Differences in Expression of Peroxisome Proliferator-activated Receptor- γ in Early-onset Preeclampsia and Late-onset Preeclampsia

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Research note

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Abstract

Objective: Preeclampsia is a leading cause of maternal and fetal morbidity and mortality. Peroxisome Proliferator-activated Receptor- γ (PPAR- γ) has been reported to be implicated in key functions in cell that were also important to ensure normal pregnancy development. In this study, we conducted primary culture of trophoblastic cells from normal pregnant women in the third trimester ($n=3$), and from patients with EOPE ($n=3$), and LOPE ($n=3$). Expression of PPAR- γ was determined by Western Blot. Data are the means $\pm$ S.E. from three separate experiments. Statistical analysis was performed by SPSS software, version 20.0, a $p < 0.05$ was considered significant (one-way ANOVA and Bonferroni's test) versus normal serum pregnancy.

Results: Serum from normal pregnant women and those with EOPE did not induce any difference in the expression of PPAR- γ ($p > 0.05$). In contrast, expression of PPAR- γ was increased in those cells with serum from LOPE ($p < 0.001$). Therefore, we conclude that hypothetically PPAR- γ might play role in the pathophysiology of LOPE but not in EOPE. Other possibility is the activity of PPAR- γ in EOPE is inversely correlated with the expression, therefore the high enzymatic activity of PPAR- γ is tightly regulated by attenuating its expression.

Introduction

Pre-eclampsia is defined as a hypertensive disorder related to pregnancy, and included among the most common medical complications of pregnancy. It carries burden around the world, causing a high maternal and perinatal mortality and morbidity.(1–2) The mechanisms involve in the genesis of this syndrome is not well understood, although several factors has been extensively well studied, such as defective placentation(3–6), ischemia of the uteroplacental circulation (5,7–9), endothelial cell dysfunction(10–11), and excessive inflammatory reactions to the invading trophoblast(12–13).

Classification of preeclampsia may be divided into two distinct entities, early- and late-onset. The early-onset preeclampsia (EOPE) occurs at < 34 weeks of gestational age, and the late onset preeclampsia (LOPE), is those preeclampsia of which manifestation present at ≥ 34 weeks of pregnancy.(14–16) Both EOPE and LOPE have been found to be associated with different biochemical markers, genetic and environmental risk factors, prognosis, heritability, and clinical features. Evidences showed that several of the markers rising prior to pregnancies, and may persisted or disappeared before clinical manifestation of preeclampsia.(17–19)

Peroxisome Proliferator-activated Receptor- γ (PPAR- γ) is a member of the nuclear receptor superfamily and has been reported to be implicated in key functions in the cells including cell proliferation and differentiation, inflammation and oxidation, glucose and lipid metabolism. These functions were also important to ensure normal pregnancy development.(20–21) In normal pregnancy, activators of PPAR- γ was increased along with increasing gestational age. A reduction in the placental expression of PPAR- γ activators has been demonstrated in some preeclamptic patient.(22) In contrast, Holdsworth-Carson et al.

(23) examined placental expression of PPAR- γ and demonstrated that placentas from women with pre-eclampsia did not demonstrate any differences in mRNA or protein expression of PPAR- γ compared with healthy controls. Here we attempted to study the possible involvement of PPAR- γ in EOPE, LOPE compared to that of normal pregnancy.

Methods

Cells

Cultured of primary trophoblastic cells obtained from term normal pregnancy was courtesy from Oncology and Stem Cell Working Group, Faculty of Medicine, Universitas Padjadjaran, Bandung, Indonesia. Cells were maintained in medium amniomax basal media (Gibco USA) added with amniomax supplement (Gibco, USA), at 37 °C/5% CO₂.

Maternal serum

Serum was courtesy from Oncology and Stem Cell Working Group, Faculty of Medicine, Universitas Padjadjaran, Bandung, Indonesia. Serum were obtained from normal pregnant women in the third trimester (n = 3), and from patients with EOPE (n = 3), and LOPE (n = 3). Pregnancies were considered normal when patients did not have medical and obstetric complications of pregnancy and delivered aterm neonate (≥ 37 weeks) who was appropriate for gestational age. Preeclampsia was defined as hypertension (systolic blood pressure ≥ 140 mmHg or diastolic blood pressure ≥ 90 mmHg on at least two occasions, 4 h to 1 week apart) and proteinuria (300 mg in a 24-h urine collection or one dipstick measurement of $> 2+$)(24–25). Patients with chronic hypertension, diabetes mellitus, systemic diseases such as antiphospholipid antibody syndrome, thrombophilia or transient blood pressure elevations were excluded from this study.

Exposure Cells to Serum

Cells were plated at 24-well and incubated for 24 h at 37 °C with 5% CO₂. The cells were followed by treatment with serum from the patients at 15% final concentration(25–26) in amniomax (Gibco USA) in the presence of amniomax supplement (Gibco, USA) for 24 h. The three groups were as follow: normal serum (normal), EOPE, and LOPE.

Whole Cell Lysate Preparation and Western Blotting

Cells were washed with ice-cold 1x PBS and then lysed in RIPA buffer in the presence of protease inhibitors and phosphatase inhibitors. Lysates were then added with SDS sample buffer (50 mM Tris-HCl pH 8.0, 150 mM NaCl, 1 mM EDTA, 1% SDS), and were centrifuged at 17,500 X g for 20 min at room temperature. The resulting supernatants with equal amount of total protein was loaded in each lane. After transfer to PVDF, the membranes were blocked with 0,25% BSA in TBS-Tween for 30 minutes at room temperature. Primary and secondary antibody incubations at 4 °C overnight, and 90 minutes at room

temperature, respectively, with antibodies diluted in blocking buffer BSA 0,1%. Antibodies used included rabbit monoclonal antibody (mAb) against PPAR- γ (Cell Signaling Technology), mouse anti-actin mAb (Thermo Fisher Scientific), HRP-conjugated secondary antibodies against anti-rabbit, and anti-mouse was from Thermo Fisher Scientific and Santa Cruz, respectively. After treatment with ECL reagent (GE Healthcare), proteins in membranes were detected by C-Digit (Licor). Expression was quantified by densitometric scanning by Image-J followed by normalizing PPAR- γ expression to that of β -actin.

Statistical Analysis

Data are the means $\pm$ S.E. from three separate experiments. Statistical analysis was performed by SPSS software, version 20.0 (SPSS Inc., Chicago), a $p < 0.05$ was considered significant (one-way ANOVA and Bonferroni's test) versus serum normal pregnancy.

Results

We attempted to study the possible involvement of PPAR- γ in EOPE, LOPE compared to that of normal pregnancy. The result of incubation of primary trophoblastic cells with serum of normal pregnancy, serum of EOPE, and LOPE, analyzed by Western Blot shown as follow in Fig. 1:

Our result showed the expression of PPAR- γ in primary trophoblastic cells treated by serum normal pregnancy compared to that of EOPE showed no expression in Western Blot. Remarkably, treatment of LOPE showed marked expression of PPAR- γ compared to that of normal pregnancy.

Next, we confirmed the significance of our result. We performed quantification and showed the result in Fig. 2. The result was there was no difference in the expression of PPAR- γ in those cells treated by EOPE compared to normal pregnancy ($p < 0.01$), while the treatment of LOPE resulted in significant high expression of PPAR- γ compared to that of normal pregnancy ($p < 0.001$).

Discussion

In our experiment, we used a primary trophoblastic cells obtained from normal pregnancy that has been exposed to either serum of normal pregnancy, EOPE, or LOPE.(7,27)

Many of in vitro experiments were done as prelude for translational researches. Cell lines have limitations for preclampsia model due to the difficulties in interpretation to in vivo conditions. Many experiments, such as by Petroff et al(29) found experimental techniques to isolate human trophoblastic cells then cultured as primary trophoblastic cells as an approach for in vivo situation. Successful isolation of cytotrophoblastic cells has been proven to resemble human condition for several placental functions, such endocrinology and immunology functions of the placenta, placental differentiation, and apoptotic conditions.(26,28–30)

Previously, others have used cultured primary trophoblastic cells given serum of preeclamptic patients as models for preeclampsia condition. Pramartirta (26) found expression of TNF- α and caspase-3, and apoptotic index in preeclampsia serum-induced trophoblast cells were higher than normal and controls. In other studies, they found structural derangement of vessels resembling disruption in interaction of trophoblastic cells with endothelial cells. (25,31) To conclude, these evidences supported the method of serum preeclampsia treatment in vitro is suitable for preeclampsia model.

The mechanisms responsible for the genesis of the preeclamptic syndrome are poorly understood(1,7,32) although roles for abnormal placentation(3–6), uteroplacental ischemia(5,7–9), endothelial cell dysfunction(10–11), and exaggerated maternal inflammatory response to deported trophoblast(3–4,33) have been proposed.

Peroxisome Proliferator-activated Receptor- γ (PPAR- γ) is a member of the nuclear receptor superfamily. Included as transcription factors, they are members of the ligand-activated nuclear hormone receptor superfamily, and play major roles in diverse aspects of energy metabolism, inflammation, and development. Following ligand binding, PPARs form heterodimers with retinoid X receptors (RXRs), and bind to PPAR-response elements (PPREs) of target genes to activate transcription.(34–36)

Three subtypes of PPAR has been identified; PPAR α , β , and γ .(20–21) All three subtypes are expressed in placenta with different levels. PPAR- γ has been reported as crucial part of normal placental development as deletion of PPAR- γ in mice is 100% lethal, with no surviving embryos born. Homozygous PPAR- γ -deletion mice embryos die due to severe placental dysfunction.(37–38)

Disrupted expression of PPAR- γ were implicated in several vital placental functions including regulation of inflammatory reactions that resulted in severe gestational disorders. Effect of PPAR- γ in the regulation of pro-inflammatory mediators such as IL-6, IL-8, and TNF- α is important for normal pregnancy to ensue. (20,38) Other evidence proofed the involvement of PPAR- γ in regulation of trophoblasts functions. Antagonists PPAR- γ used in vitro of first trimester extravillous trophoblasts resulted in increase trophoblast invasion, while agonist inhibited it. These data support the role of PPAR- γ in the regulation of trophoblasts invasion to decidua.(21,38)

Previously studies found PPAR- γ was expressed in human placenta.(40–41) PPAR- γ mRNA was expressed in term pregnancy.(39) Previously, others have found the primary culture of isolated trophoblasts from first trimester pregnancy expressed PPAR- γ . This same research strengthen the possible involvement of PPAR- γ in normal pregnancy because they found PPAR- γ expression was preserved until third trimester.(42)

In our study we proved the possible involvement of PPAR- γ in preeclampsia. We used Western Blot technique to analyze expression of PPAR- γ in primary trophoblastic cells treated by serum of normal pregnancy, EOPE, and LOPE. Treatment of the primary trophoblastic cells with LOPE was shown to induce high expression of PPAR- γ . Whereas, treatment of trophoblastic cells by EOPE resulted in similar result as treatment by normal pregnancy, with no induction of PPAR- γ expression. Handschuh et al.(43) stated

PPAR- γ activity depend on the trophoblast subpopulation, gestational age, and types of stimulating ligands. Therefore, different content of serum in EOPE and LOPE has different effect to PPAR- γ expression..

In LOPE, maternal factors such as obesity, hyperlipidemia, diabetes, chronic hypertension contribute much for emergence of the disease.(44–47) In our experiment the high expression of PPAR- γ induced by treatment with serum of LOPE may indicate PPAR- γ expression inversely correlates with its activity. As Levistka et al.(48) showed agonist PPAR- γ rosiglitazone induced reduction in the receptor expression in the human choriocarcinoma derived cell line BeWo, an established model of syncytiotrophoblast formation in vitro, and primary trophoblast cells. In contrast, inhibition of PPAR- γ activity using T0070907 causing extreme enhancement of receptor expression. Thus further study is necessary to proof the PPAR- γ expression increasement in our study is affected by the protein activity.

Above evidence suggested the activity of PPAR- γ is modulated by negative feedback.(48). Similar with Levitska et al.(48) other study by Knabl et al.(49) also stated PPAR- γ as transcription factor plays important roles in fat and glucose metabolism, an in cell growth and differentiation, so that thight autoregulation is necessary to even out its activity. A high activity will induce lower expression, while low activity result in high receptor expression.

The mechanism of EOPE in pregnancy was according to the “two-stage theory of preeclampsia”. In this theory, in the first stage of disease development, defective trophoblastic invasion result in shallow placentation, and the impair remodeling of muscular layer in spiral arteries causing the spiral arteries unable to fully dilated to supply normal pregnancy development. Further mechanism was at the second stage resulting from failure to establish an adequate uteroplacental blood flow leads to relative hypoxia in trophoblastic tissue, thereby eliciting an oxidative stress response, thus releasing placental debris that in maternal circulation and emerged as symptomatic disease. As a consequence for abnormal trophoblastic invasion at early placental development will result in earlier disease manifestation (< 34 weeks), or EOPE.(44–47) Based on this theory, in EOPE abnormal trophoblast activity may be regulated by PPAR- γ , however high activity of this receptor result in diminish receptor expression.(48–49)

Conclusion

PPAR- γ might play role in the pathophysiology of LOPE but not in EOPE. Other possibility is the activity of PPAR- γ in EOPE is inversely correlated with the expression, therefore the high enzymatic activity of PPAR- γ is tightly regulated by attenuating its expression.

Limitations

This study is limited by a possible discrepancy among gene expression, protein expression, and protein activity. Gene activities describe transcription process at mRNA level, while protein expression is highly modified at translational process or post translation, as well as protein degradation rate. Protein activity

may be affected by chemical reaction, therefore may influence by presence of catalisator or inhibitor. Further research must confirm the mechanism of higher expression of PPAR- γ in our research at pre- or post-receptor level.(50)

Abbreviations

EOPE

Early-onset Preeclampsia

LOPE

Late-onset Preeclampsia

PPAR- γ

Peroxisome Proliferator-activated Receptor- γ

PPRE

PPAR-response elements

RXR

Retinoid X receptors

Declarations

Ethics Approval

The use of the samples for research purposes was approved by the ethical reviews boards of the Health Research Ethics Committee University of Padjadjaran Bandung Indonesia.

Consent to participate was not applicable because in this research used archived biological material.

Consent to publish

All authors read and approved the final manuscript.

Availability of data and materials

Availability of data and materials are available upon request to the corresponding author. The cells and serum used as archived biological materials are courtesy from Oncology and Stem Cell Working Group, Faculty of Medicine, Universitas Padjadjaran, Bandung, Indonesia.

Competing Interests

The author(s) declared no potential conflicts of interest with respect to the research, authorship, and/or publication of this article.

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Authors' Contributions

WP, KIM, HBA did the conception and design of the study, acquisition of data, analysis and interpretation of the data, drafting the manuscript and revising the manuscript critically for important intellectual content.

AFK , UAL did the conception and design of the study, acquisition of data, and analysis and interpretation of the data.

AAR drafted the manuscript and revising the manuscript critically for important intellectual content.

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Figures

Figure 1

Expression of PPAR-γ. Primary trophoblastic cells were treated with normal serum (normal), EOPE, and LOPE. PPAR-γ proteins were subjected to immunoblot analysis with a mAb that recognizes PPAR-γ as well as with a mAb to β-actin (loading control).

Figure 2

Quantification of Expression of PPAR- γ . Expression was quantified by densitometry scanning by Image-J followed by normalizing PPAR- γ expression to that of β -actin. Data are the means $\pm$ S.E. from three separate experiments. versus serum normal pregnancy. * $p < 0.01$ (one-way ANOVA and Bonferroni's test) versus normal pregnancy. N.S., non significant.