Value of Serial Ultrasounds in Early Diagnosis and Management of Prerupture Ovarian Ectopic Pregnancy: A Case Report

Cau Van Vo1*, Carol A Major2 and Kamini Malhotra3
1Department of Obstetrics and Gynecology, Division of Clinical Education, Midwestern University, Glendale, AZ 85308, Fountain Valley Regional Hospital and Medical Center, Fountain Valley, CA 92708, USA
2Department of Obstetrics and Gynecology, Maternal-Fetal Medicine Division, University of California, Irvine, CA 92668, USA
3Department of Pathology, Fountain Valley Regional Hospital and Medical Center, Fountain Valley CA 92708, USA

Abstract

Background: Primary ovarian pregnancy is a rare form of ectopic pregnancy. Even with the advance of ultrasound techniques and the sonographic establishment of fetal development, most ovarian pregnancies are found ruptured at the time of diagnosis or are a surprise finding at the time of surgery. In these cases, the associated hemorrhage is usually heavy and in approximately 23% of cases, the patients develop hypovolemic shock requiring blood transfusion.

Case: A 30-year-old female patient at 7.4 weeks by a certain LMP presented for prenatal care. An ultrasound at the time of presentation revealed no intrauterine gestational sac. There was however, a simple cystic mass contiguous with the left ovary. Subsequently, the patient had a BhCG level drawn which was not consistent with the ultrasound image and therefore was felt to be abnormal. A primary ovarian pregnancy was suspected after serial ultrasounds revealed an enlarging cystic ovarian mass and serial quantitative human Chorionic Gonadotropin (hCG) levels were abnormal. In addition, the patient started feeling left lower quadrant abdominal discomfort and reported a scant amount of vaginal bleeding. Due to the suspicion of a possible ovarian ectopic pregnancy, a diagnostic laparoscopy was discussed with the patient. The suspicion was solidified, in order to avoid rupture, a diagnostic laparoscopy was made and the ovarian ectopic pregnancy was confirmed. A wedge resection of the ovarian cyst was performed and the final histopathology confirmed a primary ovarian pregnancy.

Conclusion: Ultrasounds have been proven safe for management of obstetrical patients. The sonographic parameters for fetal development from conception to delivery have been well established and published.2 At this time, ultrasound is the best tool in detecting abnormal fetal development in pregnancy. In this case, serial ultrasound examinations in the early stage of the first trimester and inappropriately rising serial BhCG levels provide the clues for detecting the prerupture diagnosis of ovarian ectopic pregnancy. The combination of these two modalities and pelvic examination increase the suspicion of a diagnosis of an ovarian pregnancy and will lead to a prompt intervention to prevent morbidity, mortality and preservation of future fertility.

Introduction

Primary ovarian pregnancy is an uncommon event making early diagnosis prior to its rupture very difficult. The incidence of ovarian pregnancies is estimated to be less than 1% of all ectopic pregnancies [1].

Even with the advances of ultrasound and the sonographic establishment of fetal development, most ovarian pregnancies are found to be ruptured at the time of surgery. The diagnosis is usually established after the fact, by pathological examination. Therefore, the diagnosis of an unruptured ovarian pregnancy is a big challenge for all obstetricians.

Serial ultrasounds early in the first trimester may be the most important factors contributing to the diagnosis of prerupture ovarian pregnancy. The combination of serial ultrasounds and serial serum hCG levels may lead to an accurate early diagnosis and subsequently may lead to successful minimally invasive intervention.

The following report is a case of primary unruptured ovarian pregnancy diagnosed with the use of serial ultrasounds and serial beta-hCG evaluations.

Case Report

A 30-year-old primigravid patient, presented at 7.4 weeks gestation, by her last menstrual period. The patient’s medical history was unremarkable. She had no previous sexually transmitted infections or pelvic surgeries. She also had no history of dysmenorrhea and no previous complaints of any type of abdominal or pelvic pain. Her menstrual cycle was regular. At her first prenatal examination, there were no significant physical or pelvic findings.

Her initial ultrasound showed no intra-uterine gestational sac; however, there was a cystic mass (41×39 mm) with a hyperechoic ring and a lucency in the center at the left ovary (Figure 1). The hCG was subsequently ordered and the patient’s initial beta-hCG level was 1,656.13 milli-international units/ml (mIU/ml) [2]. A repeat ultrasound and hCG four days later showed no significant change in the size of the cystic mass. There was however, a newly visualized light echoic area under the cyst wall. In addition, the repeat beta-hCG level was 1,714.29 mIU/ml [2]. A third ultrasound three days later demonstrated a significant increase in the echoic area and the ovarian cyst measured 59×43 mm (Figure 2) [2]. Again, there was no visualization of an intrauterine gestational sac nor was there any free fluid visualized in the cul-de-sac. At the time of this third ultrasound, the patient was experiencing mild left lower quadrant discomfort and also started having a very scant amount of vaginal spotting. This new development increased the suspicion of a possible ovarian ectopic pregnancy.

*Corresponding author: Cau Van Vo, Department of Obstetrics and Gynecology, Division of Clinical Education, Midwestern University, Glendale, AZ 85308, Fountain Valley Regional Hospital and Medical Center, Fountain Valley, CA 92708, USA, Tel: 714-348-4782; Fax: 714-898-2105; E-mail: cauvanvo@aol.com

Received June 03, 2014; Accepted September 22, 2014; Published September 24, 2014


Copyright: © 2014 Vo CV, et al. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.
The sonographic parameters of fetal development from conception have been well established [2]. Initially, most ovarian pregnancies would appear as a simple cyst in conjunction with an empty gestational sac [2-5]. The sac subsequently would grow larger with the wall becoming more echoic [2]. The placenta would progressively grow and form a hyperchoic structure between the gestational sac and the ovarian stroma. These changes are therefore important markers for early diagnosis of ovarian ectopic pregnancy.

In our patient, based on the ultrasonographic findings and beta-hCG levels, a diagnosis of ectopic pregnancy was suspected [3-5]. Unlike with tubal pregnancies, the ultrasound findings that we were able to visualize were strongly suggestive of an ovarian ectopic pregnancy [2,5].

A subsequent diagnostic laparoscopy revealed an unruptured left ovarian pregnancy with a purplish protrusion on the ovarian cortex (Figure 3). Both fallopian tubes appeared normal. A wedge resection of the left ovary was performed. Direct visualization of the base of the sac revealed placenta-like tissue, which was easily peeled off. A histopathological examination of the specimen confirmed the presence of chorionic villi and ovarian stroma (Figures 4-6). An endometrial curettage was also performed and revealed proliferative tissue.

The operation was performed in an outpatient surgical center. The patient remained in the recovery room for a total of three and a half hours. Her postoperative course was uneventful and her quantitative human chorionic gonadotropin level decreased from 1,714.29 to 24.9 mIU/ml at 7 days and to 0.10 mIU/ml at 27 days postoperatively.

**Comment**

Primary ovarian pregnancy is an extremely uncommon form of ectopic pregnancy. Prerupture diagnosis of an ovarian pregnancy was extremely difficult before the routine use of the diagnostic ultrasound. Even with the use of diagnostic ultrasound, making a diagnosis prior to rupture has continued to be a challenge for obstetricians. The literature illustrates that ultrasonographic appearances of ovarian pregnancy vary from the presence of a simple cyst within the ovary, to the presence of a walled cystic mass or an echocomplex mass [5,6]. There has even been a rare case report documenting the presence of an ovarian cyst with a fetal pole with cardiac activity inside the cyst [2]. In another study, there were six ovarian ectopic pregnancies diagnosed by ultrasound prior to rupture. In two of these cases, the ultrasound revealed a fetal pole with cardiac activity noted within the ovary and in the other four cases, there was just a gestational sac within the ovary. The variations in these ultrasound findings have depended on the gestational age, the degree of bleeding and whether or not there has been rupture [5].

Initial ultrasonographic examination, transabdominal or transvaginal, may not always accurately detect an ovarian pregnancy. This may be due to the proximity of the ovary to the uterine cornua, the density of the ovarian stroma, the limited acoustic window, or the patient’s body habitus. Sonographic demonstration of an intact gestational sac (Figure 1) is an important finding for the diagnosis of an ovarian pregnancy.

**Figure 1:** Sonographic image shows the intact gestational sac (C, white arrow) inside the left ovary (Figure A).

**Figure 2:** Sonographic image shows the intact gestational sac and the placenta (C, white arrow) inside the left ovary (Figure B).

**Figure 3:** Laparoscopic picture demonstrates the pregnancy with purplish bulge at the intact ovarian cortex (P, black arrow) (Figure C).

**Figure 4:** Laparoscopic picture shows the pregnancy inside the intact ovarian cortex (P, black arrow) (Figure D).
transvaginal, can start as early as 5 weeks gestation and if the pregnancy appears to be abnormal (no gestational sac visualized within the uterus), ultrasounds can be repeated at two to three day intervals for early detection prior to rupture [2]. Several authors agreed that any patient with a beta-hCG above the discriminatory level of 1,200 mIU/ml without intrauterine gestational sac visualized was clearly presumed to have an ectopic pregnancy [3,4]. This had become a significant biochemical marker of ectopic pregnancy.

Ultrasound images have been proven safe for management of obstetric patients from conception. The sonoanatomic parameter for development of the fetus from conception to delivery had been established and published [2]. Ovarian pregnancy is extremely rare, thus making it difficult to make an early diagnosis. Ovarian pregnancies also rupture earlier than tubal pregnancies. One study showed the mean gestational age of rupture of ovarian pregnancies was 45 days. Since ovarian pregnancies are also ectopic pregnancies, the BhCG marker for diagnosis of ectopic pregnancy could also be applied for ovarian pregnancy. The use of BhCG in aiding in the diagnosis of ectopic pregnancy has been well documented in the literature [4]. The majority of ovarian pregnancies develop as an empty sac and very few cases have had a live fetus with cardiac motion and a yolk sac.

Ultrasoundographic technology had emerged as a gold standard for evaluating fetal development. Ultrasound is the best tool that can detect and differentiate between normal and abnormal development of pregnancy, especially in the early stages of the first trimester. If no intrauterine gestational sac is seen at 5 weeks by a sure LMP and there is an inappropriately elevated BhCG, then an ectopic pregnancy should be suspected and followed closely. This is especially true if there is also an accompanying mass or cyst at the adnexa. If there is a suspicion of an ectopic pregnancy, ultrasounds should be repeated at 2-3 day intervals in an attempt to make an accurate diagnosis and to avoid possible rupture. Most ovarian pregnancy would appear sonographically as a simple cyst. This simple cyst is usually an anembryonic gestational sac [2,5]. It would initially show a smooth wall with lucency in the center. The gestational sac would subsequently grow larger at an appropriate rate during the initial weeks with the wall becoming more echic due to decidual reaction. The placentla development would also progress and form a hyperchoic layer between the gestational sac and ovarian tissue. These changes would be a typical development of the blighted ovum and thus an important sonoanatomic marker for early diagnosis of ovarian ectopic pregnancy. Bleeding and rupture could happen at any time leading to an immediate need for surgical intervention.

In our case, serial ultrasonographic examinations revealed significant changes in the size and the appearance of the left ovarian cystic mass. Moreover, the gestational sac size was more than 30 mm without the presence of fetal pole indicating that the ovarian cystic mass was actually a blighted ovum [2]. At the same time, serial human chorionic gonadotropin levels were not doubling in the 72-hour period. In addition, the levels of beta-hCG were above the discriminatory threshold without an accompanying intrauterine gestational sac. These were the sonoanatomic and biochemical markers for ovarian ectopic pregnancy.

The differential diagnosis in our case included a tubal pregnancy. Our sonoanatomic images were not typical for an unruptured tubal pregnancy since most unruptured tubal pregnancies appear as a small hyperechoic ring with a hypoechoic center surrounded by a distended tubal wall [2,5]. The differential also included an early abnormal intrauterine pregnancy with a corpus luteum cyst. In our case, the ovarian cyst was unlikely to be a corpus luteum because its appearance of the cyst was smooth and it had a thin echogenic border with an irregular fine internal echos [2]. The difficulty in differential diagnosis between ovarian and tubal pregnancy is increased after rupture because of ultrasonographic changes associated with blood clots or organized hematomas [2,5]. The sonoanatomic images become a complex cyst or echocomplex mass.

As expected, serial ultrasound examinations in the early stage of the first trimester coupled with serial beta-hCG allowed for more unruptured ovarian pregnancies to be diagnosed preoperatively [3,4]. With the new ultrasonographic techniques, such as high resolution transvaginal ultrasound and 3-D sonography, the success in diagnosing an ectopic, before rupture, has improved significantly. This includes the diagnosis of an ovarian pregnancy [7-9].

When a diagnosis of an unruptured ovarian ectopic pregnancy is established, a quick response is necessary. The management options depend upon the clinical presentation, upon whether or not the patient is stable and upon the clinical experience of the physician. The following section describes the case studies of “methotrexate for treatment of ovarian ectopic pregnancy” in the literature from 1988 to 2014. In that time period, there were a total of 19 cases reported that were either treated with methotrexate or surgery.

Medical management

1) There were three cases, in the literature that had unruptured ovarian pregnancies that were accurately diagnosed and documented by laparoscopy (Shamma, Chelmow and Mittal). Each of these cases was successfully managed with methotrexate.
patients with a severely damaged ovary.

In 1878 Spiegelberg established four criteria for diagnosis of ovarian ectopic pregnancy:

1) The fallopian tube on the affected side must be intact, 2) the pregnancy must occupy the normal position of the ovary, 3) the affected ovary must be attached to the uterus by the ovarian ligament and 4) the ovarian tissue must be attached to the pregnancy in the specimen [3,4,8].

Almost all articles in the literature referenced these criteria as a proof of confirmation of the diagnosis. Spiegelberg described these criteria in his article published in Germany in 1878. The results described in his article would be considered an abdominal pregnancy based on our knowledge now instead of an ovarian pregnancy. Furthermore, these criteria were based on macroscopic examination of the pelvis and the specimen at the time of the surgery. A microscopic confirmation of chorionic villi or trophoblast was not available back in 1878.

These extensive studies in the literature and the results of this ovarian ectopic pregnancy provide a new perspective into the difficulty of diagnosis and differentiation with other ectopic pregnancies. This revelation is also a compelling reason for modifying the Spiegelberg’s criteria which could be accurately applied to all future studies.

Considering the advances in laparoscopic surgery and the availability of histopathology today, we suggest that the Spiegelberg criteria for diagnosing ovarian pregnancies be reconsidered. We would recommend the following modified criteria for a diagnosis of primary ovarian ectopic pregnancy:

1) Both fallopian tubes must be intact.
2) The ectopic pregnancy must be within or attached to the ovary.
3) The chorionic villi or trophoblast and ovarian tissue must be microscopically confirmed from the surgical specimen.

In order to make a final diagnosis of ovarian ectopic pregnancy, all three of the new criteria have to be met.

Conclusion

In our case of a primigravid patient at 7.4 weeks by certain LMP who presented with an ultrasound showing no intrauterine pregnancy and a left cystic ovarian mass, there was an initial concern for an ectopic pregnancy. Our patient’s serial BhCG levels were low, but they were above the discriminatory level. In addition, the patient’s BhCG levels did not double in a 72- hour period. The serial ultrasounds in our patient revealed significant changes in size and the appearance of the ovarian cyst. These changes were not consistent with a simple ovarian cyst. The changes were most consistent with an ovarian pregnancy. Given the ultrasound findings, along with the BhCG levels, there were significant concerns regarding a diagnosis of an ovarian pregnancy. The addition to the patient’s symptoms leads to her being taken to the operating room for a diagnostic laparoscopic procedure right away.

Ultrasound is essentially the best tool that can detect ovarian ectopic pregnancy before rupture. The combination of biochemical and sonographic markers with clinical examination prompted a preoperative diagnosis before rupture and led to successful minimally invasive surgery.

Acknowledgement

The author wishes to thank Tien Tuy Vo, Ph.D., for editorial, graphic and digital art assistance.
References