Haemoperitonium : A Rare Presentation of Choriocarcinoma

Abstract
Gestational choriocarcinoma is one of the most malignant forms of group of tumors with almost 100% cure rate, develops secondary to pregnancy (Normal or ectopic) following hydatidiform mole, miscarriage or even after child birth. This starts within the uterus in the placenta but a tendency to metastasize to other parts of the body. Indicative signs are unexplained heavy and irregular uterine bleeding, raised serum \( \beta \) hCG, non normalization of \( \beta \) hCG following abortion or delivery and if it spreads, signs of metastasis. Our study presents a case of Choriocarcinoma presented with haemoperitonium, who was treated successfully with surgical and medical management. A 25 years lady Para 1+0, age of 3 years, normal delivery had an emergency admission for 2 months amenorrhoea with severe lower abdominal pain, slight vaginal bleeding suspected clinically as ruptured ectopic pregnancy. Ultrasonography suggest Invasive mole with huge peritoneal collection. After sending serum \( \beta \) hCG she was treated surgically and per operative findings were in favour of Choriocarcinoma which was confirmed by histopathological reports. Serum \( \beta \) hCG was raised and she was treated with combination (MAC protocol) chemotherapy.

Key words : Haemoperitonium; Miscarriage; Choriocarcinoma, Histopathology.

INTRODUCTION
Choriocarcinoma also known as trophoblastic malignancy is a rare form of cancer with almost 100% cure rate. The incidence of choriocarcinoma is estimated to be 0.133 per 100000 woman years. Majority of cases of choriocarcinoma arise due to malignant transformation of complete mole, although it has been reported following abortion, normal pregnancy, ectopic pregnancy or following childbirth. It is known to occur 1 in 5333 of tubal pregnancy and 1 in 1.6 million of normal intrauterine pregnancy. It can develop anytime between 5 weeks and 15 years after gestation or even after menopause. This highly malignant tumour usually starts in the uterus but can rapidly spread to other parts of body. Approximately 30% cases have metastatic disease at the time of diagnosis. Lungs (80%) are the most common sites of metastasis followed by vagina (30%) liver (10%) and involving brain in 28% of patients. So before starting treatment of molar pregnancy \( \beta \) hCG (Human Chorionic Gonadotrophin) and chest X Ray should be done. The leading symptom is heavy and irregular uterine hemorrhage sooner or later expulsion of mole or any recognized form of pregnancy. The other symptoms are vaginal bleeding, anemia, hyperemesis, hyperthyroidism, uterine and ovarian enlargement and symptoms and signs of metastasis. Because of these variable presentations, it is a diagnostic challenge. Although choriocarcinoma is very aggressive in nature, its cure rate is very high because it is extremely sensitive to chemotherapeutic agents. So early diagnosis and prompt initiation of treatment is a well known determinant of prognosis of choriocarcinoma. The cause of gestational trophoblastic disease is claimed to be genetic in origin. No environmental etiological factor has been implicated apart from deficient Vitamin A precursor carotene in diet.
CASE REPORT
A 25 years lady para 1+0 had age of 3 years, normal delivery, took an emergency admission due to 8 weeks amenorrhoea with severe lower abdominal pain and scanty vaginal bleeding and gives history of excessive vomiting and urinary pregnancy test positive at home one week prior to her admission. She was moderately anaemic, abdomen was distended and uterus was palpable about 14 weeks size. Nothing was abnormal on vaginal examination, anterior fornix was free. All the features were in favour of ruptured ectopic pregnancy only exception was hyperemesis and enlarged uterus. Emergency Ultrasonography was done which suggested Invasive mole with peritoneal collection then after counselling, informed written consent was taking blood sent for hCG, grouping and crossmatching, emergency laparotomy was done. On opening it was full of blood, uterus was about 14 weeks size, firm with multiple hemorrhagic and necrotic spots with bleeding from serosal surfaces, total abdominal hysterectomy was done, ovaries and rest of the pelvic organs were normal. On cut section uterus was congested with polypoid growth with hemorrhage and necrotic spots with a clinical suspicion of choriocarcinoma. Histopathological reports confirms the diagnosis of choriocarcinoma. hCG level was raised (12920mIU/ml ) which is a bit lower range for Choriocarcinoma. Post operative X Ray chest was done and it was normal. with the consultation of Oncologist she completed the course of combination chemotherpy. The patient was withstand well with surgery and had 3 successive normal levels of serum hCG.

DISCUSSION
choriocarcinoma almost always develops with or follows some form of recognized pregnancy, most often following Hydatidiform mole but may follow abortion, ectopic pregnancy, normal pregnancy or even after delivery. So before starting treatment of Hydatidiform mole or in case of excessive undiagnosed bleeding following any recognized pregnancy, serum hCG, X Ray chest, liver enzymes, CBC should be done. The clinical presentation of Choriocarcinoma can be from disease locally in the uterus or from distant metastasis that can cause a wide variety of symptoms. The outcome of choriocarcinoma are related to staging and risk assessment (Low risk, High risk) and not to the antecedent pregnancy. In our study the patient presented with the symptoms and signs mimicking ruptured ectopic pregnancy, the only finding in favour of Choriocarcinoma was hyperemesis and enlarged uterus. The clinical condition of the patient and ultrasonographic diagnosis of invasive mole with haemoperitonium demanded emergency laparotomy and the per operative findings were in favour of Choriocarcinoma, which was confirmed by raised serum hCG and histopathology report. The Choriocarcinoma is functional tumor and usually secretes hCG in large quantity and this causes luteinization with cyst formation in ovaries. Occasionally hCG level may be low when it is composed of non secretory elements as in our patient. Non gestational choriocarcinoma also occur and it is resistant to therapy.

CONCLUSION
A complete knowledge regarding variable clinical presentation of Choriocarcinoma helps in early diagnosis and decrease the adverse outcome of this rare and highly malignant but curable malignancy. So every pregnancy following missed period with any excessive and abnormal uterine bleeding or any abnormal presentation should be evaluated by serum hCG and ultrasonography and histopathology.

DISCLOSURE
Both the authors declared no competing interest.

REFERENCES