Daidzein induces neuritogenesis in DRG neuronal cultures

Shih-Hung Yang1,2, Chih-Chen Liao1, Ying Chen3, Jhih-Pu Syu1, Chung-Jiuan Jeng4*† and Seu-Mei Wang2*†

Abstract

**Background:** Daidzein, a phytoestrogen found in isoflavone, is known to exert neurotrophic and neuroprotective effects on the nervous system. Using primary rat dorsal root ganglion (DRG) neuronal cultures, we have examined the potential neurite outgrowth effect of daidzein.

**Methods:** Dissociated dorsal root ganglia (DRG) cultures were used to study the signaling mechanism of daidzein-induced neuritogenesis by immunocytochemistry and Western blotting.

**Results:** In response to daidzein treatment, DRG neurons showed a significant increase in total neurite length and in tip number per neuron. The neuritogenic effect of daidzein was significantly hampered by specific blockers for Src, protein kinase C delta (PKCδ) and mitogen-activated protein kinase/extracellular signal-regulated kinase kinases (MEK/ERK), but not by those for estrogen receptor (ER). Moreover, daidzein induced phosphorylation of Src, PKCδ and ERK. The activation of PKCδ by daidzein was attenuated in the presence of a Src kinase inhibitor, and that of ERK by daidzein was diminished in the presence of either a Src or PKCδ inhibitor.

**Conclusion:** Daidzein may stimulate neurite outgrowth of DRG neurons depending on Src kinase, PKCδ and ERK signaling pathway.

**Keywords:** Daidzein, DRG neuronal culture, Neurite outgrowth, Signaling mechanism

**Background**

Daidzein, found mainly in soy food products and herbs like red clover [1,2], is one of the most studied and most potent phytoestrogens. Phytoestrogens are estrogenic compounds of plant origin, and have structures and functions similar to the mammalian endogenous hormone estrogen [3]. Both estrogen and phytoestrogens can bind to intracellular estrogen receptors (ERs) to trigger downstream signal transduction pathways and achieve various biological functions [4]. Phytoestrogen acts mainly as an ER agonist. It may also function as an antagonist, by inhibition of aromatase activity in breast cancer cells, and blockage of estrogen uptake by uterine cells [5]. This mixed ER agonist/antagonist property probably explains the potential benefit of phytoestrogen in breast cancer prevention [6-8].

In response to ligand binding, ERs can signal through both genomic (classical) and non-genomic (non-classical) pathways [9,10]. In the genomic pathway, upon binding to estrogen, ERs dimerize and interact with the estrogen responsive element (ERE) in the regulatory regions of estrogen responsive genes, thereby regulating the transcription of E2-sensitive genes, e.g. c-fos, TGF-α, and angiotensinogen [9]. The non-genomic pathway involves the activation of other signal transduction pathways that lead to rapid and diverse physiological responses, including calcium and potassium influxes through cell membrane, and activation of second messenger systems such as cAMP/PKA, MAPK, PI3K/Akt, and G protein [10,11]. The precise mechanisms of non-genomic effects of estrogen are not clear and have been suggested to be mediated by membrane-associated ERα, ERβ, or the orphan G-protein-coupled receptor 30 (GPR30) [12,13]. Activation of non-nuclear ERα, for example, can stimulate endothelial cell proliferation via G protein, Src, and eNOS activation [14]. In cultured osteoblastic cells, daidzein has been suggested to activate a non-classical
Figure 1 Induction of neurite outgrowth of cultured rat DRG neurons by daidzein and NGF. DRG neuronal cultures were treated for 24 h with (A) DMSO, (B-H) various concentration of daidzein, or (F) NGF (100 ng/ml), then fixed and immunostained for NF-L. A-F, large neurons. G-H, small neurons. Scale Bar = 30 μm.
Figure 2 (See legend on next page.)
membrane ER-β pathway that involves phospholipase C-β2 (PLC-β2)/PKC and PI3K/cSrc [15].

Emerging evidence, however, indicates that for some of the phytoestrogen effects, ER activation may not be required. For instance, genistein, a rich phytoestrogen in soybeans, was shown to exert growth inhibitory effects in ER-negative breast cancer cells [16-18]. Compared to 17-β estradiol, the most biologically active estrogen in mammals, daidzein has a significantly lower affinity for both ER-α and ER-β [19]. Moreover, daidzein could induce anti anti-proliferative effects in both ER-positive and ER-negative pancreatic cells [20]. Together these observations raise the possibility that daidzein may also exert its pharmacological effect via an ER-independent signaling pathway.

Daidzein is known to exert significant neuronal protection and neurotrophic effects for a variety of cultured neuronal cells, e.g. hippocampal neurons, cortical neurons, dorsal root ganglion (DRG) neurons, and PC12 cells [21-24]. In hippocampal neuron, the neurotrophic mechanism involves ERβ-PKCa-GAP43 signaling. To further understand the diversity of the intracellular signaling mechanisms of daidzein, in the current study we focused on daidzein-induced neurite outgrowth in cultured DRG neurons. DRG culture is a well-characterized system for investigating the mechanism of neuritogenesis [25-27], and for screening neuroprotective drugs for peripheral neuropathies [28]. Studies using DRG cultures have shed light on the pathogenic mechanisms of peripheral nervous system diseases and the regeneration of spinal cord injury [29-31]. Here we showed that in cultured DRG neurons daidzein induced notable neuritogenesis via an ER-independent signaling pathway. In addition, we presented several lines of evidence suggesting that daidzein-induced neurite outgrowth in DRG neurons may be primarily mediated by the Src kinase, PKCδ and ERK signaling pathway.

Methods

Drugs

Daidzein was purchased from the Pharmaceutical Industry Technology and Development Center (New Taipei City, Taiwan). Nerve growth factor (NGF) was purchased from R&D Systems (Minneapolis, MN, USA). Dimethyl sulfoxide (DMSO), antibodies for neurofilament light chain (NF-L) and ER-α, were purchased from Sigma Chemical Co. (St. Louis, MO, USA). L-15 Leibovitz medium was purchased from Gibco (Grand Island, NY, USA). ER antagonists ICI182780, tamoxifen and G-protein coupled receptor 30 (GPR-30) antagonist G15 were obtained from TOCRIS (TOCRIS Cookson Inc., Bristol, UK). Src kinase inhibitor PP2, MEK inhibitor U0126, PKC inhibitor staurosporin, and PKCδ inhibitor rottlerin were purchased from Biomol Research Laboratory Inc. (Plymouth meeting, PA, USA).

Animals

Postnatal day 2 Wistar rat pups were purchased from the Facility for Animal Research of the National Taiwan University. All procedures were in accordance with the Guidelines for the Care and Use of Mammals in Neuroscience and Behavioral Research (National Research Council 2003) and approved by the Institutional Animal Care and Use Committee (IACUC) of National Taiwan University, College of Medicine.

Cell culture

DRG cultures were prepared as described previously [25]. Briefly, P2 rat pups were put on ice and then decapitated to harvest DRG. DRG were then dissected out under microscope and dissociated with 0.25% trypsin and 0.05% collagenase (Sigma) in HBSS solution, for 30 min at 37°C. These ganglia were then dispersed by mechanically trituration with glass pipettes. The pellet from low-speed centrifugation was re-suspended in phenol-red free L-15 Leibovitz media, supplemented with 1.2 g/L of NaHCO3, 5% fetal bovine serum, 100 IU/mL of penicillin, and streptomycin (Gibco). Cells were plated on collagen-coated coverslips for immunocytochemistry, and on 35 mm uncoated culture dishes for protein quantification by Western blot. The medium was changed to serum free L-15 for day in vitro (DIV) 2 cultured DRG cells. Cultures were maintained at 37°C in an atmosphere of 95% air and 5% CO2.

Cell survival assay

The MTT assay, a colorimetric assay for measuring the activity of mitochondrial enzymes, was used to examine whether cell viability was affected by treatment [32]. In each well of 24-well culture plates, 2 × 10^4 cells were plated and treated with 0.1% DMSO, different concentration of daidzein or different kinase inhibitors for 24 h. After treatments, cells were washed with phosphate-buffered saline (PBS; 137 mM NaCl, 2.7 mM KCl, 1.5 mM KH2PO4, 8 mM Na2HPO4, pH 7.4), and incubated in 0.5 mg/ml of 3-[4, 5-dimethylthiazol-2-yl]-2, 5-diphenyltetrazolium bromide (MTT) solution for 4 h to allow the conversion of

(See figure on previous page.)

Figure 2 Quantitative analysis of total neurite length and tip number of neurite branches per neuron following various treatments with DMSO, daidzein, or NGF. Significant neurite lengthening and branching were observed in both large (A and B) and small DRG neurons (C) treated with 30 μM daidzein or 100 nM NGF. Three independent experiments were performed. Ten neurons were chosen from each group in one representative experiment for analysis. *, p < 0.05 vs DMSO controls. **, p ≤ 0.01 vs DMSO controls.
Figure 3 Effect of different inhibitors on cell viability and neuritogenesis. DRG neurons were treated with 0.1% DMSO, 30 μM diadzein, 10 μM PP2, 10 μM U0126, 10 μM tamoxifen, 1 μM ICI182780 or 100 nM G15 for 24 h. They were then assayed by the MTT test (A), or immunostained for NF-L and analyzed for total neurite length (B) and tip numbers (C).
Figure 4 (See legend on next page.)

A

![Images of neuronal cultures with DMSO, Daidzein, Daidzein + ICI 182780, Daidzein + G15, Daidzein + PP2, and Daidzein + U0126 treatments.]

B

![Bar graphs showing the total neurite length and tip number for each treatment group.]

Legend:

- **: p < 0.01
- #: p < 0.05

(See legend on next page.)
MTT into the purple formazan product by mitochondrial dehydrogenases. The reaction medium was then removed and the cells were lysed with DMSO for 5 min. The absorbance was read at 590 nm with a spectrophotometer (Beckman Coulter Inc., Fullerton, CA).

Drug treatment
DIV 3 cultured DRG cells received either daidzein at a concentration of 10 μM, 30 μM, 50 μM, or 100 μM, vehicle solution DMSO (final concentration of 0.1%), or NGF of 100 ng/mL, in order to study the effect of daidzein on neurite outgrowth.

For inhibitor assay, one of following inhibitors was reacted 30 min before the addition of daidzein: estrogen receptor antagonist ICI182780 at 1 μM; tamoxifen at 10 μM; GPR-30 antagonist G15 at 100 nM; Src kinase inhibitor PP2 at 10 μM; PKC inhibitor staurosporin at 100 nM; PKC α/β inhibitor G66976 at 1 μM; PKCε inhibitor εV1-2 at 2 μM; PKCδ inhibitor rottlerin at 2 μM; MEK inhibitor U0126 at 10 μM.

Immunocytochemistry
After 24 h of DMSO or daidzein treatment, DRG neurons on cover glasses were fixed for 10 min with 10% formalin in PBS. After washed with PBS, cells were then permeabilized and blocked with 0.15% Triton X-100 and 5% non-fat milk in PBS for 1 h. DRG neurons were then incubated in mouse anti-NF-L antibody overnight at 4°C. After PBS wash, cells were incubated in biotin-conjugated goat anti-mouse IgG (Vector, Burlingame, CA, USA) at 1:50 dilution. Following washes with PBS containing 0.1% Tween-20, alkaline phosphatase conjugated secondary antibodies at 1:750 dilution. Following washes with PBS containing 0.1% Tween-20, alkaline phosphatase conjugated secondary antibodies at 1:750 dilution (Promega, Madison, WI, USA) were added for an hour at room temperature, and the bound antibodies visualized using enzyme-substrate reaction (substrate: 3.3 mg/ml nitro blue tetrazolium and 1.65 mg/ml 5-bromo-4-chloro-3-indoly phosphate in 100 mM NaCl, 5 mM MgCl2, 100 mM Tris-base, pH 9.5).

Quantification
Immunostained neurons were photographed at 20× or 40× (for morphological demonstration) magnification, and the images transformed into 256 gray scale images. The fields were chosen to locate individual neuron with discernible neurites from nearby neurons. Generally, an average of 2–4 neurons could be seen in a micrograph taken at 20× magnification. The total neurite length was then measured from the somata using a PC-based image analyzer software Image Pro 3.0 Plus (Media Cybernetics, Silver Spring, MD, USA). The signal intensity of bands stained on immunoblot was quantified with Gel pro 3.1 (Media Cybernetics). Student’s t-test was used for evaluating statistical differences between
the means of different groups, with p value of less than 0.05 considered significant.

**Results**

**Daidzein enhances neuritogenesis in cultured DRG neurons**

We first studied the effect of daidzein on neurite outgrowth in primary rat DRG neuronal cultures. DRG neurons were classified into large (diameter $\geq 40 \mu m$) and small (diameter $\leq 40 \mu m$) according to the criteria described by Gavazzi [33]. Based on cell diameter, small-sized unmyelinated neurons that are responsible for pain sensation, and large-sized myelinated that are for proprioception. DIV 3 neurons were incubated with different concentrations of daidzein, DMSO (negative control), or NGF (positive control) for 24 h. Compared to DMSO-treated control cultures, daidzen treatment significantly enhanced neurite extension and branching of large DRG neurons (Figures 1, 2A-B). Quantitative analysis of total neurite length and tip numbers per neuron indicated that the minimal effective concentration of daidzein in promoting neurite outgrowth was 30 $\mu M$ (length, $4180 \pm 246 \mu m$; tip number, 15.0 $\pm$ 1.4; DMSO, 2323 $\pm$ 128 $\mu m$; tip number, 7.8 $\pm$ 0.2; p $\leq 0.01$, n = 10), which had an effect similar to that of 100 ng/ml NGF (Figure 2) and was thus used for further study of the compound. The effect of daidzein-induced neuritogenesis was observed similarly in both small-sized and large-sized DRG neurons (Figures 1G-H, 2C). Furthermore, to examine whether daidzen was toxic to DRG neurons, double staining of cells with DAPI and

![Figure 5](https://www.jbiomedsci.com/content/19/1/80)
PI was performed to identify apoptotic and necrotic cells, respectively [34]. No apparent cell death was observed at 30 μM by MTT assay (Figure 3A).

**Src and ERK inhibitors block neuritogenic effect of daidzein**

To investigate the signaling mechanism underlying the neuritogenic effect of daidzein, various inhibitors were applied to DRG neuronal cultures before and during daidzein treatment. Daidzein is structurally similar to estrogen and can activate ER in other cell culture systems [15,22]. Thus, we first examined this possibility. Treatment alone with inhibitors for ER-α/β, GPR-30, Src, or MEK had no effect on total neurite length and tip numbers (Figure 3B, C). Neither the ER-α/β inhibitor ICI 182780 (length, 3778 ± 101 μm, tip number, 10.9 ± 0.9 μm; daidzein, length, 4145.8 ± 255.5 μm, tip number, 12.9 ± 0.8 μm; DMSO, length, 2363.2 ± 128.5 μm, tip number, 9.3 ± 0.9, n = 10) nor the GPR-30 antagonist tamoxifen also did not inhibit daidzein-induced neuritogenesis (length, 2256.6 ± 580 μm; DMSO, 2629.4 ± 436 μm; daidzein, 3632.7 ± 560 μm). On the other hand, both the Src inhibitor PP2 (length, 1971 ± 101 μm, tip number, 6.7 ± 0.4, n = 10, p ≤ 0.01 daidzein, length 4145.8 ± 255.5 μm, tip number 12.9 ± 0.8 control, length 2363.2 ± 128.5 μm, tip number, 9.3 ± 0.9, and the MEK inhibitor U0126 (length, 2031 ± 126 tip number, 10.0 ± 0.7, n = 10, p ≤ 0.01) significantly reduced daidzein-induced neuritogenesis (Figure 4). Treatment with PP2 or U0126 alone had no effect on cell survival, total neurite length, or tip numbers (Figure 3B, C).

**Blocking of the neuritogenic effect of daidzein by PKCδ inhibitor**

To examine whether PKC involved in the daidzein-induced neuritogenesis, we treated DRG neurons with a pan-PKC activator PMA in the presence of the src kinase inhibitor PP2. The result showed that activation of PKC by PMA could reverse the PP2-induced decrease in total neurite length (PMA + PP2 + daidzein, length, 1999 ± 85 μm; DMSO, 1768 ± 77 μm; PP2 + daidzein, 1401 ± 72 μm, p ≤ 0.05, n = 5).

To further investigate the signaling mechanism involved in the neuritogenic effect of daidzein, various PKC inhibitors were applied to DRG neuronal cultures.
PKCα inhibitor Gö6976 (length and tip number, 3853 ± 673 µm and 17.4 ± 0.6; daidzein, 3382 ± 340 µm and 18.0 ± 1.0, DMSO, 1982 ± 512 µm and 10.8 ± 0.7, n = 10) nor PKCc inhibitor eV1-2 (length and tip number, 3424 ± 482 µm and 16.1 ± 1.0, n = 10) had any effect on the daidzein-induced neuritogenesis. Only the PKCδ inhibitor rottlerin (length, 1705 ± 247 µm, n = 10, p ≤ 0.01) significantly block daidzein’s neuritogenic effect (length, daidzein 3770 ± 252 µm; DMSO; 2291 ± 193 µm, n = 10) (Figure 5). Treatment with Gö6976, eV1-2 (data not shown) or rottlerin (Figure 5) did not affect total neurite length and tip numbers.

**Daidzein treatment activates Src, PKCδ, and ERK**

We next examined whether daidzein could activate the Src, PKCδ, or the ERK signaling pathway. By using Western

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**Figure 7 Activation of ERK by daidzein treatment, and suppression of the ERK activation by both Src and PKCδ inhibitors.** Upper panel showed a representative blot from one experiment. Lower panel showed optical densities of the densitometric scans of the pERK bands. (A) DRG neuronal cultures were treated with DMSO, 30 µM daidzein (Dz) for 30 min, 10 µM the Src inhibitor PP2 for 30 min followed by 30 µM daidzein (Dz + PP2) for 30 min, or 10 µM PP2 for 60 min (PP2). The cell homogenate was analyzed for phosphorylated pERK and total ERK. *, p < 0.05 vs DMSO; #, p < 0.05 vs daidzein. n = 4. (B) DRG neuronal cultures were treated with DMSO, 30 µM daidzein (Dz) for 30 min, 2 µM the PKCδ inhibitor rottlerin for 30 min followed by 30 µM daidzein for 30 min (Dz + rott), or 2 µM rottlerin for 60 min (rott). The cell homogenate was analyzed for phosphorylated pERK and total ERK. *, p < 0.05 vs DMSO group; #, p < 0.05 vs daidzein group. n = 4.

**Figure 8 Effect of different concentrations of daidzein on activation of Src kinase and ERK.** DRG neurons were treated with 0, 30, 50 or 100 µM daidzein for 30 min. Total cell lysates were analyzed for pSrc (A) and pERK (B). GAPDH is loading control. *, p < 0.05; **, p < 0.01 vs 0-min group. n = 3.
blotting with antibodies directed against phosphotyrosine-416 of active Src (pSrc416), we found that the pSrc416 level was significantly increased by daidzein treatment (Figure 6A). Interestingly, pPKCδ levels were also up-regulated in the presence of daidzein treatment, and the Src kinase inhibitor PP2 significantly inhibited this activation of pPKCδ by daidzein (Figure 6B). Moreover, ERK1 and ERK2 were activated by daidzein, and this effect was suppressed by both the Src kinase inhibitor PP2 and the PKCδ inhibitor rottlerin (Figure 7). Interestingly, a dose-dependent study of daidzein showed that a maximal activation of Src kinase and ERK was found at 30 μM daidzein (Figure 8). This data correlates well with the previous observation that the optimal concentration of daidzein to achieve effective neuritogenic activity is 30 μM.

Discussion

Previously, in cultured osteoblastic cells, daidzein was shown to bind to cell membrane ER-β to activate the phospholipase C β2 (PLC-β2)/PKC and PI3K/cSrc pathways, leading to the expression of several groups of genes for cell differentiation, proliferation, and migration [15]. Similarly, in macrophage, daidzein inhibited the activation of STAT-1 and NF-kB proteins, thereby decreased the expression of iNOS and the production of NO [35]. Moreover, in hippocampal neurons, daidzein was demonstrated to promote neurite outgrowth via ER-β, which in turn may increase the phosphorylation of PKCa and growth associated protein 43 (GAP-43) [22]. In contrast to these ER-dependent signaling systems, here our results revealed that the neuritogenic effects of daidzein in DRG neurons did not mediate through ER α/β or GPR-30 (the third kind of ER receptor), as pretreatment of DRG neurons with specific estrogen receptor antagonists, ICI 182780, tamoxifen, or G-15, did not block the daidzein-induced neuritogenesis.

Most importantly, our studies showed that daidzein treatment increased the phosphorylation of Src kinase and ERK1/ERK2 in cultured DRG neurons. Src kinase, which is activated by various molecules like NGF, laminin, artemin, and anti-Thy-1 antibody, has been shown to be an important signaling pathway involved in the process of DRG neurite outgrowth [27,36,37]. Downstream signaling of Src includes MEK/ERK and PI3K/Akt pathways, which can be activated by NGF to induce neurite extension and branching of DRG neurons [38]. While inhibition of Src kinase by PP2 and suppression of ERK1/2 by U0126 abolished the neuritogenic effect of daidzein, inhibition of Akt by LY294002 had no effect (unpublished observation). Thus, the current study did not support the role of PI3K/Akt pathway in daidzein-induced neuritogenesis. The increased phosphorylation of ERK and daidzein-induced neuritogenesis was blocked by the Src kinase inhibitor PP2, indicating that promotion of neurite outgrowth by daidzein required Src kinase and ERK. As phosphorylated ERK could activate CREB, Cdk5, GAP-43 and other neuritogenesis-related genes [39-41], it is possible that daidzein-induced neurite outgrowth is mediated by the Src-ERK pathway. Interestingly, PKCδ has been shown to be phosphorylated and activated by Src kinases in salivary and PC12 cells [42]. Consistent with this notion, we also found that daidzein increased the phosphorylation of PKCδ, and that inhibition of PKCδ by a selective PKCδ inhibitor rottlerin resulted in the suppression of neurite outgrowth, suggesting that PKCδ may also play a role in the signaling cascade induced by daidzein. Different PKC isoforms, including PKCδ, have been found to be activated by neurotrophic agents. For example, in PC12 cells the activation of ERK by neurotogenic agents, fibroblast growth factor (FGF) and NGF, was dependent on the activation of PKCδ [43]. In diabetic rats, over-expression of PKCδ could ameliorate the retarded neurite outgrowth of DRG neurons [44]. Additional studies are therefore required to elucidate the mechanistic link between daidzein-induced activation of the Src-PKCδ-ERK pathway and the downstream signaling pathways that eventually lead to neuritogenesis.

As mentioned above, we have shown that daidzein-induced neuritogenesis in hippocampal and DRG neurons are mediated by ER-dependent [22] and ER-independent mechanisms, respectively. The precise reasons underlying this striking difference remain unclear. Intrinsic signaling pathways regulating neurite outgrowth could drastically vary among different types of neuronal cells [45,46]. For example, activation and inhibition of the small GTPase Rac1 promotes neurite outgrowths in hippocampal and DRG neurons, respectively [47,48]. Further experiments will be required to determine if the neuritogenic effect of daidzein is also differentially regulated in various regions of the nervous system.

In cultured rat hippocampal neurons, a low concentration of daidzein had neuroprotective action (at 3.9 μM), but it could not promote neuritogenesis or enhance neuronal survival [24]. Using relative affinity binding assay of cellular extracts, the affinity of daidzein for estrogen receptor was estimated to be several hundred times lower than estrogen [19]. Therefore, adequate amount of daidzein is required to achieve biological activities through the estrogen receptor pathways. Studies using daidzein at much higher concentrations (30 to 40 μM) in cultured hippocampal neurons indicate that daidzein can promote neurite extension and protect neurons from glutamate-induced cell death [22,24]. We demonstrated that daidzein at 30 μM increased neurite lengthening and branching for DRG neurons, which was in accordance with the results of previously published reports. In addition, we found 30 μM daidzein had a neuritogenic effect similar to that of NGF, indicating that daidzein had a robust neuritogenic property. Meanwhile, daidzein did not affect DRG neuronal survival at 30 μM. When daidzein was used at a concentration higher than 30 μM, the neuritogenesis was decreased. This finding is in agreement with the observed fact that both
Src and ERK achieve the highest levels of phosphorylation at 30 μM daidzein.

Our finding that daidzein facilitates neurite outgrowth of DRG and hippocampal neurons has important implications for the potential facilitation of neural regeneration [49, 50]. Preliminary study using daidzein for the treatment of optic nerve injury in rats have shown promising results [23]. Neurite outgrowth is a fundamental step in the establishment of neural connections during development and following injury. It would be interesting to see if daidzein could improve sensory and cognitive function in various animal disease models, e.g. injuries of peripheral nerves, brachial plexus, and spinal cord, as well as Alzheimer’s disease. On the other hands, the signaling mechanism of daidzein warrants further investigation. It has been shown that daidzein did not activate Src kinase via ERs, so other upstream regulators of Src could be the potential targets of daidzein. Many downstream effectors of ERK are associated with neurite outgrowth, and daidzein may preferentially activate some of them. Future research will be directed to tackle these questions.

Conclusion

Daidzein enhances neurite outgrowth of cultured rat DRG neurons which is mediated by cooperative action of Src kinase, PKCδ, and MEK.

Abbreviations

DIV: Day in vitro; DMSO: Dimethyl sulfoxide; DRG: Dorsal root ganglion; ER: Estrogen receptor; ERK: Extracellular signal-regulated kinase; GAP-43: Growth associated protein 43; GPR-30: G-protein coupled receptor 30; MEK: Mitogen-activated protein kinase/extracellular signal-regulated kinase; MPP: 1-methyl-4-phenyl pyridium; MTT: 3-[4, 5-dimethylthiazol-2-yl]-2, 5-diphenyltetrazolium bromide; NF-L: Neurofilament light chain; NGF: Nerve growth factor; PBS: Phosphate buffered saline; P38K: Phospho
erosinine 3-kinase; PLC: Phospholipase C; PKC: Protein kinase C; PKCδ: Protein kinase C subtype delta; TBS: Tris-buffered saline.

Competing interests

The authors declare that they have no competing interests.

Authors’ contributions

SHY: designed and carried out experiments, analyzed results and manuscript writing. CCL, JPS: performed DRG cultures, immunostaining. YC: western blotting analysis. CJJ: designed experiments, analyzed results and manuscript revision. All authors read and approved the final manuscript.

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Author details

1 Division of Neurosurgery, Department of Surgery, National Taiwan University Hospital, Taipei, Taiwan. 2 Department of Anatomy and Cell Biology, College of Medicine, National Taiwan University, No. 1, Section 1, Jen-Ai Road, Taipei 10051, Taiwan. 3 Department of Biology and Anatomy, National Defense Medical Center, Taipei, Taiwan. 4 Institute of Anatomy and Cell Biology, School of Medicine, National Yang-Ming University, No. 155, Section 2, Li-Non Street, Taipei 12212, Taiwan.

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References

1. Franke AA, Custer LJ, Wang W, Shi CY: HPLC analysis of isoflavonoids and other phenolic agents from foods and from human fluids. Proc Soc Exp Biol Med 1998, 217:263–273.

2. Wang SW, Chen Y, Joseph T, Hu M: Variable isoflavone content of red clover products affects intestinal disposition of biochanin A, formononetin, genistein, and daidzein. J Altern Complement Med 2008, 14:287–297.

3. Osozki AL, Kennelly EJ: Phytoestrogens: a review of the present state of research. Phytother Res 2003, 17:845–869.

4. Cheksis BJ, Greger JG, Nagpal S, Freedman LP: Signaling by estrogens. J Cell Physiol 2007, 213:610–617.

5. Hwang CS, Kwak HS, Lim HJ, Lee SH, Kang YS, Cho TB, Hur HG, Han KO: Isoflavone metabolites and their in vitro dual functions: they can act as an estrogenic agonist or antagonist depending on the estrogen concentration. J Steroid Biochem Mol Biol 2006, 101:246–253.

6. Gikas PD, Mokbel K: Phytoestrogens and the risk of breast cancer: a review of the literature. Int J Ferti Womens Med 2005, 50:250–258.

7. Hedelin M, Loj M, Olsson M, Adlercreutz H, Sandin S, Widerlund E: Dietary phytoestrogens are not associated with risk of overall breast cancer but diets rich in coumestrol are inversely associated with risk of estrogen receptor and progesterone receptor negative breast tumors in Swedish women. J Nutr 2008, 138:938–945.

8. Verheus M, van Gils CH, Keinan-Boker L, Grace PB, Bingham SA, Peeters PH: Plasma phytoestrogens and subsequent breast cancer risk. J Clin Oncol 2007, 25:6648–6655.

9. Marino M, Acconcia F, Ascenzi P: Estrogen receptor signalling: bases for drug actions. Curr Drug Targets Immune Endocr Metabol Disord 2005, 5:305–314.

10. Segars JH, Driggers PH: Estrogen action and cytoplastic signaling cascades. Part 1: membrane-associated signaling complexes. Trends Endocrinol Metab 2002, 13:349–354.

11. Falkenstein E, Tillmann HC, Christ M, Feuring M, Wehling M: Multiple actions of steroid hormones—a focus on rapid, nongenomic effects. Pharmacol Rev 2000, 52:513–556.

12. Zhao C, Dahlman-Wright K, Gustafsson JA: Estrogen signalling via estrogen receptor (beta). J Biol Chem 2010, 285:39575–39579.

13. Revankar CM, Cimino DF, Sklar LA, Atenburn JB, Prossnitz ER: A transmembrane intracellular estrogen receptor mediates rapid cell signaling. Science 2005, 307:1625–1630.

14. Chamblis KL, Wu Q, Oltman S, Konanah S, Umetani M, Korach KS, Thomas GD, Mineo C, Yuhanna IS, Kim SH, Madak-Erdogan Z, Maggi A, Dineen SP, Roland CL, Hui DY, Breken RA, Katzenellenbogen JA, Katzenellenbogen BS, Shaul PW: Non-nuclear estrogen receptor alpha signaling promotes cardiovascular protection but not uterine or breast cancer growth in mice. J Clin Invest 2010, 120:2319–2330.

15. de Wilde A, Heberden C, Chaumaz G, Bordat C, Lieberherr M: Signaling networks from Gbeta1 subunit to transcription factors and actin remodeling via a membrane-located Elf-beta receptor-protein in the rapid action of daidzein in osteoblasts. J Cell Physiol 2006, 209:786–801.

16. Constantinnou AI, Kryger AE, Mehta RR: Genistein induces maturation of cultured human breast cancer cells and prevents tumor growth in nude mice. Am J Clin Nutr 1998, 68:1426S–1430S.

17. Sloan DM, Wu J, Shen ZZ, Barsky SH: Genistein exerts multiple suppressive effects on human breast carcinoma cells. Cancer Res 1998, 58:4851–4857.

18. Vantyghem SA, Wilson SM, Postenka CO, Al-Katib W, Tuck AB, Chambers AF: Dietary genistein reduces metastasis in a posturgical orthotopic breast cancer model. Cancer Res 2005, 65:3396–3403.

19. Kuiper GG, Lemmen JG, Carlson B, Cotton JC, Safe SH, van der Saag PT, van der Burg B, Gustafsson JA: Interaction of estrogenic chemicals and phytoestrogens with estrogen receptor beta. Endocrinology 1998, 139:4252–4263.

20. Guo JM, Xiao BX, Liu Q, Ma HH: Effects of daidzein on estrogen-receptor-positive and negative pancreatic cancer cells in vitro. World J Gastroenterol 2004, 10:860–863.

21. Miller DR, Lee GM, Maness PF: Increased neurite outgrowth induced by inhibition of protein tyrosine kinase activity in PC12 pheochromocytoma cells. J Neurochem 1993, 60:2134–2144.

22. Wang P, Jeng CJ, Chien CL, Wang SM: Signaling mechanisms of daidzein-induced axonal outgrowth in hippocampal neurons. Biochim Biophys Acta 2008, 1826:393–400.
31. Di Giovanni S, Campana A, Lange PS, Lee HH, Baneejee K, Byrson JB, Mahishi L, Alam S, Giger RJ, Barnes S, Morris SM Jr, Willis DE, Twiss JL, Filbin MT, Ratan RR: A large-scale chemical screen for regulators of the arginase 1 promoter identifies the soy isoflavone daidzein as a clinically approved small molecule that can promote neuronal protection or regeneration via a CAMP-independent pathway. J Neurosci 2010, 30:759–768.

32. Zhao L, Chen Q, Diaz-Bronin R: Neuroprotective and neurotrophic efficacy of phytoestrogens in cultured hippocampal neurons. Exp Biol Med (Maywood) 2002, 227:509–519.

33. Chen CH, Chen YJ, Yang SH, Tung PY, Wang SM: Role of PKA in the anti-Thy-1 antibody-induced neurite outgrowth of dorsal root ganglionic neurons. J Cell Biochem 2007, 101:566–575.

34. Desbois C, Birge RB, Minnouri-Rongy M, Weinstein DE, Palermo JS, Neveu MK: Fas engagement induces neurite growth through ERK activation and p38 upregulation. Nat Cell Biol 2003, 5:118–125.

35. Yang SH, Chen YJ, Tung PY, Lai WL, Chen Y, Jing CJ, Wang SM: Anti-Thy-1 antibody-induced neurite outgrowth in cultured dorsal root ganglionic neurons is mediated by the c-Src-MEK signaling pathway. J Cell Biochem 2008, 103:467–77.

36. Leclere PG, Norman E, Grougi F, Coffin R, Mayer U, Pizey J, Tonge O: Impaired axonal regeneration by isoelectin B4-binding dorsal root ganglion neurons in vitro. J Neurosci 2007, 27:1190–1199.

37. Di Giovanni S, De Biasi A, Yakovlev A, Willis DE, Filbin MT, Ratan RR: A large-scale chemical screen for regulators of the arginase 1 promoter identifies the soy isoflavone daidzein as a clinically approved small molecule that can promote neuronal protection or regeneration via a CAMP-independent pathway. J Neurosci 2010, 30:759–768.

38. Harada T, Morooka T, Ogawa S, Nishida E: Impaired axonal regeneration by isolectin B4-binding dorsal root ganglion neurons. J Cell Biochem 2007, 101:566–575.

39. Schambon JC, Puschel AW: The sequential activity of the GTPases Rap1B and Cdc42 determines neuronal polarity. Nat Neurosci 2004, 7:923–929.

40. Davies AM: Neurotrophins: neurotrophic modulation of neurite growth. Curr Biol 2000, 10:R198–R200.

41. Gil OD, Zanazzi G, Struyk AF, Salzer JL: Neurotropin mediates bifunctional effects on neurite outgrowth via homophilic and heterophilic interactions. J Neurosci 1998, 18:9312–9325.

42. Ko M, Zou K, Minagawa H, Yu W, Gong JS, Yanagisawa K, Michikawa M: Cholesterol-mediated neurite outgrowth is differently regulated between cortical and hippocampal neurons. J Biol Chem 2005, 280:42759–42765.

43. Fournier AE, Takizawa BT, Stittmatter SM: Rho kinase inhibition enhances axonal regeneration in the injured CNS. J Neurosci 2003, 23:1416–1423.

44. Corbit KC, Foster DA, Rosner MR: Protein kinase Cdelta mediates neurogenic but not mitogenic activation of mitogen-activated protein kinase in neuronal cells. Mol Cell Biol 1999, 19:4209–4218.