Prevalence of preeclampsia and the associated risk factors among pregnant women in Bangladesh

Ananya Dutta Mou, Zitu Barman, Mahmudul Hasan, Rakib Miah, Jaasia Montahena Hafsa, Aporajita Das Trisha & Nurshad Ali

Preeclampsia is a multi-organ system disorder of pregnancy and is responsible for a significant rate of maternal morbidity and mortality worldwide. In Bangladesh, a large number of obstetric deaths occur every year but the exact reasons are not well investigated. The data regarding preeclampsia and its associated risk factors are scarce or limited in pregnant women in Bangladesh. Therefore, we aimed to conduct a cross-sectional study to estimate the prevalence of preeclampsia and identify the possible risk factors in a pregnant women cohort in Bangladesh. In this cross-sectional study, a total of 111 participants were enrolled and asked to include their anthropometric, socio-demographic, and other related lifestyle information in a standard questionnaire form. Blood samples were also collected from each participant to analyze serum levels of lipid profile, liver enzymes, uric acid, and creatinine by using standard methods. Logistic regression analysis was performed to identify the factors associated with preeclampsia. The overall prevalence of preeclampsia was 14.4%. About 10% of the pregnancies were found to have preeclampsia after 20 weeks of gestation without a previous history of hypertension. On the other hand, the prevalence of preeclampsia that superimposed on chronic hypertension was found to be 5.4%. Serum levels of TC, LDL-C, ALT and uric acid were significantly higher and HDL-C was significantly lower in preeclamptic pregnancies than the non-preeclamptic pregnancies. Respondents who required to take antihypertensive medications (AOR 5.45, 95% CI [1.09, 27.31]) and who never took antenatal care (AOR 6.83, 95% CI [1.00, 46.48]) were more likely to be preeclamptic. In conclusion, the present study showed a comparatively high prevalence of preeclampsia among pregnant women in Bangladesh. Some programmatic interventions such as medication for hypertension, antenatal visits to doctors, delivery and postnatal care services should be considered to reduce and prevent the hypertensive pregnancy disorders in Bangladesh.

Preeclampsia is one of the leading causes of maternal morbidity and mortality in the world. Preeclampsia is hypertension that generally occurs after 20 weeks of gestation along with proteinuria. When proteinuria is absent, preeclampsia is diagnosed in association with liver dysfunction, thrombocytopenia, pulmonary oedema, new onset of kidney dysfunction, or new-onset of cerebral or visual disturbances. It can cause severe morbidity, chronic disability, and even death of mothers and babies. Moreover, it is linked with an increased risk of cardiovascular diseases and type 2 diabetes in a mother’s later life.

In developing countries, women are at 14 times higher risk of dying from obstetric complications compared to developed countries. Pregnancy complications caused around 289,000 deaths of women worldwide in 2013 and 99% of them were from developing countries. Globally, about 12% of mothers die only from preeclampsia. As estimated by WHO, the occurrence of preeclampsia is seven times higher in developing countries compared to developed countries. The prevalence of preeclampsia ranges between 1.8 and 16.7% in developing countries.

Several studies have demonstrated that preeclampsia is linked with the failure of the trophoblastic invasion of maternal spiral arteries which leads to higher vascular resistance of uterine arteries and lower uteroplacental blood flow. If unmanaged, preeclampsia can prevent the flow of adequate blood and oxygen to the developing fetus, cause maternal liver and kidney damage and sometimes progress to eclampsia, a serious condition involving seizures. There is no known cure for preeclampsia other than the delivery of the baby or the placenta.
Early diagnosis and close monitoring can help to control preeclampsia during pregnancy. Developed countries have reduced the incidence of eclampsia and associated mortality by nearly 90% through early detection during antenatal care and by increasing access to hospital care for preeclamptic women\(^2\). The prevalence of preeclampsia is not well-documented and the associated data are scarce in Bangladesh. Thus, this study aimed to estimate the prevalence of preeclampsia, measure some biochemical variables, and identify the associated risk factors among pregnant women in the Sylhet region of Bangladesh.

**Methods**

**Study design and study participants.** A cross-sectional study was designed and conducted during the period from October 2019 to October 2020. All the relevant analyses were performed at the Department of Biochemistry and Molecular Biology, Shahjalal University of Science and Technology, Sylhet, Bangladesh. The study participants were enrolled from the Gynaec and Obstetrics unit of Sylhet Osmani Medical College, Sylhet Diabetic Hospital, and another private hospital in the Sylhet city area. About 220 pregnant women, who visited these hospitals for their regular checkups, were invited to participate in the study. Among them, 111 women (29 rural and 82 urban) were agreed to participate. Their anthropometric, socio-demographic, individual as well as family history of diabetes and hypertension and lifestyle information were noted in a standard questionnaire form. Information regarding preexisting hypertension and the necessity of antihypertensive medications were either obtained from the participant’s self-report or from their medical documents. The anthropometric measurements were made with the assistance of trained medical personnel following the standard procedure described elsewhere\(^14–17\). Briefly, The systolic blood pressure (SBP) and diastolic blood pressure (DBP) were measured using an automated blood pressure recorder (Omron M10, Japan) while they were seated in a comfortable position after at least 10 min of rest. The weight was recorded to the nearest 0.1 kg (kg) with the subject standing and wearing light clothing. Inclusion criteria: pregnant women of any trimester were included as study participants. Exclusion criteria: participants suffering from preexisting kidney disease or urinary tract infection, preexisting liver disease, and preexisting thyroid disorder were excluded from the study. During exclusion of the participants, we considered the subject’s self-reported evidence and/or medical records of their preexisting diseases.

**Sample collection and biochemical analysis.** About 5 ml of the venous blood samples were collected from each participant with the help of expert clinical assistants. Blood samples were then placed on ice and transported immediately to the laboratory. The blood samples were then kept for 10 minutes at room temperature for coagulation and centrifuged (Sorvall ST 8R, Thermo Scientific, Germany) at 4400 rpm for 10 minutes. The isolated sera were then stored in a \(-20^{\circ}\)C refrigerator until the biochemical markers were analyzed. Serum levels of TG (triglyceride), TC (total cholesterol), HDL-C (high-density lipoprotein cholesterol), LDL-C (low-density lipoprotein cholesterol), creatinine, and uric acid were measured by colorimetric methods with a biochemistry analyzer (Humalyzer 3000, USA) using available diagnostic kits (Human Diagnostic, Germany)\(^18–23\). Serum levels of ALT (alanine aminotransferase) and GGT (gamma-glutamyl transferase) were measured by kinetic methods\(^24\). The proteinuria was determined using the urine dipstick method\(^19\).

**Diagnostic criteria.** Proteinuria: women with a protein level of 1 + were grouped as having proteinuria.

- **Trimesters, gravidity, and parity:** women with gestation periods of week 1 to 12 were included as first trimester, 13 to 28 as second trimester, and 29 to 40 as third trimester. Gravidity was defined as the total number of pregnancies of a woman, regardless of the outcome. Parity was denoted as the number of pregnancies reaching ≥ 24 weeks.

- **Preeclampsia:** Preeclampsia was defined as having hypertension (≥ 140 mmHg SBP and/or ≥ 90 mmHg DBP) along with either proteinuria or elevated liver enzyme (e.g., ALT level > 40 IU/L) or kidney dysfunction (creatinine > 1 mg/dL)\(^25\). Preeclampsia that developed after 20 weeks of gestation with the previous history of normal blood pressure was denoted as de novo\(^26\). Preeclampsia with pre-pregnancy hypertension or that developed before 20 weeks of gestation was defined as preeclampsia superimposed on chronic hypertension\(^27\).

**Data processing and analysis.** Descriptive statistics were used to present the baseline data variables. P-values were obtained from the independent sample t-test for comparison between quantitative variables. Binary logistic regression was applied to determine the relationship between the dependent variables and independent variables. The independent variables that were significant at univariate analysis and some relevant variables were included in the multiple logistic regression model. IBM SPSS, version 25.0 was used for statistical data analysis. The p-value < 0.05 was considered statistically significant.

**Ethical consideration.** This study was approved by the internal review board at the Department of Biochemistry and Molecular Biology, School of Life Sciences, Shahjalal University of Science and Technology. All methods in the current study were carried out following the relevant guidelines and regulations. All the study participants were informed about the study aims and written informed consent was obtained from them before inclusion in the study.

**Results**

**Socio-demographic characteristics of the study participants.** A total of 111 pregnant women participated in the study from the Sylhet region. The age range of the subjects was 17 to 37 years, with a mean of 26.4 ± 4.9 years. Overall, the mean of systolic (SBP), diastolic (DBP), and pulse pressure (PP) were
122.2 ± 19.9 mmHg, 79.8 ± 13.6 mmHg, and 42.4 ± 12.0 mmHg, respectively, for the study cohort. Among the study participants, 22.9% had hypertension prior to pregnancy and 19.1% were diabetic, whereas 43.7% of the study subjects had a family history of hypertension, 38.4% had a family history of diabetes and 7% had a family history of hypertensive disorders in pregnancy (Table 1).

Obstetrical characteristics of the respondents. Among the total subjects, 19.6% of the participants were in the first trimester, 21.5% in the second trimester, and 58.9% were in the third trimester of their pregnancy. The percentage of primiparous pregnancies was 36.8% in total. 3.4% of the pregnancies were twin in type and the rest 96.6% were singleton. 26.6% of the study subjects were detected as having hypertensive pregnancies. None of the participants had reported convulsion during their pregnancy period (Table 2).

Prevalence of preeclampsia in study participants. The overall prevalence of preeclampsia was 14% among the study subjects. 9.9% of the pregnancies were found to have preeclampsia after 20 weeks of gestation without prior history of hypertension, whereas, the prevalence of preeclampsia that superimposed on chronic hypertension was found to be 5.4% (Table 3).

Level of blood pressure variables in the study subjects. Overall, the mean of SBP, DBP, and PP for the preeclamptic pregnancies were 148.7 ± 25.6 mmHg, 100.7 ± 16.2 mmHg, and 48 ± 17.4 mmHg, respectively. All of their values were significantly higher (p < 0.05) in preeclamptic pregnancies compared to non-preeclamptic pregnancies (Fig. 1A). Moreover, PP was found significantly higher in chronic hypertensive pregnancies with preeclampsia compared to de novo appearance after 20 weeks of gestation (Fig. 1B).

Biochemical markers in preeclamptic and non-preeclamptic pregnancies. Serum levels of TC, LDL-C, ALT and uric acid were significantly higher in preeclamptic pregnancies compared to the pregnancies without preeclampsia (at least p < 0.001 for all cases). The mean level of HDL-C was found significantly lower in preeclamptic subjects than non-preeclamptic subjects (p < 0.01) (Table 4).

Factors associated with preeclampsia. We also assessed the relationships between socio-demographic factors, obstetrical data, and the prevalence of preeclampsia. In univariate analysis, the requirement of antihypertensive medications, less intake of fruits and vegetables, and not visiting a doctor during pregnancy (antenatal care) were found to be significantly associated with the prevalence of preeclampsia. After adjusting for these significant independent factors using multiple regression analysis, the requirement of antihypertensive medications and lack of antenatal care during pregnancy were found as independent predictors for preeclampsia. Respondents who were required to take antihypertensive medications either occasionally or regularly were more likely to be preeclamptic than those who did not (AOR 5.45, 95% CI [1.09, 27.31]). Women who never consulted any doctor during their pregnancy period had about 6.8 times higher odds of being preeclamptic than women who did occasionally or regularly (AOR 6.83, 95% CI [1.00, 46.48]) (Table 5).

Discussion
The present study estimated the prevalence of preeclampsia with some biochemical variables and identified the possible associated risk factors among pregnant women in Bangladesh. The overall prevalence of preeclampsia was 14.4% in pregnancy. Lack of awareness and insufficient antenatal care in rural areas can be the reason behind this higher prevalence.

Up to now, there is limited information on preeclampsia and its associated factors among Bangladeshi women. A recent study reported about 44% preeclampsia among pregnant women who visited a Maternity Clinic in Dhaka, Bangladesh11. A probable reason for the higher prevalence of preeclampsia in that study was the inclusion of the participants from their third trimester of pregnancy, whereas in our study we included pregnant women from all three trimesters.

The prevalence of preeclampsia in the present study is higher compared with the global average, which is around 2%28. However, the prevalence ranges from 1.8 to 16.7% in developing countries4, and thus our finding falls in the upper side of this range. A study conducted in our neighboring country India reported the prevalence of preeclampsia about 28% with a variation in the prevalence across the states or regions29. A review covering larger data sets reported the prevalence of preeclampsia 0.2–6.7% in Asia, 0.5–2.3% in Africa, 2.8–5.2% in Europe, 2.8–9.2% in Oceania, 1.8–7.7% in South America and the Caribbean, and 2.6–4.0% in North America30. Another review summarized the data for the Asian regions and reported the prevalence of preeclampsia 2.07% in China, 1.19% in Japan, 2.22% in Thailand and 0.59% in Nepal31. Some possible reasons may be associated with the high prevalence of preeclampsia in Bangladesh pregnant women compared to pregnant women from other countries. For example, a large proportion of our study participants were in their third trimester of pregnancy which can be a possible underlying factor of higher prevalence of preeclampsia. Moreover, Bangladesh is a developing country with lower health concerns in the overall population especially people living in the rural regions. Furthermore, socioeconomic status and not taking treatment at the proper time may also be associated with the high prevalence of preeclampsia in Bangladeshi pregnant women.

In the present study, we also measured some biochemical variables and we observed that participants with preeclampsia had significantly increased serum levels of TC, and LDL-C and significantly decreased levels of HDL-C compared to subjects with normal pregnancy (p < 0.01), which is in line with a study finding conducted by Wang et al.32. Moreover, we found a significantly increased serum level of ALT in preeclamptic pregnancies compared to normal pregnant women. The high level of ALT may indicate liver dysfunction due to preeclamptic...
Table 1. Sociodemographic characteristics of the participants. Information about pre-pregnancy hypertension, diabetes mellitus, and amount of fruits and vegetable intake was collected from the participants. Family: it comprises of a group of kin related to each other by blood connection or by marital tie.
damage. In a previous study, serum ALT level was found significantly higher in preeclampsia/eclampsia groups compared to women with normal pregnancies\textsuperscript{33}. An elevated level of ALT was also observed in other studies which are in line with our present study\textsuperscript{34,35}.

In this study, serum uric acid levels were significantly higher in preeclamptic pregnancies compared to non-preeclamptic pregnancies, which is similar to other reports\textsuperscript{36,37}. Hyperuricemia is now considered one of the initial laboratory manifestations of preeclampsia\textsuperscript{36,37}. Several studies have demonstrated a positive relationship between serum uric acid and hypertension in the general population and pregnant women\textsuperscript{19,38}. Thus, monitoring of serum uric acid levels in the pregnancy period might be helpful in early detection and intervention of preeclampsia and in preventing associated complications.

In the present study, we identified some possible risk factors that are associated with preeclampsic pregnancies. For example, we found that women who are required to take antihypertensive medications for controlling their blood pressure level are more likely to suffer from preeclampsia. Additionally, women who consulted doctors regularly or occasionally during their pregnancy were less affected by preeclampsia compared to the women who never visited the doctor. Our finding is similar to a study conducted by Uddin et al. which showed that women

| Variables                  | n  | %    |
|----------------------------|----|------|
| Pregnancy period           |    |      |
| First trimester            | 21 | 19.6 |
| Second trimester           | 23 | 21.5 |
| Third trimester            | 63 | 58.9 |
| Hypertensive pregnancy     | 29 | 26.6 |
| Primiparous                |    |      |
| Yes                       | 39 | 36.8 |
| No                        | 67 | 63.2 |
| Pregnancy type             |    |      |
| Singleton                  | 56 | 96.6 |
| Twin                      |  2 |  3.4 |
| Gravidity                  |    |      |
| 1                         | 39 | 39.8 |
| 2                         | 21 | 21.4 |
| 3                         | 21 | 21.4 |
| ≥4                        | 17 | 17.3 |
| Parity                     |    |      |
| <2                        | 49 | 51.0 |
| 2–3                       | 37 | 38.5 |
| ≥4                        | 10 | 10.4 |
| Previous stillborn or miscarriage history |  |      |
| 0                          | 69 | 74.2 |
| 1                          | 20 | 21.5 |
| ≥2                         |  4 |  4.3 |
| Number of children         |    |      |
| <2                         | 78 | 71.6 |
| ≥2                         | 31 | 28.4 |
| Convulsion during pregnancy|  0 |  0   |

Table 2. Obstetrical characteristics of the respondents. Hypertensive pregnancy: SBP ≥ 140 mmHg and/or DBP ≥ 90 mmHg\textsuperscript{25}. Primipara means first pregnancy.

| Categories                                | n (%) |
|-------------------------------------------|-------|
| N                                         | 111   |
| Non-preeclamptic pregnancies              | 92 (83.8) |
| Preeclamptic pregnancies                  | 16 (14.4) |
| De novo                                   | 11 (9.9) |
| Superimposed on chronic hypertension      |  6 (5.4) |

Table 3. Prevalence of preeclampsia among the study participants. De novo: preeclampsia with hypertension developed after 20 weeks of gestation\textsuperscript{26}. Superimposed: preeclampsia with pre-pregnancy hypertension or that developed before 20 weeks of gestation was defined as preeclampsia superimposed on chronic hypertension\textsuperscript{27}.  

In a previous study, serum ALT level was found significantly higher in preeclampsia/eclampsia groups compared to women with normal pregnancies\textsuperscript{33}. An elevated level of ALT was also observed in other studies which are in line with our present study\textsuperscript{34,35}. In this study, serum uric acid levels were significantly higher in preeclamptic pregnancies compared to non-preeclamptic pregnancies, which is similar to other reports\textsuperscript{33,36}. Hyperuricemia is now considered one of the initial laboratory manifestations of preeclampsia\textsuperscript{36,37}. Several studies have demonstrated a positive relationship between serum uric acid and hypertension in the general population and pregnant women\textsuperscript{19,38}. Thus, monitoring of serum uric acid levels in the pregnancy period might be helpful in early detection and intervention of preeclampsia and in preventing associated complications.

In the present study, we identified some possible risk factors that are associated with preeclampsic pregnancies. For example, we found that women who are required to take antihypertensive medications for controlling their blood pressure level are more likely to suffer from preeclampsia. Additionally, women who consulted doctors regularly or occasionally during their pregnancy were less affected by preeclampsia compared to the women who never visited the doctor. Our finding is similar to a study conducted by Uddin et al. which showed that women
having poor access to antenatal care were more affected by hypertension in their pregnancy. According to a report, about 64% of pregnant women in Bangladesh aged 15–49 receive at least one antenatal care visit with a skilled health provider. The detection and counselling of preeclampsia essentially depend on the frequency of antenatal care and measurement of the blood pressure of the mother during the visit. In developing countries, many women with preeclampsia, mostly at the community level, still do not get proper antenatal care and are at higher risk of developing serious complications.

A recent study determined the possible risk factors for preeclampsia in pregnant women who went to a tertiary hospital in Dhaka, Bangladesh showed that education level, occupation, bad obstetrics history, or history of abortion were not significantly associated with preeclampsia development, which is similar to our study.
findings. According to the same study, preeclampsia was high in women having poor antenatal care, which is also in line with ours. Although obesity, primigravida, preexisting hypertension, advanced maternal age, and family history of hypertension are common risk factors for preeclampsia globally\textsuperscript{42,43}, we did not find any of them significantly associated with preeclampsia. This variation of our study with other studies may be a result of the relatively small number of participants of our study enrolled from a single region of Bangladesh.

Although the present study estimated the prevalence of preeclampsia and identified some associated risk factors, the study also had some limitations. Firstly, the study participants were included in this study from hospitals and healthcare centers which may increase the prevalence of preeclampsia. Secondly, we could not collect detailed information on participants’ access to hospital care facilities which should be considered in such a study. Thirdly, the sample size in our study was relatively small and the study was conducted in a divisional region of Bangladesh, thus our findings may not be nationally representative. However, our findings may be useful for further studies in this area in Bangladesh.

Table 5. Multiple logistic regression analysis of factors that can be associated with preeclampsia among pregnant women. *p < 0.05; **p < 0.01. \textsuperscript{a} Adjusted for the variables found significant in univariate regression analysis (Requirement of antihypertensive medication, intake of fruits and vegetables, frequency of visiting doctor during pregnancy).

| Variables                        | Preeclampsia | COR (95% CI) | AOR (95% CI)\textsuperscript{a} | p-value |
|----------------------------------|--------------|--------------|----------------------------------|---------|
| Maternal age                     |              |              |                                  |         |
| < 25                             | 34 (85.0)    | 6 (15.0)     | 1.71 (0.39–7.43)                 | 0.44 (0.07–2.98) | 0.400 |
| 25 – 29                          | 29 (90.6)    | 3 (9.4)      | 1.00                             | 1.00    |
| ≥30                              | 29 (82.9)    | 6 (17.1)     | 2.00 (0.46–8.77)                 | 0.92 (0.12–7.15) | 0.939 |
| Body weight (Kg)                 |              |              |                                  |         |
| <65                              | 47 (92.2)    | 4 (7.8)      | 1.00                             | 1.00    |
| 65 – 74                          | 13 (81.3)    | 3 (18.8)     | 2.71 (0.54–13.68)                | 0.88 (0.01–56.16) | 0.950 |
| ≥75                              | 12 (85.7)    | 2 (14.3)     | 1.96 (0.32–11.99)                | 2.19 (0.05–90.22) | 0.680 |
| Pre-pregnancy hypertension       |              |              |                                  |         |
| No                               | 54 (84.4)    | 10 (15.6)    | 1.00                             | 1.00    |
| Yes                              | 14 (73.7)    | 5 (26.3)     | 1.93 (0.57–6.56)                 | 1.03 (0.11–9.28) | 0.979 |
| Requirement of antihypertensive medication |          |              |                                  |         |
| Regularly or occasionally        | 14 (73.7)    | 5 (26.3)     | 0.24 (0.06–0.93)*                | 5.45 (1.09–27.31) | 0.039* |
| No                               | 59 (92.2)    | 5 (7.8)      | 1.00                             | 1.00    |
| Primiparous                      |              |              |                                  |         |
| Yes                              | 30 (81.1)    | 7 (18.9)     | 1.00                             | 1.00    |
| No                               | 58 (87.9)    | 8 (12.1)     | 0.59 (0.20–1.79)                 | 0.57 (0.09–3.46) | 0.536 |
| Gravidity                        |              |              |                                  |         |
| 1                                | 30 (81.1)    | 7 (18.9)     | 1.40 (0.32–6.11)                 | 1.07 (0.11–10.28) | 0.951 |
| 2                                | 20 (95.2)    | 1 (4.8)      | 0.30 (0.03–3.15)                 | 0.29 (0.02–4.94) | 0.391 |
| 3                                | 18 (85.7)    | 3 (14.3)     | 1.00                             | 1.00    |
| ≥4                               | 15 (88.2)    | 2 (11.8)     | 0.80 (0.12–5.44)                 | 1.05 (0.11–10.23) | 0.968 |
| Parity                           |              |              |                                  |         |
| <2                               | 40 (85.1)    | 7 (14.9)     | 1.44 (0.39–5.36)                 | 1.57 (0.21–11.55) | 0.659 |
| 2 – 3                            | 33 (89.2)    | 4 (10.8)     | 1.00                             | 1.00    |
| ≥4                               | 7 (70.0)     | 3 (30.0)     | 3.54 (0.64–19.45)                | 3.44 (0.34–34.65) | 0.294 |
| Previous stillborn or miscarriage history |          |              |                                  |         |
| 0                                | 57 (85.1)    | 10 (14.9)    | 1.00                             | 1.00    |
| 1                                | 17 (85.0)    | 3 (15.0)     | 1.01 (0.25–4.08)                 | 0.77 (0.11–5.39) | 0.789 |
| ≥2                               | 3 (75.0)     | 1 (25.0)     | 1.90 (0.18–20.14)                | 1.83 (0.01–380.63) | 0.825 |
| Number of children               |              |              |                                  |         |
| <2                               | 64 (85.3)    | 11 (14.7)    | 1.00                             | 1.00    |
| ≥2                               | 27 (90.0)    | 3 (10.0)     | 0.65 (0.17–2.50)                 | 1.69 (0.24–11.79) | 0.596 |
| Intake of fruits and vegetables  |              |              |                                  |         |
| Medium to high                   | 65 (90.3)    | 7 (9.7)      | 1.00                             | 1.00    |
| Low                              | 9 (56.3)     | 7 (43.8)     | 7.22 (2.05–25.42)**              | 4.29 (0.84–21.98) | 0.081 |
| Frequency of visiting doctor during pregnancy |          |              |                                  |         |
| Regularly or occasionally        | 68 (88.3)    | 9 (11.7)     | 1.00                             | 1.00    |
| Never                            | 6 (54.5)     | 5 (45.5)     | 6.30 (1.59–24.91)**              | 6.83 (1.00–46.48) | 0.050* |
Conclusion
The prevalence of preeclampsia was comparatively higher in rural areas of Bangladesh. Serum levels of TC, LDL-C, ALT and uric acid were found to be significantly higher and HDL-C was significantly lower in preeclamptic pregnancies compared to the pregnancies without preeclampsia. Participants who were required to take anti-hypertensive medications either occasionally or regularly and who never took antenatal care were found more likely to have preeclampsia. Thus, the presence of preeclampsia can also be predicted by investigating some biochemical components and proper antenatal care is an important part of prevention. Furthermore, prospective studies are needed to understand the underlying mechanisms of preeclampsia in pregnancies and their proper management strategies.

Received: 24 August 2021; Accepted: 19 October 2021
Published online: 29 October 2021

References
1. Al-Jameil, N. A brief overview of preeclampsia. J. Clin. Med. Res. 6, 1–7 (2013).
2. Vest, A. R. & Cho, L. S. Hypertension in pregnancy. Curr. Atheroscler. Rep. 16, 395 (2014).
3. American College of Obstetricians & Task Force on Hypertension in Pregnancy. Hypertension in pregnancy. Report of the American College of Obstetricians and Gynecologists’ Task Force on Hypertension in Pregnancy. Obstet. Gynecol. 122, 1122–1131 (2013).
4. Lindheimer, M. D. & Umans, J. G. Explaining and predicting preeclampsia. N. Engl. J. Med. 355, 1056–1058 (2006).
5. Tessema, G. A., Tekeste, A. & Ayele, T. A. Preeclampsia and associated factors among pregnant women attending antenatal care in Dessie referral hospital, Northeast Ethiopia: A hospital-based study. BMC Pregnancy Childbirth 15, 1–7 (2015).
6. Nour, N. M. An introduction to maternal mortality. Rev. Obstet. Gynecol. 1, 77–81 (2008).
7. Osungbade, K. O. & Ige, O. K. Public health perspectives of preeclampsia in developing countries: Implication for health system strengthening. J. Pregnancy 2011, 481095 (2011).
8. Belay, A. S. & Wudad, T. Prevalence and associated factors of pre-eclampsia among pregnant women attending anti-natal care at Mettu Karl referral hospital, Ethiopia: Cross-sectional study. Clin. Hypertens. 25, 1–8 (2019).
9. Phupong, V. et al. Predicting the risk of preeclampsia and small for gestational age infants by uterine artery Doppler in low-risk women. Arch. Gynecol. Obstet. 268, 158–161 (2003).
10. Lin, S., Shimizu, I., Suehara, N., Nakayama, M. & Aono, T. Uterine artery doppler velocimetry in relation to trophoblast migration into the myometrium of the placental bed. Obstet. Gynecol. 85, 760–765 (1995).
11. Khanum, H. Prevalance of pre-eclampsia and factors responsible among third trimester pregnant women in hospital of dhaka. Biomed. J. Sci. Tech. Res. 33, 26089–26097 (2021).
12. Wallis, A. B., Saftlas, A. F., Hsia, J. & Atrash, H. K. Secular trends in the rates of preeclampsia, eclampsia, and gestational hypertension, United States, 1987–2004. Am. J. Hypertens. 21, 521–526 (2008).
13. Goldenberg, R. L., McClure, E. M., MacGuire, E. R., Kamath, B. D. & Jobe, A. H. Lessons for low-income regions following the reduction in hypertension-related maternal mortality in high-income countries. Int. J. Gynecol. Obstet. 113, 91–95 (2011).
14. Ali, N. et al. Association between serum uric acid and metabolic syndrome: A cross-sectional study in Bangladeshi adults. Sci. Rep. 10, 1–7 (2020).
15. Ali, N. et al. Assessment of the relationship of serum liver enzymes activity with general and abdominal obesity in an urban Bangladeshi population. Sci. Rep. 11, 1–9 (2021).
16. Ali, N. et al. Hypertension prevalence and influence of basal metabolic rate on blood pressure among adults in Bangladesh. BMC Public Health 18, 1–9 (2017).
17. Rahman, S., Islam, S., Haque, T., Kathak, R. R. & Ali, N. Association between serum liver enzymes and hypertension: A cross-sectional study in Bangladeshi adults. BMC Cardiovasc. Disord. 20, 1–7 (2020).
18. Ali, N. et al. The relationship between serum uric acid and lipid profile in Bangladeshi adults. BMC Cardiovasc. Disord. 19, 1–7 (2019).
19. Ali, N. et al. Relationship between serum uric acid and hypertension: A cross-sectional study in Bangladeshi adults. Sci. Rep. 9, 1–7 (2019).
20. Islam, S. et al. Prevalence of elevated liver enzymes and its association with type 2 diabetes: A cross-sectional study in Bangladeshi adults. Endocrinol. Diabetes Metab. 3, e00116 (2020).
21. Molla, N. H. et al. Assessment of the relationship between serum uric acid levels and liver enzymes activity in Bangladesh adults. Sci. Rep. https://doi.org/10.1038/s41598-021-99623-z (2021).
22. Ali, N. et al. Prevalence of hyperuricemia and the relationship between serum uric acid and obesity: A study on Bangladesh adults. PLos ONE 13, e0206850 (2018).
23. Haque, T., Rahman, S., Islam, S., Molla, N. H. & Ali, N. Assessment of the relationship between serum uric acid and glucose levels in healthy, prediabetic and diabetic individuals. Diabetol. Metab. Syndr. 11, 49 (2019).
24. Islam, S. et al. Prevalence of elevated liver enzymes and its association with type 2 diabetes: A cross-sectional study in Bangladeshi adults. Endocrinol. Diabetes Metab. 3, 1–8 (2020).
25. Brown, M. A. et al. Hypertensive disorders of pregnancy: ISSHP classification, diagnosis, and management recommendations for international practice. Hypertension 72, 24–43 (2018).
26. Roberts, J. M., Pearson, G., Cutler, J. & Lindheimer, M. Summary of the NHLBI Working Group on research on hypertension during pregnancy. Hypertension 41, 437–445 (2003).
27. Podymow, T. & August, P. Hypertension in pregnancy. Adv. Chronic Kidney Dis. 14, 178–190 (2007).
28. Vata, P. K., Chauhan, N. M., Nallathambi, A. & Hussein, E. Assessment of prevalence of preeclampsia from Dilla region of Ethiopia. BMC. Res. Notes 8, 4–9 (2015).
29. Agrawal, S. Prevalence and risk factors for symptoms suggestive of pre-eclampsia in Indian women. J. Women’s Health Issues Care 3, 2–9 (2014).
30. Umesawa, M. & Kobashi, G. Epidemiology of hypertensive disorders in pregnancy: Prevalence, risk factors, predictors and prognosis. Hypertens. Res. 40, 213–220 (2017).
31. Abalos, E. et al. Pre-eclampsia, eclampsia and adverse maternal and perinatal outcomes: A secondary analysis of the World Health Organization Multicountry Survey on Maternal and Newborn Health. BJOG 121(Suppl), 14–24 (2014).
32. Wang, Y., Shi, D. & Chen, L. Lipid profile and cytokines in hypertension of pregnancy: A comparison of preeclampsia therapies. J. Clin. Hypertens. 20, 394–399 (2018).
33. Sarmah, J. Evaluation of serum aspartate aminotransferase (AST), Alanine aminotransferase (ALT), alkaline phosphatase (ALP), lactate dehydrogenase (LDH) and uric acid in preeclampsia. IOSR J. Dent. Med. Sci. Ver. III 14(14), 2279–2281 (2015).
34. Chaudhry, A. A. Hellp syndrome. Med. Forum Mon. 33, 32–32 (2002).
35. Munazza, B. et al. Liver function tests in preeclampsia. J. Ayub Med. Coll. Abbottabad 23, 3–5 (2011).
36. Baimbridge, S. A. & Roberts, J. M. Uric acid as a pathogenic factor in preeclampsia. Placenta 29, 67–72 (2008).
37. Roberts, J. M. et al. Uric acid is as important as proteinuria in identifying fetal risk in women with gestational hypertension. *Hypertension* **46**, 1263–1269 (2005).
38. Rajoria, L., Sonia, N. & Gidwani, C. Lactate dehydrogenase and uric acid as prognostic markers for hypertensive disorders of pregnancy. *IJSAR* 8, 12–17 (2018).
39. Uddin, A., Nessa, S., Chowdhury, S. & Banu, M. Hypertensive disorders of pregnancy and its outcome in a tertiary care hospital. *J. Armed Forces Med. Coll. Bangladesh* **9**, 38–42 (2013).
40. NIPORT. *Bangladesh Maternal Mortality and Health Care Survey 2010* (NIPORT, 2012).
41. Yeasmin, S. & Uddin, M. J. Determination of risk factors for pre-eclampsia in a tertiary hospital of Bangladesh. *Chattagram Maa-O-Shishu Hospital Med. Coll. J.* **16**, 29–32 (2017).
42. Paré, E. et al. Clinical risk factors for Preeclampsia in the 21st century. *Obstet. Gynecol.* **124**, 763–770 (2014).
43. English, F. A., Kenny, L. C. & McCarthy, F. P. Risk factors and effective management of preeclampsia. *Integr. Blood Press. Control* **8**, 7–12 (2015).
44. Abbassi-Ghanavati, M., Greer, L. & Cunningham, F. A reference table for clinicians. *Obstet. Gynecol.* **114**, 1326–1331 (2009).

**Acknowledgements**
The authors would like to thank all participants for their active participation in the study.

**Author contributions**
N.A. contributed to the conception and design of the study, data interpretation, drafting and revision of the manuscript. A.D.M. collected the samples, conducted the experiments and drafted the primary manuscript. Z.B, M.H, and R.M assisted in lab experiments and data analysis. J.M.H. and A.D.T. helped in sample and data collection. All authors contributed to manuscript revision, read and approved the submitted version.

**Competing interests**
The authors declare no competing interests.

**Additional information**

**Correspondence** and requests for materials should be addressed to N.A.

**Reprints and permissions information** is available at [www.nature.com/reprints](http://www.nature.com/reprints).

**Publisher's note** Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

[Open Access](http://creativecommons.org/licenses/by/4.0/) This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article’s Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article’s Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit [http://creativecommons.org/licenses/by/4.0/](http://creativecommons.org/licenses/by/4.0/).

© The Author(s) 2021