Tubular Deficiency of Heterogeneous Nuclear Ribonucleoprotein F Elevates Systolic Blood Pressure and Induces Glycosuria in Mice

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We reported previously that overexpression of heterogeneous nuclear ribonucleoprotein F (Hnrnpf) in renal proximal tubular cells (RPTCs) suppresses angiotensinogen (Agt) expression, and attenuates systemic hypertension and renal injury in diabetic Hnrnpf-transgenic (Tg) mice. We thus hypothesized that deletion of Hnrnpf in the renal proximal tubules (RPT) of mice would worsen systemic hypertension and kidney injury, perhaps revealing novel mechanism(s). Tubule-specific Hnrnpf knockout (KO) mice were generated by crossbreeding Pax8-Cre mice with floxed Hnrnpf mice on a C57BL/6 background. Both male and female KO mice exhibited elevated systolic blood pressure, increased urinary albumin/creatinine ratio, tubulo-interstitial fibrosis and glycosuria without changes in blood glucose or glomerular filtration rate compared with control littermates. However, glycosuria disappeared in male KO mice at the age of 12 weeks, while female KO mice had persistent glycosuria. Agt expression was elevated, whereas sodium-glucose co-transporter 2 (Sglt2) expression was down-regulated in RPTs of both male and female KO mice as compared to control littermates. In vitro, KO of HNRNPF in human RPTCs (HK-2) by CRISPR gRNA up-regulated AGT and down-regulated SGLT2 expression. The Sglt2 inhibitor canagliflozin treatment had no effect on Agt and Sglt2 expression in HK-2 and in RPTCs of wild-type mice but induced glycosuria. Our results demonstrate that Hnrnpf plays a role in the development of hypertension and glycosuria through modulation of renal Agt and Sglt2 expression in mice, respectively.

The kidney contains all components of the renin-angiotensin system (RAS)1–4. Over-activation of the intrarenal RAS appears to be involved in various kidney diseases5–7. We and others have reported that overexpression of angiotensinogen (Agt, the sole precursor of all angiotensins) in RPTCs leads to systemic hypertension and kidney injury in transgenic (Tg) mice8–10, supporting the notion that enhanced intrarenal Agt expression and RAS activation play an important role in the development of hypertension and kidney injury.

Our lab has reported that heterogeneous nuclear ribonucleoprotein F (Hnrnpf) mediates insulin inhibition of Agt gene transcription through binding to the putative insulin-responsive element (IRE) in the rat Agt promoter11,12. We recently reported that overexpression of Hnrnpf in RPTCs suppresses Agt expression, and attenuates systemic hypertension and renal injury in male Akita (type 1 diabetic murine model) Hnrnpf-Tg mice13 and db/db (type 2 diabetic murine model) Hnrnpf-Tg mice14. Since sex differences may modulate the development of systolic blood pressure (SBP)15,16, we investigated whether Hnrnpf would affect intrarenal Agt expression in a...
sex-dependent manner. We generated tubule-specific Hnrnpf KO mice by employing the Pax8-Cre/lox system and monitored the development of phenotype in both male and female mice.

Here, we report that tubule-specific (Pax8) Hnrnpf KO leads to elevated SBP and kidney injury via up-regulation of Agt and down-regulation of Sglt2 expression in RPTCs in both sexes and also results in glycosuria in a sex-dependent manner. KO of HNRNPF by CRISPR gRNA confirmed the up-regulation and down-regulation of AGT and SGLT2 expression in human RPTCs (HK-2), respectively. Treatment with canagliflozin (an inhibitor of Sglt2) had no effect on Agt and Sglt2 expression in HK-2 and in RPTCs of wild-type mice, whereas it induced glycosuria.

Results

Generation of tubular Hnrnpf KO Mice. Renal tubular Hnrnpf KO mice were generated by using Pax8-Cre/lox recombination strategy (Fig. 1A). LoxP sites were inserted to flank exon 4 of mouse Hnrnpf gene (Gene ID: 98758) which is localized on chromosome 6. Heterozygous of Hnrnpf-floxed allele mice were generated by cross-breeding male Hnrnpf-floxed mice with female Pax8-Cre mice. These mice were further crossbred to generate homozygous Hnrnpf-floxed allele and carried the Cre allele. PCR analysis of genomic DNA extracted from ear punch tissues to distinguish the genotype of Cre (392 bp), floxed (568 bp) and WT (507 bp) is shown in Fig. 1B. RT-qPCR revealed Hnrnpf mRNA expression in RPTs freshly isolated from male and female Ctrl and KO mice at the age of 8 weeks (Supplemental Fig. 1a) and 24 weeks (Fig. 1C). Hnrnpf mRNA was barely detectable in RPTs of both male and female KO mice at 8 and 24 weeks of age.

WB of isolated RPTs confirmed the expression of Hnrnpf at the age of 8 (Supplemental Fig. 1b) and 24 weeks (Fig. 1D) in Ctrl whereas Hnrnpf expression was significantly down-regulated in KO mice. No significant difference of Hnrnpf expression was observed between male and female Ctrl as well as between male and female KO mice. Double immunofluorescence of kidney sections (Fig. 1E) with an anti-Hnrnpf antibody and LTL-FITC antibody, confirmed significantly higher Hnrnpf expression in RPTs from Ctrl than in KO mice.

Physiological measurements in Hnrnpf KO mice. Deletion of renal tubular Hnrnpf did not influence body weight gain nor the non-fasting blood glucose levels in both male and female mice from the age of 6 to 24 weeks (Supplemental Fig. 1c–f, respectively). Longitudinal SBP measurements revealed consistently higher SBP in both male (Fig. 2A) and female (Fig. 2B) KO mice aged week 6 to 24 compared to Ctrl. Significant increases of Agt mRNA and protein expression were detected in both male and female KO mice compared to Ctrl at 8 weeks (Supplemental Fig. 2a) and 24 weeks of age (Fig. 2C,D, respectively). No significant difference of Agt expression in RPTs was observed between male and female Ctrl as well as between male and female KO mice. These were confirmed with immunostaining (Fig. 2E).

Increased urinary Ang II and urinary albumin/creatinine ratio (ACR) were also observed in both male and female KO mice compared to Ctrl at 24 weeks of age with no significant difference between male and female Ctrl as well as between male and female KO mice. (Fig. 2F,G, respectively). In contrast, body weight (BW), kidney weight (KW)/BW, GFR and glomerular tuft volume did not differ significantly between KO mice and Ctrl at 24 weeks of age (Table 1). Twenty-four h urine volume were significantly increased but not food and water intake in both male and female KO mice as compared to Ctrl. No differences were detected in serum and urine levels of sodium, calcium and phosphorus between male and female Ctrl and KO mice. We detected no significant differences in serum Ang II among different groups of mice (Fig. 2H).

Tubulo-interstitial fibrosis in Hnrnpf KO mice. PAS staining of kidney sections showed no obvious structural changes in KO mice at the age of 24 weeks (Fig. 3A). Increased fibrosis on Masson’s Trichrome staining (Fig. 3B) and increased expression of collagen on Sirius Red staining (Fig. 3C), and fibronectin 1 (Fn1) immunostaining (Fig. 3D) was, however, noted in glomerulo-tubular regions in KO mice as compared to Ctrl at the age of 24 weeks. Semi-quantification of tubular luminal area (Fig. 3E), RPTC volume (Fig. 3F), Masson’s Trichrome staining (Fig. 3G), Sirius Red staining (Fig. 3H) and Fn1 immunostaining (Fig. 3I) revealed an increase of tubular luminal area, RPTC volume, Masson’s Trichrome and Sirius Red staining and Fn1 immunostaining in KO mice as compared to Ctrl, respectively. These findings were associated with significant increases of mRNA expression of Fn1 (Fig. 3J) by RT-qPCR in isolated RPTs of KO mice as compared with Ctrl.

Glycosuria and Sglt2 expression in Hnrnpf KO mice. Unexpectedly, increased glucose excretion was detected in the urine using dipsticks in both male and female KO mice at age 6 weeks (Fig. 4A). From 8 weeks of age, urinary glucose levels in male KO mice steadily decreased and returned to levels similar to Ctrl mice at 12 weeks of age (Fig. 4B). In contrast, urinary glucose excretion steadily increased from week 6 in female KO mice, reached an apparent plateau at the age of 12 weeks and did not abate (Fig. 4B). No changes in urinary glucose level were detected in male and female Ctrl.

Havina observed that both male and female Hnrnpf KO mice develop glycosuria, we performed intraperitoneal glucose tolerance test (IPGTT) at the age of 23 weeks in male and female mice (Fig. 4C). KO of Hnrnpf in RPTs did not influence the glucose tolerance in either male or female KO mice.

RT-qPCR revealed lower Sglt2 expression in RPTs isolated from both male and female KO mice at 8 weeks of age (Supplemental Fig. 2b) and 24 weeks of age (Fig. 4D) as compared with Ctrl. At both 8 and 24 weeks of age, Sglt2 expression decreased by ~40% in RPTs of male Hnrnpf KO mice as compared to Ctrl, whereas persistently lower Sglt2 expression (decreased by ~60% of baseline level) was observed in RPTs from female KO mice. However, no significant difference of Sglt2 expression in RPTs was observed between male and female Ctrl as well as between male and female KO mice. WB of isolated RPTs confirmed these changes at the protein level (Fig. 4E).

Consistently, semi-quantitation of immunofluorescence staining with anti-Sglt2 antibodies and LTL-FITC confirmed reduced Sglt2 expression in RPTs of 24 week-old KO mice as compared to Ctrl (Fig. 4F,G, respectively).

LTL-FITC antibody, confirmed significantly higher Hnrnpf expression in RPTs from Ctrl than in KO mice. Female KO mice. Double immunofluorescence of kidney sections (Fig. 1E) with an anti-Hnrnpf antibody and difference of Hnrnpf expression in RPTs was observed between male and female Ctrl as well as between male and female KO mice. RT-qPCR revealed Hnrnpf mRNA expression in RPTs freshly isolated from male and female Ctrl and KO mice at the age of 8 weeks (Supplemental Fig. 1a) and 24 weeks (Fig. 1C). Hnrnpf mRNA was barely detectable in RPTs of both male and female KO mice at 8 and 24 weeks of age. WB of isolated RPTs confirmed the expression of Hnrnpf at the age of 8 (Supplemental Fig. 1b) and 24 weeks (Fig. 1D) in Ctrl whereas Hnrnpf expression was significantly down-regulated in KO mice. No significant difference of Hnrnpf expression was observed between male and female Ctrl as well as between male and female KO mice. Double immunofluorescence of kidney sections (Fig. 1E) with an anti-Hnrnpf antibody and LTL-FITC antibody, confirmed significantly higher Hnrnpf expression in RPTs from Ctrl than in KO mice.
No significant changes were detectable in Scl5a1 (Sglt1) mRNA expression in RPTs isolated from both male and female KO mice as compared to Ctrl (Supplemental Fig. 2c).

Effect of canagliflozin treatment on Agt and Sglt2 expression in RPTCs in vivo. To investigate the role of Sglt2 on Agt and Sglt2 expression in RPTCs in vivo, wild-type mice were treated with the selective Sglt2 inhibitor canagliflozin (0.2 mg/ml in drinking water). Four weeks of canagliflozin treatment had no detectable...
Figure 2. Systolic blood pressure (SBP) and intrarenal angiotensinogen (Agt) expression in tubular Hnrnpf KO mice. (A) Longitudinal average SBP measurement (performed two or three times per mouse per week in the morning without fasting) in (A) male and (B) female mice. Baseline SBP was measured daily over a 5-day period before initiation of actual measurement at week 6. Values are means ± SEM, n = 10 for each group. *P < 0.05, KO versus Ctrl. (C) Agt mRNA levels in male and female Ctrl and KO mice at the age of 24 weeks. *P < 0.05, **P < 0.01, n = 6 per group, KO versus Ctrl. (D) Representative WB of Agt protein expression and quantitation of Agt expression in Ctrl and KO groups from 24-week-old male and female mice. *P < 0.05, **P < 0.01, n = 6 per group, KO versus Ctrl. (E) Representative immunostaining for Agt in Ctrl and KO mice (original magnification ×200). Scale bars = 50 μm. G, Glomerulus; P, proximal tubule. (F) Urinary Ang II, (G) ACR and (H) serum Ang II levels at week 24 in Ctrl and KO mice. Urinary Ang II and albumin levels were normalized with urinary creatinine levels. Values are mean ± SEM, n = 8 per group. *P < 0.05, **P < 0.01 and ***P < 0.005; KO versus Ctrl.
Finally, in human cells expression of HNRNPF (Fig. 6H,I), AGT (Fig. 6H,J) and SGLT2 protein (Fig. 6H,K) and cells, indicating a lack of causality of inhibition of SGLT2 activity and confirmed by semi-quantitation of Agt (Fig. 5F) and Sglt2 (Fig. 5G) expression and qPCR of HNRNPF (Fig. 6E), immunoblots revealed that HK-2 cells with in vivo we generated HK-2 cells with HNRNPF deletion led to glycosuria with reduced expression (Fig. 6G), respectively.

Table 1. Physiological parameters of mice at 24 weeks of age. Values are mean ± SEM; n = 6/group. KW/BW: Kidney Weight/Body Weight; Na: sodium; Ca: calcium; P: phosphorus; Cr: creatinine.

|                  | Male Ctrl | Hnrnpf KO | p  | Female Ctrl | Hnrnpf KO | p  |
|------------------|-----------|-----------|----|-------------|-----------|----|
| Body weight (g)  | 34.35 ± 0.96 | 34.94 ± 0.68 | NS | 24.81 ± 0.89 | 23.55 ± 0.86 | NS |
| KW/BW (mg/g)     | 9.35 ± 0.58 | 8.38 ± 0.41 | NS | 9.97 ± 0.29 | 9.05 ± 0.45 | NS |
| GFR(μL/min)/BW(g) | 7.5 ± 0.45 | 7.7 ± 0.57 | NS | 7.9 ± 0.64 | 6.8 ± 0.34 | NS |
| Glomerular tuft volume (10^3 μm^2) | 117.3 ± 8.92 | 128.2 ± 10.94 | NS | 120.2 ± 9.90 | 129.5 ± 5.88 | NS |
| Urine volume (μL/24h) | 871.7 ± 36.1 | 1198.0 ± 102.7 | * | 265.0 ± 63.3 | 505.8 ± 41.5 | ** |
| Food intake (mg/24h) | 333.3 ± 42.2 | 333.3 ± 33.3 | NS | 566.7 ± 76.0 | 516.7 ± 54.3 | NS |
| Water intake (ml/24h) | 1.97 ± 0.08 | 2.37 ± 0.24 | NS | 2.30 ± 0.29 | 2.55 ± 0.13 | NS |
| Serum Na (mmol/L) | 147.9 ± 2.3 | 151.5 ± 1.4 | NS | 147.6 ± 2.1 * | 148.7 ± 1.4 | NS |
| Serum Ca (mmol/L) | 2.22 ± 0.03 | 2.21 ± 0.04 | NS | 2.34 ± 0.11 * | 2.20 ± 0.03 | NS |
| Serum P (mmol/L)  | 2.89 ± 0.20 | 2.95 ± 0.29 | NS | 3.37 ± 0.36 | 3.13 ± 0.14 | NS |
| Urine Na/ Cr (mmol/g Cr) | 741.0 ± 73.1 | 709.8 ± 25.0 | NS | 398.5 ± 61.1 | 353.2 ± 32.9 | NS |
| Urine P/ Cr (mmol/g Cr) | 107.2 ± 17.2 | 101.2 ± 17.9 | NS | 73.3 ± 10.9 | 79.7 ± 12.6 | NS |

AGT and SGLT2 expression in HK-2 with or without HNRNPF KO. To validate our in vivo observations, we generated HK-2 cells with HNRNPF KO by CRISPR gRNA technology. Consistent with our in vivo observation, immunoblot revealed that HK-2 cells with HNRNPF KO exhibited non-detectable HNRNPF (Fig. 6A,B), higher AGT (Fig. 6A,C) and lower SGLT2 protein expression (Fig. 6A,D) as compared to control HK-2. These findings were confirmed by RT-qPCR of HNRNPF (Fig. 6E), AGT (Fig. 6F) and SGLT2 expression (Fig. 6G), respectively. Finally, in human cells expression of HNRNPF (Fig. 6H,I), AGT (Fig. 6H,J) and SGLT2 protein (Fig. 6H,K) and mRNA (Fig. 6L-N, respectively) did not differ significantly in HK-2 cells treated with canagliflozin and untreated cells, indicating a lack of causality of inhibition of SGLT2 activity and AGT and SGLT2 expression in RPTCs.

Discussion

Our results identify a novel mechanism by which Hnrnpf affects the development of hypertension and glycosuria in mice through modulation of intrarenal Agt and Sglt2 expression, respectively.

Hnrnpf, a member of the 30 pre-mRNA-binding protein family, modulates gene expression at both transcriptional and post-transcriptional levels. Hnrnpf engages in alternative splicing of various genes and associates with TATA-binding protein, RNA polymerase II, nuclear cap-binding protein complex and various transcription factors to modulate gene expression. We have reported previously that Hnrnpf overexpression in RPTCs attenuates hypertension and kidney injury in both diabetic Akita mice and db/db mice via inhibition of intrarenal Agt expression, implying an important role for Hnrnpf in modulating the development of hypertension and nephropathy in diabetic mice.

Our present findings document that genetic deletion of Hnrnpf in tubules enhances renal Agt expression, hypertension development and kidney injury in both non-diabetic male and female mice. These observations are consistent with our hypothesis that Hnrnpf plays an important role in the development of hypertension and tubulo-interstitial fibrosis via modulation of Agt and pro-fibrotic genes in RPTCs.

Initially, we generated global Hnrnpf KO mice by cross-breeding a general Cre-deleter mouse line (CMV-Cre; B6.C-Tg(CMV-cre)1Cgn/J) with our Hnrnpf flo/fl mice on a C57BL/6 background to explore the phenotype of global Hnrnpf KO mice and found that like the global Hnrnpf deletion, global Hnrnpf KO also results in embryonic death (Supplemental Fig. 3). To circumvent this issue, we generated renal tubule-specific Hnrnpf KO mice by cross-breeding our Hnrnpf flo/fl mice with a renal tubule-specific Cre deleter (Pax8-Cre; B6.129P2(Pax8tm1(Cg)Cmu/cfmsf1)) mouse line. Several labs have also successfully employed Pax8-Cre mice to delete genes in renal tubules. Our homozygous Pax8-Hnrnpf KO mice are viable and fertile without symptoms of body weight loss, physiological imbalance and altered hearing. However, they develop hypertension and elevated ACR with increased Agt expression in RPTCs by 8 weeks of age (Supplemental Fig. 4a,b). Since Pax8 is also expressed in the thyroid gland and hindbrain, it is possible that Pax8-Hnrnpf KO mice might exhibit abnormality in thyroid gland and hindbrain thereby indirectly affecting cardiac and renal function. However, we did not detect significant changes in Hnrnpf mRNA levels or serum T4 levels in Pax8-Hnrnpf KO mice (Supplemental Fig. 5) or histological changes in thyroid gland and hindbrain. Thus, our Pax8-Hnrnpf KO mouse appears to be a valid murine model with which to study the phenotype with tubule-specific Hnrnpf KO.

An unexpected finding of our present study was that Hnrnpf deletion led to glycosuria with reduced expression of Sglt2 in RPTs of Hnrnpf KO mice. Intriguingly, serum and urine levels of Na, Ca and P did not differ effects on SBP (Fig. 5A) and blood glucose levels (Fig. 5B) in either male or female mice but enhanced the development of glycosuria in both male and female mice (Fig. 5C) as compared to non-treated mice. Immunostaining for Agt (Fig. 5D) and immunofluorescent staining for Sglt2 (Fig. 5E) revealed that canagliflozin treatment had no effect on Agt expression or Sglt2 expression in RPTCs of either male or female mice. These observations were confirmed by semi-quantitation of Agt (Fig. 5F) and Sglt2 (Fig. 5G) expression and qPCR of Agt and Sglt2 mRNA expression in isolated RPTs (Fig. 5H,I, respectively).

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between Hnrnpf KO mice and Ctrl. The phenotype of glycosuria appears to be similar to that reported in Sglt2 deficient mice\(^27\) and in patients with familial renal glycosuria (FRG)\(^28-30\) but differs from that in Sweet Pee mice, which are characterized by elevated urinary excretion of calcium and magnesium and growth retardation\(^31\), as well as from that in patients with renal Fanconi syndrome\(^32\). Intriguingly, male Hnrnpf KO mice exhibited a transient glycosuria between the ages of 6 and 12 weeks and then returned to levels similar to Ctrl. Furthermore, glycosuria correlates with reduced Sglt2 expression in RPTs of male Hnrnpf KO mice. In contrast, female Hnrnpf KO mice displayed persistent glycosuria throughout 6 to 24 weeks of age with similar inhibition of Sglt2 expression.
observed at both 8 and 24 weeks of age. These data would indicate that male sex hormones rather than female sex hormones may modulate Sglt2 expression in Hnrnpf KO mice. Indeed, this notion had been suggested by Sabolic’s group33,34 who implicated androgen, but not estradiol up-regulates Sglt2 expression and activity in mice.

Figure 4. Glycosuria and Sglt2 expression in Hnrnpf KO mice. (A) Urinary glucose in Ctrl and KO mice detected by dipstick test at the age of 6 weeks. (B) Longitudinal urinary glucose levels in male and female KO mice and Ctrl from the age of 6 weeks to 24 weeks measured by glucose colorimetric kit. Values are means ± SEM, n = 6. **P < 0.01; ***P < 0.005. KO versus Ctrl. #P < 0.01; ##P < 0.005; female KO versus male KO. IPGTT test in male and female (C) Ctrl and KO mice at the age of 23 weeks. (D) Ratio of Sglt2/Rpl13a mRNA expression quantified by RT-qPCR and (E) Representative WB of Sglt2 protein expression in male and female mouse RPTs at the age of 24 weeks. Values are means ± SEM, n = 6. ***P < 0.005. KO versus Ctrl. (F) Double immunostaining of Sglt2 and LTL (magnification x100) and semi-quantification of Sglt2/LTL immunostaining ratio (G) in male and female Ctrl and KO mouse kidneys at the age of 24 weeks. Values are Sglt2/LTL positive staining ratio, n = 6. ***P < 0.005, KO versus Ctrl.
To explore the impact of Sglt2 inhibition on Agt expression, we treated HK-2 cells and WT mice with canagliflozin. Canagliflozin had no detectable effects on SBP and blood glucose levels but enhanced the development of glycosuria in both male and female mice as compared to non-treated mice. Furthermore, canagliflozin treatment had no effect on the expression of Agt and Sglt2 expression in RPTs of mice. Thus, our data would argue against...
a causal relationship between Sglt2 inhibition and Agt expression in RPTCs; rather our data would indicate that inhibition of Hnrnpf expression modulates both Agt and Sglt2 expression in RPTCs.

Finally, to replicate our in vivo observations, we studied a human renal proximal tubular cell line (HK-2)\(^3\). By employing CRISPR gRNA technology, we obtained several clones of HK-2 cells with HNRNPF KO. Consistent with our findings in Hnrnpf KO mice, HK-2 with HNRNPF KO displayed significantly higher AGT and lower SGLT2 expression as compared to HK-2 controls. These data lend further support our previous observations that Hnrnpf down-regulates RPT Agt expression and RAS activation, leading to improve tubulo-interstitial fibrosis in the kidney. Moreover, consistent with our in vivo results, canagliflozin treatment had no effect on SGLT2 and AGT expression in HK-2 cells.

Figure 6. AGT and SGLT2 expression in HK-2 with or without HNRNPF KO. (A) WB, (B–D) semi-quantitation of WB and (E–G) RT-qPCR of HNRNPF, AGT, SGLT2 and β-ACTIN in different clones of HK-2 Ctrl and HK-2 with HNRNPF KO by CRISPR gRNA. Values are means ± SEM, n = 3. *P < 0.05, **P < 0.01; HK-2-HNRNPF KO versus HK-2 Ctrl. (H) WB, (I–K) semi-quantitation of WB and (L)(M)(N) RT-qPCR of HNRNPF, AGT, SGLT2 and β-ACTIN of expression in HK-2 with or without canagliflozin (Cana) (0.5 mM) treatment for 24 hours. Values are means ± SEM, n = 3. NS, not significant. HK-2-Cana versus HK-2 Ctrl.
At present, the underlying mechanism(s) by which genetic deletion of HNRNPF led to down-regulation of SGLT2 transcription in HK-2 cells are unclear. One possibility might be that HNRNPF affects SGLT2 transcription at the promoter activity level. This is unlikely since transcription of the HNRNPF cDNA did not affect the SGLT2 promoter activity (pGL4.2/SGLT2-N-1,986/+22 promoter) in HK-2 cells (Supplemental Fig. 6a). However, we could not rule out the possibility of putative HNRNPF-response element(s) upstream of 2 kb of the SGLT2 promoter. The second possibility is that HNRNPF deletion might alter the splicing of SGLT2 to yield mutant forms of SGLT2. This is also unlikely since only one species of SGLT2 was detectable in HK-2 cells with HNRNPF KO, which was similar to the size of SGLT2 in HK-2 (Supplemental Fig. 6b). The third possibility is that HNRNPF might affect SGLT2 mRNA stability. This notion is supported by the observations of Chu et al.,19 that HNRNPF regulates YAP expression via binding to the 3′UTR of YAP to affect its mRNA stability and of Decorsiere et al.,21 that Hnrnpf/f interacts with a G-quadruplex in maintaining p53 pre-mRNA 3′-end processing during DNA damage. The fourth possibility is that deletion of HNRNPF might suppress other un-defined signaling pathway(s) or factor(s) that might have a greater impact (stimulation) on SGLT2 expression and activity. Clearly, further studies are needed to elucidate the mechanisms underlying HNRNPF down-regulation of SGLT2 expression.

The exact mechanism(s) of Hnrnpf regulation of Agt expression is unknown. One possibility is that Hnrnpf binds to the insulin-responsive element (IRE) in the Agt promoter11,12 and functions as a negative trans-acting protein to inhibit the binding of other positive trans-acting factor(s) to TATA-binding protein (TBP) and RNA polymerase II, subsequently attenuating Agt transcription. This possibility is supported by the studies of Yoshida et al.,22 showing that Hnrnpf is associated with TBP, RNA polymerase II and nuclear cap-binding protein complex. A second possibility is that Hnrnpf is associated with Hnrnpf to form an Hnrnpf/k complex and that the Hnrnpf/k complex is more effective in inhibiting Agt transcription. Indeed, we have previously reported that Hnrnpf co-immunoprecipitated with Hnrnpk and that co-transfection of Hnrnpf with Hnrnpk was more effective in inhibiting Agt transcription than either Hnrnpf or Hnrnpk alone.35 A third possibility is that the Hnrnpf/k complex might recruit unidentified repressor molecules and subsequently repress Agt transcription. This third possibility is supported by the studies of Deniseno et al.,36 demonstrating that Hnrnpk could bind the murine repressor Zik1. Clearly, more work is needed to elucidate the precise molecular mechanism of action of Hnrnpf on Agt transcription in RPTCs.

In summary, the present study reveals a novel role for Hnrnpf in the development of hypertension, tubule-Interstitial fibrosis and glycosuria in mice via up-regulation of Agt and down-regulation of SGLT2 expression in RPTCs, respectively. With the recent development of SGLT2 inhibitors as a novel treatment for diabetic patients37–41, it would be important to understand the regulation of SGLT2 expression. This finding raises the possibility that Hnrnpf/k might affect the regulation and familial renal glycosuria in human.

**Methods**

**Chemical and reagents.** Fluorescein isothiocyanate-labeled inulin and canagliflozin (Invokana) were purchased from Sigma-Aldrich (Oakville, ON, Canada) and Janssen Inc. (Toronto, ON, Canada), respectively. Dulbecco Modified Eagle Medium (DMEM) (Cat. No. 11966-025), Ham’s F12 medium (Cat. No. 11765-054) and fetal bovine serum (FBS) were bought from Gibco (Thermo Fisher Scientific, Montreal, QC, Canada). Oligonucleotides were synthesized by Integrated DNA Technologies, Inc. (Coralville, IA) and listed in Supplemental Table 1. Restriction and modifying enzymes were purchased from New England Biolabs (Whitby, ON, Canada). The sources of antibodies used are listed in Supplemental Table 2. HK-2 (an immortalized human renal proximal tubular cell line) (Cat. No. CRL-2190) was obtained from American Tissue Cell Collection (ATCC) (Manassas, VA) (http://www.atcc.org). Human SGLT2 gene promoter (N-1,986/+) was amplified from HK-2 genomic DNA by PCR with specific primers (Supplemental Table 1) and then inserted into pGL4.20 reporter vector (Promega, Sunnyvale, CA) at XhoI and Bgl II restriction sites.

**Generation of tubular Hnrnpf KO mice.** Tubule-specific Hnrnpf KO mice were generated by cross-breeding male Hnrnpf floxed mice with female Pax8-Cre mice (Stock number: 028196; Jackson Laboratory, Bar Harbor, ME). Briefly, the murine Hnrnpf gene (Gene ID: 98758) is localized on chromosome 6: 117,900,340-117,925,622. Four exons have been identified in Hnrnpf with the ATG start codon and TAG stop codon both located in exon 4. The lox-modified Hnrnpf targeting vector was created by including 5′ and 3′ homology arms as well as two loxP sites flanking the fourth exon region amplified from SV129 BAC genomic DNA and confirmed by sequencing. C57BL/6 ES cells were used for gene targeting (Cyagen Biosciences, Santa Clara, CA). These mice allow the excision of exon 4 of Hnrnpf gene and disruption of the protein expression in the presence of Cre recombinase. By cross-breeding male Hnrnpf floxed mice with female Pax8-Cre mice, heterozygous Hnrnpf floxed allele mice were generated (genotype: Hnrnpflox/+Cre). These mice were back-crossbred to generate homozygous Hnrnpf floxed allele and carrying the Cre allele (genotype: Hnrnpflox/+Cre). The Pax8-Hnrnpf KO mice (genotype: Hnrnpflox/+Cre) and control littermates (Ctrl) (genotype: Hnrnpflox/+Cre) as well as heterozygous littermates (genotype: Hnrnpflox/+Cre and Hnrnpflox/+Cre) were used in the present studies. Experimental mice were generated from at least three different breeding couples. Offspring were genotyped by PCR to detect the Cre-recombinase as well as the presence or absence of the 5′ loxP site using specific primers (Supplemental Table 1).

**Physiological studies.** Age- and sex-matched male and female KO (genotype: Hnrnpflox/+Cre and control littermates (Ctrl) (genotype: Hnrnpflox/+Cre) were studied. Animal care and procedures followed the Principles of Laboratory Animal Care (NIH Publication No. 85-23, revised 1985 (http://grants1.nih.gov/grants/oalaw/references/ghsopol.htm) and were approved by the CRCHUM Animal Care Committee.
Weekly random blood glucose levels were measured in mice by Accu-Chek Performa (Roche Diagnostics, Laval, QC, Canada). SBP was measured with BP-2000 tail-cuff (Visitech Systems, Apex, NC) at least 2 to 3 times per week per animal in the morning without fasting as previously described. \(^{11,14}\) Baseline SBP was measured daily over a 5-day period before initiation of actual measurement at 6 weeks of age.

At 24 weeks of age, twenty-four h prior to euthanasia, mice were housed individually in metabolic cages. Food, water consumption, and urine output were recorded. Mouse serum and urine samples were extracted with C18 Sep-Pak columns (Waters, Mississauga, ON) and assayed for Ang II by specific ELISA (Bachem America, Torrence, CA) according to the recommended number III protocol. \(^{13,14,20,43}\) Urines were also assayed for levels of albumin and creatinine (ELISA, Albuwell and Creatinine Companion, Exocell, Inc., Philadelphia, PA) \(^{13,14}\) and glucose (Glucose colorimetric kit, Cayman Chemical, Ann Arbor, MI).

For tissue studies, mice were euthanized at the age of 8 or 24 weeks. Blood samples were collected by cardiac puncture. The kidneys were isolated, decapsulated and weighed. The left kidneys were processed for histology and immunostaining, and the right kidneys were used for isolation of renal proximal tubules (RPTs) by Percoll gradient. \(^{13,14,20,43}\) Aliquots of freshly-isolated RPTs from individual animals were used immediately for total RNA isolation and Western blotting.

**Serum and urine biochemical measurements.** Serum and urine sodium, phosphorus and calcium were measured by the Comparative Medicine and Animal Resources Centre, McGill University (Montreal, QC, Canada).

**Glomerular filtration rate.** The glomerular filtration rate (GFR) was estimated with fluorescein isothiocyanate inulin as recommended by the Animal Models of Diabetic Complications Consortium (http://www.diacomp.org/) with slight modifications. \(^{11,14}\)

**Intraperitoneal glucose tolerance Test.** An intraperitoneal glucose tolerance test (IPGTT) was performed after 6h fasting in non-anesthetized mice at the age of 23 weeks, as described previously. \(^{44}\)

**Real time-quantitative polymerase chain reaction.** Real time-quantitative polymerase chain reaction (RT-qPCR) analyses were performed to quantify the relative expression of \(^{11,14,20,43}\) Hnrnpf (RT-qPCR) analyses were performed to quantify the relative expression of Hnrnpf and other genes by RT-qPCR (Supplemental Table 1).

**Western blotting.** Western Blotting (WB) was performed in isolated RPTs as described previously. \(^{13,14,20,43}\) Details of the sources of antibodies and working dilutions are listed in Supplemental Table 2.

**Histology.** Kidney sections were stained with periodic acid Schiff (PAS) as previously described with specific primers (Supplemental Table 1).

**Immunofluorescence staining.** Immunofluorescence (IF) staining was performed on 3-μm tissue sections from mouse kidney fixed in formalin and embedded in paraffin followed by staining with ALEXA FLUOR-594-labeled secondary antibody (Invitrogen). Proximal tubules were identified by fluorescein-labeled lotus tetragonolobus lectin (LTL, a marker of renal proximal tubule) (Vector Labs, Burlingame, CA). Image quantification and merge were assessed by ImageJ software (http://rsb.info.nih.gov/ij/). To quantify the amount of Sglt2 expression, the pixel intensity of Sglt2 was divided by LTL intensity. To calculate the average ratio, 6 sections per mouse, 6 mice per group were analyzed.

**Human renal proximal tubular cells with or without HNRNPF.** Human renal proximal tubular cells (RPTCs) (HK-2) cells are derived from a normal adult male human kidney transfected with the human papilloma virus 16 (HPV-16) E6/E7 genes. \(^{50}\) KO of HNRNPF in HK-2 was performed by the CRISPR-Cas9 genome editing method provided by Invitrogen (TrueGuide™). Briefly, the day before transfection, HK-2 cells (2.5 × 10^5 cells per well) were cultured in a 1:1 mixture of DMEM and Ham’s F12 medium containing 10% of FBS in 6-well plate. OPTI-MEM medium with Lipofectamine Cas9 Plus™ Reagent (Cat. No. CMAX00001, Invitrogen) and the mixture of 37.5 pmol TruCut™ Cas9 Protein v2 (Cat. No. A36497, Invitrogen) and 37.5 pmol gRNA (crRNA (Cat. No. A35509, CRISPR1099776_CR, Invitrogen) / tracrRNA (Cat. No. A35506, Invitrogen)) were transfected to HK-2 and cultured for 2 days at 37 °C. Single cell clones were then isolated by using limiting dilution cloning in 96-well plates. The positive clones were identified for the absence of HNRNPF by WB of cellular extracts and confirmed by PCR of genomic sequence. The clones with HNRNPF expression were used as controls.

To test the pharmacologic effect of SGLT2 inhibition on SGLT2 and AGT expression, HK-2 cells were harvested after 24 hours of culture in serum-free normal glucose (5 mM) DMEM in the absence or presence of 0.5 mM canagliflozin as described by Pirklbauer et al. \(^{51}\). WB and RT-qPCR were used to quantify SGLT2 and AGT protein and mRNA expression, respectively.

**Canagliflozin treatment in wild-type (WT) mice.** To investigate the impact of Sglt2 inhibition and Agt expression in RPTCs in vivo, male and female WT mice were treated with or without canagliflozin (0.2 mg/ml in drinking water) at the age of 4 weeks as described previously. \(^{52}\) Body weight, blood and urinary glucose and SBP...
were monitored weekly. The mice were euthanized at the age of 8 weeks. The left kidneys were processed for histology and immunostaining, and the right kidneys were used for isolation of RTPs and were used immediately for total protein and RNA isolation to quantify protein and mRNA expression of Agt and Slgt2 by WB and RT-qPCR, respectively.

**Statistical analysis.**  The data are expressed as means ± SEM. Statistical significance between the experimental groups was analyzed by Student’s t-test or 1-way ANOVA (analysis of variance) and the Bonferroni test as appropriate. p < 0.05 values were considered to be statistically significant.

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Author contributions
J.S.D.C. and S.L.Z. are co-guarantors (principal investigators) responsible for study conception and design and, as such, have full access to all study data, taking responsibility for data integrity and the accuracy of data analysis. C.S.L. contributed to data research and discussion and drafted the manuscript. K.N.M., S.Z., A.G., S.Y.C. and I.C. contributed to the in vivo and in vitro experiments and data collection. J.G.F. and J.R.I. contributed to the discussion, and reviewed/edited the manuscript. All authors approved the final version for publication.

Competing interests
The authors declare no competing interests.

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