Preeclampsia management modalities and perinatal death: a retrospective study in Woldia general hospital

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Abstract

Background: Hypertensive disorders of pregnancy are among the most common causes of perinatal death. The disorders are highly linked to multiple factors that make prediction and prevention challenging. Early diagnosis and proper management play a crucial role in the wellbeing and life of the women and her baby. In this study, we aimed to assess the association between different management options of preeclampsia and perinatal death at a public hospital in Ethiopia.

Methods: A document review was conducted on 241 preeclamptic patients’ medical files who have been admitted and delivered in Woldia General Hospital from 2011 to 2016. The study was conducted from August 8 – September 10, 2017 in the aforementioned public hospital in Woldia town, Ethiopia. Associations were tested using Pearson chi squared test and binary logistic regression with a \( p \)-value < 0.05 considered significant.

Result: In this study, nearly 20 every 100 neonates from preeclamptic women has been died and the figure was higher (76.59% Vs 23.4%) among neonates from severe preeclamptic women than mild preeclamptic women (\( p = 0.01 \)). More than two thirds of the patients (69.3%) received magnesium sulfate to prevent convulsion. Perinatal death among women with diastolic blood pressures greater than 110 mmHg at admission was nearly 3 times (Adjusted Odds Ratio (AOR) = 2.824; 95% Confidence Interval (CI) (1.154–6.038)) higher compared to women with diastolic blood pressures below 110 mmHg.

Conclusion: In the 5-year period, the magnitude of perinatal death among inpatient preeclamptic women was remarkably high and of which stillbirths exceeded pre-discharge early neonatal death. Utilization of magnesium sulfate tended to increase across years. Maternal diastolic blood pressure at admission was significantly associated with perinatal death.

Keywords: Preeclampsia, Perinatal death, Management modalities, Woldia

Background

According to the Ethiopian Demographic and Health Survey, perinatal mortality rate is calculated as the sum of stillbirths and early neonatal deaths divided by the total sum of births expressed per 1000 pregnancies that lasted seven or more months [1]. Globally, nearly 16% of the estimated 2.6 million stillbirths annually occur in pregnancies complicated by pregnancy induced hypertension [2]. Hypertensive disorders of pregnancy (HDP) precede 10% of early neonatal deaths (8/1000 live births) [3] and a significant proportion of late neonatal deaths (3/1000 live births) [4]. Even though HDP by itself impact the fetus and its survival after
delivery, the management modalities used have also been associated with perinatal death.

Studies have shown that nationwide preeclampsia-related perinatal mortality was highly linked with the country’s economic index i.e., figures from high wealth index countries were low and disproportionately high perinatal mortality were reported from low- and middle-income countries (LMIC). In general, rates ranged from 4.7% to as high as 41.6% [5–8]. Similarly, perinatal mortality among patients with eclampsia was also reported to be 5 to 11% in High income countries (HIC) whereas it was as high as 40% in LMIC [9, 10]. Two studies conducted in public hospitals in Ethiopia; namely Jimma University Specialized Hospital and Mettu Karl Referral Hospital; revealed perinatal mortality among preeclamptic women were 317/1000 births and 120/1000 respectively [11, 12].

A study showed that inadequate and delayed initiation of treatment and preterm deliveries among preeclamptic women was found to be associated with poor fetal outcome [13]. As variation is perceived among the course of management modalities used by different institutions, its impact on perinatal death should be explored especially in resource limited areas. Therefore, the aim of this study was to assess the perinatal death rate and its association between preeclampsia management modalities and perinatal outcome among women who gave birth at Woldia General Hospital from 2011 to 2016.

Methods
Study setting, design and eligibility criteria
The study was conducted from August 8 – September 10, 2017 at Woldia General Hospital, Woldia, Ethiopia. Woldia town is the main town of North Wollo Zone. This town is located in Amhara region at 520 km North of the capital, Addis Ababa. According to the local administration report in 2016, the town has total population of 75, 446 of which 38,167 were males and 37,279 were females. Regarding the health care system, the town has one public hospital, two public health centers and 12 private pharmacies and 5 medium level clinics. The hospital provides both inpatient and outpatient services. At the time of the study, the hospital had 120 beds and 328 employees and of them there were 20 doctors, 128 nurses and 21 midwives. It has an obstetrics and gynecology department and until the data collection time, there wasn’t separate room to admit patients with preeclampsia. A hospital-based retrospective cross-sectional study was implemented. All preeclamptic women who were admitted and gave birth at Woldia General Hospital from September 1, 2011- September 1, 2016 were included for the review. Perinatal outcome which was dichotomized as alive or dead was assessed for neonates delivered after 28 completed weeks and through their early neonatal period before discharge (≤ 7 days). As the study was planned to assess the effect of different patterns of preeclampsia management on perinatal death, subjects’ medical files documented with coincidental appearance for possible causes of perinatal death were excluded. This included maternal death on arrival, during admission and labor, and multiple delivery. In addition, grossly incomplete medical files; subjects with the initial diagnosis of preeclampsia in the first-time at the emergency department and immediately delivered were also excluded. The reason for the latter was the author’s assumption that much of the inpatient management scheme that of interest for the present study would have missed and there might also be the probability of misdiagnosis. Figure 1 showed the schematic representation of sampling procedure and excluded cases from each departments (Fig. 1).

Data collection tools and quality assurance
The data were collected using a pre-tested check list which assessed socio-demographic variables, obstetrics history, signs and symptoms at presentation, laboratory results, and perinatal outcomes. The data were obtained from admission registration logbooks, medical and nursing care progress notes, delivery registration books, laboratory and drug prescriptions. The data collectors were trained a 2-days intensive training on how to review and extract the required information from the patients’ chart. They were also monitored by supervisors daily. The tool was pretested at Lalibela General Hospital and an effort was made to validate patients’ data from different folders and registration books.

Study variables
The study outcome variable was a perinatal outcome from preeclamptic women, which was dichotomized as alive and dead. The independent variables were, sociodemographic factors such as patients age and residence; obstetric related factors; gestational age (at admission and delivery), antenatal follow up for current pregnancy; current medical history (comorbidities); conditions at admission such as blood pressure measurement, level of proteinuria and signs and symptoms for end organ involvement, treatment modalities including; anti-hypertensive modalities, anti-convulsion modalities, mode of delivery; and other perinatal related factors such as intra-partum fetal heartbeat, APGAR score and birth weight.

Operational and term definitions
A patient medical file was documented as incomplete if it failed to mention perinatal outcome (as dead or alive), or if greater than 20% of the independent variables were missed. In this study, perinatal death included stillbirths (fetal death inside the uterus after 28 weeks) and early neonatal deaths (deaths within the first week after delivery). Gestational ages reported in the patient’s file was...
considered valid if it was calculated from the last normal menstrual period or using obstetric ultrasound. Fetal heart beat was usually obtained with Pinard fetoscope and a presence of any single value lower 100 bpm or above 160 bpm, classified as abnormal pattern. We have used the routine clinical classification of preeclampsia as mild and severe. Hence, cases of documented blood pressure of more than 140/90 mmHg but less than 160 mmHg systolic or 110 mmHg diastolic with qualitative dipstick proteinuria less than +2 (mild proteinuria) was labeled as mild whereas a persistent systolic blood pressure of >160 mmHg or diastolic pressure of >110 mmHg was labeled as severe.

Fig. 1 Flow chart illustrating the sampling procedure, variables extracted at each department and cases excluded from the study on preeclampsia management modalities and perinatal death at Woldia General Hospital.
mmHg with significant protein excretion of greater than +3 (marked proteinuria) or suggestive evidence for end organ involvement was recorded as severe pre eclampsia. If preeclamptic women reported at least one of these symptoms; headache, visual problem, epigastric pain and placental abruption, platelet count below 100,000/mm$^3$ or aspartate aminotransaminase (AST) > 70 IU/L, the subject was labeled as having signs of end-organ involvement. Co-morbidity status for HIV, diabetes mellitus, chronic hypertension and anemia was assessed as either ‘Yes/No’ based on patient’s self-report or laboratory result. Labor and early neonatal complications were assessed by a positive finding for at least one of the following complications: meconium aspiration, cord prolapse, vasa previa, prolonged labor (more than 24 h.), uterine rupture, neonatal tetanus, severe jaundice, neonatal sepsis, life threatening congenital defect. Women with hemoglobin level less than 11 g/dl was reported as anemic and classified as severe (Hgb < 7.0 g/dl) moderate (Hgb 7.0–9.9 g/dl) and mild (Hgb 10.0–10.9 g/dl). Time-lapse measurements were taken in hours and categorized based on their mean, median, or percentile after checking for their normality.

**Data processing and analysis**

Data were entered and analyzed by Statistical Package for the Social Science (SPSS) version 23 (RRID:SCR_002865). Percentage, rate, frequency distribution, measure of central tendency such as mean and median as well as measure of dispersion like interquartile range and standard deviation were used to describe different variables. Continuous variables such as time measurements were checked for their normality and reported using mean and standard deviation if normally distributed or median and interquartile range (IQR) if skewed. Pearson’s $\chi^2$ test were used to examine association between selected variables. The effect of each variable on perinatal death was first tested by bivariable logistic regression. Confounding bias imposed by independent variables was further controlled by multivariable logistic regression. Association was considered significant when the $p$-value was less than 0.05 and adjusted odds ratio values were reported along with their 95% confidence interval. For a better calculation of the effect of pre eclampsia and its management schemes on perinatal death, the study tried to assess the effect of other possible causes of perinatal death using the variable ‘labor and early neonatal complication’ and controlled with multivariable logistic regression.

**Result**

There were 8764 deliveries attended at Woldia General Hospital in the study period of which 248 were diagnosed as preeclamptic and gave birth in the hospital. Seven medical files were further excluded due to maternal death and incomplete documentation and therefore findings from this study were reported based on 241 reviewed medical documents. The mean maternal age in years ($\pm$SD) was

| Obstetric and medical condition at Admission | mean (SD) |
|--------------------------------------------|-----------|
| Gestational age during admission (in weeks) | 37$^{+6}$ ($\pm$2$^{+5}$) |
| Diastolic blood pressure at admission (mmHg) | 105.87 ($\pm$10.58) |
| Obstetric and medical condition at Admission | n (%) |
| Gravida | n (%) |
| Primigravida | 138 (57.3%) |
| multigravida | 103 (42.7%) |
| Antenatal follow-up for current pregnancy | |
| Yes | 168 (69.7%) |
| No | 73 (30.3%) |
| Comorbidities$^a$ (n = 98) | |
| Yes | 57 (58.2%) |
| No | 41 (41.8%) |
| Dipstick proteinuria (n = 223) | |
| Mild proteinuria | 148 (66.4%) |
| Marked proteinuria | 75 (33.6%) |
| Degree/type of preeclampsia diagnosed | |
| Mild preeclampsia | 99 (41.1%) |
| Severe preeclampsia | 139 (57.7%) |
| Preeclampsia superimposed on chronic hypertension | 3 (1.2%) |
| Sign and symptom of end organ involvement | |
| Headache (n = 228) | |
| Yes | 111 (48.7%) |
| No | 117 (51.3%) |
| Visual problem (n = 229) | |
| Yes | 30 (13.1%) |
| No | 199 (86.9%) |
| Epigastric pain (n = 213) | |
| Yes | 18 (8.3%) |
| No | 195 (91.5%) |
| Abruptio placenta (n = 117) | |
| Yes | 117 (100%) |
| No | 3 (5.2%) |
| Thrombocytopenia (n = 58) | |
| Yes | 55 (94.8%) |
| No | 52 (100%) |

$^a$assessed only for HIV, diabetes, chronic hypertension and anemia
25.73 ± 4.8 with the minimum of 15 years and maximum of 40 years. One hundred thirty-one (54.4%) of them were rural dwellers (Table 1).

The study also assessed labor and early neonatal period complications that lead to possible neonatal death. According to the review, 20 of such complications were documented of which half were meconium aspiration.

**Patterns of preeclampsia management modalities**

The median time lapsed for the patients from onset of their symptoms to arrive to the hospital was 14 h. A wide variability has been observed on the duration of time from admission to the onset of labor among patients diagnosed with severe preeclamptic and preeclampsia superimposed on chronic hypertension. Thus, the mean (±SD) hour was 44.9 (±39.1) with the minimum of 1 and maximum 192 h (8 days) duration (Table 2).

Magnesium sulfate was the most frequently (69.3%) used anticonvulsant medication used followed by diazepam (26.6%). The most frequently used antihypertensive

| Time lapses                                                                            | Median (interquartile range) in hours |
|----------------------------------------------------------------------------------------|---------------------------------------|
| Time lapsed from onset of current symptoms to reach hospital (n = 236)                | 14 (6.25–24)                          |
| Time lapsed from admission to beginning of MgSO4a (n = 139)                           | 1 (0.5–1)                             |
| Time lapsed from admission to beginning of antihypertensive medicationb (n = 101)     | 0.5 (0.28–1)                          |
| Time lapsed from admission to onset of laborc (n = 141)                               | 24 (10.25–48)                         |

*a*For patients with severe preeclampsia and preeclampsia superimposed on chronic hypertension

*b*For patients with DBP ≥ 110 mmHg

*c*For women with severe preeclampsia and preeclampsia superimposed on chronic hypertension

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**Fig. 2** Trend of convulsion management modalities among inpatient preeclamptic women admitted at Woldia General Hospital from 2011 to 2016.
medications were hydralazine, methyldopa and nifedipine that were used 112, 111 and 54 times respectively. Anticonvulsant medication was given for 221 (91.7%) during labor and its use continued for 24 h postpartum or since last convulsion for 177 (73.4%) patients. Out of 239 patients, 148 (61.9%) has commenced their labor spontaneously while it was intentionally induced for 91 (38.1%) patients. Spontaneous vaginal delivery was the major route of delivery 114 (47.5%) while 82(34.2%), 35(14.6%), and 9(3.75%) of patients managed by vacuum, caesarean section and forceps delivery respectively. Comparing the route of delivery with severity of the disease (classified as mild or severe preeclampsia), the use of vacuum delivery and caesarean section were at a statistically higher frequency among severe preeclamptic women over their counterparts \( \chi^2(3, n = 117) = 2.398, p = 0.013 \), while no significant differences were observed in the frequency of spontaneous vaginal delivery \( (p = 0.613) \) (Fig. 2).

As shown in the Fig. 2, magnesium sulfate was the drug of choice to prevent convulsion in preeclamptic women. Its use increased from 25% in 2011/12 to 93% in 2015/16 \( (P < 0.0005) \). Opposite to this, there was a decreasing trend for using diazepam from 54% in 2011/12 to 1.6% in 2015/16 \( (P < 0.0005) \). In addition, extending the use of anticonvulsant medication 24 h postpartum or after the onset of last convulsion increased across the 5 years.

### Perinatal outcome
The mean gestational age in weeks and days at birth was \( 37^{\pm 6} (\pm 2^\pm 5) \) and the minimum and the maximum value of 24 and 42 completed weeks respectively. The mean value for 1st and 5th minute APGAR score was 5.98 (±2.36) and 6.91 (±2.67) respectively. The mean value for birth weight was 2608.02 (± 558.8) gram. Low birth weight was more frequently observed among severe preeclampsia than the mild one (53.7% Vs 32%) \( p = 0.001 \). Out of 225 documents reviewed for fetal heart rate abnormalities, 24 (10.7%) abnormal patterns were documented. Forty-Seven perinatal deaths were recorded across the 5-year period yielding a case specific perinatal mortality rate 197/1000 births. Of this, 27(57.4%) were documented as stillbirth and the rest 20 (42.6%) were within the first week of their hospital stay. Perinatal death was higher (76.59% Vs 23.4%) among offspring from severe preeclampsia patients than their mild counterparts \( (p = 0.01) \) (Fig. 3).

Using logistic regression, important independent variables were checked for their possible association both in bivariable and multivariable logistic regression with a \( p \)-value less than 0.05. Three variables were significantly associated with perinatal death both in bivariable and multivariable logistic regression; namely intrapartum fetal heart rate pattern, manifestations of end organ involvement, and 5th minute APGAR score. It was observed that the odds of perinatal death among those with manifestation of end-organ involvements were 4
times (AOR = 4.404 95% CI (1.219–6.370)) higher than their counterparts (Table 3).

We found that a group of conditions operationalized as 'labor and early neonatal complications' increased the odds of perinatal death nearly by 5-fold. However, this variable is an independent cause of perinatal death even in the absence of preeclampsia. Therefore, the multivariable logistic regression was repeated in which 'labor and early neonatal complications' and other independent variables were entered into the regression model with separate blocks. From the final analysis, it was found that the diastolic blood pressure measurement and 5th minute APGAR score were associated with perinatal death. Accordingly, the odds of perinatal death was nearly 3-times (AOR = 2.824; 95% CI (1.154–6.038)) higher among women who had diastolic blood pressures greater than 110 mmHg taken at admission compared to those who had below 110 mmHg. Neonates whose 5-min APGAR score was between 0 to 3 has a 5-fold (AOR = 5.046; 95% CI (1.856–8.235)) greater risk of death than neonates whose 5-min APGAR was between 7 and 10.

**Discussion**

The time-lapsed from onset of symptoms to the patient reaching health care support is a combined reflection of patient’s health-care seeking behavior, access to health care, and the spread of disease. The time from the onset of symptoms to MgSO₄ initiation, antihypertensives, and labor onset are key indicators of the patient’s health-seeking behavior, health-care access, and the spread of disease, respectively. These time-lapses were analyzed in this study.

**Table 3** A bivariable and multivariable logistic regression analysis result testing for factors associated with perinatal outcome (death/alive) among neonates delivered from preeclamptic women from 2011 to 2016 (before the effect of labor and early neonatal complication controlled)

| Variable                                                       | COR, (95% CI)     | p-value | AOR, (95% CI)     | p-value |
|---------------------------------------------------------------|-------------------|---------|-------------------|---------|
| **Diastolic Blood pressure measurement** (n = 231)            |                   |         |                   |         |
| Less than 110 mmHg                                            | 1                 | 0.009   | 1                 | 0.335   |
| ≥ 110 mmHg                                                    | 3.688 (1.257–4.863) | 0.012   | 4.404 (1.219–6.370) | 0.035*  |
| **Manifestation of end-organ involvement**                    |                   |         |                   |         |
| Yes                                                           | 4.667 (1.332–8.001) | 0.012   | 4.404 (1.219–6.370) | 0.035*  |
| No                                                            | 1                 |         | 1                 |         |
| **Gestational age at birth**                                   |                   |         |                   |         |
| < 37 completed weeks                                          | 2.547 (1.073–4.021) | 0.030   | 1.105 (0.098–2.111) | 0.315   |
| ≥ 37 completed weeks                                          | 1                 |         | 1                 |         |
| **Time lapsed from admission to initiation of MgSO₄** (n = 139) |                   |         |                   |         |
| Less than 1 h                                                 | 1                 |         | 1                 |         |
| More than 1 h                                                | 1.254 (0.343–2.164) | 0.751   | 2.780 (0.558–5.002) | 0.427   |
| **Time lapsed from admission to initiation of antihypertensives** (n = 101) | | | | |
| Less than 17 min                                              | 2.962 (0.331–5.593) | 0.669   | 3.636 (0.231–7.014) | 0.125   |
| 17–30 min                                                    | 3.436 (0.479–6.392) | 0.397   |                   |         |
| 30 min to 1 h                                                | 2.331 (0.195–4.467) | 0.931   |                   |         |
| More than or equal to 1 h                                    | 1                 |         | 1                 |         |
| **Time lapsed from admission to onset of labor** (n = 141)    |                   |         |                   |         |
| Less than 35 h                                                | 1                 |         | 1                 | 0.412   |
| More than or equal to 35 h                                    | 1.353 (0.235–2.452) | 0.644   | 1.951 (0.896–3.006) | 0.623   |
| **Qualitative proteinuria** (n = 223)                         |                   |         |                   |         |
| Mild proteinuria                                             | 1                 | 0.776   | 1                 |         |
| Marked proteinuria                                           | 1.395 (0.547–2.243) | 0.159   | 1.159 (0.113–2.091) | 0.221   |
| **MgSO₄ continues 24 h after delivery** (n = 210)             |                   |         |                   |         |
| Yes                                                           | 1                 | 0.024   | 1                 |         |
| No                                                            | 3.132 (1.048–4.168) | 0.191   | 4.069 (0.896–6.347) | 0.112   |
| **Intrapartum Fetal heart rate** (n = 225)                     |                   |         |                   |         |
| Normal                                                       | 1                 | 0.014   | 1                 | 0.112   |
| Some abnormal records                                        | 6.328 (2.069–10.587) | 0.011   | 4.846 (0.667–9.025) | 0.017*   |
| **Onset of labor**                                            |                   |         |                   |         |
| Spontaneous                                                  | 1                 |         | 1                 | 0.119   |
| Induced                                                      | 5.580 (2.237–9.293) | 7.477   | 0.633–14.322      | 0.0005*  |
| **Labor and early neonatal complications** (n = 239)          |                   |         |                   |         |
| Yes                                                          | 3.454 (1.321–5.588) | 0.006   | 4.616 (1.365–7.867) | 0.021*   |
| No                                                           | 1                 |         | 1                 |         |
| **Fifth minute APGAR score** (n = 214)                        |                   |         |                   |         |
| 0–3                                                         | 13.962 (6.499–21.426) | 0.011   | 5.046 (1.856–8.235) | 0.021*   |
| 4–6                                                         | 7.645 (2.629–12.660) | 0.003   | 3.112 (1.223–5.001) | 0.021*   |

Key: COR: Crude odd ratio

*Show association with p value less than 0.05

*a analysis was done after excluding stillbirths

*b For patients with severe preeclampsia and preeclampsia superimposed on chronic hypertension

*c For patients with DBP > 110 mmHg

*d For women with severe preeclampsia and preeclampsia superimposed on chronic hypertension
care, infrastructure and transportation. Regarding the management modalities used, we believed that in addition to comparing this study finding with others', discussing our findings with national guideline and protocols will help to identify the magnitude of management gap.

In previous research, the rate of perinatal mortality among those mothers with preeclampsia ranged from 47/1000 to 416/1000 live births [5–8]. The perinatal mortality rate recorded in our study (197/1000 live births) was within this range but slightly higher than two studies with rates of 120/1000 and 160/1000 deaths per live births [2, 12]. Differences in death rates might arise from difference in the gestational age cut off points used among studies, the type of hypertensive disorders included, and service-delivery capacity of the hospitals where the patients were managed.

Similar to other studies, in this sample, 47.5% of all preeclamptic women had vaginal births [14, 15]. Vaginal delivery is also the preferred route of delivery for preeclamptic women according to the Ethiopian Minister of Health protocol [16]. Rates of labor induction vary a great deal across studies. Similar to a rural hospital in Western Tanzania, in this study labor was induced in 38.1% of the patients [17]. In contrast, 47.9% of patients were induced at the Mettu Karl Referral Hospital [12] and 23.4% in governmental hospitals of Addis Ababa [14]. The possible explanation for this difference could be the fact that labor induction has a variety of chief indications other than preeclampsia.

In three studies, usage of magnesium sulfate in preeclamptic women to prevent seizures ranged from 74.1 to 100% [12, 14, 17] similar to our finding of (91.7%). In addition to this, this study revealed that anti-convulsant medication was continued for 24 h postpartum for 177 (73.4%) patients. Although this finding was consistent with the aforementioned studies conducted in resource-limited areas, it was lagged behind to achieve the Ethiopian Ministry of Health management protocol which recommends the use of magnesium sulfate for all preeclamptic woman during labor. It’s also recommended that magnesium sulfate treatment should continue for 24 h postpartum [16, 18]. Unavailability of the drug or failure either to give the drug or to document could be the possible explanation for this discrepancy.

The median duration for women with severe preeclampsia between admission and labor was 24 h (IQR 10.25–48) which was shorter than other study conducted among patients with severe preeclampsia and eclampsia [17]. Shorter hospital stay for severe preeclamptic women at term was stated as a supported management option in studies and management protocols [16, 19].

Similar to our finding, other studies in low-resource settings reported more stillbirths than deaths at early neonatal period [15, 20, 21]. We found that women who had diastolic blood pressures above 110 mmHg had a 2-fold increased chance of perinatal death compared to those with lower diastolic blood pressures. Similar association has been also detected in other studies [22, 23].

Limitation
As a limitation, this study didn’t assess the effect of steroid drugs on perinatal death among preeclamptic women as only few numbers of cases received the treatment.

Conclusion
In conclusion, perinatal death was high among inpatient preeclamptic women who gave birth at Woldia General Hospital. The trend of using magnesium sulfate as an anti-convulsant increased through the 5-year period. However, there was a gap to extend its use up to 24 h postpartum or since the last convulsion as per the national protocol. From our analysis, diastolic blood pressure and 5th minute Apgar score were found associated with perinatal death. Authors recommended that future large-scale health service auditing and impact assessment studies in such resource limited-areas might serve as an input for policymakers to improve the quality of clinical care and to decrease mortality rates.

Abbreviations
APGAR: Appearance, Pulse, Grimace, Activity, Respiration; HDP: hypertensive disorders of pregnancy; Hgb: Hemoglobin

Acknowledgments
We would like to thank Woldia University for the grant allocated to conduct this study. We also extend our deepest gratitude to Woldia General Hospital Administrative bodies and staffs working at the hospital archive. Last but not the least, we would like to extend our gratefulness and heartfelt thanks to all our colleagues who gave us their unduly support, critical comments and helpful information.

Authors’ contributions
KG and ML conceptualized the study, its design, analysis, interpretation, wrote the manuscript. YA, MM and MW participated in the design, analysis of study findings and interpretation of its result. All the authors read and approved the final manuscript.

Funding
This study was funded by Woldia University with grant number WURCSVP/209/ 2017. The funding body didn’t have any role in the design of the study and collection, analysis, and interpretation of data and in writing the manuscript.

Availability of data and materials
The datasets used for analysis during the current study are available from the corresponding author on reasonable request.

Ethics approval and consent to participate
Ethical clearance was obtained from institutional review board of Woldia University. Permission was also obtained from the hospital administrative office to retrieve patients’ data from their archive. Data collectors took training how to use patients’ information confidentially. Since data were extracted by document review, written or oral consent from patients was not applicable.

Consent for publication
Not applicable.

Competing interests
The authors declare that they have no competing interests.
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Received: 20 December 2019 Accepted: 29 March 2020
Published online: 09 April 2020

References

1. Central Statistical Agency Addis Ababa Ethiopia, ICF International Calverton Maryland, USA. Ethiopia Demographic and Health Survey 2011. In: CSA, editor. Addis Ababa and Calverton; 2012.

2. Lawn JE, Blencowe H, Waiswa P. Stillbirths: rates, risk factors, and acceleration towards 2030. Lancet. 2016;387(6):17.

3. Vogel JP, Souza JP, Mori R. Maternal complications and perinatal mortality: findings of the World Health Organization multicountry survey on maternal and newborn health. BJOG. 2014;121(1):13.

4. Lawn JE, Blencowe H, Oza S. Every newborn: progress, priorities, and potential beyond survival. Lancet. 2014;384(9966):17.

5. Singhal SR, Deepika A, Nanda S. Maternal and perinatal outcome in severe preeclampsia and eclampsia. JSAFOG. 2009;13(3):8.

6. Vidhyadhar BB, Purushottam AG, Aditi SM. Maternal and foetal outcome in pregnancy induced hypertension: a study from rural tertiary care teaching hospital in India. Int J Biomed Res. 2011;2.

7. Tuffnell DJ, Jankowicz D, Lindow SW, Lyons G, Mason GC, Russell IF. Outcome of severe preeclampsia/ eclampsia. J Obstet Gynecol. 2005;112:6.

8. Al Inizi ST, Sharara H, Ahmed B. Eclampsia in Quatar: maternal and fetal outcomes, possible preventive measures. Mid East J Emerg Med. 2005;5:6.

9. Knight M. Eclampsia in the United Kingdom 2005. BJOG. 2007;114(9):8.

10. Onuh SO, Aisien AO. Maternal and fetal outcome in eclamptic patients in Benin City, Nigeria. J Obstet Gynaecol. 2004;24(7):8.

11. Wolde Z, Seghi H, Woldie M. Hypertensive disorders of pregnancy in Jimma University specialized hospital. Ethiop J Health Sci. 2010;21:147–54.

12. Seyom E, Abera M, Tesfaye M, Fernahun N. Maternal and fetal outcome of pregnancy related hypertension in Mettu Karl referral hospital, Ethiopia. J Ovarian Res. 2013;8.

13. Pannu D, Das B, Hazari P, Shilpa. Maternal and perinatal outcome in eclampsia and factors affecting the outcome: a study in North Indian population. Int J Reprod Contracept Obstet Gynecol. 2012;3:347–51.

14. Wagnew M, Desalegn M, Wonku A, Nyagero J. Trends of preeclampsia/eclampsia and maternal and neonatal outcomes among women delivering in Addis Ababa selected government hospitals, Ethiopia: a retrospective cross-sectional study. Pan Afr Med J. 2016;25(2):12.

15. Nazarius PK, Turkwesigye M, Wandabwa J, Wamuyu-Maina G, Bimenya GS, Okong P. Adverse neonatal outcomes in women with preeclampsia in Mulago Hospital, Kampala, Uganda: a cross-sectional study. Pan Afr Med J. 2009;6:17.

16. Health Federal Democratic Republic of Ethiopia Ministry of Management protocol on selected obstetrics topics. Pregnancy Induced Hypertension 2010:22.

17. Mooij R, Lugumila J, Mwashambwa MY, Mwamposa M, van Dillen J, Stekelenburg J. Characteristics and outcomes of patients with eclampsia and severe pre-eclampsia in a rural hospital in Western Tanzania: a retrospective medical record study. BMC Pregnancy Childbirth. 2012;15.

18. Sibai BM. Magnesium sulfate prophylaxis in preeclampsia: Lessons learned from recent trials. Am J Obstet Gynecol. 2004;190.

19. Barber D, Xing G, Towner D. Expectant management of severe eclampsia between 24–32 weeks gestation: a ten-year review: The 29th Annual Meeting of the Society for Maternal-Fetal Medicine; 2009.

20. Endeshaw G, Berhan Y. Perinatal Outcome in Women with Hypertensive Disorders of Pregnancy: a retrospective cohort study. Int Scholarly Res Notes. 2013;2015:18.

21. Ananth CV, Basso O. Impact of pregnancy-induced hypertension on stillbirth and neonatal mortality in first and higher order births: a population- based study. NIH Public Access. 2010;21:118–23.

22. Judy A, Chao L, Gisern A, O’Malley K, Lyell D, Blumenfield Y, Butwick A, El-Sayed Y. Maternal characteristics and neonatal outcomes in women with eclampsia versus severe preeclampsia. Am J Obstet Gynecol. 2014;192(1):5188–90.

23. Thangaratnam S, Datta A, Ismail KMM, Khan KS. What is the accuracy of blood pressure in predicting complications in pre-eclampsia? Arch Dis Child Fetal Neonatal Ed. 2011;96(1):9.