Amniotic fluid angiopoietin-2 in term and preterm parturition, and intra-amniotic infection/inflammation

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Abstract

Objective: Recent observations have revealed an interaction between inflammation and angiogenesis, which may be mediated by angiopoietins and chemokines. Given the importance of inflammation in parturition, we sought to determine whether angiopoietin-2 (Ang-2) is present in amniotic fluid (AF) and if its concentration changes with gestational age, labor, and in intra-amniotic infection/inflammation (IAI) in patients with spontaneous preterm labor and intact membranes.

Study design: This cross-sectional study included 486 patients in the following groups: 1) women in the mid-trimester of pregnancy (14–18 weeks) who underwent amniocentesis for genetic indications and delivered a normal neonate at term (n=52); 2) normal pregnant women at term with (n=48) and without (n=45) spontaneous labor; 3) patients with an episode of spontaneous preterm labor (PTL) who were classified into: a) PTL without IAI who delivered at term (n=152); b) PTL without IAI who delivered preterm (<37 weeks gestation; n=107); and c) PTL with IAI (n=82). Ang-2 concentration in AF was determined by enzyme-linked immunoassay. Non-parametric statistics were used for analysis.

Results: 1) Ang-2 was detected in all AF samples; 2) the median AF Ang-2 concentration at term was significantly lower than that in the mid-trimester (1877.4 pg/mL vs. 3525.2 pg/mL; P<0.001); 3) among patients with PTL, the median AF Ang-2 concentration was significantly higher in patients with IAI than in those without IAI (4031.3 pg/mL vs. 2599.4 pg/mL; P<0.001) and those with PTL without IAI who delivered at term (4031.3 pg/mL vs. 2707.3 pg/mL; P<0.001); and 4) no significant differences were observed in the median AF Ang-2 concentration between patients with spontaneous labor at term and those at term not in labor (1722.9 pg/mL vs. 1877.4 pg/mL; P=0.6).

Conclusions: 1) Ang-2, a protein involved in the process of vascular remodeling, is a physiologic constituent of the amniotic fluid and its concentration decreased with advancing gestation; 2) the median Ang-2 concentration in amniotic fluid is higher in patients with IAI than in those without; and 3) spontaneous parturition at term is not associated with changes in the AF concentration of Ang-2. These findings support the view of a link between angiopoietins and inflammation.

Keywords: Angiogenesis; chorioamnionitis; microbial invasion of the amniotic cavity; pregnancy; preterm delivery; preterm labor; preterm parturition syndrome, Tie-2; vasculogenesis.

Introduction

Angiogenesis is the process by which new vessels are formed from pre-existing vasculature. This process is tightly regulated by different families of growth factors. Along with the well-characterized vascular endothelial growth factor (VEGF) family, angiopoietins have been shown to be critical in orchestrating blood vessel formation [8]. Angiopoietin-1 (Ang-1) and angiopoietin-2 (Ang-2) are structurally-related endothelial growth factors. They bind with similar specificity and affinity to the tyrosine kinase receptor with immunoglobulin and epidermal growth factor homology domain 2 (Tie-2), which is expressed on endothelial cells and hematopoietic cells [12, 33, 60]. Studies in knockout mice have shown that the Ang-1/Tie-2 system plays an essential role in embryonic vascular remodeling [60, 63]. Under physiologic
conditions, Ang-1, through Tie-2 signaling, mediates vessel maturation and maintains vessel integrity by the recruitment of periendothelial cells [31].

Ang-2 is a competitive inhibitor of Ang-1 because both bind to Tie-2 tyrosine kinase receptor [12, 33, 60]. Ang-1 induces phosphorylation of Tie-2, whereas Ang-2 binding does not activate the receptor [33]. Remarkably, Ang-2 acts only as an antagonist of Ang-1 on endothelial cells, leads to loosening of cell/cell interactions and allows access to angiogenic inducers such as VEGF [33]. Ang-2 may act with VEGF to promote angiogenic sprouting from established vasculature [18, 33]. Thus, Ang-2 is specifically required for normal postnatal vessel remodeling [18].

Parturition is considered to be an inflammatory process [24, 49, 52, 54]. In addition, the expression of angiogenesis-related genes is significantly increased in the mouse uterus during spontaneous labor at term, as well as induced preterm labor by either bacteria or ovarioectomy [23]. These findings suggest that changes in gene expression of angiogenic-related genes are part of the common pathway of parturition.

Since there is a paucity of information about the role of Ang-2 in human amniotic fluid, we determined whether Ang-2 is present in human amniotic fluid and if its concentration changes with gestational age, in the presence of labor (term and preterm), and in intra-amniotic infection/inflammation (IAI) in patients with spontaneous preterm labor. Parturition.

Materials and methods

Study design and population

A cross-sectional study was designed by searching our clinical database and bank of biological samples including 486 patients in the following groups: 1) women in the mid-trimester of pregnancy (14–18 weeks) who underwent amniocentesis for genetic indications and delivered a normal neonate at term (n = 52); 2) normal pregnant women at term with (n = 48) and without (n = 45) spontaneous labor; and 3) patients with an episode of spontaneous preterm labor (PTL) and intact membranes who were classified into: a) PTL without IAI who delivered at term (n = 152); b) PTL without IAI who delivered preterm (<37 weeks’ gestation; n = 107); and c) PTL with IAI (n = 82).

Definitions

Patients were considered to have a normal pregnancy outcome if they did not have any medical, obstetrical, or surgical complications and delivered a term neonate (>37 weeks) of greater than the 10th percentile for gestational age [1, 22] without complications. Spontaneous preterm labor was defined by the presence of regular uterine contractions occurring at a frequency of at least two every 10 min associated with cervical changes before 37 completed weeks of gestation that required hospitalization. Intra-amniotic infection was defined as a positive amniotic fluid culture for microorganisms. Intra-amniotic inflammation was diagnosed by an amniotic fluid interleukin (IL)-6 concentration ≥2.6 pg/mL [68]. Histologic chorioamnionitis was diagnosed based on the presence of inflammatory cells in the chorionic plate and/or chorioamniotic membranes. Acute funisitis was diagnosed by the presence of neutrophils in the wall of the umbilical vessels and/or Wharton’s jelly, using criteria previously described [41].

Sample collection

Amniotic fluid samples were obtained from transabdominal amniocentesis performed for genetic indications, evaluation of microbial status of the amniotic cavity, and/or assessment of fetal lung maturity in patients approaching term. Women at term in labor consisted of women who were admitted for suspected preterm labor because of uncertain dates and had an amnioncentration for the assessment of fetal lung maturity. The criteria for determining that these patients were at term was derived retrospectively, if the following criteria were met: 1) spontaneous labor; 2) delivery within 24 hours from amniocentesis; 3) analysis of amniotic fluid consistent with fetal lung maturity; 4) birthweight >2500 g; 5) absence of respiratory distress syndrome or other complications of prematurity; and 6) physical examination of the newborn by pediatricians which was consistent with a term neonate. Samples of amniotic fluid were transported to the laboratory in a sterile capped syringe and cultured for aerobic/anaerobic bacteria and genital mycoplasmas. White blood cell (WBC) count, glucose concentration and Gram-stain were also performed shortly after collection, as previously described [48, 53, 55]. The results of these tests were used for clinical management. Amniotic fluid IL-6 concentrations were used only for research purposes. Amniotic fluid not required for clinical assessment was centrifuged for 10 min at 4°C, and the supernatant was aliquoted and stored at −70°C until analysis. Among patients with spontaneous preterm labor with intact membranes who delivered within 72 hours of amniocentesis, placenta, umbilical cord, and chorioamniotic membranes were collected and the presence or absence of histologic chorioamnionitis and funisitis was assessed. This period of time was selected to preserve a meaningful temporal relationship between amniotic fluid Ang-2 concentration and placental pathologic findings.

All women provided written informed consent prior to the collection of amniotic fluid. The collection of amniotic fluid and its utilization for research purposes was approved by the Institutional Review Boards of Wayne State University (Detroit, Michigan, USA), Sotero del Rio Hospital (Santiago de Chile, Chile) and the Eunice Kennedy Shriver National Institute of Child Health and Human Development, NIH, DHHS. Many of these samples have been previously used to study the biology of inflammation, hemostasis, and growth factor concentrations in normal pregnant women and those with pregnancy complications.

Determination of human Ang-2 concentration in amniotic fluid

Specific and sensitive enzyme-linked immunoassay (R&D System, Inc. Minneapolis, MN, USA) was used to determine concentrations of Ang-2 in human amniotic fluid. The concentrations of Ang-2 in amniotic fluid samples were determined by inter-
Results

Demographic and clinical characteristics of the study population

Four hundred and eighty-six patients were included in the study. Table 1 shows the demographic and clinical characteristics of the patients in the mid-trimester, term not in labor and term in labor groups. Table 2 displays the demographic and clinical characteristics of the patients with spontaneous preterm labor and intact membranes. The median body mass index (BMI), and the rates of smoking and obesity (BMI \( \geq 30 \) kg/m²) were significantly higher in patients with PTL with IAI and preterm delivery than in those without IAI and preterm delivery and those without IAI and term delivery. In addition, the gestational age at amniocentesis, the gestational age at delivery and birthweight were significantly lower in patients with PTL with IAI and preterm delivery than that...
of those without IAI and preterm delivery and those without IAI and term delivery.

**Amniotic fluid Ang-2 concentration decreases with gestational age and does not change in spontaneous labor at term**

Ang-2 was detected in all amniotic fluid samples. Women with a normal pregnancy at term not in labor had a significantly lower median Ang-2 concentration in amniotic fluid than those in the mid-trimester [term not in labor: 1877.4 pg/mL, inter-quartile range (IQR) 1322.6–2434.1 vs. mid-trimester: 3525.2 pg/mL, IQR 2574.3–4852.2; P < 0.001] (Figure 1). In contrast, no significant differences were observed in the median amniotic fluid Ang-2 concentration between patients with spontaneous labor at term and those at term not in labor (term in labor: 1722.6 pg/mL, IQR 1293.5–2499.3 vs. term not in labor: 1877.4 pg/mL, IQR 1322.6–2434.1; P = 0.6).

**Ang-2 concentration is increased in the amniotic fluid of women with spontaneous preterm labor with intact membranes with intra-amniotic infection/inflammation**

Patients with spontaneous preterm labor with IAI had a significantly higher median amniotic fluid concentration of Ang-2 than those who delivered preterm without IAI (PTL with IAI: 4041.3 pg/mL, IQR 2463.0–6385.3 vs. PTL without IAI: 2599.4 pg/mL, IQR 1822.3–4145.8; P < 0.001) and than patients with spontaneous preterm labor who delivered at term (PTL delivered at term: 2707.3 pg/mL, IQR 1791.9–4274.4; P < 0.001) (Figure 2). After adjusting for gestational age at amniocentesis and storage time, similar results were obtained (ANCOVA, P = 0.02 among subgroups comparisons). There were no differences in the median amniotic fluid Ang-2 concentration between patients with spontaneous preterm labor without IAI who delivered preterm and those who delivered at term (PTL without IAI: 2599.4 pg/mL, IQR 1822.3–4145.8 vs. PTL delivered at term: 2707.3 pg/mL, IQR 1791.9–4274.4; P = 0.6).

When the analysis was restricted to patients with spontaneous preterm labor without IAI who delivered within 72 h from the amniocentesis (n = 18), this subgroup had a similar median amniotic fluid concentration of Ang-2 compared to those with spontaneous preterm labor with intact membranes who delivered at term (PTL without IAI with delivery within 72 h: 2460.5 pg/mL, IQR 1648.7–4162.8 vs. PTL delivered at term: 2707.3 pg/mL, IQR 1791.9–4274.4; P = 0.4).

**Amniotic fluid Ang-2 concentrations in patients with histologic chorioamnionitis and funisitis**

Placental histopathologic diagnoses were available in 77% (54/70) of patients with spontaneous preterm labor who delivered within 72 h of amniocentesis. Of those,
Figure 2  Amniotic fluid concentration of angiopoietin-2 (Ang-2) among women with intra-amniotic infection/inflammation in spontaneous preterm labor with intact membranes. The median amniotic fluid concentration of Ang-2 in patients with spontaneous preterm labor with IAI was significantly higher than that of those who delivered preterm without IAI (PTL with IAI: 4041.3 pg/mL, IQR 2463.0–6385.3 vs. PTL without IAI: 2599.4 pg/mL, IQR 1822.3–4145.8; P < 0.001) and that of patients with spontaneous preterm labor who delivered at term (PTL delivered at term: 2707.3 pg/mL, IQR 1791.9–4274.4; P < 0.001). There were no differences in the median amniotic fluid Ang-2 concentration between patients with spontaneous preterm labor without IAI who delivered preterm and those who delivered at term (PTL without IAI: 2599.4 pg/mL, IQR 1822.3–4145.8 vs. PTL delivered at term: 2707.3 pg/mL, IQR 1791.9–4274.4; P = 0.6).

52% (28/54) had evidence of placental inflammation. Patients with histologic chorioamnionitis and/or funisitis (n = 28) had higher median Ang-2 concentration in amniotic fluid than those (n = 26) without histologic chorioamnionitis, but this difference was not statistically significant (histologic chorioamnionitis: median 4693.3 pg/mL, IQR 2242.3–8054.5 vs. non-histological choriomnionitis: median 3180.9 pg/mL, IQR 1690.1–5310.1, P = 0.09).

Amniotic fluid Ang-2 concentration correlated with IL-6 in spontaneous preterm labor
In patients with spontaneous preterm labor, a weak but significant correlation was observed between amniotic fluid concentration of Ang-2 and those of IL-6 (Spearman’s rho coefficient: IL-6: 0.27, P < 0.001), but not with amniotic fluid glucose concentration or WBC count (glucose: −0.09, P = 0.09 and WBC count: 0.08, P = 0.1).

Discussion
Principal findings of the study
1) Ang-2 is a physiologic constituent of the amniotic fluid; 2) Ang-2 concentration in amniotic fluid is significantly higher in the presence of IAI in patients with preterm labor with intact membranes; 3) amniotic fluid Ang-2 concentrations correlated with those of IL-6 in patients with IAI; and 4) the amniotic fluid Ang-2 concentration decreased with advancing gestation, and did not change in spontaneous labor at term.

What is Ang-2?
Ang-2 is a 66 kDa polypeptide that contains 496 amino acids, is 60% homologous to Ang-1, and promotes angiogenesis [33]. Although, both Ang-1 and Ang-2 are encoded by genes localized on chromosome 8 [9] and share a similar protein structure, their biological activities are different. Ang-2 is classically considered as a Tie-2 antagonist, and it is accepted that at sites of vascular remodeling, Ang-2 counteracts the stabilizing action of Ang-1 by exposing the endothelium to pro-angiogenic factors such as VEGF. This antagonistic role of Ang-2 was first suggested when its overexpression resulted in the impairment of blood vessel formation in transgenic mice, a phenotype similar to the one obtained in Ang-1 and Tie-2 knockout mice [33]. However, studies with Ang-2 knockout mice suggest that this role would not be restricted to counteracting Ang-1 activities. Ang-2 may...
also act as a Tie-2 agonist, being involved in postnatal vascular remodeling events [18], because Ang-2 can activate Tie-2 and stimulate both endothelial cell migration and endothelial cell capillary-like tube formation in vitro [38, 64]. In addition to their roles in angiogenesis, angiogenic factors increase vascular permeability [3, 42] and can be expressed during the course of inflammation.

**Ang-2 concentration in amniotic fluid decreases with advancing gestational age in normal human pregnancy**

Human pregnancy is characterized by angiogenesis, tissue development and remodeling [19, 67, 70]. In the early phase of development, vascular endothelial growth factor-A (VEGF-A) and placental growth factor (PIGF) play an important role in angiogenesis [13, 15]. Human trophoblast expresses PIGF, but VEGF-A is expressed in villous and decidual macrophages [11]. Vascular endothelial growth factor interacts with vascular endothelial growth factor receptor 1 (VEGFR-1; Flt-1) and vascular endothelial growth factor receptor 2 (VEGFR-2; KDR) to promote endothelial cell proliferation, cell migration, and vascular permeability [67, 70]. In a later phase, Ang-1, Ang-2 participate in angiogenesis [44, 70].

Ang-1 and Ang-2 bind with equal affinity to Tie-2, but have different functions. Ang-1 maintains vessel integrity and plays a role in the later stages of vascular remodeling [20]. Ang-2 is a functional antagonist of Ang-1, and leads to loosening of cell/cell interactions and allows access to angiogenic inducers such as VEGF [33]. When VEGF is present, Ang-2 promotes vascular growth, but when VEGF is absent, vascular regression occurs [2, 33]. Ang-2 is selectively expressed at sites of active angiogenesis, such as the ovary, uterus, and placenta [2, 33].

Our finding of higher amniotic fluid concentration of Ang-2 in mid-trimester than that of term gestation are consistent with those of Zhang et al. [70] who, performing in situ hybridization studies, have shown that Ang-2 mRNA was readily detected in the syncytiotrophoblast in the first trimester human placenta, but at term, only low levels of Ang-2 mRNA were detected [70]. Both findings suggest that the amniotic cavity is one of the most active sites of angiogenesis during the first half of pregnancy. We propose two explanations for the decrease of amniotic fluid Ang-2 concentration with gestational age. First, normal pregnancy approach to term, a switch off of the angiogenesis process might occur. Second, there might be a dynamic transfer of Ang-2 into the fetal vessels through an increase in intramembranous absorption in the amniotic cavity that occurs at term gestation [7, 14]. We have detected Ang-2 mRNA expression in human amnion, which might be a source of amniotic fluid Ang-2 (unpublished data). Since there is a vascular pathway in the fetal surface of the placenta and within the fetal membranes which serves as the primary site for transfer of water and solutes from the amniotic compartment across the amnion into fetal blood within the placenta and fetal membranes [6, 10, 57], the decreased amniotic fluid Ang-2 concentration at term might be an effect of the dilution of the protein in the amniotic cavity because there is an increased placental water flux with advancing gestation from the mother to the fetus [4]. Aquaporin-1, a cell membrane water channel that regulates the flow of water across a variety of cell membranes, has been detected in the epithelium of the chorionic plate amnion, suggesting that aquaporin-1 rapidly facilitates water transport between the amniotic cavity and the fetal circulation [35].

**Ang-2 in amniotic fluid is increased in intra-amniotic infection and inflammation**

A novel observation of this study is that the median amniotic fluid Ang-2 concentration was increased in patients with IAI who delivered preterm. This finding supports the view that Ang-2 plays a role in inflammation, and suggests that Ang-2 participates in the host response to intrauterine infection. In addition, a significant correlation was observed between the amniotic fluid concentrations of Ang-2 and that of IL-6. IL-6 is considered a sensitive and specific marker of intra-amniotic infection [56] and/or inflammation [68].

Unlike Ang-1, which protects adult peripheral vasculature from vascular leakage and inhibits the effects of proinflammatory cytokines on endothelial cells [29, 30, 66], the effect of Ang-2 on angiogenesis is more complex and is context dependent; when Ang-2 concentrations are elevated, angiogenesis is enhanced when VEGF is present, whereas vascular regression has been observed in its absence [27].

The concept that Ang-2 may promote inflammation is based on the following findings: 1) pro-inflammatory cytokines strongly activate Ang-2 transcription in endothelial cells [26, 28, 33, 34]; 2) elevated serum concentrations of Ang-2 are present in patients with sepsis [21, 39, 42]; 3) the injection of Ang-2 protein in vivo elicits a significant increase in edema formation in the mouse paw [58]; and 4) Ang-2 deficient mice have an impaired ability to express cytokine-inducible adhesion molecules on endothelial cell surfaces after inflammatory activation [16].

Since intrauterine inflammation/infection is characterized by elevated pro-inflammatory cytokine concentration in amniotic fluid [49, 50, 56, 68], the findings of our study support a role of amniotic fluid Ang-2 in the pathologic inflammation elicited by microbial agents or its
products in the process of preterm parturition. Ang-2 acts by an autocrine mechanism [17] and is stored in endothelial Weibel-Palade bodies from where it can be rapidly released upon stimulation [16]. Inflammation exists in a mutually dependent association with angiogenesis [37, 61]. Indeed, during an inflammatory process, newly formed vessels supply the inflamed tissue with nutrients and oxygen allowing the transport of inflammatory cells. Among these, neutrophils are the first cells recruited in the angiogenic bed and provide cytokines, growth factors, and proteolytic enzymes, which contribute to regulate angiogenesis [32, 45, 59]. It has been shown also that neutrophil granules contain a variety of preformed pro-angiogenic proteins and may promote the development of laser induced chorioidal neovascularization in the retina at the early stage by providing preformed pro-angiogenic factors including VEGF, Ang-1, Ang-2, and MMP-9 [71].

Since we did not find a significant correlation between amniotic fluid Ang-2 concentration and the number of amniotic fluid WBC, Ang-2 found in the amniotic fluid is most likely to originate from other tissues, such as amnion. We could detect angiopoiein-2 mRNA expression in human amnion, and its expression was increased in the presence of chorioamnionitis (unpublished data). Our preliminary observations strongly suggest that Ang-2 could be released into the amniotic cavity from the amnion.

**Ang-2 in amniotic fluid and parturition**

Human parturition has a common pathway characterized by increased uterine contractility, cervical ripening/dilatation, and membrane/decidual activation, culminating in membrane rupture [46, 47, 51]. This activation is generally a coordinated inflammatory phenomenon in normal spontaneous labor at term [40, 65, 69]. In addition, labor at term induces gene expression changes in chorioamniotic membranes consistent with a localized acute inflammatory response, despite the absence of histologic evidence of inflammation [23]. Moreover, spontaneous labor at term, as well as pathologically induced preterm labor, result in greatly increased expression of angiogenesis-related genes in the mouse uterus [23].

Our finding that Ang-2 did not change in the presence of human term labor is unexpected and may have three possible explanations. First, physiological human parturition at term may not be associated with changes in the amniotic fluid Ang-2 concentrations because, as normal pregnancy approaches to term, a switch off of the angiogenesis process might occur [43, 62]. This explanation agrees with our finding that amniotic fluid Ang-2 increases only when there is IAI. Indeed, the mean concentration of amniotic fluid Ang-2 of patients without IAI who delivered preterm was not different than that of patients with preterm labor without IAI who delivered at term (Figure 2). A physiologic inflammatory process might be different than that of pathologic inflammation. Second, the increased concentration of the angiopoietin receptor Tie-2 in term spontaneous labor might have dampened the amniotic fluid Ang-2 concentration during labor, because the expression of Tie-2 in amnion and chorion decidua of patients with spontaneous term labor is greater than that of those at term not in labor [35]. Third, human amnion is a biologically heterogeneous and compartmentalized tissue [25] and, therefore, amniotic fluid Ang-2 production in the upper compartment may not change before the onset of labor. Further research is warranted to test these hypotheses.

**Conclusions**

We report an association between Ang-2 concentrations in amniotic fluid and intra-amniotic infection and/or inflammation. These results suggest that Ang-2 plays a role in normal gestation, as well as in preterm labor with IAI.

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