An Unusual Case of Eclampsia at 21 Weeks of Gestation with Multiple Risk Factors Except Molar Pregnancy

Yırtma Birinci Gebelik Haftasında Gelişen Molar Gebelik Dışında Birçok Risk Faktörüne Sahip Nadir Bir Eklampsi Olgusu

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ABSTRACT

Preeclampsia/eclampsia is a serious complication of the pregnancy and early-onset disease is mostly associated with molar pregnancy. We present a rare case of eclampsia that occurred at very early weeks of gestation and had multiple risk factors for preeclampsia, but not molar pregnancy. A 35-year-old woman (gravida 11, parity 1) presented with eclampsia at 21 weeks of gestation. She had history of preeclampsia in previous pregnancy, chronic hypertension, recurrent pregnancy loss (a total of 9 abortions at 6-21 weeks of gestation) and methyltetrahydrofolate reductase (MTHFR) heterozygous mutation. The pregnancy was terminated by hysterotomy. Molar change was not observed in the placenta. No postoperative complication occurred and the patient was discharged with antihypertensive therapy. Eclampsia is a severe form of the preeclampsia and clinicians should keep in mind that it may be occurred at very early weeks of gestation in patients with multiple risk factors for preeclampsia.

Key Words: Eclampsia, gestational age, risk factors

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CASE REPORT

Thirty five–year-old woman (gravida 11, abortion 9, and parity 1) referred to the emergency service at 21 weeks of gestation with the complaints of unconsciousness and generalized tonic-clonic seizure two times. She had history of recurrent pregnancy loss (a total of 9 abortions at 6, 6, 16, 16, 19, 19, 21 and 21 weeks of gestation), and chronic hypertension. Despite the use of antihypertensive drugs, unregulated chronic hypertension was present in her all pregnancies. She had also history of preterm delivery with cesarean section due to severe preeclampsia at 26 weeks of gestation. Screening for thrombophilia was carried out before the current pregnancy and methyltetrahydrofolate reductase (MTHFR) heterozygous mutation was detected.
According to her medical records, she had no renal disease, diabetes mellitus, antiphospholipid antibody syndrome and other chronic autoimmune diseases. She was using metildopa, low molecular weight heparin and aspirin since the first trimester of the current pregnancy. Her blood pressure was 170/100 mmHg at admission, fetal biometry was consistent with 21 weeks of gestation. In physical examination, confusion, pretibial edema, and periorbital ecchymosis due to trauma during seizure were observed. In laboratory findings, hemoglobin level was 14 gr/dl, platelet count was 368,000 cell/mm$^3$, alanine aminotransferase (ALT) level was 21 IU/L, aspartate aminotransferase (AST) level was 26 IU/L, creatinine level was 0.9 mg/dl and there was 3+ proteinuria. Intravenous hydralazine and 2 gr/h magnesium sulphate were administered for blood pressure regulation and seizure control. The pregnancy was terminated by hysterotomy due to the eclampsia indication. Fetal weight was 300 gr and no major congenital anomaly was observed. There was not hydatiform change or another pathological finding in the placenta. No complication occurred and the patient recovered consciousness and blood pressure was discharged with oral amlodipine treatment.

DISCUSSION

Preeclampsia/eclampsia at early gestational weeks is rare and serious complication of pregnancy and mostly associated with molar pregnancy (3). A literature research revealed few published case reports of eclampsia without molar pregnancy at < 22 weeks of gestation (4-6). In these reports, 1 case had older age (4) and 2 had primigravidity alone (5,6) as a risk factor. In 1985, a 43 year old eclampsia case at 18 weeks of gestation was reported (4). In another report, a first pregnancy complicated with eclampsia at 21 weeks of gestation was terminated with prostaglandin E2 suppositories successfully (5). Gürel and Gürel (6) were also reported the eclampsia and HELLP syndrome occurred in an 18 year old, primigravida woman at 19 weeks of gestation. Unlike the previous reports, our case had multiple risk factors together: history of chronic hypertension, previous preeclampsia, recurrent pregnancy loss, and MTHFR heterozygous mutation. In all case reports including ours, patients recovered well and no neurologic or any other sequel were recorded by authors.

Presented case had a history of delivery with cesarean section due to severe preeclampsia at 26 weeks of gestation. The recurrence risk in subsequent pregnancy was 32% in patients with history of severe preeclampsia at second trimester (7). In addition, superimposed preeclampsia occurs in 40% of women with chronic hypertension (2) and three or more early pregnancy loss increases the risk of preeclampsia (8). Recurrent pregnancy loss and preeclampsia share similar etiological factors and histopathological findings in spiral arterioles (9).

Hereditary thrombophilia is also a risk factor for preeclampsia. Livinova et al (10) reported that the relative risk in patients with preeclampsia for MTHFR heterozygous mutation, and MTHFR homozygous were 1.7 and 2.73.

As a conclusion; presented case had multiple risk factors for preeclampsia. Clinicians should keep in mind that the presence of many risk factors together may lead to occurrence of severe disease and eclampsia at the early weeks of gestation, and should inform patients about the adverse maternal and perinatal outcomes of severe preeclampsia.

Conflict of interest
No conflict of interest was declared by the authors.

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