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Soluble NgR Fusion Protein Modulates the Proliferation of Neural Progenitor Cells via the Notch Pathway

Xin Li · Huanxing Su · Qing-Ling Fu · Jiasong Guo · Daniel H. S. Lee · Kwok-Fai So · Wutian Wu

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Abstract NogoA, myelin-associated glycoprotein (MAG) and oligodendrocyte myelin glycoprotein are CNS myelin molecules that bind to the neuronal Nogo-66 receptor (NgR) and inhibit axon growth. The NgR antagonist, soluble NgR1-Fc protein (sNgR-Fc), facilitates axon regeneration by neutralizing the inhibitory effects of myelin proteins in experimental models of CNS injury. Here we aim to investigate the effect of sNgR-Fc on the proliferation of neural progenitor cells (NPCs). The hippocampus cells of embryonic rats were isolated and cultured in vitro. The expression of nestin, βIII-Tubulin, GFAP and Nogo-A on these cells was observed using immunocytochemistry. In order to investigate the effect on proliferation of NPCs, sNgR-Fc, MAG-Fc chimera and Notch1 blocker were added respectively. The total cell number for the proliferated NPCs was counted. BrdU was applied and the rate of proliferating cells was examined. The level of Notch1 was analyzed using Western blotting. We identified that NogoA is expressed in NPCs. sNgR-Fc significantly enhanced the proliferation of NPCs in vitro as indicated by BrdU labeling and total cell count. This proliferation effect was abolished by the administration of MAG suggesting specificity. In addition, we demonstrate that sNgR-Fc is a potent activator for Notch1 and Notch1 antagonist reversed the effect of sNgR-Fc on NPC proliferation. Our results suggest that sNgR-Fc may modulate Nogo activity to induce NPC proliferation via the Notch pathway.

Keywords Nogo-66 receptor · Rat neural progenitor cells · Notch1 · NogoA · Myelin-associated glycoprotein

Introduction

Neural progenitor cells (NPCs) are capable of self-proliferating and differentiating into the three major cell lineages of central nervous system (CNS), and has the potential for replacement of lost or dysfunctional neurons or glial cells. Stem cell replacement therapy may 1 day become a promising strategy for CNS injuries and neurodegenerative disorders. However, the limited regenerative capacity of both endogenous and grafted NPCs is attributed to the
inhibition of NPC proliferation and differentiation in situ by local environmental factors. The proliferation and differentiation of NPCs are determined by the effects of extrinsic and intrinsic signals coming from substrates, medium components and several complex interactions among cells. Therefore, a better understanding of the role of the molecular environment to NPC neurogenesis may be crucial for developing stem cell therapy.

Several proteins associated with CNS myelin possess axon growth inhibiting properties. These include NogoA [1], myelin-associated glycoprotein (MAG) [2], and oligodendrocyte myelin glycoprotein (OMgp) [3]. All three bind the Nogo66 receptor (NgR1) [4] and the paired immunoglobulin-kuje receptor B (PirB) [5] to mediate their inhibitory influence. Multiple lines of evidence suggest that the myelin proteins and NgR1 may affect NPC activities in addition to the effects on axon regeneration. Besides being expressed in the adult neurons and weakly in adult non-neuronal cells, NgR1 is also expressed in the spinal cord, the brain of chicken and human embryo [6] and in the NPCs derived from rat spinal cords [7]. NogoA is expressed in neurons in a variety of areas of both fetal and adult human and rat brains [8]. It is also expressed in oligodendrocyte progenitor cells [9]. NogoA promotes NPCs to differentiate to the glial lineage while inhibiting their differentiation into neurons [10]. Nogo-P4 (the active segment of Nogo-66) inhibited the differentiation of NPCs derived from rat spinal cords [7]. Since the expression levels of NogoA, MAG and OMgp are upregulated after CNS injury, they may be important factors for NPC neurogenesis.

The NgR1 antagonist, recombinant rat soluble NgR-Fc fusion protein [11], effectively blocked the interaction of myelin proteins with NgR1 and has been shown to promote recovery in rodent models of CNS injuries [12–16]. Notch1 is an important signaling pathway in the embryogenesis, hematogenesis and the differentiation of NPCs during development [17, 18]. Upon activation by Notch ligands, Notch intracellular domain (NICD) is cleaved, released from the whole receptor, and activated transcription of its downstream target genes [19]. So far, Hairy/Enhancer of Split (Hes) genes appears to be the primary downstream mediators of Notch signaling. Among them, Hes5 is considered to be an essential effector of Notch-mediated activity [20]. In the developing brain, activated Notch signaling maintains NPCs and promotes proliferation of neural progenitors [21, 22]. We hypothesize that NogoA and NgR1 are involved in the proliferation of NPCs and the NgR antagonist, sNgR-Fc, may affect NPC proliferation. In this study, we examined the expression of NogoA in NPCs and investigated whether sNgR-Fc promotes the proliferation of NPCs via Notch signaling pathway in vitro.

### Methods

#### Preparation of NgR1-Fc Protein

The form of sNgR-Fc used for this study, AA-rNgR(310)-rFc [12], is an improved variant form of the NgR-ecto-Fc fusion protein reported previously [15]. This protein comprises a 310 amino acid fragment of rat NgR1 fused to a rat IgG1 Fc fragment, in which Cys266 and Cys309 were replaced with alanine residues in order to eliminate heterogenous disulfide bonds [23]. The construct was expressed in Chinese hamster ovary cells, protein was purified, and binding to Nogo66, OMgp, and MAG was verified using previously established methods [15]. This modified protein inhibits the Nogo66-NgR interaction and promotes neurite growth of rat dorsal root ganglia and cerebellar granule neurons in vitro with similar potency as the unmodified sNgR-Fc [12].

#### Primary Neurosphere Culture, Differentiation and Immunocytochemistry

The procedures for isolation of embryonic NPCs have been described previously [24]. The hippocampus of embryonic day (E) 16 embryo of Sprague–Dawley (SD) rats from Laboratory Animal Unit, The University of Hong Kong was dissected, rinsed in cold CMF-HBSS and dissociated mechanically. The cells were collected by centrifugation and resuspended in Dulbecco’s modified Eagle’s Medium/HamF12 (DMEM/F12) supplemented with epidermal growth factor (EGF; 20 ng/ml, Gibco, CA, USA), basic fibroblast growth factor (bFGF, 20 ng/ml, Sigma, MO, USA), B27 supplement (2%, Gibco, CA, USA), N2 supplement (1%, Gibco, CA, USA), penicillin (50 U/ml), and streptomycin (50 μg/ml). The cells were adjusted to $1 \times 10^5$ cells/ml for cell culture. Half of the culture medium was replaced every 3 days. To identify the cultured cells, some of neurospheres after cultured for 7 days were plated on poly-l-lysine-coated coverslips and grown in culture medium. After neurospheres were attached on the coverslips (2 h), they were fixed with 4% paraformaldehyde (PFA) for 20 min, washed, permeated in 0.25% Triton-X100/phosphate buffered saline (PBS) and blocked with 10% goat serum. Cells were then incubated with mouse anti-Nestin (1:100, Santa Cruz Biotechnology, CA, USA) at 4°C for 16 h. After washing, the cells were incubated with a fluorescent Alexa 568-conjugated secondary antibody (1:400, Molecular Probes, CA, US) for 2 h at room temperature. Cell nuclei were counterstained with 4’,6-diamidino-2-phenylindole (DAPI, Sigma, MO, USA). The above neurospheres were harvested, dissociated and plated as described above, but in culture medium without EGF and bFGF for initiating and promoting differentiation.
After 3 days of differentiation, the cells were immunostained using mouse anti-βIII-tubulin antibody (Sigma, MO, USA) for undifferentiated neural progenitor cells, rabbit anti-glia fibrillary acidic protein (GFAP) (1:1,000; Chemicon, Temecula, California, USA) for astrocytes or mouse anti-Rip (1:50, Abcam, CB, UK) for oligodendrocyte, mouse anti-βIII-tubulin antibody (1:1,000, Sigma, MO, USA) for neurons. The cells were incubated with a fluorescent Alexa 488 or 568-conjugated secondary antibody (1:400, Molecular Probes, CA, US) and Cell nuclei were counterstained with DAPI (Sigma, MO, USA).

In order to examine the expression of NogoA and Notch1 in NPCs, the above neurospheres were dissociated and seeded onto poly-L-lysine-coated coverslips in 1 × 10^5 cells/well with the above medium. The cells were then double-immunostained using rabbit anti-NogoA (1:100, Santa Cruz, Biotechnology, CA, USA) and mouse anti-Nestin antibodies as described above. The dissociated NPCs were seeded onto poly-L-lysine-coated coverslips in 1 × 10^5 cells/well with the above medium. sNgR-Fc of 50, 500, or 1,000 ng/ml was added to the cultures, respectively. Untreated cultures were used as control. In order to examine whether the sNgR-Fc promotes the proliferation of NPCs by inhibiting the effects of CNS myelin proteins, recombinant rat MAG-Fc chimera (200 ng/ml, R & D Systems, Inc, MN, USA) was added to the medium to neutralize sNgR-Fc (500 ng/ml). To confirm the role of Notch signaling pathway in the proliferation activity of sNgR-Fc on the NPCs, we used a Notch1 antagonist, MW167 (γ-Secretase Inhibitor II, 10 μmol/l, Calbiochem, CA, USA), to block Notch activity.

### Proliferation Assay for Neurospheres and NPCs in Vitro

After 7 days in culture, the neurospheres were treated with 0.25% trypsin for 10 min and then mechanically dissociated. The dissociated NPCs were seeded onto poly-L-lysine-coated coverslips in 1 × 10^5 cells/well with the 500 μl medium. sNgR-Fc of 50, 500, or 1,000 ng/ml was added to the cultures, respectively. Untreated cultures were used as control. In order to examine whether the sNgR-Fc promotes the proliferation of NPCs by inhibiting the effects of CNS myelin proteins, recombinant rat MAG-Fc chimera (200 ng/ml, R & D Systems, Inc, MN, USA) was added to the medium to neutralize sNgR-Fc (500 ng/ml). To confirm the role of Notch signaling pathway in the proliferation activity of sNgR-Fc on the NPCs, we used a Notch1 antagonist, MW167 (γ-Secretase Inhibitor II, 10 μmol/l, Calbiochem, CA, USA), to block Notch activity.

### Counting the Total Cell Number for the Proliferation Assay

To analyze neurosphere growth, the volume of living neurospheres was measured using an inverted microscope (Axiovert 100, Carl Zeiss, Jena, Germany) and ImageJ software (National Institute of Health, USA). In order to determine the cell number, the treated spheres were collected at 24, 48 and 72 h after plating and the cells were counted as described [25]. The neurospheres in one well were trypsinized with 2 ml of 1× trypsin-EDTA (ethylene-diaminetetraacetic acid) (0.05% trypsin, 0.02% EDTA-4Na) (GIBCO, Invitrogen corporation, CA, USA) for 2 min. Trypsin–EDTA was then removed and cells were incubated at 37°C for another 2 min. After incubation, cells were suspended in 1 ml medium and single-cell suspensions at 2–5 × 10^5 cells/ml were obtained. Ten μl cell suspension was loaded to the chamber of hemocytometer by capillary action and number of cells were counted using microscope. A total 72 cultures (6 cultures in each concentration of sNgR-Fc and control group every day) was used for the total cell number counting. The volume of living neurospheres after different treatments was measured using an inverse confocal microscope (LSM 510 META; Carl Zeiss, Oberkochen, Germany) and ImageJ software (National Institute of Health, USA) as previous report [25].

### BrdU Labeling for the Cells of Proliferation Assay

For bromodeoxyuridine (BrdU) labeling, 72 h after the above treatments, 10 μM BrdU was applied to the cultures for 2 h to label the nuclei of proliferating cells. Then, immunocytochemistry on the cultures was performed to detect BrdU positive cells after fixation with 4% PFA. Briefly, the cultures were denatured in 2 N HCl at 37°C for 30 min and rinsed in 0.1 M borax acid buffer (pH 8.5) at room temperature for 10 min. Then, they were blocked with 1% bovine serum albumin, 10% normal goat serum, and 0.3% Triton X-100 in 0.01 M PBS for 1 h at room temperature. After that, cultures were incubated with mouse anti-BrdU antibody (1:1,000, Sigma, MO, USA) overnight at 4°C and incubated with fluorescent Alexa Fluor® 488 or 568 goat anti-mouse secondary antibody (1:400, Molecular Probes, CA, USA) for 2 h at room temperature. Finally, the nuclei of all the cells were counterstained with DAPI (Sigma, MO, USA). A total thirty-six cultures (six cultures in each group) was used for the proliferation assay.

After immunostaining, the number of immunopositive cells in each culture was counted using a ×20 objective. Five visual fields were selected on the edge of up, down, right and left, and the central of each slide, which is a total of 30 visual fields were counted for each parameter. DAPI-stained cells, representing the total cell number, and BrdU-immunoreactive cells in each field were counted. Percentages for each antibody-immunoreactive cells were calculated.

### Western Blotting for Notch1 and Hes 5

Cells treated with sNgR-Fc, MAG or MW167 were plated at 1.0 × 10^5 cells/ml in flasks and were collected 72 h later for Notch1 analyses using Western blotting. The cells were trypsinized, washed in cold 0.01 M PBS and collected
as pellets by centrifugation at 1,500 rpm for 5 min. The cell pellet was then resuspended in 80 μl of radioimmunoprecipitation (RIPA) buffer [1% Nonidet P-40, 0.5% sodium deoxycholate, 0.1% SDS, 10 μg/ml PMSF (phenylmethylsulfonyl fluoride)], 20 μg/ml Aprotinin, 10 μg/ml Leupeptin, 100 mM sodium orthovanadate, in PBS and incubated on ice for 30 min. The protein lysate was then centrifuged at 13,500 rpm for 20 min at 4°C. The supernatant was collected and measured for the protein concentration using the DC Protein Assay Kit (Bio-Rad, Hercules, CA, USA). A 20 μg aliquot of protein from each sample was subjected to 8–12.5% SDS-polyacrylamide gel electrophoresis and transferred onto PVDF membrane. The membranes were blocked with 5% nonfat dry milk and 2% bovine serum albumin in Tris-buffered saline containing 0.1% Tween 20 (TBST) for 1 h at room temperature. Incubation with rabbit anti-Notch1 (1:1,000, Millipore, Temecula, CA, USA) antibody against NICD or rabbit anti-Hes 5 (1:1,000, Abcam, CB, UK) was performed for 16 h at 4°C. After washing, the membranes were incubated with horseradish peroxidase-conjugated goat anti-rabbit antibody (1:2,000, Dako, Denmark) in 5% non-fat dry milk and 2% bovine serum albumin in TBST for 1 h at room temperature and immunoreactive proteins were detected using the enhanced chemiluminescence method (ECL, Amersham, Piscataway, NJ, USA). Protein loading was controlled using a goat against anti-actin antibody (1:2,000, C-11, Santa Cruz Biotechnology, CA, USA). The experiments for Western blotting were repeated 3 times, and representative data is shown.

TUNEL Staining for Apoptosis

Neurospheres were prepared, cultured and collected as described above, with either 500 ng/ml sNgR-Fc or neurosphere medium alone. After fixation, an In Situ Cell Death Detection Kit FITC (Roche, Mannheim, Germany) was used for immunocytochemical detection of apoptosis, based on labeling of DNA strand breaks (TUNEL technology) according to the manufacturer’s instructions. Fluorescein isothiocyanate (FITC) fluorescence was monitored using an inverse confocal laser scanning microscope. As positive control, neurospheres were treated with DNAase (2,000 U/ml, Sigma, Deisenhofen, Germany). As negative control, labeling solution containing FITC was used.

Data and Statistical Analysis

The values presented are mean ± SEM. Statistical analysis was performed using one-way analysis of variance (ANOVA) followed by post-hoc tests (Student-Neuman-Keuls) for comparisons of results obtained from more than two groups. Data were analyzed statistically with the software SPSS 12.0. P < 0.05 was considered statistically significant.

Results

Characterization of Neural Progenitor Cells Derived from the Hippocampus

In the presence of epidermal growth factor and basic fibroblast growth factor, the hippocampus-derived neural progenitor cells proliferated quickly to form small clusters and then to larger neurospheres floating within the culture medium in 3–7 days. Immunocytochemistry revealed that the cells in neurospheres were positive for Nestin (Fig. 1a–c). The differentiation potential of NPCs was evaluated 3 days after mitogens were removed and were cultured in the presence of 10% fetal bovine serum (FBS). Neuronal cells that had differentiated from the neurospheres were confirmed by their immunoreactivity for βIII-Tubulin (Fig. 1d–f). The differentiated cells were also immunoreactive for glial fibrillary acidic protein (GFAP) (for astrocytes) (Fig. 1g–i) and Rip (for oligodendrocytes) (Fig. 1j–l). It showed that these neurospheres could differentiate into neurons, oligodendrocytes and astrocytes simultaneously.

NogoA was Expressed in NPCs in Vitro

Cultured NPCs stained strongly for NogoA (Fig. 2). We also confirmed that NogoA was coexpressed with Nestin (Fig. 2a–d). The proportion of NogoA+/Nestin+ was 98.2 ± 1.23%.

sNgR-Fc Activated the Proliferation of NPCs

We next investigated the effect of sNgR-Fc on the proliferation of NPCs in culture. NPCs were treated for up to 3 days with sNgR-Fc in different concentrations (0, 50, 500/ml and 1,000 ng/ml). There was significant difference for the treatment of sNgR-Fc on day 1 [F(3, 20) = 2.341; P < 0.01], day 2 [F(3, 20) = 4.437; P < 0.01] and day 3 [F(3, 20) = 6.674; P < 0.001]. There was significant increase in the total number of cells in the 500 ng/ml and 1,000 ng/ml groups compared to the control group on day 1, 2 and 3 respectively (Fig. 3a, P < 0.01). A significant increase in volume of the neurospheres was also found after the treatment of sNgR-Fc [F(3, 20) = 8.342; P < 0.001] (Fig. 3b). Representative photomicrographs of neurospheres at 3 days are shown in Fig. 3c (for control) and Fig. 3d (for 500 ng/ml sNgR-Fc). In order to evaluate the effects of sNgR-Fc on the induction of apoptosis, the cells were treated with sNgR-Fc for 3 days and analyzed...
Fig. 1 Identification of the cultured neural progenitor cells. **a-c** The neurosphere expressed nestin positively. **d-f** Neurospheres could differentiate into βIII-Tubulin<sup>+</sup> cells. **g-i** Neurospheres could differentiate into Rip<sup>+</sup> cells. **j-l** Neurospheres could differentiate into GFAP (glial fibrillary acidic protein) positive cells. Cultured neural progenitor cells were counterstained with 4',6-diamidino-2-phenylindole (DAPI).

Fig. 2 The expression of NogoA in the cultured neuronal progenitor cells. NogoA expressed in neural progenitor cells (*arrows*). NogoA staining (*green*) (**a**), Nestin staining (*red*) (**b**) demonstrates the co-localization of NogoA and Nestin in neural progenitor cells (**d** (*arrows*). The nucleus was counterstained with DAPI (4',6-diamidino-2-phenylindole) (**c**).
for apoptosis using a TUNEL assay (Fig. 3e). DNAse treatment as a positive control resulted in nearly 75% TUNEL-positive nuclei. Label solution as a negative control gave no positive signal. Few apoptotic nuclei were detected and there was no difference between the control and sNgR-Fc-treated groups. These results suggest that the effect of sNgR-Fc on the increase of NPCs number might be a result of proliferation, and not because of the reduction of cells through apoptosis.

Next, we examined whether the increased new cells were derived from NPCs. We analyzed progenitor cell proliferation with BrdU incorporation. This technique has been extensively used to measure DNA replication during proliferation of mammalian cells. We found that there was significant different in the cell proliferation with BrdU incorporation \[ F(5, 24) = 12.465; P < 0.001 \]. The amount of BrdU incorporation rose and was higher in 500 ng/ml and 1,000 ng/ml groups compared to the control (Fig. 4a–d, g, \( P < 0.01 \)). Here we also examined the effect of sNgR-Fc on the proliferation of NPCs via counting DAPI positive cells. The total number of DAPI showed that there was more cells after sNgR-Fc treatment (500 ng/ml or 1 µg/ml) \[ F(5, 24) = 11.128; P < 0.001 \] (Fig. 4h).

MAG Inhibits the Proliferative Effect of sNgR-Fc on NPCs

In order to investigate the mechanism by which sNgR-Fc promotes the proliferation of NPCs, we added MAG into the culture medium. The NgR-Fc fusion protein can directly bind with MAG-Fc [15]. Using the BrdU incorporation technique, we found that the ability of sNgR-Fc (500 ng/ml) to promote the ratio of BrdU incorporation was clearly inhibited after the administration of MAG-Fc (Fig. 4e, g). It suggests that sNgR-Fc may promote the proliferation of NPCs by inhibiting the action of myelin proteins such as MAG.

sNgR-Fc Activated Notch1 in NPCs

We investigated the level of Hes5 and NICD in sNgR-Fc treated NPCs using Western blotting (Fig. 5a). Significant differences among Hes 5 \[ F(5, 12) = 5.624; P < 0.001 \] and NICD \[ F(5, 12) = 8.366; P < 0.001 \] levels were subsequently revealed by ANOVA analysis. The levels of Hes5 were higher for sNgR-Fc (500 ng/ml) compared to control treatment. The levels of NICD were higher for sNgR-Fc (50 ng/ml) compared to control treatment, peaked for 500 ng/ml sNgR-Fc group and then declined slightly in the 1,000 ng/ml sNgR-Fc group. MW167, an inhibitor of the Notch1 signaling pathway, significantly blocked the activation of Notch1 by sNgR-Fc (Fig. 5a, b). Using the BrdU incorporation technique, we found that MW167 significantly inhibited the ability of sNgR-Fc (500 ng/ml) to promote the ratio of BrdU incorporation (Fig. 4f, g). Furthermore, we found that the administration of MAG-Fc blocked the activation of Notch1 with the treatment of sNgR-Fc (Fig. 5a, b). These data further suggest that sNgR-Fc may promote the proliferation of NPCs by inhibiting the activity of myelin proteins. We next demonstrated that Notch1 was coexpressed with Nestin in the NPCs using immunocytochemistry. Figure 5e, f shows

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**Fig. 3** Enhanced proliferation of neural progenitor cells with the treatment of sNgR-Fc in vitro. Neural progenitor cells (NPCs) were treated for up to 3 days with different concentration of sNgR-Fc (0, 50, 500 and 1,000 ng/ml). The total cell number was quantified. 

**a** There was an increase in the number of the cells with the treatment of sNgR-Fc. \( P < 0.01 \) compared to control group in the same time point. 

**b** Quantification of neurosphere volume. sNgR-Fc-treated neurospheres show significantly increased volume compared to untreated neurospheres (\( * P < 0.01 \)), six cultures for each group.

**c–d** Photographs of neurospheres, 3 days after plating, show larger neurospheres after sNgR-Fc treatment (500 ng/ml) (d) than in control neurospheres (c). 

**e** Apoptosis was detected using TUNEL with FITC-conjugated dUTP for the neurospheres. Positive control cultures were treated with DNase I to generate nicked DNA (e1). Terminal deoxynucleotidyl transferase was omitted for negative controls (e2). Few apoptotic nuclei were detected and there was no difference between the control (e3) and sNgR-Fc-treated (e4) groups.
Fig. 4 sNgR-Fc increased bromodeoxyuridine incorporation. Merged images of bromodeoxyuridine (BrdU) and 4',6-diamidino-2-phenylindole (DAPI). The NPCs were treated as described below for 3 days and then pulsed with BrdU (10 μM) for 2 h. a The control group, cultured neural progenitor cells (NPCs) received no further treatment. b–d Cultured NPCs received treatment of different dosage of sNgR-Fc. e Cultured NPCs received treatment with sNgR-Fc (500 ng/ml) plus myelin-associated glycoprotein (MAG). f Cultured NPCs received treatment with sNgR-Fc (500 ng/ml) plus MW167, a Notch1 inhibitor. sNgR-Fc increased the ratio of BrdU-positive neural progenitor cells in dosage-dependent manners, while MAG or MW167 decreased such increased ratio of BrdU incorporation. g A percentage of BrdU-positive cells in total number of cells with different treatments. h The total number of DAPI positive cells five visual fields in different groups. In g, h, *P < 0.05, *P < 0.01 compared with control group. Open triangle P < 0.01 compared to 500 ng/ml group. Six cultures for each group (Mean ± SEM)
representative photomicrographs of Notch1 immunolabeling in NPCs after the treatment of sNgR-Fc (500 ng/ml).

Discussion

The present study aimed to elucidate whether NgR1 fusion protein promotes the proliferation of NPCs and to examine the possible signaling mechanisms involved. Our results demonstrated that: (1) NogoA was expressed in cultured NPCs; (2) sNgR-Fc promoted NPC proliferation; (3) MAG reversed the proliferative effect of sNgR-Fc on NPCs. (4) sNgR-Fc activated Notch1 and a Notch1 inhibitor reversed the effect of sNgR-Fc on NPC proliferation.

Effect of NgR1 Fusion Protein on NPC Proliferation

NgR1 is associated with axon regeneration and neurite growth. NgR1 is expressed in neurons in different CNS regions, such as the cortex, subventricular zone and dentate subgranular zone; and upregulated after brain ischemic injury [26]. In this study we identified that the NPCs in the neurosphere and the single NPC strongly expressed NgR1. The NgR1 ecto-domain, sNgR1 is an NgR1 antagonist and promotes axon regeneration after spinal cord injury [12, 15], stroke [14] or dorsal root rhizotomy [12, 16]. Our data suggest, in addition to axon regeneration, sNgR-Fc also promotes the proliferation of NPCs.

Neural stem cells are characterized by the ability to undergo cell division and to differentiate into multiple cell types, e.g. neurons or glial cells. When stimulated by growth factors or cytokine, the NPCs could generate cleavage of asymmetry. Recently, much attention are focused on the therapeutic potential of neural stem cells to treat neurological disorders such as spinal cord injury [27, 28], stroke and Parkinson’s disease [29, 30]. The factor(s) responsible for the proliferation and differentiation of NPCs has been the subject of intense investigation because identification of regulators of NPCs may allow for their eventual manipulation in the CNS as therapies to replace the injured CNS neurons. We have confirmed including the use of BrdU incorporation experiments, a significant increase in the number of the cells after sNgR-Fc treatment. In addition, few apoptotic nuclei were detected in cells after sNgR-Fc-treatment. This suggests that the effect of sNgR-Fc on the increase in the number of NPCs may be linked to proliferative than anti-apoptotic mechanisms.
Molecular Mechanism of sNgR-Fc on NPC Proliferation

We also examined the possible mechanism of the effect of sNgR-Fc on the proliferation of NPCs. Nogo is expressed in three distinct isoforms, NogoA, NogoB and NogoC, which is known as one of the myelin inhibitors that induces growth cone collapse and inhibit neurite outgrowth. Cell proliferation was inhibited significantly in the NogoC overexpressed PC12 cell which was used as a neuronal model [31]. If NPCs derived from the spinal cords of rats were allowed to differentiate in the presence of Nogo-P4, the average neurite length and the percentage of differentiated neurons were decreased [7]. NogoA was also reported to be expressed in the oligodendrocyte progenitor cells [9]. Much NogoA was observed in all of the development periods in rat hippocampus, and especially in the embryonic and neonatal rats [32]. In this study, we identified that there was NogoA expression in the NPCs. The NogoA expressed in NPCs has the potential to inhibit the proliferation of NPCs. The Ngr1 has been shown to constitute a site of signal convergence for three axon inhibitory proteins found in CNS myelin [33]. Blockade of NgR-dependent signaling by sNgR1 allows an assessment on the significance of summated inhibition from NogoA, MAG, and OMgp rather than inhibiting individual NgR ligands as there are several reported in the literature aside from myelin molecules such as LINGO-1, p75 and Troy. Since the expression of these molecules in NPCs are unclear or found at a later stage e.g. LINGO-1 [9], it is possible that sNgR-Fc may promote the proliferation of NPCs by reversing the inhibitive effect of NogoA in particular that of the Nogo-66 loop. This is because MAG that also binds NgR1; eliminated the proliferative effect of sNgR-Fc. It will be interesting to verify if the amino-terminus of NogoA would also participate in this process. Reagents such as IN-1 antibody will be useful tools for delineating the NogoA epitopes involved in NPC proliferation.

The Signaling Pathway for sNgR-Fc Stimulated NPC Proliferation

Notch1 plays a pivotal role in the regulation of vertebrate neurogenesis [34]. During this process, Notch-mediated cell–cell interaction is essential for maintaining dividing cells and subsequent generation of cell type diversity. Upon activation by Notch ligands, NICD is cleave, released from the whole receptor, and activated transcription of its downstream target genes Hes 1 and Hes 5 [19]. Activation of Notch inhibits cellular differentiation, and abnormality of the Notch pathway leads to premature neuronal differentiation, the lack of some cell types, and severe defects of tissue morphogenesis [35]. Here we showed that the administration of sNgR-Fc activated Notch1, suggesting that sNgR-Fc may stimulate the proliferation of NPCs by activation of Notch1 or its downstream molecules. These effects could be abolished by Notch1 inhibitor suggesting specificity. It will be of interest to test if the gamma-secretase activity is directly related to NPC proliferation. Our data suggest that sNgR-Fc stimulated NPC proliferation via activation of the Notch pathway.

Our data indicate that sNgR-Fc promotes the proliferation of NPCs in vitro via the Notch1 signaling pathway. sNgR-Fc may provide an attractive therapeutic strategy of neuronal repair for treating CNS injuries and neurodegenerative disorders.

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