Water Deprivation Increases (Pro)renin Receptor Levels in the Kidney and Decreases Plasma Concentrations of Soluble (Pro)renin Receptor

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Water deprivation activates the renin-angiotensin system. We have hypothesized that the renal expression of (pro)renin receptor ((P)RR), a specific receptor for renin and prorenin, could be changed under dehydration. Moreover, plasma levels of soluble (P)RR (s(P)RR) comprising of the extracellular domain of (P)RR may reflect the renal (P)RR expression. In the present study, we therefore aimed to clarify changes of plasma s(P)RR concentrations and kidney tissue (P)RR levels using rats with dehydration. Male Wister-Kyoto rats were divided into two groups; dehydrated (DH) rats deprived of water for 72 hours with free access to food, and control rats. Plasma s(P)RR concentrations measured by enzyme-linked immunosorbent assay were significantly lower in DH rats (6.94 ± 2.08 ng/mL, mean ± SD, n = 5) than in control (12.54 ± 2.00 ng/mL, n = 5) (p < 0.05). Western blot analysis confirmed lower expression levels of s(P)RR in plasma in DH rats than in control. By contrast, western blot analysis showed higher levels of full-length (P)RR and lower levels of furin (an enzyme responsible for generation of s(P)RR from full-length (P)RR) in the kidney tissues obtained from DH rats compared to control. There was no significant difference in the renal (P)RR mRNA levels between DH rats and control. These findings suggest that water deprivation may elevate the renal full-length (P)RR levels via reducing the expression of furin. Increased full-length (P)RR may contribute to the up-regulation of the renal renin-angiotensin system and the production of concentrated urine under dehydration.

Keywords: dehydration; prorenin; receptor; renin; water deprivation

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Introduction

(Pro)renin receptor ((P)RR), a specific receptor for renin and prorenin, is a 350-amino acid protein with a single transmembrane domain (Nguyen et al. 2002; Nguyen 2011). The binding of prorenin to (P)RR activates the angiotensin I (Ang I)-generating activity of prorenin, and directly stimulates the intracellular signalling pathways, including extracellular signal-regulated kinase (ERK 1/2) (Nguyen et al. 2002; Nguyen 2011). Soluble (P)RR (s(P)RR) comprising of the extracellular domain of (P)RR, and a truncated form of (P)RR at the C-terminal region (truncated (P)RR) are generated from full-length (P)RR (f(P)RR) by furin (Cousin et al. 2009; Nguyen 2011). The truncated (P)RR is associated with vacuolar H⁺-ATPase (v-ATPase), an ATP-dependent multi-subunit proton pump (Ludwig et al. 1998). (P)RR is also related to the Wnt/β-signaling pathway (Cruciat et al. 2010).

Soluble (P)RR has been shown to be present in plasma and urine (Cousin et al. 2009; Gonzalez et al. 2011; Maruyama et al. 2013). Plasma concentrations of s(P)RR were elevated in patients with chronic kidney disease (Hamada et al. 2013), pregnant women with the development of gestational diabetes mellitus or hypertension later in pregnancy (Watanabe et al. 2012, 2013), and patients with obstructive sleep apnea syndrome (Nishijima et al. 2013).
Water deprivation is a progressive physiological condition associated with increased plasma osmolality and activation of the renin-angiotensin system (RAS) (Gottlieb et al. 2006). RAS plays an essential role in the regulation of arterial blood pressure and electrolyte balance (Laragh 1967). Because angiotensin activity can correct such disturbances, it is generally accepted that RAS is a homeostatic mechanism that maintains the quantity, composition, and distribution of body fluids.

Immunohistochemical studies showed that (P)RR was diffusely expressed in renal tubular and collecting duct cells of both human and rat kidneys (Hirose et al. 2009, 2010; Takahashi et al. 2010b). Expression levels of (P)RR in the kidney were up-regulated in rats with congestive heart failure or chronic renal failure due to 5/6 nephrectomy (Hirose et al. 2009, 2010). Ramkumar et al. (2015) showed that nephron-specific deletion of the (P)RR caused a urine concentration defect, and suggested that nephron (P)RR can potentially modulate renal water excretion. We have recently reported that high salt intake upregulates (P)RR expression in rat nephron (Rong et al. 2015). Our hypothesis is therefore that (P)RR expression in the kidney could be changed under dehydration and that plasma s(P)RR concentrations may reflect the renal (P)RR expression. The aim of the present study was therefore to clarify changes of plasma s(P)RR concentrations and kidney tissue (P)RR levels in rats with dehydration.

Materials and Methods

Water deprivation experiment

The present study was approved by the Ethics Committee for Animal Experimentation of Tohoku University School of Medicine, and performed in accordance with the guidelines for animal experimentation of Tohoku University.

Male Wister-Kyoto (WKY) rats at 11 weeks of age were obtained from Charles River Japan Ltd. (Yokohama, Japan). Rats were kept in metabolic cages for 7 days for acclimatization before the start of treatments. Rats were housed under controlled humidity (30 ± 10%) and temperature (22 ± 2°C) with a 12-h light/dark cycle, were fed a regular diet, and had free access to tap water. After 7 days, rats were randomly divided into two groups: control (n = 5) and dehydrated (DH) rats (n = 5). For control rats, water and food were supplied ad libitum for 72 h, while DH rats were deprived of water for 72 h with free access to food (Chung et al. 2008; Hindmarch et al. 2011).

Urine samples were collected during the last 24 h of 72-hour water deprivation using a metabolic cage. After 72-hour water deprivation, the rats were anesthetized with pentobarbital sodium (50 mg/kg, injected intraperitoneally), and blood samples were collected from the cannulated abdominal aorta. Kidneys were quickly removed, snap-frozen in liquid nitrogen and stored at −80°C.

Western blot analysis

To prepare the renal fraction, frozen left kidney was pulverized in liquid nitrogen (Heineke et al. 2010). One-third of the powdered sample was used as the whole kidney fraction (Chmela et al. 2001). This fraction was homogenized in 100 mmol/L potassium buffer (pH 7.25) containing 30% glycerol, 1 mmol/L dithiothreitol, and 0.1 mmol/L phenylmethylsulfonyl fluoride. Microsomal and cytosolic fractions were separated from the remaining powdered sample, as described previously (Jorgensen 1974; Chmela et al. 2001).

Western blot analysis was performed as previously reported (Hirose et al., 2009, 2010). The protein extracts of the whole kidney (12.5 µg/lane), microsomal fraction (12.5 µg/lane) and cytosolic fraction (50 µg/lane), and plasma samples (0.5 µl/lane) were loaded onto a 16% sodium dodecyl sulfate polyacrylamide gel and electrophoresed for 1 h at 150 V. The (P)RR antiserum was used at a dilution of 1:10,000 (Hirose et al. 2009, 2010). The (P)RR antiserum recognizes dual bands at 28 kDa and 35 kDa, which are compatible with the predicted sizes for s(P)RR and f(P)RR, respectively.

It was reported that expression of aquaporin 2 (AQP2) rapidly increased in the rat kidney after 1 day of dehydration, with a small further increment on the following day (Rai et al. 1997). AQP2 protein expression was therefore investigated to confirm its elevation by dehydration in the kidney tissue using a rabbit anti-AQP2 antibody (dilution, 1:200; Santa Cruz Biotechnology, California, USA) (Saito et al. 1997). In addition, furin expression was investigated in the renal microsomal fraction using a rabbit anti-furin antibody (dilution, 1:200; Santa Cruz Biotechnology) (Freyer et al. 2007). As an internal control, β-actin expression was studied using anti β-actin monoclonal antibody (1:10,000; Santa Cruz Biotechnology).

The relative intensities of the bands at 35 kDa for f(P)RR, 28 kDa for s(P)RR, and 36–45 kDa and 29 kDa for AQP2 were quantified using Image J software (version 1.40, National Institutes of Health, Bethesda, Maryland, USA).

RNA extraction and competitive, quantitative RT-PCR

Total RNA was extracted from the kidney tissue by the guanidinium isothiocyanate/cesium chloride method, and 4 µg of total RNA were reverse transcribed with 400 units of Moloney Murine Leukemia Virus reverse transcriptase (PrimeScript; TaKaRa, Otsu, Japan) using an oligo(dT) primer, as previously described (Mori et al. 2006). RAS plays an essential role in the regulation of arterial blood pressure and electrolyte balance (Laragh 1967). Because angiotensin activity can correct such disturbances, it is generally accepted that RAS is a homeostatic mechanism that maintains the quantity, composition, and distribution of body fluids.

Hormone measurement

Plasma renin activity (PRA) and plasma angiotensin II (Ang II) levels were measured by enzyme immunoassay and radioimmunoassay, respectively (BML inc., Tokyo, Japan). Plasma osmotic pressure, urine creatinine and urine osmotic pressure were measured by a standard autoanalysis technique (BML inc., Tokyo, Japan). Plasma s(P)RR levels were determined using a commercially available enzyme-linked immunosorbent assay (ELISA) kit (Immuno-Biological Laboratories Co, Fujioka, Japan) according to the manufacturer’s instructions (Maruyama et al. 2013).

Statistical analysis

Data are shown as mean ± SD. Data between two groups were compared by unpaired Student’s t-test. Statistical significance was considered at P < 0.05.
Results

Effects of water deprivation on general parameters and AQP2 expression

Effects of 72-hour water deprivation on body weight, food intake, urine output and plasma and urine chemistry are shown in Table 1. Before the water deprivation, there was no significant difference in the body weight, food intake, urine output, or urinary osmolarity between the two groups (data not shown). Water deprivation for 72 h in WKY rats significantly decreased body weight and food intake. Water deprivation also reduced urine volume, and

| Parameters                        | Control (n = 5) | Dehydration (n = 5) |
|-----------------------------------|----------------|--------------------|
| Body weight (g)                   | 295.0 ± 7.3    | 236.9 ± 7.5**      |
| Food intake (g/24h)               | 20.9 ± 2.4     | 0.9 ± 0.5**        |
| Urine output (mL/24h)             | 10.7 ± 1.9     | 1.3 ± 0.4**        |
| **Plasma chemistry**              |                |                    |
| Osmotic pressure (mOsm/kgH2O)     | 286.8 ± 2.9    | 305.6 ± 16.8*      |
| Renin activity (ng/mL/h)          | 8.5 ± 3.8      | 13.5 ± 2.8*        |
| Angiotensin II (pg/mL)            | 409.6 ± 175.6  | 3595.6 ± 1086.9**  |
| **Urine chemistry**               |                |                    |
| Osmotic pressure (mOsm/kgH2O)     | 1566 ± 381.5   | 4090.0 ± 450.6**   |

Food intake and urine output were measured for the last 24 h of 72-hour water deprivation. Data are shown as mean ± SD.

*<p < 0.05, **<p < 0.01 vs. control.

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**Fig. 1. Effects of 72-hour water deprivation on aquaporin 2 protein expression in the kidney. Upper panel: Western blot analysis of AQP2 protein and β-actin (internal control) in the whole kidney obtained from control rats and 72-hour water-deprived rats (DH). Representative two cases out of 5 rats are shown per each group. The antibody labels 29 kDa and 36-45 kDa bands, corresponding to the non-glycosylated and glycosylated forms of AQP2. Lower panel: The relative intensity of the bands corresponding to the glycosylated form of AQP2 at 36-45 kDa (left) and the non-glycosylated form of AQP2 at 29 kDa (right) (the ratio of the total intensity of AQP2 band to that of β-actin band)(mean ± SD, n = 5). Expression levels of AQP2 protein were significantly increased in DH compared to control. **<p < 0.01.
increased urine and plasma osmolality. Plasma renin activity and plasma Ang II concentrations were significantly elevated in DH rats compared to control rats.

Western blot analysis confirmed that 72-hour water deprivation increased expression of AQP2 proteins in the kidney (Fig. 1, upper panel). The intensity of both bands at 36-45 kDa and 29 kDa representing glycosylated and non-glycosylated AQP2 proteins was significantly increased by 13.3- and 3.4-fold, respectively, in the whole kidney of DH rats as compared to control (Fig. 1, lower panel).

Plasma concentrations of soluble (pro)renin receptor

Plasma s(P)RR concentrations measured by ELISA were significantly lower in DH rats (6.94 ± 2.08 ng/mL) than in control (12.54 ± 2.00 ng/mL) (p < 0.05, Fig. 2A). Western blot analysis of plasma showed a band at 28 kDa corresponding to s(P)RR (Fig. 2B). Consistent with the results in the ELISA, the intensity of the band corresponding to s(P)RR was significantly lower in DH rats than in control (26% of control, p < 0.01, Fig. 2B). There was a significant correlation between plasma s(P)RR levels measured by the ELISA, and the relative levels of plasma s(P)RR in the western blot analysis (r = 0.72, p < 0.005).

Western blot analysis of (P)RR in the kidney

We first determined the levels of 2 forms of (P)RR; f(P)RR and s(P)RR, in microsomal and cytosolic fractions by western blot analysis. As shown in Fig. 3, s(P)RR was detected primarily in the cytosolic fraction, while a lesser proportion was detected in the microsomal fraction. Full-length (P)RR was detected in all samples.

We then examined the effects of 72-hour water deprivation on (P)RR protein expression by western blot analysis. Expression levels of f(P)RR were greater in DH rats compared to control, in whole kidney (2.3-fold, p < 0.01, Fig. 4A), microsome fraction (3.0-fold, p < 0.01, Fig. 4B), and cytosolic fraction (22.9-fold, p < 0.01, Fig. 4C, upper panel), respectively. By contrast, there was no significant difference in the expression levels of s(P)RR in the cytosolic fraction between control and DH rats (Fig. 4C, lower panel).
Prorenin Receptor in Dehydration

Competitive, quantitative RT-PCR of (P)RR mRNA

The 72-hour water deprivation had no significant effects on expression levels of (P)RR mRNA in the kidney (control, 0.88 ± 0.13 mol/mol PRL32, n = 5; vs. DH rats 0.83 ± 0.19 mol/mol PRL32, n = 5, p > 0.1).

Western blot analysis of furin in the renal microsomal fraction

We finally examined the effects of the 72-hour water deprivation on the expression of furin, a protease responsible for the shedding of (P)RR, in the renal microsomal fraction. Expression levels of furin were significantly lower in DH rats than in control rats (70% of control, p < 0.05, Fig. 5).

Discussion

This study has shown increased tissue levels of f(P)RR protein in the kidney by 72-h water deprivation. By contrast, plasma s(P)RR concentrations and expression levels of furin protein in the kidney were decreased by 72-h water deprivation. Because the water deprivation had no significant effect on the expression levels of (P)RR mRNA in the kidney, the increased kidney tissue levels of f(P)RR protein may be due to the decreased cleavage of f(P)RR, possibly by furin. These findings suggested that dehydration by 72-h water deprivation decreased renal expression of furin, which resulted in decreased cleavage of f(P)RR into s(P)RR and truncated (P)RR in the kidney. The mechanism for the suppressed expression of furin in the kidney by dehydration remains to be clarified.

One possible action of increased renal f(P)RR in dehydration may be the contribution to the renal RAS activation, which results in increased Ang II generation. Increased Ang II may stimulate reabsorption of sodium and water in
suppressed the circulating RAS, whereas water deprivation activates it. It is therefore intriguing that upregulated f(P)RR levels in the kidney were found in both the conditions. Increased plasma osmolarity may be a cause and/or a target of up-regulated renal f(P)RR in both the high salt intake and dehydration. It was reported that dehydration induced a considerable increase in the daily output of sodium in urine (McKinley et al. 1983). Food and sodium intake fell during dehydration, but urinary sodium excretion increased significantly (Thrasher et al. 1984).

(P)RR is expressed, not only in the kidney, but also in various organs including brain (Hirose et al. 2009, 2010; Takahashi et al. 2010a, b; Nguyen 2011). (P)RR was co-localized with vasopressin in the paraventricular and supraoptic neurons of hypothalamus (Takahashi et al. 2010a). Shan et al. (2010) showed that (P)RR overexpression in the supraoptic nucleus stimulated vasopressin secretion and chronic knockdown of (P)RR decreased it in rats, suggesting that (P)RR plays a role in vasopressin secretion in the hypothalamus. Hypothalamic (P)RR may therefore modulate the vasopressin secretion against dehydration. Although the present study studied only plasma and kidney tissues, we could not deny the possibility that the similar alteration in the (P)RR expression levels occurred in other organs including the brain, affecting the decreased plasma concentrations of s(P)RR under dehydration.

Plasma s(P)RR concentrations were significantly decreased by 72-h water deprivation (Fig. 2), whereas the decrease of the renal s(P)RR levels in the cytosolic fraction did not reach the significance (Fig. 4C, lower panel). Newly generated s(P)RR may be secreted soon into blood by exocytosis, and therefore, the plasma levels of s(P)RR may reflect more accurately the amount of the newly-generated s(P)RR from f(P)RR by furin rather than the tissue s(P)RR levels. On the other hand, the physiological roles of s(P)RR in blood have not been clarified. Nguyen et al. (2014) reported that plasma levels of s(P)RR were independent of plasma renin, prorenin, and aldosterone concentrations in man. Moreover, plasma levels of s(P)RR showed no significant correlation with PRA or plasma aldosterone levels in patients with obstructive sleep apnea syndrome (Nishijima et al. 2014, 2016). It has been reported that most soluble receptors act as antagonists and compete with their membrane-associated counterparts for ligands (Rose-John et al. 2006; Tracey et al. 2008). There have been no reports, however, which show that s(P)RR in blood acts as an antagonist against renin. Thus, it remains to be clarified whether decreased plasma s(P)RR concentrations have some pathophysiological significance in dehydration, or just a sequence of increased tissue f(P)RR levels.

In conclusion, the present study has shown that increased tissue levels of f(P)RR protein in the kidney by water deprivation possibly via decreased furin-mediated cleavage. In contrast to the increased renal f(P)RR levels, plasma s(P)RR concentrations were decreased by water deprivation. (P)RR may play an essential role in the up-

the renal tubules and collecting ducts against dehydration. We confirmed the elevation of PRA and plasma Ang II levels under dehydration as shown in Table 1, although we could not deny the possibility that anesthesia affected these levels (Pettinger et al. 1975; Siragy et al. 1995). Furthermore, dehydration-induced (P)RR may promote the kidney dysfunction via Ang II, and the RAS-independent intracellular signalling pathways, including ERK 1/2 and transforming growth factor-β.

Another possible action of increased f(P)RR in the kidney under dehydration may be mediated by the association with v-ATPase, which plays an essential role in the secretion of H+ into urine in the type A intercalated cells of the collecting duct (Roy et al. 2015). It is now evident that intercalated cells have regulatory roles in the salt and water reabsorption by a mechanism involving v-ATPase (Chambrey et al. 2013; Roy et al. 2015). It should be noted, however, that the cleavage of f(P)RR by furin may be dispensable for the biogenesis of active v-ATPase (Kinouchi et al. 2013). Decreased furin expression may result in decreased truncated (P)RR in the kidney, and decreased v-ATPase activity. Because our (P)RR could not recognise truncated (P)RR, further studies are required to clarify whether the renal v-ATPase activity is decreased or not under dehydration.

The up-regulated f(P)RR expression in rat nephron by high salt intake was reported by us (Rong et al. 2015), and other investigators (Huang et al. 2012). High salt intake
regulation of the renal RAS and the production of concentrated urine under dehydration.

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**Conflict of Interest**

The authors declare no conflict of interest.

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