Ghrelin prevents tumour- and cisplatin-induced muscle wasting: characterization of multiple mechanisms involved

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Abstract

Background  Cachexia and muscle atrophy are common consequences of cancer and chemotherapy administration. The novel hormone ghrelin has been proposed as a treatment for this condition. Increases in food intake and direct effects on muscle proteolysis and protein synthesis are likely to mediate these effects, but the pathways leading to these events are not well understood.

Methods  We characterized molecular pathways involved in muscle atrophy induced by Lewis lung carcinoma (LLC) tumour implantation in c57/bl6 adult male mice and by administration of the chemotherapeutic agent cisplatin in mice and in C2C12 myotubes. The effects of exogenous ghrelin administration and its mechanisms of action were examined in these settings.

Results  Tumour implantation and cisplatin induced muscle atrophy by activating pro-inflammatory cytokines, p38-C/EBP-β, and myostatin, and by down-regulating Akt, myoD, and myogenin, leading to activation of ubiquitin-proteasome-mediated proteolysis and muscle weakness. Tumour implantation also increased mortality. In vitro, cisplatin up-regulated myostatin and atrogin-1 by activating C/EBP-β and FoxO1/3. Ghrelin prevented these changes in vivo and in vitro, significantly increasing muscle mass (P < 0.05 for LLC and P < 0.01 for cisplatin models) and grip strength (P = 0.038 for LLC and P = 0.001 for cisplatin models) and improving survival (P = 0.021 for LLC model).

Conclusion  Ghrelin prevents muscle atrophy by down-regulating inflammation, p38/C/EBP-β/myostatin, and activating Akt, myogenin, and myoD. These changes appear, at least in part, to target muscle cells directly. Ghrelin administration in this setting is associated with improved muscle strength and survival.

Keywords  Cachexia; Cancer; Muscle; Ghrelin; Growth hormone

Introduction

Cachexia (involuntary loss of muscle and fat mass) is a common complication of many chronic conditions including cancer, chronic heart failure, lung disease, and renal disease. In the setting of cancer, it is associated with poor tolerance to therapy, decreased quality of life, and increased mortality. In spite of its significance, there are currently no treatments for this condition. Nutritional supplements alone do not reverse cachexia, resistance training may improve muscle strength and lean body mass, but it is not commonly prescribed to patients in this setting, and medications such as progesterational agents are often used but have been shown to increase only fat mass and can cause deep vein thrombosis, adrenal insufficiency, and hypogonadism. An increase in muscle proteolysis and a decrease in protein synthesis driven...
by activation of the ubiquitin-proteasome, mitogen-activated protein (MAP) kinases, and myostatin pathways among others have been identified as central processes in tumour-induced cachexia, but the pathways leading to these have not been fully characterized.

Cisplatin is a chemotherapeutic agent that is part of the standard of care for the treatment of several cancers including lung, head and neck, ovary, testicular, and bladder cancer, and paradoxically, it is known to induce weight and muscle mass losses. We have recently reported the mechanisms by which cisplatin alters lipid metabolism thereby causing fat atrophy, including an increase in lipolysis and lipid oxidation and a decrease in lipogenesis. However, the extent to which muscle protein synthesis and degradation are affected in this setting and the mechanisms mediating these effects have not been characterized. Given that cachexia is often a dose-limiting side effect of cisplatin and other chemotherapeutic agents, treating or preventing cachexia would allow the use of higher, more effective doses of cisplatin.

Ghrelin, the endogenous ligand for the growth hormone secretagogue receptor (GHSR)-1a, has been proposed as a potential therapeutic target for cachexia, having the potential to increase appetite, fat, and muscle mass. Although its mechanisms of action have not been fully elucidated, an increase in food intake and decrease in energy expenditure via hypothalamic effects, a decrease in inflammation, an increase in GH, and direct effects in adipose tissue and skeletal muscle are thought to potentially mediate these effects.

In this study, we established the effects of ghrelin in the setting of chemotherapy- and tumour-induced muscle wasting and characterized the pathways involved. We studied these two models to determine the extent to which these mechanisms were differentially modulated, looking at gene and protein expression as well as their functional impact and their effect on survival. Because multiple pathways are likely to play a role in mediating ghrelin’s effects, we postulated that several targets would be modulated by ghrelin in these settings.

Materials and methods

Animals

Adult (age 90 ± 10 days) c57bl/6J male mice were used for all experiments. Animals were individually housed, acclimated to their cages and human handling for 1 week before the experiments, and maintained on a 12/12 light/dark cycle (lights on at 6 a.m.). All experiments were conducted with the approval of the Institutional Animal Care and Use Committee and were in compliance with the National Institutes of Health Guidelines for Use and Care of Laboratory Animals. Sample sizes of 8–10 animals/group were used for all experiments.

Acylated ghrelin was used for all in vivo and all in vitro studies. For the cisplatin-induced cachexia model, animals received vehicle (saline), cisplatin, acylated ghrelin + cisplatin, or acylated ghrelin. Cisplatin was purchased from APP Pharmaceuticals (Schaumburg, IL), and rodent acylated ghrelin was synthesized by Anaspec (Fremont, CA). The dose of cisplatin was 2.5 mg/kg given daily for 10 days at 8:30 a.m. intraperitoneally (IP), and the dose for ghrelin was 0.8 mg/kg twice daily IP at 8 a.m. and 5 p.m. for 21 days. The morning dose of acylated ghrelin was given 30 min before cisplatin. This particular dose of cisplatin and acylated ghrelin was selected based on our previous work that showed cisplatin-induced weight loss and anorexia without over toxicity, and prevention of anorexia, fat atrophy, and weight loss by ghrelin.

For the Lewis lung carcinoma (LLC)-induced cancer cachexia model, 100 μl (1 × 10^6 cells) of LLC cells (American Type Culture Collection, Manassas, VA) or an equal volume and number of heat-killed LLC cells (control) was injected subcutaneously into the right flank. Ghrelin (tumour + ghrelin, T-G group) or vehicle (saline, tumour + vehicle, T-V group) was injected subcutaneously into the left flank of mice from Day 7 after LLC implantation, when the tumour became palpable (approximately 1 cm in diameter). The dose for ghrelin was 0.8 mg/kg twice daily at 8 a.m. and 5 p.m. Mice were sacrificed on Day 28 of LLC implantation, approximately 21 days after tumour had become palpable.

Body weight and food intake were assessed daily by weighing the food and the animals before the a.m. injection. Body weight changes were expressed as change from baseline, and food intake was expressed in grammes/day. Lean body mass and fat mass were measured by nuclear magnetic resonance (NMR) with a Minispec mq NMR spectrometer (Bruker optics, The Woodlands, TX). Grip strength was measured with a grip strength metre with digital force gauge (Columbus Instruments, Columbus, OH).

Hind-leg muscles [quadriceps, gastrocnemius, soleus, tibialis anterioris (TA), and extensor digitalis longus] were collected for analyses. Comparisons between groups (i.e. muscle cross-sectional area and gene or protein expression) were always done from the same muscles. Muscle weights were normalized by lean body mass at baseline. For the LLC-induced cachexia model, body weight and lean body mass were expressed as weight after tumour weights were subtracted and expressed as change from baseline.

Cell culture

C2C12 (ATCC, Manassas, VA) myoblasts were maintained in Dulbecco’s modified Eagle medium (DMEM) (Life Technologies, Grand Island, NY) with 10% foetal bovine serum (HyClone, Logan, UT), penicillin (200 units/mL), and streptomycin (50 μg/mL) (Invitrogen, Carlsbad, CA). At 90% confluence, the media was changed to DMEM plus 2% horse

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serum (Life Technologies, Grand Island, NY) to induce myotube formation. Cells were treated with cisplatin (50 μM), acylated ghrelin (1 μM) or both on Day 3 when myotube formed and were harvested 24 h after.

Real-time quantitative polymerase chain reaction

RNA from gastrocnemii or cultured C2C12 myotubes was isolated (Trizol reagent, Life Technologies, Grand Island, NY). Transcript levels were measured by real-time PCR (7000 Sequence Detection System; Applied Biosystems). Total RNA was reverse transcribed (QuantiTect Reverse Transcription Kit, Qiagen, Valencia, CA) to cDNA. Primers for real-time PCR amplification used are from Sigma-Aldrich (St. Louis, MO; see Supporting Information, Table S1). Signals are expressed relative to glyceraldehyde-3-phosphate dehydrogenase using the standard 2ΔΔCT method.

Western blot

Protein from gastrocnemii or myotubes was isolated using tissue lysis/extraction reagent or cell lysis reagent (Sigma-Aldrich, St. Louis, MO; see Supporting Information). Nuclear protein was isolated using EpiQuik Nuclear Extraction Kit (Epigentek Farmingdale, NY). After Phosphate Buffered Saline (PBS) washing, cells were scraped and centrifuged. Cell pellets were re-suspended in 100 μL of diluted NE1 [1X, 1:1000 ratio dithiothreitol (DTT) solution and Protease Inhibitor Cocktail (PIC)] and incubated on ice for 10 min, then vortexed and centrifuged for 1 min at 24000 g. The cytoplasmic extract was removed from the nuclear pellet, and two volumes of PBS were added to the supernatant. The cytoplasmic extract was harvested 24 h after.

Protein synthesis and degradation

Protein synthesis was measured from the incorporation of L-[3,5-3H]tyrosine (5 μCi/mL; PerkinElmer, Waltham, MA) into cellular proteins during a 24 h incubation as described elsewhere.7,8 Myotubes were then washed three times with ice-cold PBS before adding 10% trichloroacetic acid (TCA) to precipitate proteins. After three additional PBS washings, the precipitate was rinsed twice with TCA and solubilized by sonication in lysis solution (1% Triton X-100 and 1 N NaOH). The incorporation of radiolabeled tyrosine radioactivity was measured using liquid scintillation (Beckman Coulter, Brea, CA), and protein concentration was measured using Pierce BCA Protein Assay. L-[3,5-3H]tyrosine (5 μCi/mL; PerkinElmer, USA) was added to C2C12 myotubes and incubated 24 h. After rinsing by fresh 2% horse serum DMEM, the myotubes were treated with vehicle (PBS), cisplatin (50 μM), and cisplatin (50 μM) + ghrelin (1 μM). Aliquots (200 μL) of culture media were taken at specified times for quantization of L-[3,5-3H]tyrosine release. Proteins were precipitated at 4°C with 10% TCA and centrifuged at 37000 g for 5 min. The precipitate was rinsed twice with TCA and solubilized by sonication in lysis solution (1% Triton X-100 and 1 N NaOH). Radioactivity in the TCA-soluble supernatant and the proteins (TCA-insoluble fraction) were measured using liquid scintillation counting. At the end of the chase period, cells were rinsed twice in PBS and precipitated at 4°C in 10% TCA, and the radioactivity in cell protein was measured as described above. Total radioactivity is the sum of the residual radioactivity in cell proteins and the TCA-soluble radioactivity at different time points. Protein degradation was expressed as L-[3,5-3H]tyrosine released as a percentage of total L-[3,5-3H]tyrosine incorporated.

Immunofluorescence staining

Frozen TA muscles sections or C2C12 myotubes were fixed with 4% paraformaldehyde. After three Tris Buffered Saline Tween-20 (TBST) washings, slides were blocked for 1 h (Dako North America, Inc., Carpentaria, CA). The primary antibody (1:100) dilution was incubated overnight at 4°C. Alexa Fluor 488 or 546 secondary antibodies (Life technologies, Grand Island, NY) were used with 1:1000 dilutions and incubated 1 h at RT. 4’6-Diamidino-2-phenylindole (Cell Signaling) was used to stain nuclei.
Ghrelin prevents cancer-related wasting

Results

Effect of ghrelin on body weight, muscle mass, and myofiber size changes caused by LLC tumour or cisplatin

Both LLC and cisplatin induced a significant decrease in body weight compared with control animals, whereas ghrelin administration prevented these changes (Figure 1A and B). Lean body mass was also decreased by LLC, and cisplatin and ghrelin also prevented these changes (for the LLC model, change from baseline for heat-killed + vehicle [HK + V] 95.70 ± 4.30%, tumour + vehicle [T + V] 83.80 ± 1.52%, and tumour + ghrelin [T + G] 96.47 ± 2.04%, P < 0.05; for the cisplatin model, change from baseline for vehicle [V] 100.39 ± 0.41%, cisplatin [C] 83.41 ± 0.78%, cisplatin + ghrelin [C + G] 93.39 ± 0.61%, and ghrelin [G] 110.20 ± 0.92%, P < 0.01) All muscles in the hind leg showed significant atrophy, and this was also prevented by ghrelin (Figure 1C and D). Paralleling the changes in muscle mass, grip strength was significantly decreased by tumour implantation and cisplatin, and these changes were also prevented by ghrelin, highlighting the functional impact of these interventions (Figure 1E and F). Cachexia in these two models was associated with a decrease in daily food intake that was also prevented by ghrelin (for the LLC model, HK + V 3.55 ± 0.13 g/day, T + V 2.74 ± 0.16 g/day, and T + G 3.23 ± 0.08 g/day, P < 0.05; for the cisplatin model, V 3.67 ± 0.05 g/day, C 2.55 ± 0.09 g/day, C + G 3.12 ± 0.05 g/day, and G 4.05 ± 0.06 g/day, P < 0.01), suggesting that one of the mechanisms mediating ghrelin’s effects is by decreasing anorexia. LLC implantation and cisplatin administration induced a significant decrease in myocyte cross-sectional area in TA muscles, and this was prevented by ghrelin (see Supporting Information, Figure S1A–D, for LLC model, P < 0.05; for the cisplatin model, P < 0.05). LLC implantation and cisplatin also decreased fat mass significantly as measured by NMR. These changes were prevented by ghrelin (for the LLC model change from baseline for HK + V 85.25 ± 0.75%, T + V 32.80 ± 5.13%, and T + G 57.53 ± 5.01%, P < 0.01; for the cisplatin model change from baseline for V 101.70 ± 1.34%, C 60.92 ± 2.51%, C + G 85.75 ± 0.93%, and G 116.66 ± 1.45%, P < 0.01).

Proteolysis and proteasome activity

Tumour implantation and cisplatin administration were associated with an increase in the expression of the ubiquitin ligases MAFbx/Atrogin-1 and muscle ring finger-1 (MuRF-1) (Figure 2A–D); whereas, the markers of muscle differentiation MyoD and myogenin were decreased by LLC and cisplatin. These changes were prevented by ghrelin suggesting that ghrelin-decreased proteolysis and increased muscle mass are mediated at least in part through these pathways.
Also, to confirm the relevance of the changes in atrogin-1/ MuRF-1 seen in these models, we measured the proteasome activity directly. As shown in Figure 2E and F, cisplatin and LLC tumour implantation increased proteasome activation, and this was also prevented by ghrelin. The transcription of Atrogin-1 and MuRF-1 is increased in part by dephosphorylation of the transcriptional factor Fox-O1-3, which is in turn regulated by the central mediator Akt. LLC inoculation and cisplatin decreased the phosphorylation of Akt (Figure 2C and D), suggesting that this pathway is involved in LLC and cisplatin-induced muscle atrophy.

Myostatin was up-regulated by LLC inoculation or cisplatin, and these changes were also abolished by ghrelin. Also, the MAP kinase p38 is believed to mediate protein degradation in cancer cachexia by activating atrogin-1 directly and through up-regulation of myostatin. Phosphorylated p38 levels were increased by LLC inoculation or cisplatin administration, and this was prevented by ghrelin.

**Pro-inflammatory cytokines**

Inflammation plays a role in muscle wasting through activation of p38 and down-regulation of Akt among other mechanisms, and ghrelin has been proposed to have anti-inflammatory effects in non-cancer settings. Serum levels of the pro-inflammatory cytokines interleukin (IL)-6, tumour necrosis factor (TNF)-α, and IL-1β were significantly increased in tumour-bearing and cisplatin-treated animals, and this was prevented by ghrelin co-administration (see Supporting Information, Figure S2).

**Effect of ghrelin on myotube breakdown, protein synthesis, and proteolysis changes induced by cisplatin in vitro**

To further characterize the mechanisms mediating the effects of cisplatin and ghrelin in cachexia, C2C12 myotubes were
treated with vehicle, cisplatin, ghrelin, or ghrelin + cisplatin. Cisplatin induced a significant decrease in myotube size and myosin heavy chain content, and these changes were prevented by ghrelin (Figure 3A–D). Cisplatin increased the expression of atrogin-1, MuRF-1, p38, and myostatin and decreased the expression of Akt, myoD, and myogenin, and these changes were prevented by ghrelin (Figure 4). Direct measurements of protein synthesis and degradation in myotubes treated with cisplatin or cisplatin + ghrelin followed the same pattern confirming the relevance of these changes (protein synthesis after 24 h measured by L-[3,5-3H]tyrosine incorporation compared to control samples: cisplatin 53.00 ± 2.00%, C + G 75.50 ± 3.50%, ghrelin 114.00 ± 2.00%, P < 0.01; protein degradation after 24 h measured by L-[3,5-3H]tyrosine release compared to control: cisplatin 127.98 ± 2.14%, C + G 102.57 ± 2.86%, ghrelin 76.57 ± 4.52%, P < 0.01).

Nuclear C/EBP-β and FoxO1/3 were significantly increased by cisplatin and prevented by ghrelin in C2C12 myotubes (Figure 5A). We then confirmed the importance of these pathways on myostatin expression using a construct expressing luciferase alongside the FoxO1/3, SMAD 2/3, and C/EBP-β binding sites in the myostatin promoter. As shown in Figure 5B, cisplatin induced activation of the myostatin promoter, and this was prevented by ghrelin. We also used a different construct with intact or mutated FoxO1/3 and C/EBP-β binding sites in the atrogin-1 promoter to test the relative contribution of these two transcriptional factors to atrogin-1 activity. As shown in Figure 5C, FoxO1/3 and C/EBP-β both significantly contribute to the activation of the atrogin-1 promoter similarly.

**Survival**

Tumour mass was no different between animals treated with vehicle and ghrelin (tumour mass for T + V 5.87 ± 1.08 g, and for T + G 6.54 ± 1.89 g, P = 0.756). Survival was decreased in tumour-bearing animals, and this was significantly improved by ghrelin administration (Figure 6). The cisplatin regimen used was not lethal.

**Discussion**

Cachexia is a devastating complication of cancer that contributes to a decrease in quality of life and early demise in individuals with this condition. Paradoxically, it is exacerbated by the administration of chemotherapeutic drugs such as cisplatin, and cachexia is often a dose-limiting side effect of these agents. Muscle atrophy contributes the most to a decrease in functionality in cancer patients, and it is associated with an increased risk of chemotherapy-induced toxicity and poor outcome. In spite of its significance, the mechanisms underlying cancer- or chemotherapy-induced muscle atrophy are incompletely understood.

The gastric hormone ghrelin causes weight gain by increasing food intake and by food intake-independent mechanisms and ghrelin or ghrelin receptor agonists have been proposed as potential therapies for cancer cachexia as they may improve anorexia, muscle mass and strength, and weight loss in patients with cancer, particularly those...
receiving cisplatin-based chemotherapy.\textsuperscript{27,11} However, their mechanisms of action in muscle remain to be fully elucidated.

In this study, tumour-bearing and cisplatin-treated animals developed marked muscle atrophy and weakness that were associated with activation of the ubiquitin-proteasome pathway. Phosphorylated Akt, which is known to down-regulate this pathway through phosphorylation of the transcriptional factor FoxO1-3,\textsuperscript{28} was decreased by tumour or cisplatin. These changes were prevented by ghrelin administration in both models suggesting that ghrelin’s effects are mediated through the ubiquitin-proteasome pathway in these settings. It is noteworthy that LLC cells from different suppliers behave somewhat differently in terms of their capacity in inducing muscle catabolism and activating related signalling pathways. For example, in this study, we used LLC cells from ATCC, which take 28 days to fully develop cachexia and induce MuRF1 up-regulation through Akt-mediated FoxO activation. However, LLC cells from National Cancer Institute (NCI) require only half of the time period to develop cachexia, yet, without activating FoxOs and up-regulating MuRF1.\textsuperscript{29} It is also important to notice that the relative contribution of this pathway to cancer cachexia in humans is not well understood with some\textsuperscript{30} but not all studies\textsuperscript{31} showing an increase in protein degradation through this pathway.

The inflammatory cytokine IL-6 is increased in the setting of cancer, and it may decrease Akt phosphorylation through activation of the suppressor of cytokine signalling-3.\textsuperscript{21} Other inflammatory cytokines such as IL-1β and TNF-α may play an important role in inducing cachexia by activating nuclear factor kappa-B and MAP kinases including p38.\textsuperscript{32,33} Ghrelin has been shown to down-regulate inflammation in the setting of LPS-induced sepsis and chronic kidney disease.\textsuperscript{13} Whether this is also true in the setting of cancer is not known. Here, we show that tumour implantation and cisplatin administration increased circulating cytokine levels, and these changes were prevented by ghrelin co-administration. These results suggest that the effects of ghrelin in this setting may be mediated through its anti-inflammatory effects, at least in part.

The MAP kinase p38 has been shown to play a role in the development of cachexia and to be activated by inflammation.\textsuperscript{32–36} However, its specific role is controversial with reports of p38 increasing in the setting of cancer cachexia\textsuperscript{29} or decreasing in the setting of dexamethasone-induced muscle wasting.\textsuperscript{16} Moreover, ghrelin was shown to decrease p38 activation in C2C12 cells.\textsuperscript{37} We found the phosphorylated levels of p38 to increase with tumour or cisplatin administration and that ghrelin prevented these changes. The increase in p38 activation we describe is consistent with previous

\textbf{Figure 3} Ghrelin improves cisplatin-induced myotubes breakdown. (A) Immunofluorescence staining for anti-myosin/myosin heavy chain (MHC) antibody in C2C12 myotubes. MHC staining outlines the myotubes (red). 4′-6-Diamidino-2-phenylindole was used to stain the nuclei (blue). (B) Myotubes size expressed as % from vehicle. (C–D) Western blot of MHC and glyceraldehyde-3-phosphate dehydrogenase (GAPDH) in C2C12 myotubes. V, vehicle-treated group; C, cisplatin-treated group; CG, cisplatin + ghrelin-treated group.
tumour models, and the down-regulation of this pathway with ghrelin suggests that its effects in the setting of muscle wasting may depend in part on the model used, down-regulating it in models where inflammation plays a role (i.e. cancer-induced cachexia) but not in other non-inflammatory models of wasting (i.e. dexamethasone-induced muscle atrophy).

Muscle wasting in cancer cachexia also has been associated recently with up-regulation of myostatin. Myostatin, a member of the Transforming growth factor beta (TGF-β) family made in muscle, is known to activate the activin receptor IIIB and to reduce Akt signalling.38,39 Moreover, targeting this pathway has been shown to ameliorate cachexia.17 In our study, myostatin was significantly up-regulated by tumour implantation and by cisplatin, and this was prevented by ghrelin administration. This is in contrast to a recent report of ghrelin not preventing the increase in myostatin induced by dexamethasone, a model where stimulation of p38 was not seen.

The myogenic regulatory factors MyoD and myogenin have been shown to play an important role in tumour-induced cachexia by regulating muscle regeneration and are regulated by Akt phosphorylation, p38, myostatin and TNF-α.41,42 Their expression levels decreased in both models suggesting that satellite cell proliferation and muscle cell differentiation and regeneration may be impaired. These changes were also prevented by ghrelin administration in vivo in both models. This points out to a potential new mechanism for ghrelin to ameliorate cachexia.43

We also developed a model of cisplatin toxicity in C2C12 cells to determine if the effects of cisplatin were due to a direct effect on muscle cells. Administration of cisplatin to C2C12 cells induced myotube atrophy, and this was associated with activation of p38, myostatin, and the ubiquitin ligases atrogin-1 and MuRF-1 along with down-regulation of Akt and the regenerative pathway (MyoD and myogenin) as we had demonstrated in vivo. This suggests that cisplatin exerts at least part of its effects directly on muscle and is in agreement with a previous report of cisplatin inducing myotube atrophy through Akt

Figure 4 Ghrelin prevents cisplatin-induced changes in protein synthesis and degradation in vitro. (A) mRNA levels for myoD, myogenin, atrogin-1/MAFbx, MuRF-1, and myostatin normalized to glyceraldehyde-3-phosphate dehydrogenase (GAPDH) and expressed as fold change from vehicle. (B) Western blot of cisplatin- and ghrelin-treated C2C12 myotubes (I). Atrogin-1/MAFbx (II), Myostatin (III), and MyoD (IV) are normalized to GAPDH, p-Akt (V) to total Akt and p-p38 (VI) to total p38. V, vehicle-treated group; C, cisplatin-treated group; CG, cisplatin + ghrelin-treated group.
down-regulation.\textsuperscript{44} The effects of cisplatin treatment on C2C12 cells also may represent disruption of the process of differentiation, and it has been suggested before that ghrelin may restore this process.\textsuperscript{37} Ghrelin increased food intake, and this is thought to mediate some but not all of its effects in this setting as we\textsuperscript{8} and others\textsuperscript{45} have recently shown. These in vitro studies suggest that ghrelin exerts its effects at least in part directly on muscle cells, independently of its orexigenic effects as suggested by these previous studies.

The only identified receptor for ghrelin to this date is the GHSR-1a, and this receptor is not expressed in muscle tissue or C2C12 cells.\textsuperscript{46,16} Nevertheless, recent reports suggest that
some of ghrelin’s effects are not mediated through this receptor.47,16 Our in vitro studies show that ghrelin may prevent atrophy induced by cisplatin in the absence of GHSR-1a. Very recently, the corticotrophin-releasing factor receptor 2 has been postulated to mediate some of ghrelin’s effect on glucose metabolism.48 Future studies should focus on this or other alternative pathways that could explain ghrelin’s effects in the absence of GHSR-1a.

Lastly, ghrelin administration lead to increased survival in tumour-bearing animals, indicating that the benefits of improving muscle mass go beyond simply an increase in functionality. This is also relevant given the potential concerns for an anabolic agent like ghrelin in inducing tumour growth and worsening outcomes. Although in vitro studies have given conflicting results with some showing an increase and some showing a decrease in cell proliferation with ghrelin administration,49,50 all in vivo models and human studies where ghrelin or ghrelin mimetics were used have not shown an increase in tumour proliferation, although none of these studies previously reported survival.51,52

In summary, we show here how multiple pathways interact and are involved in the development of cachexia, either induced by a tumour or, paradoxically, by the chemotherapeutic agent cisplatin. Activation of p38/C/EBP-β, myostatin, and inflammatory cytokines, and a decrease in Akt and Myogenin/myoD ultimately lead to increased proteolysis, decreased muscle mass, and strength. The novel hormone ghrelin prevents muscle atrophy by decreasing inflammation, by down-regulating the p38/C/EBP-β/myostatin pathway, and by increasing Akt phosphorylation and activating myogenin and myoD. The increase in muscle strength and survival induced by ghrelin in tumour-bearing animals highlights the clinical relevance of these findings.

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Supporting information

Supporting Information is available at Journal of Cachexia, Sarcopenia and Muscle online.

Figure S1. Ghrelin prevents the decrease in myofiber size caused by tumour or cisplatin. (A) Immunofluorescence staining for anti-laminin antibody of TA muscles in LLC-induced cachexia. Laminin staining (green) outlines myofibers. Dapi was used to stain the nuclei (blue). (B) Immunofluorescence staining for anti-laminin antibody of TA muscle in cisplatin-induced cachexia. (C) Average cross-sectional area of myofibers in LLC-induced cachexia. (D) Average cross-sectional area of myofibers in cisplatin-induced cachexia. HK-V represents the group injected with heat-killed (HK) LLC and vehicle. T-V and T-G group represent animals inoculated with LLC (10⁶ cells) receiving vehicle (saline) or ghrelin (0.8 mg/kg twice daily), respectively. V, vehicle-treated group, C, cisplatin-treated group; CG, cisplatin + ghrelin-treated group; G, ghrelin-treated group. *P < 0.05, **P < 0.01, compared to HK-V or vehicle group; §P < 0.05, §§P < 0.01 compared to T-V or C group.

Figure S2. Serum IL-6, TNF-α, and IL-1β levels in LLC-induced cachexia (A) and cisplatin-induced cachexia (B). Ghrelin prevents the increase in these pro-inflammatory cytokines induced.
by tumour- and cisplatin-induced cachexia. * P < 0.05 comparing to HK-V or vehicle group; $P < 0.05$ comparing to T-V or C group.

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**Conflicts of interests**

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