Functional effects of TrkA inhibition on system $x_C^-$-mediated glutamate release and cancer-induced bone pain

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Abstract

Breast cancer cells release the signalling molecule glutamate via the system $x_C^-$ antiporter, which is upregulated to exchange extracellular cystine for intracellular glutamate to protect against oxidative stress. Here, we demonstrate that this antiporter is functionally influenced by the actions of the neurotrophin nerve growth factor on its cognate receptor tyrosine kinase, TrkA, and that inhibiting this complex may reduce cancer-induced bone pain via its downstream actions on $xCT$, the functional subunit of system $x_C^-$. We have characterized the effects of the selective TrkA inhibitor AG879 on system $x_C^-$ activity in murine 4T1 and human MDA-MB-231 mammary carcinoma cells, as well as its effects on nociception in our validated immunocompetent mouse model of cancer-induced bone pain, in which BALB/c mice are intrafemorally inoculated with 4T1 murine carcinoma cells. AG879 decreased functional system $x_C^-$ activity, as measured by cystine uptake and glutamate release, and inhibited nociceptive and physiologically relevant responses in tumour-bearing animals. Cumulatively, these data suggest that the activation of TrkA by nerve growth factor may have functional implications on system $x_C^-$-mediated cancer pain. System $x_C^-$-mediated TrkA activation therefore presents a promising target for therapeutic intervention in cancer pain treatment.

Keywords
Cancer, pain, system $x_C^-$, glutamate, neurotrophin, nerve growth factor

Date Received: 16 January 2018; revised: 26 March 2018; accepted: 5 April 2018

Introduction

Breast cancer has a propensity to metastasize to the bone, causing severe cancer-induced bone pain (CIBP) and reduced quality of life. Accordingly, the effective management of CIBP is essential to improving quality of life for cancer patients. Our group has demonstrated that breast cancer cells secrete the signalling molecule glutamate via system $x_C^-$ and overexpress the gene encoding this antiporter.$^{1,2}$ When glutamate efflux becomes excessive, its character within the central nervous system switches from an excitatory transmitter to a well-characterized and pathologically relevant excitotoxin.$^{3,4}$ Considerable evidence suggests that glutamate released through system $x_C^-$ is involved in multiple physiological and pathological processes (Reviewed in Miladinovic et al.$^5$). Its dysregulation has been linked to chronic pain, an effect which is supported by animal models of CIBP.$^{6,7}$ Exogenous glutamate released from peripheral metastases may then sensitize surrounding nerves, directly acting on adjacent nociceptors within the bone environment.

The neurotrophin nerve growth factor (NGF) acts via its cognate receptor TrkA, through which it activates several downstream signalling pathways that regulate neuronal survival and differentiation. While normal breast epithelial tissue does not rely on NGF-TrkA

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signalling, breast cancer cells express the TrkA receptor,\(^8,9\) and NGF is a potent activator of the survival and proliferation of breast cancer cells.\(^8\) Furthermore, biologically active NGF is excessively synthesized and released by breast cancer cells, an effect which is blocked by the TrkA inhibitor K252, indicating the existence of an NGF autocrine loop.\(^10\)

In humans, endogenous NGF levels are elevated in pain states associated with chronic conditions,\(^11–14\) and NGF appears to be an important driver of the increased pain sensitivity characteristic of late stage metastatic cancer (reviewed in Wang et al.\(^15\)). Correspondingly, in animal models of nociception, NGF- and TrkA-knockout mice are hypoalgesic,\(^16,17\) while transgenic animals that overexpress NGF exhibit hyperalgesic behaviours.\(^18,19\)

Preclinical research suggests that anti-NGF treatment may be particularly effective in pain that originates in bone,\(^20–25\) as more than half of the nerve fibres that innervate bone are responsive to NGF.\(^26\)

To date, research on the neurotrophic consequences of metastatic cancer has largely focused on pathological nerve sprouting of sensory and sympathetic nerve fibres that innervate the tumour-bearing host tissue.\(^20,24,27,28\) Below, we propose an alternative consideration and complementary explanation of the role of NGF-TrkA in CIBP (See Figure 1 for schematic).

There is considerable evidence for bidirectional signalling between glutamate and neurotrophins. Glutamate stimulates neurotrophic expression,\(^29,30\) and the neurotrophin fibroblast growth factor-2 has been implicated in the functional upregulation of system \(x_C^-\) in mixed neuronal and glial cultures.\(^31\) Glutamate plays a key role in several neurodevelopmental and neurodegenerative disorders, and overexpression of system \(x_C^-\) has been shown to have neuroprotective effects in conditions of oxidative stress.\(^32\) Furthermore, NGF,\(^10\) TrkA\(^33\) and \(xCT\(^1,2\)\) are upregulated in breast cancer cells, supporting the notion that a functional relationship exists. The overlapping actions of \(xCT\) and NGF suggest that system \(x_C^-\) may be influenced by neurotrophic activity. To further elucidate the underlying pathophysiology of glutamate-related cancer pain, we have focused on the NGF-TrkA complex as a therapeutic target to reduce aberrant system \(x_C^-\)-mediated glutamate release that accompanies CIBP. Here, we test the hypothesis that systematically inhibiting the actions of NGF on TrkA may reduce CIBP by decreasing functional system \(x_C^-\) activity.

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**Figure 1.** Proposed mechanism by which system \(x_C^-\)-mediated glutamate release is regulated by TrkA-activated signalling pathways. NGF activates RAS-MAPK signalling through its tyrosine kinase receptor TrkA, which regulates \(xCT\) gene expression.
Materials and methods

In vitro

Cell culture. 4T1 triple-negative murine mammary gland carcinoma cells (American Type Culture Collection (ATCC)) were maintained in high glucose Roswell Park Memorial Institute (RPMI, Life Technologies); MDA-MB-231 triple-negative human breast carcinoma cells (ATCC) were maintained in high glucose Dulbecco’s Modified Eagle Medium (DMEM, Life Technologies). All cells were supplemented with 10% fetal bovine serum and 1% antibiotic/antimycotic (Life Technologies) incubated at 37°C and 5% CO2 and verified to be mycoplasma free before experimental use.

Effect of TrkA inhibition on breast cancer cell number. The selective TrkA inhibitor tyrphostin AG879 (Sigma-Aldrich), a validated and specific inhibitor of TrkA, was dissolved in dimethyl sulfoxide (DMSO) for preparation of a 79 mM stock. 4T1 and MDA-MB-231 cells were seeded at 5 × 10^5 cells/well in 96-well plates and treated with serial doses of AG879 from 50 μM through 3.125 μM dissolved in DMSO (maximum final concentration: 0.1%) or 0.1% DMSO alone (negative control).

Effect of NGF-TrkA on breast cancer cell number. Cells were seeded at 5 × 10^5 cells/well in 96-well plates and treated with various concentrations (0, 10 or 100 ng) of the recombinant human β-NGF protein (R&D Systems) in the presence of AG879 (6 μM) or control (DMSO) for 24 h, fixed with 10% neutral buffered formalin (NBF) and quantified using 0.1% Crystal Violet stain in 80% EtOH based on standard curves for each cell line. Absorbance was read on a spectrophotometer optical plate reader (BioTek) at λ = 570 nm.

Effects of NGF-TrkA on breast cancer cell metabolism. Cell lines were seeded and treated with serial doses of β-NGF (or phosphate-buffered saline (PBS)) in the presence or absence of AG879, as mentioned earlier, and incubated in 0.5 mg/mL 3-(4,5-Dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) labelling reagent (3-[4,5-dimethylthiazol-2-yl]-2,5-diphenyl tetrazolium bromide (Sigma-Aldrich) for the final 2 h of the 24-h treatment period. Absorbance was read on a spectrophotometer optical plate reader at λ = 650 nm.

Effects of NGF-TrkA on xCT protein level. Cells were treated with 100 ng β-NGF (or PBS) and 6 μM AG879 (or DMSO) for 24 h to determine the effects of NGF-TrkA activity on xCT protein levels, as quantified by Western blot analysis. Total protein concentrations were first determined using the Bio-Rad protein assay (Bio-Rad Laboratories) to ensure equal protein loading of whole cell lysates on 10% sodium dodecyl sulfate polyacrylamide gel electrophoresis gels and subsequent electrophoretic transfer to polyvinylidene difluoride membranes (Immobilon-P, Millipore Corporation). Expression of xCT was evaluated using respective primary antibody (Novus Biologicals), at a concentration of 1:1000, applied overnight at 4°C, with blocking in 5% skim milk in Tris-buffered saline with 0.1% Tween 20 (TBS-T). After washing, membranes were incubated in horseradish peroxidase-conjugated goat anti-rabbit secondary antibody (Santa Cruz) for 2 h at room temperature (RT) and visualized by enhanced chemiluminescence using an ECL-plus kit (GE Healthcare) on Amersham Hyperfilm (GE Healthcare). Calnexin (90 kDa) was used as a loading control. Processed blots were scanned and quantified using densitometry with ImageJ software (U.S. National Institutes of Health). Values were normalized to loading controls and compared to DMSO-treated control plates of the same cell line.

Effects of NGF-TrkA on xCT mRNA. mRNA was quantified using qualitative real-time polymerase chain reaction (qPCR) as previously described. Briefly, 10^6 4T1 cells were seeded in 10 cm dishes and treated with 100 ng β-NGF in the presence (6 μM) or absence (DMSO) of the TrkA inhibitor AG879 for 24 h. Total RNA isolation from cell pellets was conducted using the Qiagen RNaseasy kit (Qiagen Inc.). cDNA was then synthesized via reverse transcription using Superscript III and oligo DTs (Thermo Fisher Scientific), and qPCR assays were performed in duplicate using xCT primers (detailed in Table 1) and SsoAdvanced Universal SYBR Green Supermix (BioRad). Readings were normalized to the loading control Succinate Dehydrogenase Complex Flavoprotein Subunit A. Relative quantification analysis of gene expression data was conducted according to the 2^-ΔΔCT method.

Effects of NGF-TrkA on 14C-cystine uptake. The cellular uptake of radiolabelled 14C-cystine was quantified, as cystine uptake is mediated by an exchange with glutamate via system xCT. The cystine uptake protocol was adapted from previous reports. Briefly, 2 × 10^5 4T1 and MDA-MB-231 cells were seeded in 6-well plates and treated with 100 ng β-NGF, with or without 6 μM AG879 for 24 h. Media was then collected and stored at 4°C to quantify glutamate release (see below). Cultures were washed with Hank’s Buffered Salt Solution (HBSS) and exposed to 14C-cystine (0.015 mCi/mL) for 20 min at 37°C. Following 14C-cystine exposure, cultures were washed with ice-cold HBSS to wash away excess extracellular cystine and dissolved in 220 μL of lysis buffer (0.1% Triton-X in 0.1 N NaOH) for 30 min. A 100 μL aliquot of each cell lysate was
added to 1 mL Ecoscint-H scintillation fluid for quantification. Values were normalized to milligram of total protein using the Bradford protein assay (BioRad) and compared to 14C-cystine uptake in DMSO-treated controls on the same experimental plate.

Effects of NGF-TrkA on glutamate release. Immediately prior to 14C-cystine uptake quantification, growth media samples from 6-well plates were collected and stored at 4°C. Glutamate release was quantified using the AMPLEX Red glutamic acid assay kit (Invitrogen/Molecular Probes, Eugene, OR). This assay quantitates the fluorescence reaction product resorufin, which is produced in proportion to glutamate, for analysis on a CytoFluor Series, 4000 Fluorescence Multi-Well Plate Reader (PerSeptive Biosystems, Framingham, MA). To optimize this assay for measurement of media glutamate concentrations beyond 0.5 μM and eliminate the repeated cycling of glutamate and α-ketoglutarate, L-alanine and L-glutamate pyruvate transaminase were omitted from the reaction mixture. 37 Background glutamate from a media-only portion to glutamate, for analysis on a CytoFluor Series, 4000 Fluorescence Multi-Well Plate Reader (PerSeptive Biosystems, Framingham, MA). The Bradford protein assay, as mentioned earlier.

Immunocytochemistry. Immunocytochemistry was performed to confirm the presence of xCT on cultured 4T1 cells. Cells were seeded at 10^4 cells/well in 8-well chamber slides, fixed with 10% NBF, permeabilized with 0.1% Triton X-100, blocked with 1% bovine serum albumin, incubated in rabbit polyclonal anti-xCT primary antibody (Novus, 1:1000) for 1 h at RT and visualized using AlexaFluor-488 goat anti-rabbit secondary antibody (Life Technologies, 1:500) with 4′,6-diamidino-2-phenylindole (DAPI) counterstain using EVOS FL Cell Imaging System.

In vivo

Mice. Experimentally naive immunocompetent female BALB/c mice (Charles River Laboratories) aged 4 to 7 weeks old upon arrival were group housed in cages maintained at 24°C with a 12-h light/dark cycle and provided ad libitum access to food and water. All animal procedures were performed according to guidelines established by the Canadian Council on Animal Care under a protocol reviewed and approved by the Animal Research Ethics Board of McMaster University.

Experimental groups. Mice (N = 26) were randomly assigned to three treatment groups: sham surgery + vehicle control (n = 9), tumour + vehicle control (n = 9) and tumour + AG879 (n = 8). Mice were inoculated with either 2 × 10^4 4T1 cells (n = 17) or an equal number of C. L. killed 4T1 cell sham controls (n = 9) injected percutaneously into the right distal femur to establish tumours. Drug treatment was delayed until 7 days post-4T1 cell inoculation to allow sufficient time for the implanted cells to reliably establish a bone tumour without drug interference, more accurately mimicking clinical conditions. Osmotic pump loading and implantation were performed in accordance with the manufacturer’s specifications. AG879 was dissolved in DMSO and administered via intraperitoneally implanted Alzet model, 1002 mini-osmotic pumps (Durect) in a volume of 0.25 μL/h for a total of 5 mg/kg/day, based on the mean weight of the mice on the day of surgical implantation (20 g). Pumps were intraperitoneally implanted under isofluorane anaesthesia, and animals were systemically treated with AG879 from Day 7 post-cell inoculation through to end point.

All behavioural testing was performed with the groups being blinded, during the animals’ light cycle. Animals were randomly assigned to systemic therapy or control groups and were acclimated to the behavioural testing environment and equipment one week prior to commencing data recording and were tested twice weekly for the duration of the experiment. The progression of spontaneous and evoked pain pre- and post-4T1 cell inoculation was monitored using a battery of tests for nociceptive behaviours: The dynamic weight bearing (DWB) system (BioSeb), the dynamic plantar aesthesiometer (DPA; Ugo Basile) and open-field observations, including spontaneous guarding time and limb use. Three behavioural tests were performed prior to experiment Day 0 which served to establish a stable baseline for the animals’ normal behaviour; results are expressed for each animal as a percentage of these baseline scores. Behavioural testing occurred on days 6, 9, 13, 16 and 20 post-inoculation, and animals that had not yet reached end point were sacrificed on Day 21.
Spontaneous pain. Open-field tests, including spontaneous guarding of the affected limb and limb use, were used to visually assess ongoing and ambulatory nociception using previously validated tests. The time (s) spent spontaneously guarding the hindpaw represented ongoing pain and was recorded during a 5-min open-field observation period using a stopwatch. Guarding time was defined as the time the hindpaw was held aloft while ambulatory. During the 5-min spontaneous ambulation period, normal hindlimb use was observed and scored on a scale of 4 to 0: (4) normal use, (3) slight limp, (2) limp and guarding behaviour, (1) partial non-use of the limb in locomotor activity and (0) complete lack of limb use.

The DWB quantitates spontaneous pain-related discomfort and postural disequilibrium by recording the weight distribution of each point of contact of freely moving animals via a sensor pad covering the entire floor surface of the testing chamber. Weight-bearing data were directly captured at a sampling rate of 10 Hz, and the animal’s position on the sensor was manually validated following data collection using respective software (DWB v.1.3.4.36). To reduce experimenter bias and ensure spatial and temporal objectivity, a video camera recorded the animals’ position within the testing chamber over the capture period. Postural disequilibrium was defined as favouring the tumour-bearing limb and a compensatory shift of weight bearing to other body parts and was considered indirect evidence of nociception. Data for each test day were calculated as a mean weight recorded for all points of weight bearing and expressed as a single measurement of the weight borne by the affected limb as a ratio of the total weight recorded at each frame within the capture period. Data were normalized to the baseline scores prior to cancer cell inoculation for each mouse. A relative reduction in weight borne by the ipsilateral paw was considered evidence of postural disequilibrium.

Evoked pain. The DPA test, which is a semi-automated version of the classic von Frey test, was used to quantify mechanical allodynia and hyperalgesia in mice pre- and post-tumour cell inoculation. A mechanical metal filament was raised by an electrical actuator with variable force and acceleration. The filament stimulated the plantar surface of the affected hindpaw and contralateral hindlimb, and the threshold force and time at which each animal withdrew its paw were recorded. The average force of four independent measurements collected from each paw on each test day was calculated, and mean values were normalized to individual animal’s baseline DPA values. A reduction in force required to elicit paw withdrawal in the tumour-bearing limb was considered indicative of hypersensitivity manifested as deliberate withdrawal of the affected paw from the filament pressure.

Tumour xenografts. The 4T1 mammary carcinoma cell line is highly tumourigenic and invasive; the growth and metastatic spread of these cells in BALB/c mice very closely mimic human breast cancer in its proliferative and metastatic characteristics. Thus, this tumour was used as an animal model for stage IV human breast cancer. On experimental Day 0, mice were anaesthetized by isoflurane inhalation and injected with buprenorphine (0.05 mg/kg, subcutaneous; Schering-Plough) to mitigate surgically induced discomfort. Animals were inoculated with $2 \times 10^4$ 4T1 cells suspended in 25 μL sterile PBS percutaneously into the right distal femur. Mice were laid supine with the ipsilateral stifle joint bent to 90° to clear the patella and minimize tissue damage. A 26-gauge needle was then manually inserted between the medial and lateral condyles of the distal epiphysis by delicate rotation parallel to the longitudinal axis of the femur to breach the cortical bone. Once within the bone epiphysis, the injectate was slowly infused over the course of approximately 1 min. The contralateral hindlimb served as a negative control specific to each animal. This method of intrafemoral injection was selected to minimize damage to the surrounding tissues and has been successfully applied in mouse and rat models of bone cancer-induced nociceptive behaviour.

Transcardial perfusion and tissue collection. Throughout tumour development, each animal was monitored daily for limb use, overall health status and body weight. Animals were euthanized by transcardial perfusion when they no longer bore weight on the affected hindlimb. This experimental end point occurred on or before Day 21 post-tumour cell inoculation for all animals. Under sodium pentobarbital anaesthetic (90 mg/kg, i.p.), animals were perfused with 100 mL cold PBS, immediately followed by 100 mL cold 4% paraformaldehyde (PFA). Tumour-bearing femurs and surrounding tissues were dissected, X-rayed, post-fixed in 4% PFA, decalcified in 10% Ethylenediaminetetraacetic acid (EDTA) for 14 days and paraffin-embedded for histological analyses.

Ex vivo Radiographic analysis. High-resolution radiographic analysis was used to establish the extent of bone degradation in ipsilateral hindlimbs immediately following perfusion, with substantial bone degradation considered to be evidence of tumour invasion. X-ray scans of all mice were collected using a Faxitron MX-20 X-ray system (Faxitron X-ray Co., Wheeling, IL) on Kodak MIN-R, 2000 Mammography film (Eastman Kodak, Rochester,
NY). Affected hindlimbs were scored on a scale of 0 to 3: (0) normal bone, no visible lesion; (1) minor loss of bone density, minimal lesion; (2) Moderate to substantial loss of bone density, lesion limited to bone trabecula and cortex and (3) substantial loss of bone density, lesion includes clear periosteal involvement or fracture.

**Hematoxylin and eosin.** Ipsilateral hindlimbs were stained with hematoxylin and eosin to confirm the degree of infrafemoral tumour in animals that demonstrated lytic lesions on X-rays. Following decalcification, femurs and surrounding tissues were paraffin-embedded, sagittally sectioned at 5 μm, slide mounted, stained and xylene-cleared for microscopy.

**Immunofluorescence.** Immediately adjacent serial hindlimb sections from the same paraffin block were processed for immunofluorescence; following antigen retrieval in EDTA (pH 8, 95°C) for 30 min, sections were incubated in respective primary antibodies (xCT: Novus, 1:1000; cytokeratin 7: Santa Cruz, 1:500) overnight at 4°C, washed in PBS, incubated in 1:500 fluorescent secondary antibodies (Life Technologies AlexaFluor-488 goat anti-rabbit and AlexaFluor-647 goat anti-mouse, respectively) for 2 h at RT, counterstained with DAPI and coverslipped.

**Statistical analyses**

**In vitro.** For each cell line, a one-way analysis of variance (ANOVA) with Bonferroni post hoc comparisons was used to assess the dose response of AG879 on cell number. A two-way ANOVA with Bonferroni post hoc comparisons was then used to assess the effect of serial dilutions of β-NGF for each cell line, in the presence and absence of 6 μM AG879, on cell number and metabolism. Separate two-way ANOVAs with Bonferroni post hoc comparisons were used to assess the effect of β-NGF and AG879 on xCT protein levels and mRNA levels for each cell line. One-way ANOVAs with Bonferroni post hoc comparisons were used to assess the dose effect of β-NGF on functional system \(x_C^-\) activity, as measured by cystine uptake and glutamate release. Independent \(t\) tests were then used to assess the effect of AG879 on cystine uptake relative to cells treated with each β-NGF dose.

**In vivo.** DWB and DPA data sets were analyzed to derive survival curves comparing the time until respective behavioural scores fell irreversibly below 50% of the animals’ baseline nociceptive responses. For Kaplan–Meier curve analyses of 50% threshold pain data (time to behavioural decline) in DWB and DPA behavioural tests, chi-square analyses were used.

**Ex vivo.** Chi-square analysis of the proportion of scores within each treatment group was used to assess radiographic analysis of osteolysis. Immunocytochemical staining of xCT in 4T1 cells and immunofluorescent staining of xCT within ipsilateral hindlimbs were qualitatively considered.

All analyses were performed using GraphPad Prism 5.0a software (GraphPad Software, Inc., La Jolla, CA) and GraphPad QuickCals; \(\alpha\) was set at 0.05. Data are expressed as means ± standard error of the mean from three independent experiments, unless otherwise stated.

### Results

**In vitro**

**AG879 influences breast cancer cell number.** Selective TrkA inhibition appeared to concentration-dependently decrease cell number in 4T1 and MDA-MB-231 triple-negative cells, such that higher doses of AG879 (i.e., 12.5, 25 and 50 μM AG879) decreased cell number relative to DMSO-treated controls after 24 h treatment (see Table 2). Treatment with 6 μM AG879 did not significantly alter 4T1 or MDA-MB-231 cell growth or metabolism as measured by Crystal Violet and MTT assays. Thus, this dose was selected for all further in vitro studies.

**Selected AG879 and NGF doses do not affect 4T1 cell metabolism.** Treatment with β-NGF (0.01–100 ng) or AG879 (6 μM) for 24 h did not significantly affect breast cancer cell number (data not shown) or metabolism (Figure 2).

| Cell Line  | 3 μM  | 6 μM  | 12 μM | 25 μM | 50 μM |
|------------|-------|-------|-------|-------|-------|
| 4T1        | 1.08 ± 0.04 | 0.87 ± 0.07 | 0.8 ± 0.04* | 0.66 ± 0.07** | 0.35 ± 0.02** |
| MDA-MB-231 | 0.75 ± 0.06* | 0.85 ± 0.06 | 0.65 ± 0.11* | 0.55 ± 0.05* | 0.39 ± 0.05* |

Table 2. Values compared to DMSO control

Note: Murine 4T1 and human MDA-MB-231 carcinoma cells were seeded and treated with serial doses of AG879 for 24 h, and cell number was quantified using crystal violet staining. Values indicate mean ± SEM fold change in cell number compared to DMSO negative control (0 μM AG879), data are from three independent experiments.

Values compared to DMSO control, *p < 0.05, **p < 0.005.
AG879 attenuates β-NGF-induced increases in xCT protein in 4T1 cells. Western blotting analysis confirmed the presence of TrkA on 4T1 and MDA-MB-231 cells (data not shown). Blots also demonstrated a marked increase in xCT bands at 37 kDa in cells treated with 100 ng β-NGF in both 4T1 and MDA-MB-231 cell lines, an effect which was attenuated by AG879 in 4T1 murine carcinoma cells, with a similar, albeit modest, trend observed in MDA-MB-231 cells (Figure 3).

AG879 attenuates β-NGF-induced increases in xCT mRNA in 4T1 cells. qPCR analysis demonstrated a marked increase in xCT mRNA in 4T1 cells treated with 100 ng β-NGF, an effect which was attenuated by treatment with 6 μM AG879 (Figure 4).

AG879 decreases functional system xC⁻ activity. In 4T1 cells, ¹⁴C-cystine uptake was markedly reduced in cells treated with 6 μM AG879 for 24 h relative to DMSO- and 10 ng β-NGF-treated cells. Treatment with β-NGF appeared to marginally increase ¹⁴C-cystine uptake relative to DMSO control; however, this effect was not statistically significant (Figure 5(a) and (b)). Accordingly, glutamate release was increased in cells treated with 10 ng and 100 ng β-NGF (Figure 5(c)). Furthermore, treatment with 6 μM AG879 appeared to decrease this effect, although this trend was not statistically significant (Figure 5(c)). A similar trend was observed in triple-negative human MDA-MB-231 cells (data not shown).
**In vivo**

AG879 attenuates tumour-induced nociceptive behaviours. The TrkA inhibitor AG879 appeared to delay the onset of nociception, as measured by progressively impaired limb use and increased spontaneous guarding scores (Figure 6 (a) and (b), respectively). In both the DWB (Figure 7(a)) and the DPA (Figure 7(b)), tumour-bearing mice exhibited significantly greater pain behaviours compared to their sham controls from Day 9 post-tumour cell inoculation until end point. AG879 significantly delayed behavioural decline, as measured by postural disequilibrium in the DWB, indicating that significantly fewer tumour-bearing animals fell below 50% of their individual baseline nociceptive score when treated with AG879 (Figure 7(a)). In the DPA, a similar trend was observed, with fewer AG879-treated animals falling below 50% of their individual baseline scores, but this effect was not statistically significant (Figure 7(b)).

**Ex vivo**

AG879 impedes tumour invasion in vivo. Radiographic analysis confirmed the presence of bone degradation in cancer cell-inoculated mice at end point and AG879 appeared to partially attenuate tumour-induced osteolysis (Figure 8(a)). Hematoxylin and eosin staining of ipsilateral hindlimbs confirmed the presence of tumour cells in femurs of animals that demonstrated lytic lesions in radiographic X-rays; tumour invasion was most frequently observed in the distal epiphysis and diaphysis of ipsilateral femurs. AG879-treated mice demonstrated marginally less osteolytic activity, with generally smaller tumours observed in drug-treated animals (see Figure 8(b) and (c) for representative images), in accordance with radiographic lesion scores.

AG879 inhibits xCT expression in vivo. Immunocytochemistry confirmed the presence of xCT on 4T1 murine carcinoma cells in vitro (Figure 9(a)), and immunofluorescent staining of ipsilateral femurs demonstrated the presence of the epithelial marker cytokeratin 7 (Figure 9(b)), confirming the presence of breast cancer cells within the bone environment at end point. Double-label immunofluorescence demonstrated the

![Figure 4](image_url) AG879 attenuates β-NGF-induced increase in xCT mRNA in carcinoma cells. xCT mRNA in murine 4T1 carcinoma cells treated with DMSO and β-NGF, relative to negative control. Data are from three independent experiments, normalized to SDHA and are expressed as means ± SEM, α = 0.05. NGF: nerve growth factor; SDHA: Succinate Dehydrogenase Complex Flavoprotein Subunit A.

![Figure 5](image_url) Functional system xC⁻/C₀ activity is increased by β-NGF and attenuated by TrkA inhibition in murine 4T1 carcinoma cells. Treatment with β-NGF for 24 h demonstrated a dose-dependent trend of increased ¹⁴C-cystine uptake in 4T1 cells relative to DMSO control, although this effect was not statistically significant (a). 4T1 cells treated with 6 µM AG879 for 24 h demonstrated a relative decrease in ¹⁴C-cystine uptake relative to corresponding doses of β-NGF (b). Accordingly, a similar trend was observed in media levels of glutamate: treatment with β-NGF dose-dependently increased glutamate release, such that treatment with 10 ng and 100 ng β-NGF significantly increased media glutamate by 1.8- and 2-fold, respectively, relative to DMSO control (c). 4T1 cells treated with 6 µM AG879 demonstrated decreased glutamate release when matched to NGF doses, although the trend was not statistically significant (d). Data are from three independent experiments, normalized to protein level and are expressed as means ± SEM relative to DMSO, *significantly different from DMSO control, α = 0.05. DMSO: dimethyl sulfoxide; NGF: nerve growth factor.
overlapping distribution of xCT with breast cancer cells in ipsilateral femurs (Figure 9(c)), confirming the expression of xCT on 4T1 cells in vivo. AG879 appeared to markedly inhibit xCT expression by tumour cells (Figure 9(d)).

**Discussion**

Cancer pain is highly heterogeneous and is thought to be the result of varied pathological mechanisms. Current treatment options often induce severe dose-dependent side effects and fail to adequately manage advanced breast CIBP. Clinically, achieving analgesia often comes at the expense of patients' quality of life. The complexity of CIBP is partly due to the unique properties of the affected tissue; osteoclastogenesis, inflammation, increased neurotrophic activity, demyelination, and extracellular acidification created by the presence of bone tumours have been implicated in the sensitization of surrounding nociceptors and may

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**Figure 6.** AG879 significantly delays the onset and severity of spontaneous nociceptive behaviours. Open-field tests, including time spent spontaneously guarding the affected limb and scaled limb use, were used to visually assess ongoing and ambulatory nociception. (a) Quantification of limb use in the ipsilateral femurs throughout tumour progression. Normal hindlimb use during spontaneous ambulation was observed and scored on a scale of 4 to 0; Points indicate mean ± SEM of each stage of bone destruction as quantified by this scale. Limb use numbers represent: (4) normal use, (3) pronounced limp, (2) limp and guarding behaviour, (1) partial non-use of the limb in locomotor activity and (0) complete lack of limb use. (b) Time spent spontaneously guarding the hindpaw represented ongoing pain and was recorded during an open-field observation period. Guarding time was defined as the time (s) the hindpaw was held aloft while ambulatory. Points indicate means ± SEM; *significantly different from SHAM, significantly different from TUMOUR, α = 0.05; Vertical dashed lines indicate infratendinous injections (Day 0) and start of treatment (Day 7).

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**Figure 7.** AG879 delays time until behavioural decline. (a) The DWB quantitates the weight borne by the affected limb as a percentage of total recorded weight, standardized to baseline scores. Thus, 100% on the y-axis represents behaviour prior to tumour inoculation; less than 100% indicates a decrease in weight borne by the tumour-afflicted limb and is indicative of nociceptive behaviour. Kaplan–Meier curve analysis of the time until behavioural decline, expressed as a percentage of animals above the 50% of baseline cutoff, shows AG879 significantly delays time until the onset of postural disequilibrium in the DWB (chi square = 0.018, p = 0.724). (b) The DPA test quantitates the threshold force at which the animal withdraws from a progressive stimulus applied to the plantar surface of the affected hindpaw. Thus, 100% on the y-axis is equivalent to baseline behaviour prior to tumour inoculation; less than 100% indicates a decrease in the force withstood by the tumour-bearing limb. Kaplan–Meier curve analysis of the time until behavioural decline shows AG879 marginally, but not significantly, reduced time until decline in the DPA (chi square = 4.465, p = 0.1). Vertical dashed lines indicate start of treatment (Day 7). DWB: dynamic weight bearing; DPA: dynamic plantar aesthesiometer.
cumulatively contribute to cancer-induced hyperalgesia. Among other influences, it is clear that glutamate dysregulation and aberrant neurotrophic activity may contribute to CIBP through pathological osteoclastogenesis and sensitization. The production and vesicular secretion of NGF drives the autocrine stimulation and proliferation of breast cancer cells through its cognate receptor, TrkA. In the present study, we have demonstrated that inhibiting the NGF-TrkA complex may limit CIBP, conceivably through downstream effects on system xC\(^{-}\). Extensive research supports the hypothesis that NGF drives cancer pain through pathological nerve sprouting; this study provides evidence that the effects of NGF on breast cancer cells and surrounding cells in the tumour microenvironment may not be limited to its conventionally accepted neurotrophic actions. The discovery that NGF and system xC\(^{-}\) may be functionally interconnected provides a novel approach to indirectly targeting system xC\(^{-}\} for CIBP relief.

In vitro, AG879 dose-dependently reduced breast cancer cell growth, in accord with previous findings. More importantly, however, a dose which did not effectively alter cell number or metabolism significantly inhibited the functional activity of system xC\(^{-}\}. That is, AG879 markedly reduced cystine uptake in 4T1 carcinoma cells in the presence and absence of β-NGF. One possible explanation for the relationship between NGF and system xC\(^{-}\} is through changes that occur at the level of xCT by activation of the mitogen-activated protein kinase (MAPK)/extracellular signal-regulated kinase (ERK) signalling pathway. Through TrkA, NGF is known to activate Ras-MAPK signalling (reviewed in Huang and Reichardt). Interestingly, serine kinases, including MAPK, phosphorylate signal transducer and activator of transcription 3 (STAT3), which then homodimerizes and translocates into the nucleus where it then binds specific DNA recognition elements to regulate expression of genes, including xCT. In MDA-MB-231 breast cancer cells, for example, the MAPK inhibitor PD098059 abolishes basal xCT transcriptional activity. Furthermore, STAT3 activation is coupled with increased system xC\(^{-}\} activity. Accordingly, sustained treatment with the STAT3 inhibitor SH-4-54 decreases xCT expression and system xC\(^{-}\} activity in MDA-MB-231 cells. It is therefore plausible that phosphorylated TrkA contributes to xCT expression through MAPK and/or STAT3 signalling, although further research is required to explore the specific pathways involved in this relationship.

A consistent trend was observed across both cell lines assayed in vitro; AG879 influenced functional system xC\(^{-}\} activity in murine 4T1 and human MDA-MB-231 carcinoma cells. Thus, to mitigate the number of animals employed and avoid the complexity of using human cancer cells in an immunocompromised animal model, a syngeneic mouse model was utilized to characterize the effects of TrkA on cancer progression and CIBP in vivo. The battery of tests for nociceptive behaviours was cumulatively selected to reflect what is observed in patients who protect or suspend their afflicted limb.

Inhibition of TrkA delayed the onset of nociceptive behaviours in tumour-bearing mice, as demonstrated by spontaneous guarding and limb use scores, as well as time until behavioural decline in the DWB. No significant reduction in evoked pain was observed in the DPA, suggesting that either NGF-TrkA signalling or functional system xC\(^{-}\} activity may be more implicated in spontaneous than evoked pain. Previous work has shown that early but not late administration of an NGF sequestering antibody can prevent the onset of tumour-induced nerve sprouting in a mouse model of CIBP. Here, pharmacological TrkA inhibition began on Day 7 after intrafemoral cancer cell inoculation, corresponding with initial behavioural indications of CIBP. This time point
Figure 9. AG879 inhibits xCT expression in 4T1 murine carcinoma cells. Immunocytochemical staining of xCT in murine 4T1 mammary carcinoma cells in vitro at 60× magnification (a). Immediately adjacent serial sections to hematoxylin and eosin-stained femurs were stained for nuclei (blue), cytokeratin 7 (CK7) and xCT (green) and imaged using EVOS FL Cell Imaging System at 10× dry (b) and 60× oil (c–e) immersion lenses: double-label immunofluorescent staining of epithelial marker CK7 and xCT in representative tumour-inoculated animals (b), vehicle- (c) and AG879-treated (d) tumour-bearing femurs and sham control (e).
represents a relative late administration regimen to mimic what is seen clinically, as patients are often not diagnosed until metastases have progressed to an advanced, often painful, state. The ability of AG879 to partially mitigate, but not abolish, nociceptive behaviours when administered relatively late, with the onset of pain, suggests that irreversible damage to the bone microenvironment may have already occurred, but that this inhibitor nevertheless had subsequent effects on additional targets implicated in nociception.

The cystine-coupled export of glutamate through system $x_C^-$ bears additional significance within the nervous system, as it represents a non-vesicular route by which this excitatory neurotransmitter can activate glutamate receptors on sympathetic nerve fibres, particularly those present in extrasynaptic locales. While glutamate is recognized as a ubiquitous cell signalling molecule and critical intercellular neurotransmitter, excessive glutamate efflux produces an excitotoxic environment. One means by which glutamate may directly contribute to nociception is by acting on adjacent afferent nociceptive neurons within the bone. Another means by which glutamate evokes nociception is by influencing many aspects of metabolic homeostasis during normal bone remodelling, affecting the communication between osteoblasts and osteoclasts, which maintain a delicate balance of bone formation and resorption, respectively. The disruption of glutamatergic signalling within the bone microenvironment influences cell differentiation and function. In the presence of bone metastases, the balance between bone formation and degradation is disturbed. Disruption of this homeostasis by the aberrant increase in glutamate released from breast cancer cells into the bone microenvironment leads to excessive osteoclastogenesis and reduced bone mineralization. Given that glutamate released by system $x_C^-$ can directly stimulate adjacent afferent nociceptors, disrupt normal bone remodelling and inhibit bone mineralization, the observed delay until the time of onset of nociception is a logical consequence of inhibiting system $x_C^-$. Accordingly, the reduced osteolysis seen in AG879-treated animals, relative to untreated tumour-bearing animals, is reflective of the observed delay in nociceptive behaviours and reduction in functional system $x_C^-$ activity seen in the in vivo and in vitro data, respectively and cumulatively suggests that TrkA may be implicated in the osteolytic contribution to cancer pain.

Our group has reported on the antinociceptive effects of the steric system $x_C^-$ inhibitor, sulfasalazine (SSZ) in a validated model of CIBP in which SSZ delayed the onset of tumour-induced nociception, an outcome which has since been validated in a syngeneic mouse model. However, despite its efficacy in animal models of CIBP, SSZ has limited translational capacity given the poor bioavailability of the parent drug relative to its metabolites. Furthermore, side effects of SSZ mirror many of the symptoms already observed in cancer patients, including nausea, vomiting and anorexia, symptoms which ought to be controlled, not exacerbated. Therefore, although system $x_C^-$ is a promising pharmacological target for the treatment of CIBP, novel inhibitors are needed to better manage this pain without the negative side effects that accompany SSZ.

The present study corroborates the role of aberrant glutamate secretion through system $x_C^-$ in breast CIBP and suggests the possibility of a feedback loop between NGF and system $x_C^-$. The role of glutamate in cancer cell metabolism and the effects of NGF on system $x_C^-$-mediated glutamate output suggest that this connection may be vital to the progression of cancer pain. The relationship between TrkA and system $x_C^-$ is an important link in elucidating breast cancer-induced pain signalling. The present study supports the pharmacological inhibition of glutamate release to limit nociception and connects increases in glutamate release to previously unrelated increases in NGF. This study suggests that the actions of NGF extend beyond its conventionally recognized roles of neuronal survival and differentiation. Indirectly inhibiting system $x_C^-$ through the pharmacological modulation of TrkA therefore presents a valuable target for therapeutic intervention in the treatment of cancer pain.

Acknowledgements
The authors wish to thank Dr Mark Inman for his expert advice in statistical analyses.

Declaration of Conflicting Interests
The author(s) declared no potential conflicts of interest with respect to the research, authorship, and/or publication of this article.

Funding
The author(s) disclosed receipt of the following financial support for the research, authorship, and/or publication of this article: This work was supported by the Canadian Breast Cancer Foundation to GS.

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