Gonadotropin-releasing hormone neuropeptides and receptor in human breast cancer: Correlation to poor prognosis parameters

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A B S T R A C T

Expression of the two gonadotropin-releasing hormone homologue peptides GnRHI and GnRHHI and their receptor GnRHR has been demonstrated in a number of malignancies. In hormone-dependent breast cancer, GnRH analogs are used for therapy in premenopausal women. Gene expression of GnRHI, II and R was studied in breast biopsies from primary breast adenocarcinoma obtained from the tumor and the adjacent benign tissue. Levels were evaluated by a multiplex real-time RT-PCR. GnRHI transcripts were detected in 14.7% of the benign and 29.4% malignant biopsies and GnRHHI in 21.2% benign and 44.1% malignant biopsies. GnRHR was also more frequent in the malignant (54.2%) than in the benign (24.0%) biopsies, at similar expression levels. No transcripts were detected in biopsies from healthy individuals. There was a strong correlation between the presence of GnRHI and GnRHHI transcripts and their receptor in the benign and the malignant biopsies. GnRHI, II and R expression correlated significantly with poor prognosis pathological parameters. Immunohistochemistry for GnRHR revealed expression in malignant cells and in epithelial cells of mammary ducts of the adjacent area with pre-cancerous features. In contrast, GnRHI and II peptides were rarely expressed at low levels in breast cancer cells. In conclusion GnRHI peptides and receptor are expressed more frequently in breast tumors than in the adjacent mammary tissue, representing a malignant feature. Their expression correlated to tumor characteristics of poor prognosis and was therefore related to more aggressive malignancies. Concomitant expression of peptides and receptor supports an autocrine/paracrine regulating role.

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1. Introduction

Gonadotropin-releasing hormone (GnRH) is a decapeptide produced in the hypothalamus [10]. It interacts with a G-protein coupled receptor (GnRHR) in the anterior pituitary [23], controlling the gonadal function in both sexes. Two human isoforms have been identified, namely GnRH-I and GnRH-II. The first is the hypothalamic isoform responsible for the secretion of LH and FSH. The second differs by three amino acids [6,27] and is widely distributed in the central and peripheral nervous system. It is also expressed at significantly higher than GnRH I levels outside the brain and it has been shown to act as a neuromodulator in the behavioral components of reproduction [16,27,28].

GnRHI peptides and GnRHR have been found in extrapituitary tissues and tumors of the reproductive and other systems [4,20,29,34,2]. Extrapituitary GnRH binding sites are often associated with many novel cellular responses [7]. Furthermore, expression of GnRHR seems to be related with advanced cancer stage in ovarian carcinomas [8].

The GnRH system has been reported to play an autocrine/paracrine role in the inhibition of cellular growth and metastatic potential in breast cancer cell lines [24,35], and breast tumor regression in nude mouse [14,26]. However, its expression...
was associated with a protective effect on the chemotherapeutic drug-produced apoptosis [30]. As GnRH agonists (or antagonists) show clinical benefit when used as adjuvant pharmacotherapy in premenopausal breast cancer patients [13], the study of the GnRH system of neuropeptides and receptor in breast tumors remains emerging.

In the present study, the expression of the two GnRH neuropeptide genes (GnRH I, II) and their receptor was evaluated in a series of biopsies from primary breast cancers in a quantitative manner by multiplex real-time RT-PCR. Transcript levels from the malignant tissues were compared to these from the adjacent non-neoplastic tissue and tissues without malignancy, and were correlated to multiple clinicopathological and demographic parameters and clinical output in order to reveal potential prognostic or diagnostic value. Finally, histological mapping of peptide and receptor expression in breast cancer biopsies was done by immunohistochemistry, to reveal specific target cellular types.

2. Materials and methods

2.1. Tissues

Patients newly diagnosed with primary breast adenocarcinoma in the “Theagenio” Cancer Hospital, Thessaloniki, Greece were enrolled in the study. Biopsies were obtained from the tumor and the adjacent non-neoplastic tissue. Diagnosis was confirmed by the histological examination in all patients. Full medical history, follow-up and histopathological data were available. Patients have not been receiving any hormonal treatment chemotherapy or radiation. Patients with previous or present neoplastic disease at any other site were excluded from the study. Biopsies without signs of malignancy or other pathology obtained for diagnostic use were also used. Human term placenta was obtained by the Obstetrics and Gynecology Department of the General University Hospital in Alexandroupolis. The project was approved by the local Ethical Committee. Consent has been obtained from each patient after full explanation of the purpose and nature of all the procedures used, in accordance to the Helsinki Declaration. Tissue samples were stored in RNAlater (Invitrogen, Carlsbad, CA) at −80°C until used for RT-PCR. Breast cancer tissue sections were also taken from paraffin-embedded archival files and used for immunohistochemistry.

2.2. Cell culture

The human breast cancer cell lines MDA MB231, MCF7 and T47 were cultured in Dulbecco’s Modified Eagle Medium (DMEM) supplemented with 10% fetal calf serum (FCS) and 1% penicillin/streptomycin (all purchased from Invitrogen, UK), at 37°C in a 5% CO₂ humidified atmosphere. Cells were plated at a concentration of 2 × 10⁵ cells/ml and were harvested for total RNA extraction when they had reached approximately 80% confluence.

2.3. Multiplex real-time quantitative RT-PCR

Total RNA was extracted from biopsies using Trizol Reagent, according to the manufacturer’s instructions. Reverse transcription (RT) was performed using the SuperScript Preamplification System (Invitrogen) and random hexamers in a total volume of 20 µl. Two microliter of the same RT product was used as a template for each gene, amplified by PCR using 2 mM MgCl₂, PCR buffer, 0.2 mM of sense and antisense primers, 0.2 mM dNTPs and 2.5 U Taq Polymerase (Invitrogen) in a final reaction volume of 50 µl. Quantitative PCR was performed using the Light Cycler MX3005P (Stratagene, La Jolla, CA) with the following cycling parameters: a pre-amplification cycle (denaturation for 10 min at 95°C), 40 cycles of amplification (denaturation for 30 s at 95°C, annealing for 40 sec at 53°C, 54°C, 50°C for GnRH I, GnRH II and GnRH R respectively, and extension for 50 s at 72°C), and a final dissociation cycle (1 min at 95°C, 40 s at 57°C and 30 sec at 95°C). Primers were designed according to the GenBank published sequences as follows: for human GnRH R sense 5′-CTCTGTTGAGAACATCCGA-3′ and antisense 5′-GAGGCGTCACCTCGAT-3′ [33], for human GnRH I sense 5′-CATCTGGTGGTGCTGGA-3′ and antisense 5′-CTGGCCACCTTCTCTTCAA-3′ and for human GnRH II sense 5′-TCTGCTGTGCTGACTG-3′ and antisense 5′-CTAAGGCATCTGGG- GAT-3′ [25]. Product sizes were 319, 240 and 119 bp for GnRH R, GnRH I and GnRH II respectively. Reactions in duplicate were carried out using the SYBER Green MM QPCR Brilliant mix (Stratagene), 0.4 µM of each primer, 2 mM MgCl₂ and 0.5 µL of cDNA in a final volume of 20 µL. Results were calculated using MaxPro QPCR Software Version 4.0 (Stratagene) using the comparative threshold cycle method. Analysis of relative gene expression data was performed according to the 2−ΔΔCt method [21] using β-actin as a reference gene and RNA from human placenta as a positive control. Results are expressed as the mean from duplicate values of gene expression in relation to β-actin in the same RNA preparation. Samples with poor β-actin gene amplification were excluded from the study. Negative control samples, where no RT enzyme was added (no RT) or without DNA template (no DNA), were included in every assay in order to exclude the possibility of genomic or other DNA contamination.

2.4. Immunohistochemistry

Immunohistochemistry was conducted as previously described [32]. Briefly, tissue specimens were fixed in formalin and embedded in paraffin. Sections (4 µm) were deparaffinized, rehydrated, and treated with 0.3% H₂O₂ for 5 min in methanol. Slides were incubated for 75 min with primary mouse monoclonal antibodies for human GnRH R (ab22168, Abcam, UK), GnRH I (H111B, SantaCruz Biotechnology Inc., CA, USA) and GnRH II (D-9, SantaCruz Biotechnology Inc.) diluted 1:250, 1:100 and 1:100 respectively in 10% normal mouse serum in PBS. Negative control slides were incubated for the same period with normal mouse serum IgG. Immunostaining was detected by the Kwik Kit (Thermo Shandon, Pittsburgh, PA, USA). Finally, bound antibody complexes were stained for 10 min with 0.05% diaminobenzidine, counterstained with Mayer’s hematoxylin, mounted and examined under an Olympus BX40 microscope.

2.5. Statistical analysis

All measurements were done in duplicate. Statistical significance was assessed by Mann–Whitney U–Wilcoxon Rank Sum W Test, using the SPSS 17.0 statistical software (SPSS Inc., Chicago, IL, USA). Group differences were assessed by chi square test. Significance was set at a P value <0.05. Analysis of the data in pairs of benign and malignant biopsies from the same patient was performed by the McNemar test. Kaplan Meier survival analysis was also performed.

3. Results

3.1. Patient and tumor information

Thirty-five women with primary breast cancer were enrolled in the study, with mean age 61 ± 13 years, mean BMI 28.9 ± 5.4 kg/m² and mean age of menarche 13 ± 1.3 years. At the time of diagnosis, 27/35 (71.4%) were menopausal with mean age of menopause 48.5 ± 3.9 years. Two of them did not report any history of pregnancy, whereas for the rest the mean number of full-term pregnancies was 1.9 ± 0.8, with mean age of first pregnancy 25.1 ± 4.3
years and mean duration of lactation for all children 11.2 ± 12.8 months. Mean value for pregnancy terminations was 0.8 ± 1.0. One of them (2.85%) reported alcohol consumption (<250 ml/day of drink with 5–10% alcohol) and 4/35 (11.4%) had been administered contraceptive pills (0.25–2 years). Their education level was at 21/35 (60.0%) basic, 10/35 (28.5%) high-school and 4/35 (11.4%) university. 17/35 (48.5%) had family history of malignancy and 5/35 (14.2%) breast. 3/35 (8.5%) had been previously diagnosed with benign breast disease. None of the patients had received radiation for any reason or had history of other malignancy or syndrome.

The clinicopathological findings of the tumors by pathological examination based on the TNM system of the American Joint Committee on Cancer, used in the study are shown in Table 1.

Follow-up for 24–68 months showed disease relapse in 4/32 (12.5%) cases. In 3 patients follow-up was not possible. One patient developed endometrial adenocarcinoma 36 months after initial diagnosis, and exacerbation was reported in the patient with liver metastasis. At the completion of the study, 26/32 (81%) patients were disease-free, 4/32 (13%) still had disease and 2/32 (6%) died.

3.2. Levels of gene expression of GnRH neuropeptides and receptor in breast cancer biopsies

GnRH I, II and R gene expression was examined by comparative real-time RT-PCR in human breast cancer biopsies and in the adjacent non-malignant tissue. PCR products were denaturating at the same temperature as the product from human term placenta used as positive control.

GnRH I gene transcripts were found twice as frequently in the malignant (10/34, 29.4%) than the benign biopsies (5/35, 14.7%) examined. Transcript levels did not differ between benign and malignant biopsies in a statistically significant manner, being $92 \times 10^{-3} \pm 60 \times 10^{-3}$ and $129 \times 10^{-3} \pm 95 \times 10^{-3}$, respectively. When analysis was performed in pairs of benign and malignant biopsies from the same patient, it was found that in 18/26 (69.2%) cases both biopsies were negative, in 4/26 (15.4%) only the malignant biopsy was positive, in 2/26 (7.7%) only the benign biopsy was positive and in 2/26 (7.7%) both biopsies were positive (Fig. 1).

GnRH II transcripts were found in 7/33 (21.2%) of the benign biopsies examined and in 15/34 (44.1%) of the malignant biopsies and this difference was statistically significant ($p = 0.04$). Transcript levels, as compared to the expression levels found in human term placenta, did not differ between benign and malignant biopsies in a statistically significant manner, being $17 \times 10^{-3} \pm 11 \times 10^{-3}$ and $55 \times 10^{-3} \pm 30 \times 10^{-3}$ respectively. When analysis was performed in pairs of benign and malignant biopsies from the same patient, it was found that in 12/24 (50.0%) cases both biopsies were negative, in 2/24 (29.2%) only the malignant biopsy was positive, in 3/24 (12.5%) only the benign biopsy was positive and in 2/24 (8.3%) both biopsies were positive (Fig. 2).

GnRH R gene expression was found to be more frequent in the malignant biopsies (13/24, 54.2%) compared to the benign biopsies (6/25, 24.0%), in a statistical significant manner ($p = 0.05$). GnRH R transcript levels, expressed in relation to human term placenta, were also higher in the malignant tissues ($430 \times 10^{-3} \pm 365 \times 10^{-3}$) in comparison to the benign ($91 \times 10^{-3} \pm 54 \times 10^{-3}$) although this difference was not statistically significant. When analysis was performed in pairs of malignant and benign biopsies from the same patient, the following pattern was revealed regarding GnRH R expression; in 8/16 (50.0%) patients GnRH R transcripts were absent in both biopsies, in 4/16 (25.0%) were present in both biopsies, in 4/16 (25.0%) GnRH R was expressed only in the malignant tissues, whereas in none (0.0%) of the patients transcripts were found in the benign but not in the malignant biopsy (Fig. 3).

Table 1

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3.3. Levels of gene expression of GnRH neuropeptides and receptor in breast cancer cell lines

Expression of the three genes was also examined in the human breast cancer cell lines MDA MB231, MCF7 and T47. All cell lines expressed low levels of GnRH I (0.50 × 10⁻³, 0.06 × 10⁻³ and 34.70 × 10⁻³ units in relation to human placenta, respectively) and GnRH II (0.60 × 10⁻³, 0.05 × 10⁻³ and 16.70 × 10⁻³ units in relation to human placenta, respectively), whereas GnRH R was found in MDA MB231 and MCF7 (3.4 × 10⁻³ and 0.2 × 10⁻³ units in relation to human placenta, respectively), but not in T47.

3.4. Expression of GnRH peptides and GnRH R protein in breast cancer biopsies

Immunohistochemical analysis was used in order to detect GnRH peptides and GnRH R protein expression in 16 human breast tumor biopsies and to localize it at histological and cellular level. Immunoreactivity for GnRH I was found in breast cancer cells in 5 out of 16 tissues (31.25%), localized in perineural invasions, implants and macrophages (Fig. 4). Similarly, the majority of the tumors were negative for GnRH II. Immunoreactivity was found in 5 out of 16 tissues, being also positive for GnRH I, in breast cancer biopsies.
cells and in cancer implants, and in infiltrating macrophages (Fig. 5). Epithelial cells of cancerous implants of a mucinous carcinoma studied were positive for both GnRH I (Fig. 4C) and II (Fig. 5C). Immunoreactivity in all cases was cytoplasmic and mild, not uniform in all areas of the tissue section but rather occasional. The adjacent to the tumor area showed no immunoreactivity for both antibodies. Human term placenta, used as a positive control, showed strong cytoplasmic immunoreactivity (Figs. 4E and 5E). No specific staining was found in any cell type after replacement of the primary antibody by non-specific mouse IgG (negative control in placenta and breast cancer, Figs. 4F and 5F respectively).

Tissue sections were also stained using monoclonal antibodies against the N-terminus of human GnRH R. Representative areas of the sections are presented in Fig. 6. Positive GnRH R staining was localized in malignant cells of the tumor which varied in size and shape (A–C) and in the cancerous implants of ductal in situ carcinoma of solid type with (B) or without (A) inflammation (aggregates of periductal mononuclear inflammatory cells), and in macrophages scattered in a negative stroma. Immunostaining was clearly membranic but also cytoplasmic (C). In the adjacent to the tumor area, epithelial cells of mammary ducts presenting features of atypia, lack of polarity, enlarged nuclei, prominent nucleoli and numerous mitosis, showed some immunoreactivity for the GnRH receptor, mainly cytoplasmic (B). Finally, cancer cells of perineural invasions were negative (D).

Fig. 2. Detection of GnRH II gene transcripts in malignant and adjacent benign biopsies from breast tumors by real-time PCR following reverse transcription of total RNA extracts. (A) Presence of GnRH II transcripts and percentages of positive tissues. (B) Levels of gene expression. Bars represent means between all positive tissues and error bars the standard deviation between measurements. (C) Expression in malignant biopsies in relation to their adjacent benign tissue.
Fig. 3. Detection of GnRH R gene transcripts in malignant and adjacent benign biopsies from breast tumors by real-time PCR following reverse transcription of total RNA extracts. (A) Presence of GnRH R transcripts and percentages of positive tissues. (B) Levels of gene expression. Bars represent means between all positive tissues and error bars the standard deviation between measurements. (C) Expression in malignant biopsies in relation to their adjacent benign tissue.

4. Discussion

The two GnRH neuropeptides (GnRH I and II) and their receptor are key players of the nervous system control on the reproductive function. Their expression however has also been clearly established in many extra-pituitary organs in reproductive and non-reproductive tissues and tumors arising there, breast cancer being one of the first reported [7]. Ectopic expression of neuropeptides is frequently found in endocrine cancers. Their role seems to contribute to the cancer biology via activation of locally expressed receptors in an autocrine manner or a paracrine dialog in the tumor microenvironment between the tumor cells and the nearby located cells, such as stroma, immune cells or by innervating autonomic neurons. Breast cancer tumors are known to express multiple neuropeptides and their receptors such as CRF, GHRH and somatostatin along with GnRH, firstly studied in breast cancer by immunohistochemistry [9]. We have recently reported the expression of both CRF receptors in breast cancer and the respective benign adjacent tissue, which could serve as targets of endogenous ligands regulating cancer growth [15]. In the present study we quantify gene expression of GnRH I and II along with GnRH R in human breast tumors. It is important that two biopsies were collected by the same patient, one from the breast cancer tissue and one from the adjacent benign tissue and results in the actual tumor were compared with those from its pre-cancerous milieu.
GnRH I and GnRH II gene transcripts were found approximately twice as frequent in the malignant than in the benign biopsies, implying that they present a characteristic accompanying malignant transformation. Transcript levels showed no statistically significant differences between malignant and benign biopsies for either gene. This is in accordance with previous findings of semi-quantitative determinations of GnRH I mRNA in a small number of tissues [19]. Chen et al., 2002 [5] however, reported higher levels of GnRH I and GnRH II in the malignant biopsies of breast in comparison with the adjacent benign tissue, as detected in 6 breast cancer patients by semi-quantitative RT-PCR, attributed to the overall high protein expression and enhanced transcription machinery that exist in cancer cells. The great variability found in the expression levels between the positive samples might have prohibited statistically significant differences to emerge in our study, which however employs a strictly quantifying method correcting for the total protein content (using actin as a house-keeping gene) and a much larger number of tissues.

Fig. 4. Immunohistochemical analysis of GnRH I peptide expression in human breast tumor biopsies. Tissue sections were stained using a monoclonal antibody and representative fields are presented. Mild cytoplasmic immunoreactivity was localized in breast cancer cells of perineural invasions in some tissues (A, arrows) but not in others (D), in cancer implants (B, arrows), in macrophages (B, diamonds) and in malignant epithelial cells of cancerous implants in mucinous matrix of a mucinous carcinoma studied (C, arrows). Human term placenta, used as a positive control, showed strong cytoplasmic immunoreactivity (E). No specific staining was found in placental cells after replacement of the primary antibody by non-specific mouse IgG (F). Original magnification 200× (A, B, D, F, E), 400× (C).

Fig. 5. Immunohistochemical analysis of GnRH II peptide expression in human breast tumor biopsies. Tissue sections were stained using a monoclonal antibody and representative fields are presented. Mild cytoplasmic immunoreactivity was localized in cells of malignant implants in some tissues (A, arrows) but not in others (D), in macrophages (B, arrows) and in malignant epithelial cells of cancerous implants in mucinous matrix of a mucinous carcinoma studied (C, arrows). Human term placenta, used as a positive control, showed strong cytoplasmic immunoreactivity (E). No specific staining was found in breast cancer cells after replacement of the primary antibody by non-specific mouse IgG (F). Original magnification 200×.
At the peptide level, GnRH I and II were expressed rarely by breast cancer cells, as shown by immunohistochemical analysis using monoclonal antibodies. Low cytoplasmic immunoreactivity was found in 31.25% of the samples, a percentage similar to the mRNA positive tissues. Antigen was localized in some breast cancer cells of perineural invasions, implants and macrophages (Fig. 4). These results are in agreement to the study of Chen et al., 2002, showing GnRH I and II peptide immunoreactivity in 8 out of 14 breast cancer biopsies using polyclonal anti-serums. The adjacent to the tumor area showed no immunoreactivity for both antibodies. It is possible that post-transcriptional regulation mechanisms prevent peptide expression at the benign tissue. In fact, this was also the case in endometrial tissues and cell lines [4], in which GnRH transcripts were abundant but no secreted peptide was found by RIA. These mechanisms must be altered in certain malignant transformed cells, where peptides can be traced.

GnRH R transcripts were also found at higher frequency the malignant (54.2%) than the benign biopsies examined (24.0%), and at higher levels. Interestingly, in all cases found positive in the benign tissue, this was accompanied by transcript presence in the corresponding malignant biopsy, indicating that it could represent a pre-malignant feature of the mammary tissue. Indeed, GnRH R in the tumor biopsy correlated significantly to its presence in the corresponding adjacent area. GnRH R expression was also studied at the protein level by immunohistochemistry. Receptor was localized in the membranes but also in the cytoplasm of malignant cells of the tumor and in the cancerous implants of ductal in situ carcinoma of solid type, in accordance with previous reports [1,2,22]. Perineural invasions were negative. Adjacent to the tumor area, epithelial cells of mammary ducts presenting pre-cancerous characteristics were also positive, further supporting an early tumorigenesis event. Differential expression of GnRH R by different cell types within the tumor could explain the great variability in gene expression levels estimated by real-time RT-PCR. Mammary samples from patients without any malignancy were negative for both GnRH peptide and receptor gene expression.

GnRH I expression correlated significantly with the size and infiltration of the tumor in the cancer biopsies, while in the adjacent benign biopsies it correlated with absence of progesterone receptors and c-erbB2 positivity of the tumor. GnRH II was expressed more frequently than GnRH I in the same tissues, and its expression levels correlated with absence of ER. In addition, levels of GnRH R correlated with infiltration and higher patient BMI. All these parameters are negative biomarkers of breast cancer and therefore correlate GnRH system expression to more aggressive and poorer prognosis disease. Our results imply that GnRH system expression could hold a clinical prognostic potential for breast cancer. Fortunately, only 3 patients from our sample deceased and 6 developed metastasis in a 5-year follow up, making survival analysis in relation to GnRH gene expression inconclusive.

Specific binding sites for GnRH were demonstrated in human breast carcinomas but not in non-neoplastic breast tissue by ligand immunoblotting back in 1985 [18]. The percentage of GnRH R positive tissues here (54.2%) is in line with previous reports detecting binding sites by a multipoint assay [11]. Our study confirms that they represent the protein product of this gene. Similarly, the low affinity GnRH-binding sites described in human breast cancer cell lines (ZR-75-1, MDA-MB-231, Sk Br 3, MDA-MB-157 and MCF-7) [18] obviously represent the GnRH R gene product detected here in MDA-MB231 and MCF7 but not in T47 human breast cancer cell lines. The high GnRH neuropeptide levels detected in T47 might down-regulate receptor expression. The levels of gene expression in all the cell lines were lower that these detected in the breast cancer biopsies, but no direct comparisons can be done, due to the great variability between tissues and the heterogeneity and complexity of the cellular types in tissues (i.e. epithelial, stromal, macrophages etc. in the biopsies) vs. an epithelial cancer cell clone in the cell lines. This however might not be reflected at the translational level, as GnRH peptide expression has been shown before in MCF7 cells and extensively characterized with multiple approaches (HPLC, confocal microscopy) [5].
Co-expression of GnRH peptide and receptor in human breast tissues support an autocrine/paracrine role in human mammary gland. This is further supported by our analysis showing that receptor expression in the benign and malignant tissues correlated positively with the presence of GnRH neuroepithelium transcripts. In addition, there was a strong correlation between the presence of GnRH I and GnRH II transcripts in both the benign and the malignant biopsies, indicating the concomitant expression of both neuroepitope isoforms in breast, confirmed also by immunohistochemistry. Overall, it seems that the full neuroepitope and receptor system is simultaneously expressed in some tissues and this could be the result of inter-activation between the genes of the system, although this does not seem to be the case at least for the peptides [3]. Alternatively, co-activation of the three genes could be a concurrent response to the same stimulant. Several studies have shown differential regulation of the GnRH-I and GnRH-II genes by gonadotropins and steroid hormones in extrapituitary sites. However, other effectors and factors should be examined for their ability to transcriptionally activate GnRH genes [17] and breast cancer cell lines reported here to express both peptides and receptor could serve as a convenient in vitro model for such studies.

5. Conclusions

Growth effects of GnRH analogs on breast cancer cells depend on the amount of GnRH at cell surfaces and on receptor functionality [12,31], therefore quantification of expression levels holds a specific interest to predict cellular responses. Furthermore, GnRH receptors may contribute to the breast tumor responsiveness to pharmacotherapy with GnRH analogs and thus estimation of expression levels could define a more sensitive patient subpopulation. Our results add to the pathophysiological significance of the GnRH system in breast cancer biology possibly by contributing to the dialog between benign and malignant cell types, and imply that its quantification could complement the clinico-pathological profiling of breast cancer with potential clinical exploitation in prognosis and treatment.

Conflict of interest

All authors declare that they have no conflict of interest.

Contributors

Kalliopi Pazaitou-Panayiotou, Alexandros Kortsaris and Aikaterini Chatzaki designed the study, Christina Chemonidou, Aliki Poupi, Maria Koureta and Maria Koffa performed the experiments, Maria Lambropoulou, Theodoros C. Constantinidis and Grammati Galaktidou analyzed the results, Athina Kaprara, Anastasia Kaziiri- dou and Stylianos Kakolyris collected biologic material and clinical information, Kalliopi Pazaitou-Panayiotou, George Kolios and Aikaterini Chatzaki prepared the manuscript. All authors have approved the final article.

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