Treatment for Severe Pregnancy Induced Hypertension and Preterm Labor with Concomitant Administration of Estradiol and Progesterone under Monitoring Aminopeptidases: A Case Report

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Abstract

The knowledge for molecular mechanisms of preterm labor and preeclampsia is still premature. In the previous review, the author proposed that sex steroid treatment should be conserved for prospective clinical study in both preterm labor and preeclampsia. Unfortunately, little attention has been paid so far on the review article. Here, the authors present two cases of severe, complicated preeclampsia associated with preterm labor.

Keywords: Estradiol-progesterone therapy; Placental aminopeptidases (APs); Angiotensin II (A-II); Vasopressin (AVP); Oxytocin (OT)

Introduction

Underlying mechanism of preeclampsia and onset of labor/preterm labor, despite progress in its research on molecular basis, need to be elucidated until effective treatments are established. Historically, peptide hormones such as angiotensin II (A-II), vasopressin (AVP) and oxytocin (OT), have been investigated as essential hormones involved in preeclampsia and onset of labor/preterm labor but recently scientists' effort towards these hormones has somehow declined these days. It is important to look at these hormones in contrast to their degrading enzymes.

The authors cloned placental leucine aminopeptidase (P-LAP), the enzyme that effectively degrades AVP and OT. Serum concentrations of P-LAP increase in pregnant women as pregnancy advances [1,2]. The authors have shown existence of aminopeptidase A (APA) in human placenta as an identical substance to the A-II degrading enzyme. Later, Max Cooper cloned APA as a B lymphocyte differentiation antigen. The authors reported the fact that APA increases in maternal sera in accordance with gestational term in clinical settings [1,3]. These peptide hormones are known essential for the development of fetus and thus increase with fetal development. Concentration of these hormones in the feto-placental unit is higher than those of maternal circulation. Since the hormones are biologically active, the leak of these hormones into the maternal circulation is strictly regulated by degradation activity of placental aminopeptidases (APs) in order to maintain maternal blood pressure (BP) normal and to keep uterine tonus under control.

In 1940, Smith & Smith reported a favorable outcome of the treatment by using estradiol benzoate (E2) accompanied with P in severe preeclampsia. In 1969, taking a hint from Smith's case reports, the author had tried his own, first, sex steroids treatment in severe preeclampsia on a dose increase by gestational week method, about measuring P-LAP levels [5,6]. Owing to alleviation of clinical symptoms, blood pressure and natriuresis, prolongation of gestation at least for 3 weeks was witnessed, and the treatment was proved successful. In view of neither safe nor effective medication being practiced at present, the proposed sex steroid treatment in this report seems worthwhile for the prospective controlled studies for the treatment of both preeclampsia and preterm labor [6].

The knowledge for molecular mechanisms of preterm labor and preeclampsia is still premature. Moreover, clinical effectiveness of P and 17α-hydroxyprogesterone capronate (17-P) in preterm labor remains under debate [7]. In the previous review, the author proposed that sex steroid treatment should be conserved for prospective clinical study in both preterm labor and preeclampsia. Unfortunately, little attention has been paid so far on the review article. Here, the authors present two cases of severe, complicated preeclampsia associated with preterm labor.

Case Presentation

Case 1

The patient, 39-year-old woman (gravid-1, para-0) was conceived...
by in vitro fertilization frozen embryo transfer (IVF-ET) in 2003. Her pregnancy was a di-chorionic twin. Until this pregnancy she was without any medical complication. Her pregnancy had been uneventful until week 22, however at week 26, she was hospitalized due to severe preeclampsia (BP 190/108, proteinuria and edema) complicated with premature labor. She was treated by intravenous drip injection of beta-2 agonist (ritodorine), and intramuscular injection of terbutaline (2 mg twice/day), in addition, with indomethacin (50 mg) to control uterine contraction.

Both nifedipine (40 mg/day) and metoprolol (80 mg/day) were used concomitantly as antihypertensives. However, these agents will pass through the placenta easily thereby can exert negative effects on the fetus, against its well-being. The estimated fetal weight of twin A was 900 g and twin B was 750 g, respectively at the hospital admission. At week 27+3-day clinical symptoms had not improved and the non-stressed test (NST) showed the decreased variability and late decelerations observed. At this stage, she was transferred to our hospital. On admission, the estimated fetal weight of twin A was 1050 g without ominous sign, on the other hand, twin B was 850 g and this fetus showed the reversed umbilical blood flow by Pulse Doppler Assessment, suggesting ominous sign.

Firstly, we stopped all medicines used in the previous hospital except ritodorine. We explained her and her family about the anticipated poor prognosis of her fetuses (especially twin B), even if interruption of her pregnancy was tried by cesarean section. Therefore, on the next day (week 34+6 day) we performed elective cesarean section.

A female infant weighing 1684 g was delivered (Apgar score: 9/10). Twin B was a 400 g female stillbirth fetus which was macerated without abnormalities and two chorionic sacs were confirmed. Weight of the placenta was 470 g. None of pathological abnormality in placenta and cords was found. In puerperium, patient’s BP showed 130-150/70-90 mmHg. Proteinuria persisted for nine days. The patient was discharged eighteen days after delivery in noneventful condition. Growth of the newborn had been normal and was discharged from NICU.

**Case 2**

In this case, the patient is a 26-year-old (gravid -1, para-1), had severe preeclampsia with underlying complication of non-insulin dependent diabetes mellitus (NIDDM). Since she was diagnosed...
Figure 2: The changes of P-LAP activities in case 1.

Figure 3: The effects of estradiol and progesterone treatment in case 2.

Figure 4: The changes of P-LAP activities in case 2 were shown in black bold line. Before starting sex steroid treatment, P-LAP activity levels had been lower than of normal pregnancy. However, after the treatment initiation, P-LAP activity levels started to increase onto the range of normal pregnancy and kept increased with the treatment, though some fluctuations observed through advancement of gestation.
for NIDDM at the age of twenty, she had been treated with insulin injections until her pregnancy. In 2003, the patient was admitted to our hospital for mild preeclampsia (BP 155/80 mgHg, proteinuria: 3 g/day, and edema) complicated by preterm labor (uterine contraction, advanced shortening of uterine cervix: the effacement of uterine cervix reached to about 70% and dilation of uterine cervix: around 3 cm). The protruded bag of water from external orifice of the uterus was clearly observed. Ritodrine was administered for the suppression of uterine contraction although the drug is contraindicated to pregnancy with NIDDM. This treatment could induce the increase of blood glucose; therefore, an incremental dose of insulin was required. We proposed the patient our sex steroid treatment accompanied by ritodrine in order to sustain her pregnancy. The patient and her family accepted our treatment approach.

With sex steroid treatment, her symptoms of preeclampsia and preterm labor started to alleviated. Upon observation of symptoms, we were able to decrease the dose of ritodrine gradually and terminate administration at week 32+4 days. None of antihypertensive agents nor magnesium sulfate were administered (Figure 3).

Figure 4 shows the changes of P-LAP activity. Before starting sex steroid treatment, P-LAP activity levels had been lower than of normal pregnancy. However, after the treatment, P-LAP activities started to increase to the range of normal pregnancy and kept increased with the treatment, although fluctuations were observed throughout gestation. The condition of uterine cervix, on the other hand, did not change. At week 37-day premature rupture of membranes occurred and her body temperature (38.2°C) showed the sign of infection. On the same day, we performed cesarean section. Male infant weighing 2388 g (Apgar 6/9) was delivered alive. Her puerperium had been in good condition. The newborn was discharged in good condition.

Discussion

Serum estradiol (E2) and progesterone (P) in normal pregnant women increase with advancing of gestation and reach to about a hundred-fold or more at the point of term compared to preconception. P is the steroid hormone responsible for carrying over pregnancy by suppressing uterine contraction. Also, the role of E2 is to sustain pregnancy until reaching the timing of initiating labor, but it also has the ability to promote synthesis of P. E2 increases in the uterine blood which ensures adequate supply of oxygen and nutrients to the fetus. In this regard, both E2 and P seem to have indispensable roles in carrying over normal term pregnancy [6].

The levels of P-LAP activities under the sex steroid treatment were different between the two cases but they have demonstrated effectiveness and safety of the sex steroid treatment in sustaining a pregnancy. Alleviation of symptoms such as hypertension and uterine contraction were seen as the P-LAP and APA activities increased in maternal sera through advancing of gestation. Changes of P-LAP and APA activities in maternal sera accurately reflected improvement of clinical symptoms [5,6).

As mentioned above, we have shown the fact that both maternal P-LAP and APA activity increase more than normal pregnancy levels in mild preeclampsia, suggesting protective action against the leak of these hormones into the maternal circulation [2,4]. In the exacerbation of preeclampsia from mild to severe form, P-LAP activities in maternal sera had abruptly fallen to significantly lower levels than normal pregnancy, indicating the breakdown of the balance between increasing fetal peptide and production of placental Apo [2,4].

Conclusion

Therefore, the difference of the P-LAP levels observed between the two is specific changes of fetal peptide hormone levels before and after the sex steroid treatment. Prognosis of preeclampsia or preterm labor is dependent on P-LAP activities in maternal sera that the pattern what it shows to increase with advancing gestation [1,5]. Recently, it was shown that both APA and P-LAP could potentially be safe and effective drugs for medical treatment for preeclampsia and preterm labor [1-12]. Considering unavailability of safe - to - mothers and babies - and effective medication at present, the sex steroid treatment, as proposed in this report, seems worth proceed into comparative clinical studies on these abnormal conditions [6] so that the existing sex steroid hormone preparations could be indicated for preeclampsia and preterm labor for extensive clinical use. Effectiveness of P and 17-hydroxyprogesterone caproate (17-P) in preterm labor remains under debate [7]. These mono-progestogen approaches seem devoid of taking account dynamic aspects of estradiol and P in the placenta. A major defect is the exclusion of estradiol in their protocol [6]. Since the tragedy of diethylstilbestrol [11-13], administration of sex steroid hormones to pregnant women has been averted for a long time until emergence of clinical use in assisted reproductive technology in 1985 [14]. We should not ignore that E2 is an essential element also for the treatment with assisted reproductive technology.

In the near future, the authors expect a pharmaceutical development for both recombinant APA and P-LAP indicated for preeclampsia and preterm labor [1]. This seems the ultimate goal for the effective and safe solution for these abnormalities.

References
