The relationship between first-trimester pregnancy-associated plasma protein-A levels and intrapartum fetal distress development

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Abstract

Objective: To investigate the relationship between the development of intrapartum fetal distress and serum pregnancy-associated plasma protein-A (PAPP-A) levels measured during first-trimester aneuploidy screening tests.

Material and Methods: This retrospective study included 283 uncomplicated pregnancies that resulted in full-term live births via spontaneous labor or with the induction by oxytocin. Cases were divided into two groups based on whether their first-trimester PAPP-A multiple of the median (MoM) levels were ≤0.5 (Group 1, n=75) or >0.5 (Group 2, n=208). As primary end points, the rate of cesarean section (C/S), the rate of C/S due to fetal distress, and the umbilical artery blood pH values in cases of C/S for fetal distress were compared between the two groups. Statistical analyses were performed using the Chi-square test and independent samples t-test. P≤0.05 were considered statistically significant.

Results: The mean gestational age at birth and the birth weights were significantly lower in Group 1 than in Group 2 (p=0.002 and p=0.007, respectively). Although the rate of C/S was similar between the groups (p=0.823), the rate of C/S due to fetal distress was significantly higher in Group 1 than in Group 2 (68.4% vs. 42%, respectively; p=0.050) and the mean umbilical artery blood pH value for C/S deliveries indicated by fetal distress was lower (p=0.048) in Group 1 than in Group 2. When the mode of delivery was analyzed according to the application of labor induction, both the C/S delivery rates (31.6% in Group 1 and 31.7% in Group 2; p=0.992) and C/S delivery rates due to fetal distress (66.7% in Group 1 and 46.2% in Group 2; p=0.405) were similar in both groups.

Conclusion: Low PAPP-A levels (≤0.5 MoM) in the first trimester are associated with the risk of intrapartum fetal distress development and the likelihood of C/S for fetal distress. Nonetheless, this risk is not affected by labor induction. (J Turk Ger Gynecol Assoc 2016; 17: 139-42)

Keywords: Cesarean section, fetal distress, pregnancy-associated plasma protein A

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Introduction

Pregnancy-associated plasma protein A (PAPP-A) is one of the parameters in the first-trimester screening test that is used as a biochemical marker to detect aneuploidy in the early weeks of gestation (1). PAPP-A can be detected in the maternal blood 28 days after implantation in singleton pregnancies. The serum PAPP-A level starts to increase during the first trimester, doubling every 3 to 4 days. The rate of increase is gradual until week 36, after which it accelerates, and the maximum serum PAPP-A levels are reached at term (2).

It is thought that PAPP-A released from trophoblastic tissues at abnormal levels early during gestation affects fetal growth negatively by impairing trophoblastic invasion of the decidua, causing abnormal placentation and other pregnancy complications (3). A low PAPP-A concentration is a powerful indicator of potential pregnancy complications, including preeclampsia, intrauterine developmental retardation, gestational hypertension, fetal death, oligohydramnios, and preterm birth (4,6). Because PAPP-A levels during pregnancy are associated with obstetric complications, as well as abnormal placentation, it is thought that fetuses born to mothers with low maternal PAPP-A levels do not tolerate labor stress and also could be more likely to develop intrapartum fetal distress (7).

In this study, we compared the rate of emergency cesarean section (C/S) due to fetal distress and the umbilical arterial blood gas levels in patients with normal or low PAPP-A levels in the first trimester.

Material and Methods

We conducted a retrospective examination of 359 cases of singleton pregnancies in which the first-trimester screening for aneuploidy had been performed in our pregnancy out-
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Results

PAPP-A levels were ≤0.5 MoM in 75 (26.5%) cases (Group 1) and >0.5 MoM in 208 (73.5%) subjects (Group 2). The mean PAPP-A levels was 0.38±0.10 MoM for Group 1 and 1.14±0.63 MoM for Group 2 (p<0.001).

The age, gravidity, and parity were similar among the groups (p>0.05). (Table 1) The mean gestational age at birth and the birth weights were significantly lower in Group 1 than Group 2 (p=0.002 and p=0.007, respectively).

The cesarean delivery rate was 24.4% (69/283) in all the subjects. Although the rate of C/S were similar among the groups (p=0.823), the rate of C/S due to fetal distress was significantly higher in Group 1 (68.4% (13/19) vs 42% (21/50), p=0.050) (Table 1).

In addition, the mean umbilical artery blood gas pH value for C/S deliveries indicated by fetal distress was also significantly lower in Group 1 (p=0.048). According to the ACOG management protocol, the category III (abnormal) FHR pattern rate was 30.8% (4/13) in Group 1 and 33.3% (7/21) in Group 2 (p=0.877).

The induction of labor was applied in 19 (25.3%) subjects in Group 1 and in 41 (19.7%) subjects in Group 2 and the rate of labor induction were similar between the groups (p=0.307).

When the mode of delivery was analyzed according to applying labor induction, both C/S delivery rates ((6/19 (31.6%) in Group 1, 13/41 (31.7%) in Group 2; p=0.992)) and the rate of C/S delivery due to fetal distress ((4/6 (66.7%) in Group 1, 6/13 (46.2%) in Group 2; p=0.405) were not different in both groups.

Discussion

PAPP-A, which was first obtained from the plasma of pregnant women in 1974, is a protein released from the placenta, and its concentration in maternal blood reflects placental activity (9, 10). PAPP-A is a specific protease for insulin-like growth factor binding protein-4 (IGFBP-4), and thus plays a role in fetal growth and the development and many physiopathologic events related to IGF-1 and -2 (11).

Furthermore, it was found that PAPP-A plays an exclusive role in the autocrine and paracrine regulation of the trophoblastic invasion of the decidua (12, 13). Low PAPP-A levels are thought to be attributable to sequestration by binding proteins for free IGFs, and this can negatively affect fetal growth and cause obstetric complications (4, 12).

Ucella et al. (7) found that the rate of non-elective C/S deliveries indicated by fetal distress was higher among mothers with low PAPP-A levels (16.2%) than among mothers with normal PAPP-A (7.9%) levels. Thus, they suggested that low PAPP-A levels may be not only related to antenatal complications (preeclampsia, intrauterine growth retardation, preterm delivery, and loss of pregnancy), but also may be a risk factor for acute intrapartum fetal distress related to abnormal placentaion and placental dysfunction, leading to more emergency C/S deliveries (7).

In this study, we investigated whether fetal distress developed more frequently during labor in cases with low maternal PAPP-A levels in the first trimester and examined the umbilical arterial blood gas levels to evaluate whether there was an indirect correlation between low PAPP-A levels and the placental reserve during active labor.
Our findings are consistent with the results reported by Ucella et al. (7) and demonstrate a higher rate of C/S delivery because of fetal distress during active labor in cases with low PAPP-A levels (≤0.5 MoM) measured during the first trimester.

Ucella et al. (7) also found that the umbilical arterial blood pH was significantly lower in cases where a C/S was performed because of fetal distress in cases of low PAPP-A levels compared to cases with normal PAPP-A levels (pH: 7.19 vs pH: 7.26, respectively). We also analyzed the umbilical cord blood pH immediately postpartum in cases where a C/S was performed because of fetal distress, but we could not find any significant differences between the groups. Although the intrapartum wellbeing of the fetus during labor is generally followed with fetal electronic monitoring (FEM), in almost 50% of the fetuses where fetal distress is detected with FEM, the oxygenation of the fetus is normal (14-16).

In conclusion, PAPP-A levels between the 11th and 14th weeks of gestation likely reflect placental function, and the risk of developing intrapartum fetal distress and the risk of C/S due to fetal distress are higher in cases with low (≤0.5 MoM) PAPP-A levels. Nonetheless, this risk is not affected by labor induction by oxytocin. The PAPP-A level measured during aneuploidy screening in the first trimester of pregnancy can help predict the development of intrapartum fetal distress. Our study is limited by its retrospective design; thus, prospective studies are required to verify our results.

**Ethics Committee Approval:** Ethics committee approval was received for this study from the ethics committee of Yıldırım Beyazıt University School of Medicine.

**Informed Consent:** Written informed consent was obtained from patients who participated in this study.

**Peer-review:** Externally peer-reviewed.


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