Acetylpuerarin increases cell viability and reduces apoptosis in rat hippocampal neurons following oxygen-glucose deprivation/reperfusion

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Abstract. The effects of acetylpuerarin treatment following oxygen-glucose deprivation/reperfusion (OGD/R) were examined in rat hippocampal neurons in vitro and compared with the effects of acetylpuerarin in normoxic cells to confirm acetylpuerarin’s potential neuroprotective effects, including apoptosis inhibition. Wistar rat embryo hippocampal cells (day 18, E18) cultured for 8 days were subjected to 3 h OGD treatment, followed by reperfusion for 12, 24 or 36 h. For each time interval, a group of cells was left untreated (OGD/R-only groups) and treated with 0.1, 0.4 and 1.6 µM acetylpuerarin (OGD/R+acetylpuerarin). Neuron viability, apoptosis and caspase-8 and -3 activities were assessed by the reduction of 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT), 4′,6-diamidino-2-phenylindole (DAPI) and terminal deoxynucleotidyl-transferase-mediated dUTP nick end labeling (TUNEL) and spectrophotometric assays, respectively. Fas-ligand (Fas-L), Fas-associated death domain (FADD) and tumor necrosis factor-α (TNF-α) were determined by western blot analysis. Compared with control cells, OGD/R+acetylpuerarin cells treated with 0.1, 0.4 and 1.6 µM doses showed a concentration-dependent increase in hippocampal cell survival and viability by 69.93±2.28%, 81.49±2.13% and 85.28±2.38% at 12 h, 68.59±3.02%, 77.85±2.84% and 85.64±4.39% at 24 h and 69.70±1.70%, 77.21±3.21% and 83.90±2.12% at 36 h (P<0.05). Furthermore, OGD/R+acetylpuerarin cells exhibited a dose-dependent decrease in caspase-8 and -3 activation, TUNEL and DAPI-positive neurons and Fas-L, FADD and TNF-α expression. In conclusion, acetylpuerarin protects against OGD/R-induced neuronal apoptosis predominantly in the first 24 h following ischemia, which may be useful in mediating neuronal apoptosis in ischemic stroke patients.

Introduction

Cerebrovascular disease-related stroke affects ≤300 of every 100,000 members of the Chinese population (1) and the most common form of stroke, ischemic stroke, remains a leading cause of mortality and long-term disability in aging patients (2). Over the course of as little as 30 min, ischemia begins to produce oxygen-glucose deprivation (OGD)-induced neuronal apoptosis that may progress rapidly with increasing ischemia duration (3). Thus, the primary goal of the majority of contemporary medicines for stroke is the mediation of OGD-induced neuronal apoptosis during cerebral ischemia/reperfusion (4). Acetylpuerarin is a novel compound that has been reported to effectively attenuate the morphological changes leading to apoptosis in hippocampal cells following cerebral ischemia/reperfusion, with potential benefits even when applied a number of hours following the initial ischemic event (5). Unlike the extremely small treatment window (~3 h) of current United States Food and Drug Administration-approved therapies for stroke, including tissue plasminogen activator (6-8), acetylpuerarin may provide an alternative for stroke treatment at advanced stages (9). Prior to clinical implementation, a greater understanding of the dose-effects and mechanistic action of acetylpuerarin on hippocampal cell apoptosis during ischemia/reperfusion is required.

Acetylpuerarin is a modern synthetic derivative of puerarin (daidzein-8-C-glucoside) with higher lipid solubility than the naturally occurring compound, allowing it to better penetrate the blood-brain barrier (Fig. 1). Puerarin is an isoflavonoid compound used for treating cardiovascular disorders derived from the Chinese medical herb Radix puerariae, known as ‘Ge Gen’, from the root of the kudzu plant. Treatment of puerarin has been reported to effectively reduce symptoms following acute ischemic stroke with few side effects in a number of preliminary in vivo and in vitro studies (10-12). A number of studies have reported that acetylpuerarin exerts protective effects against ischemia-reperfusion injury in the hippocampus by mediating the cascade of events leading to OGD-induced damage (5,13,14). Thus, acetylpuerarin may be capable of mediating the degree
Materials and methods

Animal subjects and study design. Hippocampal cells were isolated from Wistar rat embryos (day 18, E18) purchased from the Laboratory Animal Center of the Shandong University (Shandong, China; Grade II, Certificate No. 20021015) and cultured for 8 days.

Cells were subjected to 3 h OGD treatment followed by reperfusion for 12, 24 or 36 h. For each time interval, a group of cells was left untreated (OGD/R) and OGD/R groups underwent OGD, as previously published (17). Each experiment was performed in cultures from three E18 rats and was repeated in triplicate.

Terminal deoxynucleotidyl-transferase-mediated dUTP nick end labeling (TUNEL) staining. Upon completion of reperfusion, cultured neurons were incubated with 5 µg/ml fluorescent DNA-binding dye DAPI (Sigma-Aldrich) at 37°C for 1 min. Staining solution was removed and apoptotic cells were visualized by fluorescence microscopy (Olympus DP-72; Olympus Corporation, Tokyo, Japan). Apoptotic cells were morphologically identified by cytoplasmic or nuclear shrinkage, chromatin condensation and DNA fragmentation. The percentage of apoptotic cells was determined in 5 randomly selected fields of ~500 neurons per field (magnification, x100). Experimental data was pooled from three coverslips and each experiment was conducted three times.

Western blot analysis for Fas-L, FADD and TNF-α expression. Total protein (50 µg) was extracted from samples from each group, separated by 12% sodium dodecyl sulfate polyacrylamide gel electrophoresis and blotted onto polyvinylidene fluoride membranes (Beijing Solarbio Science and Technology Co., Ltd, Beijing, China). Western blot analysis showed that the expression of Fas-L, FADD and TNF-α was significantly decreased in the OGD/R+acetylpuerarin 9 groups III-V, VII-IX and XI-XIII, respectively, according to the dosage concentrations suggested by Liu et al (5). Each experiment was performed in cultures from three E18 rats and was repeated in triplicate. The following formula (18) was used to calculate cell viability as percentage: (absorbance of treated cells / absorbance of normal cells) x 100%.

Acetylpuerarin administration. Acetylpuerarin was dissolved in DMSO ≤0.1% and administered at the onset of the ischemic period in concentrations of 0.1, 0.4 and 1.6 µM in the OGD/R+acetylpuerarin 9 groups III-V, VII-IX and XI-XIII, respectively, according to the dosage concentrations suggested by Liu et al (5). Each experiment was performed in cultures from three E18 rats and was repeated in triplicate.

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Membranes were probed with primary antibodies against Fas-L (1:400), FADD (1:500) and TNF-α (1:300; Santa Cruz Biotechnology, Inc., Santa Cruz, CA, USA) and peroxidase-conjugated secondary antibody (1:5,000; ZSGB-Bio, Beijing, China). β-actin (1:2,000; ZSGB-Bio) was used as a control. Band intensities were quantified using AlphaEaseFC software (Genetic Technologies, Miami, FL, USA).

Caspase-8 and -3 activity assay. Caspase-8 and -3 activities were assessed using FLICE/caspase-8 and caspase-3/CPP32 colorimetric assay kits, respectively, (BioVision, Inc., Milpitas, CA, USA), according to the instructions suggested by the manufacturer. Briefly, 100 or 50 µg of total protein were incubated with 10 mM dithiothreitol and IETD-pNA for caspase-8 and -3 activity quantification (caspase-8 substrate; 200 µM final concentration) or MDEVD-pNA (caspase-3 substrate; 200 µM final concentration) at 37˚C for 2 h. Samples were determined at 405 nm in a microplate reader (Therm MK3; Finland). The final reading did not include the background reading of cell lysates and buffers. Enzymatic activity was expressed in arbitrary units of OD per mg protein.

Statistical analysis. All data were analyzed using SPSS version 16 (SPSS, Inc., Chicago, IL, USA) and expressed as means ± SEM. Between group and multiple group differences were analyzed using Student’s t-tests. P<0.05 was considered to indicate a statistically significant difference.

Results

Acetylpuerarin increases neuron viability following OGD/R. The MTT assay revealed no significant changes in viability in the control and normoxic control groups treated with acetylpuerarin treatment at all times (P>0.05, data not shown). Compared with the OGD/R-only group, cell survival was 61.94±2.73% (OGD/R, 12 h), 60.61±3.29% (OGD/R, 24 h) and 57.77±0.66% (OGD/R, 36 h). Compared with the control group, OGD/R+acetylpuerarin groups treated with acetylpuerarin doses of 0.1, 0.4 and 1.6 µM increased cell survival and viability by 69.93±2.28%, 81.49±2.13% and 85.28±2.38% at 12 h, respectively; 68.59±3.02%, 77.85±2.84% and 85.64±4.39% at 24 h, respectively and 69.70±1.70%, 77.21±3.21% and 83.90±2.12% at 36 h, respectively (P<0.05; Fig. 2). Higher acetylpuerarin concentrations enhanced neuron survival more efficiently in a concentration-dependent manner. Similar results were obtained by morphological analysis. The control group exhibited round cell bodies with clear edges and fine dendritic network. OGD/R groups exhibited a decreasing number of neurons, shrinkage of cell bodies and disruption of dendritic networks. Acetylpuerarin mitigated the morphological manifestations of cell damage (data not shown).

Acetylpuerarin reduces apoptosis following OGD/R. DAPI staining revealed that <8% of control-only cells showed signs of apoptosis at any time point; however, OGD/R treatment resulted in significant time-dependent increases in apoptotic cell numbers, peaking at 24 h (P<0.01; Fig. 3). In the OGD/R-only group at 24 h, 75.85±7.59% of remaining cells showed signs of apoptosis. Comparatively, OGD/R+acetylpuerarin groups treated with 0.4 µM acetylpuerarin doses exhibited reductions in DAPI positive neurons to 63.01±7.35%, 59.06±5.98% and 51.05±5.98%, respectively (all P<0.05). Thus, increasing the concentration of acetylpuerarin exhibited no significant effects on apoptosis rates in cells treated with OGD/R following 24 h reperfusion (all P>0.05).

These results were confirmed with TUNEL staining, indicating that 5% of the control-only group cells exhibited apoptosis under basal conditions. TUNEL-positive cells were markedly increased in the OGD/R-only group at 24 h and the AI was 68.79±10.01%. Acetylpuerarin treatment decreased the number of TUNEL-positive cells to 52.30±9.73%, 46.08±13.42% and 39.04±7.29% in all OGD/R+acetylpuerarin groups treated with 24 h reperfusion (all P<0.05; Table I).

Acetylpuerarin inhibited OGD/R-induced caspase-8 and -3 activation. Caspase-8 was significantly increased in hippocampal neurons following OGD/R treatments at 24 h and acetylpuerarin treatment led to a concentration-dependent decrease in the expression of enzymatic activity of caspase-8.
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Table I. Effects of acetylpuerarin on AI (24-h reperfusion).

<table>
<thead>
<tr>
<th>Group</th>
<th>Acetylpuerarin concentration, µM</th>
<th>AI, %</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>0</td>
<td>4.57±2.48</td>
</tr>
<tr>
<td>OGD/R</td>
<td>0</td>
<td>68.79±10.06a</td>
</tr>
<tr>
<td>OGD/R+acetylpuerarinb</td>
<td></td>
<td></td>
</tr>
<tr>
<td>VII</td>
<td>0.1</td>
<td>52.30±9.73c</td>
</tr>
<tr>
<td>VIII</td>
<td>0.4</td>
<td>46.08±13.42c</td>
</tr>
<tr>
<td>IX</td>
<td>1.6</td>
<td>39.04±7.29c</td>
</tr>
</tbody>
</table>

aP<0.05 vs. control-only group. bTreated with 3 h oxygen-glucose deprivation and 24 h reperfusion. cP<0.01, vs. control-only group. Values are expressed as the mean ± SD of three independent experiments. AI, apoptotic index; OGD/R, oxygen-glucose deprivation/reperfusion.

As caspase-3 is considered to be a prototypical caspase and a key effector of programmed cell death (19), enzymatic activity of caspase-3 was also determined. As shown in Table II, exposure to OGD significantly enhanced caspase-3 proteolytic activity by 24 h, while acetylpuerarin significantly diminished the activation of caspase-3 activity in a concentration-dependent manner.

Acetylpuerarin inhibited OGD/R-induced Fas-L, FADD and TNF-α protein expression. Western blot analysis indicated that Fas-L levels were induced by OGD treatment in hippocampal neurons by 24 h post-reperfusion (Fig. 4A). This Fas-L induction was significantly attenuated by acetylpuerarin treatment in a concentration-dependent manner. High levels of FADD protein were detected in neurons following OGD/R 24 h. However, FADD expression was significantly downregulated by acetylpuerarin (Fig. 4B). OGD/R-induced apoptosis was accompanied by a significant increase in TNF-α (Fig. 4C). Western blot analysis also revealed that acetylpuerarin inhibited TNF-α expression induced by OGD/R 24 h in a dose-dependent manner.

Discussion

A model of ischemic OGD/R in hippocampal neurons was used to assess the potential protective role of acetylpuerarin in ischemic injury. Acetylpuerarin not only increased neuron viability, but also decreased DAPI and TUNEL-positive cells, caspase-8 and -3 activity and Fas-L protein expression, FADD and TNF-α induced by OGD/R. Thus, acetylpuerarin may be capable of mediating the protection to hippocampal cells by decreasing neuronal apoptosis during acute ischemia and may be useful in clinical treatment development, pending further assessment of the clinical effects of the drug.

The cell-extrinsic apoptosis pathway Fas→FADD→caspase-8→caspase-3 is activated by binding of pro-apoptotic ligands, including Fas/CD95L (TNFSF6) or Apo2L/TRAIL (TNFSF10) to their cognate death domain-containing receptors on the surface of target cells (20-25). In a previous study,
acetylpuerarin was shown to alleviate morphological damage and increase neuron viability, consistent with the current findings (5). The current study is also consistent with reports indicating that TNF-α activation, followed by the inhibition of subsequent inflammatory responses, apoptosis formation (active caspase-3) and neutrophil activation are central to the

Table II. Effects of acetylpuerarin on caspase-3 and caspase-8 activities (24-h reperfusion).

<table>
<thead>
<tr>
<th>Group</th>
<th>Acetylpuerarin concentration, µM</th>
<th>Caspase-3</th>
<th>Caspase-8</th>
</tr>
</thead>
<tbody>
<tr>
<td>OGD/R</td>
<td>0</td>
<td>4.36±0.26</td>
<td>4.27±0.29</td>
</tr>
<tr>
<td>OGD/R + acetylpuerarin&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.1</td>
<td>3.78±0.13&lt;sup&gt;b&lt;/sup&gt;</td>
<td>3.40±0.28&lt;sup&gt;c&lt;/sup&gt;</td>
</tr>
<tr>
<td></td>
<td>0.4</td>
<td>3.14±0.26&lt;sup&gt;b&lt;/sup&gt;</td>
<td>2.80±0.32&lt;sup&gt;c&lt;/sup&gt;</td>
</tr>
<tr>
<td></td>
<td>1.6</td>
<td>2.28±0.27&lt;sup&gt;b&lt;/sup&gt;</td>
<td>2.55±0.23&lt;sup&gt;c&lt;/sup&gt;</td>
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<sup>a</sup>Treated with 3 h oxygen-glucose deprivation and 24 h reperfusion. <sup>b</sup>P<0.05, vs. caspase-3 activity in the OGD/R 24 h group. <sup>c</sup>P<0.05, vs. caspase-8 activity in the OGD/R 6 h group. Values are expressed as the mean ± SD of three independent experiments. OGD/R, oxygen-glucose deprivation/reperfusion.

Figure 4. Effects of acetylpuerarin on OGD/R-induced Fas-L, TNF-α and FADD protein expression. Relative levels of (A) Fas-L, (B) FADD and (C) TNF-α in hippocampal neurons were determined by western blot analysis. β-actin was used as a protein loading control. Values are expressed as the mean ± SD of three independent experiments. OGD/R, oxygen-glucose deprivation/reperfusion.
In considering these results, it is important to consider that in vivo results may vary and that systemic side effects may be associated with treatment. Thus, the effects of acetylpuerarin in vivo merit further clinical investigation. Since the effects of acetylpuerarin were examined over a relatively short 24 h period following reperfusion, it is also important to determine, in future studies, whether acetylpuerarin is capable of mediating neuronal death and promoting functional neurological recovery over longer periods, a current topic of research in our laboratory.

The current findings demonstrated that acetylpuerarin treatment induced dose-dependent neuroprotection against OGD/R-induced neuronal cell death by inhibiting Fas-L, FADD and TNF-α pathway-mediated apoptosis. Furthermore, caspase-3 and -8 activities were inhibited, providing further evidence that acetylpuerarin impacts a number of pathways known to induce apoptosis following OGD/R. Thus, acetylpuerarin may be a promising future candidate for the clinical treatment of acute ischemic stroke, pending further mechanistic verification and clinical investigations.

Acknowledgements

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References


