High glucose stimulates expression of aldosterone synthase (CYP11B2) and secretion of aldosterone in human adrenal cells

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Aldosterone synthase is the key rate-limiting enzyme in adrenal aldosterone production, and induction of its gene (CYP11B2) results in the progression of hypertension. As hypertension is a frequent complication among diabetes patients, we set out to elucidate the link between diabetes mellitus and hypertension. We examined the effects of high glucose on CYP11B2 expression and aldosterone production using human adrenal H295R cells and a stable H295R cell line expressing a CYP11B2 5'-flanking region/luciferase cDNA chimeric construct. D-glucose, but not its enantiomer L-glucose, dose-dependently induced CYP11B2 transcription and mRNA expression. A high concentration (450 mg/dL) of D-glucose time-dependently induced CYP11B2 transcription and mRNA expression. Moreover, high glucose stimulated secretion of aldosterone into the media. Transient transfection studies using deletion mutants/NBRE-1 point mutant of CYP11B2 5'-flanking region revealed that the NBRE-1 element, known to be activated by transcription factors NGFIB and NURR1, was responsible for the high glucose-mediated effect. High glucose also induced the mRNA expression of these transcription factors, especially that of NURR1, but NURR1 knockdown using its siRNA did not affect high glucose-induced CYP11B2 mRNA.
expression. Taken together, it is speculated that high glucose may induce CYP11B2 transcription via the NBRE-1 element in its 5’-flanking region, resulting in the increase of aldosterone production although high glucose-induced NURR1 is not directly involved in the effect. Additionally, glucose metabolism and calcium channels were found to be involved in the high glucose effect. Our observations suggest one possible explanation for the high incidence of hypertension in diabetic patients.

Keywords: Aldosterone synthase, NURR1, Hypertension, Diabetes mellitus

Abbreviations: NURR1, Nur-related factor 1; NGFIB, nerve growth factor-induced clone B; NBRE, NGFIB response element; SF-1, steroidogenic factor-1; CREB, cAMP-response element binding protein; CREM, cAMP- response element modulator; COUP-TF, chicken ovalbumin upstream promoter transcription factor; ATF, activating transforming factor; ARB, angiotensin II receptor blocker; CCB, calcium channel blocker.

Running heading: Effects of high glucose on CYP11B2 expression

1. Introduction

The number of patients with diabetes mellitus is increasing every year, and 382 million people in the world were estimated to be affected in 2013 [1]. Among diabetic patients, hypertension is one of the most frequently observed complications. In Japan,
the incidence of hypertension in diabetic patients is approximately 60%, which is twice that in non-diabetic people [2]. The etiology of hypertension in diabetic patients is partially explained by the effect of hyperinsulinemia on renal proximal tubules due to insulin resistance [3]. Additionally, endothelial dysfunction and atherosclerosis induced by diabetes mellitus may also contribute to the progression of hypertension [4]. However, the direct involvement of high glucose on the etiology of hypertension in diabetic patients still remains uncertain.

The renin-angiotensin-aldosterone system (RAAS) is known as the main humoral pathway involved in the etiology of hypertension, and aldosterone, the final product of the pathway, plays an important role in the progression of hypertension and vascular damages in combination with sodium [5]. Aldosterone is synthesized in the zona glomerulosa of the adrenal cortex from cholesterol catalyzed via side chain cleavage enzyme (CYP11A1), 3β-hydroxysteroid dehydrogenase (3β-HSD), steroid 21-hydroxylase (CYP21), and aldosterone synthase (CYP11B2), which is the key rate-limiting enzyme in aldosterone production [6]. Aldosterone synthase gene (CYP11B2) expression is mainly regulated by angiotensin II (AII) and potassium via transcription factors including NURR1 [7]. Recently, genetic analyses of KCNJ5, ATP1A1, ATP2B3, and CACNA1D have revealed that chronic overexpression of CYP11B2 induces not only aldosterone hypersecretion but also the formation of primary aldosteronism [8], resulting in the progression of severe hypertension. Moreover, aberrant WNT signaling caused by mutations in CTNNB1 has also been recognized to be involved in the formation of primary aldosteronism [9]. In order to investigate the direct link between hypertension and diabetes mellitus, we here examined the effects of high glucose on CYP11B2 expression and aldosterone secretion using human adrenal
2. Materials and methods

2.1. Reagents

D-glucose was purchased from Wako (Osaka, Japan), and L-glucose, used for the adjustment of osmolality, was purchased from Sigma (St. Louis, MO). 2-deoxy-D-glucose, D-sorbitol, D-fructose, and 3-O-methyl-D-glucose were purchased from Sigma. Olmesartan (olmesartan medoxomil) was purchased from Toronto Research Chemicals (North York, Canada). Losartan (losartan potassium) was purchased from LKT Laboratories (St. Paul, MN). Valsartan was purchased from Cayman Chemical (Ann Arbor, MI). Candesartan (trityl candesartan cilexetil) was purchased from Sequoia Research Products (Pangbourne, UK). Nifedipine and efonidipine (efonidipine hydrochloride monoethanolate) were purchased from Sigma. Amlodipine was purchased from Cayman Chemical. Benidipine (benidipine hydrochloride) was kindly provided by Kyowa Hakko Kirin Pharma (Tokyo, Japan). Human AII was purchased from Sigma.
2.2. Plasmids

Subcloned chimeric constructs containing the human CYP11B2 genomic DNA and luciferase cDNA (pGL3-Basic, Promega, Madison, WI) [7, 10] were used for the transient transfection studies: -1521/+2-luc (harboring the CYP11B2 5'-flanking region from -1521 to +2 relative to the transcription start site upstream of the luciferase cDNA in pGL3-Basic), -747/+2-luc; -135/+2-luc; -106/+2-luc; -65/+2-luc. The NBRE-1 mutant construct of -1521/+2-luc (NBRE-1 mut) was also used [11]. In some experiments, a previously described stable H295R cell line expressing CYP11B2 promoter (-1521/+2)/luciferase chimeric reporter construct (CYP11B2-H295R cells) was used [7]. β-galactosidase control plasmid in pCMV (pCMV-β-gal) was purchased from Clontech (Palo Alto, CA).

2.3. Cell culture

H295R cells or CYP11B2-H295R cells were grown with a 1:1 mixture of DMEM and Ham’s F12 medium supplemented with 10% fetal bovine serum (FBS), Insulin-Transferrin-Selenium-G Supplements (Invitrogen, Carlsbad, CA), 1.25 mg/mL BSA (Sigma), 5.35 μg/mL linoleic acid (Sigma), 100 U/mL penicillin, 100 μg/mL streptomycin. Cells were cultured in a humidified incubator at 37°C with 5% CO₂. Since the D-glucose concentration in the media was approximately 100 mg/dL, we added either concentrated D-glucose or L-glucose solution to adjust the final concentration. For example, the 450 mg/dL D-glucose concentration was composed of 100 mg/dL D-glucose from the media and 350 mg/dL D-glucose from the concentrated
D-glucose solution, and its osmolality-adjusted control was composed of 100 mg/dL D-glucose from the media and 350 mg/dL L-glucose from the concentrated L-glucose solution. In some experiments, 2-deoxy-D-glucose, 3-O-methyl-D-glucose, D-sorbitol, or D-fructose was used instead of L-glucose. Moreover, CYP11B2-H295R cells were incubated either with angiotensin II receptor blockers (ARBs) or calcium channel blockers (CCBs) in the presence of 450 mg/dL D-glucose.

2.4. RNA Preparation and quantitative real-time PCR

When H295R cells were grown to 60% confluence in 24-multiwell plates, they were exposed to several concentrations of D-glucose or D-glucose plus L-glucose for the indicated times, and their total RNA was extracted using Sepasol®-RNA I Super G (Nacalai Tesque, Kyoto, Japan) according to the manufacturer’s instructions. Total RNAs were subjected to reverse transcription (RT) reaction using PrimeScript Reverse Transcriptase (Takara Bio, Ohtsu, Japan) with random 6mer and oligo dT primers according to the manufacturer’s instructions. Thereafter, the obtained templates were used for quantitative real-time PCR (95 °C, 3 min for 1 cycle; 95 °C, 15 sec; 60 °C, 10 sec; 72 °C, 20 sec for 40 cycles) either with iQ Supermix (Bio-Rad, Hercules, CA) (for CYP11B2, CYP11B1, HSD3B2, and CYP17) or THUNDERBIRD® SYBR® qPCR Mix (TOYOBO, Osaka, Japan) (for others) by DNA Engine thermal cycler attached to Chromo4 detector (Bio-Rad). The sequences of the primers and TaqMan probes are shown in Table 1.

2.5. Transient transfection and luciferase assay

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H295R cells were plated to 60% confluence in 24-multiwell plates. Thereafter, they were transiently transfected with 200 ng luciferase reporter plasmids and 100 ng pCMV-β-gal using Lipofectamine® 2000 Transfection Reagent (Life Technologies, Carlsbad, CA) for 24 hours. The cells were then exposed to D-glucose or L-glucose for the indicated times and concentrations. They were thereafter washed with PBS, and the cell extracts were prepared using Glo Lysis Buffer (Promega). Luciferase activity was measured using Bright-Glo reagents (Promega), and β-galactosidase activity was simultaneously measured. Data were normalized by the β-galactosidase activities. When the stable CYP11B2-H295R cells [7] were used, only the luciferase activity was measured.

2.6. Small interfering RNA transfection

Small interfering RNA (siRNA) for NURR1 (s9785) [12] was obtained from Thermo Fisher Scientific (Waltham, MA), and negative control siRNA (SI03650318) was obtained from Qiagen (Hilden, Germany). H295R cells were plated to 60% confluence in 12-multiwell plates. Thereafter, they were transiently transfected with 10 pmol of each siRNA by electroporation using Nucleofector 4D™ (Lonza, Basel, Switzerland) as previously described [13].

2.7. Measurement of aldosterone/cortisol concentration

H295R cells were plated to 60% confluence in 24-multiwell plates. Thereafter, they
were exposed to either 100 mg/dL D-glucose, 450 mg/dL D-glucose, 100 mg/dL D-glucose plus 100 nmol/L AII (for aldosterone), or 450 mg/dL D-glucose plus 100 nmol/L AII (for aldosterone) for 72 hours. The aldosterone and cortisol concentrations of the media were thereafter measured by Aldosterone EIA Kit and Cortisol EIA Kit (Cayman Chemical), respectively after their extraction with dichloromethane according to the manufacturer's instructions. The obtained data were normalized by the protein concentrations measured by Protein Assay Kit (Bio-Rad).

2.8. Statistical analyses

All data are presented as mean ± SEM. For the statistical analyses, ANOVA followed by post hoc Tukey test was performed. *P*<0.05 was considered statistically significant.

3. Results

3.1. Effects of high glucose on CYP11B2 expression and aldosterone secretion

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We first examined the effects of high glucose on CYP11B2 mRNA expression using H295R cells. As shown in Fig. 1A, D-glucose levels above 270 mg/dL significantly induced CYP11B2 mRNA expression. Time course experiments in the presence of 450 mg/dL D-glucose demonstrated that high glucose induced CYP11B2 mRNA expression after 48 hours (Fig. 1B). We next examined the effect of high glucose on CYP11B2 transcription using stable CYP11B2-H295R cells [7], and also observed similar stimulatory effects in both the dose response (Fig. 1C) and time course (Fig. 1D) experiments. We also examined the effect of high glucose (450 mg/dL D-glucose) on the mRNA expression of other enzymes/protein involved in adrenal steroidogenesis. As shown in Fig. 2, high glucose treatment significantly induced the mRNA expression of 11β-hydroxylase gene (CYP11B1) (A) and steroidogenic acute regulatory protein gene (StAR) (E) after 48 hours incubation, while high glucose treatment significantly decreased that of CYP11A1 (D) after 24 hours incubation. High glucose treatment did not affect the mRNA expression of 3β-HSD gene (HSD3B2) (B) and CYP21 (C), while it tended to decrease, although not significantly, the expression of 17α-hydroxylase/17,20 lyase gene (CYP17) (F). We then examined the effect of high glucose on aldosterone secretion from H295R cells. As shown in Fig. 3A, incubation of the cells in the presence of 450 mg/dL D-glucose for 72 hours significantly induced aldosterone secretion into the media, which was comparable to the AII-induced aldosterone secretion. Incubation with 450 mg/dL D-glucose plus AII did not further increase the aldosterone secretion (Fig. 3A). It can be concluded that high glucose induces CYP11B2 transcription and mRNA expression resulting in the increase of aldosterone secretion. In contrast, although high glucose induced the mRNA expression of CYP11B1 (Fig. 2A), it did not induce cortisol secretion into the media (Fig. 3B).
probably due to the decreasing trend of *CYP17* mRNA expression (Fig. 2F).

### 3.2. Identification of the element(s) responsible for the high glucose-induced CYP11B2 transactivation

In order to identify the element(s) responsible for the high glucose-induced *CYP11B2* transactivation, we examined the effects of high glucose on the promoter activity of *CYP11B2* 5'-flanking region deletion mutants by comparing the effects between D-glucose (450 mg/dL) and L-glucose (100 mg/dL D-glucose and 350 mg/dL L-glucose) using H295R cells. As shown in Fig. 4A, although high glucose-induced *CYP11B2* transactivation was significantly observed in -1521/+2-luc, it was not observed in -747/+2-luc, -135/+2-luc, -106/+2-luc, or -65/+2-luc. These data indicate that the region between -1521 and -747 may be responsible for the high glucose effect. Since NBRE-1 element [11], which is known to be transactivated by NURR1 and NGFIB [11, 14], is located within the region (-766/-759), we next examined the effect of high glucose on the element. As shown in Fig. 4B, point mutation of NBRE-1 element (NBRE-1 mut) completely abolished the high glucose effect. These data indicate that the element responsible for the high glucose-induced *CYP11B2* transactivation may possibly be the NBRE-1 element.

### 3.3. Effects of high glucose on the expression of transcription factors involved in CYP11B2 transcription

We next examined the effects of high glucose on the mRNA expression of
transcription factors that are known to regulate CYP11B2 promoter [14] using H295R cells. As shown in Fig. 5A, D-glucose (450 mg/dL), but not control (100 mg/dL D-glucose) or L-glucose (100 mg/dL D-glucose and 350 mg/dL L-glucose), significantly induced the expression of NURR1 mRNA. D-glucose also induced the mRNA expression of NGFI-B (Fig. 5B), but not that of SF-1 (Fig. 5C), CREB (Fig. 5D), CREM (Fig. 5E), COUP-TF (Fig. 5F), ATF-1 (Fig. 5G), or ATF-2 (Fig. 5H). Since NURR1 is known to bind to NBRE-1 and activate it [11], it is indicated that high glucose-induced NURR1 may transactivate CYP11B2 expression via the NBRE-1 element.

3.4. Effects of NURR1 siRNA on the high glucose-induced CYP11B2 mRNA expression

In order to examine the involvement of NURR1 in the high glucose-induced CYP11B2 mRNA expression, we next transfected either the control or NURR1 siRNA into H295R cells, and thereafter treated the cells with 450 mg/dL D-glucose for 48 hours. As shown in Fig. 6A, NURR1 mRNA expression was significantly decreased by the transfection of NURR1 siRNA as compared to that of control siRNA in the presence of either 100 mg/dL D-glucose or 450 mg/dL D-glucose, suggesting the efficient knockdown of endogenous NURR1 mRNA. However, NURR1 knockdown by its siRNA transfection did not affect the high glucose-induced CYP11B2 mRNA expression in comparison to control siRNA transfection (Fig. 6B). These data indicate that other NR4A family members or other transcription factors may be involved in the high glucose-induced CYP11B2 mRNA expression via the NBRE-1 element.
3.5. Effects of 2-deoxy-D-glucose, 3-O-methyl-D-glucose, D-sorbitol, and D-fructose on CYP11B2 mRNA expression

We next examined the involvement of glucose metabolism on the high glucose-induced CYP11B2 expression. When we treated H295R cells with either 2-deoxy-D-glucose, which could be phosphorylated but could not be metabolized further [15, 16], or 3-O-methyl-D-glucose, which could not be phosphorylated [16], the induction of CYP11B2 mRNA expression was not observed (Fig. 7). These data suggest that D-glucose metabolization may be more necessary for the induction than glucose 6-phosphate. Moreover, incubation with D-sorbitol or D-fructose, both of which are D-glucose metabolites via the polyol pathway [17], did not affect CYP11B2 mRNA expression (Fig. 7) indicating that the pathway may not be involved in the induction.

3.6. Effects of ARBs and CCBs on the high glucose-induced CYP11B2 transcription

We next examined the effects of ARBs and CCBs on the high glucose-induced CYP11B2 transcription. As shown in Fig. 8A, each ARB at 1 μmol/L did not affect the high glucose effect. In contrast, each CBB dose-dependently inhibited the high glucose-induced CYP11B2 transactivation (Fig. 8B). It is therefore suggested that calcium channels, but not AII type 1 receptor, are involved in the high glucose effect.
4. Discussion

In the present study, we first demonstrated the stimulatory effect of high glucose on

*CYP11B2* transcription and mRNA expression as well as aldosterone secretion in human

adrenal cells (Fig. 1, Fig. 3A). The high glucose-induced *CYP11B2* mRNA expression

was not observed when we used glucose analogs, 2-deoxy-D-glucose and

3-O-methyl-D-glucose [15, 16] (Fig. 7), suggesting that it is necessary for D-glucose to

be metabolized within the cells for the stimulatory effect. Since we also observed the

high glucose-induced *StAR* mRNA expression (Fig. 2E), increased *StAR* and *CYP11B2*

may coordinately induce aldosterone production. Aldosterone not only induces

hypertension and vascular damage in combination with sodium [5], but is also known to

inhibit glucose-induced insulin secretion in pancreatic β-cells [18] as well as insulin

signaling in peripheral tissues [19, 20]. Therefore, the high glucose-induced aldosterone

may induce “a vicious cycle” in terms of the exacerbation of glucose

intolerance/diabetes mellitus. Although the plasma aldosterone concentration in diabetic

patients has long been controversial, it has recently been confirmed to be significantly

higher than that of normal subjects by fixing sodium/potassium intake and the time for
drawing blood samples [21]. Therefore, based on our present observation, high
glucose-induced aldosterone production may possibly contribute to the increased
plasma aldosterone level in diabetic patients.

Transient transfection experiments using CYP11B2 5'-flanking region deletion
mutants and NBRE-1 point mutant revealed that the NBRE-1 element, which is known
to be activated by NURR1/NGFIB binding [11, 14], was responsible for the high
glucose-induced CYP11B2 transactivation (Fig. 4). Additionally, high glucose was
demonstrated to induce the mRNA expression of NURR1 significantly compared with
that of NGFIB (Fig. 5). However, since NURR1 knockdown did not affect the high
glucose-induced CYP11B2 mRNA expression (Fig. 6), other NR4A family members or
other transcription factors may bind to and activate NBRE-1 element to induce
CYP11B2 transactivation. In human adrenocortical neoplasms, CYP11B2 mRNA
expression significantly and positively correlated with NURR1 mRNA expression, but
not with NGFIB mRNA expression [22]. Since H295R cells are also derived from
human adrenocortical carcinoma, it is plausible that NURR1 also plays an indispensable
role in CYP11B2 transactivation in the cells. Interestingly, high glucose was
demonstrated to suppress SF-1 mRNA expression, although not significantly (Fig. 5C).
Since SF-1 is known to suppress CYP11B2 transcription [23, 24], the high
glucose-mediated SF-1 decrease may also contribute to the induction of CYP11B2
transactivation. The mechanisms by which high glucose induce NURR1 mRNA
expression remain uncertain. AII and potassium are two major factors that regulate
CYP11B2 transcription [14, 25]. AII is known to bind to AII type 1 receptor and activate
phospholipase C to increase inositol 1,4,5-trisphosphate (IP₃), and IP₃ induces the
release of intracellular calcium from the endoplasmic reticulum, while potassium causes
depolarization of the membrane allowing extracellular cytoplasmic calcium influx through the T- and L-type calcium channels [14, 25]. In both cases, increased calcium leads to the activation of calcium/calmodulin-dependent kinase, resulting in the induction of NURR1 mRNA expression [14, 25]. When we treated the stable CYP11B2-H295R cells with several ARBs, the high glucose-induced CYP11B2 transactivation was not affected (Fig. 8A). In contrast, when we treated the cells with several CCBs, they, especially benidipine that blocks both T- and L-type calcium channels [26], dose-dependently inhibited the high glucose-induced CYP11B2 transactivation (Fig. 8B). These data indicate that high glucose may affect at least pathway(s) mediated via T- and/or L-type calcium channels, but not pathway(s) mediated via AII type 1 receptor. Interestingly, we have recently observed the high glucose-induced mRNA expression of T-type calcium channel subunits (CaV3.1, CaV3.2, and CaV3.3) [27], which may also be involved in the high glucose effect. Further studies are needed to clarify the precise molecular mechanisms of the high glucose-induced CYP11B2 transactivation.

In summary, we here demonstrated high glucose-induced CYP11B2 transcription and mRNA expression as well as aldosterone secretion via NURR1 induction. Since our observation provides a novel insight in the etiology of hypertension in diabetic patients, it may also lead to novel therapeutics, such as an inhibitor of CYP11B2 transcription, for diabetic patients complicated with hypertension.

**Conflict of interest statement**

The authors declare that there are no conflicts of interest.
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Author contributions

AU, AY, and AS conceived and designed the experiments, HS, NK, EN, KS, IS, KS, MK, DS, and TSI performed the experiments, MK, RP, TSI, AU, ASH, WER, and AY analyzed the data, WER contributed reagents/materials/analysis tools, and HS, NK, TSI, SI, AY, and AS wrote the paper.
References


glucose uptake by degradation of insulin receptor substrate (IRS) 1 and IRS2 via a reactive oxygen species-mediated pathway in 3T3-L1 adipocytes. Endocrinology 150 (2009) 1662-1669.


Figure legends

Fig. 1. Effects of high glucose on CYP11B2 transcription and mRNA expression. (A) Dose-response effects of high glucose on CYP11B2 mRNA expression. H295R cells were incubated either with 100 mg/dL D-glucose, 100 mg/dL D-glucose plus 350 mg/dL L-glucose, 180 mg/dL D-glucose plus 270 mg/dL L-glucose, 270 mg/dL D-glucose plus...
180 mg/dL L-glucose, 350 mg/dL D-glucose plus 100 mg/dL L-glucose, or 450 mg/dL D-glucose for 48 hours. Data represent mean ± SEM (n = 4), percent of 100 mg/dL D-glucose (control), normalized by β-actin mRNA levels. (B) Time-course effects of high glucose on CYP11B2 mRNA expression. H295R cells were incubated with 450 mg/dL D-glucose for the indicated times. Data represent mean ± SEM (n = 4), percent of 0 hour (control), normalized by β-actin mRNA levels. (C) Dose-response effects of high glucose on CYP11B2 transcription. CYP11B2-H295R cells were incubated with several concentrations of D-glucose as in (A) for 48 hours. Data represent mean ± SEM (n = 4), percent of 100 mg/dL D-glucose (control). (D) Time-course effects of high glucose on CYP11B2 transcription. CYP11B2-H295R cells were incubated with 450 mg/dL D-glucose for the indicated times. Data represent mean ± SEM (n = 4), percent of 0 hour (control). (A-D) * P < 0.01, vs. control. ** P < 0.05, vs. control.

Fig. 2. Effects of high glucose on mRNA expression of enzymes/protein involved in adrenal steroidogenesis. Effects of high glucose on CYP11B1 mRNA expression (A), HSD3B2 mRNA expression (B), CYP21 mRNA expression (C), CYP11A1 mRNA expression (D), StAR mRNA expression (E), and CYP17 mRNA expression (F). H295R cells were incubated with 450 mg/dL D-glucose for the indicated times. Data represent mean ± SEM (n = 4), percent of 0 hour (control), normalized by β-actin mRNA levels. (A, D, E) * P < 0.01, vs. control. ** P < 0.05, vs. control.

Fig. 3. Effects of high glucose on aldosterone and cortisol secretion. (A) Effects of high glucose and/or AII on aldosterone secretion. H295R cells were incubated with either 100 mg/dL D-glucose, 450 mg/dL D-glucose, 100 mg/dL D-glucose plus 100 nmol/L AII.
AII, or 450 mg/dL D-glucose plus 100 nmol/L AII for 72 hours. Data represent mean ± SEM (n = 4), percent of 100 mg/dL D-glucose (control), normalized by the protein concentrations. (B) Effects of high glucose on cortisol secretion. H295R cells were incubated with either 100 mg/dL D-glucose or 450 mg/dL D-glucose for 72 hours. Data represent mean ± SEM (n = 4), percent of 100 mg/dL D-glucose (control), normalized by the protein concentrations. Ang II; AII. (A) ** P < 0.05, vs. control.

Fig. 4. Effects of 5’-flanking region mutants on the high glucose-induced CYP11B2 transactivation. (A) Effects of CYP11B2 5’-flanking region deletion mutants. Either -1521/+2-luc, -747/+2-luc, -135/+2-luc, -106/+2-luc, -65/+2-luc, or pGL3-Basic (control plasmid) was transiently transfected with pCMV-β-gal into H295R cells, and the cells were incubated either with 450 mg/dL D-glucose (D-glucose, D-glu) or 100 mg/dL D-glucose plus 350 mg/dL L-glucose (L-glucose, L-glu) for 48 hours. Data represent mean ± SEM (n = 4), percent of control (L-glucose), normalized by β-galactosidase activities. (B) Effects of NBRE-1 point mutant. Either -1521/+2-luc, NBRE-1 mut, or pGL3-Basic (control plasmid) was transiently transfected with pCMV-β-gal into H295R cells, and the cells were incubated either with 450 mg/dL D-glucose (D-glucose, D-glu) or 100 mg/dL D-glucose plus 350 mg/dL L-glucose (L-glucose, L-glu) for 48 hours. Data represent mean ± SEM (n = 4), percent of control (L-glucose), normalized by β-galactosidase activities. (A, B) * P < 0.01, vs. control.

Fig. 5. Effects of high glucose on mRNA expression of transcription factors. Effects of high glucose on NURR1 mRNA expression (A), NGFIB mRNA expression (B), SF-1 mRNA expression (C), CREB mRNA expression (D), CREM mRNA expression (E),
COUP-TF mRNA expression (F), ATF-1 mRNA expression (G), and ATF-2 mRNA expression (H). H295R cells were incubated either with 100 mg/dL D-glucose (control), 100 mg/dL D-glucose plus 350 mg/dL L-glucose (L-glu), or 450 mg/dL D-glucose (D-glu) for 48 hours. Data represent mean ± SEM (n = 4), percent of 100 mg/dL D-glucose (control), normalized by the β-actin mRNA levels. (A) * \( P < 0.01 \), vs. control. (B) ** \( P < 0.05 \), vs. control.

**Fig. 6.** Effects of NURR1 siRNA. Effects of NURR1 siRNA on NURR1 mRNA expression (A) and CYP11B2 mRNA expression (B). H295R cells transfected either with control siRNA (ctrl) or NURR1 siRNA (si NURR1) were incubated with either 100 mg/dL D-glucose or 450 mg/dL D-glucose for 48 hours. In some experiments, untransfected H295R cells were incubated with 100 mg/dL D-glucose for 48 hours (base). Data represent mean ± SEM (n = 11), percent of 100 mg/dL D-glucose (base), normalized by GAPDH mRNA levels. (A, B) * \( P < 0.01 \). ** \( P < 0.05 \).

**Fig. 7.** Effects of 2-deoxy-D-glucose, 3-O-methyl-D-glucose, D-sorbitol, and D-fructose on CYP11B2 mRNA expression. H295R cells were incubated either with 100 mg/dL D-glucose plus 350 mg/dL L-glucose, 450 mg/dL D-glucose, 100 mg/dL D-glucose plus 350 mg/dL 2-deoxy-D-glucose, 100 mg/dL D-glucose plus 350 mg/dL 3-O-methyl-D-glucose, 100 mg/dL D-glucose plus 350 mg/dL D-sorbitol, or 100 mg/dL D-glucose plus 350 mg/dL D-fructose for 72 hours. Data represent mean ± SEM (n = 4), percent of 100 mg/dL D-glucose plus 350 mg/dL L-glucose, normalized by the β-actin mRNA levels. * \( P < 0.01 \), vs. 100 mg/dL D-glucose plus 350 mg/dL L-glucose. ** \( P < 0.01 \), vs. 450 mg/dL D-glucose.
Fig. 8. Effects of ARBs (A) and CCBs (B) on the high glucose-induced CYP11B2 transactivation. In (A), CYP11B2-H295R cells were incubated either with 100 mg/dL D-glucose (control), 450 mg/dL D-glucose (D-glu), 450 mg/dL D-glucose plus 1 µmol/L losartan (D-glu + Los), 450 mg/dL D-glucose plus 1 µmol/L valsartan (D-glu + Val), 450 mg/dL D-glucose plus 1 µmol/L olmesartan (D-glu + Olm), or 450 mg/dL D-glucose plus 1 µmol/L candesartan (D-glu + Can) for 48 hours. Data represent mean ± SEM (n = 3), percent of control. * P < 0.01, vs. control. In (B), CYP11B2-H295R cells were incubated either with 100 mg/dL D-glucose, 450 mg/dL D-glucose, 450 mg/dL D-glucose plus 0.01 µmol/L amlodipine, 450 mg/dL D-glucose plus 0.1 µmol/L amlodipine, 450 mg/dL D-glucose plus 0.01 µmol/L benidipine, 450 mg/dL D-glucose plus 0.1 µmol/L benidipine, 450 mg/dL D-glucose plus 1 µmol/L benidipine, 450 mg/dL D-glucose plus 0.01 µmol/L efonidipine, 450 mg/dL D-glucose plus 0.1 µmol/L efonidipine, 450 mg/dL D-glucose plus 1 µmol/L efonidipine, 450 mg/dL D-glucose plus 0.01 µmol/L nifedipine, 450 mg/dL D-glucose plus 0.1 µmol/L nifedipine or 450 mg/dL D-glucose plus 1 µmol/L nifedipine for 96 hours. Data represent mean ± SEM (n = 4), percent of 100 mg/dL D-glucose. * P < 0.01, vs. 100 mg/dL D-glucose. ** P < 0.05, vs. 450 mg/dL D-glucose. *** P < 0.01, vs. 450 mg/dL D-glucose.
Fig. 1.

A) Relative mRNA expression of CYP11B2

B) Relative mRNA expression

C) Relative luciferase activity

D) Relative luciferase activity

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Fig. 2.

A. Relative mRNA expression (%) of CYP11B1 over time (hour).

B. Relative mRNA expression (%) of HSD3B2 over time (hour).

C. Relative mRNA expression (%) of CYP21 over time (hour).

D. Relative mRNA expression (%) of CYP11A1 over time (hour).

E. Relative mRNA expression (%) of StAR over time (hour).

F. Relative mRNA expression (%) of CYP17 over time (hour).
**Fig. 3.**

**A**

Relative aldosterone concentration (%)

- 100 mg/dL D-glucose
- 450 mg/dL D-glucose
- 100 mg/dL D-glucose
- 450 mg/dL D-glucose

**B**

Relative cortisol concentration (%)

- 100 mg/dL D-glucose
- 450 mg/dL D-glucose

+ 100 nmol/L Ang II

* * *

**N.S**

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Fig. 4.

A

B

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Fig. 5.

A. NURR1 expression
B. NGFIB expression
C. SF-1 expression
D. CREB expression
E. CREM expression
F. COUP-TF expression
G. ATF-1 expression
H. ATF-2 expression

* p < 0.05
** p < 0.01

Relative mRNA expression (%)
control L-glu D-glu control L-glu D-glu control L-glu D-glu control L-glu D-glu control L-glu D-glu control L-glu D-glu control L-glu D-glu
**Fig. 6.**

**A**

![Graph showing NURR1 relative mRNA expression](feb4_12277_f6.ai)

**B**

![Graph showing CYP11B2 relative mRNA expression](feb4_12277_f6.ai)
**Fig. 7.**

<table>
<thead>
<tr>
<th>Sugar</th>
<th>CYP11B2 Relative mRNA expression (%)</th>
</tr>
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<tbody>
<tr>
<td>D-glucose</td>
<td>100</td>
</tr>
<tr>
<td>L-glucose</td>
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</tr>
<tr>
<td>2-deoxy-D-glucose</td>
<td>350</td>
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<td>3-O-methyl-D-glucose</td>
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<tr>
<td>D-sorbitol</td>
<td>350</td>
</tr>
<tr>
<td>D-fructose</td>
<td>350</td>
</tr>
</tbody>
</table>

(mg/dL)
Fig. 8.

A

B

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