Characterization of ghrelin receptor activity in a rat pituitary cell line RC-4B/C

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Abstract

Ghrelin, a 28 amino acid, octanoylated peptide, is an endogenous ligand for the growth hormone secretagogue receptor (GHS-R). In addition to various endocrine functions, including stimulation of GH release, ghrelin has been characterized as an important regulator of energy homeostasis. Ghrelin administration has been shown to increase adiposity in rodents and stimulate food intake in humans. Studies suggest that these orexigenic effects are mediated primarily through GHS-R expression in hypothalamic and pituitary neuronal pathways. In this context, GHS-R has been recognized as a potential target for the treatment of GH deficiency and body weight disorders. Cell lines provide convenient in vitro systems to identify and characterize potential pharmacophores and to analyze GHS-R functional activity. While recombinant cell lines that overexpress GHS-R have served as effective research tools for these studies, such cell lines may differ in signaling response to ghrelin compared with hypothalamic or pituitary cells expressing GHS-R. We show here that a cell line derived from a rat anterior pituitary adenoma, RC-4B/C, expresses endogenous GHS-R as judged by reverse transcriptase-PCR. In a Ca\(^{2+}\) mobilization assay, RC-4B/C cells demonstrate a dose-dependent increase in intracellular [Ca\(^{2+}\)] on stimulation with rat ghrelin and a related peptide agonist, hexarelin (EC\(_{50}\), 1·0 nM and 1·7 nM respectively), but are unresponsive to treatment with inactive des-octanoyl rat ghrelin. A subclone, RC-4B/C.40, with a more robust and stable ghrelin response, was isolated from the parental population of cells to allow further analysis of GHS-R signal transduction. Using pertussis toxin and the phospholipase C inhibitor U-73122, we show that ghrelin signals through the Gq pathway in the RC-4B/C.40 cells. We also demonstrate that the ghrelin-induced rise of intracellular [Ca\(^{2+}\)] in RC-4B/C.40 cells involves initial Ca\(^{2+}\) release from intracellular stores followed by a sustained elevation that occurs via influx of extracellular Ca\(^{2+}\) through ion channels. In addition, unlike observations reported in recombinant cell systems, the RC-4B/C.40 cells do not exhibit a high level of GHS-R constitutive activity as determined in a phosphatidylinositol hydrolysis assay. Overall, the data presented here suggest that the RC-4B/C parental and RC-4B/C.40 cells provide novel in vitro systems for the characterization of GHS-R pharmacophores and ghrelin signaling.

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Introduction

The pulsatile secretion of growth hormone (GH) by the anterior pituitary is a complex process governed mainly by two hypothalamic hormones (Giustina & Veldhuis 1998), growth hormone-releasing hormone (GHRH) and its counter-regulatory hormone, somatostatin. In addition, there are synthetic, peptidyl, and non-peptidyl molecules termed GH secretagogues (GHSs) known to possess growth hormone-releasing activity (Smith et al. 1997). These molecules were first described 20 years ago in efforts to develop alternative treatments for GH deficiency. They were recently shown to act through the GH secretagogue receptor (GHS-R) (Howard et al. 1996, Smith et al. 1996). GHSs and GHRH have synergistic effects, indicating that they probably act via different mechanisms (Smith et al. 1997, Ghigo et al. 2001).

Recently, a natural ligand for GHS-R was identified. Ghrelin is a 28-amino acid, octanoylated peptide secreted primarily by the upper intestinal tract (Kojima et al. 1999). It is a potent dose-dependent GHS in rodents (Saito et al. 2003) and humans (Peino et al. 2000, Takaya et al. 2000). However, ghrelin function is not restricted to GH release, as it has been shown to be an important regulator of food intake and body weight (Horvath et al. 2003). Ghrelin plasma levels rise before meals and fall following feeding (Cummins et al. 2001). Ghrelin infusion in near-physiological doses increases hunger in humans (Wren et al. 2001) and adiposity in rodents (Tschöp et al. 2000). Consistent with high sequence homology to motilin, ghrelin stimulates gastric contractility and acid secretion (Masuda et al. 2000).

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Although ghrelin was isolated and purified from the oxyntic mucosa of the stomach (Kojima et al. 1999, Date et al. 2000), it is produced in other tissues, albeit at lower levels. These include not only the hypothalamus (Cowley et al. 2003) and pituitary (Korbonits et al. 2001), the sites of GHS-R expression, but also the pancreas, kidney, liver, heart, lung, ovary, prostate, and the placenta (Gnanapavan et al. 2002). The full-length ghrelin receptor, GHS-R1a, is expressed mainly in the GH-producing cells of the pituitary (Adams et al. 1998, Korbonits et al. 1998, Skinner et al. 1998, Korbonits et al. 2001). It is also found in the hypothalamus where it is thought to mediate the orexigenic (appetite stimulating) effects of ghrelin in addition to stimulating GHRH-neurons (Howard et al. 1996). Recently, GHS-R expression has also been identified in afferent vagal neurons. It was shown that ghrelin-induced increase of feeding in rats is dependent on an intact vagal nerve (Howard et al. 1997, Kojima et al. 1998). It is also found in the hypothalamus where it is characterized in a variety of cell systems (Smith et al. 2001). It is also found in the hypothalamus where it is thought to mediate the orexigenic (appetite stimulating) effects of ghrelin in addition to stimulating GHRH-neurons (Howard et al. 1996). Recently, GHS-R expression has also been identified in afferent vagal neurons. It was shown that ghrelin-induced increase of feeding in rats is dependent on an intact vagal nerve (Date et al. 2002). Therefore, ghrelin appears to be a physiological link between the stomach, the pituitary, and the hypothalamus.

GHS-R expression and activity have been well characterized in a variety of cell systems (Smith et al. 1997, Kojima et al. 1999). Activation of the receptor by ghrelin, peptidomimetics or small-molecule agonists is coupled via Gq-proteins to the phospholipase C pathway, leading to inositol trisphosphate (IP) and diacylglycerol (DAG) production. The elevation of IP produces a subsequent increase in cytosolic intracellular Ca2+ concentration ([Ca2+]i) from intracellular stores. In parallel, DAG stimulates protein kinase C (PKC), leading to activation of the mitogen-activated protein kinase (MAPK) pathway. The MAPK pathway has been identified as responsible for mitogenic and antiapoptotic effects of ghrelin (Kim et al. 2004, Nanzer et al. 2004). GHS-R activation also results in activation of L-type Ca2+ channels and influx of Ca2+ from the extracellular milieu (Smith et al. 1997, Glavaski-Joksimovic et al. 2003).

There is substantial evidence that GHS-R in such systems displays a high level of constitutive activity (Holst et al. 2003, 2004; Holst & Schwartz 2004). While treatment with inverse agonists such as substance P and related analogs has been shown to effectively suppress this activity, it remains to be determined whether the constitutive activity seen in recombinant GHS-R expression has physiological relevance in vivo.

As cell lines are much more convenient for receptor pharmacology studies than either primary cell preparations or direct in vivo studies, our aim was to identify and characterize the functional activity of endogenously expressed GHS-R in a physiologically relevant cell line. We reasoned that such a cell line might provide a useful intermediate between recombinant cell lines and primary cell preparations for elucidating GHS-R signaling. We identified RC-4B/C cells (Hurbain-Kosmath et al. 1990) derived from an anterior pituitary adenoma of a male rat as a suitable in vitro system for this purpose. We demonstrate here that RC-4B/C cells respond to ghrelin treatment with an increase in [Ca2+]i that is blocked by GHS-R antagonists. Additionally, we isolated a subclone of the parental RC-4B/C cells, RC-4B/C.40, which demonstrated a more robust and stable ghrelin response in a Ca2+ mobilization assay. As a more homogeneous cell population with enhanced GHS-R expression, the subclone allowed for further analysis of ghrelin signaling, including characterization of the ghrelin-induced Ca2+ pathways. It appears that the RC-4B/C.40 cells signal through Gq in a manner similar to Chinese hamster ovary (CHO)-K1 cells expressing human GHS-R (CHO-hGHS-R). However, when using a phosphatidylinositol (PI) hydrolysis assay, we did not observe the level of constitutively active GHS-R in the RC-4B/C.40 cells that occurs in the recombinant CHO line. This observation seems particularly interesting in that it may be the first reported example of a nonconstitutively active, functional GHS-R in a relevant pituitary cell line. Overall, our results indicate that RC-4B/C parental and RC-4B/C.40 cells are potentially useful in vitro systems for the characterization of GHS-R agonists and antagonists. Furthermore, these cell lines may provide a better model of ghrelin signaling in vivo.

Materials and methods

Cell culture

CHO-hGHS-R (Euroscreen s.a. Brussels, Belgium) were cultured in Ultra-CHO medium from Cambrex Bio Science (Walkersville, MD, USA) supplemented with 1% dialyzed fetal bovine serum (FBS), 50 μg/ml gentamicin, and 400 μg/ml G418 (all from Invitrogen) at 37 °C in a humidified cell incubator containing 5% CO2. Rat pituitary adenoma cells, RC-4B/C (CRL-1903; ATCC, Manassas, VA, USA), were cultured in a 1:1 mixture of Dulbecco’s Modified Eagle’s Medium (Sigma) and Minimum Essential Alpha Medium (Invitrogen) supplemented with 0·01 mM non-essential amino acids (Sigma), 15 mM HEPES (Sigma), 2·5 ng/ml epidermal growth factor (Invitrogen), and 10% dialyzed, heat-inactivated FBS at 37 °C in a humidified cell incubator containing 5% CO2. Sub-cloning of the parental population was performed by limiting dilution cloning in 96-well tissue culture plates using a conditioned RC-4B/C media. Conditioned media was prepared by incubation on a confluent monolayer of RC-4B/C cells for 24 h, after which time the media was decanted, filtered through a 0·22 μm filter, and stored at 4 °C until use.
Reagents

GHS-R agonist hexarelin (His-D-2-methyl-Trp-Ala-Trp-δ-Phe-Lys) was purchased from American Peptide Inc. (Sunnyvale, CA, USA), rat ghrelin (Gly-Ser-Ser (n-octanoyl)-Phe-Leu-Ser-Pro-Glu-His-Gln-Lys-Ala-Gln-Gln-Arg-Lys-Glu-Ser-Lys-Lys-Pro-Pro-Ala-Lys-Leu-Gln-Pro-Arg) was synthesized at Abbott, and rat des-octanoyl ghrelin was either purchased from Phoenix Pharmaceuticals Inc. (Belmont, CA, USA) or synthesized at Abbott. GHS-R antagonist [D-Lys3]-GHRP-6 (H-His-D-Trp-δ-Lys-Trp-δ-Phe-Lys) and the inverse agonist (D-Arg1, D-Phe5, D-Trp7,9, Leu11)-substance P analog were purchased from Bachem Bioscience Inc. (King of Prussia, PA, USA). The PLC inhibitor U-73122 was purchased from Bioprobe. Thapsigargin was purchased from Alexis Biochemicals (Lausen, Switzerland). [3H]myoinositol was purchased from Perkin-Elmer (Shelton, CT, USA). AG1-X8 anion-exchange resin columns were purchased from Perkin-Elmer (Shelton, CT, USA). ATP, GTP, MgSO4, and CaCl2 were obtained from Sigma-Aldrich (St. Louis, MO, USA). 

myo-inositol was purchased from BioRad. The L-type calcium channel blocker, nifedipine, Ptx, and myoinositol were purchased from Sigma. Compounds A and B were synthesized at Abbott.

Reverse transcriptase PCR (RT-PCR)

Total RNA was isolated from cell pellets using TRI-reagent (Sigma) according to the manufacturer’s specifications. First-strand cDNA was generated using the SuperScript First-Stand Synthesis System (Invitrogen) with oligo (dT) primers and 5 μg total RNA. The target cDNA was then amplified by PCR. The reaction contained 10×PCR buffer (Invitrogen), 200 nM dNTP mix, 1·5 mM MgCl2, 200 nM primers to human GHS-R1a, 5′-TCTTGGTTCTTCCCTCTCTGTC3′ (sense) and 5′-AGCTGAGACTGACCACG3′ (antisense), and platinum Taq DNA polymerase (Invitrogen) in 50 μl total volume. Thermal cycling consisted of 5 min at 95 °C followed by 40 cycles of 30 s at 95 °C, 30 s at 50 °C, 2 min at 72 °C, and was concluded with 10 min at 72 °C.

Fluorescent calcium indicator (FLIPR) assay

Agonist EC50/antagonist IC50 determinations

RC-4B/C parental or RC-4B/C.40 cells were plated in black 96-well plates with clear bottoms (Corning Inc. Corning, NY, USA) and cultured to confluence overnight in growth medium at 37 °C in a humidified cell incubator containing 5% CO2. Growth medium was replaced with 100 μl Dulbecco’s PBS (DPBS) containing 1000 mg/l D-glucose, 36 mg/l sodium pyruvate (Invitrogen) supplemented with 1·14 mM Fluo-4 AM (Molecular Probes, Eugene, OR, USA) and 2·5 mM probenecid (Sigma) for 1–3 h in the dark at room temperature. For EC50 determinations, various concentrations of agonist were prepared at 4× final concentrations in DPBS containing 0·1% BSA.

After aspirating the dye solution, cells were washed three times in DPBS using an ELx-405 Automatic Plate Washer (BioTek Instruments, Winooski, VT, USA). Following the last wash, cells were placed in 150 μl/well of DPBS and the plates were then transferred to the FLIPR unit (Molecular Probes) where 50 μl of 4× agonist doses were added and the agonist-induced rise of [Ca2+], detected by Fluo-4AM fluorescence was followed for 3 min. Fluorescence emissions from 96 wells were measured simultaneously at excitation and emission wavelengths of 488 and 520 nm respectively for 3 min in 1-s intervals for the first minute and 5-s intervals thereafter. Agonist activity was thus expressed as percent activation with an EC50 being the concentration of agonist causing a 50% activation of a maximal ghrelin response. For IC50 determinations, after loading with dye solution, washing, and replacing with 100 μl DPBS, cell plates were placed in FLIPR where 50 μl/well of 4× antagonist concentrations were added and measurements taken for 3–6 min, followed by the addition of 50 μl/well of 4× ghrelin ligand in DPBS/0·1% BSA (final concentration of 3 or 10 nM for RC-4B/C cells and 1 nM for CHO-hGHS-R cells). Fluorescence measurements were then taken for another 5 min. IC50s were determined as the concentration of antagonist causing a 50% reduction of the ghrelin-induced calcium signal. Sigmoidal curves were fitted by GraphPad Prism software (GraphPad Software, Inc., San Diego, CA, USA) and EC50 or IC50 values were calculated based on results from triplicate determinations.

Effects of pertussis (Ptx)

RC-4B/C.40 cells were exposed to 1000 ng/ml Ptx for 4 h, 37 °C. At the end of the 4 h incubation, the culture medium was aspirated, 100 μl DPBS containing glucose and pyruvate were added to each well along with 100 μl calcium assay reagent (Molecular Devices, Sunnyvale, CA, USA) containing 2·5 mM probenecid. Cells were analyzed in the FLIPR for their response to rat ghrelin as described in the previous subsection.

Effects of extracellular and intracellular Ca2+

The role of extracellular Ca2+ in ghrelin signaling in RC-4B/C.40 cells was determined as described above, except that after loading with Fluo-4 AM dye solution, cells were washed a minimum of three times in Ca2+/Mg2+-free DPBS, and then stimulated with rat ghrelin in either standard Ca2+/Mg2+-containing or Ca2+/Mg2+-free DPBS. To further define the role of extracellular Ca2+, cells were also pretreated with the L-type calcium channel blocker, nifedipine. Briefly, after washing three times in Ca2+/Mg2+-free DPBS,
cells were transferred to the FLIPR unit where 150 μl of a 1:5× dose of nifedipine was added (for a final concentration of 100 μM), and the signal was monitored for 6 min. Following pretreatment with nifedipine, cells were stimulated with 10 nM ghrelin and the effects on [Ca\(^{2+}\)]\(_i\) were monitored for an additional 3 min. To assess the role of Ca\(^{2+}\) release from intracellular stores, RC-4B/C.40 cells were washed in Ca\(^{2+}\)/Mg\(^{2+}\)-free DPBS and pretreated with 1 μM thapsigargin in Ca\(^{2+}\)/Mg\(^{2+}\)-free DPBS (causing Ca\(^{2+}\) release from intracellular stores and preventing re-uptake into the endoplasmic reticulum). The thapsigargin-induced release of intracellular Ca\(^{2+}\) was monitored in FLIPR for a minimum of 6 min (allowing for the return of signal to baseline). Cells were then stimulated with rat ghrelin in either Ca\(^{2+}\)/Mg\(^{2+}\)-free or Ca\(^{2+}\)/Mg\(^{2+}\)-containing DPBS. The ghrelin-induced fluorescence signal was monitored on FLIPR for an additional 5 min. The resultant fluorescent signal tracings were normalized to the basal signal in Ca\(^{2+}\)/Mg\(^{2+}\)-free DPBS using FLIPR software and further analyzed in GraphPad Prism (GraphPad Software).

### PI hydrolysis assay

The constitutive activity of GHS-R in RC-4B/C.40 and CHO-hGHS-R cells was determined using a PI hydrolysis assay that measures the accumulation of [\(^{3}\)H]inositol phosphates following activation of GHS-R. Briefly, cells were seeded at 5 × 10\(^5\) cells/well in 24-well plates and cultured for 24 h. Cells were then labeled for 16–18 h with [\(^{3}\)H]myoinositol (2 μCi/well). After washing with room temperature DPBS, cells were treated with 200 μl/well of 10 mM LiCl in Earle's/25 mM HEPES/0.0073 mM pepstatin/0.1 mM PMSF buffer for 45–60 min at 37°C in a 5% CO\(_2\) incubator to block endogenous myoinositol mono-phosphatase activity. Next, cells were stimulated with either ghrelin or [\(\alpha\)-Arg\(^1\), \(\beta\)-Phe\(^5\), \(\delta\)-Tryp\(^7\), Leu\(^11\)]-substance P analog for 30 min at 37°C. Cells were lysed by the addition of 50 μl/well of 1 N NaOH followed by an equal volume of 1 N HCl. Samples were then transferred to 12 × 75 mm glass tubes, extracted by adding 1.5 ml/tube of chloroform/methanol (1:2, v/v), vortexed gently, and incubated for 1 h at room temperature. Additional 0.5 ml/tube volumes of chloroform were added followed by 0.4 ml/tube of double distilled (dd) H\(_2\)O. Following centrifugation at 1100 × g for 20 min at room temperature, the aqueous phase was diluted to 3 ml with ddH\(_2\)O. Samples were loaded onto AG1-X8 columns and washed with 20 ml ice-cold, non-radioactive myoinositol, followed by 6 ml 60 mM sodium formate/5 mM sodium tetraborate. The [\(^{3}\)H]inositol phosphate fraction was eluted with 5 ml 1 M ammonium formate/0.1 N formic acid. Total counts per minute (CPM) were determined in a Beckman LS 6500 scintillation counter.

### Results

#### GHS-R mRNA expression in RC-4B/C cells

A single product of the expected size (349 bp) was generated by RT-PCR using cDNA derived from the RC-4B/C cell line (Fig. 1, lane c). The corresponding negative control reaction lacking the RT did not yield any products (Fig. 1, lane d), thus ruling out potential contamination by genomic DNA in the RNA isolation. PC3, a prostate cancer cell line that was previously shown to express GHS-R (Volante et al. 2002), yielded a product of expected size (Fig. 1, lane b). In addition, cDNA from CHO-GHS-R cells and a vector encoding GHS-R1a were used as positive controls. Taken together, these observations suggest that RC-4B/C cells express GHS-R1a mRNA.

#### Ca\(^{2+}\) Flux response to GHS-R agonists and antagonists

**RC-4B/C parental cells**

We next investigated whether the RC-4B/C cell line, expressing an endogenous GHS-R1a, could be used as a reagent for screening of GHS-R agonists and antagonists *in vitro*. To determine the functional activity of GHS-R, we measured [Ca\(^{2+}\)]\(_i\) using fluorescence imaging. RC-4B/C cells were treated with ghrelin, hexarelin, and des-octanoyl ghrelin, an inactive form of ghrelin serving as a negative control. These peptides were tested at half-log concentrations and their activity expressed as percent maximal activation. Both ghrelin and hexarelin demonstrated a robust dose-dependent response in the RC-4B/C cells with EC\(_{50}\) values of 1·04 nM and 1·67 nM respectively (Fig. 2A). Des-octanoylated ghrelin, on the other hand, did not elicit a signal at any of the doses tested (Fig. 2A), indicating that the ghrelin response in the RC-4B/C cells is specific to its active form. We also tested the effects of GHS-R antagonists in this assay. Preincubation of RC-4B/C cells with a known peptidyl antagonist [\(\beta\)-Lys\(^3\)]-GHRP-6 and compound A resulted in a dose-dependent inhibition

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**Figure 1** GHS-R1a mRNA expression in various cell lines. RT-PCR products amplified from (a) CHO-hGHS-R cells, (b) PC3 cell line, (c) RC-4B/C cell line, (d) negative control, RC-4B/C total RNA preparation without the addition of reverse transcriptase, (e) positive control, vector encoding the human GHS-R1a gene.
of ghrelin-induced increase in Ca\textsuperscript{2+} flux, with IC\textsubscript{50} values of 3.86 µM and 73.1 nM respectively (Fig. 2B). Compound B, a structural homolog of compound A, was also tested and found to be inactive (Fig. 2B). This finding is in agreement with the poor affinity that compound B displayed for GHS-R in a binding assay (data not shown).

**RC-4B/C.40 subclone**

The RC-4B/C cells are a heterogeneous population of corticotropic and gonadotropic, as well as somatotropic cells. While it has been possible to evaluate GHS-R agonists and antagonists using the parental cells, further characterization of ghrelin signaling is complicated by the presence of these other cell types. Another factor limiting the utility of the parental RC-4B/C cells has been the greatly diminished or even loss of ghrelin responsiveness routinely observed after only ten culture passages. Consequently, an effort to identify a stable subclone was initiated which resulted in the screening of 60 subclones displaying various morphologies. Among these, clone 40 (RC-4B/C.40) demonstrated a significantly more robust ghrelin response in the Ca\textsuperscript{2+} mobilization assay that was more stable over multiple passages compared with the parental RC-4B/C cells (Fig. 3). Thus, the subclone RC-4B/C.40 was selected for further ghrelin signaling studies. RTqPCR analysis indicated that GHS-R mRNA expression is approximately twofold higher in the RC-4B/C.40 cells than in the parental RC-RB/C cells (data not shown).

![Figure 2](image1.png)

**Figure 2** Effects of GHS-R agonists and antagonists in RC-4B/C cells. (A) Agonist effects – cells were treated with various concentrations of ghrelin (■), hexarelin (◇), and des-octanoyl ghrelin (○), and their effects on [Ca\textsuperscript{2+}] were measured by fluorescence imaging using fluorescent calcium indicator (FLIPR) with activity expressed as a percentage of a maximal, 10 nM ghrelin response. (B) Antagonist effects – cells were treated with various concentrations of [d-Lys-3]-GHRP-6 (◇), compound A (■), or compound B (○) prior to stimulation with ghrelin. Their effects were expressed as percent inhibition of maximal ghrelin response (10 nM). A representative of at least two independent experiments is shown with triplicate measurements at each peptide concentration.

![Figure 3](image2.png)

**Figure 3** Ghrelin dose–response in multiple passages of the RC-4B/C.40 subclone versus the parental RC-4B/C cells. Ghrelin dose–response curves for separate passages of the subclone RC-4B/C.40 cells are depicted (P.5 cells, EC\textsubscript{50} = 21.2 nM, solid line; P.9 cells, EC\textsubscript{50} = 7.2 nM, dotted line; and P.12 cells, EC\textsubscript{50} = 14.3 nM, dashed line). A representative ghrelin dose–response for the parental RC-4B/C cells (P.7 cells, EC\textsubscript{50} = 109 nM, dashed gray line) is shown for comparison. RFU, relative fluorescence units.
As observed in the parental cells (Fig. 2), treatment of the RC-4B/C.40 cells with both rat ghrelin and the related peptide agonist hexarelin resulted in robust, dose-dependent responses, yielding EC₅₀ values of 5·5 and 2·7 nM respectively (Fig. 4A). Treatment with des-octanoyl rat ghrelin did not elicit a signal at any dose of peptide, indicating that the RC-4B/C.40 cells respond specifically to active ghrelin (Fig. 4A). The RC-4B/C.40 cells were also treated with the peptidic GHS-R antagonist [d-Lys⁵] GHRP-6 as well as the small molecule inhibitor, indicating that the RC-4B/C.40 cells respond specifically to active ghrelin (Fig. 4A). The RC-4B/C.40 peptide, indicating that the RC-4B/C.40 cells respond specifically to active ghrelin (Fig. 4A). Both [d-Lys³] GHRP-6 and the small molecule compound A effectively inhibited the 10 nM ghrelin response, with IC₅₀ values of 4·09 µM and 195 nM respectively, while compound B was inactive, with an IC₅₀ > 10 µM (Fig. 4B). Again, the results for [d-Lys³] GHRP-6 agree well with literature-reported values and the effects of all three agents in the RC-4B/C.40 cells are similar to the effects observed in both the parental RC-4B/C cells (Fig. 2B) and the CHO-K cells that stably overexpress human GHS-R (data not shown).

**Characterization of ghrelin Ca²⁺ signaling in the subclone RC-4B/C.40 cells**

In order to further characterize ghrelin signaling in the RC-4B/C.40 subclone, we pretreated cells with Ptx, which specifically inhibits Gₛ and Gₒ coupling, or with the PLC inhibitor U-73122, which blocks the signal downstream of Gₜ, activation that ultimately causes the rise of [Ca²⁺], measured in FLIPR. Pretreatment with 1 µg/ml Ptx for 4 h at 37 °C did not affect the ghrelin dose–response, although the maximal signal attained was slightly higher (Fig. 5A). On the other hand, pretreatment with 1 µM, 3 µM, and 10 µM U-73122 resulted in a dose-dependent inhibition of the ghrelin response (Fig. 5B). Taken together, these results suggest that ghrelin signaling in RC-4B/C.40 cells occurs primarily through Gₛ and not Gₒ or Gₒ coupling.

We next determined the relative contributions of extracellular Ca²⁺ influx and release of Ca²⁺ from intracellular stores in the ghrelin-induced Ca²⁺ signal. RC-4B/C.40 cells were stimulated with 3 nM and 100 nM ghrelin in either the presence or absence of extracellular Ca²⁺. The ghrelin response in the presence of extracellular Ca²⁺ was of greater magnitude (with a peak signal 22% greater than the signal without extracellular Ca²⁺) and sustained duration (with approximately 50% of the signal remaining at 300 s after ghrelin stimulation) than that observed in the absence of extracellular Ca²⁺ (Fig. 6A). To further demonstrate the role of extracellular Ca²⁺ in ghrelin signaling, we treated cells with the L-type channel blocker, nifedipine. Pretreatment with 100 µM nifedipine in the presence of extracellular Ca²⁺ resulted in a 43% reduction of the ghrelin-induced rise in [Ca²⁺], as compared with the untreated cells (Fig. 6B). Next, to address the role of Ca²⁺ release from intracellular stores in the ghrelin signal, we treated cells with thapsigargin. Thapsigargin causes the release of Ca²⁺ from

![Figure 4](https://example.com/figure4.png)

**Figure 4** Effects of GHS-R agonists and antagonists on the subclone RC-4B/C.40 cells. (A) Agonist treatment: cells were treated with various concentrations (half-log dose increments ranging from 0·01 to 100 nM) of rat ghrelin (▴), hexarelin (▽), and des-octanoyl rat ghrelin (▪). (B) Antagonist treatment: RC-4B/C.40 cells were treated with various, half-log concentrations (ranging from 0·3 nM to 100 µM) of the peptide antagonist [d-Lys³] GHRP-6, the small molecule compound A, and the structurally related analog, compound B. Both [d-Lys³] GHRP-6 (▴, dashed line) and the small molecule compound A (‴, solid line) dose dependently inhibited the 10 nM ghrelin response (IC₅₀ of 4·09 µM and 195 nM respectively) while compound B (▪ solid line) was essentially inactive (IC₅₀ > 10 µM). The results shown are representative of at least two independent experiments with triplicate measurements at each peptide concentration.
intracellular stores and prevents re-uptake into the endoplasmic reticulum (ER). Pretreatment with 1 μM thapsigargin was shown to effectively block the ghrelin response in the absence of extracellular Ca\textsuperscript{2+} (Fig. 7). This effect, however, was not observed in the presence of extracellular Ca\textsuperscript{2+}, possibly reflecting the drive to restore intracellular Ca\textsuperscript{2+} tone (Fig. 7). Together, these results suggest that the ghrelin-induced rise of [Ca\textsuperscript{2+}]\textsubscript{i} involves an initial release from intracellular stores followed by an influx through L-type Ca\textsuperscript{2+} channels that contributes to both the magnitude and sustained duration of the signal.

**Constitutive activity of GHS-R in RC-4B/C.40 and CHO-hGHS-R cells**

The constitutive activity of GHS-R in the RC-4B/C.40 and the recombinant CHO-hGHS-R cells was evaluated.

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**Figure 5** Effects of Ptx and PLC inhibitor U-73122 on ghrelin signaling in RC-4B/C.40 cells. (A) Pretreatment with 1 μg/ml Ptx for 4 h showed no effects on the ghrelin dose–response (dashed line) compared with the untreated control dose–response (solid line), with EC\textsubscript{50} values of 6.15 and 8.56 nM respectively. (B) Pretreatment with 1, 3, and 10 μM PLC inhibitor U-73122 resulted in dose-dependent inhibition of a 10 nM ghrelin response, with >90% inhibition observed at the maximum 10 μM dose. Ctrl, control; RFU, relative fluorescence units.

**Figure 6** Ghrelin signaling in RC-4B/C.40 cells in the presence and absence of extracellular Ca\textsuperscript{2+} and following pretreatment with the L-type channel blocker nifedipine. (A) Representative Ca\textsuperscript{2+} signal tracings measured in FLIPR are shown following stimulation with 3 nM ghrelin (▲) and 100 nM ghrelin (■) in the presence of extracellular Ca\textsuperscript{2+}, and for 3 nM ghrelin (○) and 100 nM ghrelin (●) in the absence of extracellular Ca\textsuperscript{2+}. (B) Effects of pretreatment with 100 μM nifedipine on the 10 nM ghrelin response are shown. The Ca\textsuperscript{2+} signal tracing for the untreated, control cells is shown as a black line with solid black squares (■). The signal tracing for the ghrelin response following pretreatment with 100 μM nifedipine is shown as a gray line with open gray triangles (△). RFU, relative fluorescence units.
using a PI hydrolysis assay that measures the accumulation of \[^3H\]inositol phosphates from \[^3H\]myo-inositol turnover. In accordance with the G-protein coupled receptor (GPCR) signaling paradigm, ghrelin or basal stimulation of GHS-R presumably activates G\(_{\text{q}}\), which, in turn, activates PLC, resulting in the formation of 1,4,5-triphosphate (IP\(_3\)) from phosphatidylinositol 4,5-bisphosphate (PI (4,5) P\(_2\)). The increase in IP\(_3\) then causes intracellular Ca\(^{2+}\) release. CHO-hGHS-R cells exhibit a basal level of constitutive GHS-R activity, which increased by 40% upon stimulation with 100 nM ghrelin (Fig. 8A). Treatment with 50 nM [D-Arg\(^1\), D-Phe\(^5\), D-Trp\(^7,9\), Leu\(^11\)]-substance P analog, a known GHS-R inverse agonist (Holst et al. 2003), resulted in a statistically significant 37% reduction of the basal activity (\(P \text{ value } < 0.005\)), indicating that GHS-R overexpressed in the CHO line constitutively exhibits about 50% of maximal activity under basal conditions, as reported in other recombinant cell systems (Holst et al. 2003). In contrast, endogenous GHS-R in RC-4B/C40 cells does not exhibit significant constitutive activity (Fig. 8B). Stimulation with 100 nM ghrelin resulted in an 88% increase of \[^3H\]inositol phosphate over that detected under basal conditions as expected. However, unlike in CHO-hGHS-R cells, treatment with 10 nM and 100 nM [D-Arg\(^1\), D-Phe\(^5\), D-Trp\(^7,9\), Leu\(^11\)]-substance P analog did not result in any significant reduction of basal activity, suggesting that constitutive activity of GHS-R in the RC-4B/C.40 cells is undetectably low.

**Discussion**

The GHS family of peptides and small molecules was known long before the receptor mediating their GH-releasing activity was identified. However, based on its function, it was suspected that the receptor would...
be found in the pituitary and hypothalamus. Indeed, GHS-R was first characterized in the plasma membrane fraction of pituitary and hypothalamic tissue by binding of [35S]MK0677, a small molecule GHS (Smith et al. 1997). Subsequently, the expression of GHS-R and ghrelin, by then its newly identified natural ligand, was demonstrated in the normal and abnormal human pituitary using RT-PCR (Korbonits et al. 1998, 2001). Furthermore, recent studies involving double immunofluorescence localized ghrelin in specific cell populations contained within the rat anterior pituitary. It is expressed in somatotrophs, lactotrophs and thyrotrophs, but not in the corticotrophs or gonadotrophs (Caminos et al. 2003). GHS-R mRNA expression has also been shown to be the highest in human somatotroph tumors and essentially non-detectable in gonadotroph tumors (Korbonits et al. 2001). As the expression and function of ghrelin and its receptor in the anterior pituitary is well established, we investigated whether a tumor cell line derived from the pituitary could be used as a routine reagent for measurements of GHS-R activity in vitro. The RC-4B/C cell line was established in The Jackson Laboratory (Bar Harbor, ME, USA) from an aged male rat with pituitary adenoma. It is a heterogeneous cell line containing all known anterior pituitary hormone-secreting cell types as demonstrated by immunocytochemistry (Hurbain-Kosmath et al. 1990). We demonstrate here using RT-PCR that RC-4B/C cells express GHS-Rα1a mRNA.

The elucidation of the second messenger system involved in GHS-R signaling is an area of great interest. Earlier studies involving activation of GHS-R by artificial GHSs showed a variety of responses such as an increase in intracellular cAMP and free Ca2+ levels as well as activation of PKA and PKC (Chen 2000). Recently, the ghrelin-stimulated GHS-R signal transduction pathway has been investigated in isolated porcine somatotrophs. It was shown that ghrelin increased the [Ca2+]i, in 98% of the cells that responded to GHRH, an effect that was decreased in the presence of a GHS-R peptidic antagonist [d-Lys3]-GHRP-6 (Glavaski-Joksimovic et al. 2003). In addition, another report demonstrated a ghrelin dose-dependent increase in the [Ca2+]i in rat-derived single hypothalamic neuropeptide Y-neurons, known for their critical role in the stimulation of food intake (Kohno et al. 2003).

Ghrelin and hexarelin have previously been shown to be equipotent in a binding assay using pitutitary membrane preparations (Muccioli et al. 2001). We show here that ghrelin and hexarelin, a non-natural peptidyl GHS (Arvat et al. 2001), activated GHS-R in RC-4B/C cells with similar potency, as judged by an increase in the [Ca2+]i. Their EC50 values in RC-4B/C cells are comparable to those obtained using CHO cells overexpressing GHS-R (data not shown). In contrast, des-octanoyl ghrelin, a precursor that is devoid of the endocrine activity of acylated ghrelin (Broglio et al. 2003), had no effect on the [Ca2+]i, in RC-4B/C cells. In addition, antagonists of GHS-R blocked the ghrelin-induced increase in [Ca2+]i. Compound A was more potent than [d-Lys-3]-GHRP-6, consistent with their relative affinities for GHS-R in a radiolabeled ghrelin binding assay using membrane preparations of CHO cells overexpressing GHS-R (data not shown). Compound B, a structural homolog of compound A but with a weak binding affinity for GHS-R, was inactive in the [Ca2+]i flux assay.

While effective as an in vitro tool for screening GHS-R pharmacophores, the RC-4B/C cells proved to be unsuitable for further characterization of ghrelin signal transduction, possibly due to being a heterogeneous cell population. Moreover, a consistent pattern of diminished ghrelin responsiveness during cell culture was observed, which further limited their utility as an in vitro screening tool. In an attempt to overcome these limitations, a subclone RC-4B/C.40, was generated. It demonstrated the most stable and highest magnitude of ghrelin response compared with both the parental RC-4B/C cells and other subclones. Like the parental cells, RC-4B/C.40 cells showed a robust response to both rat ghrelin and the peptide agonist hexarelin, while no response was observed with des-octanoyl ghrelin treatment. Likewise, treatment with the peptide antagonist [d-Lys3]-GHRP-6 or the small molecule inhibitor, compound A, effectively blocked the ghrelin response, while the structural analog, compound B, was essentially inactive.

A number of reports have identified molecules potentially involved in the ghrelin-induced Ca2+ signaling pathway. For example, using enzyme inhibitors, it was shown that adenyl cyclase and PLC are involved in GHS-R signaling in somatotropes (Glavaski-Joksimovic et al. 2003), and PKA in the hypothalamic neurons (Kohno et al. 2003). Using calcium channel blockers, it was also demonstrated that L-type channels are essential for Ca2+ influx in somatotropes (Glavaski-Joksimovic et al. 2003) and N-type channels in hypothalamic neurons (Kohno et al. 2003). As a homogeneous cell population, the RC-4B/C.40 subclone provided the means to investigate ghrelin signal transduction. We demonstrate here that treatment with Ptx, which inhibits Gi and Go coupling, had no effect on the ghrelin signal, while pretreatment with the PLC inhibitor U-73122 dose-dependently blocked the ghrelin signal in RC-4B/C.40 cells. Similarly, treatment with the PLC inhibitor U-73122 dose-dependently blocked the ghrelin signal in RC-4B/C.40 cells, suggesting that signaling occurs primarily through Gq coupling, as expected. We also demonstrated that the ghrelin-induced [Ca2+]i flux in RC-4B/C.40 cells is dependent on both influx of extracellular Ca2+, consistent with observations in isolated somatotropes, and release from intracellular stores. Removal of Ca2+ from the cell medium
greatly reduced the overall magnitude and duration of the ghrelin-induced increase in $[\text{Ca}^{2+}]_i$, as did incubation with the L-type channel blocker nifedipine, indicating the involvement of calcium channels in the ghrelin response. The release of intracellular $\text{Ca}^{2+}$ stores by thapsigargin also effectively blocked the ghrelin response in the absence of extracellular $\text{Ca}^{2+}$. Taken together, these observations indicate that ghrelin signaling through GHS-R involves the release of $\text{Ca}^{2+}$ from intracellular stores as well as influx of extracellular $\text{Ca}^{2+}$ through plasma membrane ion channels.

Recent studies involving GHS-R expression in recombinant cell systems have reported a high degree of constitutive activity, similar to that observed with several other GPCRs, including the neurotensin receptor 2 (NT-R2) and the viral ORF-74 receptor (Holst et al. 2003, Holst & Schwartz 2004). Using inositol phosphate turnover and a cAMP response element-dependent transcriptional reporter assay to assess constitutive activity, Schwartz and colleagues demonstrated that GHS-R transiently expressed in COS-7 and HEK 293 cells signals at about 50% maximal activation in the absence of ghrelin (Holst et al. 2003). Stimulation with ghrelin results in a twofold increase in activity, while treatment with the inverse agonist [D-Arg1, D-Phe5, D-Tryp7,9, Leu11]-substance P analog potently suppresses basal activity. The constitutive activity of GHS-R in these recombinant cell systems may be an artifact of receptor overexpression. However, the structurally homologous motilin receptor does not show measurable constitutive activity when expressed in a recombinant system, suggesting that high constitutive activity may in fact be an intrinsic, physiologically relevant property of GHS-R (Holst et al. 2003, Holst & Schwartz 2004).

We decided to investigate the constitutive activity of GHS-R in the RC-4B/C.40 cells using a PI hydrolysis assay and compare it with the level of activity in the recombinant CHO-hGHS-R cells. The CHO-hGHS-R cells exhibited a high degree of constitutive activity, which was reduced upon treatment with the inverse agonist [D-Arg1, D-Phe5, D-Tryp7,9, Leu11]-substance P analog, in agreement with the effects observed in other recombinant cell systems (Holst et al. 2003, 2004, Holst & Schwartz 2004). In contrast, the RC-4B/C.40 cells showed little, if any, constitutive activity. The basal inositol phosphate level was threefold less than in the CHO cells and was not further reduced by treatment with the inverse agonist [D-Arg1, D-Phe5, D-Tryp7,9, Leu11]-substance P analog.

Thus, the endogenously expressed GHS-R in RC-4B/C.40 cells does not exhibit constitutive activity in our hands. As this runs counter to what has recently been reported in several recombinant cell lines, our observations raise important questions as to the physiological relevance of GHS-R constitutive activity. A possible explanation for the disparity between the levels of constitutive activity in the recombinant CHO line versus the RC-4B/C.40 cells is the presence of an intracellular regulator of GHS-R signaling, which functions to suppress GHS-R constitutive activity. Presumably, such an intracellular regulator would not be expressed in the recombinant CHO-hGHS-R line where its absence may account for the high level of constitutively active GHS-R observed, while a more physiologically relevant pituitary cell line, RC-4B/C, may express the regulator, accounting for the low level of constitutive activity in this cell line. Alternatively, as with the MC-4 receptor, where AgRP functions as an inverse agonist, there may be an unidentified endogenous inverse agonist that attenuates GHS-R constitutive activity and may play a critical role in mediating the effects on the complex network of neuroendocrine inputs to ghrelin effector cells (Nijenhuis et al. 2001, Holst et al. 2003). However, such an endogenous inverse agonist would most likely only be observed in an in vivo setting and not in a homogeneous cell population in vitro. Nonetheless, the RC-4B/C.40 cells may ultimately provide a more relevant cell line in which to model and investigate ghrelin signaling than what can be currently achieved using recombinant GHS-R cell systems.

In conclusion, we have identified a suitable cell line for GHS-R pharmacology and signaling studies. The rat anterior pituitary RC-4B/C cells express endogenous GHS-R that responds to ghrelin as judged by increased $[\text{Ca}^{2+}]_i$. GHS-R antagonists blocked the ghrelin response with the anticipated rank order of potency. Additionally, we were able to identify and characterize a subclone, RC-4B/C.40, which demonstrated a more robust and stable ghrelin response than that observed in the parental cells. Along with showing agonist and antagonist effects similar to those observed in the parental RC-4B/C cells and in CHO-hGHS-R cells, the RC-4B/C.40 subclone also provided the means to more fully characterize ghrelin signaling. The mechanism of ghrelin-induced increase in $[\text{Ca}^{2+}]_i$ in the RC-4B/C.40 cells involves Gs coupling along with both extracellular $\text{Ca}^{2+}$ flux and release of $\text{Ca}^{2+}$ from intracellular stores. The lack of constitutive activity of GHS-R in RC-4B/C.40 cells indicates that GHS-R functions differently in this more physiologically relevant cell type than in recombinant cell lines. Thus, the RC-4B/C parental and RC-4B/C.40 cell lines described here represent useful in vitro tools for the identification of GHS-R agonists or antagonists with potential therapeutic applications in the treatment of growth hormone deficiency or body weight disorders, and, more significantly, for examination of the mechanism of GHS-R activation and signaling, including the basis of constitutive activity.
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