Extracts of *Glycyrrhiza uralensis* and Isoliquiritigenin Counteract Amyloid-β Toxicity in *Caenorhabditis elegans*

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**Affiliations**

**Key words**

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amyloid-β

*Caenorhabditis elegans*

**Abstract**

Alzheimer’s disease is a rising threat for modern societies as more and more people reach old age. To date, there is no effective treatment for this condition. In this study, we investigated the potential of *Glycyrrhiza uralensis* to counteract amyloid-β toxicity, one of the key features of Alzheimer’s disease. An LC-MS/MS analysis revealed glycyrrhizic acid and glycosylated forms of isoliquiritigenin and liquiritigenin as major constituents of water and methanol extracts of *G. uralensis*. These extracts and the pure compounds were tested for their activity in two *Caenorhabditis elegans* models of amyloid-β aggregation and amyloid-β toxicity, respectively. The number of amyloid-β aggregates decreased by 30% after treatment with isoliquiritigenin, the methanol extract could reduce the number by 14%, liquiritigenin and glycyrrhizic acid by 15%, and the aglycon of glycyrrhizic acid, glycyrrhetinic acid, by 20%. Both extracts and isoliquiritigenin also showed significant activity against acute amyloid-β toxicity in transgenic *C. elegans* that express human amyloid-β peptides, delaying the paralysis in this model by 1.8 h and 1.1 h, respectively. We conclude that secondary compounds of *G. uralensis* may become interesting drug candidates for the treatment of Alzheimer’s disease, which, however, need further analysis in other model systems.

**Abbreviations**

- Aβ: amyloid-beta
- AD: Alzheimer’s disease
- EGCG: (−)-epigallocatechin gallate
- GA: glycyrrhizic acid
- GUE: *Glycyrrhiza uralensis* extract
- GRA: glycyrrhetinic acid
- ILG: isoliquiritigenin
- LG: liquiritigenin
- NGM: nematode growth medium
- NMDA: N-methyl-D-aspartate
- PT50: median paralysis time
- TCM: traditional Chinese medicine

**Introduction**

AD occurs with a prevalence of 5% in people over 60 and the risk increases with age. About 36 million people were estimated to suffer from this disease in 2010 according to Alzheimer’s disease International [1]. Despite these facts, there is still no effective cure for this disease today. Therefore, it is important to continue research for new possible disease modulating substances. Rich sources for potential therapeutic compounds are medicinal plants and their secondary metabolites, which have evolved as a means of protection against herbivores and microbes for plants producing them [2]. *Glycyrrhiza* species (eng. licorice; Fabaceae) have been known as medicinal plants and sweets for centuries both in Europe and Asia. The traditional uses include conditions of the respiratory system and the gastrointestinal tract [3]. Licorice is also used in many formulations of TCM and is the most frequently used herb in the Chinese Formulae Database [4]. Several studies indicate that *Glycyrrhiza* can be beneficial in the treatment of AD [5–7]. The most abundant secondary metabolite found in *Glycyrrhiza* species is the triterpene saponin GA. GA and its aglycone GRA have been shown to have anti-inflammatory and neuroprotective effects [8,9]. Next to the triterpenes, the plant also produces many flavonoids. These polyphenols can interact with biomolecules via their phenolic hydroxyl groups and thereby modify the function of many proteins. The flavonoid ILG shows anti-inflammatory [10–12] and neuroprotective ef-
fects [13, 14]. It is an NMDA receptor antagonist [15]. Additionally, ILG and its isomer LG exhibit antidepressant properties in mice [16] and both have been shown to inhibit neurotoxicity caused by Aβ in rat neurons [17, 18].

Aβ is one of the key proteins in AD and a potential drug target [19]. This peptide of 38–43 amino acids can take different conformations, build aggregates, and interact with cellular processes. Monomeric Aβ is not very stable in aqueous solutions and builds oligomers, which are toxic to cells [20]. Further aggregation leads to so-called senile plaques that are abundant in the brains of AD patients.

To better understand Aβ aggregation and its toxic effect in vivo, transgenic Caenorhabditis elegans models have been developed [21]. These worms express the human Aβ_{3–42} peptide in their muscles, where it aggregates and forms plaques. The toxicity to the surrounding muscle cells manifests in a paralysis phenotype. This in vivo model can assess the bioavailability of the compounds in contrast to in vitro aggregation experiments or toxicity assays with cell cultures. In the present study, the effect of Glycyrrhiza uralensis Fisch. ex DC and its secondary metabolites on Aβ aggregation and acute Aβ toxicity were studied using transgenic C. elegans strains.

**Results**

**A**

The HPLC analysis revealed several compounds (Fig. 1), 18 of which were further analyzed (Table 1). Substances 1–9, 11, 12, 15, and 18 could be identified according to their mass spectra and to published data for GUES [22–24]. GA and glycosylated forms of LG and ILG are among the most abundant compounds in this extract. In the water extract (Fig. 1A), fewer compounds were found. The more lipophilic substances 15, 17, and 18 were missing; 12, 14, and 16 were only present in traces. Also, the other compounds were less concentrated in the water extract, except for the saponins 7 and 8 that have a slightly higher abundance in the water extract compared to the methanol extract.

CL2006, a transgenic C. elegans strain that constitutively expresses the human Aβ peptide, was used to test the effect on Aβ aggregation in vivo. Aβ aggregates can be visualized by thioflavin S staining. Fig. 2 shows typical pictures for the thioflavin S staining of a control (A) and a GUE-treated worm (B). GUE (500 µg/mL) and the pure substances GA, GRA, LG, and ILG (50 µg/mL) all significantly reduced the number of Aβ aggregates (Fig. 2C). The effect of ILG (30% reduction) was similar to the positive control EGCG (a polyphenol from green tea) (100 µg/mL) that reduced the number of Aβ aggregates by 35%. GUE, GA, GRA, and LG had a weaker effect (a reduction by 14% for GUE, 15% for GA and LG, and 20% for GRA). For all treatments, the low concentrations had no significant effect; a significant effect was reached at a concentration of 50 µg/mL for GRA, LG, and ILG and at a concentration of 500 µg/mL for GUE, as can be seen on Fig. 2D. Therefore, the treatments can be considered dose-dependent.

The paralysis assay with strain CL4176 reveals the effect of the treatment on Aβ toxicity. Since the methanol that was used as a solvent for methanol GUE, ILG, LG, and ILG also had a paralysis delaying effect itself, the results of the compounds were compared to a methanol-treated control (Fig. 3A, B). EGCG and water GUE that were solved in water were compared to a water-treated control (Fig. 3C). EGCG was used as a positive control in a concentration of 100 µg/mL. It increased the median time to paralysis (PT_{50}) by 2.7 h (Table 2). Treatments with 500 µg/mL methanol GUE, 200 µg/mL GUE (both methanol and water), and 50 µg/mL ILG could also significantly increase the time until paralysis compared to the control. The water extract was less effective than the positive control, but with a 1.7 h increase in PT_{50}, it was more effective than the same amount of methanol extract.
Treatments with LG, GRA, and GA showed no significant increase. The control strain CL802 showed no paralysis when treated with extracts or pure compounds.

Discussion

All the tested substances had a significant effect on Aβ aggregation and lowered the number of plaques in the C. elegans strain CL2006, but the effects were much weaker in the test for acute Aβ toxicity. Only GUE and ILG could delay the paralysis in CL4176. In this C. elegans strain, amyloid toxicity was observed despite the small number of amyloid deposits, suggesting that

<table>
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<th>λ&lt;sub&gt;max&lt;/sub&gt; [min]</th>
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* Only present in the methanol extract

**Fig. 2** Effects of G. uralensis extract, glycyrrhizic acid, liquiritigenin, and isoliquiritigenin on the amyloid-β aggregation in C. elegans strain CL2006. A, B Typical images of the methanol-treated control worm (A) and a worm treated with 500 µg/mL methanol GUE (B), both stained with thioflavin S. Arrowheads point out the Aβ plaques. C Reduction in number of Aβ aggregates relative to the control. Control treated with 1% methanol, positive control with 100 µg/mL EGCG, samples for GUE with 500 µg/mL methanol extract, and other samples with 50 µg/mL of respective substance. D Dose-dependence of the treatments. *P < 0.05, **P < 0.01 compared to control.
inability of this study to reproduce the positive effects of LG seen in mice and cell cultures might be due to lack of a corresponding target or pathway in the chosen model organism.

The beneficial effect of ILG against Aβ toxicity might be achieved by NMDA receptor antagonism, antioxidative, or antiinflammatory properties of this compound. The reports about antioxidant properties of ILG are somewhat inconsistent. ILG has poor radical scavenging activity compared to other phenolics in licorice, but it can prevent LDL oxidation [29] and has shown beneficial effects in in vivo experiments [30]. Thus, the antioxidant properties in the given model system should be determined before any conclusions about the involvement of this mechanism can be made. It has also been shown that ILG can inhibit Aβ aggregation [31]. Possibly, all these effects contribute to the observed results, but the exact mechanism of action needs further elucidation.

ILG was only found in traces in GUE, but its glycosylated forms isoliquiritin and isoliquiritin apioside were far more abundant. Suggesting that the effect of GUE is based on ILG, it is possible that the glycosylated forms found in the extract have similar effects as ILG or are cleaved either by the Escherichia coli used as a food source for C. elegans or by the enzymes of C. elegans itself. Still there is the possibility that this compound alone is not responsible for the effect observed by the treatment with GUE. Therefore, some other not yet tested substances in this extract might either have a toxicity ameliorating effect against Aβ themselves or can synergistically contribute to the effect of ILG.

The observation that the water extract had an even stronger effect against the paralysis supports the suggestion that ILG is not the only active component of the extracts. The water extract contains less derivatives of ILG, but the suggestion that ILG is not the only active component of the extract would be strengthened if these substances were found to have a beneficial effect as ILG or are cleaved by the enzymes of C. elegans itself. The results shown here suggest that GUE or ILG might be potential drugs. The results for ILG are in accordance with earlier results in cell cultures [18]. Interestingly, the isomer LG fails to delay the paralysis in CL4176, although it could prevent apoptosis of cultured rat neurons at even lower concentrations than ILG [17]. The suggested mechanism of action for LG involves the estrogen receptor β and Notch-2 signaling, resulting in lower astrogliosis [28]. Although C. elegans expresses homologues of an estrogen receptor and Notch-2, the nervous system of C. elegans is much simpler. For example, it does not contain astrocytes. Therefore, the
the low bioavailability may also account for the low effects observed in the present study. It is also possible that part of the drugs were metabolized by the bacteria used as a food source, thereby lowering the effective drug concentration in the experiments. For further studies in C. elegans, it is therefore recommended to use dead bacteria.

In conclusion, it can be stated that ILG and GUE exhibit significant effects, counteracting the pathological effects of Aβ in C. elegans. While this nematode is a good tool for identifying new drug candidates and studying specific targets, it lacks the complexity of the vertebrate body. Therefore, ILG and possibly other constituents of G. uralensis should be further studied as possible drug candidates against AD in vertebrate systems.

Materials and Methods

Chemicals
EGCG (from green tea, purity ≥ 95%) and thioflavin S were purchased from Sigma-Aldrich Co. ILG (purity > 99%), GA (purity 75%), and glycyrrhetinic acid (GRA) (purity > 96%) were isolated by Prof. Dr. Yujie Fu. LG (purity > 98%) was a kind gift from Dr. Qiujuan Lu, Wangjing Science and Technology Park. The purity of ILG, GA, GRA, and LG was confirmed by HPLC as described in [35].

Plant material
Dried roots of G. uralensis were purchased in China and provided to our Institute by Prof. Dr. Thomas Efferth. The authenticity of the plant material was confirmed by DNA barcoding of the ITS sequence (GenBank accession number KF588200). Voucher specimens with the registration number P6873 are deposited at the Department of Biology, Institute of Pharmacy and Molecular Biotechnology, Heidelberg University, Germany.

Fifty g of the dried plant material were pulverized and extracted with 300 ml of water or methanol at moderate heat for 4 h. The water extract was lyophilized (DER 13.8 : 1), the methanol extract was reduced in a rotary evaporator to dryness (DER 11.1 : 1), and both extracts were stored at −20°C.

LC-MS/MS analysis of the extract
The composition of the methanol extract was analyzed on an LCQ-Duo ion trap mass spectrometer with an ESI source (Thermoquest) coupled to a Beckman Gold HPLC system (solvent module 125P, PDA detector 168) with a LiChroCART RP18 column (5 µm, 250 × 4 mm, Merck). A gradient of water and acetonitrile with 0.1% formic acid each was applied from 20% to 80% ACN in 20 min and isocratic for 10 min with the latter conditions. The flow rate was 1 ml/min throughout the whole run. The absorption maxima were determined in background-subtracted spectra by 32 Karat™ software (Beckman Coulter, Inc.). The MS operated in the negative mode with a capillary voltage of −10 V, a source temperature of 200°C, and high purity nitrogen as a sheath and auxiliary gas at a flow rate of 80 and 40 (arbitrary units), respectively. The ions were detected in a mass range of 50–2000 m/z. Major peaks were identified by comparison with published data [22–24].

Caenorhabditis elegans strains and culturing conditions
The C. elegans strains CL2006 (constitutively expressing Aβ in its muscle cells) [36], CL4176 (a temperature inducible strain expressing Aβ in its muscle cells) [25], and CL802 (standard control for CL4176) were obtained from Caenorhabditis Genetics Center. The worms were kept on NGM at 20°C (CL2006) or 16°C (CL4176 and CL802) and fed with E. coli OP50. For all experiments, C. elegans eggs, gained from gravid hermaphrodites by sodium hypochlorite treatment, were used to obtain an age-synchronized population. The Aβ aggregation assay was conducted in S-medium with E. coli OP50 (1 × 10⁶ cells/ml) as a food source [37].

Amyloid-β aggregation in Caenorhabditis elegans CL2006
The worms were treated with 500 µg/ml methanol extract of G. uralensis or 50 µg/ml pure substances (solved in methanol) on day two after hatching. The concentrations were chosen based on a dose-dependence experiment, where various concentrations between 5–100 µg/ml for pure compounds and 50–500 µg/ml for the extract were tested. A treatment with 100 µg/ml EGCG served as a positive control and the solvent methanol (1% of the final volume) as a negative control. On day six, the worms were fixed, followed by thioflavin S staining of the Aβ aggregates as described before [38, 39]. In contrast to the original method, 0.013% of thioflavin S in 50% ethanol was used. The Aβ aggregates in the head region of the worms were quantified using Nikon Eclipse Ni-E with an FITC filter and 40× objective. Pictures were acquired with a DS-Q1Mc camera, and deconvolution was performed with Huygens software by SVI.

Paralysis assay in Caenorhabditis elegans CL4176
The paralysis assay was conducted as described by Dostal and Link [40]. The methanol extract of G. uralensis (200 µg/ml and 500 µg/ml), the water extract of G. uralensis (200 µg/ml), or one of the pure substances, ILG, LG, GA, or GRA (50 µg/ml), was added to each NGM plate. EGCG (100 µg/ml) and the respective solvent (1%) were used as controls. The worms were kept at 16°C for 36 h, and then the temperature was upshifted to 25°C to induce the transgene expression. Paralysis was scored every 2 h for 12 h or, in the case of GUE 50 µg/ml, for 14 h starting at 24 h after the temperature upshift. Worms that failed to move as a response to a touch with a platin wire were counted as paralyzed.

Statistical analysis
Data are expressed as the mean ± SD of at least three independent experiments (n = 4 for the Aβ aggregation experiments, n = 3 for the paralysis assay). With the data from the paralysis assays, a survival analysis using the life table method was conducted to calculate the median paralysis times (PT₅₀). Differences were analyzed in StatView software using ANOVA and Dunnet’s test with p < 0.05 as the threshold for significance.

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Conflict of Interest

The authors declare no conflict of interest.

References

2 Wink M. Evolution of secondary metabolites from an ecological and molecular phylogenetic perspective. Phytochemistry 2003; 64: 3–19
8 Wang CY, Kao TC, Lo WH, Yen GC. Evolution of secondary metabolites from an ecological and molecular phylogenetic perspective. Phytochemistry 2003; 64: 3
22 Montoro P, Maldini M, Russo M, Postorino S, Placente S, Piazza C. Metabolomic profiling of roots of licorice (Glycyrrhiza glabra) from different geographical areas by ESI/MS/MS and determination of major metabolites by LC/ESI/MS and LC/ESI-MS/MS. J Pharm Biomed Anal 2011; 54: 535–544