SHORT COMMUNICATION

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Isolation and chromosomal mapping of a novel human gene showing homology to Na⁺/PO4 cotransporter

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Abstract We isolated a cDNA clone which shows a significant similarity with the renal Na⁺/phosphate cotransporter (NPT) from a human intestine mucosa cDNA library. The cDNA is 2626 bases long, with one open reading frame encoding a protein of 497 amino acids. The deduced amino acids sequence shows an overall homology of 48% with the human renal NPT1 protein. This gene is expressed in intestine, colon, liver, and pancreas. Thus, this gene may code for intestinal type NPT or closely related proteins. The chromosomal location of the gene was determined on the chromosome 6p21.3-p22 region by polymerase chain reaction-based analysis with both a human/rodent mono-chromosomal hybrid cell panel and a radiation hybrid mapping panel.

Key words Na⁺/Phosphate cotransporter (NPT) · Intestine · Chromosome 6p21.3-p22 · RT-PCR · cDNA library

Introduction

Phosphate homeostasis in humans is maintained by the balance between intake, intestinal absorption, bone deposition and resorption, and renal excretion. The central molecule for the control of kidney excretion is the Na+/phosphate cotransporter (NPT), which is located in the proximal tubule (Biber 1989, Biber et al. 1993). This molecule uses the transmembrane electrochemical potential gradient of so-

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Genome Research Group, National Institute of Radiological Science, Chiba, Japan dium to transport phosphate across the cell membrane (Murer et al. 1991). A similar NPT also plays an important role in the absorption of the phosphate in the intestine mucosa (Murer et al. 1994). While human NPT of the kidney has been cloned (Chong et al. 1993) and studied, the molecular structure of human intestinal NPT is yet to be elucidated. Here, we report the isolation of a novel cDNA with significant homology to the renal NPT1 from the human intestinal mucosa cDNA library.

Results and discussion

A full length-enriched cDNA library was constructed, using the oligo-capping method (Maruyama and Sugano 1994, Suzuki et al. 1997) with mRNA isolated from normal mucosa of ileum. The sequences of the 5' ends of the cDNA clones from the library were determined for 3150 clones. One clone, kaia2138, showed significant homology to human renal NPT1 cDNA. We determined the entire sequence of this clone. The nucleotide sequence data reported here will appear in the DDBJ, EMBL, and GenBank nucleotide sequence databases, with accession number AB020527. The cDNA was 2626 bp in length and has one open reading frame encoding a protein of 497 amino acids. A homology search revealed that it is homologous to the NPT family. Recently, human NPT3 and NPT4 cDNAs, homologous to human NPT1, were isolated from a 1.1-megabase region of the hereditary hemochromatosis locus of human chromosome Their nucleotide sequences have been submitted to GenBank. The multiple alignments of amino acid sequences of the kaia2138 protein and NPTs are shown in Fig. 1. The kaia2138 protein was most homologous to the human NPT3 protein, having 54% identity at the amino acid level. The amino acid sequence of kaia2138 was 48% identical to human NPT1 and 43.5% identical to human NPT4.

The expression level of this gene was not detectable by Northern blot analysis, so we examined the tissue distribution of the transcript in various human adult tissues by reverse transcription-coupled polymerase chain reaction (RT-PCR). Primers used for RT-PCR were to amplify the

Fig. 1. Multiple alignment of kaia2138 (accession number AB020527) and Na¹/phosphate cotransporter (NPT) families. The NPT sequences are human NPT1 (NPT1_HUMAN; accession number Q14916), rabbit NPT1 (NPT1_RABIT; accession number Q28722), rat NPT1 (NPT1_RAT; accession number Q62795), mouse NPT1 (NPT1_MOUSE; accession number Q61983), human NPT3 (NPT3_HUMAN; accession number O00624) and human NPT4 (NPT4_HUMAN; accession number O00476). Identities are indicated by shadowed background. Asterisks denote the terminal codon

kaia2138 NPT1_HUMAN NPT1_RABIT NPT1_RAT NPT1_MOUSE NPT3_HUMAN NPT4_HUMAN	1:MSTGPDVKATVGDISSDGNLNVAQEECSRKGFCSVRHGLALILQLCNFSIYTQOMNLSIA 1:	41 41 41 41 41 41 43
NPT1_HUMAN NPT1_RABIT NPT1_RAT NPT1_MOUSE NPT3_HUMAN NPT4_HUMAN	42: MVMWNSTDPHGLPNTSTKKLLDNIKNPMYNWSPDIQGIILSSTSYC 42: MVAMVNNTNLHGSPNTSAEKRLDNTKNPVYNWSPDVQGIIFSSIFYG 42: MVAMVNNTGSPHLSNKSVAEMLDNVKNPVHSWSLDIQGLVLSSVFLG 42: MVAMVNNTGSPHLSNESVVEMLDNVKNPVYSWSPDIQGLILSSVFFG 42: IIAMVNTTQQQGLSNASTBGPVADAFNNSSISIKEFDTKASVYQWSPETQGIIFSSINYG 44: MVAMVNSTSPQSQLNDSSE	88 88 88 88 101
kaia2138 NPT1_HUMAN NPT1_RABIT NPT1_RAT NPT1_MOUSE NPT3_HUMAN NPT4_HUMAN	121:SFLAPIPSGYVAGIFGAKYVVGAGLFISSFLTLFIPLAANAGVALLIVLRIVQGIAQVMV 89:VIIIQVPVGYFSGIYSTKKMIGFALCLSSVLSLLIPPAAGIGVAWVVVCRAVQGAAQGIV 89:AFLIQIPVGYISGIYSIKKLIGFALFLSSLVSIFIPQAAAVGETWIIVCRVVQGITQGTV 89:MVVIQVPVGYLSGAYPMEKIIGSSLFLSSLVSLLIPFAAQVGAALVIVCRVLQGIAQGTV 89:MVVVQAPVGYLSGIYPMKRIIGSSLFLSSLMSLLIPPAAQVGAALVIVCRVLQGIAQGTV 102:IILTLIPSGYLAGIFGAKKMLGAGLLISSLLTLFTPLAADFGVILVIMVRTVQGMAQGMA 63:VLPVDSFGGLSKAPKSL	148 148 148 148 161
kaia2138 NPT1_HUMAN NPT1_RABIT NPT1_RAT NPT1_MOUSE NPT3_HUMAN NPT4_HUMAN	181:LTGQYSIWVKWAPPLERSQLTTIAGSGSMLGSFIVLLAGGLLCQTIGWPYVFYIFGGIGG 149:ATAQFETYVKWAPPLERGRLTSMSTSGFLLGPFIVLLVTGVICESLGWPMVFYIFGACGC 149:TTAQHEIWVKWAPPLERGRLTSMSLSGFLLGPFIVLLVTGIICESLGWPMVFYIFGACGC 149:STGQHGIWVKWAPPLERGRLTSMTLSGFVMGPFIALLVSGFICDLLGWPMVFYIFGIVGC 149:STGQHEIWVKWAPPLERGRLTSMTLSGFVMGPFIVLLVSGFICDLLGWPMVFYIFGIVGC 162:WTGQFTIWVKWAPPLERGRLTSMTLSGFSFIILCVGGIISQALSWPFIFYIFGSTGC 86:LGGQFAIWEKWGPPQERSRLCSIALSGMLLGCFTAILIGGFISETLGWPFVFYIFGGVGC	208 208 208 208 208 221
kaia2138 NPT1_HUMAN NPT1_RABIT NPT1_RAT NPT1_MOUSE NPT3_HUMAN NPT4_HUMAN	241: ACCPLWFPLIYDDPVNHPFISAGEKRYIVCSLAQQDCSPGWSLPIRAMIKSLPLWAILVS 209: AVCLLWFVLFYDDPKDHPCISISEKEYITSSLVQQVSSSRQSLPIKATLKSLPVWAISIG 209: AVCLLWFVLYYDDPKDHPCVSLHEKEYITSSLIQQGSSTRQSLPIKAMIKSLPLWAISIG 209: VLSLFWFILLFDDPNNHPYMSSSEKDYITSSLMQQVHSGRQSLPIKAMLKSLPLWAIILN 209: VLSLSWFFLFFDDPKDHPYMSSSEKDYITSSLMQQVSSGRQSLPIKAMLKSLPLWAIILN 222: VCCLLWFTVIYDDPMHPCISVREKEHILSSLAQQPSSPGRAVPIKAMVTCLPLWAIFLG 146: VCCLLWFVVIYDDPFSYPWISTSEKEYIISSLKQQVGSSKQPLPIKAMLRSLPIWSICLG	268 268 268 268 268 281
kaia2138 NPT1_HUMAN NPT1_RABIT NPT1_RAT NPT1_MOUSE NPT3_HUMAN NPT4_HUMAN	301:YFCEYWLFYTIMAYTPTYISSVLQANLRDSGILSALPFVVGCICIILGGLLADFLLSRKI 269:SFTFFWSHNIMTLYTPMFINSMLHVNIKENGFLSSLFYLFAWICGNLAGQISDFFLTRNI 269:CFAYLWTYSRLIVYTPTLINSMLHVDIRENGLLSSLFYLFAWICGVIAGHTADFLMSRNM 269:SFAFIWSNNLLVTYTPTFISTTLHVNVRENGLLSSLFYLLAYICGIVAGQMSDFLLSRKI 269:SFAFIWSNSLLVTYTPTFISTVLHVNVRENGLLSSLFYLLAYICGILAGGMSDFFLTRRI 269:FFSHFWLCTIILTYLFTYISTLHVNIRDSGVLSSLFFIAAASCTILGGQLADFLLSRNI 282:FFSHFWLCTIILTYLFTYISTVHVNIRDNGLLSALFFIVAWVIGMVGGYLADFLLTK-K	328 328 328 328 328 341
kaia2138 NPT1_HUMAN NPT1_RABIT NPT1_RAT NPT1_MOUSE NPT3_HUMAN NPT4_HUMAN	361: LRLITIRKLFTAIGVLFPSVILVSLPWVRSSHSMTMTFLVLSSAISSFCESGALVNFLDI 329: LSVTAVRKLFTAIGVLFPATFGVCLPYLSSTFYSIVIFLILAGATCSFCLGGVFINGLDI 329: LSLTAIRKLFTAIGLLLPIVFSMCLLYLSSGFYSTITFLILANASSSFCLGGALINALDI 329: FSVVAVRKLFTTLGIFCPVIFVVCLLYLSYNFYSTVIFLTLANSTLSFSFCGQLINALDI 329: FSIVTYRKLFTTLGSFCPVIFIMCLLYLSYNFYSTVIFLTLANSTLSFSYCGQLINALDI 342: LRLITVRKLFSSLDMQVSSWESQ	388 388 388 388 388 382
kaia2138 NPT1_HUMAN NPT1_RABIT NPT1_RAT NPT1_MOUSE NPT3_HUMAN NPT4_HUMAN	421: APRYTGFLKGLLQVFAHIAGAISPTAAGFFISQDSEFGWRNVFLLSAAVNISGLVFYLIE 389: APRYFGFIKACSTLTCMIGGLIASTLTGLILKQDPESAWFKTFILMAAINVTGLIFYLIV 389: APRYYVFIKGVTTLIGMTGGMTSSTVAGLFLSQDPESSWFKIFLLMSIINVISVIFYLIF 389: APRYYGFLKAVTALIGIFGGLISSTLAGLILNQDPEYAWHKIFFLMAGINVTCLAFYLLF 389: APRYYGFLKAVTALIGMFGGLISSTLAGLILNQDPEYAWHKISFLMAGINVTCLYFYLF 383: SVRILSLVGGMSFSCLLQSTCLAMSFTSRLDKGNFKTOPKGOPLPASEDLKLQT* 325: APRYSSFLMGASRGFSSIAPVIVPTVSGFLLSQDPEFGWRNVFFLLFAVNLLGLLFYLIF	448 448 448 448 436
kaia2138 NPT1_HUMAN NPT1_RABIT NPT1_RAT NPT1_MOUSE NPT3_HUMAN NPT4_HUMAN	481:GRADVQDWAKEQTFTHL* 449:ATAEIQDWAKEKQHTTL* 449:AKGEIQDWAKEKQHTTL* 449:AKGDIQDWAKETKTTRL* 449:AKGEIQDWAKEIKTTRL* 437: 385:GEADVQEWAKERKLTLL*	497 465 465 465 465 465

407 bp of the 3'-untranslated region of the cDNA (Fig. 2). The *kaia2138* gene was mainly expressed in liver, pancreas, small intestine, and colon.

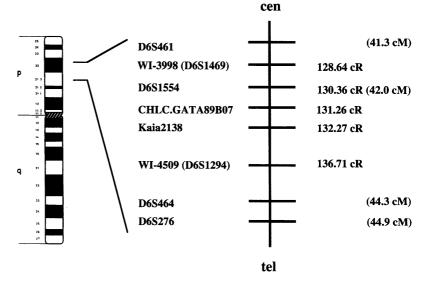
Chromosomal assignment of the *kaia2138* gene was done by PCR analysis of a human/rodent somatic cell hybrid panel

and a radiation hybrid panel, as described previously (Saito et al, 1997). The specific amplified PCR primers were designed at the 3'-untranslated region of the gene (5'-TTCTGG-CCATCTTGACTTCTG-3', 5'-CCAACAGTCAGAGGGG-CAAAC-3'; PCR product size was 178bp). First, a specific

Fig. 2. Reverse transcription-coupled polymerase chain reaction (RT-PCR) analysis of kaia2138 in multiple human tissues. Primers used for RT-PCR amplify 407 bp of the 3'-non-coding region of the cDNA. The primers were (5'-TACCTGTGCTCCAGAGTTAGC-3') and (5'-CTTATGATCCTCCATGC-TGC-3'). The cDNA templates for RT-PCR were from the human tissues of poly (A)+ RNAs and purchased from Clontech (Palo Alto, CA, USA). PCR was carried out in a final volume of 50 μl containing 1 × GC buffer I (Takara, Kyoto, Japan), 0.2 μM each primer, 400 µM each dNTP, 5 µl (~1 ng) of template cDNA and 2.5 units of LA-Taq DNA polymerase (Takara, Kyoto, Japan). For the positive control, 50 pg of the kaia2138 clone from a full-length enriched cDNA library was used for the template. The temperature and time schedules were: 35 cycles at 94°C for 1 min, 55°C for 1 min, and 72°C for 1 min. PCR products were separated on 2.0 % agarose gel. The lower panel shows the result with a Human G3PDH Control Amplimer Set The control G3PDH cDNA was also purchased from Clontech and used for PCR (5 µl [~1ng])

Control
Heart
Brain
Placenta
Lung
Lung
Liver
Skeletal muscle
Kidney
Pancreas
Spleen
Thymus
Prostate
Testis
Ovary
Small intestine
Colon
Leukocyte

Fig. 3. Chromosomal placement of the *kaia2138* gene at a relative distance to framework markers on the WICGR (Whitehead Institute for Biomedical Research/MIT Center for Genome Research) radiation hybrid map of the human genome (http://www.genome.wi.mit. edu/). The approximate corresponding cytogenetic location of the gene on the long arm of the telomeric (*tel*) region of human chromosome 6 is indicated. Distances of the markers are in centirays (cR) and centimorgans (cM) from the top of the chromosome 6 linkage group. *Cen*, Centromeric region



Chromosome 6

amplified product for humans was detected only from the hybrid containing human chromosome 6 (data not shown). Then, we performed further mapping analysis using a PCRbased radiation hybrid panel (Genebridge 4; Research Genetics) with the same primers as those used in the assay for the human/rodent somatic cell hybrid panel. Statistical analysis of the radiation hybrid data was performed using RHMAPPER software package (http://wwwgenome.wi.mit.edu/cgi-bin/contig/rhmapper.pl). The data vector for the gene was 1000000100 0010000000 1100000000 1001010100 1110010000 0001000001 0110000010 0011001100 1111010000 001 and the consequent report indicated that the gene was placed to 1.01 cR proximal from CHLC.GATA89B07 (lod>3.0), which was cytogenetically mapped to the 6p21.3-p22 region (Fig. 3). The human renal NPT1 gene was also mapped to 6p21.3-p23 (Chong et al. 1993). Furthermore, NPT3 and NPT4 were identified within a 1.1-megabase region of the hereditary hemochromatosis locus (Lauer et al. 1997), which is located at 6p21.3p22. It is possible that human NPT genes are clustered around the chromosome 6p21-p23 region.

References

Biber J (1989) Cellular aspects of proximal tubular phosphate reabsorption. Kidney Int 36: 360-369

 Biber J, Custer M, Werner A, Kaissling B, Murer H (1993) Localization of NaPi-1, a Na/Pi cotransporter, in rabbit kidney proximal tubules. II. Localization by immunohistochemistry. Pflugers Arch 424: 210-215

Chong SS, Kristjansson K, Zoghbi HY, Hughes MR (1993) Molecular cloning of the cDNA encoding a human renal sodium phosphate transport protein and its assignment to chromosome 6p21.3-p23. Genomics 18: 355-359

Lauer P, Meyer NC, Prass CE, Starnes SM, Wolff RK, Gnirke A (1997) Clone-contig and STS maps of the hereditary hemochromatosis region on human chromosome 6p21.3-22. Genome Res 7: 457-470

Maruyama K, Sugano S (1994) Oligo-capping: a simple method to replace the cap structure of eukaryotic mRNAs with oligo-ribonucleotides. Gene 138: 171-174

Murer H, Werner A, Reshkin S, Wuarin F, Biber J (1991) Cellular mechanisms in proximal tubular reabsorption of inorganic phosphate. Am. J Physiol 260: C885-899

Murer H, Markovich D, Biber J (1994) Renal and small intestinal sodium-dependent symporters of phosphate and sulphate. J Exp Biol 196: 167-181
Saito T, Seki N, Ishii H, Ohira M, Hayashi A, Kozuma S, Hori T (1997) Complementary DNA cloning and chromosomal mapping of a novel phosphatidylinositol kinase gene. DNA Res 4: 301-305

Suzuki Y, Yoshitomo-Nakagawa K, Maruyama K, Suyama A, Sugano S (1997) Construction and characterization of a full length and 5'-end-enriched cDNA library. Gene 200: 149-156