Effect of activation of canonical Wnt signaling by the Wnt-3a protein on the susceptibility of PC12 cells to oxidative and apoptotic insults

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Abstract

Wnt proteins are involved in tissue development and their signaling pathways play an important role during embryogenesis. Wnt signaling can promote cell survival, which is beneficial for neurons, but could also lead to tumor development in different tissues. The present study investigated the effects of a Wnt protein on the susceptibility of a neural tumor cell line (PC12 cells) to the cytotoxic compounds ferrous sulfate (10 mM), staurosporine (100 and 500 nM), 3-nitropropionic acid (5 mM), and amyloid β-peptide (Aβ25-35; 50 µM). Cells (1 x 10^6 cells/mL) were treated with the Wnt-3a recombinant peptide (200 ng/mL) for 24 h before exposure to toxic insults. The Wnt-3a protein partially protected PC12 cells, with a 6-15% increase in cell viability in the presence of toxic agents, similar to the effect measured using the MTT and lactate dehydrogenase cell viability assays. The Wnt-3a protein increased protein expression of β-catenin by 52% compared to control. These findings suggest that Wnt signaling can protect neural cells against apoptosis induced by toxic agents, which are relevant to the pathogenesis of Alzheimer’s and Huntington’s diseases.

Key words: Wnt-3a; PC12 cells; Beta-catenin; NF-κB

Introduction

In 1982, Nusse and Varmus (1) identified a proto-oncogene called Wnt-1 as a signaling molecule involved in breast cancer development. It is now well established that Wnt proteins, which were first studied in Drosophila and Xenopus, are important mediators of intercellular communication and the signaling mediated by these proteins is vital to normal organogenesis in several systems including the central nervous system (CNS) (for reviews, see Refs. 1-3). It has been demonstrated that Wnts regulate several cellular processes including cell proliferation, cell polarity and movement, apoptosis, neuronal circuit development, and neuronal plasticity in adult animals (4).

Wnt proteins can be divided into two classes based on the signaling pathways they activate: canonical Wnts (Wnt-1, Wnt-3a and Wnt-8), which act through β-catenin, and non-canonical Wnts (Wnt-4, Wnt-5a and Wnt-11), which are not related to β-catenin, and instead signal via planar cell polarity (PCP) and calcium (5,6). Wnt proteins bind to receptors called Frizzled (Fz), which are transmembrane proteins (7). So far 19 Wnt genes and 10 Fz genes have been identified in mice. The Wnt-Fz activates a cytosolic protein called Dishevelled (Dvl) (8). Once activated, Dvl can engage three pathways: canonical or Wnt/β-catenin, PCP and Wnt/calcium. Evidence suggests crosstalk among these pathways (4,9).

Activated Dvl inhibits the complex formation by AXIN, adenomatous polyposis coli protein (APC), glycogen synthase kinase-3β (GSK-3β), and β-catenin. In cells where Wnt is not activated, GSK-3β phosphorylates β-catenin, activating its degradation in the proteasome. On the other hand, Wnt activation causes GSK-3β inhibition that leads to an increase of cytosolic β-catenin that migrates to the nucleus to form a complex with the T-cell factor (TCF), a transcription factor that regulates many different genes (for a review, see Ref. 10).
Recent findings suggest that Wnt signaling may play a neuroprotective role, for example, protecting cultured hippocampal neurons against amyloid β-peptide (Aβ) (11). In addition, it has been proposed that a decrease in Wnt activity may play a role in CNS diseases such as schizophrenia (12), retinal neurodegeneration (13), autism (14), and Alzheimer’s disease (15,16).

Other evidence has shown the involvement of Wnt signaling in tumor development in humans, especially in colon and liver cancers (17,18). Therefore, a better understanding of the mechanisms by which Wnt signaling affects the susceptibility of cells to toxic insults could be useful in the development of novel interventions for the treatment of neurodegenerative disorders. In the present study, we evaluated the effects of Wnt signaling on the viability of neural cells exposed to an array of cytotoxic compounds including ferrous sulfate, staurosporine, 3-nitropropionic acid, and Aβ25-35.

Material and Methods

Reagents

Dulbecco’s minimum essential medium (DMEM), fetal calf serum, penicillin, and streptomycin were purchased from Cultilab (Brazil). Horse serum was purchased from Invitrogen (USA). Aβ25-35 was obtained from Bachem (USA). Wnt-3a was purchased from R&D Systems (USA) and staurosporine from Tocris (USA). Reagents for SDS-PAGE and immunoblotting were purchased from Bio-Rad Laboratories (USA), and ethidium bromide from Promega (USA). Antibody to β-actin was obtained from Santa Cruz Biotechnology (USA) and antibody to β-catenin from Cell Signaling (USA). Ferrous sulfate (FeSO4), 3-nitropropionic acid (3-NP) acid, and Aβ25-35 were obtained from Sigma-Aldrich (USA). A lactate dehydrogenase (LDH) kit was obtained from Roche Diagnostics (Germany).

PC12 cells

The PC12 cell line (cells derived from rat pheochromocytoma) was obtained from the American Type Culture Collection, USA. Cells were cultured (1 x 10⁶ cells/mL) in DMEM plus 10% horse serum and 5% fetal calf serum and kept at 37°C in a humidified 5% CO₂ atmosphere until 90% confluence (19).

Treatment of PC12 cell with recombinant Wnt-3a and toxic agents

Aβ25-35 was diluted in distilled water to 1 mM and kept at -20°C until use. Before use, the peptide was diluted in sterile PBS to 200 µM and incubated at 37°C for 5 days to allow peptide aggregation (20). FeSO₄ was dissolved in water, staurosporine in DMSO and 3-NP acid in ethanol; all these agents were prepared as 1 M stock solutions. DMSO or ethanol vehicles (0.1% final concentration) were added to control cultures. Aβ25-35, 50 µM, 10 mM FeSO₄, 100 and 500 ng/mL staurosporine, and 5 mM 3-NP acid were added to the culture medium 24 h after the addition of mouse recombinant Wnt-3a peptide dissolved in saline at three concentrations (50, 100, and 200 ng/mL). Cell viability and protein expression by immunoblot were measured 24 h later. During the treatment period cells were kept in DMEM with 1% horse serum and 0.5% fetal calf serum (19).

MTT reduction and LDH release assay

Cell viability was estimated by the MTT reduction (21), which changes the color of MTT only in the presence of live cells since the ring of the tetrazolium salt is cleaved by several dehydrogenase enzymes present in active mitochondria. After incubation of the cells with Wnt-3a, staurosporine, 3-NP acid and FeSO₄ or Aβ25-35 for 24 h, the culture medium was changed, and MTT (final concentration 0.5 mg/mL) was added to the cells. Absorbance (570 nm) was measured after 30 min in a multiwell plate reader (Bio-Tek Instruments, USA). Results are reported as percent of control values.

Cytotoxicity was also estimated by measuring the release of LDH into the medium (22). After treatment of the cells, the LDH released in the culture medium was measured using a colorimetric LDH cytotoxicity assay kit (Roche Applied Sciences, USA) according to manufacturer instructions. Briefly, LDH oxidizes lactate to pyruvate with conversion of NAD⁺ to NADH, which reacts with iodonitrotetrazolium to form formazan. The results are reported as percent of control absorbance at 490 nm measured after 15-20 min in a multiwell plate reader (Bio-Tek Instruments).

Electrophoretic mobility shift assay

Nuclear extracts were prepared as described previously (23). Briefly, cells were scrapped in cold PBS (supplemented with 2 µg/mL leupeptin, 2 µg/mL antipain, and 0.5 mM PMSF) and centrifuged at 4°C for 2 min at 13,000 g. Pellets were resuspended in lysis buffer (10 mM HEPES, pH 7.9, 1.5 mM MgCl₂, 10 mM KCl, 0.1 mM EDTA, 0.5 mM PMSF, 2 µg/mL leupeptin, 2 µg/mL antipain, 3 mM sodium orthovanadate, 30 mM sodium fluoride, and 20 mM sodium pyrophosphate) and incubated on ice for 15 min. After the addition of NP-40 (to a final concentration of 0.5%), samples were vigorously mixed and centrifuged for 30 s at 13,000 g. Supernatants were kept at -80°C for immunoblot analysis. Nuclei were resuspended in extraction buffer (20 mM HEPES, pH 7.9, 25% glycerol, 1.5 mM MgCl₂, 300 mM NaCl, 0.25 mM EDTA, 0.5 mM PMSF, 2 µg/mL leupeptin, and 2 µg/mL antipain), incubated for 20 min on ice, and centrifuged for 20 min at 13,000 g at 4°C. The remaining supernatants containing nuclear proteins were aliquoted and stored at -80°C. Protein concentration was determined using the Bio-Rad protein reagent (Bradford).

Electrophoretic mobility shift assay (EMSA) for NF-κB was performed using a gel shift assay kit from Promega.
NF-κB double-stranded consensus oligonucleotide (5'-AG TTGAGGGGACCTTTCCAGGC-3') was end-labeled with γ-[32P]-ATP using T4 polynucleotide kinase. Unincorporated nucleotides were removed by passing the reaction mixture through a Sephadex G-25 spin column (Amersham-Pharma,

cia, Uppsala, Sweden). A purified [32P]-labeled probe (25,000 cpm) was incubated with 5 µg nuclear extracts in a 20-µl binding reaction mixture containing 50 mM NaCl, 0.2 mM EDTA, 0.5 mM DTT, 4% glycerol, 10 mM Tris-HCl, pH 7.5, and 0.05 µg poly(dI-dC) for 30 min at room temperature. DNA-protein complexes were separated by electrophoresis on a 6% nondenaturing acrylamide:bis-acrylamide (37.5:1) gel in 0.5% Tris-borate/EDTA (TBE) for 2 h at 150 V. Gels were vacuum-dried and analyzed by autoradiography.

For competition experiments, unlabeled double-stranded consensus NF-κB oligonucleotide was included in 2.5- to 10-fold molar excess over the amount of the [32P]-NF-κB probe to detect specific and nonspecific DNA-protein interactions. Unlabeled oligonucleotides were added to the reaction mixture 20 min before the radioactive probe. The composition of the complexes was determined by supershift assays; antibodies (1:10 dilution) against p50 and p65 NF-κB subunits were added before the incubation of nuclear extracts with the labeled oligonucleotide. Autoradiographs were quantified with the Chemilumigtaion detection system (Alpha-Innotech Corporation, USA), and several exposure times were analyzed to ensure linearity of the band intensities.

Immunoblot analyses

Electrophoresis was performed using a Bio-Rad mini-Protein III apparatus. Briefly, the proteins present in the cytosolic (10 µg) fraction were size-separated by SDS-PAGE (10% acrylamide). The proteins were blotted onto a nitrocellulose membrane (Bio-Rad) and incubated with the β-catenin antibody. The Ponceau-staining method was used to control for differences in protein loading (24). Proteins recognized by antibodies were developed by the ECL technique, following manufacturer instructions (Amersham, USA). A Chemilumigtaion detection system (Alpha-Innotech Corporation) was used to standardize and quantify the immunoblots. Several exposure times were analyzed to ensure the linearity of the band intensities. β-actin antibody was used as an internal control. Data are reported in relation to the intensity of the β-actin band.

Measurement of Frizzled mRNA levels by RT-PCR

Total RNA was isolated using the Trizol reagent (Invitrogen) according to manufacturer instructions. Semiquanitative reverse transcription-PCR (RT-PCR) amplification was performed using the ThermoScript RT kit (Invitrogen) according to manufacturer instructions. The primer sequences were: Fz 1 (170 bp), 5'-GCC AAG CCT AGT CAC TCG AG-3' (sense) and 5'-TAG TAT CGG AGG GGC TAC ATT CT-3' (antisense); Fz 2 (287 bp), 5'-CCT GGA GGT GCA TCA ATT CT-3' (sense) and 5'-GGT GAG TAG CGC AGG AGT TC-3' (antisense); Fz 3 (167 bp), 5'-GAA GCA AAG CAG GGA GTG TC-3' (sense) and 5'-GAG TGA TCT GTC AGC CGT GA-3' (antisense); Fz 4 (110 bp), 5'-AGC TGA CTG GCC TGT GCT AT-3' (sense) and 5'-CAA GCC TGC AGCAAT GAA TA-3' (antisense); Fz 5 (275 bp), 5'-TGT GCA CAC TCG TCT TCC TC-3' (sense) and 5'-CCA AGA CAA AGG CTC GTA GC-3' (antisense); Fz 6 (164 bp), 5'-TGG GCT GTC TCT CCT CT-3' (sense) and 5'-GCC AAG AAG CGT CAC TAA GG-3' (antisense); Fz 7 (268 bp), 5'-TCT GGG TAG GTG TGT GGT CA-3' (sense) and 5'-AAG CAC CAT GAA GAG GAT GG-3' (antisense); Fz 8 (154 bp), 5'-TCC GTT CAG TCATCAACGAG-3'sence) and 5'-CGG TTG TGC TGC TCA TAG AA-3'(antisense); Fz 9 (105 bp) 5'-AAG TCC ATG TTG AGG CGT TC-3' (antisense) and 5'-AAG GCT GGA AA GGT CAT GG-3' (antisense), and Fz 10 (190 bp) 5'-CAA GAC ACC TGA CTG CCT GA-3' (sense) and 5'-TCC CCA AGG TGA GGT TTT TG-3' (antisense). The PCR conditions were 5 min at 94°C, 35 cycles of 94°C for 45 s, 55°C for 45 s and 72°C for 90 s and a final extension at 72°C for 10 min. Gel electrophoresis of the PCR product was performed using an ethidium bromide-containing agarose gel (2%), and the resulting bands were visualized under UV light. The photographs were obtained with a Vilber Lourmat system (Alpha-Innotech Corporation).

Statistical analysis

Data are reported as means ± SEM for the indicated number of experiments. Statistical comparisons were performed by the unpaired t-test, or one-way ANOVA followed by the Newman-Keuls test (P < 0.05 was considered to be as statistically significant). All analyses were performed using a Prism4 software package (GraphPad Software, USA).

Results

Regular RT-PCR was used to evaluate which Fz receptors are expressed in non-differentiated PC12 cells. This cell line expressed mRNA of Fz receptors 1 to 7, but not 8, 9, or 10 (Figure 1). Mouse recombinant Wnt-3a protein increased

![Figure 1](www.bjournal.com.brBraz J Med Biol Res 45(1) 2012 www.bjournal.com.br)
the protein levels of β-catenin in PC12 cells, indicating the presence of the Wnt canonical signaling pathway (Figure 2). The results of densitometric analyses of the immunoblot are shown. The results of the MTT reduction (control = 0.56 ± 0.005; Wnt-3a = 0.61 ± 0.02) and LDH release (control = 0.56 ± 0.01; Wnt-3a = 0.56 ± 0.02) assays showed that the concentration of 100 ng/mL Wnt-3a protein was not toxic to the PC12 cells (P < 0.05, unpaired Student t-test).

In addition, at the same concentration, the Wnt-3a protein prevented about 10% of the cell damage induced by staurosporine at two toxic concentrations (100 and 500 nM; Figure 3A,B). To evaluate the concentration-response effect of Wnt-3a against the toxic effect of the higher (500 nM) concentration of staurosporine, three concentrations of this protein were tested (50, 100, and 200 ng/mL). The results showed that the low Wnt-3a concentration prevented about 6.5% of the cell damage, while the concentrations of 100 and 200 ng/mL were able to induce a significant increase (10-15%) in the protection of PC12 cells against this toxic compound (Figure 3C,D). Based on these results that showed no difference in the effect induced by 100 and 200 ng/mL Wnt-3a peptide, the concentration of 100 ng/mL was used in all subsequent experiments (Figure 3C,D). A similar result was obtained when cells were incubated with 10 mM FeSO4, since the toxic effect of this compound on PC12 cells was prevented by 100 ng/mL Wnt-3a, yielding about 10% protection of cells from iron toxicity (Figure 4).

When PC12 cells were incubated with 3-NP acid, 75% of the cells died. Wnt-3a (100 ng/mL) prevented about 7% of the cell toxicity caused by 3-NP acid (Figure 5). A higher concentration of Wnt-3a (200 ng/mL) protected about 9% of the cells against 3-NP acid toxicity, but the effect was not significantly different from the dose of 100 ng/mL (Figure 5).

Aβ25-35 was toxic to the PC12 cells at a concentration of 50 µM, and Wnt-3a (100 ng/mL) was able to prevent 10% of the cell death (Figure 6).

EMSAs were performed to study the effects of the Wnt-3a protein in the absence or presence of two toxic agents (staurosporine and FeSO4) on NF-kB-binding activity in PC12 cells. Staurosporine (500 nM) and 10 mM FeSO4 reduced the basal levels of NF-kB DNA-binding activity (Figure 7A,B). The pre-incubation of PC12 cells with Wnt-3a partially reversed this effect since this protein activated NF-kB DNA-binding activity when compared to the activity obtained in cells exposed to staurosporine or FeSO4 alone.

Cell nuclear extracts presented a similar pattern of the DNA-NF-kB protein complex. The complex was displaced by an excess of unlabeled NF-kB, demonstrating the specificity of NF-kB-DNA interaction (Figure 7C). Supershift analysis indicated that the antibody against the p65 subunit was able to shift the DNA-protein interaction observed in the complex. In addition, the antibody against the p50 subunit was also able to shift the NF-kB/DNA complex (Figure 7D). The presence of antibodies against the p52 and c-rel subunits did not affect the DNA-protein complex (data not shown). Therefore, taken together, these results suggest that p50/p65 heterodimers were included in NF-kB protein complexes.

Finally, the effect of Wnt-3a in the presence or absence of staurosporine and FeSO4 on the expression of β-catenin protein was evaluated in PC12 cells by immunoblot. Exposure of the cells to staurosporine and FeSO4 caused a decrease in β-catenin protein levels compared to control cultures. In addition, data revealed that when the cells were pre-incubated with Wnt-3a and then exposed to staurosporine or FeSO4, β-catenin protein expression increased compared to the cells treated with toxic agents alone (Figure 8).

Discussion

Fz has been known to function as a Wnt receptor. Although 10 isoforms of Fz receptors have been identified in mammals, their binding specificities with Wnt and functions are poorly understood. Here, we demonstrated that
PC12 cells express most of the Fz receptors, except Fz 8, 9, and 10, which is in accordance with a previous study in which Wnt signaling was observed in PC12 cells (25). Fz receptors seem to have a functional role in PC12 cells since their activation by Wnt-3a recombinant protein induced an increase in β-catenin expression.

When discovered, the Wnt protein was associated with tumors in vertebrates. Wnt signaling pathways are now well known for their involvement in tumor development in humans, especially in colon and liver cancers.
Figure 4. Wnt-3a peptide reduced the loss of cell viability induced by 10 mM FeSO₄ (Fe). A, B, PC12 cells were incubated with 10 mM FeSO₄ 24 h after the addition of mouse recombinant Wnt-3a peptide (100 ng/mL). Twenty-four hours later, cell viability was measured by A, MTT reduction (570 nm) and B, by lactate dehydrogenase (LDH) release (490 nm). Data are reported as means ± SEM for 6 independent experiments. *P < 0.001 vs control (vehicle: water); **P < 0.05 vs Fe (one-way ANOVA followed by the Newman-Keuls test).

Figure 5. Effect of two concentrations of Wnt-3a peptide against 3-nitropropionic acid (3-NP) toxicity. PC12 cells were incubated with Wnt-3a (100 or 200 ng/mL) for 24 h followed by incubation with 5 mM 3-NP for 24 h. Twenty-four hours later, cell viability was measured by A, MTT reduction (570 nm) and by B, lactate dehydrogenase (LDH) release (490 nm). Data are reported as means ± SEM for 6 independent experiments. *P < 0.001 vs control (vehicle: 0.08% ethanol), **P < 0.05 vs 3-NP (one-way ANOVA followed by the Newman-Keuls test).

Figure 6. Effect of recombinant Wnt-3a protein on loss of cell viability induced by Aβ₂₅-₃₅. PC12 cells were incubated with Wnt-3a (100 ng/mL) for 24 h followed by 50 µM Aβ₂₅-₃₅ for 24 h. Twenty-four hours later, cell viability and protein expression were evaluated by A, MTT reduction (570 nm) and by B, lactate dehydrogenase (LDH) release (490 nm) assays. Data are reported as means ± SEM for 6 independent experiments. *P < 0.01 vs control (vehicle: PBS); **P < 0.05 vs Aβ (one-way ANOVA followed by the Newman-Keuls test).
Figure 7. Effect of Wnt-3a on NF-κB suppression induced by staurosporine and FeSO₄ on PC12 cells. Nuclear proteins (2.5 µg) were extracted from PC12 cells pre-incubated (24 h) with mouse recombinant peptide Wnt-3a (100 ng/mL) followed by 24-h incubation with A, 500 nM staurosporine or B, 10 mM FeSO₄. Densitometric analyses (arbitrary units, AU) of the NF-κB band are presented in Panels A and B. Data are reported as means ± SEM for 3 independent experiments *P < 0.001 vs control; **P < 0.05 vs staurosporine and FeSO₄ (one-way ANOVA followed by the Newman-Keuls test). C, Competition studies were performed using nuclear extracts (2.5 µg) from PC12 cells pre-incubated (24 h) with mouse recombinant protein Wnt-3a (100 ng/mL) followed by 24-h incubation with saline in the absence and in the presence of unlabeled specific oligonucleotide (2.5- and 10-fold molar excess), as indicated. D, Supershift assays were performed with nuclear proteins (2.5 µg) extracted from PC12 cells pre-incubated (24 h) with mouse recombinant peptide Wnt-3a (100 ng/mL) followed by 24-h incubation with FeSO₄ (10 mM) in the absence or presence of antibodies against subunits p50 (1:10 dilution), p65 (1:10 dilution), as indicated. Antibodies were added 20 min prior to the addition of the radiolabeled NF-κB consensus oligonucleotide. The position of the specific NF-κB/DNA binding complex p50/p65 (band) is indicated. The localization of the probe is also indicated. Results are representative of 3 independent experiments.
However, Wnt signaling has recently been linked to diseases related mainly to the CNS, including Alzheimer’s disease (26), Parkinson’s disease (27), bipolar disorder (28), and schizophrenia (29).

Chong and Maiese (30) demonstrated that overexpression of Wnt-1 in cultured rat primary hippocampal neurons protected the cells against oxidative stress and the toxicity of Aβ. In another study, Wnt-3a protected a rat primary neuronal hippocampal culture against Aβ peptide toxicity (11).

Our findings confirm a cytoprotective effect of Wnt-3a against Aβ toxicity, and extend previous observations by demonstrating that activation of the canonical Wnt signaling pathway protects neural cells against a broad range of oxidative and apoptotic insults including exposure to FeSO₄, the mitochondrial toxin 3-NP acid, and staurosporine, which triggers mitochondria/caspase-3-mediated apoptosis. These findings shed light on the roles of Wnt signaling in neuroprotection against adverse conditions relevant to the pathogenesis of multiple neurodegenerative disorders.

Previous studies have shown that β-catenin is proteolytically cleaved during staurosporine-induced apoptosis (31). Here we observed a similar effect since exposure of PC12 cells to staurosporine resulted in decreased β-catenin expression in PC12 cells as revealed by immunoblot analysis. However, when the cells were pre-treated with Wnt-3a recombinant peptide the amount of β-catenin was increased compared to the cells treated with staurosporine alone, demonstrating a protective role of Wnt-3a peptide against staurosporine-induced apoptosis. This cytoprotective effect of Wnt-3a was confirmed by MTT reduction and LDH release assays. Staurosporine also decreased NF-κB DNA-binding activity, and the increased activity of this transcription factor in cells pre-incubated with Wnt-3a peptide probably reflects the protective action of this peptide. Indeed, previous studies have shown that activation of NF-κB can protect neurons against death induced by Aβ (32), FeSO₄ (33), and excitotoxicity (34).

An abnormally long polyglutamine or polyalanine tract in different proteins often results in protein aggregation that can be pathological. A well-known example is polyglutamine expansion in the huntingtin protein as the cause of Huntington’s disease (HD) (35). It has been demonstrated that lithium can protect against the polyglutamine toxicity of the HD mutation in cell models (36) and in a Drosophila model of HD (37) by inhibiting GSK-3β. We observed that 3-NP acid decreased β-catenin levels in PC12 cells and that pre-incubation of the cells with Wnt-3a peptide increased the β-catenin level compared to the cells treated with 3-NP acid alone, partially protecting PC12 cells against 3-NP acid toxicity.

Figure 8. Effect of recombinant Wnt-3a protein on β-catenin protein suppression induced by staurosporine and FeSO₄. PC12 cells were pre-incubated with Wnt-3a recombinant protein (100 ng/mL) 24 h before treatment with 500 nM staurosporine and 10 mM FeSO₄. Twenty-four hours later, cytosolic proteins (10 µg) were isolated and β-catenin levels were determined by immunoblot. The respective densitometric analyses (arbitrary units, AU) of the β-catenin/β-actin ratio bands are presented in Panels A and B. Data are reported as means ± SEM for 3 independent experiments (one-way ANOVA followed by the Newman-Keuls test). *P < 0.05 vs control; **P < 0.01 vs staurosporine + Wnt-3a; †P < 0.05 vs control and FeSO₄. β-actin was used as internal control.
toxicity. Some studies have also demonstrated that lithium, which activates Wnt signaling pathways, can have a protective action against toxicity in cell models of HD (36).

It has been reported that Aβ causes alterations in the Wnt signaling components, including a reduction of β-catenin levels in cultured rat hippocampal neurons (11) (38) and GSK-3β activation (39), which could favor tau protein hyperphosphorylation (40). Our findings suggest that Wnt-3α may counteract such adverse effects of Aβ and other oxidative and apoptotic insults. On the other hand, we cannot rule out the possibility that protection induced by Wnt signaling does not involve pathways different from the canonical cascade, which is linked to β-catenin activation. Based on our results, we suggest that Wnt-3α protein has a protective effect against different insults including staurosporine, iron, 3-nitropropionic acid, and amyloid-β, and possibly against different neurodegenerative diseases.

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