

Pregnancy of Unknown Location: A Case Report and Review of the Literature

Deepa C Rajan^{1*} and Suzy JF Matts²

¹Specialist Registrar, Department of Obstetrics and Gynaecology, George Eliot Hospital, Nuneaton, UK

²Consultant Obstetrician and Gynaecologist, Department of Obstetrics and Gynaecology, George Eliot Hospital, Nuneaton, UK

Abstract

We present a case of pregnancy of unknown location with high serum β hcg levels. This case posed a diagnostic difficulty in the management of pregnancy of unknown location. The patient underwent two laparoscopic procedures to establish the diagnosis. The case highlights the importance of close surveillance of patients who present with pregnancy of unknown location.

Keywords: Pregnancy of unknown location; Diagnostic laparoscopy

Introduction

Management of Pregnancy of Unknown Location (PUL) has always been challenging. Expectant management is an option for clinically stable women with minimal symptoms and a pregnancy of unknown location [1]. Since there is always a possibility of ectopic pregnancy these women should be monitored with caution. The prevalence of ectopic pregnancy in PUL is 14-28% [2]. This is an interesting case of pregnancy of unknown location with serum β hcg levels above the discriminatory levels.

Case Report

A 27 yr old G3P2 presented to the early pregnancy clinic at 6 weeks gestation with lower abdominal pain and mild spotting per vaginum. Her serum β hcg level was 5574 iu/l and Transvaginal Scan (TVS) reported no evidence of intrauterine pregnancy with normal adnexae. The patient was admitted with a diagnosis of PUL and a diagnostic laparoscopy was done. On laparoscopy there was no evidence of an extrauterine pregnancy and the pelvis was normal. She was then followed up with serial β hcg levels as an outpatient. A repeat β hcg level in 48hrs was 6096 iu/l. A repeat TVS revealed an endometrial thickness of 14 mm, no intrauterine sac, normal ovaries and small area of mixed echogenicity suggesting possible early pregnancy. She was managed conservatively with monitoring of β hcg levels and ultrasound pelvis. The follow up TVS after a week showed no evidence of Intrauterine Pregnancy (IUP) and 16×19×15 mm hyperechoic focal lesion in the left ovary. The serum β hcg level was 8015 iu/l. The patient continued to remain stable without any symptoms. She then had a second diagnostic laparoscopy and endometrial curettage. The laparoscopy showed no evidence of extrauterine pregnancy. The histology report from the curettings was consistent with products of conception. Her β hcg levels following the endometrial curettage dropped to 922 iu/l (Figure 1). The final diagnosis was a silent miscarriage which had presented in this unusual way.

Discussion

This case demonstrates the difficulty in managing cases of PUL. About 8-13% of women who present to the early pregnancy unit are classified as having a PUL [3-5]. Although conservative management is an option, there is always a possibility of missing an ectopic pregnancy especially with such high β hcg levels. β hcg levels are routinely used to monitor outcomes of PUL. Discriminatory zones of β hcg are defined as levels above which IUP must be visualised on ultrasound. There are various discriminatory zones based on the quality of the ultrasound machine, skill of the sonographer, the symptoms of the woman,

presence of uterine fibroids and multiple pregnancy [1]. Varying the discriminatory zone does not significantly improve the detection of ectopic pregnancy in a PUL population [6]. Although β hcg titres are normally expected to double in 48 hrs this can vary according to the gestation [7]. When the rise or the fall in hCG is suboptimal [8-11], the most likely outcome is ectopic pregnancy. Approximately 71% of women with ectopic pregnancy have a rise in β hCG that is slower than the minimal rise for a viable pregnancy or a decline that is slower than the minimal rate of fall in spontaneous miscarriage [9]. However, failing PULs and 15% of normal pregnancies [12,13] will have an abnormal doubling time. Serum progesterone can be a useful adjunct when ultrasound suggests pregnancy of unknown location. In the presence of pregnancy of unknown location serum progesterone level less than 20nmol/l predicts spontaneous pregnancy resolution with a sensitivity of 93% and specificity of 94% [7]. Serum hCG and progesterone levels at defined times can be used to predict the immediate viability of a PUL, but cannot be used reliably to predict its location [5]. According to the Association of Early Pregnancy Units guidelines [8], if no intrauterine or ectopic pregnancy or retained products of conception are seen on transvaginal ultrasound and the woman is asymptomatic

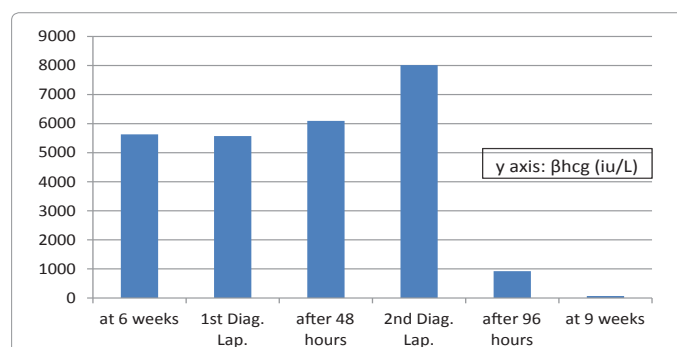


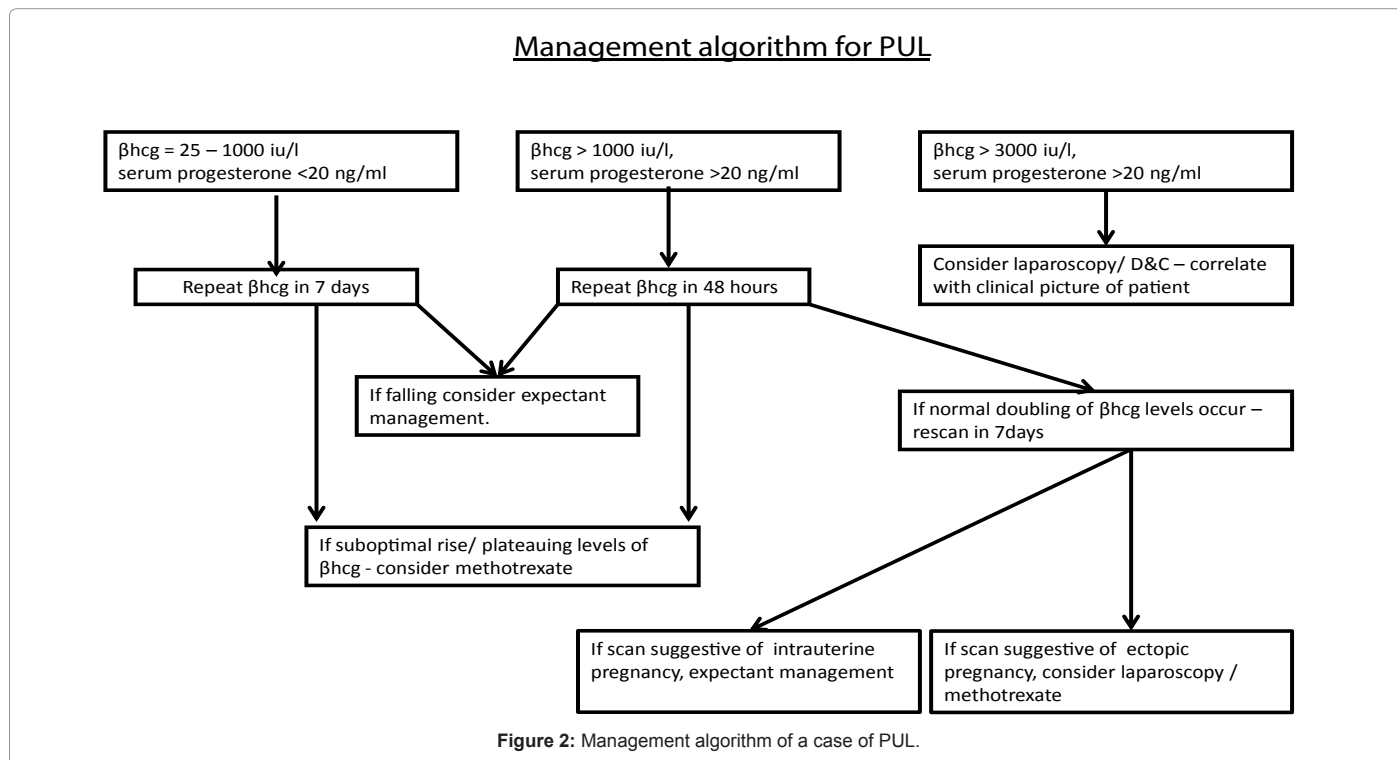
Figure 1: Graph demonstrating the β hcg levels with progression of pregnancy.

*Corresponding author: Dr. Deepa Christy Rajan, Department of Obstetrics and Gynaecology, George Eliot Hospital, Nuneaton, UK, Tel: 0044-7846335763; E-mail: deepaherman@yahoo.co.uk

Received September 07, 2012; Accepted September 26, 2012; Published September 28, 2012

Citation: Rajan DC, Matts SJF (2012) Pregnancy of Unknown Location: A Case Report and Review of the Literature. J Clin Case Rep 2:207. doi:10.4172/2165-7920.1000207

Copyright: © 2012 Rajan DC, 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.



at initial assessment, she can be managed conservatively. It is advisable to follow up with β hCG and transvaginal ultrasound assessments until the pregnancy is located accurately or intervention becomes necessary [14]. Women should be considered for active intervention if symptoms of ectopic pregnancy occur, serum β hcg levels rise above the discriminatory level or levels start to plateau [4,15]. D&C remains valuable to differentiate EP from nonviable IUP and to avoid misdiagnosis and unnecessary exposure to methotrexate. Low initial hCG values and ultrasound findings such as a thin endometrial echo complex and the presence of free fluid are associated with but are not diagnostic of an ectopic pregnancy [16]. Women should be followed up until the final pregnancy outcome is known which includes: failing PUL, intra-uterine pregnancy, ectopic pregnancy and persisting PUL [17]. We suggest a possible management algorithm for patients with unusual presentation (Figure 2).

In conclusion this case highlights the importance of cautious monitoring in case of PUL and timely intervention. Also clear information should be given to the women with PUL regarding the management, follow up and outcomes.

References

1. RCOG greentop guidelines No.21:May 2004
2. Banerjee S, Aslam N, Woelfer B, Lawrence A, Elson J, et al. (2001) Expectant management of early pregnancies of unknown location: a prospective evaluation of methods to predict spontaneous resolution of pregnancy. *BJOG* 108: 158–163.
3. Hahlin M, Thorburn J (1995) *Hum Reprod* 10:1223-1227
4. Banerjee S, Aslam N, Zosmer N, Woelfer B, Jurkovic D (1999) The expectant management of women with early pregnancy of unknown location *Ultrasound. Obstet Gynecol* 14: 231-236.
5. Condous G, Lu C, Van Huffel SV, Timmerman D, Bourne T (2004) Human

chorionic gonadotrophin and progesterone levels in pregnancies of unknown location. *Int J Gynaecol Obstet* 86: 351-357.

6. Condous G, Kirk E, Lu C, Van Huffel S, Gevaert O, et al. (2005) Diagnostic accuracy of varying discriminatory zones for the prediction of ectopic pregnancy in women with a pregnancy of unknown location. *Ultrasound Obstet Gynecol* 26: 770-775
7. RCOG greentop guideline No.25(2006)
8. Guidelines (2007) Association of Early Pregnancy Units.
9. Seeber BE, Barnhart KT (2006) Suspected ectopic pregnancy. *Obstet Gynecol* 107: 399–413.
10. Seeber BE, Sammel MD, Guo W, Zhou L, Hummel A, et al. (2006) Application of redefined human chorionic gonadotropin curves for the diagnosis of women at risk for ectopic pregnancy. *Fertil Steril* 86: 454–459.
11. Barnhart K, Sammel MD, Chung K, Zhou L, Hummel AC, et al. (2004) Decline of serum human chorionic gonadotropin and spontaneous complete abortion: defining the normal curve. *Obstet Gynecol* 104: 975–981.
12. Kadar N, Caldwell BV, Romero R (1981) A method of screening for ectopic pregnancy and its indications. *Obstet Gynecol* 58: 162–166.
13. Ling FW, Stovall TG (1994) Update on the diagnosis and management of ectopic pregnancy. Chicago: Mosby Year Book, Inc 55–83.
14. Sagili H, Mohamed K (2008) Pregnancy of unknown location: An evidence – based approach to management. *The Obstetrician & Gynaecologist* 10: 224-230.
15. Hajenius P, Mol B, Ankum W, Van der Veen F, Bossuyt P, et al. (1995) Suspected ectopic pregnancy: expectant management in patients with negative sonographic findings and low serum β -hCG concentrations. *Early Pregnancy* 1: 258–262.
16. Chung K, Chandavarkar U, Opper N, Barnhart K (2011) Reevaluating the role of dilation and curettage in the diagnosis of pregnancy of unknown location. *Fertil Steril* 96: 659-662.
17. Condous G, Okaro E, Bourne T (2005) Pregnancies of unknown location: diagnostic dilemmas and management. *Curr Opin Obstet Gynecol* 17: 568-573.