

Screening of traditional European herbal medicines for acetylcholinesterase and butyrylcholinesterase inhibitory activity

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Acetylcholinesterase (AChE) inhibitors are widely used for the symptomatic treatment of Alzheimer's disease (AD) to enhance central cholinergic transmission. On the other hand, butyrylcholinesterase (BuChE) inhibitors were reported to produce a significant increase in brain extracellular AChE without triggering severe peripheral or central side effects. In the present study, we selected twelve plants used in traditional European medicine to treat different central nervous system (CNS) disorders or to improve memory.

Methanolic and hexane extracts of these plants were tested for the AChE and BuChE inhibitory activity using Ellman's colorimetric method. The most potent AChE and BuChE inhibition was observed in the hexane extracts of the flowers of *Arnica chamissonis* Less. subs. *foliosa* and *Ruta graveolens* L. herb at a concentration of 400 µg mL⁻¹. However, methanolic extracts of the flowers of *Arnica chamissonis* Less. subs. *foliosa* and the *Hypericum perforatum* L. herb demonstrated at the same concentration, selective inhibition only against AChE but not against BuChE. The other extracts did not show any significant AChE or BuChE inhibitory activity. Our results show that further investigations of the extracts of arnica, rue and St. John's Wort are needed to identify the compounds responsible for the AChE and BuChE inhibitory activity.

Keywords: acetylcholinesterase inhibition, butyrylcholinesterase inhibition, traditional medicine

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Alzheimer's disease (AD) is one of the most widespread neurodegenerative diseases that involve dementia and mainly afflict people over 65 years of age. The therapy of early and moderate stages of AD is mainly based on acetylcholine esterase inhibitors such as synthetic donepezil and galanthamine isolated from the bulbs of daffodils. However, these licensed medicines have drawbacks of inducing severe peripheral and central

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side effects, including gastrointestinal disturbances, insomnia, fatigue or depression. On the other hand, since the BuChE activity in AD increases progressively as the severity of dementia progresses, researchers have investigated selective BuChE inhibitors in the treatment of AD as well (1). The serious side effects caused by licensed drugs used to treat AD have forced researchers to investigate safer AChE- or BuChE inhibitors from natural sources.

One of the best sources of new substances to treat AD are natural products and their derivatives. Traditionally, plants have been used to enhance memory and to alleviate other symptoms associated with AD (2). The biologically active plant-derived substances that may be considered as a source of new anticholinesterase drugs come from different classes of compounds and are characterized by the diversity of their structures. The majority of bioactive substances are indole-, steroidal-, piperidine- and *Amaryllidaceae* alkaloids, phenylpropanoids (furanocoumarins, xanones, and flavonoids) and terpenoids (diterpenes) (2).

The aim of this study was to investigate possible AChE or BuChE inhibitors in plants traditionally used in European medicine and to point to the role of these plants as potential sources for the development of therapeutic agents of AD. Selection of the species screened in this study was based on their use as remedies for the central nervous system diseases, as antidotes for plant and animal poisoning or to enhance memory.

EXPERIMENTAL

Plant material and chemicals

Plants were collected from The Botanical Garden of the Polish Academy of Science. A specimen of each raw material is available in the herb collection of the Department of Pharmacognosy and Molecular Basis of Phytotherapy, Faculty of Pharmacy, Medical University of Warsaw, Poland. Each plant material was dried and ground in a grinder. Plants and their parts used in this study are presented in Table I.

Electric eel AChE (type VI-S), horse-serum BuChE, acetylthiocholine iodide, butyrylthiocholine chloride, 5,5'-dithio-bis[2-nitrobenzoic acid], Tris-HCl [Tris(hydroxymethyl)aminomethane hydrochloride], 1-naphtyl acetate, bovine serum albumin (BSA, albumin fraction V from bovine serum), 3,3'-dimethoxybiphenyl-4,4'-di(diazonium) zinc chloride (Fast Blue B salt), physostigmine salicylate, galanthamine hydrobromide, anisaldehyde were purchased from Sigma-Aldrich (USA). Methanol, toluene, ethyl acetate, hexane and sulphuric acid were purchased from Chempur (Poland). All reagents used in the study were of analytical grade.

Extract preparation

For extraction, 2 g of air-dried and powdered plant material was extracted over a water bath at 60 °C in 40 mL of methanol or hexane under reflux for 30 min. The extraction was done in triplicate. The resulting liquid extract was filtered and concentrated to dryness under reduced pressure at 60 °C. At 60 °C, the components in plant extracts were

Table 1. Anti-AChE and anti-BuChE activity of plant extracts

Plant species	Family	Plant part analyzed	Solvent	AChE inhibition (%)		BuChE inhibition (%)	
				100 µg mL ⁻¹	400 µg mL ⁻¹	100 µg mL ⁻¹	400 µg mL ⁻¹
<i>Anchusa officinalis</i> L.	Boraginaceae	herb	methanol	6.4 ± 5.8	10.0 ± 7.9	1.9 ± 1.2	34.0 ± 4.7
			hexane	15.6 ± 3.8	39.6 ± 8.7	13.8 ± 6.3	33.5 ± 4.5
<i>Arnica chamissonis</i> Less. ssp. <i>foliosa</i> (Nutt.) Maguire	Asteraceae	flower	methanol	89.2 ± 3.1	95.0 ± 1.1	27.7 ± 3.1	59.7 ± 8.1
			hexane	76.5 ± 2.1	97.4 ± 1.8	63.9 ± 4.8	88.2 ± 3.6
<i>Ballota nigra</i> L.	Lamiaceae	herb	methanol	6.5 ± 5.1	35.7 ± 2.6	12.3 ± 5.8	38.7 ± 7.2
			hexane	26.7 ± 2.4	42.9 ± 3.5	19.0 ± 1.9	35.2 ± 0.9
<i>Cnicus benedictus</i> L.	Asteraceae	herb	methanol	2.7 ± 3.5	21.6 ± 6.1	7.9 ± 3.2	21.8 ± 5.6
			hexane	5.5 ± 2.3	27.0 ± 0.8	22.2 ± 1.7	47.5 ± 2.1
<i>Galium odoratum</i> L. Scop.	Rubiaceae	herb	methanol	11.2 ± 5.5	32.3 ± 6.7	15.3 ± 4.5	41.8 ± 5.6
			hexane	27.3 ± 3.6	53.1 ± 1.1	21.7 ± 3.3	49.1 ± 2.9
<i>Hypericum perforatum</i> L.	Hypericaceae	herb	methanol	29.3 ± 2.6	72.1 ± 6.3	23.2 ± 5.0	62.7 ± 5.1
			hexane	3.1 ± 1.9	40.5 ± 4.5	18.0 ± 3.3	47.8 ± 2.6
<i>Hyssopus officinalis</i> L.	Lamiaceae	herb	methanol	5.2 ± 8.2	13.2 ± 5.6	11.5 ± 0.5	31.8 ± 7.3
			hexane	29.6 ± 2.3	55.0 ± 1.7	23.2 ± 2.0	51.7 ± 1.7
<i>Menyanthes trifoliata</i> L.	Menyanthaceae	leaf	methanol	8.6 ± 7.4	21.5 ± 1.7	14.6 ± 2.8	35.7 ± 2.0
			hexane	17.1 ± 1.6	47.6 ± 10.2	22.5 ± 3.6	43.8 ± 3.9
<i>Primula officinalis</i> (L.) Hill.	Primulaceae	flower	methanol	12.5 ± 3.1	49.6 ± 4.3	13.1 ± 1.2	55.6 ± 5.3
			hexane	14.5 ± 3.7	24.6 ± 6.3	14.1 ± 3.2	23.4 ± 0.3
<i>Ruta graveolens</i> L.	Rutaceae	herb	methanol	28.2 ± 3.4	59.1 ± 4.3	13.7 ± 8.0	39.8 ± 6.6
			hexane	71.3 ± 3.1	94.9 ± 2.1	51.1 ± 5.1	86.0 ± 1.9
<i>Tanacetum parthenium</i> (L.) Schulz-Bip.	Asteraceae	herb	methanol	16.6 ± 2.8	32.4 ± 6.6	3.2 ± 2.5	39.8 ± 2.6
			hexane	13.0 ± 3.3	33.5 ± 4.8	20.9 ± 1.0	40.9 ± 2.3
<i>Verbena officinalis</i> L.	Verbenaceae	herb	methanol	2.8 ± 3.4	25.5 ± 4.6	9.4 ± 5.6	38.8 ± 3.2
			hexane	11.3 ± 3.5	25.3 ± 1.0	14.7 ± 2.5	32.8 ± 3.1

stable. The dry extracts were stored at 4 °C and dissolved in an appropriate solvent just before the test.

Determination of enzyme activity

All extracts were tested for AChE and BuChE inhibitory activity at concentrations of 100 and 400 $\mu\text{g mL}^{-1}$ by the modified spectrophotometric method developed by Ellman (3). For herbal extracts that were proven to exert significant inhibition and for positive controls, dose-dependent inhibitory assays were performed. The plant extracts were tested in a concentration range of 12.5 to 400 $\mu\text{g mL}^{-1}$. Galanthamine hydrobromide and physostigmine salicylate were used as AChE and BuChE positive controls. They were tested in a concentration range between 0.01 and 100 $\mu\text{g mL}^{-1}$.

The assay was performed in a 1.5-mL Eppendorf tube as follows: 1 mg of methanolic extract was dissolved in 1 mL Tris-HCl (50 mmol L^{-1} , pH 7.8). Hexane extract (1 mg) was dissolved in 100 μL methanol and diluted to 1 mL with Tris-HCl (50 mmol L^{-1} , pH 7.8). Positive controls, namely galanthamine and physostigmine, were dissolved in methanol (1 mg mL^{-1}).

To determine the AChE and BuChE inhibitory activity, the following reaction was performed. An appropriate amount of extract or positive control and 40 μL of AChE (0.45 U mL^{-1}) or BuChE (0.45 U mL^{-1}) was mixed and diluted to 1 mL with Tris-HCl. After 30 min of incubation at 4 °C, the reaction was initiated by addition of 20 μL of 5,5'-dithio-bis[2-nitrobenzoic acid] (3 mmol L^{-1}) and 20 μL of acetylthiocholine iodide or butyrylthiocholine chloride, respectively (both 15 mmol L^{-1} , dissolved in Tris-HCl). The samples were incubated for another 20 min over a water bath at 37 °C and the reactivity was terminated by addition of 20 μL physostigmine salicylate (0.1 mmol L^{-1} in methanol). Absorbance of the produced yellow 5-thio-2-nitrobenzoate anion was measured at a wavelength of 412 nm using a Shimadzu UV 160A (Japan) spectrophotometer. Tris-HCl buffer was used as a blank.

Thin layer chromatography (TLC) bioautographic assay

TLC bioautographic assay was performed as described by Marston *et al.* (4). Extract of the *Arnica chamissonis* flowers, 20 μL (0.2 g of dried extract dissolved in 1 mL methanol) was applied to a silica gel plate (Kieselgel G, F254, type 60, Merck, Germany) and eluted with toluene/ethyl acetate (50:50, V/V). After drying, the plate was sprayed with 13 U mL^{-1} AChE (dissolved in 50 mmol L^{-1} Tris-HCl with 0.1 % BSA, pH 7.8) and kept in a water bath at 37 °C for 20 minutes. The chromatogram was dried until complete mobile phase was removed. Finally, the plate was sprayed with 0.25 % 1-naphthyl acetate (dissolved in methanol) and 0.25 % Fast Blue B salt (dissolved in deionized water). After a few minutes, a purple background appeared with white spots for AChE inhibiting compounds.

TLC method

Methanolic and hexane extracts of *Arnica chamissonis* ssp. *foliosa*, 10 μL and 20 μL , respectively (0.2 g of dried extract dissolved in 1 of solvent) were applied to the silica gel

plate (Kieselgel G, F254, type 60, Merck) and eluted with toluene/ethyl acetate (50:50, V/V). Visualization of sesquiterpene lactones was done by spraying with methanolic solutions of 0.5 % anisaldehyde and 5 % sulphuric acid followed by heating at 105 °C for 5 minutes.

Date processing

The extent of the enzymatic reaction was calculated based on the following equation: $E = 100 - [(T - C_1/C_0) \times 100]$, where E is the activity of the enzyme. E value conveys the effect of the plant extract or the positive control on AChE or BuChE enzyme activity expressed as the percentage of the remaining activity in the presence of plant extract or positive control, T (test) is the absorbance of the tested sample (plant extract or positive control in the solvent) in the presence of enzyme, C_1 (control 1) is the absorbance of the tested sample (plant extract or positive control in the solvent) in the absence of enzyme, C_0 (control 0) is the absorbance of the solvent in the presence of enzyme.

Data are expressed as mean \pm standard error (SEM) and the results were taken from at least three independent experiments performed in duplicate.

Estimation of IC₅₀ values

The IC_{50} values (concentration of test compounds that inhibits the hydrolysis of substrates by 50 %) were determined by spectrophotometric measurement of the effect of increasing concentrations of test compounds (plant extracts and positive controls) on the AChE or BuChE activity. IC_{50} values were obtained from dose-effect curves by linear regression.

RESULTS AND DISCUSSION

Screening of methanolic and hexane extracts showed that 4 out of 24 plant extracts were able to inhibit the enzymatic activity of either AChE or BuChE or both. It was found out that all the hexane and methanolic extracts had dose-dependent inhibitory activity. Significant reduction in the activity of AChE was observed at the concentration of 400 $\mu\text{g mL}^{-1}$ for *Arnicae flos* (hexane and methanolic extract), *Rutae herba* (hexane and methanolic extract) and *Hyperici herba* (methanolic extract). The strongest BuChE inhibitors that showed inhibition higher than 50 % at the concentration of 100 $\mu\text{g mL}^{-1}$ were found in hexane extracts of *Rutae herba* and *Arnicae flos*. The results on the effects of the tested herbal extracts on AChE and BuChE activity are summarized in Table I. The dose-dependent AChE inhibitory activity of these three herbs was further studied and the IC_{50} values of inhibition are tabulated in Table II.

The most potent extract was the hexane extract of *Arnica chamissonis* with IC_{50} of 29 $\mu\text{g mL}^{-1}$. At a concentration of 100 $\mu\text{g mL}^{-1}$, it reduced the enzymatic activity of AChE to 23.6 % and BuChE to 36.1 %. Methanolic extract of *Arnica chamissonis* caused similar reduction of AChE activity at a concentration of 100 $\mu\text{g mL}^{-1}$ ($IC_{50} = 43 \mu\text{g mL}^{-1}$). However, no BuChE inhibitory activity was found for the hexane extract of *Arnica chamissonis*. The AChE inhibitory effect of these extracts was less potent than that of the refer-

Table II. Anti-AChE and BuChE activity of the most active plant extracts

Sample	IC ₅₀ (AChE) (μg mL ⁻¹)	IC ₅₀ (BuChE) (μg mL ⁻¹)
<i>Arnicae flos</i> (hexane extract)	29	88
<i>Arnicae flos</i> (methanolic extract)	43	–
<i>Hyperici herba</i> (methanolic extract)	178	–
<i>Rutae herba</i> (hexane extract)	34	61
Positive control	IC ₅₀ (anti-AChE)	IC ₅₀ (anti-BuChE)
Galanthamine	0.14 μg mL ⁻¹ (0.37 μmol L ⁻¹)	0.54 μg mL ⁻¹ (8.29 μmol L ⁻¹)
Physostigmine	0.064 μg mL ⁻¹ (0.21 μmol L ⁻¹)	3.13 μg mL ⁻¹ (1.73 μmol L ⁻¹)

– Not evaluated.

IC₅₀ values were obtained from the dose-effect curves by linear regression.

ences galanthamine (IC₅₀ of galanthamine for AChE and BuChE was 0.14 and 3.13 μg L⁻¹, respectively) and physostigmine (IC₅₀ of physostigmine for AChE and BuChE 0.064 and 0.54 μg L⁻¹, respectively).

The components found in *Arnica chamissonis* include sesquiterpene lactones, flavonoids and essential oil with terpenes. Detailed chemical composition of the tested plant parts is presented in Table III. Out of these compounds, terpenes like cineol, borneol, geraniol, 3-carene, α-caryophyllene or limonene were previously found to express some AChE or BuChE inhibitory activity (2, 5). However, no sesquiterpene lactones inhibitory

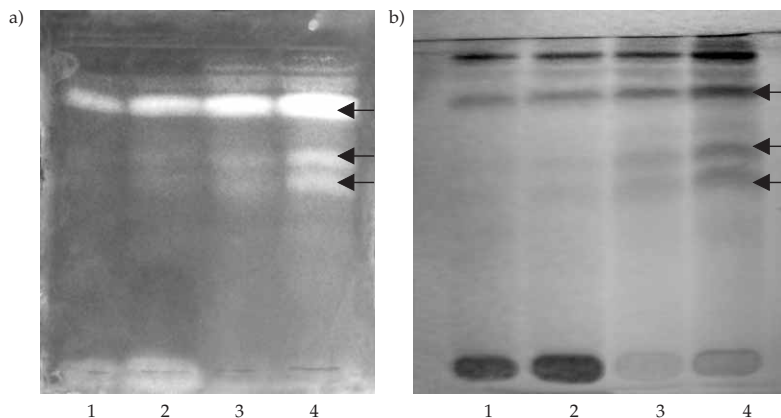


Fig. 1. TLC of extracts of *Arnica chamissonis* Less. ssp. *foliosa* showing acetylcholinesterase inhibitory activity: a) white spots indicate inhibition, arrows indicate the AChE inhibiting constituents, b) TLC plate sprayed with methanolic solution of 0.5 % anisaldehyde and of 5 % sulphuric acid followed by heating at 105 °C for 5 minutes. Black spots indicate the presence of sesquiterpene lactones; arrows indicate the AChE inhibiting constituents. 1–4 are methanolic extract 10 μL, methanolic extract 20 μL, hexane extract 10 μL, hexane extract 20 μL, respectively.

Table III. Detailed chemical composition of the tested plants (literature survey)

Plant species	Plant part	Chemical composition	References
<i>Anchusa officinalis</i> L.	herb	pyrrolizidine alkaloids, triterpene saponins, polyphenolic acids	10
<i>Arnica chamissonis</i> Less. ssp. <i>Foliosa</i> (Nutt.) Maguire	flower	sesquiterpene lactones of pseudoguaianolide type (0.2–1.5 %)-helenelin, flavonoids (0.4–0.6 %)-flavonols and flavones, essential oil (0.15–0.28 %)-fatty acids, sesquiterpenes, monoterpenes, caffeoylquinic acids-chlorogenic acid, cynarin, polysaccharides	11, 13
<i>Ballota nigra</i> L.	herb	phenylpropanoids (5.5 %)-verbascoside, forsythoside B, arenarioside and ballotetroside, diterpenoids, flavonoids luteolin 7-lactate, 7-glucosyllactate	11
<i>Cnicus benedictus</i> L.	herb	bitter sesquiterpene lactone ester cnicin (0.2–0.7 %); other germacrane sesquiterpenes include salonitenolide and artemisiifolin; the bitter lignans trachelogenin, arctigenin, and nortracheloside are also present, essential oil (0.03 %)	12
<i>Galium odoratum</i> L. Scop.	herb	coumarins (1.06 %)-coumarin, <i>o</i> -coumaric acid glucoside; iridoids asperuloside (0.28 %), monotropein (0.042 %), scandoside (0.042 %)	13
<i>Hypericum perforatum</i> L.	herb	phloroglucinol derivatives (0.2–0.4 %), principally hyperforin, adhyperforin, naphthodiantrones (0.1–0.3 %)-hypericin (0.08 %), pseudohypericin, flavonoids (2–4 %) include hyperoside, rutin, isoquercitrin, biflavones and others, tannins-procyanidins, xanthones, essential oil	11, 12
<i>Hyssopus officinalis</i> L.	herb	essential oil (0.3–1 %)-mainly isopinocampone, limonene, β -pinene; flavonoids (5–6%)-principally diosmin (3–6%), tanins (5–8%), phenylpropanoids including rosmarinic acid, caffeic acid and isoferulyl-D-glucose ester, triterpenes-ursolic acid	11, 13
<i>Menyanthes trifoliata</i> L.	leaf	secoiridoid glycosides dihydrofoliamenthin, menthiafolin and loganin, monoterpenoid alkaloids (0.035 %) gentianine and gentianidine, gentialutein, gentiatibetin, flavonoids	12
<i>Primula officinalis</i> (L.) Hill.	flower	saponins (2 %), flavonoids (3 %), carotinoids, essential oil	12
<i>Ruta graveolens</i> L.	herb	essential oil (0.2–0.4 %) with nonan-2-one, flavonoids (2–5 %) – rutine, coumarins-furanocoumarines – bergapten, psoralen, xanthoxanthin, xanthotoxin, isopimpinellin and rutamarin, alkaloids (0.4–1.4 %) are furoquinoline alkaloids gamma-fagarine, skimmianine and acridone alkaloids-arborinine	14
<i>Tanacetum parthenium</i> (L.) Schulz-Bip.	herb	sesquiterpene lactones (0.2–1.8 %)-parthenolid, eudesmenolide, guajanolide, flavanoids-apigenin, luteolin, chrysoeriol derivatives, essential oil (0.2–0.7 %)	12

Table III. Continued

Plant species	Plant part	Chemical composition	References
<i>Verbena officinalis</i> L.	herb	iridoid glycosides – verbenalin (0.15 %), hastatoside (0.08 %), dihydrocornine (0.01 %) and others, hydroxycinnamic acid derivatives – verbascoside (0.8 %), isoverbascoside, martynosid, eukovoside, flavanoids – flavone glycosides, especially luteolin 7-diglucuronide and apigenin 7-glucuronide, triterpenes- ursolic acid	11

activity against AChE was reported. Therefore, we performed a TLC- and TLC bioautographic assay to rule out this action.

The TLC bioautographic assay demonstrated that AChE inhibiting activity of *Arnica chamissonis* was due to compounds which in the TLC bioautographic assay appeared as white spots (Fig. 1). To determine whether the inhibition of AChE activity by arnica extracts was mediated by sesquiterpene lactones, TLC analysis was performed. The compounds that reacted as dark purple-grey spots with anisaldehyde sulfuric acid reagent indicated sesquiterpene lactone derivatives. The location of the white and dark spots on the TLC plate presented in Fig. 1 implicated that the compounds reacting as AChE inhibitors were of sesquiterpene lactones origin. The presented findings have never been reported earlier and we suggest that *Arnica chamissonis* should be considered for further studies to isolate compounds responsible for the anti-AChE activity.

In the present study, the hexane extract of *Rutae herba* exhibited moderate AChE and BuChE inhibitory activity with IC_{50} 34 and 61 $\mu\text{g mL}^{-1}$, respectively. Adersen (6) found AChE inhibitory activity of the methanolic extract of *Ruta graveolens* (100 $\mu\text{g mL}^{-1}$), which caused 39 % inhibition; on the other hand, no reports about the activity of *Ruta graveolens* hexane extract against BuChE inhibition have been published. Compounds present in rue that could be considered to have cholinergic activity included alkaloids and furanocoumarins. It has been previously shown that alkaloids belonging to different groups such as physostigmine, (-)-hupercin A or galanthamine (4) and some furanocoumarins like isoimperatorin, xanthotoxin and marmesin exhibited AChE and BuChE inhibitory effects (7). Kang (7) suggested that coumarin skeleton contained a pyrone moiety, which played an important role in the inhibitory activity against AChE. The presence of coumarins in *Ruta* could partly explain the anti-AChE activity of the hexane rue extract proven in this study.

The lowest IC_{50} value was obtained with the methanolic extract of *Hypericum perforatum* herb. The methanolic extract was found to exhibit a weak inhibitory effect against AChE compared to galanthamine and physostigmine. It reduced AChE activity to 28 % at a concentration of 400 $\mu\text{g mL}^{-1}$ with IC_{50} of 178 $\mu\text{g mL}^{-1}$. It was previously reported that the ethanolic extract of another *Hypericum* species, namely *Hypericum undulatum*, inhibited AChE activity by 69 % at 500 $\mu\text{g mL}^{-1}$ (8). Pharmacological actions of the ethanolic extract of *Hypericum perforatum* on CNS were well documented in previous reports (9). The inhibiting AChE activity of this extract may belong to these biological actions.

The other tested plant extracts did not exhibit any significant AChE or BuChE inhibitory activity.

CONCLUSIONS

The present investigation has shown that hexane and/or methanolic extracts from the flowers of *Arnica chamissonis* Less. ssp. *foliosa*, the herb *Ruta graveolens* L. and the herb *Hypericum perforatum* L. could inhibit the activity of AChE or BuChE or both. AChE inhibitory activity of *Arnica* extracts was among others, due to sesquiterpene lactones.

The extracts of arnica, rue and St. John's Wort were proved to have a great potential and should be considered for further studies to identify the constituents responsible for the AChE and BuChE inhibitory activity, which can be eventually utilized in the treatment of AD.

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S A Ž E T A K

Pretraživanje tradicionalnih europskih ljekovitih biljaka na inhibiciju acetilkolinesteraze i butirilkolinesteraze

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Inhibitori acetilkolinesteraze (AChE) povećavaju kolinergičku transmisiju u mozgu, pa se koriste za simptomatsko liječenje Alzheimerove bolesti (AD). S druge strane, inhibitori butirilkolinesteraze (BuChE) značajno povećavaju ekstracelularnu količinu AChE u mozgu, a da pri tome ne uzrokuju snažne nuspojave ni u središnjem ni u perifernom živčanom sustavu. Galantamin, jedan od odobrenih AChE inhibitora, alkaloid iz lukovica narcisa, pokazuje da su biljke značajni izvor novih potencijalnih AChE- i BuChE-inhibitora. U ovom radu, ispitan je učinak dvanaest biljaka koje se koriste u tradicionalnoj europskoj medicini na različite poremećaje središnjeg živčanog sustava i na poboljšanje pamćenja. Pomoću Ellmanove kolorimetrijske metode praćen je inhibitorni učinak metanolnih i heksanskih ekstrakata tih biljaka na AChE i BuChE. Najjači inhibitorni učinak pokazali su heksanski ekstrakti cvjetova *Arnica chamissonis* Less. subsp. *foliosa* i nadzemnih dijelova *Ruta graveolens* L. u koncentraciji od 400 $\mu\text{g mL}^{-1}$. Međutim, metanolni ekstrakti cvjetova *Arnica chamissonis* Less. subsp. *foliosa* i nadzemnih dijelova *Hypericum perforatum* L. u istim koncentracijama pokazuju selektivnu inhibiciju samo na AChE. Ostali ekstrakti bili su nedjelotvorni. Rezultati ukazuju na potrebu daljnjih ispitivanja ekstrakata arnike, rute i gospine trave da se utvrdi koji su sastojci ekstrakata odgovorni za inhibiciju AChE i BuChE.

Ključne riječi: inhibicija acetilkolinesteraze, inhibicija butirilkolinesteraze, tradicionalna medicina

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