

RABMS Journal of Research in Applied and Basic Medical Sciences

# Effect of Antihypertensive Drugs on Placental Villi of Preeclamptic Patients: A Histopathological Study

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#### Abstract

**Background & Aims:** The placenta has been described as the most accurate record of the infant's prenatal experience. The present study was a quantitative study and it was carried out to see the effect of pre-eclampsia on histopathology of placenta and the comparative changes in placenta of pre-eclamptic patients taking antihypertensives.

*Materials & Methods*: A total of 60 patients were selected from the Obstetrics and Gynecology Department of Jawahar Lal Nehru Medical College & Hospital, AMU, Aligarh, UP, India and were grouped into control, untreated, and treated groups. Placenta of each group was fixed in 10% formalin solution and processed for section cutting. 5-micron thick sections were prepared and stained with Hematoxylin-eosin and Wiegerts & Van gieson stain and observed under the light of the microscope. A total of 200villi/hpf was counted and percentages for each parameter were expressed.

*Result*: It has been shown that there was a significant reduction (<0.05) in the percentage of placental villi showing syncytial knot, cytotrophoblast proliferation, Basement membrane thickening, fibrinoid necrosis, and stromal fibrosis.

*Conclusion*: It has been concluded that antihypertensive drugs were not only effective in controlling the symptoms of pre-eclampsia but also effective in minimizing the placental histopathology and helpful in continuation of pregnancy under low utero-placental flow without completely normalizing the placental changes.

Keywords: Pre-eclampsia, Methyldopa, Chorionic villi, Basement membrane

#### Received 10 May 2022; accepted for publication 19 June 2022

#### Introduction

The presence of the fetus in the womb depends on a vital organ called the placenta. It is the most accurate data on the prenatal experience of an infant (1).

Hypertensive diseases that complicate pregnancy are common and, along with hemorrhage and infection, form a deadly triad, leading to numerous maternal mortalities and thus fetal death. Although the cause

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remains unknown, preeclampsia can only occur if there is a placenta. In preeclampsia, maternal blood flow to the fetal placental unit is significantly reduced compared to normal pregnancy (2).

The placenta fails to develop appropriately in preeclampsia. There is a resultant placental endothelial dysfunction and reduced maternal uteroplacental blood flow, which can indirectly constrict fetal stem arteries, resulting in changes found in placenta (3). Majumdar et al. (4) have noted significant morphological changes in placenta of preeclamptic patients.

The  $\alpha$ -methyldopa ( $\alpha$ MD) is most commonly used to treat preeclampsia. Trophoblast cells carry the  $\alpha$ 2adrenergic receptor, which is the primary target of  $\alpha$ methyldopa (5).

#### **Materials & Methods**

This study is a quantitative study to confirm the effect of preeclampsia on the histopathology of placental villi and to see how effective antihypertensive drugs are in restoring normal histopathological changes in the hypertensive placenta.

Ethical clearance was taken from Institutional Ethics Committee (IEC).

A total of 60 cases were taken and categorized into three categories: control, untreated, and treated.

#### **Inclusion Criteria:**

- Only cases with blood pressure less than 140/90mm Hg without proteinuria or edema were included in the control group.
- Only cases with or without edema and / or proteinuria, with blood pressure in the range of 140/90 mm Hg or higher were included in the untreated group.
- Only cases with preeclampsia who took αmethyldopa 250 mg TDS or QID for approximately

an equal duration were included in the treated group.

## **Exclusion Criteria:**

- Patients with other comorbidities such as diabetes, etc.
- Those who did not take methyldopa on a regular basis, and
- Those who did not regularly attend prenatal clinics. After birth, the placenta was collected and fixed with

10% formalin. Processing was done. Sections (5  $\mu$  thick) were prepared and then stained with hematoxylin eosin and Wiegert and Van gieson stain, and viewed under a light microscope. 200 villi / hpf was counted and percentages of each parameter were expressed.

## Statistical analysis:

All statistical data were analyzed using the SPSS software version 16 statistics package for Windows. Continuous variables are expressed as mean  $\pm$  standard deviation. Depending on the normal distribution, we used the independent samples' unpaired "t" test to compare the continuous variables between the two groups. The p-value less than 0.05 was considered significant.

## Results

The obtained data were compared as follows.

After statistical analysis, a significant increase (p<0.05) of histopathological findings in the placental villi was observed in the untreated group compared to the control group (Table 1). When the data between untreated and treated groups were analyzed, there was significant reduction (p<0.05) also а of histopathological changes in the placental villi of the treated group (Table 2). But the reductions in histopathological changes in patients receiving antihypertensives were not significantly close to those in controls (Table 3).

#### **Table 1:** Comparison of control group and untreated group

Histological findings features	Hypertensive untreated group	Hypertensive treated group	Statistical significance
(PER HPF)	$(\text{mean} \pm \text{sd})$	$(\text{mean} \pm \text{sd})$	(p<0.05)
Syncytial knots	$56.21\pm3.01$	$49.42\pm 6.06$	Significant

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Cytotrophoblastic cells	$56.74\pm3.04$	$46.36\pm8.53$	Significant
proliferation			
Thickening of Basement	$41.20\pm2.42$	$18.95\pm3.71$	Significant
membrane			
Fibrinoid necrosis	$33.25\pm3.01$	$8.40\pm2.09$	Significant
Stromal fibrosis	$58.95 \pm 2.32$	$22.35\pm5.95$	Significant

# Table 2. Comparison of untreated group and treated group

Histological feature (PER HPF)	Control group	Hypertensive untreated group	Statistical significance (p<0.05)
	$(mean \pm sd)$	$(mean \pm sd)$	
Syncytial knots	$17\pm2.32$	$56.21\pm3.09$	Significant
Cytotrophoblastic cells proliferation	$12.55\pm2.06$	$56.74\pm3.04$	Significant
Thickening of Basement Membrane	0.00	$41.20\pm2.42$	Significant
Fibrinoid necrosis	$2.50 \pm 1.23$	$33.25\pm3.01$	Significant
Stromal fibrosis	$1.95\pm0.76$	$58.85 \pm 2.32$	Significant

## Table 3. Comparison of treatment group and control group

	6 1		
Histological findings features (PER HPF)	Control group	Hypertensive treated	Statistical significance
	(mean ±sd)	group (mean ±sd)	(p<0.05)
Syncytial knots	$17\pm2.32$	$49.42\pm 6.05$	Significant
Cytotrophoblastic cells proliferation	$12.55\pm2.06$	$46.36\pm88.53$	Significant
Thickening of Basement membrane	0.00	$18.95\pm3.71$	Significant
Fibrinoid necrosis	2.50±1.23	$8.40\pm2.09$	Significant
Stromal fibrosis	$1.95\pm0.76$	$22.35\pm5.95$	Significant

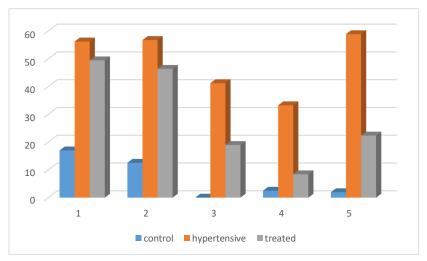


Fig. 1. comparison among 3 groups

1. Syncytial knot formation 2. Cytotrophoblast proliferation 3. Basement membrane thickening

4. Fibrinoid necrosis 5. Stromal fibrosis

# Discussion

This study was a quantitative study of histopathology of placentas of three groups, i.e. controlled hypertension, untreated hypertension, and treated hypertension.

The histopathology of placenta of hypertensive mothers showed a significant increase in the percentage of syncytial knot formation, cytotrophoblast proliferation, basement membrane thickening, fibrinoid necrosis, stromal fibrosis, and villous edema in comparison to control group. These results are completely consistent with previous studies by Anderson and Mackey (6) and Jones and Fox (7).

Mallik et al. (8), Masodkar et al. (9), Sodhi et al. (10), and Majumdar et al (11) also observed increased syncytial knot formation, cytotrophoblast formation, fibrinoid necrosis, and stromal fibrosis in the placentas of preeclamptic patients.

According to Fox (12), excessive syncytial knot formation is a good indicator of the degree of hypoperfusion of the villi.

The degree of cytotrophoblast hyperplasia is associated with the degree of syncytial damage. Therefore, it serves as a rough quantitative indicator of the ischemic severity to which the villi have been exposed (13).

Genset(14) also reported that an overall reduction of fetal perfusion of the placenta results in stromal fibrosis and extensive syncytial formation in generalized form.

Hypertensive pregnancies have placental lesions such as infarctions, villous fibrinod necrosis, and villous hypermaturity that are significantly related to the severity of the hypertension. In addition, when hypertension is high, angiogenesis and vascularization of placenta are significantly impaired (15).

In mild or severe preeclampsia/eclampsia, the placenta shows significant histological evidence of ischemia, and the degree of placental involvement due to infarction is inversely proportional to the birth weight of the fetus. The fetal placental ratio increased as the severity of the disease increased, but the average body weight decreased (16). Placental villi of hypertensive mothers taking  $\alpha$ Methyldopa showed a significant reduction in the percentages of all the histopathological findings. No parallel data supported our study on the histopathology of placental villi of treated patients.  $\alpha$ -Methyldopa may have specific effects on placental and/or endothelial cell function in patients with pre-eclampsia by altering angiogenic proteins. In preeclampsia,  $\alpha$ -methyldopa can directly affect placental synthesis and/or secretory function (17).

#### Conclusion

Hence, it has been concluded that antihypertensive drugs were effective in controlling the symptoms of preeclampsia and minimizing the placental histopathology and also helpful in continuation of pregnancy under low uteroplacental flow without completely normalizing the placental changes.

#### **Financial/ support**

Nil

#### **Conflicts of interest**

There are no conflicts of interest

## Ethical Statement

This study was conducted in accordance with the ethical principles of the Declaration of Helsinki and the guidelines for ethical conduct in the care and use of human participants in research. The study protocol was approved by the Institutional Ethics Committee (IEC).

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