (150 mg/kg, 300 mg/kg) plays an important role in spatial memory formation, especially on working and reference memories	[93]
	<i>Arachis hypogaea</i>	Seed, leaf	Refer to the text above	
	<i>Cassia obtusifolia</i> / <i>Senna obtusifolia</i>	Seed	Ethanollic seed extract (50 mg/kg) attenuates memory impairment induced by scopolamine or 2VO and that these effects are mediated by enhancing the cholinergic nervous system <i>via</i> acetylcholinesterase inhibition	[94]
	<i>Clitoria ternatea</i>	Root, whole plant	Refer to the text above	
	<i>Glycyrrhiza glabra</i>	Root	Aqueous extracts (150 mg/kg ) significantly improved learning and memory of mice	[95]
	<i>Piliostigma thonningii</i>	Stem, Bark	Aqueous and methanolic stem bark extract showed remarkable cognitive-enhancing activities which were reflected in significantly shorter transfer latencies, navigation distances, longer time spent in platform quadrant, and lower MDA levels compared to the negative control mice	[96]
	<i>Pueraria candollei</i> var. <i>mirifica</i>	Roots, tuber	Refer to the text above	



Lamiaceae	<i>Hyptis suaveolens</i>	Whole plant (Except root)	Refer to the text above	
	<i>Melissa officinalis</i>	Leaf	Administration of ethanolic extract (200 mg/kg) could significantly enhance learning and memory in memory-impaired rats	[97]
	<i>Prunella vulgaris var. lilacina</i>	Flower	Ethanolic flower extract (25 mg/kg) significantly shortened escape latencies in training-trials. Furthermore, swimming times within the target zone during the probe-trial were significantly increased as compared with scopolamine-treated mice	[98]
	<i>Salvia rosmarinus</i>	Leaf	Refer to the text above	
	<i>Salvia miltiorrhiza</i>	Root	Subchronic administration of root extract (200 mg/kg) led to an improvement of long-term memory of rats	[99]
	<i>Salvia officinalis</i>	Leaf	<i>S. officinalis</i> aroma group performed significantly better than the control group on the quality of memory and secondary memory primary outcome factors from the test battery	[100]
	<i>Scutellaria baicalensis</i>	Root	Extracts (30 mg/kg) improved spatial memory functions and rescued neuronal cells immune-reactive to ChAT and the NMDA receptor subunit, NR2A, in the hippocampus of memory deficient rat model	[101]
	<i>Vitex negundo</i>	Leaf	Refer to the text above	
Lauraceae	<i>Cinnamomum zeylanicum</i>	Bark	Aqueous bark extract-treated animals exhibited an improved discrimination between a familiar object and a novel object, indicating the reversal of extract induced memory impairment. Extract also restored alteration in AChE activity and oxidative stress parameters in both brain parts	[102]
Lythraceae	<i>Lawsonia inermis</i>	Leaf	Extract at doses of 200 mg/kg and 400 mg/kg significant increased the inflexion ratio in elevated plus maze and increase in percentage alternation in Y-maze model compared to negative control animals	[103]

Malvaceae	<i>Abelmoschus manihot</i>	Flower	Refer to the text above	
	<i>Abelmoschus moschatus</i>	Seed	Oral administration of ethanolic seed extract (100 mg/kg, 200 mg/kg) for seven days demonstrated a dose-dependent improvement in memory in young mice and effectively reversed diazepam-induced memory deficits	[104]
	<i>Thespesia populnea.</i>	Bark	Refer to the text above	
Menispermaceae	<i>Cissampelos pariera</i>	Root	The dose of 400mg/kg of CPE significantly improved learning and memory of mice	[105]
	<i>Tiliacora triandra</i>	Leaf	Leaf extract at doses of 300 mg/kg and 600 mg/kg significantly enhances spatial learning and learning flexibility. Only the 300 mg/kg dose showed a significant improvement in spatial memory	[106]
Moraceae	<i>Ficus racemosa</i>	Bark	Administration of the extract at two levels, 250 mg/kg and 500 mg/kg extract resulted in significant reduction in transfer latency on elevated plus-maze, which was used as an exteroceptive behavioural model to evaluate memory in rats	[107]
Myristicaceae	<i>Myristica fragrans</i>	Seed	Extract at the lowest dose of 5 mg/kg P.O. administered for 3 successive days significantly improved learning and memory of young and aged mice	[108]
Plantaginaceae	<i>Bacopa monnieri</i>	Whole plant/leaf	Refer to the text above	
Poaceae	<i>Cymbopogon citratusx</i>	Leaf	After the inhalation, the lemongrass essential oil enhanced their cognitive performance for the domains of the continuity of attention and the quality of memory, whereas the mood in terms of alertness and calmness was also increased	[109]
Pleurotaceae	<i>Pleurotus eryngii</i>	Fruiting Body	The ethanol extract of <i>P. ryngii</i> exhibits estrogen-like effects, providing potential benefits for alleviating depression and improving memory impairment in rats  Feeding with <i>P. eryngii</i> for six weeks has promoted memory and learning capacity in an AD mouse model	[110, 111]

Polygalaceae	<i>Polygala japonica</i>	Root	A traditional medicine for treatment of variety of ailments, including anti-inflammatory, antibacterial, sedative and nootropic agent  Compound saponins can improve the learning and memory in mice	[112]
	<i>Polygala tenuifolia</i>	Root	The impaired spatial memory of the aged mice was partly reversed by extract (100 mg/kg and 200 mg/kg) as compared with the aged control mice. In stepdown tests, the nonspatial memory of the aged mice was improved by extract (100 mg/kg and 200 mg/kg)	[113]
Ranunculaceae	<i>Nigella sativa</i>	Seed	Hydro-alcoholic extract of NS improved the LPS-induced learning and memory impairments in rats	[114]
Rutaceae	<i>Lunasia amara</i>	Bark	A traditional Ayurveda medicine as a brain tonic, memory and intelligence enhancer	N/A
Solanaceae	<i>Withania somnifera</i>	Root	Known as Ashwagandha, it has been traditionally used in Ayurvedic medicine as a substance to strengthen the nervous system  Root extract alleviated hypobaric hypoxia -induced memory impairment and neurodegeneration in the hippocampus by modulating corticosterone levels through nitric oxide pathways	[115,116]
Xylariaceae	<i>Xylaria nigripes</i>	Mycelium	Water extracts (300 mg/kg/day, 6 weeks) improved Rapid Eye Movement Sleep Deprivation (REMSD)-induced memory impairment in rats	[117]
Rubiaceae	<i>Morinda citrifolia</i>	Fruit	Extract prevented memory impairment induced by amyloid- $\beta$ peptide in mice	[118,119]

## Conclusion

Southeast Asia, with its rich tradition of medicinal practices and reliance on natural remedies, offers a diverse array of natural substances with potential memory-enhancing properties. This review has summarized 57 natural substances that originate from, are cultivated in, or are traditionally used in the region, many of which could serve as alternatives to nootropics or have already been incorporated into nootropic formulations. These substances span 31 different families such as the Lamiaceae, Fabaceae and Malvaceae families. The review discusses about the traditional knowledge and scientific evidence related to their memory-enhancing effects. While most evidence is derived from animal studies, human clinical studies of effect of rosemary (*S. rosmarinus*) peanut (*A. hypogaea*), arecanut (*Areca catechu*) and Bacopa (*Bacopa monnieri*) are also discussed. This review also discusses the bioactive compounds of the highlighted substances. However, the specific compounds responsible for memory enhancement and their underlying pharmacological mechanisms in many natural substances remain unclear. Further research is needed to elucidate these mechanisms, identify the compounds responsible for memory enhancement, evaluate their safety profiles and explore the potential of these natural substances as alternatives to existing nootropics.

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## References

1. Radvansky GA. Human memory. Routledge. 2021.
2. Atkinson RC, Shiffrin RM. Human memory: A proposed system and its control process. Psychol Learn Motiv-Adv Res Theory. 1968;2:189-195.
3. Andrew B, Paul S. Memory loss: A practical guide for clinicians. 2011.
4. Halder S, Anand U, Nandy S, Oleksak P, Qusti S, et al. Herbal drugs and natural bioactive products as potential therapeutics: A review on pro-cognitive and brain booster's perspectives. Saudi Pharm J. 2021;29(8):879-907.
5. Gur RC, Gur RE. Memory in health and in schizophrenia. Dialogues Clin Neurosci. 2013;15(4):399-410.
6. Jahn H. Memory loss in Alzheimer's disease. Dialogues in clinical neuroscience. 2013;15(4):445-454.
7. Nichols E, Steinmetz JD, Vollset SE, Fukutaki K, Chalek J, et al. Estimation of the global prevalence of dementia in 2019 and forecasted prevalence in 2050: An analysis for the Global Burden of Disease Study 2019. Lancet Public Health. 2022;7(2):e105-e125.
8. Malik M, Tlustos P. Nootropics as cognitive enhancers: Types, dosage and side effects of smart drugs. Nutrients. 2022;14(16):3367.
9. Joshi Pranav C. A review on natural memory enhancers (Nootropics). Unique Journal of Engineering and Advanced Sciences. 2013;1(01):8-18.
10. Malik R, Sangwan A, Saihgal R, Paul Jindal D, Piplani P. Towards better brain management: nootropics. Curr Med Chem. 2007;14(2):123-131.
11. Schifano F, Catalani V, Sharif S, Napoletano F, Corkery JM, et al. Benefits and harms of 'smart drugs'(nootropics) in healthy individuals. Drugs. 2022;82(6):633-647. 1.
12. Garcia HS, Reula LM, Fernandez AP, Sanchez VP, Lopez NO, et al. Nootropics: Emergents drugs associated with new clinical challenges. Eur Psychiatr. 2017;41(S1):s877-s878. 1.
13. Farooqui AA, Farooqui T, Madan A, Ong JH, Ong WY. Ayurvedic medicine for the treatment of dementia: Mechanistic aspects. Evid Based Complement Alternat Med. 2018;2018(1):2481076.
14. Mehla J, Gupta P, Pahuja M, Diwan D, Diksha D. Indian medicinal herbs and formulations for Alzheimer's disease, from traditional knowledge to scientific assessment. Brain Sci. 2020;10(12):964.
15. Xu Z, Deng M. Identification and Control of Common Weeds: Volume 2. Dordrecht: Springer Netherlands; 2017.
16. Abdelhalim A, Hanrahan J. Biologically active compounds from Lamiaceae family: Central

- nervous system effects. Studies in natural products chemistry. 2021;68:255-315.
17. Seidemann J. World spice plants. Berlin, Heidelberg: Springer Berlin Heidelberg. 2005.
  18. Li R, Tang G, Liu X, Li J, Wang D, et al. An ethnopharmacological review of *Hyptis suaveolens* (L.) Poit. Trop J Pharm Res. 2020;19(7):1541-1550. 1.
  19. Aguele FO, Oke EO, Abam FI, Nnabodo D, Agbana AS. Biological and antioxidant activities, extraction methodology and prospects of essential oil from *Hyptis suaveolens* (L.): A review. Clean Eng Technol. 2023;17:100685. 1.
  20. Sharma PP, Roy RK, Anurag DG, Vipin KS. *Hyptis suaveolens* (L.) poit: A phyto-pharmacological review. Int J Chem Pharm Sci. 2013;4(1):1-11.
  21. Perusomula R, Thamannagari S, Paturi G, Alluri R. Evaluation of anti-arthritis activity of *Hyptis suaveolens* seeds in rats. GSC Biol Pharm Sci. 2024;27(3):077-98. 1.
  22. Rahbardo MG, Hosseinzadeh H. Therapeutic effects of rosemary (*Rosmarinus officinalis* L.) and its active constituents on nervous system disorders. Iran J Basic Med Sci. 2020;23(9):1100-1112.
  23. De Guzman CC, Siemonsma JS. Plant resources of South-East Asia. Backhuys Publ. 1999.
  24. Borrás-Linares I, Stojanović Z, Quirantes-Piné R, Arráz-Román D, Švarc-Gajić J, et al. *Rosmarinus officinalis* leaves as a natural source of bioactive compounds. Int J Mol Sci. 2014;15(11):20585-20606.
  25. Okamura N, Fujimoto Y, Kuwabara S, Yagi A. High-performance liquid chromatographic determination of carnosic acid and carnosol in *Rosmarinus officinalis* and *Salvia officinalis*. J Chromatogr A. 1994;679(2):381-386.
  26. Nematollahi P, Mehrabani M, Karami-Mohajeri S, Dabaghzadeh F. Effects of *Rosmarinus officinalis* L. on memory performance, anxiety, depression, and sleep quality in university students: A randomized clinical trial. Complement Ther Clin Pract. 2018;30:24-28.
  27. Pengelly A, Snow J, Mills SY, Scholey A, Wesnes K, et al. Short-term study on the effects of rosemary on cognitive function in an elderly population. J Med Food. 2012;15(1):10-17.
  28. *Vitex negundo* L. Germplasm resources information network. Agricultural Research Service, United States Department of Agriculture. 2011.
  29. Kanwal A, Mehla J, Kuncha M, Naidu VG, Gupta YK, et al. Anti-amnesic activity of *Vitex negundo* in scopolamine induced amnesia in rats. Pharmacology & Pharmacy. 2010;1(01):1.
  30. Otari KV, Bichewar OG, Shete RV, Upasani CD. Effect of hydroalcoholic extract of *Vitex negundo* Linn. Leaves on learning and memory in normal and cognitive deficit mice. Asian Pac J Trop Biomed. 2012 ;2(1):S104-S1011.
  31. Raj SP, Solomon PR, Thangaraj B, Raj SP, Solomon PR, et al. Fabaceae. Biodiesel from Flowering Plants. 2022:291-363.
  32. Çiftçi S, Suna GÜ. Functional components of peanuts (*Arachis hypogaea* L.) and health benefits: A review. Future foods. 2022;5:100140.
  33. Parilli-Moser I, Domínguez-López I, Trius-Soler M, Castellví M, Bosch B, et al. Consumption of peanut products improves memory and stress response in healthy adults from the ARISTOTLE study: A 6-month randomized controlled trial. Clin Nutr. 2021;40(11):5556-5567.
  34. Cook BG, Pengelly BC, Brown SD, Donnelly JL, Eagles DA, et al. Tropical Forages: An interactive selection tool. 2005.
  35. Jamil N, Zairi MN, Nasim NA, Pa'ee F. Influences of environmental conditions to phytoconstituents in *Clitoria ternatea* (butterfly pea flower)-A review. J Sci Technol. 2018;10(2).
  36. Damodaran T, Cheah PS, Murugaiyah V, Hassan Z. The nootropic and anticholinesterase activities of *Clitoria ternatea* Linn. root extract: Potential treatment for cognitive decline. Neurochem Int. 2020;139:104785.
  37. Kongkaew C, Scholfield NC, Dhippayom T, Dilokthornsakul P, Saokaew S, et al. Efficacy and safety of *Pueraria candollei* var. *mirifica* (Airy Shaw & Suvat.) Niyomdham for menopausal women: A systematic review of clinical trials and the way forward. J Ethnopharmacol. 2018;216:162-174.
  38. Malaivijitnond S. Medical applications of phytoestrogens from the Thai herb *Pueraria*



- mirifica. *Front Med.* 2012;6(1):8-21.
39. Cherdshewasart W, Kitsamai Y, Malaivijitnond S. Evaluation of the estrogenic activity of the wild *Pueraria mirifica* by vaginal cornification assay. *J Reprod Dev.* 2007;53(2):385-393.
40. Chansakaow S, Ishikawa T, Seki H, Sekine K, Okada M, et al. Identification of Deoxymiroestrol as the Actual Rejuvenating Principle of “Kwao Keur”, *Pueraria mirifica*. The Known Miroestrol May Be an Artifact. *J Nat Prod.* 2000;63(2):173-175. 1.
41. Passwater R. *Pueraria mirifica*: Just for menopause or the herb of the decade? Part I, *Whole Foods Magazine.* 2007.
42. Chulikhit Y, Sukhano W, Daodee S, Putalun W, Wongpradit R, et al. Effects of *Pueraria candollei* var *mirifica* (Airy Shaw and Suvat.) Niyomdham on ovariectomy-induced cognitive impairment and oxidative stress in the mouse brain. *Molecules.* 2021;26(11):3442.
43. Xu Z, Deng M. Identification and Control of Common Weeds: Volume 2. Dordrecht: Springer Netherlands; 2017.
44. Wien HC. Plant resources of South-East Asia. No. 8. vegetables: JS Siemonsma and Kasem Paluek (Editors), Pudoc, Wageningen, 1993, 412 pp., ISBN 90-220-1058-9. *Sci Hortic.* 1995;62(3):202-203.
45. Mandey JS, Soetanto H, Sjoftan O, Tulung B. Genetics characterization, nutritional and phytochemicals potential of gedi leaves (*Abelmoschus manihot* (L.) Medik) growing in the North Sulawesi of Indonesia as a candidate of poultry feed. *Res J Biol.* 2014;4(2):1276-1286.
46. Luan F, Wu Q, Yang Y, Lv H, Liu D, et al. Traditional uses, chemical constituents, biological properties, clinical settings, and toxicities of *Abelmoschus manihot* L.: A comprehensive review. *Front Pharmacol.* 2020;11:1068.
47. Zhou HR, Wu JR, Bei L, Wang BX, Xu H, et al. Hydroalcoholic extract from *Abelmoschus manihot* (Linn.) Medicus flower reverses sleep deprivation-evoked learning and memory deficit. *Food Funct.* 2020;11(10):8978-8986.
48. Orwa C. Agroforestry Database: A tree reference and selection guide, version 4.0. 2009.
49. Khare CP. *Indian Medicinal Plants.* 2007.
50. Rastogi RM. Compendium of Indian medicinal plants. Central Drug Research Institute, Lucknow, India. 1990;1:388-389.
51. Kumar SP, Bhushan MS, Trishna D, Arun M. Phytochemical investigation and antimicrobial activity of stem bark extracts of *Thespesia populnea*. *Int J Pharm Res.* 2010;1(1):24-29.
52. Puckhaber LS, Stipanovic RD. Thespesenone and dehydroxoperezinone-6-methyl ether, new sesquiterpene quinones from *Thespesia populnea*. *J Nat Prod.* 2004;67(9):1571-1573.
53. Vasudevan M, Parle M. Pharmacological actions of *Thespesia populnea* relevant to Alzheimer's disease. *Phytomedicine.* 2006;13(9-10):677-687.
54. Plants of the World Online. The Royal Botanic Gardens. 2024.
55. Zumbroich TJ. The origin and diffusion of betel chewing: A synthesis of evidence from South Asia, Southeast Asia and beyond. *E- Indian J Med.* 2008;1(3):87.
56. Orwa C. Agroforestry Database: *Areca catechu* Arecaceae L, version 4.0. 2009.
57. Staples GW, Bevacqua RF. *Areca catechu* (betel nut palm). Species profiles for Pacific Island agroforestry. 2006;1(13):1-9.
58. Sun H, Yu W, Li H, Hu X, Wang X. Bioactive Components of Areca Nut: An Overview of Their Positive Impacts Targeting Different Organs. *Nutrients.* 2024;16(5):695.
59. Joshi M, Gaonkar K, Mangoankar S, Satarkar S. Pharmacological investigation of *Areca catechu* extracts for evaluation of learning, memory and behavior in rats. *Int Curr Pharm J.* 2012;1(6):128-132.
60. Bhat SK, Ashwin D, Mythri S, Bhat S. Arecanut (*Areca catechu* L) decreases Alzheimer's disease symptoms: Compilation of research works. *J Med Plants Stud.* 2017;5(5):4-9.
61. Sitaram N, Weingartner H, Gillin JC. Human serial learning: Enhancement with arecholine and choline impairment with scopolamine. *Science.* 1978;201(4352):274-276.
62. Christie JE, Shering A, Ferguson J, Glen AI. Physostigmine and arecoline: Effects of intravenous infusions in Alzheimer presenile

- dementia. Br J Psychiatry. 1981;138(1):46-50.
63. Prasansuklab A, Brimson JM, Tencomnao T. Potential Thai medicinal plants for neurodegenerative diseases: A review focusing on the anti-glutamate toxicity effect. J Tradit Complement Med. 2020 ;10(3):301-308.
64. GK S, MS 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(1):33-49.
65. Chaudhari KS, Tiwari NR, Tiwari RR, Sharma RS. Neurocognitive effect of nootropic drug Brahmi (*Bacopa monnieri*) in Alzheimer’s disease. Ann Neurosci. 2017;24(2):111-22.
66. Aguiar S, Borowski T. Neuropharmacological review of the nootropic herb *Bacopa monnieri*. Rejuvenation Res. 2013;16(4):313-326.
67. Pase MP, Kean J, Sarris J, Neale C, Scholey AB, et al. The cognitive-enhancing effects of *Bacopa monnieri*: A systematic review of randomized, controlled human clinical trials. J Altern Complement Med. 2012;18(7):647-652.
68. Fatima U, Roy S, Ahmad S, Al-Keridis LA, Alshammari N, et al. Investigating neuroprotective roles of *Bacopa monnieri* extracts: Mechanistic insights and therapeutic implications. Biomed Pharmacother. 2022;153:113469.
69. Zhou Y, Peng L, Zhang WD, Kong DY. Effect of triterpenoid saponins from *Bacopa monnieri* on scopolamine-induced memory impairment in mice. Planta Med. 2009;75(06):568-574.
70. Njan AA, Adenuga FO, Ajayi AM, Sotunde O, Ologe MO, et al. Neuroprotective and memory-enhancing effects of methanolic leaf extract of *Peristrophe bicalyculata* in rat model of type 2 diabetes mellitus. Heliyon. 2020;6(5):e04011.
71. Brimson JM, Brimson SJ, Brimson CA, Rakkhitawatthana V, Tencomnao T. *Rhinacanthus nasutus* extracts prevent glutamate and amyloid- $\beta$  neurotoxicity in HT-22 mouse hippocampal cells: Possible active compounds include lupeol, stigmasterol and  $\beta$ -sitosterol. Int J Mol Sci. 2012;13(4):5074-5097.
72. Esfandiari E, Ghanadian M, Rashidi B, Mokhtarian A, Vatankhah AM. The effects of *Acorus calamus* L. in preventing memory loss, anxiety, and oxidative stress on lipopolysaccharide-induced neuroinflammation rat models. Int J Prev Med. 2018;9(1):85.
73. Mukherjee PK, Kumar V, Mal M, Houghton PJ. *Acorus calamus*: Scientific validation of ayurvedic tradition from natural resources. Pharm Biol. 2007;45(8):651-666.
74. Shin JW, Cheong YJ, Koo YM, Kim S, Noh CK, et al.  $\alpha$ -Asarone ameliorates memory deficit in lipopolysaccharide-treated mice via suppression of pro-inflammatory cytokines and microglial activation. Biomol Ther (Seoul). 2014;22(1):17-26.
75. Li Y, Zhang XL, Huang YR, Zheng YY, Zheng GQ, et al. Extracts or active components from *Acorus gramineus* aiton for cognitive function impairment: Preclinical evidence and possible mechanisms. Oxid Med Cell Longev. 2020;2020(1):6752876.
76. Cho N, Lee KY, Huh J, Choi JH, Yang H, et al. Cognitive-enhancing effects of *Rhus verniciflua* bark extract and its active flavonoids with neuroprotective and anti-inflammatory activities. Food Chem Toxicol. 2013;58:355-361.
77. Asuquo OR, Udonwa UN, Eluwa MA, Ekanem TB. Effects of *Spondias mombin* leaf extract on the cytoarchitecture of the cerebral cortex and on learning and memory in Wistar Rats. Int J Sci Res. 2013;2(9):5-8.
78. Porwal M, Kumar A. Neuroprotective effect of *Annona squamosa* & (-) Annonaine in decreased GABA receptor of epileptic rats. J Appl Pharm Sci. 2015;5(1):018-023.
79. Soumyanath A, Zhong YP, Henson E, Wadsworth T, Bishop J, et al. *Centella asiatica* extract improves behavioral deficits in a mouse model of Alzheimer’ s Disease: Investigation of a possible mechanism of action. Int J Alzheimers Dis. 2012;2012(1):381974.
80. Pham HN, Vuong QV, Bowyer MC, Scarlett CJ. Phytochemicals derived from *Catharanthus roseus* and their health benefits. Technologies. 2020;8(4):80.
81. Chattipakorn S, Pongpanparadorn A, Pratchayasakul W, Pongchaidacha A, Ingkaninan K, et al. *Tabernaemontana divaricata* extract inhibits neuronal acetylcholinesterase activity

- in rats. J Ethnopharmacol. 2007;110(1):61-68.
82. Yamauchi Y, Ge YW, Yoshimatsu K, Komatsu K, Kuboyama T, et al. Memory enhancement by oral administration of extract of *Eleutherococcus senticosus* leaves and active compounds transferred in the brain. Nutrients. 2019;11(5):1142.
83. Kim CJ, Ryu HY, Lee S, Lee HJ, Chun YS, et al. Neuroprotective effect and antioxidant potency of fermented cultured wild ginseng root extracts of *Panax ginseng* CA meyer in mice. Molecules. 2021;26(10):3001.
84. Sharma K, Bhatnagar M, Kulkarni SK. Effect of *Convolvulus pluricaulis* Choisy. and *Asparagus racemosus* Willd on learning and memory in young and old mice: A comparative evaluation. Indian J Exp Biol. 2010;48(5):479-485.
85. Akbaba E, Hassan S, Mohammed Sur T, Bagci E. Memory enhancing, anxiolytic and antidepressant effects of *Achillea biebersteinii* (Asteraceae) essential oil on scopolamine-induced rats. J Essent Oil-Bear Plants. 2018;21(3):825-839.
86. Cho N, Lee HK, Jeon BJ, Kim HW, Kim HP, et al. The effects of *Betula platyphylla* bark on amyloid beta-induced learning and memory impairment in mice. Food Chem Toxicol. 2014;74:156-163.
87. Sumanth M, Yashashwini YC. Evaluation of learning and memory enhancing activity of *Commiphora caudata* leaves in rats. World J Pharm Res. 2016;5(5):1223-1236.
88. Bhandari R, Gyawali S, Aryal N, Gaire D, Paudyal K, et al. Evaluation of Phytochemical, Antioxidant, and Memory-Enhancing Activity of *Garuga pinnata* Roxb. Bark and Bryophyllum pinnatum (Lam) Oken. Leaves. Scientific World Journal. 2021;2021(1):6649574.
89. Kim JJ, Jeon SG, Kim KA, Kim JJ, Song EJ, et al. *Platycodon grandiflorus* root extract improves learning and memory by enhancing synaptogenesis in mice hippocampus. Nutrients. 2017;9(7):794.
90. Cho J, Kang JS, Long PH, Jing J, Back Y, et al. Antioxidant and memory enhancing effects of purple sweet potato anthocyanin and cordyceps mushroom extract. Arch Pharm Res. 2003;26:821-825.
91. Wahid S, Alqahtani A, Khan RA. *Cucurbita maxima* seeds reduce anxiety and depression and improve memory. Behav Neurol. 2023;2023(1):7509937.
92. Patro G, Bhattamisra SK, Mohanty BK. Effects of *Mimosa pudica* L. leaves extract on anxiety, depression and memory. Avicenna J Phytomed. 2016;6(6):696-710.
93. Beppe GJ, Dongmo AB, Foyet HS, Tsabang N, Olteanu Z, et al. Memory-enhancing activities of the aqueous extract of *Albizia adianthifolia* leaves in the 6-hydroxydopamine-lesion rodent model of Parkinson's disease. BMC Complement Altern Med. 2014;14:142.
94. Kim DH, Yoon BH, Kim YW, Lee S, Shin BY, et al. The seed extract of *Cassia obtusifolia* ameliorates learning and memory impairments induced by scopolamine or transient cerebral hypoperfusion in mice. J Pharmacol Sci. 2007;105(1):82-93.
95. Dhingra D, Parle M, Kulkarni SK. Memory enhancing activity of *Glycyrrhiza glabra* in mice. J Ethnopharmacol. 2004;91(2-3):361-365.
96. Moriasi GA, Ileri AM, Ngugi MP. *In Vivo* Cognitive-Enhancing, *Ex Vivo* Malondialdehyde-Lowering Activities and Phytochemical Profiles of Aqueous and Methanolic Stem Bark Extracts of *Piliostigma thonningii* (Schum.). Int J Alzheimers Dis. 2020;2020(1):1367075.
97. Soodi M, Naghdi N, Hajimehdipoor H, Choopani S, Sahraei E. Memory-improving activity of *Melissa officinalis* extract in naïve and scopolamine-treated rats. Res Pharm Sci. 2014;9(2):107-114.
98. Park SJ, Kim DH, Lee IK, Jung WY, Park DH, et al. The ameliorating effect of the extract of the flower of *Prunella vulgaris* var. *lilacina* on drug-induced memory impairments in mice. Food Chem Toxicol. 2010;48(6):1671-1676.
99. Ozarowski M, Mikolajczak PL, Piasecka A, Kujawski R, Bartkowiak-Wieczorek J, et al. Effect of *Salvia miltiorrhiza* root extract on brain acetylcholinesterase and butyrylcholinesterase activities, their mRNA levels and memory evaluation in rats. Physiol Behav. 2017;173:223-230.

100. Moss L, Rouse M, Wesnes KA, Moss M. Differential effects of the aromas of *Salvia* species on memory and mood. *Hum Psychopharmacol*. 2010;25(5):388-396.
101. Heo H, Shin Y, Cho W, Choi Y, Kim H, et al. Memory improvement in ibotenic acid induced model rats by extracts of *Scutellaria baicalensis*. *J Ethnopharmacol*. 2009;122(1):20-27.
102. Malik J, Munjal K, Deshmukh R. Attenuating effect of standardized lyophilized *Cinnamomum zeylanicum* bark extract against streptozotocin-induced experimental dementia of Alzheimer's type. *J Basic Clin Physiol Pharmacol*. 2015;26(3):275-285.
103. Rajesh V, Riju T, Venkatesh S, Babu G. Memory enhancing activity of *Lawsonia inermis* Linn. leaves against scopolamine induced memory impairment in Swiss albino mice. *Orient Pharm Exp Med*. 2017;17(2):127-142.
104. Nandhini S, Vadivu R, Jayshree N. Memory strengthening activity on seeds of *Abelmoschus moschatus*. *IJRPC*. 2014;4(2).
105. Kulkarni PD, Ghaisas MM, Chivate ND, Sankpal PS. Memory enhancing activity of *Cissampelos pariera* in mice. *Int J Pharm Sci*. 2011;3(2):206-211.
106. Wachiryah TA, Hathaipat L. Enhancing effect of *Tiliacora triandra* leaves extract on spatial learning, memory and learning flexibility as well as hippocampal choline acetyltransferase activity in mice. *Avicenna J Phytomed*. 2018;8(4):380-388.
107. Ahmed F, Chandra JN, Manjunath S. Acetylcholine and memory-enhancing activity of *Ficus racemosa* bark. *Pharmacognosy research*. 2011;3(4):246-249.
108. Parle M, Dhingra D, Kulkarni SK. Improvement of mouse memory by *Myristica fragrans* seeds. *J Med Food*. 2004;7(2):157-161.
109. Sriraksa N, Kaewwongse M, Phachonpai W, Hawiset T. Effects of lemongrass (*Cymbopogon citratus*) essential oil inhalation on cognitive performance and mood in healthy women.
110. Thai Pharmaceutical and Health Science Journal. 2018;13(2):80-88.
111. Minami A, Matsushita H, Horii Y, Ieno D, Matsuda Y, et al. Improvement of depression-like behavior and memory impairment with the ethanol extract of *Pleurotus eryngii* in ovariectomized rats. *Biol Pharm Bull*. 2013;36(12):1990-1995.
112. Liang CH, Huang PC, Mau JL, Chiang SS. Effect of the king oyster culinary-medicinal mushroom *Pleurotus eryngii* (Agaricomycetes) basidiocarps powder to ameliorate memory and learning deficit in ability in A $\beta$ -induced Alzheimer's disease C57BL/6J mice model. *Int J Med Mushrooms*. 2020;22(2).
113. Zhou Y, Ma C, Li BM, Sun C. *Polygala japonica* Houtt. reverses depression-like behavior and restores reduced hippocampal neurogenesis in chronic stress mice. *Biomed Pharmacother*. 2018;99:986-996.
114. Li Z, Liu Y, Wang L, Liu X, Chang Q, et al. Memory-enhancing effects of the crude extract of *Polygala tenuifolia* on aged mice. *Evid Based Complement Alternat Med*. 2014;2014(1):392324.
115. Norouzi F, Hosseini M, Abareshi A, Beheshti F, Khazaei M, et al. Memory enhancing effect of *Nigella sativa* hydro-alcoholic extract on lipopolysaccharide-induced memory impairment in rats. *Drug Chem Toxicol*. 2019;42(3).
116. Baitharu I, Jain V, Deep SN, Hota KB, Hota SK, et al. *Withania somnifera* root extract ameliorates hypobaric hypoxia induced memory impairment in rats. *J Ethnopharmacol*. 2013;145(2):431-441.
117. Mikulska P, Malinowska M, Ignacyk M, Szustowski P, Nowak J, et al. Ashwagandha (*Withania somnifera*)-Current research on the health-promoting activities: A narrative review. *Pharmaceutics*. 2023;15(4):1057.
118. Zhao Z, Li Y, Chen H, Huang L, Zhao F, et al. *Xylaria nigripes* mitigates spatial memory impairment induced by rapid eye movement sleep deprivation. *Int J Clin Exp Med*. 2014;7(2):356-362.
119. Muralidharan P, Kumar VR, Balamurugan G. Protective effect of *Morinda citrifolia* fruits on  $\beta$ -amyloid (25–35) induced cognitive dysfunction in mice: An experimental and biochemical study. *Phytother Res*. 2010;24(2):252-258.

**Corresponding author:** *Aya Wada, Department of Natural Product Chemistry and Biological Research, Delightex Pte. Ltd, Singapore, Singapore*

**E-mail:** *aya@delightexplorers.com*

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