Characterization of the human and mouse genes encoding the tuberoinfundibular peptide of 39 residues, a ligand of the parathyroid hormone receptor family

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Abstract

The polypeptide TIP39 (tuberoinfundibular peptide of 39 residues) is a potent activator of the parathyroid hormone (PTH)-2 receptor (P2R) and an antagonist of the PTH-1 receptor (P1R). To clarify its possible physiological function(s), we studied its interaction with the human P1R and P2R and examined the expression of TIP39 in man and mouse. To find out possible sites of this ligand interaction in the organism, we identified the genes encoding the TIP39 protein precursors of Homo sapiens and Mus musculus in the databases of the human and mouse genome projects respectively. We then obtained the full-length cDNAs of both species by RACE-PCR. The deduced TIP39 preprohormones consist of an N-terminal 30 amino acid (aa) signal peptide followed by a 29 aa TIP39 precursor-related peptide, an Arg-Arg processing site, and the actual 39 aa TIP39 sequence. The first 23 aa of the actual TIP39 sequence, thought to contain the P2R receptor activation site, are identical in man and mouse and thus phylogenetically conserved. By contrast, the 16 aa C-terminal portion showed a higher degree of diversity (75% aa identity). By using RT-PCR, TIP39 was found to be highly expressed in human central nervous system tissues, trachea, fetal liver, and, to a lesser degree, in human heart and kidney. Using in situ hybridization, TIP39 mRNA expression was revealed in various areas of the mouse brain. In a homologous human cell model using human embryonic kidney 293 cells stably transfected with human P1R and P2R, human TIP39 did bind to P1R with moderate affinity (IC₅₀ ~ 10⁻⁷–10⁻⁶ M), but showed higher affinity binding to P2R (IC₅₀ ~ 10⁻⁵ M), comparable to the affinity of human N-terminal PTH (hPTH(1–34)) to this receptor. In P2R-transfected cells, the cAMP pathway was activated more efficiently (~10-fold) by TIP39 as a ligand compared to hPTH(1–34). In P1R-transfected cells, only hPTH(1–34) but not TIP39 was able to elicit a cAMP response, but TIP39 was able to directly antagonize the cAMP-stimulating effect of hPTH(1–34) on this receptor. In conclusion, we could show a possible function of TIP39 for the human organism as a potent activator of P2R (e.g. in brain) as well as an antagonist of the action of PTH and/or PTH-related protein on P1R (e.g. in bone and kidney). The physiological role of TIP39 in calcium metabolism with regard to these actions remains to be determined. The tools developed in this work will allow us to investigate the possible role of TIP39 as a locally or systemically secreted ligand modulating the function of the PTH receptor family.


Introduction

The polypeptide TIP39 (tuberoinfundibular peptide of 39 residues) has been discovered recently in bovine hypothalamus (Usdin et al. 1999b) as a third member of the parathyroid hormone (PTH) ligand family, which now consists of PTH, PTH-related protein (PTHrP) and TIP39 (Usdin et al. 2000). Whereas PTH plays a major role in calcium metabolism and PTHrP regulates bone and cartilage development (among other functions), the biological function of TIP39 is largely unknown. It shows a limited homology with PTH, only 9 out of 39 residues are identical in the bovine amino acid (aa) sequence (Usdin et al. 1999b, Piserchio et al. 2000). TIP39 is a potent activator of the PTH-2 receptor (P2R), the physiological function of which is also unknown at present (Usdin 2000). However, TIP39 also acts as an antagonist on the PTH-1 receptor (P1R), to an extent which varies in different species (Hoare et al. 2000). TIP39 has been isolated from bovine hypothalamus due to its capability of activating the P2R. Since the publication of this discovery in November 1999 (Usdin et al. 1999b), there are still only very few data about the possible function(s) of this peptide and where it might be expressed in the organism, besides
the hypothalamus. A possible function of TIP39 in this hypothalamic region has been suggested recently by experiments showing some influence on hypothalamo–pituitary axes (Ward et al. 2001). To clarify its possible physiological function(s), we studied the interaction of human TIP39 in a homologous human model system with the human P1R and human P2R and examined the expression of TIP39 in man and mouse.

Materials and Methods

Cell lines, peptides and animals

The cDNA of P1R (kindly provided by Dr Harald Jüppner, Boston, MA, USA) and P2R (kindly provided by Dr T Usdin, Bethesda, MD, USA) were subcloned into the expression vector pCEP4 (Invitrogen, San Diego, CA, USA) and used to generate human embryonic kidney (HEK) 293 cells stably expressing the receptor proteins, as described (Blind et al. 1995). The cells were maintained in DMEM medium with Glutamax, containing 10% fetal calf serum, 100 U/ml penicillin G, and 100 µg/ml streptomycin in a humidified atmosphere of 95% air/5% CO2 at 37 °C. Human PTH(1–34) (hPTH(1–34)) (hPTH(1–34)) was purchased from Bachem (Heidelberg, Germany). TIP39 was custom synthesized by Immundiagnostik (Bensheim, Germany) using the published aa sequence (Usdin et al. 1999b).

Adult male CD-1 mice were purchased from Charles River Breeding Laboratories (Sulzfeld, Germany). Animals were kept under standard laboratory conditions with a 12 h light:12 h darkness cycle. All efforts were made to minimize both the suffering and number of animals used and all procedures were performed according to the accepted standards of good animal care.

Radioligand binding assays

Competitive binding studies were carried out in intact cell assays by displacement of the radioactive ligand [125I]-Nle8,21-Tyr34-rat PTH(1–34)-amide (NEN, Boston, MA, USA) and used to generate human embryonic kidney (HEK) 293 cells stably expressing the receptor proteins, as described (Blind et al. 1995). The cells were maintained in DMEM medium with Glutamax, containing 10% fetal calf serum, 100 U/ml penicillin G, and 100 µg/ml streptomycin in a humidified atmosphere of 95% air/5% CO2 at 37 °C. Human PTH(1–34) (hPTH(1–34)) was purchased from Bachem (Heidelberg, Germany). TIP39 was custom synthesized by Immundiagnostik (Bensheim, Germany) using the published aa sequence (Usdin et al. 1999b).

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Measurement of total inositol phosphate (IP) turnover and of accumulated intracellular cAMP

The determination of accumulated total IPs was carried out in cells metabolically prelabeled with [3H]myoinositol (Amersham, Freiburg, Germany) as described previously (John et al. 2001). To improve detectability of rather small degrees of activation of this pathway, the protein kinase inhibitors H-89 (30 µM) and GF 109203X (6 µM) (both from Bachem) were added, since we had shown previously that blocking protein kinases A and C with these substances resulted in an enhanced IP response (Blind et al. 1996).

To stimulate intracellular formation of cAMP, cells were incubated in 12-well plates with DMEM medium containing 1% BSA, 20 mM Hepes buffer and 1 mM 3-isobutyl-1-methylxanthine at 37 °C for 15 min together with the test substances. After removal of the supernatant, accumulated intracellular cAMP was extracted with 1 ml 95% ethanol, pH 3. After 2 h, the alcohol was removed by evaporation and cAMP measured by RIA (Beckmann Coulter, Unterschleissheim, Germany).

Data analysis

Data for ligand-stimulated second messenger accumulation and inhibition of radioligand binding were analyzed using the software package Prism (GraphPad Software, Inc., San Diego, CA, USA), which was also used to calculate EC50 and IC50 values.

Isolation of RNA from solid tissues and RACE

Total RNA was isolated by a commercially available modification (TRIzol; Invitrogen, Karlsruhe, Germany) of the one-step phenol/guanidinium thiocyanate method (Chomczynski & Sacchi 1987). Poly A+ RNA was isolated using the Oligotex mRNA Mini Kit (Qiagen, Hilden, Germany).

RACE was performed using the SMART RACE cDNA Amplification Kit (Clontech, Alameda, CA, USA) following the manufacturer’s instructions. One microgram of poly A+ RNA was used as starting material. The following synthetic oligonucleotides were used as primers for RACE-PCRs: Homo sapiens 5’-RACE (5’-CTGCAC GTTAGGGACTGTGCGGGAAGCTTG-3′), Homo sapiens 3’-RACE (5’-CTGCAC GTTAGGGACTGTGCGGGAAGCTTG-3′), Mus musculus 5’-RACE (5’-CTGCAC GTTAGGGACTGTGCGGGAAGCTTG-3′), Mus musculus 3’-RACE (5’-CTGCAC GTTAGGGACTGTGCGGGAAGCTTG-3′). RACE products were analyzed in 1.5% agarose gels. Isolated bands were cut out of the gel and agarose was removed using Ultrafree-DA spin columns (Millipore, Eschborn, Germany). The cDNAs were inserted into the PCRII-TOPO vector (Invitrogen) following the manufacturer’s instructions. Sequencing was performed by Toplab ( Martinsried, Germany).

Labeling of RNA probes with digoxigenin-UTP (DIG-UTP) by in vitro transcription

In vitro transcribed RNA (cRNA) was synthesized from cDNA fragments cloned into the plasmid PCRII-TOPO...
(Invitrogen) using T7 or SP6 polymerase respectively. Plasmids were linearized by digestion with an appropriate restriction enzyme. Reactions were performed according to protocols of the DIG RNA labeling kit (Roche, Basel, Switzerland).

In situ hybridization on cryosections

Total brains of adult mice were dissected, carefully frozen with dry ice powder and stored at −80 °C. Cryosections (10 µm) were transferred to SuperFrostPlus microscope slides (Roth, Karlsruhe, Germany) and fixed for 15 min in 4% para-formaldehyde/PBS at 4 °C. In situ hybridization was performed as described in the Nonradioactive In Situ Hybridization Application Manual (Roche Molecular Biochemicals, 2nd edition). The reactive structures were visualized by colorimetric reaction (4-nitroblue tetrazolium chloride/5-bromo-4-chloro-3-indolyl-phosphate). The specimens were mounted in Aquamount (Dako, Hamburg, Germany) and analyzed under the microscope.

RT-PCR

The reactions were carried out using total RNA (250 µg) as template and the Qiagen OneStep RT-PCR Kit (Qiagen) according to the manufacturer’s instructions. The PCR products were analyzed in 1.5% agarose gels stained with ethidium bromide. The following synthetic oligonucleotide pairs were used as primers for Homo sapiens, and for mouse genome database was identified. Starting with mouse genome a sequence derived from the NCBI locus on the long arm of chromosome 19 at band 19q13.3. A database search in the human genome using the aa sequence described in the literature revealed a TIP39 gene locus on the long arm of chromosome 19 at band 19q13.3. In the mouse genome a sequence derived from the NCBI mouse genome database was identified. Starting with templates of mouse and fetal human brain, we obtained human and murine cDNAs by RACE-PCR (submitted to GenBank and available under the GenBank accession numbers: AY037555 and AC073740). An alignment of the nucleotide sequences of the human and murine cDNAs revealed 80% identity. The human cDNA consists of a 5′-untranslated sequence of 102 bp, an open reading frame of 300 bp, and 55 bp of 3′-untranslated sequence containing a polyadenylation signal (Fig. 1A and B). The genes of both man and mouse consist of two exons separated by an intron at corresponding sites. Exon 1 encodes 43 aa of the TIP39 precursor protein, exon 2 encodes the remaining 57 aa (Fig. 1C).

A computational analysis (Nielsen et al. 1997) of the 100 aa prepro sequence indicates that the first 30 aa most probably function as a signal peptide that directs the polypeptide chain to the endoplasmatic reticulum. The TIP39 precursor also contains two possible cleavage sites (Arg-Arg motif with compatible adjacent residues). The first separates an intercalated peptide from the secreted peptide (Fig. 1D), the second is found at position 22/23 within the actual TIP39 sequence, suggesting that they may play a role in the processing of TIP39.

Comparison of the primary structure of the human and mouse preproTIP39 and the partially known bovine sequence using the Clustalw software shows a 100% identity between Homo sapiens and Bos taurus within the actual TIP39 sequence. Man and mouse share 79% overall identity and 89% identity within the secreted TIP39 peptide itself (Fig. 1F). The identities of the signal peptides comprise 77%, the intercalated peptides 68%. The actual TIP39 sequence shows the highest degree of identity in the N-terminal region.

Distribution of TIP39 mRNA in man

By RT-PCR we screened human tissues for TIP39 mRNA expression (Fig. 2). Using templates of fetal and adult tissues we obtained strong bands in fetal and adult brain, cerebellum and trachea. Furthermore, there was evidence for TIP39 mRNA synthesis in spinal cord, fetal liver, kidney and heart. No response was detected in adult liver, lung, placenta and adrenal gland.

Expression of murine TIP39 mRNA in mouse brain

To evaluate the expression pattern of TIP39 mRNA, a DIG-labeled RNA probe was constructed and the distribution examined by in situ hybridization on cryosections from mouse brain (Fig. 3). A wide range of hybridization intensities was observed, with many positive neurons throughout all regions of the nervous system. In the cerebral cortex and subcortical areas, e.g. septal nuclei and caudate-putamen, many neurons were densely labeled whereas the glia cell-rich corpus callosum remained unstained. In the cerebellum, hybridization signals were found in Purkinje cells, and in cells of the molecular layer (probably basket and stellate cells), but were nearly absent in the neuron-rich granular cell layer.
Comparison of PTH and TIP39 actions on the human PTH receptors

We directly compared ligand binding of human TIP39 and hPTH(1–34) to the human P2R and P1R in an intact cell radioligand assay as described above. In both P1R-transfected and P2R-transfected cell lines, the peptides were able to displace the radioactive ligand (Fig. 4A and B). Non-specific binding was less than 20% of the total amount of tracer bound. Untransfected HEK 293 cells showed no specific binding of the radioligand (data not shown). hPTH(1–34) did bind to both human PTH receptor-transfected cell lines with similar affinities (P1R: IC$_{50}$ = 19 nM; P2R: IC$_{50}$ = 16 nM). Binding affinity of TIP39 was moderate with P1R (IC$_{50}$ = 333 nM). It showed higher binding affinity to P2R (IC$_{50}$ = 11 nM), comparable to the affinity of hPTH(1–34) for both receptors (Fig. 4A and B).
The activation of the cAMP signaling pathway of P1R and P2R was measured by standard RIA after incubation with increasing concentrations of hPTH(1–34) and TIP39. The P1R could only be activated by hPTH(1–34) (EC50=0·21 nM), whereas TIP39 in concentrations of up to 1 µM showed no detected increase in accumulated intracellular cAMP (Fig. 4C). In P2R-transfected cells, the cAMP dose–response curve was shifted about 10-fold to the left with TIP39 as a ligand, compared to hPTH (1–34) (EC50=0·36 nM and 3·50 nM respectively). The maximal increase in cAMP accumulation was similar with both peptides (Fig. 4D).

Additionally, the activation of the phospholipase C pathway was estimated by measuring ligand-induced IP hydrolysis. Incubation with 3 µM hPTH(1–34) caused a strong increase in intracellular total IP (about 10-fold) in P1R-transfected cells and a small increase (about 2-fold) in P2R-transfected cells. TIP39, however, was not able to elicit a response of the IP signaling pathway in either cell line (data not shown).

To test the antagonistic activity of TIP39 on P1R directly, we exposed P1R-transfected cells to increasing amounts of TIP39 up to 10 µM in the presence of 0·25 nM hPTH(1–34). TIP39 almost completely inhibited agonist-induced cAMP accumulation with an IC50 value of half-maximal inhibition of approximately 1 µM (Fig. 4E).

Discussion

TIP39 may exert its biological function via two different mechanisms, either by inhibiting the action of PTH and/or PTHrP on P1R, or by acting independently on P2R. We have shown in our homologous human cell model system that human TIP39 acts as a competitive antagonist to the biologically active part of hPTH (hPTH(1–34)) on the human P1R. Additionally, we showed for the first time that TIP39, unlike PTH on P1R, does not activate the IP signaling pathway in P1R or P2R. Our results are in accord with published results of similar model systems, often using heterologous systems with ligands, cells or receptors from other species, however (Hoare et al. 1999, 2000, Jonsson et al. 2001). It is thus conceivable that locally secreted TIP39 could inhibit the action of systemically circulating PTH, especially since the physiological levels of the bioactive part of this peptide are rather low in the circulation, in the range 0·5–3 pM (Gao et al. 2001). This would be especially relevant for the classic target organs of PTH in calcium homeostasis, bone and kidney. However, there are no published data on whether TIP39 is expressed in these organs. We were able to show that TIP39 mRNA is indeed present in the kidney, when looked at by RT-PCR. Whether TIP39 acts as a competitor of P1R, or as an activator for P2R in these tissues is unclear at present. Some evidence comes from investigations with P2R, however. Although P2R mRNA was not detected by Northern blot in renal tissue (Usdin et al. 1995), Usdin et al. (2000) could show small numbers of cells near the vascular pole of kidney glomeruli to be P2R-positive by antibody staining and by in situ hybridization for P2R mRNA. These cells possibly belong to the juxtaglomerular apparatus, leading to speculations that their function might involve regulation of blood pressure as a target of a renin-releasing factor, which has been shown to be released from the hypothalamus and which is indistinguishable from TIP39 by size (Urban et al. 1992). We have no data yet on whether TIP39 is expressed in bone. It is also not clear at present whether P2R, as the other possible target besides the osteoblastic P1R, is present in bone, since no such study has been performed in this tissue so far. Attempts to detect P2R mRNA by RT-PCR in bone-derived cells have yielded equivocal results (Usdin et al. 2000).

Thyroid C-cells are another tissue linked to calcium metabolism where P2R was found to be expressed (Usdin...
et al. 1999a). C-cells were specifically labeled by a P2R-specific antibody whereas the surrounding majority of follicular thyroid cells were negative. Since C-cells also express the calcium sensing receptor and respond with secretion of bioactive peptides upon an increase in serum calcium, there might be a link of P2R to the regulation of calcium metabolism. We have no data yet, however, on whether TIP39 is involved in this process.

As estimated by Northern blot, P2R is expressed at particularly high levels in brain. Unlike in peripheral tissue, TIP39 seems to be the single probable ligand of this receptor in brain, since investigators have failed to detect PTH mRNA in rat brain tissue (Usdin 1997), and since most regions of cerebral P2R expression would not be reached by PTH from the blood stream. The pattern of P2R distribution in the rat nervous system has been extensively studied recently (Wang et al. 2000). It shows a widespread expression, most often in discrete groups of neurons, with an especially high level of expression in hypothalamic, limbic and sensory areas. The pattern of distribution of TIP39 expression found in our studies seemed to differ significantly, with TIP39 showing a distinct distribution pattern, which was, however, also widespread within various cerebral regions. A more detailed study is, therefore, needed to locate TIP39 expression in comparison to the P2R expression pattern.

We could detect TIP39 mRNA in human spinal cord. The rat P2R was shown to be expressed there, too, localized to the superficial dorsal horn, spinal trigeminal tract and nucleus. Because these areas receive their principal input from sensory neurons involved in pain perception (Usdin et al. 1999b), this colocalization might hint at a possible TIP39/P2R interaction modulating nociceptive function.

An analysis of the deduced aa sequence of the TIP39 preprohormones of man and mouse clearly shows the

**Figure 3** Distribution of TIP39 mRNA in coronal sections of mouse brains visualized by *in situ* hybridization. A broad range of hybridization intensities was observed, with many positive neurons throughout all regions of the central nervous system. (a) Overview of a mouse brain cryosection (× 12.5). CCA, corpus callosum; CCI, cingulate cortex; Cpu, caudate-putamen; CFr, frontal cortex; DS, dorsal septal nucleus. (b) Inset of (a), showing cortical neurons in different layers stained with various intensity (× 100). (c) Inset of (a), showing absence of TIP39 mRNA in glia cells of the corpus callosum (× 100). (d) Overview of mouse cerebellum cryosection (× 12.5). GL, granular cell layer; PL, Purkinje cell layer; ML, molecular layer. (e) Inset of (d), showing distinct staining of Purkinje cells and of cells in the molecular layer, whereas staining was largely absent in the granular cell layer (× 100). (f) Cortical area corresponding to the area shown in (b) (× 100). *In situ* hybridization was performed using a DIG-labeled TIP39 sense RNA probe as a negative control (× 12.5).
features of secreted neuropeptides with a characteristic signal peptide for the secretory pathway and a dibasic cleavage site for subtilisin-like endoproteases to process the prohormone (Hosaka et al. 1991).

The overall structural features of the preprohormones of TIP39, PTH and PTHrP – signal peptide, intercalated peptide and hormone – are identical. However, the intercalated peptide of TIP39 comprises 31 aa, and is much longer than the predicted intercalated peptides of PTH and PTHrP (8 aa). The overall aa sequence similarity between these three ligands is very low, whereas the tertiary structure of the portion interacting with P1R

Figure 4  (A, B) Radioreceptor assays. HEK 293 cells stably expressing P1R (A) or P2R (B) were exposed to varying amounts of human TIP39 (●) or hPTH(1–34) (○) to displace the radiolabeled ligand. (C, D) Activation of the cAMP signaling pathway. Ligand-induced accumulation of intracellular cAMP in HEK 293 cells stably expressing P1R (C) or P2R (D). Cells were exposed to varying amounts of human TIP39 (●) or hPTH(1–34) (○). (E) Competition of TIP39. Ligand-induced accumulation of intracellular cAMP in HEK 293 cells stably expressing P1R was measured with varying amounts of human TIP39 in the presence of 0.25 nM hPTH(1–34). The x-axis shows the log of the ligand concentration. Data are the means of triplicate determinations ± S.D.
seems similar (Piserchio et al. 2000). PTH, PTHrP and TIP39 thus seem to have only a distant phylogenetic relationship.

The N-terminal alpha helix of TIP39, which seems to be responsible for P2R activation, is identical in man and mouse, whereas the C-terminal alpha helix, which is important for receptor binding (Piserchio et al. 2000), differs in four aa positions, being apparently responsible for the different binding properties of murine and human TIP39 (Gould et al. 2001).

There is no experimental evidence yet that the predicted dibasic cleavage site found between the two alpha helices is functional. One might speculate that it is involved in the degradation and/or inactivation of TIP39, or that the possible cleavage products have bioactivity on their own. While it seems certain that TIP39 is in fact a secreted peptide, it is entirely unknown whether detectable amounts of TIP39 can be found in plasma and whether therefore this peptide could act systemically.

In conclusion, we could show that in the human organism TIP39 is expressed in various tissues and might function as a potent activator of P2R (e.g. in brain) as well as an antagonist of the action of PTH and/or PTHrP on P1R (e.g. in bone and kidney). The physiological role of TIP39 in calcium metabolism with regard to these actions, e.g. as a locally secreted ligand modulating the function of the PTH receptor family, remains to be determined, however. The tools developed in this work will allow further investigation of the function of this new ligand of the PTH receptor family.

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