Heterogeneity of carboxyl-terminal parathyroid hormone circulating forms in patients with hyperparathyroidism due to end stage renal disease

Heterogeneidade das formas carboxi-terminal circulantes de paratormônio em pacientes com hiperparatiroidismo devido à insuficiência renal crônica terminal

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ABSTRACT

Objective: To study carboxyl-terminal (COOH) parathyroid hormone (PTH) circulating forms in patients with hyperparathyroidism due to end stage renal disease (ESRD). Methods: An immunometric assay that recognizes both intact and COOH PTH forms was developed. The assay, in conjunction with an intact assay, was used to measure PTH in serum samples obtained from 25 patients with hyperparathyroidism due to ESRD. Samples were also submitted to gel filtration chromatography in a Superdex® 30 1.6 x 60 cm column, and the PTH content in the elution tubes, measured using both assays. Results: Values from 39.000 to 232.300 ng/mL (mean ± sd = 101.680 ± 45.330 ng/mL) were found using the COOH assay (PTH 39-84 was used as standard). Values obtained by the intact PTH assay ranged from 318 to 3.307 ng/mL (1.769 ± 693 ng/mL) with a correlation between assays of 0.462 (p = 0.02). The elution profile obtained using the COOH assay showed a preponderance of forms with MW ranging from 8.500 to 4.500 daltons. The profiles obtained from the 25 patients were very similar. Conclusions: In patients with hyperparathyroidism due to ESRD circulating PTH levels contain a broad range of molecular forms including COOH with MW ranging from 8.500 to 4.500 daltons. These forms are not recognized by the standard intact PTH assays. The correlation of these findings to the clinical aspects of bone disease in ESRD patients remains to be studied.

Keywords

PTH assays; carboxy-terminal fragments of PTH; end-stage kidney disease; chromatography, gel

RESUMO

Objetivo: Estudar as formas carboxi-terminal (COOH) circulantes de paratormônio (PTH) em pacientes com hiperparatiroidismo devido à insuficiência renal crônica (IRC) terminal. Métodos: Foi desenvolvido um ensaio imunométrico que reconhece formas intactas e COOH longas de PTH. Esse ensaio foi utilizado, em conjunto com um ensaio para molécula intacta de PTH, em amostras de 25 pacientes com hiperparatiroidismo devido à IRC. As amostras também foram submetidas à cromatografia de gel filtração em coluna de Superdex® 30 de 1,6 x 60 cm, e o conteúdo de PTH nos tubos de eluato foi medido, empregando-se os dois ensaios. Resultados: Valores entre 39.000 e 232.300 ng/mL (média ± dp = 101.680 ± 45.330 ng/mL) foram obtidos usando-se o ensaio COOH (PTH 39-84 foi utilizado como padrão). O perfil cromatográfico obtido com o ensaio COOH mostrou predominio de formas com PM entre 8.500 e 4.500 daltons. Os perfis cromatográficos dos 25 pacientes foram bastante semelhantes. Conclusões: Em pacientes com hiperparatiroidismo devido à IRC, os níveis circulantes de PTH contêm um espectro de formas moleculares que incluem formas carboxi-terminais, com PM entre 8.500 e 4.500 daltons. Essas formas não são conhecidas pelos ensaios de rotina utilizados para a medida de PTH intacto. A correlação entre esses achados e os aspectos clínicos da doença óssea em pacientes com IRC necessita de maiores estudos.

Descritores

Ensaios para PTH; formas carboxi-terminais de PTH; insuficiência renal crônica; cromatografia em gel
COOH-terminal fragments of PTH in renal patients

INTRODUCTION

Parathyroid hormone (PTH) is a linear 84 amino acid peptide secreted by the parathyroid glands. The biological activity of PTH, defined by binding to the classical PTH1 receptor (PTH1R), depends on the integrity of the amino-terminal sequence of the peptide, mainly its first four amino acids (1).

Methods for measuring circulating PTH have evolved significantly since the introduction of the first radioimmunoassay in 1963 (2,3). Several competitive immunoassays were described during the 1970’s and 1980’s. Reference ranges as well as diagnostic sensitivity and specificity obtained with those assays were markedly different. The differences mirrored the wide spectrum of circulating forms of PTH recognized by the several polyclonal antibodies used in the competitive assays. With the description, in the late 1980’s, of the so-called “second-generation non-competitive immunometric assays” (“sandwich”-type), supposedly measuring only “intact” PTH (sequence 1-84), there was a clear improvement of diagnostic sensitivity and specificity, as well as a better standardization of assay results (4).

Nonetheless, new studies suggested that, in some clinical circumstances, mainly in patients with end stage renal disease (ESRD), additional molecular forms, besides the sequence 1-84, were also measured by these intact PTH assays (5). Included were carboxyl-terminal (COOH) fragments lacking only the first amino-terminal amino acids (e.g. peptide 7-84), also known as “long” COOH fragments. These forms are part of a broad spectrum of COOH-terminal forms which includes the quantitatively more important shorter forms (not recognized by the second-generation assays), known to circulate in high concentration in patients with ESRD (6).

Later “third-generation” assays were introduced (7) based on the employment of amino-terminal specific antibodies with strict specificity for the first four amino acids of the PTH molecule. Consequently, they measure only the bioactive molecule, via PTH1R, PTH-(1-84). These observations brought the possibility that, mainly in patients with ESRD, results with better correlation to PTH action (via PTH1R) would be obtained with these third-generation assays (7). In patients with ESRD, the new assay could mean a more specific tool for measuring PTH action on bone cells, since, in these patients, PTH is used as a surrogate marker for metabolic bone disease. Some publications showed that this could indeed be the case, in special if one measures PTH using both second- and third-generation assays and use both results to calculate a relationship between the PTH1R active ((1-84)-PTH) and non-active ((7-84)-PTH) forms (8).

In this paper, we describe an immunofluorometric assay that recognizes an ample spectrum of COOH-terminal forms of PTH. The levels of PTH found in patients with ESRD using the assay are compared with those obtained using an intact (second-generation) PTH assay. We also present the results obtained using both assays to measure PTH immunoreactivity in the eluate obtained by applying serum samples of ESRD patients to a gel filtration chromatography column.

METHODS

Samples

Twenty-five dialysis patients with ESRD submitted to parathyroidectomy in order to control severe hyperparathyroidism were studied. Blood samples were collected just prior to surgery and serum samples stored at -20 °C until studied. The protocol was approved by the Ethics Committee of our institution (Escola Paulista de Medicina, Universidade Federal de São Paulo) and was part of a large study on surgical and parathyroid tissue autotransplantation optimization protocols.
COOH assay

To measure COOH-terminal forms of circulating PTH, an immunofluorometric assay based on monoclonal and polyclonal antibodies was developed. The monoclonal antibody (H5P10) was produced by conventional fusion protocol based on mice immunized against synthetic human PTH-(53-84) and its specificity was directed against the region around amino acids 68-69 (14). The polyclonal antibody was developed in rabbits immunized against bovine PTH and had a broad specificity, including amino and COOH-terminal regions of human PTH. In the assay design, the COOH-terminal specific monoclonal antibody is adsorbed to a solid phase (microtiter plates, FluoroNunc™, Nunc, Roskilde, Denmark) at 10 µg/mL in PBS, 20 mM, pH 7.4 (200 µL/well). After an overnight incubation at 4 °C, the plates are washed and, in order to lower nonspecific binding, treated with 200 µL/well of TrisHCl 50 mM solution containing 0.5% bovine albumin and 0.05% bovine γ-globulin (assay buffer) for one hour at 37 °C. After a new wash, the plates were ready for the assay.

As standard, synthetic hPTH-(39-84) (Bachem AG, Bubendorf, Switzerland) was used diluted in assay buffer with 2% mouse serum. Aliquots of 100 µL of the standard samples (diluted 1:10 in assay buffer) or column eluate were added, together with 100 µL of the rabbit polyclonal antibody in a 1:2000 dilution. After an overnight incubation followed by washing, 200 µL of a 1:1 000 dilution of europium-labeled (Europium labeling kit, PerkinElmer, Turku, Finland) monoclonal anti-rabbit IgG antibody, produced at our laboratories, was added. After an additional a two-hour incubation, plates were washed and time-resolved fluorescence was read. The assay presented a sensitivity of 50 ng/L (hPTH-(39-84)), an inter-assay CV of less than 10%, and a cross-reactivity (based on a 100% cross-reactivity with hPTH-(39-84)) of 52.8% with hPTH-(7-84), 28.4% with hPTH-(1-84) and no cross-reactivity with the sequences 53-84, 44-68 and 69-84 of hPTH. The study of samples obtained from 30 adults with normal renal function showed values inferior to 500 ng/L.

Intact PTH assay

A monoclonal antibody-based immunofluorometric assay developed at our laboratory was employed. The assay employed two monoclonal antibodies, one amino-terminal specific (labeled with europium) and one COOH-terminal specific (the same H5P10 used in the long COOH assay). The assay had an analytical sensitivity of 3 ng/L and a cross-reactivity of 50% with hPTH-(7-84) (PTH-(1-84) was used as reference). Upper normal range for serum PTH using this assay was 46 ng/L (15).

Gel filtration chromatography

Serum samples were submitted to gel filtration chromatography on a Hiload Superdex 30, 1.6 x 60 cm column coupled to a FPLC Akta system (GE Healthcare). The gel column was chosen due to its fractio-nation range (globular proteins) with V0 of < 10,000 daltons. Serum samples of 0.5 mL were applied to the system, eluted (1 mL/min) with a PBS 0.05M, pH 7.2, 0.15M NaCl buffer and 1.0 mL aliquots were collected for PTH analysis using both assays. Column calibration included blue dextran (V0), synthetic human PTH-(1-84)(MW 9 425) was measured using the intact PTH assay, and synthetic human PTH-(39-84)(MW 4 962) was measured using the COOH immunometric assay. PTH measurement using the COOH and intact assays were performed in tubes number 40 to number 70 in all chromatographies. These tubes were chosen after calibration studies, and results of pilot serum samples (from patients with secondary hyperparathyroidism due to ESRD) indicated that they included all PTH forms detectable by both assays.

RESULTS

Correlation between serum values of intact and COOH-terminal PTH assays

PTH values in serum samples from the 25 patients with ESRD ranged from 318 to 3,307 ng/L (mean ± sd: 1,769 ± 693 ng/L) using the intact assay and from 39,000 to 232,300 ng/L (101,680 ± 45,350 ng/L) using the COOH assay. The coefficient of correlation (Pearson’s r) between the values obtained with the two methods was 0.462 (p = 0.02). Data are presented in figure 1.

Chromatography results

In figure 2, the chromatography profile obtained with the serum of one representative patient with intact PTH of 1,389 ng/L and COOH-terminal PTH of 118,450 ng/L was depicted. In the left Y-axis (ng/L), the PTH values measured in the eluate tubes, using both assays, are plotted. The void volume of the column (V0) as well as the elution of PTH-(1-84) (MW 9 425 daltons) and PTH-(39-84) (MW 4 982 daltons) are indicated by arrows, and the scale of MW (daltons) represented in the right Y-scale.
All immunoreactive PTH measured with the intact assay appeared in tubes 45 to 55, which is compatible with column calibration and assay specificity, since PTH (1-84) eluted with a peak around tube 50, as seen in figure 2. The elution of PTH forms detected with the COOH assay ranged from tube 45 to 65, encompassing forms with molecular weight from 10,000 to 4,500, according to the calibration profile. In order to evaluate the distribution of PTH forms recognized by the long COOH assay along the elution profile of the column, the elution profile obtained with the COOH assay was arbitrarily divided into four quartiles. The first one included tubes 45 to 50; the second, tubes 50 to 55; the third, from 55 to 60; and the fourth, from 60 to 65. The percent (mean ± sd) quartile distribution in the 25 samples was 7.1 ± 5.7 for the first quartile; 30.5 ± 10.8 for the second; 51.0 ± 11.1 for the third; and 11.4 ± 6.4 for the forth. In accordance to the molecular weight distribution, the intact assay recognized only PTH eluted in the first and second quartiles, and the COOH assay forms with MW ranging from 10,000 to 4,500 daltons, included in the four quartiles, with higher values found in the third.

**DISCUSSION**

PTH in circulation is quite heterogenous, with forms encompassing the complete 1-84 peptide to small fragments consisting mainly of COOH sequences. These last ones are known to accumulate in patients with progressive degrees of renal failure since their metabolic clearance depends heavily on glomerular filtration. The possibility that PTH fragments lacking the amino-terminal sequence and, consequently, biological action via PTHR1 can have some biological action is not new (16,17). Experimental evidence of the presence of a COOH receptor is summing up (18,19). The clinical importance of measurement of these COOH fragments is still not clear but is based mainly on data provided by intact PTH assays that measure, besides the 1-84 form, only molecular forms with small amino-terminal deletions, e.g. PTH (7-84) (20). The other methodological alternative is the use of old COOH specific competitive assays that measure almost all circulating forms, from the intact molecule to small COOH fragments (6).

In the setting of ESRD, the question is particularly complex and is still subject to research and controversy (21,22). Our results confirm the presence of high levels of COOH fragments of significant molecular weight (> 4500 daltons) in serum samples obtained from patients with ESRD. These forms include the PTH 7-84 (MW 8871 daltons), but also include large quantities of COOH molecular form fragments, shorter than 7-84. One interesting observation is the profile of forms detected by the COOH assay running from intact PTH to forms around 4500 daltons. These COOH PTH molecules have MW ranging from 8500 to 4500 and correspond to deletions of 10 to 40 amino acids. They are represented in the chromatographic system employed by forms eluting in the third and fourth quartiles. These peptides are not recognized by the second-generation intact assays, have no activity in the PTH1R, but bind to
the C-PTH receptor with the same affinity of PTH-(1-84) and (7-84) (12). One additional point that must be taken into account in the analysis of the elution profile is the specificity of the assay developed that spans from 100% with PTH-(39-84) to 28% with PTH-(1-84).

The present findings are in accordance with those described by Zhang and cols. (23) in a recent publication, which used immunoaffinity and mass spectrometry to detect COOH fragments in samples from patients with ESRD. They described peptides ranging from 34-84 to 45-84 to be the most abundant. These forms are, at least in part, detected by our COOH-specific assay. Methodological details could explain some potential differences in the detected forms, and can be related to the recognition of the antibodies used in the present assay or in the affinity purification used by Zhang and cols. (23).

The relative uniformity in the PTH-elution profile observed in the 25 patients with ESRD, confirmed by the low CV’s of the percent distribution in the percentiles, indicate that it may be difficult to use this protocol to classify the bone disease presented by these patients. More studies correlating bone histomorphometry studies to PTH levels measured using assays like the one described here are necessary in order to define a potential clinical utility for the measurement of these long COOH terminal fragments of PTH in these patients.

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