Effect of β-carotene on cancer cell stemness and differentiation in SK-N-BE(2)C neuroblastoma cells

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Abstract. Neuroblastoma is a solid tumor often diagnosed in childhood. While there have been intense efforts to develop a treatment for neuroblastoma, current therapies remain unsuccessful due to high rate of resistance and metastasis. Most cancers originate from a subset of self-renewing cells, primarily cancer stem cells (CSCs), which establish a tumor through continuous self-renewal and differentiation. The successful elimination of CSCs is an important goal in the development of effective strategies to achieve complete remission for cancers. Although β-carotene has been associated with several anticancer mechanisms, the efficacy of β-carotene against CSCs remains unclear. In the present study, β-carotene was shown to reduce cell growth and induce neuronal cell differentiation, concomitant with a marked increase in the phosphorylation of extracellular signal-regulated kinases (ERK) (p42/p44). More importantly, β-carotene inhibited self-renewal characteristics of CSCs and decreased expression of several stem cell markers. Levels of mRNA and protein of Drosophila delta-like 1 homolog (Drosophila) (DLK1) were downregulated following treatment with β-carotene. In addition, knockdown of DLK1 by siRNA enhanced the inhibitory effect of β-carotene on colony formation of neuroblastoma cells. β-carotene also potentiated the effect of cisplatin on the self-renewal characteristics of CSCs in neuroblastoma, revealing that β-carotene has the capacity to resensitize cells to cisplatin cytotoxicity by directly targeting CSCs. In conclusion, β-carotene was shown to strongly increase the anticancer efficacy against neuroblastoma cancer stem-like cells. Moreover, these results suggest that the targeting of CSCs is a novel mechanism of β-carotene. Thus, β-carotene is a potential chemotherapeutic reagent for this cancer.

Introduction

Neuroblastoma (NB) is the most common solid tumor diagnosed in children, and it originates from immature neuronal cells of the sympathetic nervous system. NBs are also comprised of multiple cell phenotypes, including neuroblasts, non-neuronal cells and melanocytes (1). In NB cell lines, as well as in primary NB tumors, three distinct cell types have been identified: Schwannian/melanoblastic precursors (S-type), neuroblastic/neuroendocrine precursors (N-type), and morphologically intermediate stem cells (I-type). I-type stem cells represent precursors of both N- and S-type cells, and also represent the most immature and highly malignant population of NB cells (2). Based on significant similarities in the gene expression profiles of normal neuroblasts and malignant NB, as well as the morphological and functional differentiation that can be achieved with NB cells in culture, the differentiation process of NB can be studied in vitro.

A growing body of evidence has demonstrated that many types of cancers are driven by a small population of cancer cells, termed cancer stem cells (CSCs), or cancer-initiation cells (3). CSCs have the capacity to undergo differentiation, self-renewal and tumor generation. Moreover, CSCs have the potential to induce tumor growth, metastasis and resistance to radiation and chemotherapy (4). Understanding the signaling pathways that mediate functional characteristics is important in preclinical models, including animal models or in vitro cell lines. Drosophila delta-like 1 homologue (DLK1) is a member of the epidermal growth factor-like homoeotic protein family. DLK1 has been known to modulate differentiation signaling in adipocytes and hematopoietic stem cells and has been characterized as a bona fide stem cell gene in neuronal tumors (5,6). Mitogen activated protein kinase (MAPK) signaling pathways have been shown to play pivotal roles in growth, differentiation and stress responses (7,8). It has been reported that MAPK signaling is affected by the expression of DLK1 (9). Correspondingly, extracellular signal-regulated kinases (ERK)(p42/44) are phosphorylated during neuronal differentiation and the activation of MEK/ERK signaling was found to decrease the self-renewal capacity of embryonal stem cells (8,10).

During tumor treatment, most cancer cells, including a majority of differentiated tumor cells and actively replicating cells, respond to cytotoxic treatments. However, acquired...
multidrug resistance to conventional therapies can develop in CSCs. Therefore, the successful elimination of both actively replicating tumor cells, and slowly dividing CSCs, may represent a more effective strategy for the complete treatment of various types of cancer.

β-carotene (BC) is a provitamin A carotenoid, and a source of retinoids in dark green and orange fruits and vegetables. Observational studies have reported that an increase in the intake of dietary BC with increased BC plasma concentrations is associated with a decreased risk of several types of cancers (11,12). Accumulating evidence also suggests that mechanisms involving antioxidant potential, alterations in intracellular signaling pathways, and regulation of cell proliferation and apoptosis, contribute to the anticarcinogenic effects associated with BC (13,14). Retinoic acid (RA) has also been shown to promote the differentiation of embryonic stem cells and CSCs (15), and is used as a chemotherapeutic drug for various types of cancers (16). Moreover, direct, or indirect effects of several dietary compounds, including sulforaphane and curcumin on self-renewal pathways, have been identified (17,18).

Currently, little is known about the role of BC in the differentiation and self-renewal characteristics of CSCs. Therefore, in the present study, the chemopreventive effects of BC on the induction of differentiation and attenuation of cancer cell stemness were investigated. Moreover, based on the potential for CSCs to contribute to chemoresistance, the chemotherapeutic effect of administering BC and cisplatin in combination was investigated, as well as the hypothesis that the effects of BC are mediated through the targeting of NB cancer stem cell-like cells.

Materials and methods

Reagents and cell culture. BC, all-trans-retinoic acid (RA), tetrahydrofuran (THF) and cis-diaminedichloroplatinum(II) (cisplatin) were purchased from Sigma Chemical Co. (St. Louis, MO, USA). U0126 and PD98059 were obtained from Cell Signaling Technology Inc. (Danvers, MA, USA). Two human NB cell lines, SK-N-BE(2)C [BE(2)C] and SH-SY5Y (SY5Y), were obtained from the American Type Culture Collection (ATCC, Rockville, MD, USA) and maintained in a 1:1 mixture of Minimum Essential Medium (MEM) and Ham's F-12 medium supplemented with 10% fetal bovine serum (FBS) and antibiotics. BC was freshly prepared in THF containing 0.01% butylated hydroxytoluene (BHT), then added to the culture medium to a final concentration of 5-20 µM. The concentration of THF in culture medium was adjusted to be 0.01%.

Culture and reagents. Minimum Essential Medium (MEM) was obtained from VWR International (Bridgeport, PA, USA) and supplemented with 10% FBS and antibiotics. Ham's F-12 medium supplemented with 10% FBS and antibiotics was obtained from GIBCO (Grand Island, NY, USA). FBS was purchased from HyClone (Logan, UT, USA). U0126 and PD98059 were obtained from Cell Signaling Technology Inc. (Danvers, MA, USA). U0126 (20 µM) and PD98059 (20 µM) were added to the culture medium to assess their effect on cell proliferation. The concentration of THF in culture medium was adjusted to be 0.01%.

Measurement of neurite outgrowth. BE(2)C cells were plated in triplicate in 6-well plates (4x10^4 cells/well) and treated with varying doses of BC for 5 days. A process having at least twice the diameter of the associated cell body, and possessing a terminal growth edge, was considered as a neurite. The percentage of cells with outgrowing neurites in ~50 cells/well was counted. The mean and standard error of the mean (SEM) is reported. Three independent experiments for each treatment were performed.

Clonogenic assays. To determine the self-renewal characteristics of cells, clonogenic assays were performed as previously described (20). Briefly, BE(2)C cells were plated in 6-well plates (500 cells/well) and treated with varying concentrations of BC for 10 days. After fixing the cells with 0.9% NaCl, colonies were stained with crystal violet (Sigma). Colonies containing at least 50 stained cells were counted. Plating efficiency (PE) was calculated as follows: PE = (colony number/total cell number) x 100%.

Sphere formation assays. To facilitate sphere formation, poly-(2-hydroxyethyl methacrylate) (polyHEMA; Sigma-Aldrich)-coated plates were prepared by applying 500 µl of 10% polyHEMA stock solution [12% polyHEMA solution in 95% ethanol (w/v)] into each well of a 6-well plate (21). Sphere medium was prepared with DMEM-F12 (3:1; Invitrogen, Carlsbad, CA, USA) containing 2% B27 (Invitrogen), 20 ng/ml recombinant human epidermal growth factor and 40 ng/ml recombinant human fibroblast growth factor basic (PeproTech, Rocky Hill, NJ, USA). After 6 days, the number of tumor spheres containing >50 cells was counted.

Antibodies and western blot assay. Western blot assays were performed as previously described (22). Briefly, cell lysates were separated by SDS-PAGE and transferred onto polyvinylidene fluoride (PVDF) membranes (Millipore, Billerica, MA, USA). The membranes were incubated with 5% non-fat dried milk, followed by antibodies against DLK1, SOX2 (Millipore), Notch1 (Novus Biologicals, Littleton, CO, USA), β-tubulin III (Sigma), phospho-ERK (p42/p44), total ERK (Cell Signaling), or β-actin (Santa Cruz Biotechnology, Santa Cruz, CA, USA). Bound antibodies were detected using enhanced chemiluminescence reagents (Animal Genetics Inc., Suwon, Kyonggi-do, Korea).

RNA isolation and real-time PCR. Cells were treated with various doses of BC for 4 days and total RNA extraction was performed on each sample using TRIzol reagent (Invitrogen). cDNA of 1 µg total RNA was synthesized by reverse transcription with RevertAid First Strand cDNA synthesis kit (Thermo Fisher Scientific, Waltham, MA, USA), and real-time PCR was performed with StepOnePlus™ (Applied Biosystems, Foster City, CA, USA) to examine the relative DLK1 mRNA levels among the samples according to the manufacturer's instructions. PCR primers were as follows: human DLK1, 5'-CTG...
morphological and functional differentiation of NB cells that possess cancer cell stemness and undergo morphological and functional differentiation in vitro (20). To examine the effect of BC on NB cell differentiation, cells were treated with various concentrations of BC. Following treatment, BE(2)C cells underwent morphological differentiation in a dose-dependent manner manifested by an increase in neurite growth, which is a hallmark of neuronal differentiation (Fig. 2A and B). In these assays, RA was used as a positive control to induce neuronal differentiation. The observed increase in neuronal differentiation following treatment with BC was confirmed with an upregulation of β-tubulin III that was detected by western blotting (Fig. 2C). Neurite elongation and protein expression profiles determined following BC treatment were also very similar to those obtained following RA treatment.

**BC-induced neuronal cell differentiation involves ERK phosphorylation.** It has been reported that ERK1/2 (p42/p44) is phosphorylated during the differentiation of neuronal progenitor cells (8). Similarly, when phosphorylation of ERK1/2 was assessed following treatment with BC, a marked increase was observed, whereas the expression of total ERK was unchanged (Fig. 3A). To examine whether MAPK pathways are involved in BC-induced NB cell differentiation, the MAPK-specific inhibitors, U0126 and PD98059, were used. In these studies, phosphorylation of ERK1/2 was completely blocked following treatment with U0126 and 10 or 20 µM BC, while phosphorylation of ERK1/2 was only partially blocked by PD98059 at both concentrations of BC (Fig. 3B). Consistent with these results, BC-induced neurite elongation was also completely blocked by U0126 treatment, and only partially blocked by PD98059 (Fig. 3C). In combination, these data clearly demonstrate that BC induces neuronal differentiation of NB, and plays an important role in the MAPK phosphorylation that is involved in this process.
Figure 2. BC induces neuronal cell differentiation. (A) Phase contrast microscopy was used to examine the morphological features of differentiated BE(2)C cells following treatment with BC for 5 days. RA (1 µM) was used as a positive control. Neuronal differentiation was evaluated based on the number of lengthened neurites (indicated with arrows) that were counted. Magnification, x100. (B) The percentage of neurite-bearing cells was calculated from samples plated in triplicate. At least three independent experiments were performed, and the means ± SEM are presented. Different letters at the top of the given bars of the histogram indicate that those values are significantly different from each other (P<0.05). (C) Expression of neuron-specific β-tubulin III was analyzed by western blotting with detection of β-actin used as a loading control. BC, β-carotene; RA, all-trans-retinoic acid.

Figure 3. BC-induced neuronal differentiation involves ERK phosphorylation. BE(2)C cells were treated with BC for 5 days and (A) ERK phosphorylation was analyzed by western blotting with detection of total-ERK used as a loading control. (B and C) Cells were treated with various doses of BC, then with each MEK/ERK inhibitor (U0126 and PD98059) on the next day. After 3 days of incubation, immunoblots (B) and cell morphology (C) were analyzed. β-actin was used as a loading control, and arrows indicate lengthened neurites representative of neuronal differentiation. BC, β-carotene.
a clonosphere and following treatment with BC exhibited a marked decrease in clonosphere formation compared to the controls (P<0.05), and this decrease was dose-dependent (Fig. 4C and D). To further verify the effect of BC on neuronal stem cell markers, BE(2)C cells were treated with BC for 4 days and the protein expression of several well-known stem cell markers, including Sox2 and Notch1 was assessed. The expression of these stem cell markers was found to be downregulated (Fig. 4E). Taken together, these results suggest that BC is highly effective in decreasing the self-renewal capacity of NB cells, and in eliminating most NB stem cells.

**BC downregulates DLK1 expression.** DLK1 is recognized as a neuronal stem cell marker and has been hypothesized to play an important role in the growth of stem/progenitor cells and cancer cells (20). Here, in cells treated with BC, mRNA and protein levels of DLK1 were found to be downregulated (Fig. 5A and B). To determine whether a combination of BC treatment and knockdown of DLK1 inhibits cancer cell stemness, BE(2)C cells were transfected with siRNA oligonucleotides designed to target DLK1. Subsequently, endogenous expression of DLK1 was found to be completely abrogated (Fig. 5C). Moreover, in the clonogenicity assay, treatment with BC or knockdown of DLK1 significantly inhibited colony formation (22 and 49%, respectively, P<0.05). Furthermore, the combination of BC treatment and DLK1 knockdown resulted in a 65% inhibition in clonogenicity compared to the control (Fig. 5D). Collectively, inhibition of DLK1 by siRNA was found to potentiate BC-mediated inhibition of the self-renewal capacity of NB cells. Consequently, BC treatment with DLK1 knockdown may provide a more effective chemotherapeutic effect by regulating CSCs.

**BC potentiates the anticancer effect of cisplatin by inhibiting the self-renewal capacity of NB cells.** CSCs resist conventional chemotherapeutic drugs and are likely to play an important role in cancer relapse. Therefore, it is important to develop cancer therapeutic agents targeting CSCs. Furthermore, most chemotherapeutic agents have undesirable side-effects. Therefore, various agents are administered in combination to increase their therapeutic efficacy while decreasing associated toxicity. Cell growth was examined using MTT assays to assess whether BC treatment in the presence of cisplatin has an additive effect on BE(2)C cells (Fig. 6A and B). For these assay,
Figure 5. DLK1 expression is downregulated by BC. (A) Protein expression of DLK1 was assessed by western blotting 4 days after BE(2)C cells were treated with BC or RA. (B) Real-time PCR was performed to analyze levels of DLK1 mRNA following the treatment of BE(2)C cells with BC or RA for 3 days. β-actin was used as an internal control. Different letters at the top of the given bars of the histogram indicate that those values are significantly different from each other (P<0.05). (C) BE(2)C cells were transfected with DLK1-specific or non-targeting control siRNA. After 48 h, cell lysates were prepared, and expression of DLK1 was analyzed by western blotting using β-actin as a loading control. (D) Clonogenic assays were performed to examine the self-renewal capacity of the cells described in Fig. 4. BC, β-carotene; RA, all-trans-retinoic acid.

Figure 6. Combination treatment with BC and cisplatin decreases stem cell characteristics. (A) Cell growth inhibition by cisplatin was determined using MTT assays. BE(2)C cells were treated with various doses of cisplatin for 4 days. (B) Cell growth inhibition in response to treatment with BC and cisplatin was analyzed. BE(2)C cells in 6-well plates were treated with cisplatin (0.5 μM), BC (10 μM) or both agents (0.5 μM cisplatin + 10 μM BC). After 4 days, cell viability was analyzed using MTT assay. (C and D) Sphere formation assay was performed for BE(2)C cells treated with a combination of cisplatin and BC as described in Fig. 4. Spheres were imaged and counted. Magnification, x100. BC, β-carotene; cis, cisplatin.
cells were treated with increasing doses of BC and cisplatin over 4 days. IC_{50} values calculated from the dose-response data obtained were ~0.59 µM for cisplatin (Fig. 6A) and ~10.8 µM for BC (Fig. 1A). To evaluate the antitumor effects of BC in combination with cisplatin, NB cells were treated with 0.5 µM cisplatin, 10 µM BC, or both. After 4 days, cell growth was analyzed using MTT assay. Whereas treatment with BC or cisplatin alone significantly decreased cell viability, the combination of BC with cisplatin further suppressed NB cell growth (Fig. 6B). Furthermore, to evaluate the effects of BC and cisplatin on the self-renewal characteristics of CSCs, a sphere formation assay was also conducted. The combination treatment had an additive effect on reducing sphere formation although either BC or cisplatin showed inhibition of sphere formation (Fig. 6C and D). These results suggest that BC has the potential to suppress the growth of CSCs, and the combination of BC with conventional chemotherapeutic agents could represent a more effective regimen for the treatment of this deadly childhood cancer.

Discussion

Neuroblastoma is the most common malignant tumor of the neural crest and arises within the sympathetic nervous system. In addition, the non-differentiated stem cell phenotype associated with NB has been identified as a potential independent risk factor for NB malignancy. Therefore, NB cells represent a model system applicable to the study of cancer stem cell-like cells. In the present study, treatment with BC was observed to induce the differentiation of NB cells and to decrease the self-renewal characteristics of CSCs. Therefore, BC may represent a novel therapeutic strategy for the targeting of CSCs associated with NB, thereby preventing recurrence and metastasis.

RA is a widely used chemotherapeutic drug that induces cell differentiation. Similarly, carotenoids have also been shown to induce cell differentiation (23,24). For example, in a study by Gross et al (23), both BC and lutein were shown to induce the differentiation of HL-60 cells, β-cryptoxanthin, one of the major carotenoids present in blood, has also been shown to induce the morphological differentiation of mouse Neuro2a cells, in vitro (24). Prasad et al (25) also successfully differentiated the murine NB cell line, NBA2, by treating cells with BC for 3 days, followed by irradiation with X-rays at a dose of 20 Gy and higher. However, evidence regarding the differentiation signaling induced by BC, and the efficacy of BC in regard to CSCs, remains unclear. In this present study, phosphorylation of ERK (p42/p44) was observed to markedly increase following treatment with BC. Moreover, the use of specific ERK inhibitors, PD98059 and U0126, confirmed the involvement of the ERK/MEK1/2 pathway in BC-induced neuronal differentiation. It has previously been proposed that U0126 directly inhibits MEK1/2 activity, whereas PD98059 is an inefficient inhibitor of MEK2 (26). The differential effects of these inhibitors on osteogenesis and adipogenesis have also been characterized (27). Similarly, in a previous study of RA-induced neurite growth, growth was completely blocked by U0126, yet only partially blocked by PD98059 (28). Collectively, these data suggest that BC-induced neuronal differentiation requires the inhibition of both MEK1 and MEK2. Furthermore, BC-induced neuronal differentiation appears to be partly mediated via a pathway also used for RA-induced neuronal differentiation. Therefore, the mechanistic details of the regulation of BC on ERK phosphorylation, as well as on the phosphorylation of other MAPKs require further analysis. However, our results suggest that ERK is an important mediator of BC-mediated neuronal differentiation.

In the present study, BC induced the neuronal differentiation of human NB cells, and several mechanisms could account for this observation. Following are examples proposed for NB differentiation induced by BC: i) BC could be converted to RA since BC is a precursor of RA and then cleaved via a central cleavage pathway (29); ii) differentiation may be induced by nuclear receptor-binding activities, subsequently leading to the induction of several transcription factors and signaling pathways involved in differentiation; or iii) RA signaling could be promoted by metabolites of BC. Further studies are needed to investigate these possibilities, as well as to determine the contributions of BC to the differentiation of CSCs and RA precursor function.

The self-renewal capacity of CSCs is associated with their tumorigenic potential (4). For example, the ability of CSCs to form colonies as well as anchorage-independent spheres is directly proportional to the proportion of self-renewing cells present. Correspondingly, an evaluation of these two characteristics can determine the effect of a particular treatment on CSCs. In the present study, NB cells were observed to form colonies and spheroids in the absence of BC, yet in the present of BC, the formation of colonies and spheroids was reduced. A decrease in the levels of stem cell markers, DLK1, Notch1 and Sox2 was also noted in BC-treated cells.

DLK1 is a transmembrane protein that is expressed extensively in immature cells with regenerative potential (30,31). It has previously been reported that various tumors express higher levels of DLK1 (32,33), and exogenous expression of DLK1 increases the self-renewal capacity of NB cells. In contrast, knockdown of DLK1 has been shown to enhance neuronal differentiation (20). In the present study, the combination of BC treatment with knockdown of DLK1 resulted in a significantly higher inhibition of colony formation than treatment with either agent alone. Since DLK1 is involved in MAPK signaling, it is hypothesized that BC and DLK1 also have roles in the same ERK pathway. Taken together, these results suggest that BC inhibits the self-renewal capacity of CSCs. Moreover, poorly differentiated tumors have recently been shown to have similar gene expression profiles as normal embryonic stem cells or progenitor cells (34,35). However, studies are ongoing to further elucidate the mechanisms by which stem cell genes regulate tumor progression and metastasis, as well as the role of nutrients and tumor microenvironments.

Although initially responsive to cytotoxic treatment, most tumors acquire a multidrug-resistant (MDR) phenotype and relapse. CSCs are likely to play a major role in chemotherapy resistance and cancer relapse. Therefore, it is important to develop therapies that bypass the drug resistance of CSCs. Currently, the most promising strategy for the treatment of cancer is the administration of two or more drugs with different mechanisms of action to produce synergistic, or at least additive anticancer effects. Cisplatin was the first platinum-derived drug produced, and is still used as a first-line
therapeutic agent against several types of cancers, including NB (36). While the chemotherapeutic effects of cisplatin are based on the formation of an adduct with DNA and proteins, as well as the disruption of several signaling pathways (37,38), the administration of cisplatin is also associated with serious side-effects, including nephrotoxicity and peripheral neuropathy due to immune suppression (39). Due to these toxicities and undesirable side-effects, equally effective and safer treatments are needed. It has been reported that a combination treatment of minocycline with BC or 13-cis-retinoic acid, along with cisplatin, can result in decreased cytotoxicity (40). Furthermore, a mixture of three or four vitamins containing BC showed the potential to enhance the growth-inhibitory effects of cisplatin on melanoma cells (41). In the present study, we demonstrated for the first time that BC can potentiate the cytotoxic effects of cisplatin against NB CSCs. In detail, BC increased the self-renewal activity of NB cells in vitro and mediated a reduction in sphere formation. In the latter case, we hypothesize that this role for BC contributed to the resensitization of CSCs to the chemotherapeutic agent, cisplatin. Further study of the molecular mechanisms of BC in relation to the effects of chemotherapeutic agents are needed to confirm this hypothesis, and are warranted to develop more potent and less toxic combination therapies. However, our results are based on in vitro cell culture rather than in vivo. Therefore, further in vivo studies using animals should be performed to support and confirm the present results.

In conclusion, the present study revealed that BC exhibits excellent anti-CSC qualities, and can induce neuronal cell differentiation via phosphorylation of MAPK and downregulation of DLK1. For example, treatment with BC decreased the expression of stem cell markers and suppressed the self-renewal characteristics of the NB cells assayed, including their colony and spheroid formation potential. Furthermore, our results suggest that BC may represent a potent medical therapeutic agent against several types of cancers, including NB. Further studies using animals should be performed to support and confirm the present results.

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References


