TrkB antibody elicits cytotoxicity and suppresses migration/invasion of transitional cell carcinoma cells

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Received April 29, 2010; Accepted June 2, 2010

DOI: 10.3892/ijo_00000745

Abstract. BDNF (brain-derived neurotrophic factor) and its receptor TrkB (tropomyosin receptor kinase B) play important roles in the progression of cancer, including transitional cell carcinoma (TCC) cells reported in our previous investigation. In this study, we used a specific TrkB antibody (Ab) to evaluate its effects on survival, proliferation and migration/invasion in three TCC cell lines (BFTC905, T24 and TSGH8301) in vitro. The TrkB Ab at 1 and 3 μg/ml, but not the TrkA or TrkC Abs, significantly elicited cytotoxicity in TCC cells. The TrkB Ab at 3 μg/ml also induced apoptosis of TCC cells, which may result from up-regulation of phospho-p38 plus down-regulation of survivin and securin expression. The TrkB Ab at 0.5 μg/ml, which did not show cytotoxicity, suppressed migration of TCC cells and invasion of BFTC905 cells, possibly mediated through increased E-cadherin, decreased BDNF-stimulated phospho-PLCγ1 and reduced MMP-9 activity. These results indicate that TrkB blockade may be a new strategy for TCC therapy.

Introduction

Trk (tropomyosin receptor kinase) family of tyrosine kinase receptors consists of TrkA, TrkB and TrkC. Each Trk is activated by its preferred neurotrophin ligands, e.g., NGF (nerve growth factor) binds to TrkA, BDNF (brain-derived neurotrophic factor) and NT4/5 (neurotrophin 4/5) bind to TrkB, and NT3 binds to TrkC. These Trk receptors and their ligands play important roles in neuronal survival and differentiation (1).

Apart from neuronal functions, Trk receptors also possess oncogenic character (2,3). Some cancer cells have been reported to express Trk receptors (4). For example, BDNF protects neuroblastoma cells from chemotherapeutic agent induced cytotoxicity (5). However, high level of TrkA expression in neuroblastoma specimens is associated with favorable outcome (6). Trk receptors may be potential targets for cancer therapy, but specific Trk blockade has only been investigated in few studies (7). Our previous study demonstrated the existence of BDNF and TrkB in transitional cell carcinoma (TCC) cell lines (8). BDNF and TrkB are overexpressed in human TCC specimens compared to normal urothelium (unpublished results). TrkB activation mediated by BDNF is a survival signal for TCC (8). Therefore, we hypothesize that selective TrkB blockade may be exploited as an alternative new treatment for TCC cells. Our results demonstrate that TrkB antibody inhibits proliferation, and migration/invasion, and induces apoptosis in TCC cells.

Materials and methods

Cell culture. Three human TCC cell lines including BFTC905, TSGH8301 and T24 were chosen for the investigation. The cells were maintained as described previously (8). Grading of BFTC905 (9) and TSGH8301 (10), both originated from Taiwanese TCC patients, T24 from ATCC (American Type Culture Collection, Manassas, VA) are 3 and 2, and 2/3, respectively.

Cell proliferation and MTT cytotoxicity assay. TCC cells were plated at a density of 8x10^4 cells in 6-well dishes with RPMI medium containing 10% FBS. The plating day was designated as day 1. One or 3 μg/ml of TrkB antibody (Ab) (sc-8316, Santa Cruz Biotechnology Inc., Santa Cruz, CA), 3 μg/ml of human IgG (Sigma, St. Louis, MO), or PBS was added separately to the culture dishes on day 2. Calculation of the cell counts in a hemocytometer was performed on day 5 (72 h after TrkB Ab administration).

For cytotoxicity assay, TCC cells with 1x10^4 cells per well were seeded in 96-well plates. Time designation was the same as cell growth assay. Different concentrations (1 or 3 μg/ml) of TrkB Ab (AF175, R&D Systems, Minneapolis, MN), TrkAb (sc-8316, Santa Cruz), or TrkC Ab (AF373, R&D Systems) were administered on day 2. Human IgG antibody treatment served as a negative control. Conventional MTT (3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide, Sigma) assay followed 72 h (day 5) after Ab administration (8).
Western blot. Standard procedures were followed (8). Primary antibodies used included anti-phospho-p38 (no. 9211, Cell Signaling, Danvers, MA), anti-p38 (sc-535, Santa Cruz), anti-phospho-PLC\(\gamma\)1 (no. 2821, Cell Signaling), anti-PLC\(\gamma\)1 (no. 2822, Cell Signaling), E-cadherin (sc-7870, Santa Cruz), anti-survivin (GF029, Millipore, Billerica, MA), and anti-securin (ab3305, Abcam, Cambridge, MA). Expression of ERK2 (sc-154, Santa Cruz), or actin (sc-1616, Santa Cruz) was used as the internal standard. Can Get Signal™ immunostain was used for enhancement of phosphorylated protein signals (Toyobo, Osaka, Japan). The intensity of immunoreactive proteins was calculated with NIH software ImageJ V.1.40.

TUNEL (terminal deoxynucleotidyl transferase dUTP nick end-labeling) assay. The TACS-XL™ in situ apoptosis kit (R&D Systems) was used for the detection of DNA fragmentation of TCC cells. Cells were treated with 3 μg/ml of IgG or TrkB Ab for 24 h and fixed with 4% paraformaldehyde for 10 min. Subsequent procedures suggested by the assay kit were followed. Cells were chosen randomly from 10 fields and photographed under x400 light microscopy for semiquantitation. The number of positive nuclear stain indicating DNA fragmentation and thus apoptosis was counted. The ratio of apoptotic cells divided by total cells in a chosen field was calculated, and the data were expressed as number of apoptotic cells per 100 cells counted.

Invasion/migration assay. Wound healing assay was used for the migration study. BFTC905, T24 and TSGH8301 cells were cultured separately in 3-cm dishes in an RPMI medium containing 0.5% FBS. Several straight wounds were established by the tip of a 2-μl pipette on the dish filled with cancer cells. TrkB Ab at 0.1 or 0.5 μg/ml, or human IgG at 0.5 μg/ml were delivered into the medium on the day of wound development. After 24 h, cells were stained with hematoxylin and the wounds were observed and photographed under x40 magnification.

The BioCoat™ growth factor reduced Matrigel™ Invasion Chamber (BD Biosciences, MA) was used for in vitro invasion study. TCC cells (2.5x10^4) in 500 μl of a medium containing 1% FBS were delivered into the chamber. TrkB Ab (0.5 μg/ml) was added to the chamber, and IgG (0.5 μg/ml) served as control. Subsequently, the chambers were put into 24-well plate containing 750 μl of culture medium plus 10% FBS. After 48-h incubation, invading cells on the bottom

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Figure 1. Cytotoxicity of Trk Ab in TCC cells. (A) Viabilities of TCC cells were measured by MTT assay 72 h after TrkA, TrkB, or TrkC Ab treatment. The viability ratio was calculated as the ratio of O.D. value from Ab treatment divided by that of PBS. TrkB Ab elicited significant cytotoxicity on 3 TCC cell lines with concentration of 1 μg/ml (0.64±0.04, 0.61±0.05 and 0.69±0.05 for BFTC905, TSGH8301 and T24, respectively; n=3-4) and 3 μg/ml (0.61±0.02, 0.59±0.04 and 0.52±0.07 for BFTC905, TSGH8301 and T24, respectively; n=3-4). (B) Apoptotic BFTC905 cells were detected by the TUNEL assay. Dark staining of nuclei indicated DNA fragmentation of cells after 3 μg/ml of TrkB Ab or IgG treatment for 24 h (x400). (C) Significantly increased TUNEL-positive cells were observed after TrkB Ab treatments compared to IgG treatments (BFTC905, TSGH8301 and T24; left to right).
surface of the membrane were counted under microscope. Cell numbers per field in both chambers (x200) counted from 10 randomized fields by two staff members were averaged.

**MMP-9 activity assay.** Matrix metalloproteinase-9 (MMP-9) activity in culture medium was measured by ELISA method (Biotrak activity assay system, code RPN2634, Amersham Pharmacia Biotech, Little Chalfont, UK). TrkB Ab at 0.5 μg/ml was added to 1x10^6 BFTC905 cells cultured in 24-well dishes with serum-free medium. After 24 h, MMP-9 activity in the medium was assayed according to manufacturer's instructions.

**Statistical analysis.** Data are presented as mean ± SEM (standard error of mean). The difference between experimental and control group was evaluated by Student's t-test. In all experiments, p<0.05 was considered statistically significant and labeled by the symbol * in the figures.

**Results**

TrkB Ab elicited cytotoxicity and suppressed the proliferation of TCC cells. Cytotoxicity was measured by MTT assay 72 h after TrkA, TrkB, or TrkC Ab treatment. As shown in Fig. 1A, only TrkB Ab (both 1 and 3 μg/ml for 3 TCC cell lines) exerted significant toxicity on TCC cells while TrkA Ab, TrkC Ab, and IgG was without cytotoxic effects. Fig. 1B shows that more TUNEL-positive BFTC905 cells were observed after 3 μg/ml TrkB Ab when compared to IgG treatment. Apoptotic TCC cells were increased significantly 24 h after 3 μg/ml TrkB Ab treatment (4.0±1.2 vs. 15.3±1.5, 5.0±1.5 vs. 16.7±2.4, and 3.6±1.8 vs. 16.7±2.3 for BFTC905, TSGH8301 and T24 cells after IgG vs. TrkB Ab treatment, respectively; n=3) (Fig. 1C). Furthermore, proliferation of TCC cells was significantly suppressed 72 h after administration of 3 μg/ml TrkB Ab (53.3±6.1 vs. 24.0±4.5, 61.0±2.8 vs. 34.0±4.0, 90.6±1.6 vs. 47.0±7.4x10^4 in BFTC905, TSGH8301 and T24 cells after PBS vs. TrkB Ab treatment, respectively; n=3) (Fig. 2). No differences in cell numbers among PBS, IgG and 1 μg/ml TrkB Ab treatments were observed.

TrkB Ab induced phospho-P38, and inhibited survivin and securin expression in TCC cells. The effect of TrkB Ab on the expression of phospho-p38 mitogen-activated protein kinase (MAPK) was examined because activation of p38 by several chemotherapeutic agents has been reported to mediate cancer cells apoptosis (11). After the addition of 3 μg/ml of TrkB Ab to TCC cells, a time-dependent increase in phospho-p38 expression was found (Fig. 3A). The increased expression was obvious as early as 2 h post-treatment, the earliest time studied, and continued to increase for 24 h. The intensity ratio of phospho-p38/total p38 was increased from zero (in the absence of treatment) to ~2 after 24 h treatment (Fig. 3B).

Survivin, a member of the inhibitor of apoptosis protein family, has been shown to be a promising biomarker for diagnosis and prognosis of bladder cancer. Decreased expression of survivin in bladder cancer specimens has been reported to be a good prediction of response to therapies (12,13). As shown in Fig. 3C and D, expression of survivin decreased significantly 24 h after the addition of 3 μg/ml TrkB Ab in TCC cells. Securin, also known as putative tumor-transforming gene (PTTG), plays important roles in tumorigenesis and invasiveness in various neoplasms (14). The expression of securin decreased significantly 24 and 48 h after the addition of 3 μg/ml TrkB Ab in TCC cells (except 24-h expression in TSGH8301 cells, Fig. 3E and F).

TrkB Ab inhibited migration and invasion of TCC cells. To evaluate further the effect of TrkB Ab on bladder cancer cell migration and invasion, two relatively low concentrations of TrkB Ab at 0.1 and 0.5 μg/ml were chosen for wound healing assay because these concentrations were found to be devoid of effect on the proliferation in culture medium containing 0.5% FBS. These three bladder cancer cell lines did not proliferate within 24 h in this starved condition (results not shown). The time-dependent changes of wound healing are shown in Fig. 4A. Narrowing and even disappearance of wounds were observed 24 h after human IgG treatment. Delayed healings were observed after 0.5 μg/ml TrkB Ab treatment, exerting the greatest inhibitory effects on migration, especially for BFTC905 and TSGH8301 cells. In addition, 0.5 μg/ml TrkB Ab significantly inhibited the invasive ability of BFTC905 cells in Matrigel assay (Fig. 4B). Although trends of invasive suppression for TSGH8301 and T24 cells were observed after TrkB Ab treatment, the difference, when compared to IgG treatment, did not reach statistical significance.
TrkB Ab increased E-cadherin expression, inhibited phosphorylation of PLCγ, and decreased MMP-9 activity in BFTC905 cells. Epithelial-to-mesenchymal transition (EMT) through loss of E-cadherin mediates migration of TCC cells and is associated with poor outcome (15). Fig. 5A shows the changes of E-cadherin in three cell lines 24 h after 0.5 μg/ml TrkB Ab treatments. Increased E-cadherin expression was observed in BFTC905 cells compared to IgG treatment. E-cadherin was not expressed in T24 cells after IgG and TrkB Ab treatments. Weak E-cadherin expression in TSGH8301 cells was noted, but was not changed after TrkB Ab treatment.

TrkB has been demonstrated to mediate neuronal migration via PLCγ (16), which has also been found to play a role in the migration and invasion of cancer cells after growth factor (e.g., epidermal growth factor) stimulation (17,18). Due to significant inhibition of invasive activity in BFTC905 cells after TrkB Ab treatment, the changes of phospho-PLCγ were measured in BFTC905 cells by Western blotting.
Addition of 10 nM rhBDNF for 10 min increased the phosphorylation of PLC$_{1}$. IgG (0.5 μg/ml) or TrkB Ab (0.5 μg/ml) alone was without effect on the phospho-PLC$_{1}$ level. Pre-treatment with 0.5 μg/ml of human IgG did not alter the response elicited by 10 nM rhBDNF. In contrast, pre-treatment with TrkB Ab (0.5 μg/ml) for 24 h almost completely inhibited phospho-PLC$_{1}$ expression 10 min after the addition of 10 nM rhBDNF.

The level of MMP-9 is higher in high grade and more invasive type of urothelial carcinoma (19). Thus, we examined the MMP-9 activity in culture medium of BFTC905 cells by ELISA assay. MMP-9 activity was significantly inhibited by 40% after TrkB Ab treatment for 24 h (2.08±0.07 ng/ml vs. 1.26±0.05 ng/ml, IgG vs. TrkB treatment, respectively).

Discussion

BDNF and TrkB have been found in various cancer tissues, e.g., prostate cancer (20), and hepatocellular carcinoma (HCC) (21). However, there are only few reports investigating TrkB blockade as a new target therapy. Indolo-carbazole alkaloid K252a was widely used as the Trk family inhibitor in neuroscience and cancer studies (22,23). However, K252a is also a potent inhibitor of protein kinases, e.g., protein kinase C (PKC) (24), and myosin light chain kinase (MLCK) (25). It also inhibits the phosphorylation of c-met protein (MET) (26). Moreover, PKC (27), MLCK (28) and MET (29) also play important roles in cancer progression. Thus, the therapeutic effects of K252a on cancers may result from acting on multiple targets. RNA interference of TrkB is a specific method for TrkB inhibition (7,30), but the clinical application of systemic siRNA administration has just begun (phase I trial) (31). Neutralizing antibodies targeting on RTK (receptor tyrosine kinase), such as trastuzumab, have been widely used in cancer patients (32). Therefore, we used TrkB antibody for its blocking effect in TCC cells.

The wild-type TrkB protein contains five domains. A recent structure-activity study demonstrated that tyrosine kinase activity of TrkB plays critical functions for anti-anoikis, tumorigenesis and metastasis (33). Deletion of LRM (leucine-rich motif) or Ig-2 (IgG-like domain 2) eliminated TrkB functions, indicating that they play important roles in mediating BDNF response. In our study, the sc-8316 antibody binding to the extracellular domain of TrkB is against amino acids 160-340, which covers the extracellular Ig-1 and Ig-2 domain. Consequently, it will inhibit the binding of BDNF to the extracellular domain of TrkB and block the downstream signaling pathways. This may explain why sc-8316 antibody inhibited proliferation and migration/invasion of TCC cells in vitro, as well as blocked the phosphorylation...
of TrkB and PLCγ1 despite rhBDNF stimulation in our current and previous studies (8). The TrkB inhibition also induced p38 phosphorylation, suppressed survivin and securin expression, and finally caused apoptosis of TCC cells in our experiments. As far as we know, our results are the first report to elucidate the signaling mechanism among p38, survivin, securin and TrkB in cancers. Cytotoxicity and apoptotic effect have been observed after survivin siRNA treatment in T24 cells (34,35). Survivin inhibition may partly explain the TrkB Ab-induced apoptosis in TCC cells. Overexpressed securin in glioma specimens is associated with an unfavorable consequence (36). Our study also demonstrated the expression of securin in TCC cells, and the inhibition of survivin expression by TrkB blockade. The detailed signalings among survivin, securin and TrkB should be investigated in the future.

We also studied the effect of TrkB Ab on the migration and invasion in TCC cells. Migration of BFTC905 and TSGH8301 cells was delayed when TrkB receptor was blocked with concentration of TrkB Ab that was without anti-proliferative effect but weak inhibitory effect was observed in T24 cells. The discrepancy between 3 TCC cell lines might have resulted from the E-cadherin expression. BDNF activation of TrkB receptors mediated the migration and invasion of head and neck squamous cell carcinoma cells via EMT promotion, which upregulated twist and down-regulated E-cadherin (37). No dynamic twist changes following TrkB treatment in 3 TCC cell lines were observed (results not shown). Upregulated E-cadherin in BFTC905 cells could explain the migration and invasion inhibition after TrkB blockade. Our study is consistent with a previous report which demonstrated the absence of E-cadherin protein and mRNA level in T24 cells (38). It may explain the weak inhibition of migration and the absence of inhibition on invasion in T24 cells after TrkB blockade. Furthermore, it has been shown that PLCγ1 mediates neuronal migration after TrkB activation (17) and cancer cell migration/invasion after EGF stimulation (18). Thus, we examined the phosphorylation of PLCγ1 in BFTC905 cells after rhBDNF stimulation. Phospho-PLCγ1 activation was evident 5 min after rhBDNF stimulation and lasted for 30 min (results not shown). The activation was inhibited when cells were pre-treated with TrkB Ab for 24 h. Thus, PLCγ1 phosphorylation may also play important roles in the migratory and invasive effects of BDNF-TrkB interaction in TCC cells. BDNF has been shown that PLCγ1 activation was inhibited when cells were pre-treated with K-252a (39). Decrease of extracellular MMP-9 activity in BFTC905 cells after TrkB Ab treatment results in poor degradation of the matrigel, suggesting that TrkB Ab elicited inhibitory effect on invasion.

In conclusion, TrkB blockade by TrkB Ab elicited cytoxicity, induced apoptosis, and suppressed migration/invasion of TCC cells in vitro. Agents aimed at blocking TrkB may have potential as a new therapy for bladder cancer.

Acknowledgements
This study was partially supported by grant-in-aid from Tzu Chi University (THC) and Tzu Chi General Hospital (YTH), Hualien 970, Taiwan, and a grant from NSC (NSC-97-2314-B-303-016, YTH), Taiwan.

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