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**Title**

**Regulation of Tissue-Specific Expression of the Human and Mouse Urate  
Transporter 1 Gene by Hepatocyte Nuclear Factor 1  $\alpha/\beta$  and DNA Methylation**

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**Running title**

**Regulation of the URAT1 Expression**

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Abbreviations: OAT, organic anion transporter; URAT1, urate transporter 1; HNF1, hepatocyte nuclear factor 1; PCR, polymerase chain reaction; EMSA, electrophoretic mobility shift assay; Gapdh, glyceraldehyde-3-phosphate dehydrogenase; T-DMR, tissue-dependent differentially methylated region; MODY, maturity onset diabetes of the young.

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## Abstract

Expression of Urate transporter 1 (*URAT1/SLC22A12*) is restricted to the proximal tubules in the kidney, where it is responsible for the tubular reabsorption of urate. In order to elucidate the mechanism underlying its tissue-specific expression, the transcriptional regulation of the *hURAT1* and *mUrat1* genes was investigated. Hepatocyte Nuclear Factor 1  $\alpha$  (HNF1 $\alpha$ ) and HNF1 $\beta$  positively regulate minimal promoter activity of the *URAT1* gene as shown by reporter gene assays. Electrophoretic mobility shift assays revealed binding of HNF1 $\alpha$  and/or HNF1 $\beta$  to the HNF1-motif in the *hURAT1* promoter. Furthermore, the mRNA expression of *Urat1* is reduced in the kidneys of *Hnfl* $\alpha$ -null mice compared with wild-type mice, confirming the indispensable role of HNF1 $\alpha$  in the constitutive expression of *URAT1* genes. It was also shown that the proximal promoter region of *mUrat1* was hypermethylated in the liver and kidney medulla, whereas this region was relatively hypomethylated in the kidney cortex. These methylation profiles are in a good agreement with the proximal tubule-restricted expression of *mUrat1* in the kidney cortex. Taken together, these results strongly suggest that tissue-specific expression of the *URAT1* genes is coordinately regulated by the transcriptional activation by HNF1 $\alpha$ /HNF1 $\beta$  heterodimer and repression by DNA methylation.

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## Introduction

Urate is an end product of purine metabolism in humans and other higher primates. It is generally recognized that urate works as a scavenger of potentially harmful radicals in the human body. However, hyperuricemia can be associated with health problems such as gout, nephrolithiasis, hypertension, and vascular disease, while hypouricemia is primarily characterized by exercise-induced acute renal failure. Thus, the serum urate level must be tightly regulated through complex renal handling processes, which is historically explained by a four-component model, i.e. glomerular filtration, tubular secretion, and pre- and post-secretory reabsorption. Following these processes, approximately 90% of the urate filtered through the glomerulus is reabsorbed in humans (Enomoto and Endou, 2005; Hediger et al., 2005; Rafey et al., 2003).

Urate transporter 1 (URAT1/*SLC22A12*) is a membrane transporter responsible for the reabsorption of urate in the apical membrane of the renal proximal tubules (Enomoto et al., 2002). Mutations in the *URAT1* gene, causing functional impairment, are associated with idiopathic renal hypouricemia (Enomoto et al., 2002; Iwai et al., 2004). In addition, fractional excretion of urate in patients with homozygous or compound heterozygous *SLC22A12* mutations is not sensitive to uricosuric (probenecid and benzbromarone) and anti-uricosuric (pyrazinamide) drugs, suggesting that these drugs target URAT1 to exert their effects in vivo (Ichida et al., 2004). The mouse homologue of hURAT1, mUrat1, shows the same tissue distribution as that of hURAT1, and is probably involved in the reabsorption of urate in the kidney (Hosoyamada et al., 2004). It can also be an exit pathway for several anionic drugs in the proximal tubules together with the basolateral uptake transporters (Organic anion transporter (Oat) 1 and Oat3) (Imaoka et al., 2004).

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A previous report suggested the presence of a Hepatocyte Nuclear Factor 1 (HNF1) binding motif in the minimal promoter region of the *hURAT1* gene, which is conserved in the proximal promoters of rodent *Urat1* genes (Li et al., 2004). HNF1 is known to regulate transcription of many hepatic genes by forming homodimers or heterodimers between two isoforms, HNF1 $\alpha$  and HNF1 $\beta$  (Mendel and Crabtree, 1991; Tronche and Yaniv, 1992). Their expression in the kidney and a renal Fanconi syndrome or polycystic kidney disease in *Hnf1 $\alpha$* - or kidney-specific *Hnf1 $\beta$* -null mice, respectively, also imply a role for these transcription factors in the kidney (Gresh et al., 2004; Pontoglio et al., 1996). We have recently found that HNF1 $\alpha/\beta$  is a potent transactivator of the human and mouse *OAT3* promoter (Kikuchi et al., 2006). Subsequently, it was reported that the mRNA levels of several drug transporters, including Oat1 and Oat3, are reduced in the kidney of *Hnf1 $\alpha$* -null mice, further confirming the involvement of HNF1 $\alpha$  in the transcriptional regulation of drug transporters in the kidney (Maher et al., 2006). The functional importance of HNF1 in the regulation of the *URAT1* gene has yet to be investigated. However, considering the fact that the serum levels of urate are reduced in *Hnf1 $\alpha$* -null mice while the renal fractional excretion is increased (Pontoglio et al., 1996), it is possible that HNF1 $\alpha$  directly regulates URAT1 expression.

HNF1 $\alpha$  and HNF1 $\beta$  are expressed in the polarized epithelia of a variety of tissues, including liver, kidney, intestine, stomach, and pancreas (Blumenfeld et al., 1991; De Simone et al., 1991). Thus, positive regulation by HNF1 alone cannot account for the predominant expression of URAT1 in the kidney. Our previous study suggests that the expression of hOAT3 is negatively regulated by DNA methylation in addition to the positive regulation by HNF1 $\alpha$  and HNF1 $\beta$  (Kikuchi et al., 2006). DNA methylation is one of the most well characterized mechanisms underlying the epigenetic regulation of

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gene expression (Bird, 2002; Shiota, 2004). Methylation of the cytosine residue in the CpG dinucleotide negatively regulates gene transcription through the recruitment of chromatin remodeling factors, and many reports highlight the key role of DNA methylation in tissue-specific gene transcription. Since several CpG dinucleotides were found in the 5'-flanking region of *hURAT1* and *mUrat1* (Fig. 1), it would be of great interest to determine whether the concerted effect of DNA methylation and HNF1 $\alpha/\beta$  regulates tissue-specific expression of these genes.

The present study was aimed to clarify the involvement of HNF1 $\alpha$  and HNF1 $\beta$  as genetic factors and DNA methylation as an epigenetic factor in the tissue-specific expression of human and mouse *URAT1* genes.

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## Materials and Methods

### Isolation of the 5'-flanking sequence of *hURAT1* and *mUrat1* gene

The minimal promoter region of *hURAT1* gene (-253/+83) (Li et al., 2004) was amplified by PCR using human genomic DNA as a template and the forward (h-253) and reverse (h+83) primers with an artificial KpnI and HindIII site, respectively (Table 1). The PCR product was subcloned into the KpnI and HindIII restriction sites of pGL3-Basic (Promega, Madison, WI), yielding the *hURAT1* minimal promoter construct, *hURAT1*<sub>-253/+83-HNF1wt</sub>. The transcriptional start site of the *mUrat1* gene was determined using the public database Database of Transcriptional Start Sites (<http://dbtss.hgc.jp/>) with the ref sequence identification for mUrat1 (NM\_009203), and based on high homology with the 5'-flanking region of *hURAT1* (Fig. 1). Since the position of transcriptional start site in the clone MS2926204 is nearest to that of *hURAT1* transcriptional start site which was experimentally determined in the previous report, we chose this site as the transcriptional start site of *mUrat1* gene. The *mUrat1* putative promoter region was isolated from mouse genomic DNA by a PCR-based approach using the forward (m-261) and reverse (m+80) primers (Table 1). The PCR product was subcloned into pGL3-Basic as described above, yielding *mUrat1*<sub>-260/+80</sub>. The sequences of all constructs were verified by DNA sequencing.

### Site-directed Mutagenesis

Mutations in the HNF1-motif located within the *hURAT1* minimal promoter were introduced using the QuickChange XL Site-Directed Mutagenesis Kit (Stratagene, La Jolla, CA) with internal mutated oligonucleotides (Table 1). The introduction of mutations was verified by DNA sequencing.

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### **RNA isolation and quantitative PCR**

Total RNA was isolated from the kidneys of 7- to 14-week-old male ( $n = 3$ ) and female ( $n = 4$ ) wild-type or *Hnf1 $\alpha$* -null adult mice (Lee et al., 1998) and treated with DNase I to remove the contaminated genomic DNA, followed by reverse-transcription using a random-nonamer primer (Takara, Shiga, Japan). To quantify the mRNA expression of mUrat1 in wild-type or *Hnf1 $\alpha$* -null mice, real-time quantitative PCR was performed as described previously using the primers shown in Table 1 (Kikuchi et al., 2006). The mRNA expression of mUrat1 was normalized by the mRNA expression of Gapdh, and statistically analyzed by Student's t-test. Asterisks (\* and \*\*) represent significant differences ( $P < 0.05$  and  $P < 0.01$ , respectively) between wild-type and *Hnf1 $\alpha$* -null mice of each gender.

### **Cell culture and transfections**

Cell culture, transfections, and luciferase assays were performed as described previously (Kikuchi et al., 2006). For transactivation assays in HEK293 cells, 5 ng empty pcDNA3.1(+) vector, 5 ng HNF1 $\alpha$  expression vector, 2.5 ng HNF1 $\alpha$  and HNF1 $\beta$  expression vectors, or 5 ng HNF1 $\beta$  expression vector was cotransfected with 0.5  $\mu$ g of the corresponding promoter construct and 0.05  $\mu$ g internal standard pRL-SV40. The promoter activity was measured as relative light units of firefly luciferase per unit of Renilla luciferase.

### **Preparation of nuclear extracts and electrophoretic mobility shift assay**

Nuclear extracts were prepared from HepG2, Caco-2, and HEK293 parent cells, or HEK293 cells transiently transfected with pcDNA3.1(+), either HNF1 $\alpha$  or HNF1 $\beta$  expression vector, or both expression vectors as described previously (Kikuchi et al., 2006). Transient transfection of the expression vectors into HEK293 cells was



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performed using FuGENE6 (Roche Diagnostics, Indianapolis, IN) according to the manufacturer's instructions. Three kinds of double-stranded oligonucleotides (wt, per, and mut) were obtained by hybridizing the single-stranded complementary oligonucleotide with sense sequences shown in Table 1. Sequence 'wt' corresponds to the wild-type HNF1-motif in the *hURATI* promoter, and 'per' corresponds to the perfect consensus sequence for the HNF1-motif, while 'mut' denotes the wild type sequence mutated within the motif. Five  $\mu\text{g}$  of nuclear extracts from HepG2, Caco-2, or HEK293 parent cells, or 3  $\mu\text{g}$  of nuclear extracts from HEK293 cells transfected with several expression vectors were used in the electrophoretic mobility shift assays. Competition and supershift assays were performed as described previously with Dig Gel Shift Kit, 2<sup>nd</sup> Generation (Roche Diagnostics).

#### **Sodium bisulfite genomic sequencing**

Genomic DNA from liver, kidney cortex, or kidney medulla of ddY male mice at 8-weeks of age was extracted using a Get pure DNA Kit (Dojindo Molecular Technologies, Gaithersburg, MD). One microgram of genomic DNA digested with EcoRI was denatured by adding NaOH to give a final concentration of 0.3 M and incubating for 15 min at 37 °C. After the incubation, sodium metabisulfite (pH 5.0) and hydroquinone were added to give final concentrations of 2.0 M and 0.5 mM, respectively, and the mixture was incubated for 16 h at 55 °C in the dark. The modified DNA was purified using the Wizard DNA Clean-Up System (Promega), and the bisulfite reaction was terminated by adding NaOH to give a final concentration of 0.3 M and incubating for 15 min at 37 °C. The solution was neutralized by adding NH<sub>4</sub>OAc (pH 7.0), and the DNA was ethanol-precipitated, dried, and resuspended in TE buffer (10 mM Tris-HCl, 1 mM EDTA, pH 8.0). The DNA fragment covering the proximal promoter region of the

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*mUrat1* gene was amplified by PCR using the following primers: -419-B-F and +134-B-R (Table 1). The PCR products were cloned into pGEM-T Easy vector (Promega), and 10 clones randomly picked from each of two independent PCRs were sequenced to determine the presence of methylated cytosines.

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## Results

### Involvement of HNF1 $\alpha$ / $\beta$ in the promoter activity of *URAT1* genes.

A previous report suggested the presence of *cis*-elements required for the basal promoter activity of the *hURAT1* gene in the region from –253 to –39 bp relative to the transcriptional start site (Li et al., 2004). The minimal promoter construct of *hURAT1* was transfected into three human-derived cell lines, and luciferase activities were measured. The *hURAT1* promoter showed a significant increase in luciferase activity compared with the promoterless pGL3-Basic plasmid in HepG2 and Caco-2 cells, while the promoter activity was negligible in HEK293 cells (Fig. 2A).

In order to investigate the involvement of HNF1 $\alpha$  or HNF1 $\beta$  in the *hURAT1* minimal promoter, the HNF1-motif found in the *hURAT1* promoter was disrupted by site-directed mutagenesis and the promoter activity was measured in HepG2 and Caco-2 cells where endogenous HNF1 $\alpha$  and/or HNF1 $\beta$  are expressed (Kikuchi et al., 2006). Mutation in the HNF1-motif completely abolished the promoter activity in both cell lines (Fig. 2B). Direct confirmation of the importance of HNF1 $\alpha$  or HNF1 $\beta$  for the promoter activity of *URAT1* genes was obtained by cotransfection assays in HEK293 cells where neither HNF1 $\alpha$  nor HNF1 $\beta$  is endogenously expressed (Fig. 2C). Exogenous expression of HNF1 $\alpha$  and/or HNF1 $\beta$  markedly increased luciferase activity of the *hURAT1* wild-type reporter (–253/+83-HNF1wt) compared with the pcDNA3.1(+)-transfected control, whereas the luciferase activity of pGL3-Basic was unaffected. The luciferase activity driven by *hURAT1* HNF1-mutated reporter (–253/+83-HNF1mut) was not transactivated by exogenously-expressed HNF1 $\alpha$  and/or HNF1 $\beta$  to the same degree as that of the wild-type construct as expected. The activity of *mUrat1* promoter (mUrat1<sub>–261/+80</sub>) was also stimulated by forced-expression of

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HNF1 $\alpha$  and/or HNF1 $\beta$ . These results strongly suggest that the minimal promoter activity of human and mouse *URAT1* genes predominantly depends upon the function of HNF1 $\alpha$  or HNF1 $\beta$ .

### **Interaction of HNF1 $\alpha/\beta$ with the *hURAT1* promoter**

Interaction of digoxigenin-labeled oligonucleotide probes (wt, per, and mut) with nuclear proteins of HepG2, Caco-2, or HEK293 cells was assessed by electrophoretic mobility shift assays. The wt probe corresponds to the wild-type HNF1-motif in the *hURAT1* promoter, and per probe corresponds to the consensus sequence for the HNF1-motif, while the mut probe has the same mutation in the HNF1-motif as that used in the luciferase assays. A non-specific band, which was formed with every probe and abolished by 25-fold molar excess of both per and mut competitor, was observed in all cell lines. In the competition assays (Fig. 3A), one (band *a*) or two shifted bands (bands *a* and *b*) were formed with nuclear extracts derived from HepG2 or Caco-2 cells, respectively, when the wt and per probe were used (lanes 1, 4, 6, and 9). The bands *a* and *b* were abolished by an excess of unlabeled per but not by mut oligonucleotide (lanes 2, 3, 7, and 8), and these bands were not formed when the mut probe was used (lanes 5 and 10). These results suggest that bands *a* and *b* represent the binding of HNF1 $\alpha$  or HNF1 $\beta$  to the HNF1-motif in the *hURAT1* promoter. In contrast, there were no specific bands in HEK293 cells (lanes 11 and 12), consistent with the lack of HNF1 $\alpha$  and HNF1 $\beta$  in this cell line.

Supershift analysis with antibodies against HNF1 $\alpha$  or HNF1 $\beta$  revealed specific interaction of HNF1 $\alpha$  and/or HNF1 $\beta$  with the HNF1-motif in the *hURAT1* promoter (Fig. 3B). Addition of an anti-HNF1 $\alpha$  antibody supershifted band *a* in HepG2 cells (lanes 3 and 4) and bands *a* and *b* in Caco-2 cells (lanes 8 and 9). Addition of an anti-HNF1 $\beta$

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antibody resulted in the supershift of band *b* in Caco-2 cells (lanes 10 and 11), whereas band *a* was not supershifted (lanes 5, 6, 10, and 11). These results suggest that the bands *a* and *b* reflect the binding of HNF1 $\alpha$ /HNF1 $\alpha$  homodimer and HNF1 $\alpha$ /HNF1 $\beta$  heterodimer, respectively. Binding of exogenously-expressed HNF1 $\alpha$  or HNF1 $\beta$  to the HNF1-motif was also demonstrated. Another shifted band showing faster mobility than HNF1 $\alpha$ /HNF1 $\beta$  heterodimer was detected when the nuclear extracts of HEK293 cells transfected with HNF1 $\beta$  alone were incubated with the labeled probe, and this band was supershifted by anti-HNF1 $\beta$  antibody but not by anti-HNF1 $\alpha$  antibody (data not shown). This result suggests that HNF1 $\beta$ /HNF1 $\beta$  homodimer can also interact with the HNF1-motif in the *hURAT1* promoter.

#### **Impaired expression of mUrat1 in *Hnf1 $\alpha$* -null mice.**

Real-time PCR analysis revealed that the expression of mUrat1 mRNA in the kidney is much lower in *Hnf1 $\alpha$* -null mice compared with wild-type mice of both genders (Fig. 4). In contrast to a previous report (Hosoyamada et al., 2004), there was no significant difference in the expression level of mUrat1 between male and female wild-type mice. On the other hand, the expression of mUrat1 mRNA in male was significantly higher than that in female in *Hnf1 $\alpha$* -null mice. The difference in *Hnf1 $\alpha$* -dependent expression (wild-type minus *Hnf1 $\alpha$* -null mice) between male and female did not reach statistical significance.

#### **Epigenetic regulation of the *mUrat1* gene.**

Eight CpG dinucleotides, primary targets of DNA methylation in the vertebrate genome, are located in the 5'-flanking sequence up to -500 bp of *mUrat1* gene; seven of them are within the minimal promoter region (Fig. 1). In order to elucidate the role of DNA methylation in the tissue-specific expression of the *mUrat1* gene, the methylation

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status of each CpG site was analyzed by sodium bisulfite genomic sequencing in the liver, kidney cortex, and kidney medulla, and the total methylation profiles in these tissues were compared (Fig. 5). These CpGs were heavily methylated in the liver and kidney medulla, where there is no expression of mUrat1. In contrast, the region was relatively hypomethylated in the kidney cortex, where mUrat1 is predominantly expressed. These results suggest that tissue-specific expression of *mUrat1* gene is regulated through DNA methylation-mediated gene silencing, and the proximal promoter region of the *mUrat1* gene can be regarded as a tissue-dependent differentially methylated region (T-DMR).

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## Discussion

In the present study, the involvement of both genetic and epigenetic mechanisms in the transcriptional regulation of human and mouse *URAT1* genes was demonstrated. A pivotal role for HNF1 $\alpha$  and HNF1 $\beta$  in the minimal promoter activity of *URAT1* genes was confirmed by reporter gene assays (Fig. 2), and interaction of HNF1 $\alpha$ /HNF1 $\alpha$  homodimer, HNF1 $\alpha$ /HNF1 $\beta$  heterodimer, and HNF1 $\beta$ /HNF1 $\beta$  homodimer with the HNF1-motif in the *hURAT1* promoter was shown by electrophoretic mobility shift assays (Fig. 3).

In the kidney, HNF1 $\alpha$  normally exists as a heterodimer with HNF1 $\beta$ , although the HNF1 $\alpha$ /HNF1 $\alpha$  homodimer is a more potent transactivator than the heterodimer in vitro. HNF1 $\beta$ /HNF1 $\beta$  homodimer is also detectable in nuclear extracts from kidney, but the transactivation potency is the lowest amongst the three species (Pontoglio et al., 1996; Rey-Campos et al., 1991). Because expression of HNF1 $\alpha$  is confined to the proximal tubules, while that of HNF1 $\beta$  is observed along the entire nephron (Lazzaro et al., 1992; Pontoglio et al., 1996), formation of HNF1 $\alpha$ /HNF1 $\beta$  heterodimer will be restricted to the proximal tubules in the kidney and that of HNF1 $\beta$ /HNF1 $\beta$  homodimer is possible in the entire nephron. The expression of human and mouse *URAT1* genes is predominantly in the proximal tubules (Enomoto et al., 2002; Hosoyamada et al., 2004), which is consistent with the distribution of the HNF1 $\alpha$ /HNF1 $\beta$  heterodimer. This suggests that the HNF1 $\alpha$ /HNF1 $\beta$  heterodimer is required for constitutive expression of human and mouse *URAT1*. In contrast, the contribution of HNF1 $\beta$ /HNF1 $\beta$  homodimer is debatable. Although the HNF1 $\beta$ /HNF1 $\beta$  homodimer can also transactivate the *mUrat1* promoter, the expression of *mUrat1* was markedly reduced in the kidney of *Hnf1 $\alpha$* -null mice (Fig. 4). This supports an essential role for the HNF1 $\alpha$ /HNF1 $\beta$  heterodimer, which could not

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be compensated for by the HNF1 $\beta$ /HNF1 $\beta$  homodimer.

Eight CpG dinucleotides in the 5'-flanking region of the *mUrat1* gene appeared hypermethylated in the liver and kidney medulla, whereas they were relatively hypomethylated in the kidney cortex (Fig. 5). Two mechanisms are proposed for DNA methylation-dependent gene silencing: 1) DNA methylation directly interrupts the binding of transcription factors to their recognition sequences including the CpG dinucleotide, and 2) methyl CpG binding proteins bound to the methylated cytosine recruit chromatin remodeling factors, such as histone deacetylases, and cause the neighboring chromatin configuration to condense, thereby preventing many transcription factors from accessing their recognition sequences within that region (Shiota, 2004). The HNF1-motif in the *mUrat1* promoter is located within the T-DMR, but the motif itself does not contain the CpG dinucleotide. It is possible that methylation of DNA in the T-DMR indirectly inhibits the binding of HNF1 $\alpha$  or HNF1 $\beta$  to the promoter through chromatin remodeling events. Hypermethylation of the T-DMR may explain the absence of *mUrat1* expression regardless of the formation of HNF1 $\alpha$ /HNF1 $\alpha$  homodimers in the liver. In the kidney, the promoter region is hypomethylated in the cortex but hypermethylated in the medulla. Together with the proximal tubule-restricted distribution of the HNF1 $\alpha$ /HNF1 $\beta$  heterodimer, the regional difference in the methylation status seems to lead to the constitutive expression of *mUrat1* in the proximal tubules in the cortex. In humans, the methylation profiles of *hURAT1* promoter remain unknown. Considering the similar frequency of CpG dinucleotides in the minimal promoter region between human and mouse (Fig. 1), DNA methylation will be also involved in the tissue/region-specific expression of *hURAT1*.

Mice lacking HNF1 $\alpha$  suffer from severe renal Fanconi syndrome, a dysfunction in



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renal proximal tubular reabsorption (Pontoglio et al., 1996). The renal fractional excretion of urate is higher in mutant animals compared with wild-type controls, resulting in a lower serum urate levels. In the present study, in vitro and in vivo evidence of the importance of HNF1 $\alpha$  in the expression of *URAT1* genes was provided; accurate tissue-specific expression of this gene is essential for the reabsorption of filtered urate in the kidney. The altered renal excretion of urate in mutant mice is thus accounted for by impaired expression of *Urat1*. It was reported that expression of transporters responsible for reabsorption of filtered glucose, phosphate, and bile acids is also absent or lower in *Hnfl*  $\alpha$ -null mice (Cheret et al., 2002; Pontoglio et al., 2000; Shih et al., 2001). Taken together, the expression of transporters required for the reabsorption of filtered solutes in the kidney proximal tubules is largely impaired in mutant mice, leading to renal Fanconi syndrome.

In addition to renal Fanconi syndrome, *Hnfl*  $\alpha$ -null mice exhibit a phenotype similar to non-insulin-dependent diabetes mellitus (Lee et al., 1998; Pontoglio et al., 1998). In humans, mutations in HNF1 $\alpha$  and HNF1 $\beta$  cause maturity onset diabetes of the young (MODY) types 3 and 5, respectively (Horikawa et al., 1997; Yamagata et al., 1996). MODY is an autosomal dominant inherited disease that is responsible for 2-5% of non-insulin-dependent diabetes mellitus (Ledermann, 1995), and MODY3 accounts for 20-75% of MODY patients, while MODY5 is a rare condition (Froguel and Velho, 1999; Ryffel, 2001; Winter and Silverstein, 2000). It was reported that the serum urate levels are low in patients with both type I and type II diabetes mellitus because of the elevated renal clearance (Golik et al., 1993; Magoula et al., 1991; Shichiri et al., 1987). The increase in the urate clearance has been ascribed to an elevated glomerular filtration rate and/or a defect in tubular urate reabsorption. It is thus possible that the expression of

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URAT1 mRNA is reduced in MODY3 and MODY5 patients, resulting in hypouricemia.

In conclusion, clear evidence was provided for the involvement of both genetic (HNF1 $\alpha$  and HNF1 $\beta$ ) and epigenetic (DNA methylation) mechanisms in establishing the tissue-specific expression of mouse, and probably, human *URAT1* genes. This is the first demonstration of the presence of T-DMR in the promoter region as far as transporters in the kidney are concerned.

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## Legends for figures

Fig. 1

### **Alignment of the proximal promoter of human and mouse *URAT1*.**

Nucleotide sequences of the proximal promoter region of human and mouse *URAT1* genes were aligned using Genetyx ver. 8 to show the high homology of the 5'-flanking region between species. Nucleotide numberings are relative to the transcriptional start sites indicated by an arrow (+1), and homologous sequences between species are boxed. Canonical HNF1 binding motif in the promoter is shaded and CpG dinucleotides in each sequence are reverse-colored.

Fig. 2

### **HNF1 $\alpha$ and HNF1 $\beta$ predominantly regulate *URAT1* promoters.**

A, Analysis of *hURAT1* minimal promoter. HepG2 (white bars), Caco-2 (gray bars), and HEK293 cells (black bars) were transiently transfected with promoterless pGL3-Basic plasmid or minimal promoter construct of *hURAT1* (hURAT1<sub>-253/+83</sub>-HNF1wt) together with the internal standard pRL-SV40 to normalize the transfection efficiency. B, Mutational analysis of the HNF1-motif. HepG2 (white bars) and Caco-2 cells (gray bars) were transfected with pGL3-Basic, wild-type (hURAT1<sub>-253/+83</sub>-HNF1wt), or HNF1-mutated (hURAT1<sub>-253/+83</sub>-HNF1mut) promoter construct. C, Exogenous expression of HNF1 $\alpha$  and HNF1 $\beta$ . HEK293 cells were transfected with pGL3-Basic, wild-type or the HNF1-mutated promoter construct of *hURAT1* (hURAT1<sub>-253/+83</sub>-HNF1wt or hURAT1<sub>-253/+83</sub>-HNF1mut, respectively), or the *mUrat1* promoter construct (mUrat1<sub>-261/+80</sub>), together with empty pcDNA3.1(+) vector (white bars), HNF1 $\alpha$  expression vector (black bars), HNF1 $\alpha$  and HNF1 $\beta$

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expression vectors (gray bars), or HNF1 $\beta$  expression vector (light gray bars). The promoter activity was measured as described under Materials and Methods, and was shown as the induction factor over the background activity measured in cells transfected with pGL3-Basic in each cell line (A and B), or cells transfected with pGL3-Basic together with pcDNA3.1(+) (C). All results are presented as the mean  $\pm$  S.E. of triplicate samples.

Fig. 3

**HNF1 $\alpha$  and HNF1 $\beta$  interact with the *hURAT1* promoter.**

A, Competition assays. Three kinds of digoxigenin-labeled probe (Table 1) were incubated with nuclear extracts of HepG2, Caco-2, or HEK293 cells, with or without a 25-fold excess of unlabeled competitor (per or mut) as indicated. B, Supershift assays. The wt probe was incubated with nuclear extracts of HepG2 or Caco-2 cells with or without an increasing amount of specific antibody against HNF1 $\alpha$  ( $\alpha$ ) or HNF1 $\beta$  ( $\beta$ ) as indicated. The DNA-protein complex was detected as described under Materials and Methods.

Fig. 4

**Relative mRNA expression of mUrat1 in wild-type and *Hnf1 $\alpha$* -null mice.**

mRNA expression of mUrat1 in the kidneys of male or female wild-type (WT, white bars) and *Hnf1 $\alpha$* -null mice (null, black bars) was measured by real-time quantitative PCR using specific primers (Table 1), and the data were normalized by the mRNA expression of Gapdh. The relative mRNA expression was given as a ratio with respect to the mRNA expression of mUrat1 in male wild-type mice that was taken as 100%. Results are



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presented as the mean  $\pm$  S.E. of three (male) or four (female) mice. \* $P < 0.05$  and \*\* $P < 0.01$ , significantly different between wild-type and *Hnf1 $\alpha$* -null mice.

Fig. 5

**DNA methylation profile of the *mUrat1* promoter.**

Top, a schematic diagram of the *mUrat1* 5'-flanking region. The vertical lines and numbers indicate the positions of cytosine residues of CpGs relative to the transcriptional start site (TSS, +1). The position of the HNF1-motif is shown by a rectangle. Bottom, DNA methylation status of individual CpGs. Bisulfite sequencing analysis was performed with genomic DNAs extracted from mouse liver, kidney cortex, and kidney medulla. Open and filled circles represent unmethylated and methylated cytosines, respectively.

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## Tables

**Table 1: Oligonucleotides used for the production of promoter fragments, mobility shift assays, site-directed mutagenesis, quantitative PCR, and bisulfite PCR**

Regarding the oligonucleotides used for the mobility shift assays and site-directed mutagenesis, the HNF1-motif in the *hURAT1* promoter region is underlined. Bold type indicates the difference in the sequence of the per and mut compared with the wild-type sequence found in the *hURAT1* promoter.

Oligonucleotide	Orientation	Sequence (5' to 3')
<b>Primers used for the cloning of 5'-flanking regions</b>		
<i>hURAT1</i>		
h-253	Forward	CGGGGTACCTTGCTCAGCCACTCTGGGAGGT
h+83	Reverse	CCCAAGCTTAGAGAGGCAGCTGCTCCAGACC
<i>mUrat1</i>		
m-261	Forward	CGGGGTACCCTGGGTTAGCTCACAGTACAG
m+80	Reverse	CCCAAGCTTAGCGACACAGCAGAGTCTG
<b>Oligonucleotides used for the construction of EMSA probe and competitor or site-directed mutagenesis</b>		
wt	Sense	CTCAACGCGGGTAAACTTTGACCAAGGAAATG
per	Sense	CTCAACGCGGGTAA <b>TCATTA</b> ACCAAGGAAATG
mut	Sense	CTCAACGCGGG <b>GGCG</b> AACTGTGACCAAGGAAATG
<b>Primers used for quantitative PCR</b>		
mUrat1	Forward	GAGGGAGACACGTTGACCAT
	Reverse	AAGTCCACAATCCCGATGAG
mGapdh	Forward	AACGACCCCTTCATTGAC
	Reverse	TCCACGACATACTCAGCAC
<b>Primers used for bisulfite PCR</b>		
<i>mUrat1</i>		
-419-B-F	Forward	GGAATTAATAAGGGAGTGTAGGAA
+134-B-R	Reverse	TCACCATAAAACCTAAAACCCTCT

Fig. 1

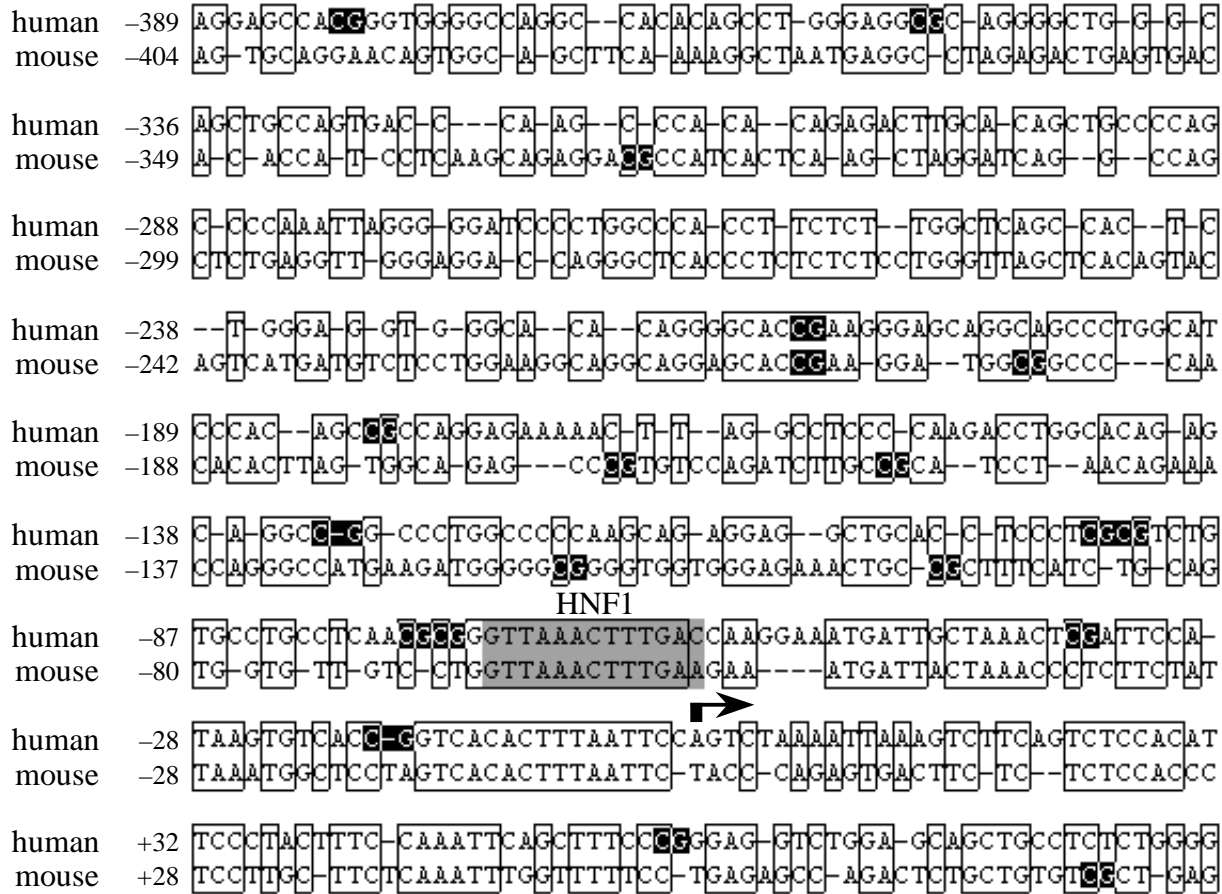
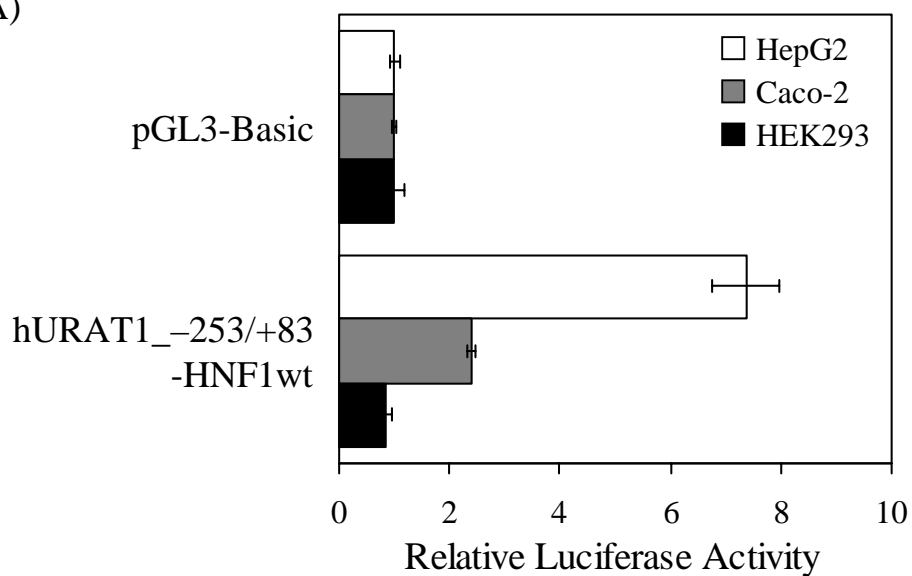
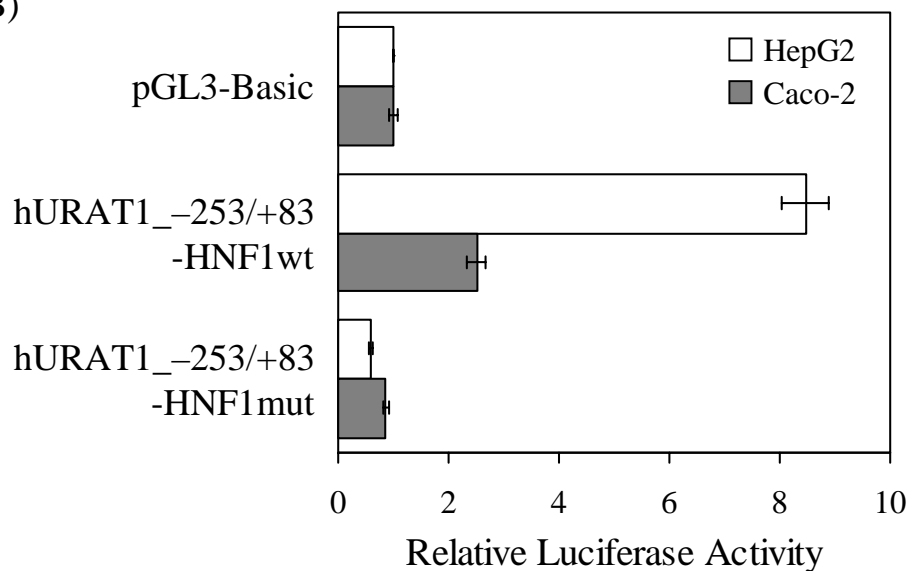


Fig. 2 (A)



(B)



(C)

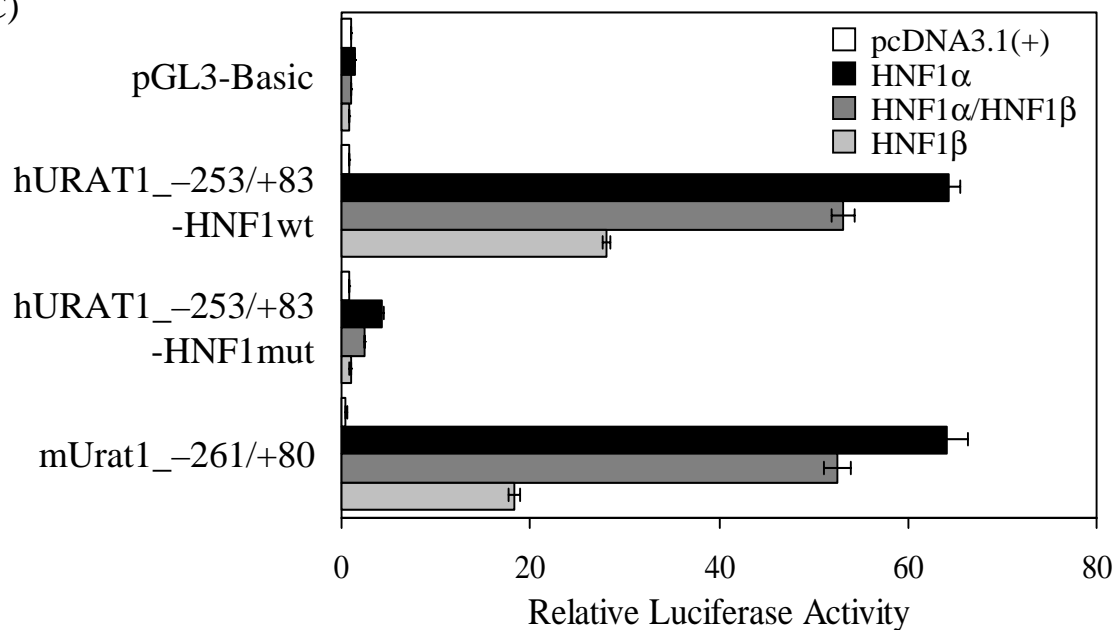


Fig. 3

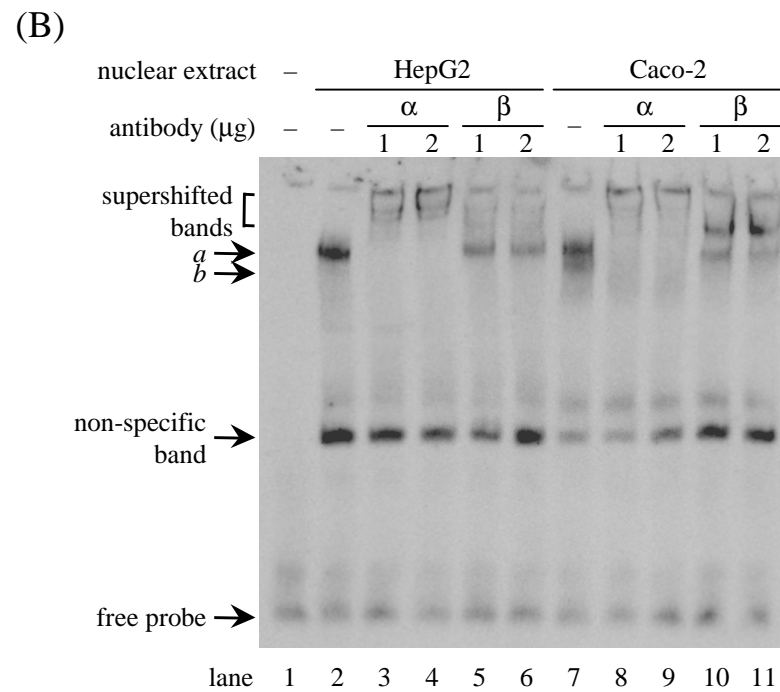
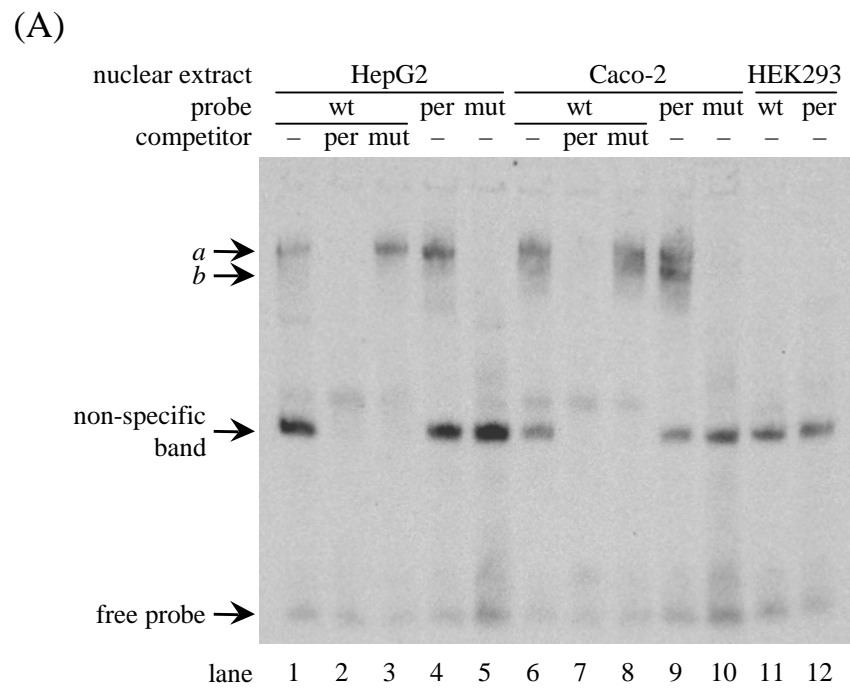


Fig. 4

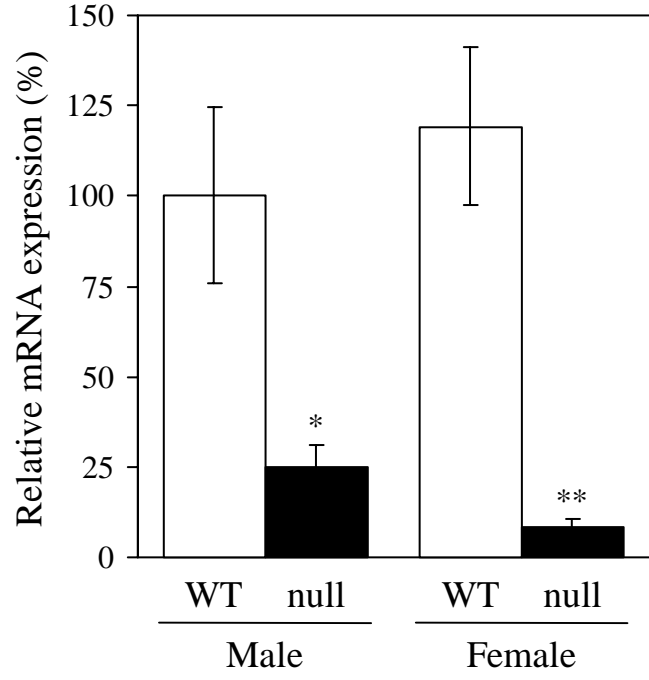


Fig. 5

