Introduction

Alzheimer’s disease (AD) is a multifarious neurodegenerative disease that causes cognitive impairment and gradual memory loss. Senile plaques and neurofibrillary tangles (NFTs) comprised of amyloid β (Aβ) peptides and hyper-phosphorylated tau proteins, respectively, are the classic pathological hallmarks of AD [1]. Several hypotheses have been put forward to reveal the pathobiology of the disorder including cholinergic hypothesis, inflammation hypothesis, oxidative stress, mitochondrial cascade hypothesis, and metabolic hypothesis [2, 3]. According to the World Health Organization, AD is the most common form of dementia and accounts for about 60–70% of all dementia cases [4]. There is more prevalence for people aged over 65 y [5]. Epigenetic variations on the genetic material of neurons can result in neurodegenerative disorders and these variations in AD genes change from region to region [6]. Familial Alzheimer’s disease (FAD) and sporadic Alzheimer’s disease are the 2 specific types of AD [7]. The combined action of environmental and genetic factors might be culpable for the SAD and FAD types [8]. Depending on the age of onset, AD can be further grouped into early onset (EOAD) and late onset (LOAD) [9]. Mutations in some specific genes like amyloid precursor protein (APP), presenilin I, and presenilin II are responsible for EOAD whereas apolipoprotein E is the only gene identified that causes LOAD [10]. Approximately 5% of the AD cases is familial early onset type and it develops before 65 y of age, and the remaining 95% accounts for sporadic late onset type and it develops after 65 y of age [11, 12].

The neuropathophysiology of the neurodegeneration in AD is triggered by anomalous deposition of amyloid plaques and NFTs in varied regions of the brain involved in cognition and memory [13]. Amyloid plaques are protein fragments composed of Aβ
ABSTRACTS

ABTS 2,2-azinobis-3-ethylbenzthiazoline-6-sulfonic acid
ACH acetylcholine
ACHE acetylcholinesterase
AD Alzheimer’s disease
Aβ amyloid β
APP amyloid precursor protein
BChE butyrylcholinesterase
DPPH 1,1-diphenyl-2-picrylhydrazyl
EOAD early onset Alzheimer’s disease
EO essential oil
FAD familial Alzheimer’s disease
ICV intracerebroventricular
LD50 lethal dose 50%
LOAD late onset Alzheimer’s disease
MCFA medium-chain fatty acid
MWM Morris water maze
NFTs neurofibrillary tangles
SAD sporadic Alzheimer’s disease
SHXW SuHeXiang Wan
ZM Zataria multiflora Boiss

Out of all above pathogenetic pathways, the mechanisms of essential oils (EOs) are associated with cholinergic hypothesis and oxidative stress hypothesis. As stated by cholinergic hypothesis, loss of cholinergic neurotransmission in the cerebral cortex and the destruction of cholinergic neurons in the basal forebrain disturb cognitive function [17]. Depletion of acetylcholine (ACH) level is usually noticed in dementia patients and exploratory studies in animals and humans have also confirmed the role of ACh in cognitive function [17]. Depletion of acetylcholine (ACh) and the destruction of cholinergic neurons in the basal forebrain and oxidative stress hypothesis. As stated by cholinergic hypothesis [16].

EOs constitute a blend of enormously complex, volatile, naturally derived compounds that are obtained as secondary metabolites from plants. They are found to be copious in leaves, seeds, flowers, barks, and rhizomes and are commonly obtained through cold pressing, hydro-distillation methods [24]. Its major constituents include monoterpenes hydrocarbon, sesquiterpenes hydrocarbon, oxygenated sesquiterpenes, oxygenated monoterpenes, and esters [25]. A review on therapeutic and pharmaceutical use of EOs related to cardiovascular disease, cancer, anti-diabetic, penetration enhancer, antimicrobial, massage therapy, neuroprotective, aromatherapy, and anti-aging effects revealed the immense medicinal benefits of EOs [26]. Dobetzberger and Buchbauer [27] pointed out the analgesic, anxiolytic, anticonvulsive, and anti-AD effects. Recently, the anti-aging and neuroprotective effects of numerous EOs and their involvement in various brain disorders have been outlined [24]. Studies revealed that EOs are classic naturally derived antioxidants and most of them exhibits cholinesterase inhibitory potential and are relevant in AD pathogenesis [28].

Table 1 shows the list of EOs with respect to their anti-Alzheimer’s intervention.

Methods and Results

Scientific databases such as PubMed, ScienceDirect, Scopus, and Google Scholar from April 1998 to June 2018 were explored to collect data. The keywords searched were Alzheimer’s disease, EOs, amyloid hypothesis, NFTs, antioxidants, acetylcholinesterase, and dementia. We have conducted wide search on various EOs used in different models of AD. A total of 540 papers were obtained by scientific search, and 89 papers found to be relevant to the review. Studies of about 55 EOs were identified for Alzheimer’s disease,Twenty-eight EOs were finally included based on the quality of the study of which short descriptions of in vivo studies of 13 EOs together with clinical trial data of Salvia officinalis L., S. officinalis L. ssp. lavandulifolia (vahl) Gams, Melissa officinalis L., Lavandula angustifolia Mill., and Rosmarinus officinalis L. were highlighted. Chemical constituents and medicinal benefits of the EOs, animal models, doses and time of administration, study duration, behavioral parameters, and biochemical and histological assessments were also summarized. In vitro studies of 15 EOs that possess antioxidant and anticholinesterase potential are also mentioned.

Coriandrum sativum

Coriandrum sativum L. (coriander) of Umbelliferae/Apiaceae family has been reported to have anti-anxiety, anti-convulsive, antioxidative, anti-spasmodic, hypoglycemic, anti-inflammatory, antimutagenic, diuretic, antihypertensive, carminative, hypolipidemic,
antispasmodic, and antidepressant effects [29]. The major constituents of its EO include linalool (60–80%), terpinen-4-ol (3%), r-cymene (3.5%), α-terpinene (1–8%), α-pinene (0.2–8.5%), linalyl acetate (0–2.7%), α-terpineol (0.5%), camphor (0.9–4.9%), camphene (1.4%), myrcene (0.2–2.0%), geranyl acetate (0.1–4.7%), and geranial (1.2–4.6%) [30]. The effects of EO from dried fruits of C. sativum in AD has been studied in an in vivo study by Cionanco et al. [31] AD model was developed in rats by ICV Aβ(1–42) injection and the animals of the test group were made inhale C. sativum EO (1% and 3%) daily for 21 d. Cognitive performances were evaluated using radial arm maze and Y-maze tests. Aβ(1–42)-treated rats showed notable cognitive dysfunction, which was significantly reversed by EO inhalation. Coriander oil inhalation significantly decreased levels of SOD, LDH, and MDA and increased glutathione peroxidase levels in rat hippocampus. Presence of amyloid deposits were abundant in rats brain induced with ICV Aβ(1–42) and the deposits were less in rats treated with both doses of C. sativum EO. Authors concluded that exposure of C. sativum EO in Aβ(1–42)-induced rats could capably rebuild brain antioxidant status and confer neuroprotective effect possibly due to suppression of oxidative stress caused by Aβ(1–42).

### Syzygium aromaticum

*Syzygium aromaticum* (L.) Merr. & L. M. Perry. (clove) is an aromatic flower bud of Myrtaceae family. Clove oil is a mind stimulant used for medical care in anxiety, insomnia, and depression. The active constituents of its EO are eugenol (87.34%) and smaller amounts of eugenol acetate (5.18%) and β-caryophyllene (2.01%) [32].

### Table 1 List of EOs with respect to their anti-Alzheimer’s intervention.

<table>
<thead>
<tr>
<th>Plant/source</th>
<th>Study design</th>
<th>Results</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>C. sativum</em></td>
<td>ICV Aβ(1–42)-induced AD in rats</td>
<td>Reduced oxidative stress induced by Aβ(1–42) and confer neuroprotective effects</td>
<td>[31]</td>
</tr>
<tr>
<td><em>S. aromaticum</em></td>
<td>ICV colchicine-induced AD in rats</td>
<td>Reversed memory impairment, restored the levels of ACh and antioxidants, reduced neuroinflammation and mitochondrial dysfunction in colchicine treated rats.</td>
<td>[33]</td>
</tr>
<tr>
<td><em>Z. multiflora</em></td>
<td>ICV Aβ(25–35)-induced AD in rats</td>
<td>Reversed Aβ-induced learning deficits, exhibited antioxidant, anti-inflammatory and anti-cholinesterase activities.</td>
<td>[35]</td>
</tr>
<tr>
<td>SuHeXiang Wan</td>
<td>Drosophila model of AD</td>
<td>Neuroprotective effects</td>
<td>[39]</td>
</tr>
<tr>
<td>A. nemorosa</td>
<td>Scopolamine-induced spatial memory impairment in rats</td>
<td>Prevented cognitive defects.</td>
<td>[42]</td>
</tr>
<tr>
<td><em>S. officinalis</em></td>
<td>Single blind trial in healthy volunteers</td>
<td>Enhanced mood and cognitive performance</td>
<td>[45]</td>
</tr>
<tr>
<td><em>S. officinalis</em> ssp. <em>lavandulifolia</em> (vahl) Gams</td>
<td>Single blind trial in healthy volunteers</td>
<td>No significant effects.</td>
<td>[45]</td>
</tr>
<tr>
<td><em>M. officinalis</em></td>
<td>Clinical study in healthy volunteers for mood and cognition.</td>
<td>Enhanced mood and cognition</td>
<td>[50]</td>
</tr>
<tr>
<td><em>M. officinalis</em></td>
<td>Clinical trial in mild/moderate AD patients</td>
<td>Improved memory and cognition</td>
<td>[51]</td>
</tr>
<tr>
<td><em>J. communis</em></td>
<td>ICV Aβ(1–42)-induced AD in rats</td>
<td>Reduced memory impairment, antioxidant and anti-cholinesterase potential</td>
<td>[59, 60]</td>
</tr>
<tr>
<td><em>C. nucifera</em></td>
<td>Prospective study in AD patients</td>
<td>Improved memory status in AD patients.</td>
<td>[62]</td>
</tr>
<tr>
<td></td>
<td>In vitro neurotoxicity study in rat cortical neurons</td>
<td>Neuroprotective effects</td>
<td>[63]</td>
</tr>
<tr>
<td><em>P. peregrina</em></td>
<td>Scopolamine-induced memory deficits, depression, and anxiety in a rat model of AD</td>
<td>Hindered depression, anxiety, and memory deficits.</td>
<td>[65]</td>
</tr>
<tr>
<td><em>L. angustifolia</em></td>
<td>Scopolamine-induced amnesia model in rats</td>
<td>Hindered depression, anxiety, and memory deficits.</td>
<td>[69]</td>
</tr>
<tr>
<td></td>
<td>Clinical trials</td>
<td>Improved mood and cognitive performance.</td>
<td>[70, 71]</td>
</tr>
<tr>
<td><em>R. officinalis</em></td>
<td>Scopolamine-induced dementia model in mice</td>
<td>Reduced memory impairment.</td>
<td>[75]</td>
</tr>
<tr>
<td></td>
<td>Memory performances in aged and young mice.</td>
<td>Improved memory performances</td>
<td>[76]</td>
</tr>
<tr>
<td></td>
<td>Clinical trials</td>
<td>Improved mood and cognitive performances</td>
<td>[70, 71, 76–78]</td>
</tr>
</tbody>
</table>

AD: Alzheimer’s disease, ICV: intra-cerebroventricular, Aβ: amyloid β
neuroprotective potential of clove oil in ICV colchicine-induced memory impairment in rats has been reported in a study by Kumar et al. [33]. Male SD rats were subjected to intraperitoneal injection of minocycline (25 and 50 mg/kg), clove oil (0.05 mL/kg and 0.1 mL/kg), and their combinations for 21 d. ICV colchicine-treated rats showed significant cognitive impairment in Morris water maze (MWM) test and showed an increase in AChE activity, oxidative stress, neuro-inflammation, and mitochondrial dysfunction as compared to control group. Treatment with clove oil was found to reverse the memory impairment, restored the levels of ACh and antioxidants, and reduced neuroinflammation and mitochondrial dysfunction in colchicine-treated rats. Authors highlighted that the favorable potential of clove oil in reversing memory impairment and cognitive dysfunction is due to its antioxidative, mitochondrial-restoring, and microglial inhibitory mechanisms.

**Zataria multiflora**

*Zataria multiflora* Boiss. (ZM), belonging to Lamiaceae family, is a traditional Iranian herb with anti-inflammatory, antibacterial, antifungal, spasmyloytic, anticoagulant, antioxidative, and immune stimulant properties. The main constituents of its EO include thymol (40.94%), carvacrol (22.23%), p-cymene (7.73%), linalool (7.92%), caryophyllene (3.95%), and y-terpinene [36]. Majlessi et al. [35] mentioned the beneficial effects of EO from the leaves of ZM in ICV Aβ(25–35)-induced AD model in rats. Seven days after induction of disease, EO was administered in a dose of 50, 100, and 200 µg/mL (i.p.) 30 min prior to MWM task. Aβ injection in the rat brain remarkably impaired the ability of rats to locate the hidden platform in MWM task. Administration of ZM EO significantly reversed Aβ-induced learning deficits. Also, LD50 of EO was determined in male NMRI mice and was found to be 1264.9 µL/kg. Authors concluded that EO administration remarkably reversed Aβ(25–35)-induced cognitive dysfunction in rats due to its antioxidant, anti-inflammatory, and anticholinesterase activities. Further studies are needed to detect the presence of amyloid peptides and neurological alterations and more significant results can be obtained.

**SuHeXiang Wan**

SuHeXiang Wan (SHXW) is a conventional Chinese medicinal mixture of about 15 crude herbs including *Cyperus rotundus* L., *Arctotyloses macrolephala* Koidz., *Boswellia sacra* Flueck., *Bubalus bubalis* Linnaeus L. horn powder concentrate, artificial moschus, *Salvia officinalis* L. (common sage/sage) and *cinalis* of Lamiaceae family have prominent medicinal benefits [43]. The active constituents of *S. officinalis* EO are camphor (95.5%), 1,8-cineole (50.9%), cis-thujone (65.5%), trans-thujone (40.1%), a-humulene (33.7%), and linalool (35.0%). *S. lavandulifolia* EO contains 1,8 cineole (34.5%), α-pinene (23.2%), β-pinene (19.2%), limonene (16.6%), camphor (15.4%), and β-caryophyllene (8.1%) as main constituents [44]. In a single-blind randomized controlled trial led by Moss et al. [45], the reputed effects of these EOs on mood and cognition in healthy volunteers have been evaluated. *S. officinalis* EO-treated group showed better positive performance compared to control group. Results confirmed an enhancement in mood and cognitive performance delivered through Bond-Lader mood scales and cognitive drug research system, respectively. EO of *S. lavandulifolia* does not exhibit any significant effects. EO of *S. lavandulifolia* and its constituents reported antioxidant [46], anti-inflammatory, oestrogenic [47], CNS depressant, and AChE inhibitory potential in bovine erythrocyte and human brain tissue [48] and in rat brain [49]. In a clinical trial conducted in healthy volunteers, EO treatment reported successive results on mood and cognition [50]. Oral administration of *S. lavandulifolia* EO in patients with mild/
moderate AD reported significant improvement in memory and cognition in a pilot open-label study [51]. Even though S. lavandulifolia EO possesses antioxidant, anti-inflammatory, anti-AChE, and memory enhancement potential, it fails in a clinical study [45] and that might be due to lack of standardization and pharmacokinetics studies of this EO. The concentration of bioactive components varies in plants based on its region, climate, etc., and this variation remains a major drawback concerning the properties of EO and is likely to remain so.

**Melissa officinalis**

*Melissa officinalis* L. (lemon balm) of Labiatae has been used traditionally in nervous system diseases, cognitive dysfunction, and sedation [52]. Geranial (65.4%), geranyl acetate (7.4), nerol (24.7%), (Z)-carveol (0.1%), linalool (0.8%), myrtenol (0.1%), farnesene (0.1%), 2,3-dehydro-1,8-cineole (0.1%), and carvophyllene (0.8%) are the active components that have been isolated from the leaves of *M. officinalis* EO [53]. Agitation is one of the main symptoms of AD. Aromatherapy of *M. officinalis* EO or placebo (sunflower oil) in a double-blind placebo-controlled trial in 72 Alzheimer’s patients (agitation with severe dementia) has been reported by Ballard et al. [54]. Patients were advised to apply oil combined with base lotion in faces and arms twice daily for 4 wk. Treatment analysis like Cohen-Mansfield Agitation Inventory score and quality of life indices were assessed and the results reflected 35% and 11% improvement in agitation in EO and placebo-treated groups, respectively. Later in 2011, a clinical trial report revealed that massaging EO in hands and upper arms of dementia patients with agitation does not produce any significant results compared to standard drug donepezil or placebo oil [55]. However, positive results of *M. officinalis* oil for its anti-agitation potential have also been reported [56,57]. While all these trials comprised comparatively short duration and small sample sizes, the significant improvement in agitation with *M. officinalis* oil appear to explain the importance for further long-term clinical studies, assessment of adverse effects, appropriate doses, and efficacy compared to standard drugs for AD.

**Juniperus communis**

*Juniperus communis* L. (juniper) belongs to the family Cupressaceae. The chemical compositions of its EO are α-pinene (41.13%), α-thujene (3.78%), myrcene (10.16%), sabinene (11.73%), γ-terpinene (1.57%), and limonene (8.63%) followed by lower quantities of β-elemene (1.1%), terpinen-4-ol (3.42%), β-caryophyllene (1.03%), eremophyline (1.17%), and germacrene D (1.51%). The EO obtained from needles and berries of *J. communis* has been reported to have carminative, diuretic, anti-anxiety, astringent, rubefacient, antispasmodic, stimulating, stomachic, blood purifier, antihemorrhagic, antiseptic, and tonic effects. [58,59] Cioanco et al. [59] reported about the multiple exposure of juniper oil inhalation boosted memory defects in an Aβ-induced rat model of AD. In this study, rats were split up into 4 groups, of which control group received saline treatment (0.9% NaCl), disease group was subjected to ICV Aβ(1–42) injection, and the other 2 treatment groups were subjected to inhalation of juniper volatile oil 1% and 3% for 20 d after ICV Aβ(1–42) injection. Y-maze and radial arm maze tests were used for the evaluation of memory status. Juniper oil treated groups showed significantly less memory impairment in both behavioral tests as compared to Aβ(1–42) alone treated group. Later Cioanco et al. [60] demonstrated that antioxidant and anti-AChE potential of juniper volatile oil ameliorates cognitive dysfunction in Aβ(1–42)-induced rat model of AD.

**Cocus nucifera**

*Cocus nucifera* L. (coconut) belonging to Areccaceae family, consists of saturated fatty acids (SFA) 92% and medium-chain fatty acids (MCFA) 62–70% as major constituents. MCFA can be transformed into ketone bodies, which are alternative energy source in the brain in conditions wherein disturbance in brain glucose level and can be favorable to humans progressing or earlier with cognitive dysfunction [61]. Hu Yang et al. [62] conducted a prospective study in Alzheimer’s patients in which the test group was provided with coconut oil (40 mL/day) for 3 wk. The memory test such as lobo cognitive test and mini test scores pre- and post-involvement were performed in test and control groups. A statistically appreciable elevation in test score was noticed in patients taking the oil and therefore a positive improvement in memory status was noted. An in vitro study conducted by Nafer et al. [63] highlighted the neuron protecting effects of coconut oil in Aβ-injected neurotoxicity in rat cortical neurons. Cells were initially induced with Aβ for 1, 6, or 24 h and then treated with coconut oil further for 24 h. Survival of neurons and various cellular parameters (synaptophysin labeling, cleaved caspase 3, and reactive oxygen species) were assessed. Reports unfolded that coconut oil protected against Aβ-induced neurotoxicity, inhibits Aβ-induced elevation of cellular stress markers, and in addition provides neuron protecting effects by activating Akt and ERK signaling pathways.

**Pimpinella peregrina**

*Pimpinella peregrina* L. is an aromatic herb belonging to Apiaceae family. The major constituents of its EO are trans-pinocarveol (35.1%), α-cubebene (12.4%), pregejerene (15.1%), (+)-epi-bicyclohexylsphenellandrene (7.5%), and α-terpineol (6.7%) [64,65]. EO is widely used in flavored drinks like raki, ouzo, pastis, arak etc. Adyn et al. [65] reported the effects of inhalation of EO from aerial parts of *P. peregrina* in scopalamine-induced memory deficits, depression, and anxiety in a rat model of AD. Rats were exposed to inhalation of EO (1% and 3%) continuously for 21 d. Scopolamine was injected (i. p.) 30 min before the behavioral tests. Memory impairments were observed through radial arm-maze and Y-maze tests. Also, elevated plus-maze and forced swimming tests were used for checking anxiety and depressive behavior, respectively. Successive results were obtained in EO-treated groups as compared to respective scopolamine-treated groups, suggesting the neuroprotective effects of *P. peregrina* EO in the treatment of anxiety, depression, and memory deficits. In conclusion, more biochemical and histological findings are needed to prove the effect of *P. peregrina* EO on AD.

**Lavandula angustifolia**

*Lavandula angustifolia* Mill. (lavender) belongs to the family Lamiaceae. The dominant components of its EO are caryophyllene (24.12%), terpinen-4-ol (9.57%), β-phellandrene (16%), and euca-

lymptol (15.69%) [66]. It has been used in the treatment of anxiety, migraines, stress, irritability, exhaustion, depression, headaches, digestion, colds, flatulence, insomnia, loss of appetite, stomach upset, liver diseases, nervousness, and aromatherapy [67,68]. Hritcu et al. [69] reported the effects of 2 distinct EOs from Lavandula hybrida Rev. and L. angustifolia subsp. angustifolia on neurological capability in scopolamine (0.7 mg/kg)-induced amnesia model in rats. Lavender EO inhalation for 7 consecutive days notably hindered depression, anxiety and memory impairment in scopolamine treated rats as observed through various behavioural tests. A human study led by Moss et al. [70] unveiled that exposure of EOs of Rosmarinus officinalis L. and L. angustifolia significantly affects the mood and cognitive performances of sound volunteers. Aromatherapy of 4 different EOs from R. officinalis, Citrus limon (L.), Citrus sinensis (L.) Osbeck, and L. angustifolia reported significant improvement in cognitive function in a clinical trial led in 28 elderly people [71].

**R. officinalis**

Rosmarinus officinalis L. (rosemary) belongs to the family Lamiaceae. The EO of R. officinalis consist of α-pinene (21.3%), 1,8 cineole (28.5%), camphor (27.7%), camphene (8.7%), β-pinene (4.7%), and borneol (2.5%) as major constituents. [72] Antioxidant, antiproliferative, antibacterial, antimutagenic, antiphlogistic, antiallergic, and chemopreventive, and ANS stimulant potentials of its EO have been reported [73,74]. Satiou et al. [75] have recently studied the beneficial effects of EO from leaves and flowers of R. officinalis on scopolamine-induced Alzheimer’s-type dementia model in mice. The results showed that inhalation of R. officinalis EO (4 µL/L or 8 µL/L air) have shown a positive result in spontaneous alternation behavior in Y-maze test, and 1,8 cineole, α-pinene, and β-pinene were found in dominant concentration in the mice brain. Asadi et al. [76] have tested the potential of EO from leaf and aerial part of R. officinalis on memory in aged and young mice. The test group were administered EO (200,400,600 and 800 mg/kg i.p.) daily for 7 d and improvement in memory performances were found in all test groups. Filipstora et al. [77] reported the effects of R. officinalis EO on the short-term and numerical memory in a study conducted in 79 school students (aged 13–17 y). The result showed positive results as evidenced by increased image and number memory compared to control groups. The positive correlation of 1,8 cineole on mood and cognitive performances following inhalation of rosemary oil has been described in a clinical study conducted in healthy volunteers [78]. As mentioned above, R. officinalis EO reported memory enhancement potential in many clinical studies [70,71]. Further studies are essential for this oil to verify possibility for drug interactions and adverse effects in long duration trials and to determine appropriate dose that can mediate relevant neuroprotective effects.

**In vitro studies**

Ayaz et al. [79] reported the cholinesterase inhibitory activities and reactive oxygen species scavenging potential of EOs from the flowers and leaves of Persicaria hydropiper (L.) Delarbre. In AChE inhibition assay, EO of leaf exhibited IC50 of 120 µg/mL and flower of 220 µg/mL, whereas flower and leaf EOs reported IC50 of 225 and 130 µg/mL, respectively, in butyrylcholinesterase (BChE) inhibitory assay. In 1,1-diphenyl-2-picrylhydrazyl (DPPH) antiradicals assay, these EOs showed remarkable radical scavenging potentials. Also, IC50 of 50 and 45 µg/mL for flower oil and 60 and 180 µg/mL for leaf oil was obtained in hydrogen peroxide and 2,2-azinobis(3-ethylbenzthiazoline-6-sulfonic acid (ABTS) scavenging assays, respectively. Ahmed et al. [80] reported the antioxidant and anticholinesterase activities of EO from Rumex hastatus D. Don. It exhibited significant BChE and AChE inhibitory activities with IC50 of 97.38 and 32.54 µg/mL, respectively. Also, IC50 values of 6.29 and 3.71 µg/mL were obtained for ABTS and DPPH assays, respectively. Salvia ch ionanha Boiss EO was analyzed for its antioxidant activity and reported 77.4 ± 0.5% inhibition in ABTS assay. The anti-AChE and BChE activities were found to be 56.7 ± 1.9% and 41.7 ± 2.9%, respectively [81]. In vitro anti-AChE activity of EO from Acorus calamus L. rhizomes has been studied and exhibited significant AChE inhibitory potential (IC50 = 10.61 µg/mL) [82]. Ertas et al. [83] reported the anticholinesterase activities of EOs from Anchnusa arvensis subsp. Orientalis (L.) Nordh. and Tragopogon latafolius Boiss. Both of them betrayed moderate AChE and BChE inhibitory activities.

EO isolated from leaf and stem of Gymnura bicolor (Roxb.ex Willd.) DC. has been screened for anti-AChE activity. These oils have been evaluated and compared with standard AChE inhibitor pulegone. The leaf and stem oil inhibited AChE (53%) at a concentration of 0.50 µg/mL with IC50 values of 85 and 92 µg/mL, respectively. Both oils showed better results than pulegone [84]. EO obtained from fresh Chaerophyllum aromaticum L. root and aerial parts exhibited significant anticholinesterase and antioxidant activities [85]. Bonesi et al. [86] reported the cholinesterase inhibitory activities of EOs from Pinus nigra J.F. Arnold and Pinus heldreichii Christ. P. heldreichii was more active with IC50 of 80.6 and 51.1 µg/mL toward BChE and AChE inhibitory assays, respectively. EOs from Cistus libanotis L., Cistus salvi folius L., Cistus creticus L., Cistus villosus L., and Cistus monspeliensis L. were evaluated for antiradicals and anti-AChE activities. In ferric reducing antioxidant assay, C. libanotis exhibited most potent value of 19.21 MFe (II)/g. In AChE inhibitory assay C. salvi folius revealed IC50 of 58.1 µg/mL. Meanwhile, C. c reticus, C. libanotis, and C. salvi folius exhibited significant anti-BChE effects with IC50 values of 29.1, 23.7, and 34.2 µg/mL, respectively. In β-carotene bleaching test, C. monspeliensis EO exhibited better results with IC50 of 54.7 µg/mL [87]. Eminent results of all this data suggest the anticholinesterase and antioxidant potential of EOs for the development of therapeutic remedies targeting AD pathogenesis. **Table 2** shows the list of *in vitro* studies of EOs that possess antioxidant and anticholinesterase activities.

**Discussion**

This literature review displayed promising evidence that supports the use of EOs for reversing cognitive and memory impairment of AD. EOs obtained from numerous medicinal plants are reported to possess anti-AD potential. EOs of *S. officinalis*, *S. officinalis* ssp. lavandulifolia (vahl), *M. officinalis*, L. angustifolia, and *R. officinalis* have proven their putative effects in clinical studies. EOs are rich in various bioactive markers in different concentrations and it changes from place to place. Linalool, thymol, β-caryophyllene, 

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**Note:** The text has been abridged for brevity and clarity. The full document is available upon request.
carvacrol, α-pinene, α-terpinen-1, α-terpinene, and eugenol are some of the components that are common in above-mentioned oils. ▶ Fig. 1 depicts the chemical structure of some of the main constituents of EOs. Some are present in higher amount like linalool (60–70%) in C. sativum, caryophyllene (23.6%) in A. nemorosa, and 1,8 cineole (46.0%) in R. officinalis while others are in minor amounts. Monoterpenes like pinene, limonene, sabinene, etc., are found to be responsible for the antioxidant and anticholinesterase properties [88]. Therefore, it seems to be important to screen the main constituents of EOs based upon the strength of the biologic potential and more findings are needed to establish the required dose and proportions that are necessary to protect the brain from degenerative diseases.

Majority of the clinical as well as animal studies prefer inhalation as the route of administration of EO for AD therapy. Aromatherapy for AD are used since long back and found to be very effective. The aroma of EO acquired through inhaling or massaging directly affects mood and cognition. These oils are supposed to enter the blood stream through lung mucosa, nasal, or even diffuse directly into the olfactory nerve and reach limbic system in the brain [89]. The EOs mentioned in this review is mostly non-toxic at the recommended doses. Most of the EOs possess anticholinesterase property. In vitro studies of 15 EOs that displayed both AChE and BChE inhibitory potentials are mentioned. ACh is the brain neurotransmitter involved in regulating memory and cognition and cholinesterase inhibitors prevent degradation of ACh and maintain its synaptic concentration. Hence, assessment of the potential shown by EOs on cholinesterase adds up a useful approach for developing new drugs for AChE/BChE inhibitor with fewer cholinergic side effects. Oxidative stress plays a major role in the pathophysiology of AD. Imbalance in oxidants/antioxidants defense generates free radical that causes cell/neuron damage.

Antioxidant-rich EOs can hence counteract the free radical mediated oxidative stress and thereby protects brain form neuronal damage. The ability of EOs to cross blood brain barrier remains as a main reason for activity in brain regions

However, it is important to initiate further studies on pharmacokinetics and toxicities of EOs and their bioactive markers responsible for anti-Alzheimer’s action. More cell culture and in vivo studies are essential. All these findings result in the establishment of good interventions and subsequent development of safer and more effective anti-Alzheimer’s drug.

▶ Table 2 List of in vitro studies of essential oils that possess antioxidant and anti-cholinesterase activities.

<table>
<thead>
<tr>
<th>Plant/source</th>
<th>Study design</th>
<th>Results</th>
<th>References</th>
</tr>
</thead>
<tbody>
<tr>
<td>P. hydropiper</td>
<td>AChE, BChE inhibition DPPH, ABTS, H2O2 assay</td>
<td>↓ AChE, BChE, DPPH, ABTS, H2O2 activity</td>
<td>[79]</td>
</tr>
<tr>
<td>R. hastatus</td>
<td>AChE, BChE inhibition DPPH, ABTS, H2O2 assays</td>
<td>↓ AChE, BChE, DPPH, ABTS activity</td>
<td>[80]</td>
</tr>
<tr>
<td>S. chionantha</td>
<td>AChE, BChE inhibition ABTS assays</td>
<td>↓ AChE, BChE, ABTS activity</td>
<td>[81]</td>
</tr>
<tr>
<td>A. calamus</td>
<td>AChE inhibition</td>
<td>↓ AChE activity</td>
<td>[82]</td>
</tr>
<tr>
<td>A. arvensis subsp. Orientalis T. latifolius</td>
<td>AChE, BChE inhibition</td>
<td>↓ AChE, BChE activity</td>
<td>[83]</td>
</tr>
<tr>
<td>G. bicolor</td>
<td>AChE inhibition</td>
<td>↓ AChE activity</td>
<td>[84]</td>
</tr>
<tr>
<td>C. aromaticum</td>
<td>AChE inhibition Antioxidant assay</td>
<td>↓ AChE activity Significant antioxidant activity</td>
<td>[85]</td>
</tr>
<tr>
<td>P. nigra P. heldreichii</td>
<td>AChE, BChE inhibition</td>
<td>↓ AChE, BChE activity</td>
<td>[86]</td>
</tr>
<tr>
<td>C. crenatus C. lornotis C. salvifolius C. montspeliensis C. villosus</td>
<td>AChE, BChE inhibition FRAP assay, β-carotene bleaching assay</td>
<td>↓ AChE, BChE activity Significant antioxidant activity</td>
<td>[87]</td>
</tr>
<tr>
<td>P. nigra</td>
<td>AChE inhibition</td>
<td>↓ AChE activity</td>
<td>[88]</td>
</tr>
<tr>
<td>P. heldreichii</td>
<td>AChE, BChE inhibition</td>
<td>↓ AChE, BChE activity</td>
<td>[89]</td>
</tr>
</tbody>
</table>


▶ Fig. 1 Some of the main bioactive constituents of EOs.
There are numerous natural resources with reputed ethnobotanical uses shown to ameliorate memory impairment in different Alzheimer’s studies. However, comparatively only few are explored to determine any therapeutic support for their putative effects. Our literary survey revealed encouraging results regarding various EOs being studied in preclinical and clinical studies of AD with significant effects in modulating the pathology through anti-amyloid, antioxidants, anti-AChE, and memory enhancement activity. The literature studies disclosed many drugs that showed positive results in animal studies but failed in clinical trial. This demands more studies focused on human physiology, pharmacokinetics, dose, and route of administration. Thus, the establishment of potential targets and optimization of safety and efficacy of EOs in AD remain as a promising area for future research.

Conflict of Interest

There are no conflicts of interest among the authors.

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