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Naphthazarin Protects Against Glutamate-Induced Neuronal Death Via Activation of the Nrf2/ARE pathway

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Abstract

Nuclear factor E2-related factor 2 (Nrf2)/ antioxidant response element (ARE) pathway is an important cellular stress response pathway involved in neuroprotection. We previously screened several natural phytochemicals and identified plumbagin as a novel activator of the Nrf2/ARE pathway that can protect neurons against ischemic injury. Here we extended our studies to natural and synthetic derivatives of plumbagin. We found that 5,8-dimethoxy-1,4-naphthoquinone (naphthazarin) is a potent activator of the Nrf2/ARE pathway, up-regulates the expression of Nrf2-driven genes in primary neuronal and glial cultures, and protects neurons against glutamate-induced excitotoxicity.

Keywords

Nrf2; glutamate; naphthazarin; antioxidant response element

1. Introduction

Extracellular accumulation of glutamate has been linked to neuronal death associated with acute damage (i.e. stroke, spinal cord and head trauma), as well chronic neuropathologies such as Alzheimer's, Parkinson's and Huntington's diseases [1]. Glutamate excitotoxicity results from the over-stimulation of postsynaptic glutamate receptors with massive calcium influx leading to rapid mitochondrial calcium overload and mitochondrial generation of reactive oxygen species (ROS) [1,2]. The increased ROS levels cause oxidative damage within the postsynaptic neurons, and can be released from them propagating the damage to surrounding cells [1,2].

The nuclear factor E2-related factor 2 (Nrf2)/ antioxidant response element (ARE) pathway is a critical signaling pathway regulating antioxidants and phase II detoxification enzymes, such as heme oxygenase 1 (HO1), NAD(P)H quinone oxidoreductase 1 (NQO1), and

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thioredoxin reductase 1 (TrxR1). The importance of Nrf2 in protecting cells against toxins and in pathological conditions characterized by enhanced oxidative stress has been clearly demonstrated using siRNA methodologies and Nrf2 knockout mice [3]. Furthermore, activation of the Nrf2/ARE pathway has beneficial roles in cell culture and animal models of neurodegenerative diseases, including Alzheimer's and Parkinson's diseases [3–5].

We previously screened natural botanical pesticides for their ability to activate adaptive stress response pathways, and identified plumbagin as a novel Nrf2/ARE activator and anti-ischemic compound [6]. In the present study we further our findings testing natural and synthetic plumbagin analogues with the aim to identify stronger Nrf2 activators and potential neuroprotective agents. Among the various analogues examined, naphthazarin showed higher levels of Nrf2/ARE activation than plumbagin in stable ARE reporter HepG2 cells. Naphthazarin induced expression of endogenous ARE-mediated target genes in primary neuronal and astrocytes cultures. Furthermore, naphthazarin protected neurons against glutamate-induced cell death and reduced α -fodrin cleavage in primary neurons.

2. Materials and Methods

2.1 Cell culture

The Nrf2 reporter cell line ARE-*bla* HepG2 was purchased from Invitrogen (Carlsbad, CA, USA) and maintained in Dulbecco's Modified Eagle's Medium (DMEM) supplemented with 10% dialyzed fetal bovine serum, 2 mM GlutaMAX, 0.1 mM non essential amino acids, 25 mM HEPES, 1% penicillin/streptomycin, and 5 μ g/ml of blasticidin (Invitrogen) at 37°C in a humidified 5% CO₂ atmosphere. Blasticidin was removed from the medium during experimental treatments. Neuronal cultures were generated as previously described [6] and plated at 0.1×10^6 cells/cm² in Neurobasal medium supplemented with B27 (Invitrogen). Experiments were performed using 9- to 10-day-old cultures. Primary astrocytes cultures were generated from postnatal Sprague-Dawley rats as described [7]. Cultures were maintained in DMEM with 10% serum for 2–3 weeks, at which time microglia were removed by shaking and adherent astrocytes were passaged.

2.2. Cell viability and Nrf2/ARE reporter assay

ARE-*bla* HepG2 cells (5000 cells in 100 μ l of medium) were seeded in 96-well plates 24 hours before treatment with the indicated concentration of 5,8-dihydroxy-1,4-naphthoquinone (naphthazarin) (Sigma, St. Louis, MO). Cell viability was measured 24 hours post-treatment using Celltiter 96[®] AQUEOUS One Solution reagent (Promega, Madison, WI, USA). β -lactamase activity was measured according to the manufacturer's instructions (Invitrogen).

2.3. Western blot analysis

Cell lysates were processed for Western blot analysis as previously described [6]. Equal amounts of protein were separated by SDS-PAGE, transferred onto nitrocellulose membranes, and probed with antibodies against heme oxygenase 1 and thioredoxin reductase 1 (Santa Cruz Biotechnology, Santa Cruz, CA), glial fibrillary acid protein (GFAP) (Abcam, Cambridge, MA), α -fodrin (Millipore, Billerica, MA), and GAPDH (clone 6C5; Advanced Immunochemical Inc, Long Beach, CA). Immunoreactive bands were detected using HRP-conjugated secondary antibody (Amersham Bioscience, Piscataway, NJ) and chemiluminescence (Pierce). Autoradiograms were quantified using image J software.

2.4. Statistical analysis

Statistical analysis was performed by Student's *t*-test or analysis of variance (ANOVA) as appropriate. The results are expressed as mean + S.E.M. Values of *p*<0.05 were considered statistically significant.

3. Results

3.1. Naphthazarin activates the Nrf2/ARE pathway in stable ARE –*bla* Hep G2 cell line

Plumbagin (5-hydroxy-2-methyl-1,4-naphthoquinone) (Figure 1A structure 1) is a natural quinoid with several therapeutic properties [6,8,9]. We investigated the ability of various structural analogs of plumbagin (Fig. 1A) for their ability to induce ARE-mediated transcription using the stable ARE reporter cell line ARE–*bla* Hep G2. None of the compounds tested showed toxicity at the concentration utilized for the reporter assay (Fig. 1B). Compounds 7-Hydroxy-1-Tetralone, 5-Methoxy-1-methyl-2-Tetralone, 4,7-Dihydroxy-3-methyl-indan-1-one, 1,4-Dihydroxy-7,8-dihydro-6H-benzocycloheptene-5,9-dione and 2-Hydroxy-2-(2-methyl-2,3-dihydro-1H-indol-5-yl)-1H-indene-1,3(2H)-dione showed similar or lower ARE-inducing activity compared to plumbagin (Fig. 1C). However 5,8-dihydroxy-1,4-naphthoquinone (naphthazarin) displayed significantly higher activity than plumbagin (Fig. 1C).

3.2. Naphthazarin induces endogenous ARE-mediated target genes in primary neuronal cultures

We then tested the ability of naphthazarin to induce the expression of endogenous ARE-mediated target genes in primary neuronal cultures. Neurons and astrocytes were treated with different concentrations of naphthazarin for 6 and 24 hours and levels of HO1 and TrxR1, respectively, were determined by immunoblotting. In both cell types naphthazarin increased the expression levels of the proteins in a concentration-dependent fashion (Fig. 2B) validating its use as an Nrf2/ARE activator.

3.3. Naphthazarin protects against neuronal cell death and reduces α -fodrin cleavage induced by glutamate in rat primary neurons

In order to test the neuroprotective potential of naphthazarin we used an acute excitotoxic cell culture model. Neuronal cultures were treated with the indicated concentrations of naphthazarin for 6 hours and were then subjected to glutamate challenge for 16 hours. A significant decrease in cell survival (Fig. 3A) paralleled by enhanced α -fodrin cleavage (Fig. 3B) was observed in glutamate-treated cells. However, naphthazarin pretreated cells displayed significantly higher resistance to glutamate excitotoxicity (Fig. 3A), as well decreased α -fodrin cleavage (Fig. 3B).

4. Discussion

Activation of the Nrf2/ARE pathway has been shown to counteract neurotoxicity resulting from glutathione depletion, intracellular calcium overload, mitochondria dysfunction, increased ROS and lipid peroxidation in different experimental models [10–11]. Given its multiple roles in regulating cell homeostasis the Nrf2/ARE pathway is considered a promising therapeutic target for neurodegenerative disorders [3,4]. Over the past decades several phytochemicals have been identified as Nrf2 inducers and neuroprotective agents. For example, sulforaphane protects against hydrogen peroxide-, glutamate-, and 6-hydroxydopamine-induced toxicity in cultured neurons and *in vivo* [12–13]. The naphthoquinones plumbagin and curcumin reduce brain damage and ameliorate the associated neurological deficits in a model of focal ischemia through Nrf2-dependent

mechanisms [6, 14]. Licochalcone has anti-inflammatory and cytoprotective effects that are attenuated by siRNA-mediated Nrf2 silencing [15].

In most cases the study of potential neuroprotective agents is restricted to pure neuronal cultures. However astrocytes play a critical role in maintaining normal brain physiology and responding to injury or diseases. In addition to structural and metabolic support, astrocytes are important regulators of extracellular neurotransmitters and ion concentrations [16]. Furthermore, Nrf2/ARE responsive genes appear to be preferentially activated in astrocytes, which consequently may have stronger detoxification and antioxidant potential than neurons [17–19]. It is reasonable that agents able to modulate the Nrf2/ARE pathway both in neurons and astrocytes may provide a better protective spectrum. Here we show that, contrary to many known Nrf2 activators, naphthazarin can induce the expression of endogenous HO-1 and TrxR1 proteins both in neurons and astrocytes. Furthermore, we found that naphthazarin protects neurons against glutamate-induced cell death. Glutamatergic intracellular calcium mobilization in neuronal cells is crucial for neurotoxicity-induced death. In addition to impairment of mitochondrial functions and induction of oxidative stress, the elevation of intracellular calcium levels leads to the overstimulation of proteolytic enzymes such as calpains and caspases [1, 20]. We found that in our experimental conditions glutamate induced the accumulation of the calpain-specific 150 kDa α -fodrin cleavage product (Fig. 3B). Notably, we found that naphthazarin attenuates the generation of the α -fodrin cleavage product suggesting that naphthazarin may reduce intracellular calcium levels. Although it is not known whether naphthazarin regulates neuronal calcium homeostasis via Nrf2/ARE-responsive genes, previous observations revealed an important role for Nrf2 in the maintenance of calcium homeostasis by showing that Nrf2^{-/-} neurons are more sensitive to increased intracellular calcium concentrations induced by ionomycin [21].

In conclusion, our findings identify naphthazarin as a novel potent Nrf2/ARE activator in neurons and astrocytes. In addition to inducing detoxification and antioxidant proteins, naphthazarin attenuates intracellular calcium-dependent protease activity and protects cortical neurons against glutamate-mediated excitotoxicity. These findings imply that naphthazarin may have therapeutic potential for various neurological diseases.

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Abbreviations

ARE	antioxidant response element
HO1	heme oxygenase 1
Nrf2	Nuclear factor E2-related factor 2
NQO1	NAD(P)H quinine oxidoreductase 1
ROS	reactive oxygen species
TrxR1	thioredoxin reductase 1

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Highlights

- Naphthazarin activates the Nrf2/ ARE pathway
- Naphthazarin induces Nrf2-driven genes in neurons and astrocytes
- Naphthazarin protects neurons against excitotoxicity

Figure 1. Naphthazarin activates the Nrf2/ARE pathway in ARE-*bla* Hep G2 cells

A. Chemical structure of plumbagin and its analogs tested in the study. 1: Plumbagin, 2: 2-Methyl-1-Tetralone, 3: 5-Methoxy-1-Tetralone, 4: 5,8-Dimethoxy-1-Tetralone, 5: 5-Hydroxy-1-Tetralone, 6: 7-Hydroxy-1-Tetralone, 7: 5-Methoxy-1-methyl-2-Tetralone, 8: 5,8-Dihydroxy-1,4-naphthoquinone, 9: 1,2,3,4-Tetraoxo-1,2,3,4-tetrahydronaphthalene Dihydrate, 10: 4,7-Dihydroxy-3-methyl-indan-1-one, 11: 1,4-Dihydroxy-7,8-dihydro-6H-benzocycloheptene-5,9-dione, 12: 2-Hydroxy-2-(2-methyl-2,3-dihydro-1H-indol-5-yl)-1H-indene-1,3(2H)-dione, 13: 4-Benzhydryl-N-phenyl-1-piperazinecarboxamide, **B and C.** ARE-*bla* Hep G2 cells were treated with 5 μ M sulforaphane (SF) or 1 μ M of the various compounds. After 24 hours, cell viability (**B**) and ARE reporter activity (**C**) were assessed using MTS and β -lactamase activity assays respectively. Data are mean and S.E.M. * $p < 0.05$, ** $p < 0.01$ and *** $p < 0.001$ vs. control.

Figure 2. Naphthazarin induces ARE-mediated target genes in rat primary cultures
Representative immunoblots and quantification data of HO1 and TrxR1 expression in neurons (A) and astrocytes (B) following treatment with the indicated concentrations of naphthazarin. Data are mean and S.E.M. (n=3). *p<0.05, **p<0.01 and ***p<0.001 vs. control.

Figure 3. Naphthazarin protects neurons against glutamate excitotoxicity

Primary rat neuronal cultures were pretreated with the indicated concentration of naphthazarin for 6 hours and exposed to 200 μM of glutamate for 16 hours. Cell viability was determined by MTS assay (A), and α-fodrin cleavage by western blot (B). GAPDH was used as control for equal loading. Data are mean and S.E.M. (n=3). ***p<0.001 vs. control; # p<0.05 and ### p< 0.001 vs. glutamate treated cells.