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Evaluation of ADAM-12 as a Diagnostic Biomarker of Ectopic Pregnancy in Women with a Pregnancy of Unknown Location

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Abstract

Background: Ectopic pregnancy (EP) remains the most life-threatening acute condition in modern gynaecology. It remains difficult to diagnose early and accurately. Women often present at emergency departments in early pregnancy with a ‘pregnancy of unknown location’ (PUL) and diagnosis/exclusion of EP is challenging due to a lack of reliable biomarkers. Recent studies suggest that serum levels of a disintegrin and metalloprotease protein-12 (ADAM-12) can be used to differentiate EP from viable intrauterine pregnancy (VIUP). Here we describe a prospective study evaluating the performance of ADAM-12 in differentiating EP from the full spectrum of alternative PUL outcomes in an independent patient cohort.

Methodology/Principal Findings: Sera were collected from 120 patients at their first clinical presentation with a PUL and assayed for ADAM-12 by ELISA. Patients were categorized according to final pregnancy outcomes. Serum ADAM-12 concentrations were increased in women with histologically-confirmed EP (median 442 pg/mL; 25%–75% percentile 232–783 pg/mL) compared to women with VIUP (256 pg/mL; 168–442 pg/mL) or miscarriage (192 pg/mL; 133–476 pg/mL). Serum ADAM-12 did not differentiate histologically-confirmed EP from spontaneously resolving PUL (srPUL) (416 pg/mL; 154–608 pg/mL). The diagnostic potential of ADAM-12 was only significant when ‘ambiguous’ PUL outcomes were excluded from the analysis (AORC = 0.6633; P = 0.03901).

Conclusions/Significance: When measured in isolation, ADAM-12 levels had limited value as a diagnostic biomarker for EP in our patient cohort. The development of a reliable serum biomarker-based test for EP remains an ongoing challenge.

Introduction

The diagnosis of ectopic pregnancy (EP) continues to present a major clinical challenge in obstetrics and gynecology, with patients often asymptomatic or presenting with non-specific symptoms that do not readily differentiate EP from miscarriage or viable intrauterine pregnancy.

Whilst in many cases, an EP will be detected by transvaginal ultrasonography (TVUSS) at the first clinic visit [1], TVUSS is often inconclusive and the pregnancy has to be initially classified as a “pregnancy of unknown location” (PUL) [2]. In patients with a PUL, subsequent diagnosis of EP relies on the serial measurement of serum human chorionic gonadotrophin (hCG) levels (and, in some centers, progesterone), together with follow-up TVUSS [3–5]. This approach significantly delays the diagnosis and management of EP and is resource intensive and expensive [6]. There remains an unmet clinical need for a serum biomarker capable of identifying EP at first clinical presentation [5,7].

Recently, Rausch et al [8] found a statistically significant decrease in a disintegrin and metalloprotease protein-12 (ADAM-12) in the sera of patients with EP (median 2.5 ng/mL), when compared to women with viable intrauterine pregnancy (median 18.6 ng/mL). The authors demonstrated this difference in a large cohort of 199 patients in the United States presenting with pain or bleeding in the first trimester of pregnancy. There appeared to be good discrimination between the groups as assessed by receiver operating characteristics (Area under ROC curve = 0.82; P<0.0001). They concluded that serum ADAM-12 was a promising biomarker for the diagnosis of ectopic pregnancy in women with symptoms in the first trimester.

However, there is debate as to the specificity of ADAM-12 with regard to differentiating EP from outcomes other than VIUP [9] due to the fact that other conditions, such as trisomy 21 can also present with alteration of ADAM-12 [10,11]. Furthermore, the promising findings reported by Rausch et al needed independent verification. We therefore set out to validate Rausch et al’s
findings, measuring ADAM-12 in a cohort of women prospectively recruited in the United Kingdom with a PUL.

**Results**

A total of 120 Caucasian women (aged 18–45 years) with a PUL were recruited to the study and grouped according to final pregnancy outcomes. Patient recruitment: 120 patients with an initial diagnosis of a PUL were recruited to the study and grouped according to final pregnancy outcomes.

<table>
<thead>
<tr>
<th>Group</th>
<th>Inclusion criteria</th>
<th>HCG (mU/ml)</th>
<th>Age (years)</th>
<th>Weight (Kg)</th>
<th>BMI</th>
<th>n</th>
</tr>
</thead>
<tbody>
<tr>
<td>dVIUP</td>
<td>Definite viable intrauterine pregnancy: TVUSS confirmation of intrauterine gestational sac with yolk sac and embryo with cardiac activity.</td>
<td>6844±217</td>
<td>28±1</td>
<td>70±4</td>
<td>26±2</td>
<td>28</td>
</tr>
<tr>
<td>dNVIUP</td>
<td>Definite nonviable intrauterine pregnancy: USS confirmation of intrauterine gestational sac with yolk sac and/or embryo without cardiac activity seen prior to uterine evacuation.</td>
<td>4022±1904</td>
<td>32±1</td>
<td>74±3</td>
<td>27±1</td>
<td>26</td>
</tr>
<tr>
<td>dEP</td>
<td>Definite ectopic pregnancy: intervention prompted by adnexal mass on TVUSS or by abnormal rise in serum hCG levels and confirmed at surgery and by histopathology.</td>
<td>1151±238</td>
<td>29±1</td>
<td>70±4</td>
<td>25±1</td>
<td>17</td>
</tr>
<tr>
<td>NP</td>
<td>Not pregnant: positive home pregnancy test result subsequently not confirmed by serum hCG measurement.</td>
<td>&lt;5</td>
<td>26±2</td>
<td>70±8</td>
<td>27±3</td>
<td>11</td>
</tr>
<tr>
<td>srPUL</td>
<td>Spontaneously resolving PUL: PUL with spontaneous resolution of serum hCG levels.</td>
<td>428±114</td>
<td>32±1</td>
<td>74±4</td>
<td>28±1</td>
<td>27</td>
</tr>
<tr>
<td>tPUL</td>
<td>Treated persistent PUL: abnormal rise in serum hCG levels but no adnexal mass or IU sac seen on TVUSS after monitoring, managed medically with methotrexate.</td>
<td>400±188</td>
<td>32±4</td>
<td>83±15</td>
<td>28±5</td>
<td>3</td>
</tr>
<tr>
<td>pEP</td>
<td>Probable ectopic pregnancy: inhomogenous adnexal mass or extraterine sac-like structure on TVUSS managed medically with methotrexate.</td>
<td>597±200</td>
<td>33±1</td>
<td>63±4</td>
<td>25±1</td>
<td>8</td>
</tr>
</tbody>
</table>

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**Discussion**

ADAM-12 may have some potential as a serum biomarker of dEP. However, we were unable to verify the findings of Rausch et al who had concluded ADAM-12 was a highly promising marker of ectopic pregnancy with strong diagnostic marker performance. In fact, we found that serum ADAM-12 concentrations in our UK cohort were elevated in patients with dEP compared to dVIUP (Figure 1A), rather than decreased as reported previously [8]. Furthermore, ADAM-12 did not appear to perform well as a biomarker of ectopic pregnancy.

We can only speculate why we were unable to replicate the promising findings described by Rausch et al [8]. They used a dissociation-enhanced lanthanide fluoroimmunoassay platform DELFIA/AutoDELFIA ADAM-12 research kit (PerkinElmer), compared to the ADAM12 Quantikine ELISA (R&D systems) used in the current study, and it is possible that the conflicting findings are due to this difference. However, it seems unlikely that this could account for the trend reversal observed between the two cohorts and differences in study design offer a more plausible explanation.

Gestational age is likely to be a key factor in ADAM12 levels, as it rises exponentially from around week 5 of the first trimester [13]. Therefore, it seems possible that the lower levels of ADAM12 we report reflect the gestational age of our prospectively collected first presentation cohort. Our study population was also slightly smaller.
than that of Rausch et al. and we only recruited from a single UK center whereas Rausch et al. recruited from multiple US sites. This could potentially explain the disparity in our findings.

Another key difference in the design of our study and that published by Rausch et al. [8], is the inclusion of the entire range of PUL outcomes in the study design, not just dEP and dVIUP. Crucially, we found that serum ADAM-12 concentrations in patients who required surgical intervention for dEP were very similar to those observed in patients with final outcome of srPUL, who did not require surgical or medical intervention (Figure 1A).

Regardless of these differences, the discrepancy in the findings of our study and those of Rausch et al. [8] demonstrates the importance of verifying potential EP biomarkers in independent cohorts, and preferably from multiple international centers. The development of non-invasive blood biomarker test that reliably diagnoses EP remains an ongoing challenge.

Materials and Methods

Patient samples

Ethical approval for this prospective study was obtained from the Lothian Research Ethics Committee (LREC 04/S1103/20 and 09/S1103/39), with informed written consent obtained from all patients. Whole blood was obtained from women during their first clinical presentation with a positive home pregnancy test and abdominal pain and/or bleeding and a TVS that had been unable to locate the site of the pregnancy. After clotting for 2 hrs at RT, sera were collected and stored at $-80^\circ$C in multiple aliquots. The women were monitored until their discharge from hospital and their final pregnancy outcomes were classified according to the recent PUL consensus statement [12].

Ultrasound assessments

The ultrasound system used was the Toshiba Aplio XG and all of the ultrasound assessments were performed by a team of trained, qualified and experienced ultrasonographers.

ADAM-12 ELISA

Sera were assayed using the ADAM12 Quantikine ELISA kit (R&D systems, Abingdon, UK) according to the manufacturers’ instructions. Comprehensive details of the assay’s performance parameters are available from the manufacturer [http://www.rn dsystems.com/pdf/DAD120.pdf].

Statistical Analysis

Statistical analyses, ELISA standard curve formulae and receiver operating characteristic (ROC) curves were generated using Prism 5.0 (GraphPad Software, La Jolla, USA).

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Author Contributions

Conceived and designed the experiments: AWH ST TK. Performed the experiments: JKB. Analyzed the data: AWH JKB. Contributed reagents/materials/analysis tools: AWH. Wrote the paper: AWH JKB ST TK.
References