

Growth Hormone Secretagogue Binding Sites in Peripheral Human Tissues*

MAURO PAPOTTI, CORRADO GHÈ, PAOLA CASSONI, FILOMENA CATAPANO,
ROMANO DEGHENGI, EZIO GHIGO, AND GIAMPIERO MUCCIOLI

Departments of Biomedical Sciences and Oncology (M.P., P.C.), Anatomy, Pharmacology and Forensic Medicine (C.G., F.C., G.M.) and Internal Medicine (E.G.), University of Turin, Turin, 10125 Italy; and Europeptides (R.D.), Argensteuil 95108, France

ABSTRACT

The family of GH secretagogues (GHS) includes peptidyl (hexarelin) and nonpeptidyl (MK 0677) molecules possessing specific receptors in the brain, pituitary, and thyroid. GHS receptor subtypes have also been identified in the heart; and a gastric-derived peptide, named ghrelin, has recently been proposed as a natural ligand. Our aim was to investigate the presence of GHS receptors in a wide range of human tissues, by radioreceptor assay with [¹²⁵I]Tyr-Ala-hexarelin. GHS receptors were detected mainly in the myocardium, but they were also present (in order of decreasing binding activity) in adrenal, gonads, arteries, lung, liver, skeletal muscle, kidney, pituitary, thyroid, adipose tissue, veins, uterus, skin, and lymphnode. In contrast,

negligible binding was found in parathyroid, pancreas, placenta, mammary gland, prostate, salivary gland, stomach, colon, and spleen. Hexarelin, MK 0677, and human ghrelin completely displaced the radioligand from binding sites of endocrine tissues, but MK 0677 and ghrelin were less potent than hexarelin. In nonendocrine tissues, both MK 0677 and ghrelin were inactive in displacement of [¹²⁵I]Tyr-Ala-hexarelin, whereas hexarelin was as active as a displacing agent in endocrine tissues. This study provides the first detailed analysis of the tissue localization of GHS receptors and suggests that a still unknown receptor subtype, specific for peptidyl GHS, may exist in the heart and in other tissues. (*J Clin Endocrinol Metab* 85: 3803–3807, 2000)

GH SECRETAGOGUES (GHS) are synthetic, peptidyl [GH-releasing peptides (GHRPs)], and nonpeptidyl molecules that possess strong, dose-dependent, and reproducible GH-releasing activity *in vivo* in several species and in man. They are active by the iv, sc, intranasal route and even after oral administration (1–3). Both peptidyl and nonpeptidyl compounds also possess significant PRL- and ACTH/cortisol-releasing effect (4, 5). The neuroendocrine activities of GHRPs are mediated by specific receptors, which have originally been identified in the pituitary and the hypothalamus in humans (6), as well as in rats (7, 8), using radiolabeled peptidyl ([¹²⁵I]Tyr-Ala-hexarelin) or radiolabeled nonpeptidyl GHS ([³⁵S]MK 0677).

A specific animal and human GHS receptor has recently been cloned (9). It is encoded by a rare messenger RNA (mRNA) with a predicted open reading frame of 366 amino acids with a transmembrane topology typified by the G-protein-coupled receptor family. Receptor transcripts are expressed in the pituitary and the hypothalamus (4, 9), and their sequence shows partial homology with the neurotensin receptor and other orphan receptors, such as GPR38, GPR39, and FM-3 (10, 11).

The hypothalamus and the pituitary gland display the highest specific GHS binding in humans and in animals (6–9), though a high level of specific binding has also been found in other areas of the central nervous system, such as the cerebral (but not cerebellar) cortex, hippocampus, medulla oblongata, choroid plexuses, thalamus, striatum, and substantia nigra (4, 6). The existence of GHS receptors in the brain and pituitary gland probably accounts for the endocrine and central activities of GHS (1–3). Recent evidence indicates that the distribution of the GHS receptor is not restricted to central organs, and the expression of type I GHS receptor mRNA has been demonstrated in the human pancreas (12) and in neuroendocrine tumors (13, 14). Radioiodinated peptidyl GHS are also able to label specific binding sites in the rat and human heart (15–17). There is already evidence, both in animals and in humans (16, 18–23), that cardiac GHS receptors could mediate biological activities that are probably independent from the effect on GH secretion. Very recently, a gastric-derived peptide, named ghrelin, has been proposed as a natural ligand for GHS receptor (24). It has been shown that ghrelin has a stimulatory effect on GH secretion in the rat, but its capacity to selectively bind the GHS receptor subtype identified with radiolabeled peptidyl GHS has not yet been elucidated.

Based on the foregoing, the aims of the present study were: 1) to investigate the presence of GHS binding sites in a wide range of peripheral endocrine and nonendocrine human tissues, by radioreceptor assay with [¹²⁵I]Tyr-Ala-hexarelin; and 2) to evaluate the ability of different unlabeled GHS (including ghrelin) to compete with the radioligand for binding sites in different tissues.

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Address correspondence and requests for reprints to: G. Muccioli, Ph.D., Division of Pharmacology, Department of Anatomy, Pharmacology and Forensic Medicine, University of Turin, Via P. Giuria 13, 10125 Turin, Italy. E-mail: mucchioli@medfarm.unito.it.

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Subjects and Methods

Chemicals

Hexarelin (His-D-2Me-Trp-Ala-Trp-D-Phe-Lys-NH₂), Tyr-Ala-hexarelin, human ghrelin [Gly-Ser-Ser-(O *n*-octanoyl)-Phe-Leu-Ser-Pro-Glu-His-Gln-Arg-Val-Gln-Gln-Arg-Lys-Glu-Ser-Lys-Lys-Pro-Pro-Ala-Lys-Leu-Gln-Pro-Arg-NH₂], and MK 0677 (N-[1(R) {[1,2-dihydro-1-methanesulphonylspiro-(3H-indole-3, 4'-piperidin)-1'-yl]-2-(phenylmethoxy)-ethyl]-2-amino-2-methylpropanamide methane sulphonate) were supplied by Europeptides (Argenteuil, France). Human GHRH (GHRH 1–44) and SRIF (SRIF 1–14) were purchased from Bachem A. G. Feinchemikalien, Bubendorf, Switzerland. [¹²⁵I]Tyr-Ala-hexarelin (SA 1800–2100 Ci/mmol) was iodinated using a lactoperoxidase method and purified by reverse-phase high-performance liquid chromatography, as previously described (6).

Tissue samples

Various autoptotic and surgical human tissue specimens were included in this study. They comprised a wide range of peripheral tissues, both from endocrine (adrenal gland, ovary, pancreas, parathyroid, pituitary gland, testis, and thyroid gland) and nonendocrine organs (adipose tissue, aortic endothelium, aortic smooth muscle, carotid, colon, coronary, kidney, liver, lung, lymphnode, mammary gland, myocardium, prostate, salivary gland, skeletal muscle, skin, spleen, stomach, uterus, and vena cava). All postmortem tissues were obtained at autopsy from 10 patients [5 males, ranging in age from 24–63 yr (median age, 53 yr); and 5 females, ranging in age from 27–54 yr (median age 46 yr)] who died of trauma or neoplasms and were submitted to autopsy for diagnostic purposes, in years 1998 to 1999, in the Department of Pathology, University of Turin. Tissue removal had been given ethical approval by our hospital committee. Five samples of placenta from at-term spontaneous deliveries and 2 fragments each of adipose tissue, adrenal gland, colon, liver, lung, kidney, lymphnode, mammary gland, ovary, pituitary, prostate, salivary gland, skeletal muscle, skin, spleen, stomach, testis, thyroid, and uterus were also collected from surgical specimens received in the above Department in the same period. Of each specimen, a small (less than 1 cm³) tissue fragment (adjacent to the one fixed in formalin and used for histopathology) was immediately frozen at –30 C and stored for 1–14 months until processed for membrane preparation and binding studies. Although care was taken to collect all tissues from apparently normal organs, the absence of major pathological abnormalities was also confirmed by microscopic examination of adjacent tissue blocks of each specimen.

Binding studies

GHS binding assay with tissue membranes was performed using [¹²⁵I]Tyr-Ala-hexarelin as radioligand (6, 25). The membrane fractions were prepared from frozen pieces of tissue using the method previously described for brain and pituitary gland (6). The thawed tissues were homogenized in 5–10 vol sucrose 0.3 mol/L. The homogenate was first centrifuged at 500 × *g* at 4 C for 10 min, and the supernatant was carefully decanted and subjected to a second centrifugation at 30,000 × *g* for 30 min. The resulting pellet was resuspended in ice-cold buffer (50 mmol/L Tris, 2 mmol/L EGTA, 0.03% bacitracin, titrated with HCl to pH 7.3) and immediately used to determine protein content (26) and for binding studies.

In preliminary experiments, it was found that equilibrium binding conditions for the different tissues were similar to those found for binding to human hypothalamus and pituitary gland (6). For saturation binding studies, tissue membranes (corresponding to 100 μg protein) were incubated in triplicate, at 0 C for 60 min, with increasing concentrations (0.25–20 nmol/L) of [¹²⁵I]Tyr-Ala-hexarelin in a final vol of 500 μL assay buffer (50 mmol/L Tris, 2 mmol/L EGTA, 0.1% BSA, 0.03% bacitracin, titrated with HCl to pH 7.3). Parallel incubations, where 2.5 μmol/L unlabeled Tyr-Ala-hexarelin was also present, were used to determine nonspecific binding, which was subtracted from total binding to yield specific binding values. The binding reaction was terminated by adding ice-cold assay buffer followed by filtration through Whatman GF/B filters (Merck Eurolab s.r.e., Milan, Italy). Filters were rinsed three times with assay buffer, and the radioactivity remaining bound to the filters was measured by a Packard gamma counter A5003 (Packard

Bioscience s.r.e., Milan, Italy). Specific binding was calculated as the difference between binding in the absence and in the presence of excess unlabeled Tyr-Ala-hexarelin and was expressed as fmol/mg protein. Precautions were taken to minimize variations in the binding of [¹²⁵I]Tyr-Ala-hexarelin to tissue fractions. Thus, all binding studies related to one membrane preparation were carried out using the same batch of radiotracer. Saturation isotherms were transformed using the method of Scatchard (27), and the number of binding sites (maximal binding capacities) and the dissociation constant (K_d) for each tissue preparation were calculated with the Prism 3 program (GraphPad Software, Inc., San Diego, CA). To establish binding site specificity, increasing concentrations of various competitors (hexarelin, human ghrelin, MK 0677, GHRH, and SRIF) were tested in displacement assays with [¹²⁵I]Tyr-Ala-hexarelin, and the IC₅₀ values were calculated by iterative nonlinear curve-fitting program.

Statistical analysis

Values are expressed as median and range unless otherwise noted. In saturation and competition binding experiments, they are expressed as mean ± SEM. The number of subjects is indicated by *n*. Significant differences between groups were assessed by one-way ANOVA followed by the Mann-Whitney test or Duncan's multiple-range test, depending on the experiments. *P* < 0.05 was chosen as the level of significance.

Results

Binding of [¹²⁵I]Tyr-Ala-hexarelin to membranes from various human tissues

Wide variations of binding were observed among the tissues studied (Table 1). The highest [¹²⁵I]Tyr-Ala-hexarelin binding activity was observed in the myocardium of both

TABLE 1. Specific binding of [¹²⁵I]Tyr-Ala-hexarelin to membranes from various autoptotic human tissues

Tissue	Specific binding of [¹²⁵ I]Tyr-Ala-hexarelin (fmol/mg protein)	
	Male subjects	Female subjects
Myocardium	4302 (3897–4856)	3987 (3712–4564)
Adrenal gland	2853 (2656–3024)	2786 (2442–3223)
Testis	2561 (2243–2856)	—
Aortic smooth muscle	2434 (1862–2521)	2217 (2034–2543)
Aortic endothelium	2366 (1897–2658)	2187 (1816–2488)
Coronary	2245 (1925–2452)	2108 (1686–2356)
Carotid	2005 (1824–2206)	1989 (1665–2221)
Lung	1896 (1789–2114)	1865 (1765–2023)
Ovary	—	1865 (1786–2012)
Liver	1602 (1435–1634)	1498 (1421–1558)
Skeletal muscle	1523 (1404–1725)	1478 (1345–1723)
Kidney	1427 (1285–1610)	1356 (1229–1532)
Pituitary gland	1184 (990–1345)	1205 (1002–1353)
Thyroid gland	912 (689–1095)	1002 (879–1121)
Adipose tissue	487 (376–654)	323 (233–612)
Vena cava	406 (267–546)	434 (312–673)
Uterus	—	302 (223–482)
Skin	282 (200–432)	289 (211–512)
Lymphnode	210 (134–423)	212 (108–376)
Parathyroid gland	<10 (0–5)	<10 (0–7)
Pancreas	<10 (2–7)	<10 (0–4)
Placenta ^a	—	<10 (3–6)
Mammary gland	—	<10 (4–7)
Prostate	<10 (0–8)	—
Salivary gland	<10 (6–9)	<10 (0–8)
Stomach	<10 (0–3)	<10 (0–3)
Colon	<10 (0–5)	<10 (3–5)
Spleen	<10 (6–7)	<10 (2–9)

^a From at-term spontaneous deliveries. Values from five subjects are expressed as median (range).

sexes and ranged from 3712–4856 fmol/mg protein. The specific binding in the myocardium represented 70–82% of the total radioactivity bound. Adrenal gland, testis, aortic smooth muscle, aortic endothelium, coronary, carotid, lung, and ovary possessed intermediate binding, with values ranging from 1665–3223 fmol/mg protein; liver, skeletal muscle, kidney, pituitary gland, and thyroid gland also showed significant binding, in the range of 689–1725 fmol/mg protein. Low binding, ranging from 108–673 fmol/mg protein, was detected in membranes from adipose tissue, vena cava, uterus, skin, and lymphnode. By contrast, negligible binding (<10 fmol/mg protein) was seen in the remaining tissues (parathyroid gland, pancreas, placenta, mammary gland, prostate, salivary gland, stomach, colon, and spleen). Differences in the specific binding values were statistically significant when comparing the myocardium with the other tissues ($P < 0.001$ in both sexes); likewise the values obtained in adrenal gland, testis, aortic smooth muscle, aortic endothelium, coronary, carotid, lung, and ovary differed significantly ($P < 0.01$ in both sexes) from those of liver, skeletal muscle, and kidney; differences in the binding values were also statistically significant for liver, skeletal muscle, and kidney *vs* pituitary gland and thyroid gland ($P < 0.05$ in both sexes) and for pituitary gland and thyroid gland *vs*. adipose tissue, vena cava, skin, uterus, and lymphnode ($P < 0.001$ in both sexes). No sex differences in the Tyr-Ala-hexarelin binding to various organs were observed. Determinations performed with the same batch of radiolabeled Tyr-Ala-hexarelin, on membranes from surgical or autopsy specimens of the same tissues, yielded overlapping binding values.

Representative saturation isotherms and Scatchard plots of [125 I]Tyr-Ala-hexarelin binding to membranes from those tissues (myocardium and adrenal gland) that displayed the highest specific binding activity are shown in Fig. 1. Experiments using increasing concentrations of radiotracer revealed evidence of saturable specific binding in the myocardium and adrenal gland (Fig. 1a). Scatchard analysis of these data (Fig. 1b) demonstrated the existence in both tissues of a single class of high-affinity sites with K_d values not substantially different from one another, being 5.5 ± 1.3 nmol/L for males ($n = 5$) and 5.4 ± 1.6 nmol/L for females ($n = 5$) in the myocardium and 6.5 ± 1.2 nmol/L ($n = 5$) for males and 5.9 ± 1.4 nmol/L for females ($n = 5$) in the adrenal gland. K_d values very close to those found in the above tissues were also detected in the other GHS-receptor positive tissues of both sexes.

Specificity of binding

The specificity of [125 I]Tyr-Ala-hexarelin binding to tissue membranes was assessed by competitive binding studies with different GHS. Table 2 shows the IC_{50} values of hexarelin, MK 0677 and human ghrelin calculated from the respective displacement curves in various endocrine and nonendocrine organs. Unlabeled hexarelin completely displaced radiolabeled Tyr-Ala-hexarelin from binding sites. The concentration of hexarelin required to inhibit radiotracer binding by 50% did not significantly differ among the various tissues examined and ranged from 2.7 to 5.5×10^{-8} mol/L. When human ghrelin and MK 0677 were tested in the dis-

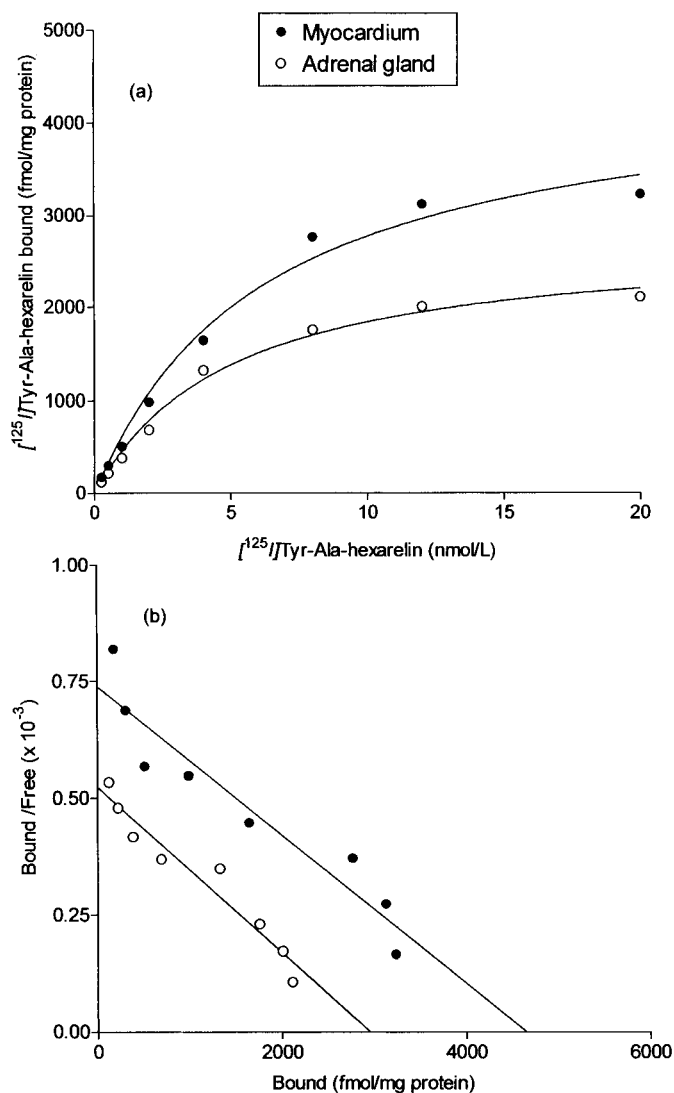


FIG. 1. Representative saturation isotherms (a) and Scatchard plots (b) of [125 I]Tyr-Ala-hexarelin binding to membranes from myocardium and adrenal gland of male subjects. Experiments were performed by incubating a fixed amount of membrane protein (100 μ g/tube) with increasing concentrations of radiolabeled Tyr-Ala-hexarelin alone (total binding) or plus 2.5 μ mol/L unlabeled Tyr-Ala-hexarelin to define nonspecific binding. Specific binding values were obtained by subtracting nonspecific binding from total binding. The saturation curves of specific binding were analyzed by Scatchard analysis in order to calculate the maximal binding capacities and the K_d values.

placement studies, a dose-dependent inhibition of binding was seen with both compounds only in the adrenal gland (Fig. 2) and in various endocrine organs such testis, ovary, pituitary and thyroid gland. In these tissues human ghrelin and MK 0677 completely displaced [125 I]Tyr-Ala-hexarelin with equal efficacy, but they were significantly ($P < 0.05$) less potent than hexarelin (4–5 times). By contrast, in the myocardium (Fig. 2), as well as in the other nonendocrine tissues, human ghrelin and MK 0677 (tested at concentrations of 0.1 nmol/L–2 μ mol/L) were able to displace only 11–31% of the specifically bound radiolabeled Tyr-Ala-hexarelin. Other peptides (GHRH, SRIF) that have a known influence on GH release had no effect on the binding of [125 I]Tyr-Ala-hexarelin

TABLE 2. Concentrations of hexarelin, human ghrelin, and MK 0677 required to inhibit by 50% (IC₅₀) the specific binding of [¹²⁵I]Tyr-Ala-hexarelin to membranes from various human tissues

Tissue	IC ₅₀ (×10 ⁻⁸ mol/L)		
	Hexarelin	Ghrelin	MK 0677
Adrenal gland	3.4 ± 0.6	17.0 ± 2.5 ^a	18.3 ± 1.4 ^a
Testis	2.7 ± 1.0	19.7 ± 1.2 ^a	22.0 ± 3.2 ^a
Ovary	3.9 ± 1.5	15.7 ± 3.8 ^a	19.3 ± 3.2 ^a
Pituitary gland	3.7 ± 1.1	19.3 ± 3.3 ^a	18.0 ± 1.7 ^a
Thyroid gland	4.6 ± 0.8	16.3 ± 2.6 ^a	17.3 ± 2.0 ^a
Myocardium	3.7 ± 0.8	(27 ± 6)	(16 ± 3)
Aorta	2.8 ± 1.0	(26 ± 5)	(20 ± 3)
Coronary	3.1 ± 0.2	(24 ± 6)	(18 ± 5)
Carotid	5.0 ± 1.0	(27 ± 8)	(20 ± 2)
Lung	4.8 ± 0.8	(23 ± 9)	(19 ± 4)
Liver	4.5 ± 1.6	(24 ± 6)	(17 ± 7)
Skeletal muscle	5.1 ± 2.0	(21 ± 5)	(14 ± 4)
Kidney	3.8 ± 0.9	(17 ± 3)	(27 ± 4)
Adipose tissue	3.8 ± 1.0	(19 ± 7)	(20 ± 6)
Vena cava	5.0 ± 1.1	(15 ± 5)	(16 ± 5)
Uterus	4.8 ± 1.3	(16 ± 5)	(23 ± 5)
Skin	5.5 ± 1.2	(11 ± 4)	(12 ± 5)
Lymphnode	5.3 ± 3.3	(31 ± 8)	(22 ± 5)

The values in parentheses indicate the percentage of displacement produced by 2 μmol/L ghrelin or MK 0677. Values are means ± SEM of three separate experiments.

^a P < 0.05 vs. hexarelin.

to receptors of endocrine and nonendocrine tissues (data not shown).

Discussion

Peptidyl and nonpeptidyl GHS possess specific receptor subtypes mostly distributed at the level of the pituitary gland and the hypothalamus (6–8), where they probably mediate the classical endocrine activities of GHS, *i.e.* strong GH-releasing effect but also significant stimulatory effect on PRL and ACTH secretion (4, 5, 28). On the other hand, the presence of appreciable amounts of GHS binding sites and GHS-receptor mRNA in the central nervous system may account for the central activities of GHS, such as their influence on food intake and sleep (29–32). The existence of GHS binding sites in some peripheral tissues had also been reported in animals and in humans using *in vitro* radioreceptor assay (15–17, 25) and GHS-receptor mRNA analysis (12). In particular, specific GHS binding sites had been demonstrated in the rat (16) and human heart (15, 17) and even in human thyroid (25). Interestingly, other G-protein-coupled orphan receptors, sharing significant homology with the GHS receptors, have recently been cloned and characterized from different peripheral tissues such as thyroid, stomach, colon, pancreas and bone marrow (10, 11).

In the present study we have found that several endocrine and nonendocrine peripheral human tissues show specific binding values for Tyr-Ala-hexarelin which are even higher than found in the pituitary gland. This was found to be the case in the myocardium, adrenal, testis, arteries, lung, ovary, liver, skeletal muscle, and kidney. Specific binding values quite close to those detected in the pituitary were found in the thyroid gland. Lower levels of binding were observed, on the other hand, in adipose tissue, veins, uterus, skin and lymphnode, while negligible binding was found in parathy-

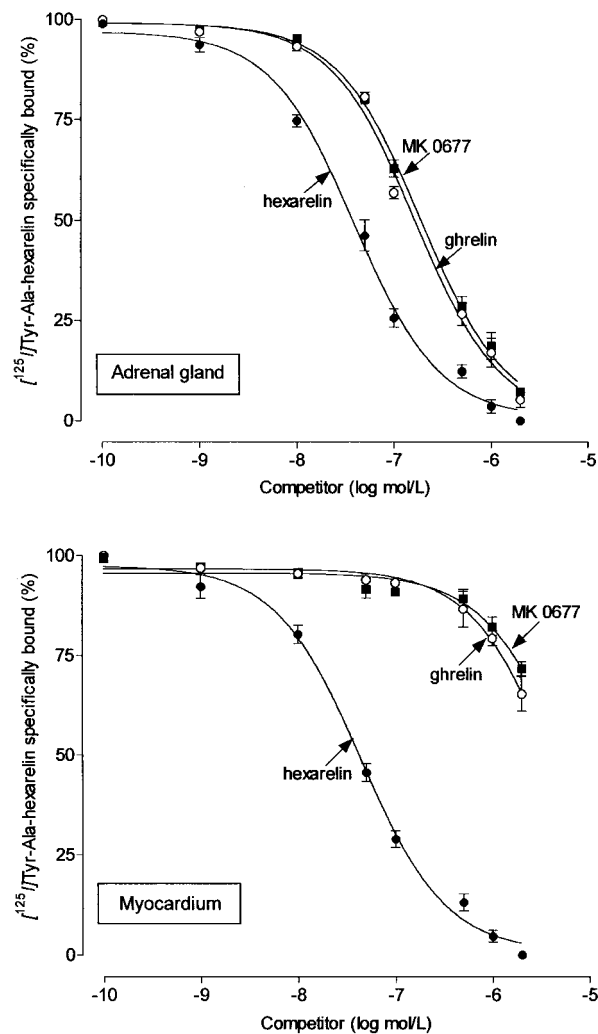


FIG. 2. Displacement of [¹²⁵I]Tyr-Ala-hexarelin from membranes of adrenal gland and myocardium by hexarelin, human ghrelin and MK0677. Binding assays were conducted as described in *Subjects and Methods*. The ordinate represents binding as a percentage of control (specific binding in the absence of unlabeled competitor). Each point represents the mean ± SEM of three separate experiments.

roid glands, pancreas, placenta, mammary gland, prostate, salivary gland, stomach, colon, and spleen. These findings indicate that GHS have widely spread receptors in some (but not all) peripheral endocrine and nonendocrine human tissues which could mediate effects other than classical endocrine and central activities (33). Evidence has been provided that treatment with peptidyl GHS exerts cardiovascular activities. In fact, coronary vasoconstriction (16) or protective effect against ischemia depending on dose and experimental conditions (18–20) and improvement of cardiac performances after myocardial infarction (23) have been observed in rats, while an increase in the left ventricular ejection fraction has been reported in humans (21). It will be noted that the myocardial effects of peptidyl GHS are elicited also in animals (18) and humans (20, 22) with severe GH deficiency. In addition, no cardiac effects were noted after giving non-peptidyl GHS. One can therefore argue that the GHS binding sites in the heart are specific for peptidyl GHS only and

mediate cardiovascular activities independently from their GH-releasing effect. This is not the case of the thyroid where both peptidyl and nonpeptidyl GHS displaced Tyr-Ala-hexarelin from binding sites of normal and follicular-derived neoplastic tissues and carcinoma cell lines. The thymidine incorporation and the proliferation of the latter is inhibited by both peptidyl and nonpeptidyl GHS (25). Finally, it is noteworthy that among peripheral tissues lacking GHS binding there was the stomach which has been shown as the major site of ghrelin synthesis and release (24) and the pancreas where GHS-receptor mRNA has been demonstrated (12). These data further indicate the complexity of the GHS compound/receptor interactions. Even the discovery of ghrelin as a natural GHS-like ligand (24) has not completely clarified the whole matter.

The binding of [¹²⁵I]Tyr-Ala-hexarelin to membranes from peripheral human tissues showed many of the properties typical of the ligand-receptor interaction. These included high affinity, saturability and structural specificity. The specificity of radioiodinated Tyr-Ala-hexarelin binding was very similar to that observed in other human peptidyl GHS target tissues such as the hypothalamus and the pituitary gland (6), since the binding of radioligand was displaced by unlabeled Tyr-Ala-hexarelin, hexarelin, but not by peptides (GHRH or SRIF) structurally unrelated to peptidyl GHS.

The peripheral GHS binding sites do not necessarily reflect the peripheral distribution of the classic GHS receptor cloned by Howard *et al.* (4, 8, 9). At least in some tissues, they could even reflect the existence of GHS receptor subtypes different from that previously characterized in the pituitary (6, 7). In our study binding specificity showed remarkable differences among tissues displaying binding sites. In fact, the binding of [¹²⁵I]Tyr-Ala-hexarelin was inhibited by ghrelin as well as by the nonpeptidyl spiroindoline MK 0677 in membranes from endocrine but not from nonendocrine tissues. It is worth noticing that in membranes from endocrine tissues ghrelin and MK 0677 inhibited the binding of radiolabeled Tyr-Ala-hexarelin but to a lower extent than that shown by peptidyl GHS. In all, these data clearly point toward the existence of new GHS receptor subtypes which in peripheral nonendocrine tissues seem specific for peptidyl GHS and do not bind ghrelin nor the spiroindoline MK 0677.

In conclusion, the present data demonstrate that GHS have specific receptors in a wide range of endocrine and nonendocrine human tissues and suggest that a still unknown receptor subtype, specific for peptidyl GHS, may exist in the heart and in other tissues.

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