High aldosterone-to-renin variants of \( CYP11B2 \) and pregnancy outcome

Geneviève Escher\(^1\), Martino Cristiano\(^1\), Maja Causevic\(^1\), Marc Baumann\(^2\), Felix J. Frey\(^1\), Daniel Surbek\(^2\) and Markus G. Mohaupt\(^1\)

\(^1\)Department of Nephrology/Hypertension and \(^2\)Department of Obstetrics and Gynecology, University of Bern, Berne, Switzerland

Correspondence and offprint requests to: Markus G. Mohaupt; E-mail: markus.mohaupt@insel.ch

Abstract

Background. Increased aldosterone concentrations and volume expansion of normal pregnancies are hallmarks of normal pregnancies and blunted in pre-eclampsia. Accordingly, we hypothesized an active mineralocorticoid system to protect from pre-eclampsia.

Methods. In pregnant women (normotensive \( n = 44 \); pre-eclamptic \( n = 48 \)), blood pressure, urinary tetrahydro-aldosterone excretion and activating polymorphisms (SF-1 site and intron 2) of the aldosterone synthase gene \( CYP11B2 \) were determined; 185 non-pregnant normotensive individuals served as control. Amino acid-changing polymorphisms of the DNA- and agonist-binding regions of the mineralocorticoid receptor were evaluated by RT-PCR, SSCP and sequencing.

Results. Urinary tetrahydro-aldosterone excretion was reduced in pre-eclampsia as compared to normal pregnancy \( (P < 0.05) \). It inversely correlated with blood pressure \( (r = 0.99, P < 0.04) \). Homozygosity for activating \( CYP11B2 \) polymorphisms was preferably present in normotensive as compared to pre-eclamptic pregnancies, identified (intron 2, \( P = 0.005 \); SF-1 site, \( P = 0.016 \)). Two mutant haplotypes decreased the risk of developing pre-eclampsia (RR 0.16; CI 0.05–0.54; \( P < 0.001 \)). In contrast, intron 2 wild type predisposed to pre-eclampsia \( (P < 0.0015) \). No functional mineralocorticoid receptor mutant has been observed.

Conclusions. High aldosterone availability is associated with lower maternal blood pressure. In line with this observation, gain-of-function variants of the \( CYP11B2 \) reduce the risk of developing pre-eclampsia. Mutants of the mineralocorticoid receptor cannot explain the frequent syndrome of pre-eclampsia.

Keywords: aldosterone synthase; aldosterone; arterial hypertension; pre-eclampsia; pregnancy outcome

Introduction

Pre-eclampsia (PE), as defined by maternal arterial hypertension and proteinuria in pregnancy beyond the 20th week of gestation, causes severe morbidity often in pre-viously healthy individuals [1]. In addition to the immediate effects of the disease that may include cerebral accidents, liver disease and renal failure, PE is associated with remote cardiovascular and metabolic disease [2,3]. Thus, the ability to predict PE, especially with tests based on pathophysiological mechanisms, would be an important step towards eventually preventing and managing the disease.

Extracellular and plasma volume expansion of normal pregnancy is linked to changes in renal haemodynamics and of circulating levels of several steroid hormones including markedly increased circulating aldosterone concentrations [4–10]. Following production by the enzyme aldosterone synthase (\( CYP11B2 \)), aldosterone acts in target tissues via the mineralocorticoid receptor (MR).

Catalyzed by \( CYP11B2 \), aldosterone is synthesized from deoxycorticosterone in a three-step reaction with 18-methyl oxidation being rate limiting. In a group of pre-eclamptic women, we recently observed a reduced \( CYP11B2 \) activity characterized by a compromised 18-methyl oxidation step due to a loss-of-function mutation of \( CYP11B2 \) [11].

In non-pregnant subjects, essential hypertension with a high aldosterone-to-renin ratio, suggesting a high activity of \( CYP11B2 \), has previously been attributed to gain-of-function mutants representing biallelic gene polymorphisms of the \( CYP11B2 \). This is a -344C/T exchange located in the putative steroidogenic binding factor-1 (SF-1) within the 5′ regulatory region and an intron 2 conversion [Int2 (c)] resulting from the replacement of this region with that of the neighbouring gene encoding \( CYP11B1 \) [12].

With respect to our initial finding of an increased prevalence of loss-of-function mutants of \( CYP11B2 \) in PE, such gain-of-function alleles might protect from developing PE. Several functional relevant mutants of the aldosterone effector, the MR, are known to lead either to a loss or to a gain of function, which could as such further affect the aldosterone response [13].

We hypothesized that in pregnancy, gain-of-function mutants of the \( CYP11B2 \) gene in the absence of loss-of-function polymorphisms of the MR sustain a lower blood pressure and reduce the risk of developing PE.
Table 1. Characteristics of the pregnant women studied

<table>
<thead>
<tr>
<th>Type of pregnancy</th>
<th>Normotensive</th>
<th>Pre-eclampsia</th>
<th>P &lt;</th>
</tr>
</thead>
<tbody>
<tr>
<td>N</td>
<td>44</td>
<td>48</td>
<td></td>
</tr>
<tr>
<td>Age (years)</td>
<td>33.2 ± 1.3</td>
<td>31.2 ± 0.7</td>
<td></td>
</tr>
<tr>
<td>BMI</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>At delivery (kg/m²)</td>
<td>27.4 ± 0.9</td>
<td>29.0 ± 0.9</td>
<td></td>
</tr>
<tr>
<td>Blood pressure</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Systolic (mmHg)</td>
<td>117 ± 3</td>
<td>159 ± 16</td>
<td></td>
</tr>
<tr>
<td>Diastolic (mmHg)</td>
<td>75 ± 2</td>
<td>97 ± 2</td>
<td></td>
</tr>
<tr>
<td>Fetal birth weight (g)</td>
<td>3268 ± 98</td>
<td>1774 ± 179</td>
<td></td>
</tr>
<tr>
<td>Birth weight percentiles corrected for gestation</td>
<td>42.3 ± 4.8</td>
<td>26.2 ± 4.1</td>
<td></td>
</tr>
<tr>
<td>Low birth weight &lt;2.5 kg (%)</td>
<td>7</td>
<td>79</td>
<td></td>
</tr>
<tr>
<td>SGA &lt;10th birth weight percentile (%)</td>
<td>21</td>
<td>42</td>
<td></td>
</tr>
<tr>
<td>Gestation at time of sampling (days)</td>
<td>221.9 ± 9.1</td>
<td>224.7 ± 4.9</td>
<td></td>
</tr>
<tr>
<td>Proteinuria (g/day)</td>
<td>n.d.</td>
<td>2.3 ± 0.5</td>
<td></td>
</tr>
<tr>
<td>Time to delivery (day)</td>
<td>44 ± 6</td>
<td>6 ± 4</td>
<td></td>
</tr>
<tr>
<td>Total number of pregnancies (n/woman)</td>
<td>1.8 ± 0.2</td>
<td>1.8 ± 0.2</td>
<td></td>
</tr>
<tr>
<td>Primiparous (%)</td>
<td>53</td>
<td>57</td>
<td></td>
</tr>
<tr>
<td>Multiparous (%)</td>
<td>47</td>
<td>43</td>
<td></td>
</tr>
<tr>
<td>History of pre-eclampsia (%)</td>
<td>0</td>
<td>22</td>
<td></td>
</tr>
<tr>
<td>History of HELLP syndrome (%)</td>
<td>0</td>
<td>4</td>
<td></td>
</tr>
<tr>
<td>Fetal loss (n/woman)</td>
<td>0</td>
<td>0.4 ± 0.1</td>
<td></td>
</tr>
<tr>
<td>Premature birth (n/woman)</td>
<td>0</td>
<td>1.1 ± 0.1</td>
<td></td>
</tr>
<tr>
<td>Normal birth weight (n/woman)</td>
<td>1.8 ± 0.2</td>
<td>0.4 ± 0.1</td>
<td></td>
</tr>
</tbody>
</table>

n.d. = not detectable on either dipstick urinalysis or quantitative assessment of proteinuria, n.s. = not significant, SGA = small for gestational age.

Material and methods

Patients

Ninety-two pregnant women, 44 with normal pregnancies and 48 with PE, and 185 normotensive control subjects (115 males and 70 females) were investigated. Approval by the appropriate hospital ethics review committee and informed consent were obtained from all individuals. Basic demographic data, including blood pressure, information on concurrent diseases and the present medication, were obtained. Women with pre-existing arterial hypertension, diabetes mellitus or renal disease were excluded from the study. Patients were included in the PE group if blood pressure was ≥140/90 mmHg and proteinuria > 0.3 g/day in the absence of urinary tract infection beyond the 20th week of gestation (Table 1) [14]. Healthy pregnant women were selected as case controls according to the gestational age at initial presentation of pre-eclamptic women. Both groups of women were followed to delivery.

Collection of urine and blood samples and isolation of genomic DNA

Twenty-four-hour urine samples were collected from all women with PE at initial presentation. These samples were matched with those from women with a normal pregnancy at the same gestational age (± 2 weeks); however, analysis was postponed until delivery in order to establish the presence of an uncomplicated pregnancy. Ten millilitres of blood samples were drawn into tubes containing EDTA, mixed and stored at 4°C. Genomic DNA was extracted from peripheral blood leukocytes following the protocol supplied by the nucleon BACC3 kit (Bioscience; Piscataway, USA). Purified DNA was stored at 4°C in ddH₂O.

(A) hMR

DNA-binding domain

Steroid-binding domain

(B) CYP11B2

Screening of genomic DNA for the gain-of-function mutants of CYP11B2 [SF-1–344CT and Int2 (w/c)g] and functionally relevant mutants of the MR (Figure 1)

Polymerase chain reaction (PCR) for SF-1 and Int2 polymorphisms of CYP11B2 was performed using the primers and conditions reported by Davies et al. [16]. The region of DNA containing the HaeIII/SF-1 polymorphism was amplified by PCR by use of conditions similar to those previously described. A PCR product of 228 bp was amplified. This was digested with HaeIII and subjected to electrophoresis in 3% MetaPhor agarose. The 228 bp ampiclon contains two HaeIII restriction enzyme sites (GG GC). The presence of a C-T transition at position -344 (GG CT) removes one of these sites. After digestion, individuals homozygous for the transition (TT) produce two bands of 175 and 53 bp, individuals homozygous for the wild-type (CC) three bands of 104, 71 and 53 bp, and heterozygous individuals (TC) four bands.

The Int2 conversion was genotyped by using two separate PCR reactions, one that amplifies the normal gene (WT) and one that amplifies the conversion. The size of the amplicon in each reaction is ∼418 bp.

The functionally important domains of the MR for DNA and agonist binding, exons 3 to 9, respectively, were amplified by using the following primers: forward 5'-TGT GCA GCA TGT TAA ATG TGC-3', 5'-TTC AAG GCA GGA AAG TAG-3', 5'-TGG CCT CAT TAT CAT TGG TTT C-3', 5'-TTC CTG GAA TGT ACA ATG C-3', 5'-CAG TCT GGG TTT GAT AAT G-3', 5'-TTC GGA CTT GAA CAT TAT CTC TG-3' or 5'-AAA TGC AGA AGG CAG AGG GAT C-3', respectively. As reverse primers served: 5'-AAC ACC CAC ATG AAC ATT TAC C-3', 5'-TGC TGT TGC ATT ACC TAT TAA C-3', 5'-TGC AGC CTG TGA AAG GAG AG-3', 5'-AAA TCA TAA CCG ATC ACT CTG C-3', 5'-TAG AAA CAG...
haplotypes [SF-1 T/T and Int2 (c)/(c)]. Significance was assigned at
lotype of both genes [SF-1–344C/T and Int2 (w)/(c)] and two mutated
follows: no mutated haplotype [SF-1 C/C, Int2 (w)/(w)], one mutated hap-
Each allele contributing, for each gene, and by grouping haplotypes as
Comparisons for the number of mutated haplotypes were performed for
Fisher’s exact test were used, taking the appearance of PE as threshold.
ized calculations of relative risk (RR) and confidence intervals (CI) were
pressure as related to urinary tetrahydro-aldosterone/creatinine). Standard-
were not normally distributed. Linear regression analysis was performed
metric analysis and the Wilcoxon or Kruskal–Wallace test for variables that

term. Four
10 (SPSS Inc., Chicago, IL, USA) or GraphPad Prism version 4.01 for
0.05. All statistical analyses were performed using either SYSTAT version

Statistical analysis
All data are presented as means ± SEM. To test for statistically significant
differences, Student’s t-test was used for continuous variables by nonpara-
metric analysis and the Wilcoxon or Kruskal–Wallace test for variables that
were not normally distributed. Linear regression analysis was performed
for the continuous parameters (birth weight and systolic/diastolic blood
pressure [mm Hg]). Correlation of tetrahydro-aldosterone secretion with
blood pressure was performed using either SYSTAT version 10 (SPSS Inc.,
Chicago, IL, USA) or GraphPad Prism version 4.01 for Windows (GraphPad Software,
San Diego, CA, USA).

Results
Patients characteristics
Basic demographic, clinical and biochemical data of the 92 pregnant women,
dichotomized as to the occurrence of PE, are reported in Table 1. There was no difference in age
and body mass index in the two groups of women. The clinical
status was confirmed to be uneventful within the current
and all former pregnancies in the 44 women included as
normally pregnant. PE was diagnosed in 48 women by the
presence of proteinuria and hypertension as defined above.
The birth weight of children from pre-eclamptic pregnan-
cies was 40% lower than that of children from normal
pregnancies at similar gestational weeks (Table 1). Com-
parable numbers of women with either normal pregnancy
or PE were primiparous (53 versus 57%). Twenty percent
of multiparous women experienced a first pre-eclamptic
episode, and 22% were exposed to recurrent PE. Women
with a PE in the current pregnancy had former pregnan-
cies complicated by fetal losses, premature births and low
birth weight of their children (Table 1). Antihypertensive
treatment in PE consisted of labetalol, α-methyldopa and
nifedipine.

Fig. 2. Correlation between blood pressure and urinary tetrahydro-
aldosterone. Tetrahydro-aldosterone was measured by GC-MS in 24-
hr urine of pregnant women experiencing either a normal (n = 44)
or a pre-eclamptic (n = 48) pregnancy. Systolic and diastolic blood
pressure is depicted as circles and triangles, respectively, with values
from normotensive and pre-eclamptic pregnant women closed and open,
respectively.

Correlation of tetrahydro-aldosterone secretion with
clinical patterns
Urinary tetrahydro-aldosterone excretion was significantly
reduced in pre-eclamptic women when compared to nor-
mal pregnancies (13.4 ± 1.8 µg/mmol versus 38.0 ±
9.8 µg/mmol creatinine) (Figure 2). Lower blood pressure
correlated with a higher urinary tetrahydro-aldosterone ex-
cretion (systolic r = 0.99, P < 0.04; diastolic r = 0.99,
P < 0.04) (Figure 2). High urinary tetrahydro-aldosterone
excretion was found in the presence of higher birth weights
in normal pregnancies as compared to the lower tetrahydro-
aldosterone excretion and birth weights in PE (38.0 ± 9.8
versus 13.4 ± 1.8 µg/mmol creatinine; P < 0.05 and 3286 ±
98 g versus 1774 ± 179 g; P < 0.05).

Genotype and haplotype analysis of MR and CYP11B2
gene polymorphisms (Figure 1)
A PCR-based screening for mutations in the DNA and
steroid-binding domains of the MR was performed, and
no amino acid-changing mutation leading to structural or
functional changes was identified.

The CYP11B2 genotype distribution within the popula-
tion of the pregnant women and the controls studied were
in Hardy–Weinberg equilibrium. Normotensive pregnan-
cies more likely took place in the presence of homozy-
gosity for Int2 (c)/(c) (P = 0.0046), but less frequently
with Int2 (w)/(w) (P = 0.015). Homozygosity for SF-1
T/T was associated with reduced occurrence of hyperten-
sive pregnancies (P = 0.016, Figure 3), yet heterozygosity
did not protect from PE (P = 0.012). Combining alleles
associated with high aldosterone-to-renin ratios indicated
an increased frequency of these alleles in normotensive

| TGC CAG AAT G-3 |
|-----------------
| 5′-TCA AGG AAC CAA GGA G-3′ |
| 5′-CTT AAG GCA AGG TTC TTT G-3′ |
| 5′-TTT TGG G-3′ |

| 175 |
| 150 |
| 125 |
| 100 |
| 75 |
| 50 |
| 25 |
| 0 |

| 0 |
| 10 |
| 20 |
| 30 |
| 40 |
| 50 |

| systolic normal |
| systolic preeclamptic |
| diastolic normal |
| diastolic preeclamptic |

40% lower
when compared to pre-eclamptic pregnancies (2.6 ± 0.3 versus 1.9 ± 0.2 alleles; \( P = 0.023 \)). So far, these results point towards a relationship between pregnancy outcome and the number of mutant haplotypes. The presence of two, yet not one, mutant haplotypes for each gene locus was highly significantly linked to uneventful pregnancies \( (P < 0.001, \text{Figure 4}) \). The relative risk of developing PE was not reduced with one mutant haplotype, but with two (compared to no and one mutant haplotype RR 0.16; CI 0.05–0.54).

**Discussion**

This study shows an association of maternal blood pressure and fetal birth weight with aldosterone availability. As proposed, in normal pregnancies a high aldosterone production, determined via the measurement of urinary tetrahydro-aldosterone excretion, the most sensitive marker of circadian aldosterone production, was related to low maternal blood pressure levels [19]. Confirming our hypothesis, the number of high aldosterone-to-renin ratio alleles was increased in normal pregnancies. The absence of a high aldosterone-to-renin ratio allele of intron 2 was associated with an augmented frequency of PE. Furthermore, the presence of both mutants significantly favoured normal pregnancies.

The clarification of mechanisms of PE and hypertension in pregnancy is of practical relevance for multiple reasons with preventing the disease being the most important one [20–25]. If high aldosterone levels and the increased circulating plasma volume found in normal pregnancies represent a physiological adaptation crucial to support an adequate circulating fluid volume in pregnancy, two observations should be possible: first, a phenotypical adaptation in response to the aldosterone available, and second, mechanisms within the mineralocorticoid system either leading to or mimicking high aldosterone availability should prove beneficial to avoid PE [4–9,26].

If the primary impact of sufficient aldosterone availability would be to benefit pregnancy by avoiding PE, a genetic predisposition to produce high quantities of aldosterone or to appropriately respond to a given aldosterone concentration via the MR would be reasonable. Such a predisposition has been recognized by two gene loci of Mendelian forms of the CYP11B2 gene, the promoter SF-1 single nucleotide polymorphism -344C/T and a conversion in intron 2, which...
are variably linked to arterial mineralocorticoid hypertension with increased aldosterone-to-renin ratios in the non-pregnant population [12]. With these findings we stay in line with our initial finding of a compromised aldosterone synthesis being associated with PE [11].

In our study, the most pronounced differences between normal pregnancies and PE could be observed, if either an increased number of high ARR alleles or homozygosity for the mutant form were present. The RR to develop PE was reduced by >80%, if women were homozygous for both mutant alleles. Thus, combinations of activating polymorphisms seem to add to an uneventful pregnancy outcome, most likely due to an amplified aldosterone production as has been demonstrated during postural changes, when the maximal aldosterone response was found in subjects with two mutant haplotypes [12]. This observation is in line with our results and supports the view that variations in CYP11B2 may well lead to an intermediate phenotype of aldosterone excess supporting arterial hypertension in non-pregnant subjects and normotension during pregnancy due to an improved utero-placental perfusion.

One of these polymorphisms, the SF-1–344C/T conversion, had been studied in a pre-eclamptic population from Turkey with a lack of association [27]. A direct comparison of these data with our current findings is limited by the missing information on primiparity, which was high in our study group, and the ethnic divergence between the study groups. Also, we investigated a group of pregnant women as controls without any abnormalities in their for-mer pregnancy history, which may have resulted in a bias towards genotypes of the CYP11B2 with a high aldosterone production. Furthermore, PE appears to be a multifactorial disease with each factor potentially contributing to a different extent, even more in a genetically divergent context. Interestingly, the low expression of high aldosterone-to-renin ratio variants in PE resembled the pattern seen in the healthy, normotensive, non-pregnant subjects and normotension during pregnancy due to an improved utero-placental perfusion.

This study supports the assumption that an uncompro-mised aldosterone production is importantly contributing to an uneventful pregnancy outcome. If aldosterone production is diminished due to a polymorphism within the CYP11B2 gene, the reduced aldosterone availability in the presence of increasing concentrations of the partial aldosterone antagonist progesterone throughout gestation might reduce the compensatory potential for shifts in circulating volume warranting the genotypes discussed here with a high capacity to produce aldosterone [11]. Our current observations well support the concept of a genetic susceptibility of PE as has been repetitively demonstrated in family and twin studies, though stable environmental factors must also be considered in recurrent PE and have not been addressed here [28–30].

The exact mechanisms determining the magnitude of aldosterone secretion in the presence of the CYP11B2 variants have not yet been elucidated [31]. Adrenal release of aldosterone is reduced in hypoxia as shown by Raff and Kohandarvish [32] in vitro and by Lawrence et al. [32,33]. Thus, one might speculate that during established PE factors such as adrenal hypoxemia further contribute to a diminished aldosterone release.

Earlier, we ruled out aldosterone levels to be decreased due to an excess of alternate mineralocorticoid-like steroid hormones [11]. Yet, aldosterone concentrations might be affected by an altered activity of the MR by either gain- or loss-of-function mutations within functionally relevant regions of the MR gene [13,17,34]. Our present data do support the assumption that structural or functional mutations of the MR are extremely rare and will not have to be judged as a contributing major factor to explain the frequently occurring PE, a finding supporting observations in very small subsets of pre-eclamptic patients by other groups [35,36].

The present investigation revealed a close relationship between aldosterone availability, gestational maternal blood pressure and fetal birth weight. High aldosterone-to-renin ratio variants of the CYP11B2, which provide for an increased aldosterone availability, were associated with a lower frequency of pre-eclamptic pregnancies. Most strikingly, double homozygosity for both high aldosterone-to-renin ratio variants, predicted a significantly reduced risk of developing PE. Though the exact regulatory events for CYP11B2 in pregnancy still have to be elucidated more in detail, this observation is of practical importance since it might help to define subgroups of patients with an altered chance to develop pregnancies complicated by PE. Thus, women could be given a tailored follow-up throughout their pregnancies according to their CYP11B2 genotype in addition to further risk predictors. These results should be verified in a larger cohort also including women with an intermediate phenotype.

Acknowledgements. This work has been supported in part by a grant for scientific research from the Swiss National Science Foundation (No. 3200B0-113902/1 to M.M.) and a Marie-Heim Vögelein grant (PMPDB-106128 to G.E.).

Conflict of interest statement. None to be declared.

(See related article by M. D. Lindheimer and P. August. Aldosterone, maternal volume status and healthy pregnancies: a cycle of differing views. Nephrol Dial Transplant 2009; 24: 1712–1714.)

References
Mineralocorticoid system and pregnancy


Received for publication: 26.9.08; Accepted in revised form: 19.12.08