Role of peroxiredoxin III in the pathogenesis of pre-eclampsia as evidenced in mice

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As a member of peroxiredoxin (Prx) family, PrxIII has been demonstrated to play an important role in scavenging intracellular reactive oxygen species (ROS). Since PrxIII knockout mice exhibited oxidative stress in placenta resembling pathophysiologic changes in placenta of human pre-eclampsia, we measured blood pressure through the carotid artery and detected oxidative status by western blotting in pregnant mice. We did not notice hypertension in pregnant PrxIII knockout mice as compared with wild-type littermates, although endothelin-1 was overexpressed in PrxIII-deficient placenta. Our results indicate that PrxIII is not involved in pre-eclamptic development. Instead, PrxIII is an indispensable antioxidant in placenta where oxidative stress exists.

Introduction

As a unique organ that connects mother and fetus, the placenta provides the place for materno-fetal exchange of gases, nutrients and metabolic products. Since placenta is abundant in mitochondria, large amount of reactive oxygen species (ROS) is continuously produced during metabolic processes. During pregnancy, ROS production is gradually elevated with the increase of energy demand due to both fetal and maternal requirements. Although small amount of ROS is considered to be a second messenger, excessive ROS would cause oxidative injuries on DNA, lipids, and proteins. Organisms have developed multiple antioxidant defenses against oxidative damage. For example, placental

antioxidant activity was increased with gestation progression,¹ and higher activity of superoxide dismutase was observed in maternal erythrocytes.² More importantly, mitochondrion possesses a primary antioxidant chain that is composed of peroxiredoxin III (PrxIII), thioredoxin II and thioredoxin reductase.³

As a member of peroxiredoxin family, PrxIII was found to be responsible to oxidative stress in placenta.^{4,5} We have recently observed placental oxidative damage in PrxIII knockout (PrxIII^{-/-}) mice.⁶ Interestingly, our findings in PrxIII-deficient placenta were quite similar to the major pathophysiologic changes in placenta of pre-eclampsia (PE),⁷⁻⁹ a serious pregnancy complication in human beings. Since it is speculated that increased placental lipid peroxides and/or tumor necrosis factor alpha (TNFalpha) cause activation of leukocytes which serve as circulating mediators and contribute to endothelial dysfunction and subsequent hypertension,¹⁰ we have investigated whether placental oxidative damage causes hypertensive disorders in mice resembling PE.

Results

Blood pressure in pregnant mice. Since oxidative damage in PrxIII-deficient placenta of mice resembled pathological findings in pre-eclamptic placenta,⁶⁻⁹ we measured mean arterial blood pressure in pregnant mice trying to elucidate pathogenic role of PrxIII in pre-eclamptic development. As indicated in Table 1, there was no significant difference of blood pressure

Key words: peroxiredoxin, placenta, oxidative stress, hypertension, pre-eclampsia, mouse, reactive oxygen species

Abbreviations: ROS, reactive oxygen species; Prx, peroxiredoxin; PE, pre-eclampsia; TNFα, tumor necrosis factor alpha; ET-1, endothelin-1

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Table 1. Blood pressure in PrxIII^{+/+}, PrxIII^{+/-} and PrxIII^{-/-} mice before and after pregnancy

Blood Pressure	PrxIII ^{+/+}	PrxIII ^{+/-}	PrxIII ^{-/-}
Before Pregnancy	95 ± 2	95 ± 2	95 ± 3
After Pregnancy	95 ± 6	93 ± 5	97 ± 5

Blood pressure in mice was measured as described in “Materials and methods” section (n = 10 for PrxIII^{+/+}, PrxIII^{+/-} and PrxIII^{-/-} mice respectively). There was no significant difference of blood pressure among individual groups before or after pregnancy (p > 0.05).

Figure 1. Western blotting analysis for ET-1 expression in PrxIII^{+/+} (A) and PrxIII^{-/-} (B) mouse organs including (from 1 to 7) brain, heart, lung, liver, kidney, placenta and skeletal muscle. Signals for ET-1 protein were significantly enhanced in PrxIII knockout placenta and kidney as compared with wild-type organs.

before or after pregnancy among PrxIII^{+/+}, PrxIII^{+/-} and PrxIII^{-/-} mice.

Systemic oxidative status in pregnant mice. To further understand oxidative status in pregnant mice, we detected expression of endothelin-1 (ET-1) systemically. As shown in Figure 1, ET-1 protein was evidently enhanced in placentas derived from PrxIII^{-/-} mice, which was in accordance with our previous report.⁶ Nevertheless, ET-1 expression was not significantly elevated in other PrxIII-deficient organs except kidney as compared with wild-type samples.

Discussion

We recently reported that PrxIII-deficiency resulted in excessive lipid peroxidation and TNFα production in placentas of mice,⁶ which was quite similar to the pathological changes in pre-eclamptic placentas.⁷⁻⁹ We then measured blood pressure in pregnant mice and found serious oxidative stress in placentas of PrxIII knockout mice without subsequent dysfunction of peripheral endothelium and hypertensive disorder.

PE is one of the most important pregnancy complication characterized by hypertension, proteinuria, and edema. Although the precise mechanism for pre-eclamptic occurrence remains unclear, oxidative stress was considered to be

one of the main causative factors.^{13,14} Malperfusion of the placenta was implicated as a cause of oxidative stress in complications of human pregnancy,¹⁵ and subsequent endothelial activation and peripheral vascular contraction was considered to be the basic pathophysiology of pre-eclampsia.^{16,17} Both TNFα and ET-1 were reported to play important roles in linking placental oxidative damage and systemic endothelial dysfunction in PE.^{17,18} We detected high levels of TNFα and ET-1 in PrxIII-deficient placentas of mice, implying serious oxidative stress and poor perfusion in the organs. Nevertheless, we did not notice the increase of ET-1 in other organs except kidney and subsequent hypertension in pregnant PrxIII^{-/-} mice. Combining with our results and another report that PrxIII is significantly upregulated in pre-eclamptic placentas,⁵ we suppose that PrxIII is not involved in the pathogenesis of pre-eclamptic development. Instead, it is very likely that PrxIII is an indispensable ROS scavenger in oxidation-threatened placentas and contributes to the inhibition of PE progression.

Materials and Methods

Animals. Experimental mice were generated by mating C57BL/6 mice with PrxIII^{-/-} mice to generate PrxIII^{+/-} offspring, which were then intercrossed to produce PrxIII^{+/+}, PrxIII^{+/-} and PrxIII^{-/-} littermates for experiments. PrxIII genotypes were determined by PCR analysis as previously described.¹¹ Use of the experimental animals was approved by the institutional ethics committee of Tohoku University, Japan.

Measurement of blood pressure. Mean arterial blood pressure of mice was measured through the carotid artery.¹² After anesthesia with diethyl ether the carotid artery of maternal mouse was isolated and a catheter was inserted into the artery

through a tiny incision. Another end of the catheter was connected with a pressure transducer (Spectramed Medical Products, Singapore) whose output was fed into an amplifier (NEC-Sanei Co., Tokyo, Japan) and recorded. Age-matched non-pregnant female mice were used as controls.

Systemic detection of oxidative stress.

Brains, hearts, lungs, livers, kidneys, skeletal muscles and placentas were removed from maternal mice that were pregnant for 18 days. Samples from age-matched non-pregnant mice were used as controls. The samples were homogenized through ultrasonic disruption and protein concentration was detected by Bio-Rad protein assay kit. After separation on 15% SDS-PAGE and transfer to Immobilon membrane (Millipore Corporation), proteins were recognized with primary antibody anti-ET-1 (Santa cruz). Visualization was achieved with ECL western blotting detection system (Amersham). Anti-β-actin antibody (Sigma) was used as internal control.

Statistic analysis. The value of blood pressure was presented as mean ± SEM and was compared among PrxIII^{+/+}, PrxIII^{+/-} and PrxIII^{-/-} mice by analysis of variance. p < 0.05 was considered to be statistically significant.

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