LETTER TO THE EDITOR

Large Subchorionic Hematoma: Breus’ Mole

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Abstract

A case of large subchorionic hematoma complicated by intrauterine growth retardation and oligohydramnios diagnosed at 32 weeks’ gestation with twin pregnancy after ICSI is reported below. The patient was on clexane injection during pregnancy for mitral valve replacement. She was managed with tocolysis using progesterone therapy and antibiotic with followup until delivery. At 34 weeks, a male baby weighing was delivered without complication by caesarean section because of single fetal demise of twin pregnancy.

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Introduction

Large subchorionic hematoma, which was first described by Breus [1] in 1892 (Breus’ mole), is a serious condition that is frequently complicated by intrauterine growth retardation (IUGR) and intrauterine fetal death (IUFD). We encountered a case complicated by IUGR and oligohydramnios diagnosed at 32 weeks’ gestation in twin pregnancy after ICSI on anticoagulant.

Case report

A 30 year-old, G2P0+1, history of previous one hysterotomy, was admitted to our hospital for ANC. There was no complaint. She was about 28 weeks of gestation with twin pregnancy after ICSI. She was hypothyroid on L-Thyroxine and with mitral valve replacement on clexane injection of 80 mg every 12 h and PGDM on insulin. Ultrasonographic examination revealed twin dichorionic diamniotic pregnancy with IUGR of first baby with oligohydramnios. A large homogenous mass about 14 cm in dimensions was detected beside the fetal placenta of the IUGR fetus (Fig. 1). No blood flow was detected in the mass on color Doppler. The patient was admitted for tocolysis with progesterone and follow-up delivery with antibiotic. The patient was managed conservatively for 4 weeks, where IUGR fetus died. A male baby was delivered without complication by caesarean section after steroid therapy for lung maturation at 32 weeks. During removal of the placenta, large amount of dark brown blood with no clots was noted.

This hematoma might be related to minor trauma with high dose of anticoagulant therapy.

Discussion

It is a rare disease, with an incidence of 1 in 1200 placentas. Women with cardiac problems, disorders of circulation,
monosomy, hypertension and diabetes are predisposed to Breus’ mole. The mole is formed as a subchorionic hematoma, formed out of the intervillous blood, causing progressive accumulation of the clotting substance called fibrin with increasing gestational age [2].

Breus mole is reported to be found in the placenta of macerated stillborn foetuses, indicating that massive subchorionic hematoma could have been the cause of their demise. A massive Breus’ mole can cause disturbances in blood flow in the spiral arteries and might result in intrauterine growth restriction of the foetus [3].

Although the reported frequency of subchorionic hematoma is very low, some cases of IUGR associated have been investigated [4–6]. The mechanism of IUGR is due to utero-placental insufficiency. The location of hematoma has been suggested to impede the cord insertion [5,7]. The hematoma can easily compress the umbilical vessels. Tocolysis and antibiotic therapy are common for the management because likely to result in abortion or premature delivery associated with chorioamnionitis [8]. In 1991, Baxi and Pearlstone [9] suggested that antibodies are an etiologic factor for subchorionic hematoma. The antibodies have been suggested to increase the platelets to aggregate, which leads to thrombosis and vasculitis. It is a condition in which a large amount of blood mainly of maternal origin collects and separates the chorionic plate from the villous chorion [6].

Its etiology and pathogenesis remained a dilemma in the 18th and early 19th century. The term Breus’ ‘Mole’ does not actually mean molar degeneration as seen in hydatiform moles, but is related to “mass” or coagulated blood. Their significance depends not on their size, but on their site of appearance. For eg: if near cord insertion they may lead to cord compression and decreased fetal perfusion, fetoplacental insufficiency or hematoma dissected into base of the umbilical cord causing umbilical venous obstruction.

Author’s contribution

All the authors contributed to protocol development, data collection and management, Data analysis and Manuscript writing/editing.

Ethical disclosure

The authors declare that the procedures followed were in accordance with the regulations of the relevant clinical
research ethics committee and with those of the Code of Ethics of the World Medical Association (Declaration of Helsinki).

Confidentiality of data

The authors declare that they have followed the protocols of their work center on the publication of patient data.

Right to privacy and informed consent

The authors have obtained the written informed consent of the patients or subjects mentioned in the article. The corresponding author is in possession of this document.

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