

Growth hormone-releasing hormone (GHRH) and its agonists inhibit hepatic and tumoral secretion of IGF-1

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ABSTRACT

The role of hypothalamic growth hormone-releasing hormone (GHRH) in the release of growth hormone (GH) from the pituitary is well established. However, direct effects of GHRH and its agonistic analogs on extra-pituitary cells and tissues have not been completely elucidated. In the present study, we first demonstrated that human and rat hepatocytes express receptors for GHRH. We then showed that GHRH(1-29)NH₂ and GHRH agonist, MR-409, downregulated mRNA levels for IGF-1 in human cancer cell lines and inhibited IGF-1 secretion *in vitro* when these cancer lines were exposed to rhGH. Another GHRH agonist, MR-356, lowered serum IGF-I and inhibited tumor growth in nude mice bearing xenografted NCI-N87 human stomach cancers. GHRH(1-29)NH₂ and MR-409 also suppressed the expression of mRNA for IGF-1 and IGF-2 in rat and human hepatocytes, decreased the secretion of IGF-1 *in vitro* from rat hepatocytes stimulated with rhGH, and lowered serum IGF-I levels in hypophysectomized rats injected with rhGH. Vasoactive intestinal peptide had no effect on the release of IGF-1 from the hepatocytes. Treatment of C57BL/6 mice with MR-409 reduced serum levels of IGF-I from days 1 to 5. These results show that GHRH and its agonists can, by a direct action, inhibit the secretion of IGF-1 from the liver and from tumors. The inhibitory effect of GHRH appears to be mediated by the GHRH receptor (GHRH-R) and GH receptor (GHR), with the involvement of JAK2/STAT5 pathways. Further studies are required to investigate the possible physiopathological role of GHRH in the control of secretion of IGF-1.

INTRODUCTION

During the past years, our group reported the synthesis of multiple antagonistic analogs of growth hormone releasing hormone (GHRH) and the biological evaluation of their inhibitory effects on growth of various tumors [1, 2]. We have also synthesized powerful agonistic analogs of GHRH [3] and demonstrated that in rodents, they improve cardiac function after experimental myocardial infarction and reduce cardiac hypertrophy [4-6], stimulate the proliferation and regeneration of pancreatic islet cells [7, 8], and accelerate wound healing

[9]. Although for many years the sole role of GHRH was thought to be the regulation of the release of growth hormone (GH) from the pituitary, these findings show that GHRH and its analogs can exert direct effects on extra-pituitary cells/tissues.

Insulin-like growth factor 1 (IGF-1), plays an essential physiological role on the growth and development, [10-13]. The functions of IGF-1 in the development and progression of cancers are also well documented. IGF-1 has been shown to stimulate cancer cell proliferation, inhibit apoptosis [14, 15] and facilitate metastasis by promoting cell migration and invasion [16].

Most circulating IGF-1 is produced by the liver, following the stimulation with pituitary GH [17]. High levels of circulating IGF-1 have been linked to an increased risk of neoplasia and poor prognosis in many clinical studies [18, 19]. Various studies have been carried out on secretion of IGF-1 in physiological, clinical and pathological conditions [20-25], but to the best of our knowledge the involvement of a neurohumoral component in these situations has not been previously identified. This paper describes our investigations of the effects of GHRH and its agonists on the synthesis and release of IGF-1 in normal liver tissue and in tumors.

RESULTS

Human and rat hepatocytes express GHRH receptor

The presence of GHRH receptor in primary human and rat hepatocytes was determined by a PCR-based method and Western blots. Human or rat pituitaries were used as positive controls. PCR primers, (F) GATGAGAGTGCCTGTCTACAAGCA, (R) TCTGAGCTGAAGTGAGAGAAGAAATC, were designed to target a unique region between exon 2 and exon 3 of mRNA for human GHRH-R (Genebank: NM_000823). The PCR primers, (F) CCTTCCAGGGTTTTGTTGTTG, (R) GGTGAGCACCTTCACTCTCGAT, were used to target mRNA for rat GHRH-R (Genebank: NM_012850). The PCR products amplified from the cDNA of human/rat hepatocytes and their respective pituitary control exhibited

the expected size (Figure 1A, 1B). The specificity of the results of PCR was further verified by DNA sequencing (data not shown). The expression of GHRH receptor at the protein level was determined by Western blots. The GHRH-R antibody recognized a band of 47 kD (Figure 1C, 1D) in the pituitary and the hepatocytes from both human and rats, which matched the calculated size of the unglycosylated pituitary type GHRH receptor. Both the PCR and Western blot findings confirmed the presence of GHRH receptor in primary human and rat hepatocytes.

GHRH(1-29)NH₂ and its agonist, MR-409 suppress the expression of mRNA for IGF-1 and IGF-2 in human and rat hepatocytes

Direct effects of GHRH(1-29)NH₂ and its analog, MR-409, on the expression of IGF-1 in primary hepatocytes was investigated *in vitro*. Human or rat hepatocytes were treated with rhGH alone or in combination with GHRH(1-29)NH₂ or MR-409. As shown in the Figure 2A, in human hepatocytes, both GHRH(1-29)NH₂ and MR-409 inhibited the IGF-1 expression induced by treatment with rhGH (0.5 µg/ml). The peptides by themselves, however, shows no effect (Supplementary Figure 1A). In the human hepatocytes, IGF-1 expression stimulated by rhGH was decreased by 16.2±2.2% ($p < 0.05$), at 12 hours in the presence of 1 µM GHRH(1-29)NH₂, but the decrease at 24 hours (5.1±0.8%) and 48 hours (1.5±2.3%) was not significant. In the cells treated with rhGH and 1 µM MR-409, mRNA for IGF-1 decreased by 27.7±3.2% ($p < 0.01$) at 12 hours, by 19.7±2.3% ($p < 0.05$) at 24 hours, and by 13.6±3.7% ($p < 0.05$) at 48 hours, compared to cells treated with rhGH alone. In rat hepatocytes, an inhibition was observed as early as 2 hours after treatment by 1 µM MR-409 (Figure 2B). The mRNA for IGF-1 decreased significantly by 22.3±1.8% ($p < 0.01$) at 2 hours, 35.5±2.1% ($p < 0.01$) at 4 hours, 37.1±2.2% ($p < 0.01$) at 8 hours, and 37.2±3.6% ($p < 0.01$) at 24 hours. These results indicate a significant inhibition of IGF-1 synthesis in hepatocytes by GHRH(1-29)NH₂ and its analog, MR-409.

Interestingly, the expression of IGF-2 was also decreased by GHRH(1-29)NH₂ and MR-409 (Supplementary Figure 1B). In human hepatocytes treated by rhGH and GHRH(1-29)NH₂, the expression of IGF-2 was reduced by 11.9±3.4% ($p < 0.05$) at 12 hours, and 25.8±5.9% ($p < 0.05$) at 24 hours, compared to the cells treated only with rhGH. In human hepatocytes treated with both rhGH and 1 µM MR-409, the reduction in mRNA for IGF-2 was 23.9±3.7% ($p < 0.01$) at 12 hours, 22.3±2.9% ($p < 0.05$) at 24 hours, and 40.0±9.4% ($p < 0.01$) at 48 hours. In general, the MR-409 induced a greater and more protracted inhibitory effect than GHRH(1-29)NH₂. This may be linked to its much greater activity and the increased resistance to proteases of MR-409 (3), compared

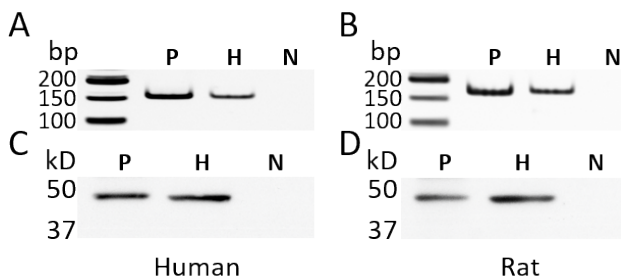


Figure 1: The expression of GHRH receptor (GHRH-R) in the primary human hepatocytes and rat hepatocytes. A., B. The PCR-based amplification of a fragment from GHRH-R cDNA. The size of the PCR product from the human cDNA (A) was 155 bp and from rat cDNA (B) 164 bp. C., D. Western blots using a rabbit polyclonal antibody (Abcam 76263) against human GHRH-R (C) and rat GHRH-R (D). P: human or rat pituitary, positive control; H: human or rat hepatocytes; N: negative control; The reaction without cDNA was used for the negative control of PCR. In the Western blots, the primary antibody against GHRH-R was replaced by normal rabbit IgG. bp: base pair; kD: kilodalton.

to GHRH(1-29)NH₂.

GHRH(1-29)NH₂ and its agonist MR-409 inhibit synthesis and release of IGF-1 *in vitro* from rat hepatocytes

To test the effects of GHRH(1-29)NH₂ and MR-409 on the synthesis and release of IGF-1 *in vitro*, rat hepatocytes were treated with 1 μg/ml of rhGH alone or in combination with 1 μM GHRH(1-29)NH₂ or MR-409. As shown in Figure 3A, MR-409 not only inhibited the secretion of IGF-1 by the hepatocytes, but also decreased the concentration of IGF-1 in the cells. Thus,

MR-409 significantly suppressed the release of IGF-1 induced by rhGH by 24.3±1.4% ($p < 0.01$) at 24 hours and by 21.1±1.2% ($p < 0.01$) at 48 hours. In addition, the intracellular IGF-1 level in the cells treated with MR-409 was 44.0±1.0% ($p < 0.01$) lower than in the cells exposed to rhGH alone. In a similar experiment using GHRH(1-29)NH₂, rat hepatocytes treated with both rhGH (1 μg/ml) and 1 μM GHRH(1-29)NH₂ secreted less IGF-1 compared to cells exposed only to rhGH. GHRH(1-29)NH₂ significantly decreased the levels of IGF-1 in the medium by 19.6±3.3% ($p < 0.05$) at 24 hours and by 24.4±1.3% ($p < 0.01$) at 48 hours. GHRH(1-29)NH₂ also lowered the intracellular concentration of IGF-1 by 23.1±1.1% ($p < 0.01$) (Figure 3B). Thus, both GHRH(1-29)NH₂ and

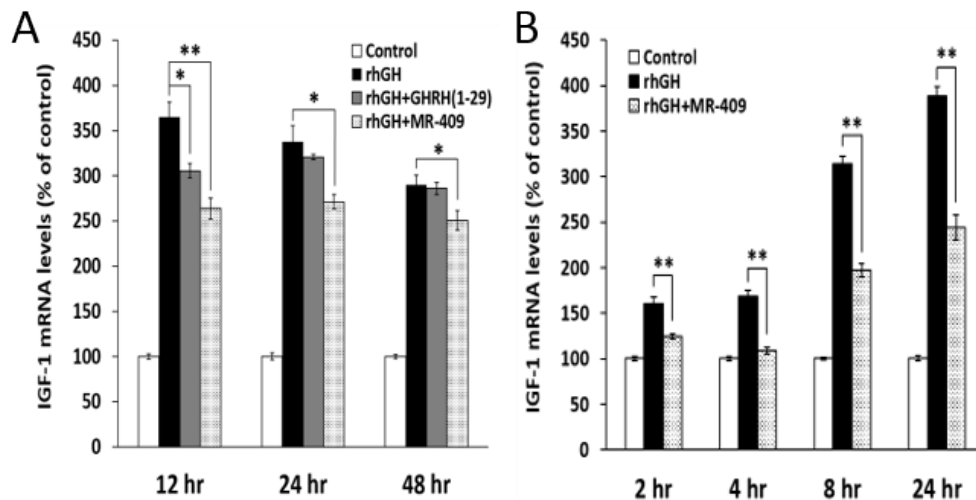


Figure 2: GHRH(1-29)NH₂ and its analog, MR-409, inhibit the expression of IGF-1 in primary hepatocytes. Human A. or rat B. hepatocytes were treated with either recombinant human growth hormone (rhGH) alone (0.5 μg/ml for human hepatocytes and 1 μg/ml for rat hepatocytes) or a combination of rhGH with 1 μM GHRH(1-29)NH₂ or MR-409. The RNAs were isolated at selected times and the levels of mRNA for IGF-1 were determined by quantitative RT-PCR. (Error bars indicate ±SEM; * $p < 0.05$, ** $p < 0.01$).

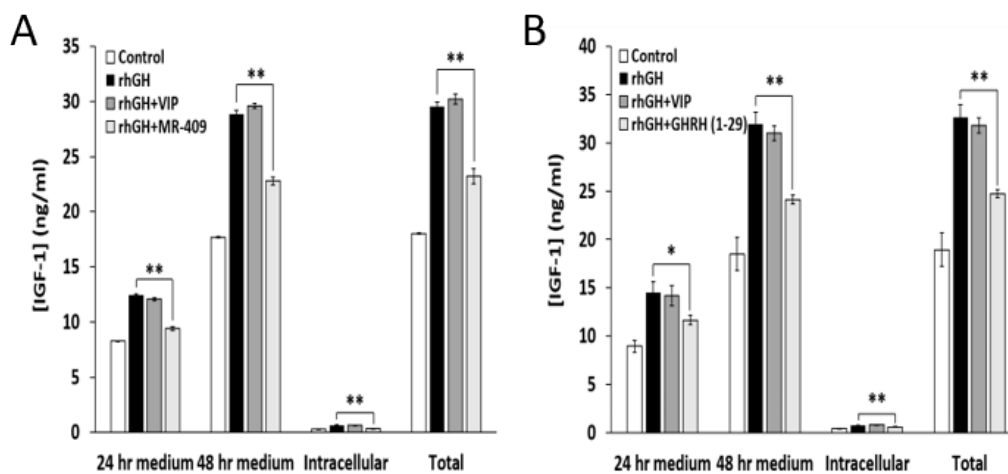


Figure 3: GHRH(1-29)NH₂ and MR-409, but not VIP, inhibit synthesis and release of IGF-1 by rat hepatocytes. Rat hepatocytes were treated by rhGH (1 μg/ml) alone or combined with A. 1 μM MR-409 or B. 1 μM GHRH(1-29)NH₂. The concentrations of IGF-1 in the cell culture medium and in the cells were measured at 24 and 48 hours respectively by ELISA. Both MR-409 and GHRH(1-29)NH₂ significantly inhibited the stimulatory effect of rhGH on the release of IGF-1. In both experiments (A) and (B), 1 μM VIP was tested as a peptide control and had no effect. (Error bars indicate ±SEM; * $p < 0.05$, ** $p < 0.01$).

Table 1: Effect of GHRH(1-29)NH₂ and its agonist MR-409 on the secretion of IGF-1 by cancer cells *in vitro*

Treatment	IGF-1 (pg/ml) ^a			
	MCF7 breast cancer		J82 bladder cancer	
	Measured level	Stimulation ^b	Measured level	Stimulation ^b
Control (no treatment)	529.9±53.5	-	240.3±24.7	-
rhGH (1 µg/ml)	1081.0±113.1*	551.1	365.5±45.0	125.2
rhGH (1 µg/ml) plus MR-409 (1 µM)	594.6±62.9*	64.7	284.6±30.5	44.3
rhGH (1 µg/ml) plus GHRH(1-29) (1 µM)	692.7±78.4*	162.8	310.7±31.6	70.4

a. All values were normalized to the total protein in the dishes. The results are representative of three independent experiments.

b. Values were calculated against the value of the control.

Compared to the control, the measured IGF-1 level was significantly increased in the MCF7 cells treated with rhGH, **p* < 0.05; MR-409 and GHRH(1-29) significantly decreased the IGF-1 secretion induced by rhGH, **p* < 0.05.

its agonist, MR-409, showed the capability to inhibit the synthesis and release of IGF-1 in rat hepatocytes treated with rhGH. In order to confirm the specificity of the inhibitory effects of GHRH(1-29)NH₂ and MR-409 on the release of IGF-1, the vasoactive intestinal peptide (VIP), containing 28 amino acid residues, was used as a peptide control. There is a structural similarity between VIP and GHRH as well as considerable homology between their receptor proteins, which belong to the family of G protein coupled receptors. As shown in Figure 3A and 3B, VIP showed no effect on the IGF-1 secretion induced by rhGH in rat hepatocytes in contrast to GHRH(1-29)NH₂ and GHRH agonist MR-409.

GHRH(1-29)NH₂ and MR-409 inhibit hepatic IGF-1 secretion in hypophysectomized rats

In the week prior to the experiment, the baseline levels of IGF-1 in the blood of hypophysectomized

rats were measured to be 58.8±8.5 ng/ml, indicating a successful elimination of the endocrine GH/IGF-1 axis. The animals were then treated daily for three days with rhGH, at a dose of 0.5 mg/kg bw, alone or together with the agonist MR-409, at 5 µg/25 g bw, as described in the legend to Figure 4. As shown in Figure 4A, rhGH strongly stimulated serum levels of IGF-1 immediately after the first injection, while MR-409 significantly inhibited this IGF-1 surge. The serum IGF-1 values were reduced by MR-409 from 410.6±17.4 ng/ml to 231.5±22.5 ng/ml, (*p* < 0.01), on day 1; from 605.8±43.9 ng/ml to 347.4±37.8 ng/ml, (*p* < 0.01), on day 2; and from 618.8±35.0 ng/ml to 396.4±28.6 ng/ml, (*p* < 0.01), on day 3. Compared to the effect of rhGH alone, the serum levels of IGF-1 in animals treated with both rhGH and MR-409 declined by 43.6±10.4% on day 1, 42.6±9.5% on day 2, and 36.0±7.8% on day 3. This result clearly demonstrates the inhibitory effect of MR-409 on the hepatic secretion of IGF-1 in response to the stimulus with GH.

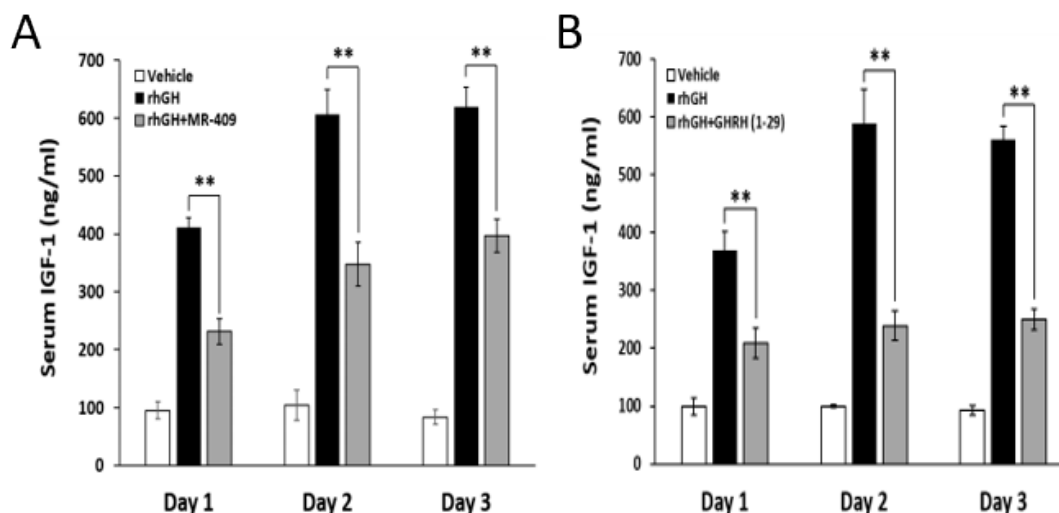


Figure 4: GHRH(1-29)NH₂ and MR-409 suppress the hepatic IGF-1 secretion induced by rhGH. Hypophysectomized rats (6 animals/group) were given daily i.p. injections of vehicle, rhGH alone or in combination with GHRH(1-29)NH₂ or MR-409 for three continuous days. Group I, received only the vehicle; Group II, was injected with rhGH (0.5 mg/kg bw); Group III, received rhGH (0.5 mg/kg bw) plus either A. MR-409 (5 µg/25 g bw) or B. GHRH(1-29)NH₂ (200 µg/25 g bw). Blood samples were collected at 24 hours after each injection. Serum IGF-1 was analyzed by ELISA. (Error bars indicate ±SEM; ***p* < 0.01).

In a parallel study, in which we used GHRH(1-29)NH₂, at a high dose of 200 µg/25 g bw, about 40 times greater on the weight and molecular basis than that of MR-409, GHRH(1-29)NH₂ diminished the rhGH-induced increase in IGF-1 levels by 43.3±7.1% (*p* < 0.01) on day 1, by 59.5±4.3% (*p* < 0.01) on day 2, and 55.4±3.2% (*p* < 0.01) on day 3 (Figure 4B). In our earlier pilot tests, GHRH(1-29)NH₂ at dosages lower than 50 µg/25 g bw failed to significantly lower IGF-1 levels in animals treated with rhGH (0.5 mg/kg bw). This is likely due to a very short half-life of GHRH(1-29)NH₂ in circulation [26].

MR-409 suppresses serum IGF-1 in C57BL/6 mice

The effect of MR-409 on serum IGF-1 was also tested in C57BL/6 mice. As shown in Figure 5A, daily injections of MR-409 significantly decreased serum IGF-1 from 24 hours on day 1 to day 5. At 24 hours, serum IGF-1 in mice treated with MR-409 was reduced from 382.1±12.1 ng/ml (in control mice) to 307.8±13.8 ng/ml (*p* < 0.05), and on day 5, from 396.5±14.4 ng/ml to 338.1±18.1 ng/ml, (*p* < 0.05). MR-409, however, showed a small stimulatory effect on serum IGF-1 within the first 6

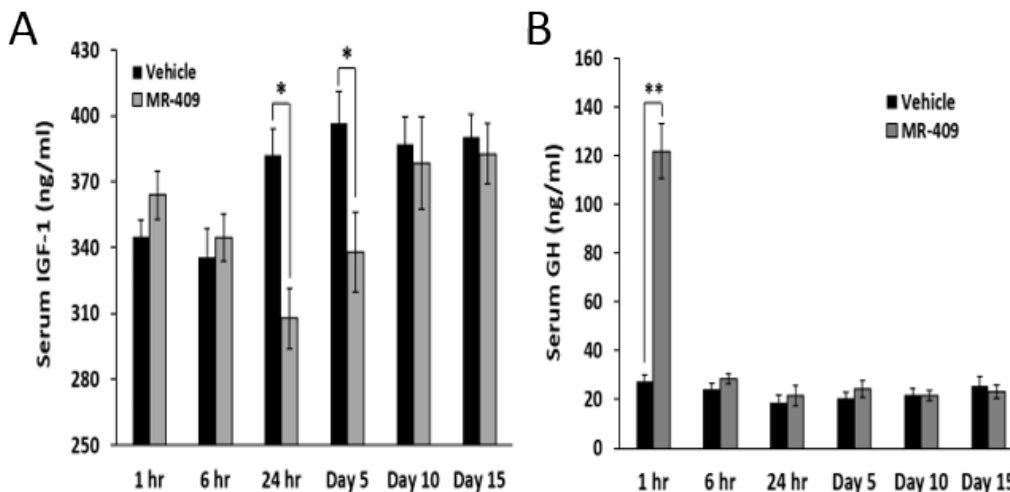


Figure 5: MR-409 suppresses the serum IGF-1 in C57BL/6 mice. Wild type C57BL/6 mice (8 animals/group) were treated daily with either 5 µg/day MR-409 s.c. or vehicle for 15 days. Blood or serum samples were collected at selected times and used for **A.** IGF-1 and **B.** GH ELISAs. (Error bars indicate ±SEM; **p* < 0.05, ***p* < 0.01).

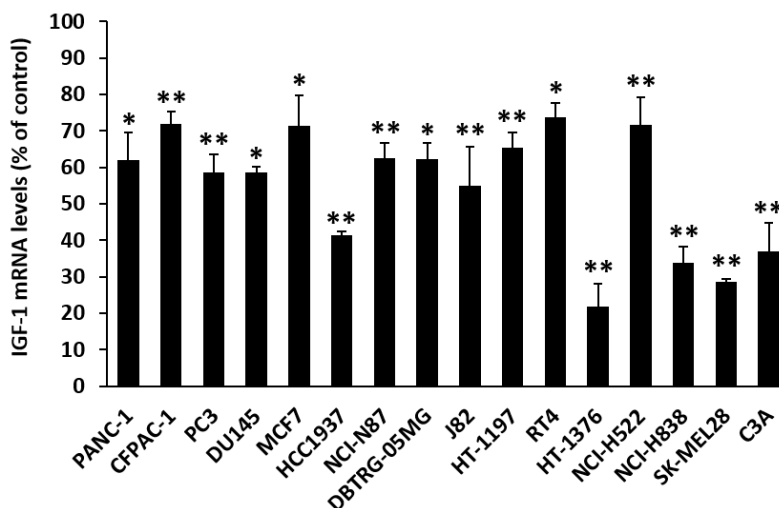


Figure 6: MR-409 down-regulates IGF-1 expression in cancer cells. Cancer cells were treated with 10-1000 nM MR-409 in respective serum-reduced (1% FBS) medium for 48 hours. RNAs were isolated and analyzed by quantitative real-time PCRs. The inhibition of IGF-1 expression for each cancer cell was shown as % of the control (Error bars indicate ±SEM; **p* < 0.05, ***p* < 0.01). (The definition of each cancer cell line is as follows; PANC-1, CFPAC-1: pancreatic cancer; PC3, DU145: prostate cancer; MCF7, HCC1937: breast cancer; NCI-N87: gastric cancer; DBTRG-05MG: glioblastoma; J82, HT-1197, RT4, HT-1376: bladder cancer; NCI-H522, NCI-H838, lung cancer; SK-MEL28: skin cancer, and C3A: liver cancer).

hours after the injection (not significant). After day 10, the serum levels of IGF-1 in the animals treated with MR-409 became similar to those in controls. The same sera were also used to measure the concentrations of GH. Except for the first hour, no significant difference in GH levels was found between the animals treated with MR-409 and the control (Figure 5B).

MR-409 down-regulates mRNA for IGF-1 in cancer cells

GHRH receptors are present in many tumors [1, 2, 27-33]. In order to determine whether GHRH agonist, MR-409, has an effect on the expression of IGF-1 in cancer cells, 16 human cancer cell lines were examined. The treatment of cancer cells consisted of *in vitro* exposure to concentrations of 10 nM to 1000 nM MR-409 for 48 hours as pre-determined by the cell proliferation assays. The mRNA levels for IGF-1 were analyzed at 12, 24, and 48 hours after the treatment. The effect on each cancer cell line is summarized in Figure 6. MR-409 exerted significant inhibitory effect on IGF-1 expression in all cancer cells tested. The mean mRNA level for IGF-1 was reduced by $46.0 \pm 16.5\%$, ($p < 0.05$) as compared to the control, the range being from $26.3 \pm 3.9\%$ to $78.2 \pm 6.3\%$. Treatment of cancer cells with GHRH(1-29)NH₂ also led to a decrease in mRNA levels for IGF-1. Thus GHRH(1-29)NH₂ at 10 nM-1000 nM concentration reduced the mRNA for IGF-1 in J82, HT-1197, and HT-1376 bladder cancer cells by $28.3 \pm 9.5\%$ ($p > 0.05$), $42.0 \pm 2.3\%$ ($p < 0.01$), and $19.0 \pm 4.2\%$ ($p < 0.01$), respectively; in CFPAC-1 and PANC-1 pancreatic cancer cells by $25.3 \pm 5.2\%$ ($p < 0.05$) and $13.5 \pm 10.5\%$ ($p > 0.05$), respectively; and in MCF7 breast cancer cells by $27.6 \pm 5.0\%$ ($p < 0.01$).

GHRH(1-29)NH₂ and MR-409 inhibit the secretion of IGF-1 by cancer cells

We then determined the effects of GHRH(1-29)NH₂ and its agonist, MR-409, on the secretion of IGF-1 by cancer cells *in vitro*. As summarized in Table 1, in both MCF7 breast cancer cells and J82 bladder cancer cells, the IGF-1 secretion was greatly stimulated by rhGH. The addition of MR-409 or GHRH(1-29)NH₂ to culture medium induced a strong inhibition of the response to rhGH in both MCF7 and J82 cancer cells. In breast cancer lines, the stimulated level of rhGH (551.1 pg/ml) was reduced by $88.2 \pm 11.4\%$ ($p < 0.05$) in the presence of MR-409 and by $70.4 \pm 14.2\%$ ($p < 0.05$) after exposure to GHRH(1-29)NH₂. In J82 bladder cancer cells, the stimulation induced by rhGH (125.2 pg/ml) was diminished by $64.6 \pm 24.4\%$ ($p > 0.05$) in the presence of MR-409 and by $43.8 \pm 12.0\%$ ($p > 0.05$) following the exposure to GHRH(1-29)NH₂.

MR-356 suppresses serum IGF-1 in nude mice bearing NCI-N87 stomach cancer

Nude mice (12 animals per group) bearing xenografts of NCI-N87 stomach cancer, were treated with daily s.c. injections of 5 µg/day of MR-356, a potent agonistic analog of GHRH(1-29)NH₂. Blood levels of IGF-1 were monitored throughout the treatment. As shown in Figure 7, serum levels of IGF-1 in the group treated with MR-356 were significantly lower than those of controls. The mean serum levels of IGF-1 in the treated group were reduced by $21.8 \pm 4.4\%$, ($p < 0.05$) on day 21, by $19.6 \pm 4.0\%$, ($p < 0.05$) on day 47, and by $17.9 \pm 9.2\%$ ($p > 0.05$) on day 63. In addition, the treatment with MR-356 markedly decreased tumor growth as shown by the significant reduction of the mean volume of gastric tumors in the treated group to 360 ± 111 mm³ on day 63 from that found in the control (1289 ± 416 mm³) ($p < 0.02$).

Effects of GHRH(1-29)NH₂ and its analog, MR-409, on the GH-induced GHR/JAK2/STAT5 signaling

The GH receptor (GHR)-mediated JAK2/STAT5 pathway is a major mechanism, which regulates the IGF-1 production induced by GH [34], while members of the GH-inducible suppressor of cytokine signaling (SOCS) family and cytokine-inducible SH2 (CISH) proteins negatively regulate the GH-induced activation of JAK2/STAT5 signaling [35, 36]. Consequently, we investigated the role that GHRH(1-29)NH₂ and its analog, MR-409, may play in these events. As shown in Figure 8A, both GHRH(1-29)NH₂ and MR-409, reversed the activation of GHR/JAK2/STAT5 signaling in rat hepatocytes induced by GH. The phosphorylation levels of the signaling molecules were all significantly decreased ($p < 0.05$).

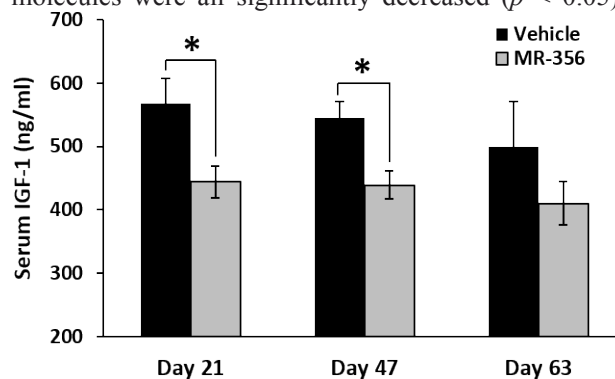


Figure 7: MR-356 suppresses serum IGF-1 in nude mice. Nude mice were xenografted with NCI-N87 stomach cancer. For the treatment, animals were injected s.c. with 5 µg MR-356/day for 9 weeks. The blood from the control and treated groups (12 mice/group) were collected at selected days. Concentrations of serum IGF-1 were measured by ELISA. (Error bars indicate \pm SEM; * $p < 0.05$).

Meanwhile, in the C3A human liver cancer cells, MCF-7 breast cancer cells and CFPAC-1 pancreatic cancer cells treated by MR-409, there was also a dramatic decrease in phosphorylation of GHR/JAK2/STAT5 (Figure 8B-8D). In addition, we evaluated the effects of GHRH(1-29)NH₂ and MR-409 on the SOCSs and CISH. Treatment of human hepatocytes with GHRH(1-29)NH₂ and MR-409 escalated the expression of SOCS1 by 192.6±19.7% (*p* < 0.05) and 206.3±17.6% (*p* < 0.01), respectively; mRNA for SOCS-2 by 122.4±22.5% (*p* > 0.05) and 146.9±29.9% (*p* < 0.05), respectively; and the expression of SOCS-3 by 156.9±2.3% (*p* < 0.05) and 196.9±24.4% (*p* < 0.05), respectively, compared to human hepatocytes exposed to rhGH. The expression of CISH, however, was not significantly changed (Supplementary Figure 2).

DISCUSSION

Hypothalamic GHRH regulates the release of GH from the pituitary, and in turn GH stimulates the secretion of IGF-1 from the liver [1, 17, 25, 37-39]. The regulation of the secretion of IGF-1 is incompletely understood and appears to be mediated by multiple mechanisms [20, 23-25, 37, 40, 41]. Since IGF-1 is under the control of GH, the suppression of the release of GH from the pituitary by antagonistic analogs of GHRH disrupts the pituitary GH/hepatic IGF-1 axis and leads to a reduction in the blood levels of GH, which in turn results in a lowering of circulating IGF-1 [1, 2, 20, 24, 37, 42]. This inhibitory action of GHRH antagonists on IGF-1 levels in serum has been demonstrated by our group in nude mice bearing

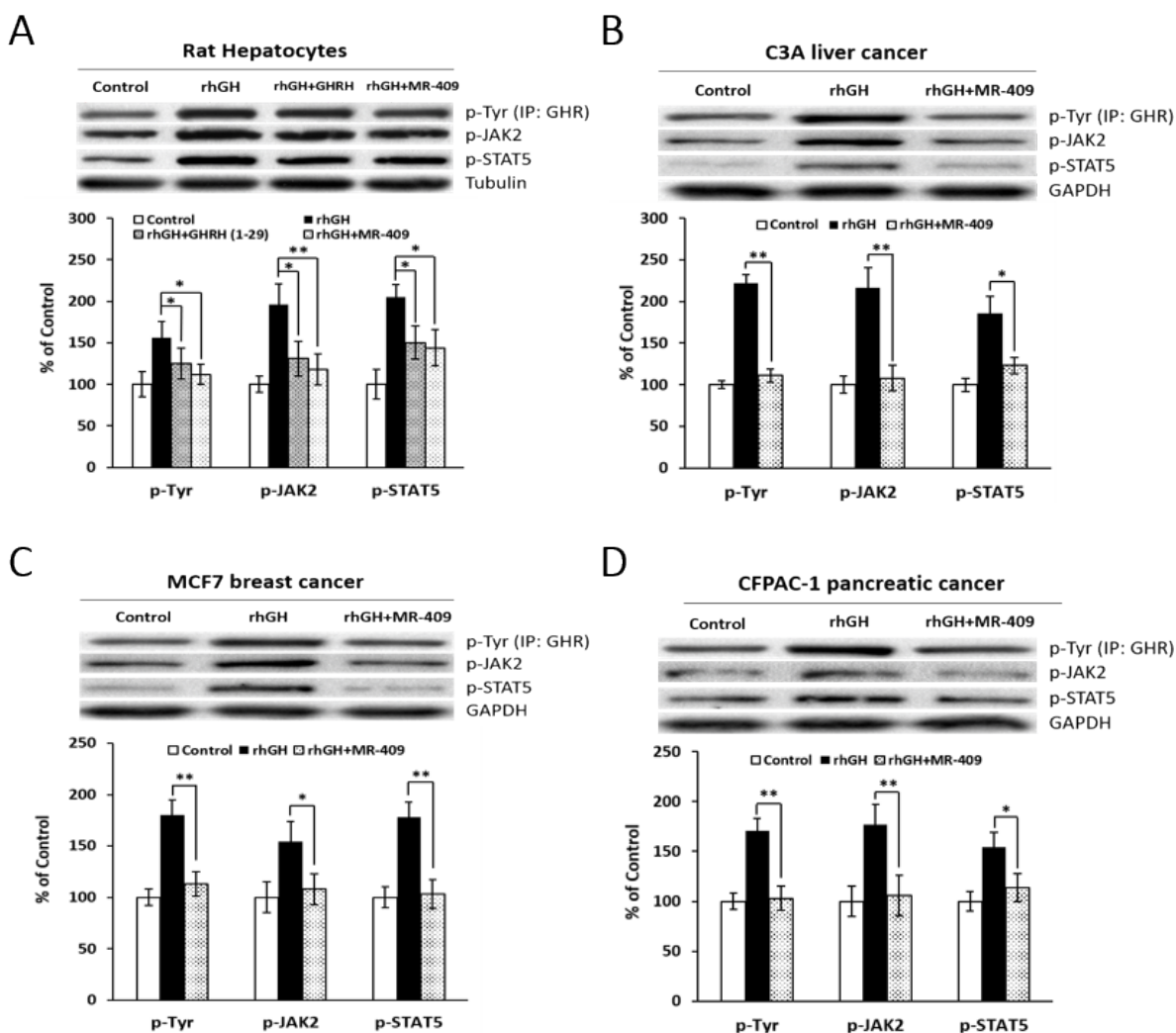


Figure 8: The effect of GHRH(1-29)NH₂ and MR-409 on GHR/JAK2/STAT5 signaling. A. Rat hepatocytes were treated by either 1 µg/ml rhGH alone or its combination with 1 µM GHRH(1-29)NH₂ or MR-409 in serum-free William's Medium E containing 0.1% BSA for 30 mins. The phosphorylation levels of GHR, JAK2, and STAT5 were analyzed by Western blots. B. C3A Human liver cancer cells, C. MCF7 breast cancer cells, and D. CFPAC-1 pancreatic cancer cells were treated by 1 µg/ml rhGH alone or a combination with 0.25 µM MR-409 for C3A and CFPAC-1, or 1 µM MR-409 for MCF7, in their respective serum-free basic medium for 30 mins. The protein levels for p-GHR, p-JAK2, and p-STAT5 were analyzed by Western blots. For quantification, expression levels of tubulin (for rat hepatocytes) or GAPDH (for human cancer cells) were used as references. (Error bars indicate ±SEM; **p* < 0.05; ***p* < 0.01).

various human cancers [1, 27, 31, 43-46]. Nevertheless, the main inhibitory effect of GHRH antagonists on tumor growth appears to be exerted through GHRH receptors which are present in various human cancers [1, 2, 27-33].

In elaborate studies, Gahete, Kineman et al. and others documented that IGF-1 inhibits the release and synthesis of GH *in vitro* and *in vivo* by a feedback action [20]. However, the involvement of a neurohormonal component in control of IGF-1 release, has not been demonstrated to date. Recently, in the evaluations of the effects of our GHRH agonists on normal tissues and tumors, we have detected their inhibitory action on IGF-1 secretion. To the best of our knowledge, this phenomenon has not been previously reported. The demonstration of the inhibitory effects of GHRH agonists on the secretion of IGF-1, described herein, appears to be therefore a relevant contribution to knowledge on the regulation of GH/IGF-1 axis.

IGF-1 is a potent growth factor for various cancer cells [14-16, 18, 19, 21, 22, 24, 47-49]. Various experimental and clinical studies have demonstrated the role of IGF-1 in growth and metastasis of many cancers [16, 18, 19, 21, 22, 24]. The findings of epidemiological studies expertly reviewed by Samani et al [16] also indicate that IGF-1 is involved in the development of cancers. These clinical investigations have revealed that high levels of serum IGF-1 and/or decreased concentrations of IGF-1 binding proteins are linked to an augmented risk for several malignancies [16], including breast cancer [19, 50, 51], prostate [18, 52], lung [53], and colorectal cancer [54-56]. In our studies on GHRH antagonists, we have previously shown an association between the levels of serum or tumoral IGF-1 and tumor growth [27, 43-46]. We were aware that the reduction in the levels of IGF-1 was mostly induced by the inhibition of GH secretion by the GHRH antagonists. Our findings that GHRH(1-29)NH₂ and its agonist, MR-409, can also reduce the secretion of hepatic and tumoral IGF-1 are unexpected. Nevertheless, the observations that the GHRH(1-29)NH₂ and GHRH agonist, MR-409, inhibited the expression of mRNA for IGF-1 as well as IGF-1 secretion by human and rat hepatocytes and human cancer cells *in vitro*, after stimulation with rhGH, imply that GHRH agonists may exert these effects by direct action on the cells. The specificity of the effect of GHRH and its agonists was demonstrated by the fact that vasoactive intestinal peptide, structurally similar to GHRH, showed no effect. GHRH(1-29)NH₂ and agonist MR-409 lowered serum IGF-1 levels in hypophysectomized rats injected with rhGH. These results and findings that GHRH agonists, MR-356 or MR-409, reduced serum IGF-1 levels in nude mice xenografted with human NCI-N87 stomach cancer as well as in C57BL/6 mice show that the inhibitory effects occur also *in vivo*. The inhibition of tumor growth which accompanied the treatment of NCI-N87 stomach cancer and other tumors by the GHRH agonists *in vivo*

is the subject of our intense investigations, which will be reported elsewhere.

The mechanisms through which GHRH(1-29)NH₂ and its agonist, MR-409, suppress IGF-1 production and release in hepatic and cancer cells is not completely understood. In this study, wild type C57BL/6 mice treated with MR-409 demonstrated a decrease in serum IGF-1 at 24 hours after the injection of the GHRH agonist. The GH levels, however, increased only within the first hour post injection and subsequently returned to the baseline levels 6 hours after injection. Thus, MR-409 showed only a transient stimulatory effect on GH release. These observations, suggest that in animals treated with GHRH agonists under these experimental conditions, serum IGF-1 may not be related to a fall in circulating GH levels. This action is different from that of GHRH antagonists which decrease the release of GH from pituitary somatotropes after binding to the GHRH receptors and block the production and release of GH [1, 37]. The decrease in circulating IGF-1 is then a consequence of blocking the secretion of GH as stated above. Such a mechanism, however, is not applicable to the GHRH agonists. Instead, the secretion of IGF-1 appears to be suppressed by a direct action of GHRH and its agonists on the hepatocytes and the tumors. This is supported by our studies using hepatocytes *in vitro* as well as hypophysectomized animals. In both cases, GHRH(1-29)NH₂ and MR-409 suppressed the IGF-1 secretion stimulated by rhGH.

Incidentally, these are not the first findings of direct effects of GHRH agonists on extra-pituitary cells/tissues. GHRH(1-29)NH₂ and its agonists have been demonstrated to exert direct stimulatory effects on cardiovascular endothelial cells [4-6], pancreatic β cells [7, 8], and fibroblasts [9]. These observations are in accord with direct effects of GHRH and its agonists on hepatocytes and cancer cells as found in our studies. In an effort to determine the mechanisms of action for these observations, we investigated whether the GH receptor (GHR)-mediated JAK2/STAT5 signaling transduction could be involved. In the course of stimulation with the GH, GHR/JAK2/STAT5 pathway plays a critical role in hepatic synthesis and release of IGF-1 [34, 57]. In cancer cells, this pathway was also found to be associated with tumoral expression of mRNA for IGF-1 [58] and the stimulation of proliferation [59] and survival of cancer cells [60]. We demonstrated that GHRH(1-29)NH₂ and its agonist, MR-409, can suppress the GH-induced activation of GHR/JAK2/STAT5 pathway, which suggests that this mechanism of action is different from that of GHRH antagonists. Thus, the anti-neoplastic effect of GHRH antagonist, MIA-602, was found to be associated with blockage of GHRH-R-mediated PAK1/STAT3/NF- κ B pathways in gastric cancers [28]. One possible explanation could be that the actions of GHRH agonists on the hepatocytes may involve a hypothetical unidentified factor, whose recruitment to the GH/GHR complex, after binding of GH to GHR, could be

required for the activation of GHR and/or the maintenance of its kinase activity for downstream signaling. When GHRH or its agonists bind to the GHRH receptor, this leads to the dissociation of this unknown factor from the GHR, resulting in a quiescent complex with restrained capability of making transduction signals for the synthesis and release of IGF-1. This hypothesis, however, does not exclude a possibility that GHRH agonists may also bind to an unknown homologous G-protein coupled receptor and directly inhibit the IGF-1 production in both hepatic and tumor cells as proposed by Kineman for the actions of GHRH antagonists [37].

IGF-1 exerts its functions through the IGF-1 receptor (IGF1-R) [24]. In our preliminary studies on J82 bladder cancer, GHRH agonist MR-409 downregulated the expression of IGF1-R, inhibited the phosphorylation of IGF1-R and its downstream AKT and ERK pathways. In addition, GHRH(1-29)NH₂ and MR-409 also induced an inhibition of mRNA for IGF-2 in human hepatocytes. An investigation on the effect of GHRH and its agonists on IGF-2 production may be also worth considering.

A major challenge to the use of GHRH *in vivo* has been its short half-life due to an enzymatic cleavage at the N-terminal [26]. To overcome this problem, various GHRH agonists, designated as the “MR” series, have been synthesized and evaluated by our group [3]. Some of these agonists, such as MR-409 and MR-356, showed significantly greater potency, higher binding affinities, and longer half-life *in vivo* due to increased resistance to degradation by proteases, compared to GHRH(1-29)NH₂ [3]. Their effects on various cells/tissues have been reported in our previous publications [3-9, 61].

It is difficult to envision that GHRH could be involved in the pathophysiologic control of IGF-1 secretion since the circulating levels of hypothalamic GHRH are very low. In contrast, the levels of GHRH in the tumors can be very high [1, 38, 39]. The studies on this complex topic would require further extensive investigations. The results reported herein reveal the inhibitory effects of GHRH(1-29)NH₂ and its agonists on the hepatic and tumoral secretion of IGF-1, which occurred under the conditions described.

MATERIALS AND METHODS

Peptide and chemicals

Synthesis and purification of MR-409 ((N-Me-Tyr¹, D-Ala², Orn^{12,21}, Abu¹⁵, Nle²⁷, Asp²⁸)-GHRH(1-29)NHCH₃) and MR-356 ((N-Me-Tyr¹, Ala², Phe⁶, Gln⁸, Orn¹², Abu¹⁵, Orn²¹, Nle²⁷, Asp²⁸, Arg²⁹)-GHRH(1-29)NH₂) have been previously described [3]. Human GHRH(1-29)NH₂ was supplied by AnaSpec (Fremont, CA). All other chemicals, if not specified, were purchased from Sigma-Aldrich (St.

Louis, MO, USA).

Experiments using hepatocytes

(I) Human and rat hepatocytes were treated with 0.5 µg/ml or 1 µg/ml rhGH respectively, or the combination of rhGH with GHRH(1-29)NH₂ or its analog MR-409 (1 µM) in serum-free William's Medium E containing 0.1% BSA. RNAs were isolated at selected times. The amounts of mRNA for target genes were analyzed by quantitative RT-PCR. (II) Rat hepatocytes (1×10⁶ cells/6-well) were treated with rhGH (1 µg/ml) alone or in combination with 1 µM GHRH(1-29)NH₂ or MR-409. The culture media at 24 and 48 hours were collected for IGF-1 ELISA. Cell lysates at 48 hours were used for measurement of intracellular IGF-1 levels.

IGF-1 secretion by cancer cells *in vitro*

Cancer cells (5×10⁶ cells per 10 cm culture dish) were pre-starved in serum-free medium for 6 hours, and then treated with 1 µg/ml rhGH alone or combined with 1 µM MR-409 or GHRH(1-29)NH₂ for 48 hours. Cell culture media were then freeze-dried and reconstituted in distilled water. The salt was removed by using Amicon Ultra-2 column (Millipore). The levels of IGF-1 in the samples were measured by the Human IGF-1 Quantikine ELISA kit (R&D Systems).

Animal studies *in vivo*

(I) Wild-type C57BL/6 mice (9 weeks old, 8 animals/group) were treated daily with 5 µg/day MR-409 s.c. or the vehicle for 15 days. Blood samples were collected from the tail vein at selected times. Serum IGF-1 and GH were determined by ELISAs. (II) Male Wistar Han IGS rats (CrI:WI), 6-weeks old, were hypophysectomized by Charles River Laboratories. Two weeks later, animals were then divided into three groups of six animals each, and were injected i.p. daily with rhGH at 0.5 mg/kg bw, or rhGH plus MR-409 at 5 µg/25 g bw, for three consecutive days. Control animals received the vehicle only. In a similar experimental setting, GHRH(1-29)NH₂ (200 µg/25 g bw) was used instead of MR-409. At 24 hours after each injection, blood samples were collected for measurement of serum IGF-1. (III) Female athymic nude mice, bearing xenografted NCI-N87 human stomach cancers, were assigned to two groups of 12 mice each. Experimental group received 5 µg/day of MR-356 s.c. for 9 weeks and the control group the vehicle only (10% 1,2-propanediol). Tumor size was monitored on a weekly basis and blood samples were collected for measurement of IGF-1.

Statistical analysis

Data are shown as mean \pm SEM. To determine statistical significance, one-way ANOVA followed by unpaired two-tailed Student's *t*-test were used (Origin v6.0). Differences were considered significant if $p < 0.05$.

More detailed information on materials and methods is provided in Supplementary Materials.

Author contributions

TC and AVS designed research; TC performed research; TC and AVS analyzed the data; AVS contributed new reagents; TC and AVS wrote the paper.

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CONFLICTS OF INTEREST

Andrew V. Schally is listed as a co-inventor on the patent for GHRH agonists, assigned to the University of Miami and the Veterans Affairs. However, the investigation of the effects of GHRH and its agonists on IGF-1 was an academic endeavor without any commercial interests. Tengjiao Cui declares no conflict of interest.

REFERENCES

- Schally AV, Varga JL, Engel JB. Antagonists of growth-hormone-releasing hormone: an emerging new therapy for cancer. *Nat Clin Pract Endocrinol Metab.* 2008; 4:33-43.
- Zarandi M, Cai R, Kovacs M, Popovics P, Szalontay L, Cui T, Sha W, Jaszberenyi M, Varga J, Zhang X, Block NL, Rick FG, Halmos G, Schally AV. Synthesis and structure-activity studies on novel analogs of human growth hormone releasing hormone (GHRH) with enhanced inhibitory activities on tumor growth. *Peptides.* 2017; 89:60-70.
- Cai R, Schally AV, Cui T, Szalontay L, Halmos G, Sha W, Kovacs M, Jaszberenyi M, He J, Rick FG, Popovics P, Kanashiro-Takeuchi R, Hare JM, et al. Synthesis of new potent agonistic analogs of growth hormone-releasing hormone (GHRH) and evaluation of their endocrine and cardiac activities. *Peptides.* 2014; 52:104-12.
- Kanashiro-Takeuchi RM, Tziomalos K, Takeuchi LM, Treuer AV, Lamirault G, Dulce R, Hurtado M, Song Y, Block NL, Rick F, Klukovits A, Hu Q, Varga JL, et al. Cardioprotective effects of growth hormone-releasing hormone agonist after myocardial infarction. *Proc Natl Acad Sci U S A.* 2010; 107:2604-9.
- Kanashiro-Takeuchi RM, Takeuchi LM, Rick FG, Dulce R, Treuer AV, Florea V, Rodrigues CO, Paulino EC, Hatzistergos KE, Selem SM, Gonzalez DR, Block NL, Schally AV, Hare JM. Activation of growth hormone releasing hormone (GHRH) receptor stimulates cardiac reverse remodeling after myocardial infarction (MI). *Proc Natl Acad Sci U S A.* 2012; 109:559-63.
- Gesmunro I, Miragoli M, Carullo P, Trovato L, Larcher V, Di Pasquale E, Brancaccio M, Mazzola M, Villanova T, Sorge M, Taliano M, Gallo MP, Alloati G, et al. Growth hormone-releasing hormone attenuates cardiac hypertrophy and improves heart function in pressure overload-induced heart failure. *Proc Natl Acad Sci U S A.* 2017; 114:12033-12038.
- Zhang X, Cui T, He J, Wang H, Cai R, Popovics P, Vidaurre I, Sha W, Schmid J, Ludwig B, Block NL, Bornstein SR, Schally AV. Beneficial effects of growth hormone-releasing hormone agonists on rat INS-1 cells and on streptozotocin-induced NOD/SCID mice. *Proc Natl Acad Sci U S A.* 2015; 112:13651-6.
- Ludwig B, Ziegler CG, Schally AV, Richter C, Steffen A, Jabs N, Funk RH, Brendel MD, Block NL, Ehrhart-Bornstein M, Bornstein SR. Agonist of growth hormone-releasing hormone as a potential effector for survival and proliferation of pancreatic islets. *Proc Natl Acad Sci U S A.* 2010; 107:12623-8.
- Cui T, Jimenez JJ, Block NL, Badiavas EV, Rodriguez-Menocal L, Vila Granda A, Cai R, Sha W, Zarandi M, Perez R, Schally AV. Agonistic analogs of growth hormone releasing hormone (GHRH) promote wound healing by stimulating the proliferation and survival of human dermal fibroblasts through ERK and AKT pathways. *Oncotarget.* 2016; 7:52661-52672.
<https://doi.org/10.18632/oncotarget.11024>.
- Salmon WD Jr, Daughaday WH. A hormonally controlled serum factor which stimulates sulfate incorporation by cartilage *in vitro*. *J Lab Clin Med.* 1957; 49:825-36.
- Laron Z. Laron syndrome (primary growth hormone resistance or insensitivity): the personal experience 1958-2003. *J Clin Endocrinol Metab.* 2004; 89:1031-44.
- Baker J, Liu JP, Robertson EJ, Efstratiadis A. Role of insulin-like growth factors in embryonic and postnatal growth. *Cell.* 1993; 75:73-82.
- Elders MJ, Garland JT, Daughaday WA, Fisher DA, Whitney JE, Hughes ER. Laron's dwarfism: studies on the nature of the defect. *J Pediatr.* 1973; 83:253-63.
- Macaulay VM. Insulin-like growth factors and cancer. *Br J Cancer.* 1992; 65:311-20.

15. Pollak MN, Schernhammer ES, Hankinson SE. Insulin-like growth factors and neoplasia. *Nat Rev Cancer*. 2004; 4:505-18.
16. Samani AA, Yakar S, LeRoith D, Brodt P. The role of the IGF system in cancer growth and metastasis: overview and recent insights. *Endocr Rev*. 2007; 28:20-47.
17. Lund PK. *Handbook of Physiology: Hormonal control of growth*, eds. Kostyo JL, Goodman HM. (Oxford Univ. Press, New York). 1999; 537-571.
18. Chan JM, Stampfer MJ, Giovannucci E, Gann PH, Ma J, Wilkinson P, Hennekens CH, Pollak M. Plasma insulin-like growth factor-I and prostate cancer risk: a prospective study. *Science*. 1998; 279:563-6.
19. Hankinson SE, Willett WC, Colditz GA, Hunter DJ, Michaud DS, Deroo B, Rosner B, Speizer FE, Pollak M. Circulating concentrations of insulin-like growth factor-I and risk of breast cancer. *Lancet*. 1998; 351:1393-6.
20. Gahete MD, Córdoba-Chacón J, Lin Q, Brüning JC, Kahn CR, Castaño JP, Christian H, Luque RM, Kineman RD. Insulin and IGF-I inhibit GH synthesis and release *in vitro* and *in vivo* by separate mechanisms. *Endocrinology*. 2013; 154:2410-20.
21. Gallagher EJ, LeRoith D. Minireview: IGF, Insulin, and Cancer. *Endocrinology*. 2011; 152:2546-51.
22. LeRoith D. IGF-I: panacea or poison? *J Clin Endocrinol Metab*. 2010; 95:4549-51.
23. Faje AT, Barkan AL. Basal, but not pulsatile, growth hormone secretion determines the ambient circulating levels of insulin-like growth factor-I. *J Clin Endocrinol Metab*. 2010; 95:2486-91.
24. Humbel RE. Insulin-like growth factors I and II. *Eur J Biochem*. 1990; 190:445-62.
25. Clemmons DR. Modifying IGF1 activity: an approach to treat endocrine disorders, atherosclerosis and cancer. *Nat Rev Drug Discov*. 2007; 6:821-33.
26. Frohman LA, Downs TR, Williams TC, Heimer EP, Pan YC, Felix AM. Rapid enzymatic degradation of growth hormone-releasing hormone by plasma *in vitro* and *in vivo* to a biologically inactive product cleaved at the NH2 terminus. *J Clin Invest*. 1986; 78:906-13.
27. Braczkowski R, Schally AV, Plonowski A, Varga JL, Groot K, Krupa M, Armatis P. Inhibition of proliferation in human MNNG/HOS osteosarcoma and SK-ES-1 Ewing sarcoma cell lines *in vitro* and *in vivo* by antagonists of growth hormone-releasing hormone: effects on insulin-like growth factor II. *Cancer*. 2002; 95:1735-45.
28. Gan J, Ke X, Jiang J, Dong H, Yao Z, Lin Y, Lin W, Wu X, Yan S, Zhuang Y, Chu WK, Cai R, Zhang X, et al. Growth hormone-releasing hormone receptor antagonists inhibit human gastric cancer through downregulation of PAK1-STAT3/NF- κ B signaling. *Proc Natl Acad Sci U S A*. 2016; 113:14745-14750.
29. Klukovits A, Schally AV, Szalontay L, Vidaurre I, Papadia A, Zarandi M, Varga JL, Block NL, Halmos G. Novel antagonists of growth hormone-releasing hormone inhibit growth and vascularization of human experimental ovarian cancers. *Cancer*. 2012; 118:670-80.
30. Fahrenholtz CD, Rick FG, Garcia MI, Zarandi M, Cai RZ, Block NL, Schally AV, Burnstein KL. Preclinical efficacy of growth hormone-releasing hormone antagonists for androgen-dependent and castration-resistant human prostate cancer. *Proc Natl Acad Sci U S A*. 2014; 111:1084-9.
31. Pópulo H, Nunes B, Sampaio C, Batista R, Pinto MT, Gaspar TB, Miranda-Alves L, Cai RZ, Zhang XY, Schally AV, Sobrinho-Simões M, Soares P. Inhibitory effects of antagonists of growth hormone-releasing hormone (GHRH) in thyroid cancer. *Horm Cancer*. 2017; 8:314-324.
32. Chu WK, Law KS, Chan SO, Yam JC, Chen LJ, Zhang H, Cheung HS, Block NL, Schally AV, Pang CP. Antagonists of growth hormone-releasing hormone receptor induce apoptosis specifically in retinoblastoma cells. *Proc Natl Acad Sci U S A*. 2016; 113:14396-14401.
33. Jimenez JJ, DelCanto GM, Popovics P, Perez A, Vila Granda A, Vidaurre I, Cai RZ, Rick FG, Swords RT, Schally AV. A new approach to the treatment of acute myeloid leukaemia targeting the receptor for growth hormone-releasing hormone. *Br J Haematol*. 2018; 181:476-485.
34. Woelfle J, Billiard J, Rotwein P. Acute control of insulin-like growth factor-I gene transcription by growth hormone through Stat5b. *J Biol Chem*. 2003; 278:22696-702.
35. Ram PA, Waxman DJ. SOCS/CIS protein inhibition of growth hormone-stimulated STAT5 signaling by multiple mechanisms. *J Biol Chem*. 1999; 274:35553-61.
36. Hansen JA, Lindberg K, Hilton DJ, Nielsen JH, Billestrup N. Mechanism of inhibition of growth hormone receptor signaling by suppressor of cytokine signaling proteins. *Mol Endocrinol*. 1999; 13:1832-43.
37. Kineman RD. Antitumorigenic actions of growth hormone-releasing hormone antagonists. *Proc Natl Acad Sci U S A*. 2000; 97:532-4.
38. Schally AV, Varga JL. Antagonistic analogs of growth hormone-releasing hormone: new potential antitumor agents. *Trends Endocrinol Metab*. 1999; 10:383-91.
39. Frohman LA, Kineman RD. *Handbook of Physiology: Hormonal control of growth*, eds. Kostyo, JL, Goodman, HM. (Oxford Univ. Press, New York). 1999; 189-221.
40. Laron Z. Insulin-like growth factor 1 (IGF-1): a growth hormone. *Mol Pathol*. 2001; 54:311-6.
41. Daughaday WH, Rotwein P. Insulin-like growth factors I and II. Peptide, messenger ribonucleic acid and gene structures, serum, and tissue concentrations. *Endocr Rev*. 1989; 10:68-91.
42. Yakar S, Liu JL, Fernandez AM, Wu Y, Schally AV, Frystyk J, Chernausk SD, Mejia W, Le Roith D. Liver-specific igf-1 gene deletion leads to muscle insulin insensitivity. *Diabetes*. 2001; 50:1110-8.
43. Jungwirth A, Schally AV, Pinski J, Halmos G, Groot

- K, Armatis P, Vadillo-Buenfil M. Inhibition of *in vivo* proliferation of androgen-independent prostate cancers by an antagonist of growth hormone-releasing hormone. *Br J Cancer*. 1997; 75:1585-92.
44. Jungwirth A, Schally AV, Pinski J, Groot K, Armatis P, Halmos G. Growth hormone-releasing hormone antagonist MZ-4-71 inhibits *in vivo* proliferation of Caki-I renal adenocarcinoma. *Proc Natl Acad Sci U S A*. 1997; 94:5810-3.
 45. Pinski J, Schally A, Jungwirth A, Groot K, Halmos G, Armatis P, Zarandi M, Vadillo-Buenfil M. Inhibition of growth of human small cell and non-small cell lung carcinomas by antagonists of growth hormone-releasing hormone (GH-RH). *Int J Oncol*. 1996; 9:1099-105.
 46. Pinski J, Schally AV, Groot K, Halmos G, Szepeshazi K, Zarandi M, Armatis P. Inhibition of growth of human osteosarcomas by antagonists of growth hormone-releasing hormone. *J Natl Cancer Inst*. 1995; 87:1787-94.
 47. Pollak M. Insulin and insulin-like growth factor signalling in neoplasia. *Nat Rev Cancer*. 2008; 8:915-28.
 48. Yu H, Rohan T. Role of the insulin-like growth factor family in cancer development and progression. *J Natl Cancer Inst*. 2000; 92:1472-89.
 49. Chhabra Y, Waters MJ, Brooks AJ. Role of the growth hormone IGF-1 axis in cancer. *Expert Rev Endocrinol Metab*. 2011; 6:71-84.
 50. Toniolo P, Bruning PF, Akhmedkhanov A, Bonfrer JM, Koenig KL, Lukanova A, Shore RE, Zeleniuch-Jacquotte A. Serum insulin-like growth factor-I and breast cancer. *Int J Cancer*. 2000; 88:828-32.
 51. Allen NE, Roddam AW, Allen DS, Fentiman IS, Dos Santos Silva I, Peto J, Holly JM, Key TJ. A prospective study of serum insulin-like growth factor-I (IGF-I), IGF-II, IGF-binding protein-3 and breast cancer risk. *Br J Cancer*. 2005; 92:1283-7.
 52. Stattin P, Bylund A, Rinaldi S, Biessy C, Déchaud H, Stenman UH, Egevad L, Riboli E, Hallmans G, Kaaks R. Plasma insulin-like growth factor-I, insulin-like growth factor-binding proteins, and prostate cancer risk: a prospective study. *J Natl Cancer Inst*. 2000; 92:1910-7.
 53. Yu H, Spitz MR, Mistry J, Gu J, Hong WK, Wu X. Plasma levels of insulin-like growth factor-I and lung cancer risk: a case-control analysis. *J Natl Cancer Inst*. 1999; 91:151-6.
 54. Probst-Hensch NM, Yuan JM, Stanczyk FZ, Gao YT, Ross RK, Yu MC. IGF-1, IGF-2 and IGFBP-3 in prediagnostic serum: association with colorectal cancer in a cohort of Chinese men in Shanghai. *Br J Cancer*. 2001; 85:1695-9.
 55. Palmqvist R, Hallmans G, Rinaldi S, Biessy C, Stenling R, Riboli E, Kaaks R. Plasma insulin-like growth factor 1, insulin-like growth factor binding protein 3, and risk of colorectal cancer: a prospective study in Northern Sweden. *Gut*. 2002; 50:642-6.
 56. Ma J, Pollak MN, Giovannucci E, Chan JM, Tao Y, Hennekens CH, Stampfer MJ. Prospective study of colorectal cancer risk in men and plasma levels of insulin-like growth factor (IGF)-I and IGF-binding protein-3. *J Natl Cancer Inst*. 1999; 91:620-5.
 57. Herrington J, Smit LS, Schwartz J, Carter-Su C. The role of STAT proteins in growth hormone signaling. *Oncogene*. 2000; 19:2585-97.
 58. Gan Y, Buckels A, Liu Y, Zhang Y, Paterson AJ, Jiang J, Zinn KR, Frank SJ. Human GH receptor-IGF-1 receptor interaction: implications for GH signaling. *Mol Endocrinol*. 2014; 28:1841-54.
 59. Koppikar P, Lui VW, Man D, Xi S, Chai RL, Nelson E, Tobey AB, Grandis JR. Constitutive activation of signal transducer and activator of transcription 5 contributes to tumor growth, epithelial-mesenchymal transition, and resistance to epidermal growth factor receptor targeting. *Clin Cancer Res*. 2008; 14:7682-90.
 60. Ahonen TJ, Xie J, LeBaron MJ, Zhu J, Nurmi M, Alanen K, Rui H, Nevalainen MT. Inhibition of Transcription factor Stat5 induces cell death of human prostate cancer cells. *J Biol Chem*. 2003; 278:27287-92.
 61. Thounaojam MC, Powell FL, Patel S, Gutsaeva DR, Tawfik A, Smith SB, Nussbaum J, Block NL, Martin PM, Schally AV, Bartoli M. Protective effects of agonists of growth hormone-releasing hormone (GHRH) in early experimental diabetic retinopathy. *Proc Natl Acad Sci U S A*. 2017; 114:13248-13253.