

RESEARCH ARTICLE

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Comparison of placental growth factor and fetal flow Doppler ultrasonography to identify fetal adverse outcomes in women with hypertensive disorders of pregnancy: an observational study

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Abstract

Background: Hypertensive disorders of pregnancy and intrauterine growth restriction (IUGR) are leading causes of maternal and perinatal morbidity and mortality. Failure to detect intrauterine growth restriction in women at high risk has been highlighted as a significant avoidable cause of serious fetal outcome. In this observational study we compare fetal flow using Doppler ultrasonography with a new test for placental growth factor (PIGF) to predict fetal adverse events.

Methods: Eighty-nine women with hypertensive disorders of pregnancy (24 with chronic hypertension, 17 with gestational hypertension, 12 with HELLP syndrome, 19 with preeclampsia and 17 with superimposed preeclampsia) were enrolled. A single maternal blood sample to measure free PIGF (Alere Triage) taken before 35 weeks of pregnancy was compared to the last Doppler ultrasound measurement of fetal flow before delivery. PIGF was classified as normal ($\text{PIGF} \geq 100$ pg/ml), low ($12 < \text{PIGF} < 100$) or very low ($\text{PIGF} \leq 12$ pg/ml). A positive test for abnormal fetal flow was defined as either signs of centralisation of the fetal circulation or diastolic block or reverse flow in the umbilical artery or descending aorta; this was a criterion for delivery. Fetal outcomes were intrauterine growth restriction and birth before 37 weeks of pregnancy.

Results: In total 61/89 women had a preterm birth and 22 infants had IUGR. Of those who delivered preterm, 20/20 women with abnormal fetal flow and 36/41 (87.8%) women with normal fetal flow had low or very low PIGF. Of those infants with IUGR, 22/22 had low or very low maternal PIGF and 10/22 had abnormal fetal flow.

Conclusions: PIGF may provide useful information before 35th gestational week to identify fetuses requiring urgent delivery, and those at risk of later adverse outcomes not identified by fetal flow Doppler ultrasonography.

Keywords: Hypertension, Placental growth factor, PIGF, Preeclampsia, IUGR, Doppler, Growth restriction

Background

Hypertensive disorders are one of the most common complications of pregnancy, with a prevalence of 6–22% [1,2]. These conditions are responsible for the majority of maternal and perinatal morbidity and mortality, including intrauterine growth restriction (IUGR). A recent study has shown that unrecognised IUGR is the single largest risk factor to pregnancies that end in stillbirth [3]. Research focus is now on improving antenatal

detection of fetuses at risk, to allow selective management, timely delivery and minimisation of serious outcomes.

Placental growth factor (PIGF) is a member of the vascular endothelial growth factor (VEGF) family. It is produced mainly by the placenta, and has potent pro-angiogenic effects. In normal uncomplicated pregnancy, PIGF levels rise until approximately pregnancy week 32 and then fall until delivery [4]. In pregnancies complicated by preeclampsia before the 37th week with or without IUGR, PIGF levels are significantly lower [5].

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The role of determination of sFlt-1, PlGF and other angiogenic factor levels in maternal peripheral blood in the prediction and diagnosis of preeclampsia has been extensively studied in recent years [6-20]. We recently added to the limited information available about the levels of these factors in other forms of hypertension in pregnancy [21-24] and found that free PlGF measured before 35 weeks of pregnancy may predict preterm delivery in all forms of hypertensive disorders of pregnancy [25]. There is still little information to date on their ability to predict fetal outcomes such as intrauterine growth restriction through their assessment of placental function [26].

In this study, we used a new method, the Alere Triage® PlGF test, for measuring free PlGF levels in the peripheral blood of hypertensive pregnant women before the 35th gestational week of pregnancy. We examined its prognostic efficiency regarding adverse fetal outcomes in all forms of hypertensive disorders of pregnancy, compared with currently recommended assessment of fetal wellbeing.

Methods

In this observational study, 89 women with hypertensive disorders of pregnancy (19 women with preeclampsia, 12 with hemolysis, elevated liver enzymes and low platelet count (HELLP) syndrome, 17 with superimposed preeclampsia (SIPE), 24 with chronic hypertension, and 17 with gestational hypertension) were enrolled. The study subjects were selected from the previously reported groups of hypertensive pregnant women [25] based on availability of Doppler ultrasound results. All subjects were Caucasian and resided in the same geographic area. Blood samples were taken in the First Department of Obstetrics and Gynecology, Semmelweis University, Budapest, Hungary between May 2008 and October 2010. The blood draw occurred between the 22nd and 34th completed gestational week at the time of the first routine clinical blood test and repeated only if women were reclassified. An interview was carried out with every subject after the diagnosis and again 12 weeks after the delivery, and if necessary, reclassification was done independently of the PlGF level. All subjects gave written informed consent prior to the involvement in the study. The study protocol was approved by the Regional and Institutional Committee of Science and Research Ethics of the Semmelweis University (TUKÉB 52/2008). The study was conducted in accordance with the Declaration of Helsinki.

Hypertension was diagnosed if systolic blood pressure (BP) was ≥ 140 mmHg, or diastolic BP ≥ 90 mmHg, on two occasions, six hours apart. According to the definitions of the ACOG and NHBPEP we classified the subjects' diagnoses into four groups: chronic hypertension,

gestational hypertension, preeclampsia, and superimposed preeclampsia. We separately categorised any women with hypertension, proteinuria and the syndrome of hemolysis, elevated liver enzymes and low platelet count as HELLP.

Chronic hypertension was diagnosed if high blood pressure developed prior to pregnancy or before the 20th week of gestation, or if hypertension persisted for more than 12 weeks postpartum. Gestational hypertension was applied to women who developed new-onset hypertension after the 20th week of gestation, in the absence of proteinuria, confirmed after delivery. Patients who later progressed to preeclampsia were excluded from this group, and following repeated blood draw, they were included in the pre-eclamptic group. Preeclampsia was defined as hypertension and proteinuria (≥ 300 mg / 24 hours, or $\geq +$ by urine dipstick) with an onset after the 20th gestational week. Severe preeclampsia was diagnosed if one of the following occurred: systolic BP ≥ 160 mmHg, diastolic BP ≥ 110 mmHg, proteinuria ≥ 5000 mg per 24 hours or $\geq +++$ by urine dipstick, partial HELLP syndrome, signs of renal insufficiency, pulmonary edema or threatening eclampsia. The diagnosis of HELLP syndrome was made based on characteristic laboratory findings (platelet count < 150 G/l, SGOT and SGPT > 70 U/l, LDH > 600 U/l). All of the subjects in this group also met the criteria for severe preeclampsia. Subjects with chronic hypertension who developed proteinuria after 20 weeks of gestation were categorised as superimposed preeclampsia. Subjects with preeclampsia, superimposed preeclampsia or HELLP syndrome were further grouped into early-onset (disease onset before 34th completed gestational week). Women with multiple gestations were not enrolled in this study.

A newborn was considered small for gestational age (SGA) if the birth weight was below the 10th percentile for gestational age and gender, according to a Hungarian birth weight percentile table. All neonates with SGA had an asymmetric size (normal length, but low weight for gestational age at delivery), indicating that they had intrauterine growth restriction and were not constitutionally small. In addition, none of them had fetal structural abnormalities or genetic diseases. For the determination of the abnormalities of fetal circulation we evaluated the last Doppler ultrasound results prior to delivery. Blood flow was examined in the umbilical and in the fetal middle cerebral arteries, as well as in the descending aorta of the fetus. Abnormal fetal flow was diagnosed if diastolic block or reverse flow in the umbilical artery or descending aorta, or if signs of the centralisation of the fetal circulation (increased resistance in the umbilical artery and/or descending aorta with decreased resistance in the middle cerebral artery) were present.

Our protocol for delivery dictates birth by Cesarean section, as clinically required, for women at any gestational age with HELLP syndrome, severe uncontrollable hypertension, severe proteinuria, with signs of renal insufficiency, pulmonary edema, threatening eclampsia (persistent headache and visual disturbance), low platelet count ($< 100\,000/\mu\text{l}$), elevated liver enzymes (SGOT > 70 U/l) with epigastric pain, or evidence of fetal compromise (abnormal fetal flow or pathological CTG), severe IUGR or severe oligohydramnios. In the absence of any of these factors, we continue conservative management until the 37th completed gestational week.

After each blood draw, the EDTA-anticoagulated plasma samples were immediately centrifuged at 3000 g for 10 min at 4°C and the supernatants were kept frozen at -80°C until assayed. Plasma was analysed for free PIGF using the Alere Triage® PIGF assay according to the manufacturer's instructions. Using fluorescently-labelled monoclonal antibodies against PIGF for PIGF quantification, this immunoassay is run with a single-use disposable plastic assay test cartridge in conjunction with the Triage Meter. Briefly, 250 microliter of thawed plasma (room temperature) is pipetted into the sample port of a new test cartridge. The cartridge is inserted into the meter and results are displayed in approximately 15 minutes in pg/ml. The cartridge contains chemistries for on-board positive and negative control systems. Control systems at the level of the cartridge and meter ensure that the quantitative PIGF result is valid. Calibration information is supplied by the manufacturer in the form of a lot-specific EPROM chip that is contained within each kit of devices. The measurable range of the assay is 12–3000 pg/ml. Concentrations below 12 pg/ml are value assigned based on the calibration curve, but this value is displayed to the user as a qualitative result "<12 pg/ml". Women were tested up to 34 completed weeks, as recommended in the manufacturer's product insert. The intra/ inter-assay coefficients of variation at mean concentrations of 85.2 and 1300 pg/ml were 12.1/12.8% and 11.7/13.2%, respectively.

Although PIGF concentrations fluctuate during pregnancy [4,27], original cut-off levels based on the 5th percentile of a PIGF level in a normal healthy pregnant population do not differ significantly from an absolute threshold of 100 pg/ml in women presenting with signs and symptoms of preeclampsia before the 35th week of pregnancy [Chappell et al., personal communication, submitted for publication, see Appendix for full list of researchers]. Therefore levels were classified as normal (PIGF ≥ 100 pg/ml), low (12 pg/ml $<$ PIGF < 100 pg/ml) or very low (PIGF ≤ 12 pg/ml).

Descriptive statistics were used to present the clinical characteristics. Outcomes were broken out into three PIGF groups and abnormal/ normal fetal flow,

as described above. All statistical analyses were conducted using the MATLAB version 8.0. P-values were calculated using a two-tailed Fisher Exact test of each 2 \times 2 contingency table (in this case a single PIGF cut-off of 100 pg/ml was used).

Results

In total 89 women with hypertensive disorders of pregnancy were recruited into the study. The demographics and clinical characteristics of the study participants are presented in Table 1.

By outcome, 61/89 women had a preterm birth and 22 neonates had IUGR. Table 2 shows the results of the fetal flow and PIGF tests by outcome (preterm birth and IUGR) and by diagnoses. All 20 women with abnormal fetal flow had a PIGF < 100 pg/ml so that the overall concordance between fetal flow and PIGF was high (p-value=0.0023, two tailed Fisher Exact test of the contingency table dividing the 89 women by PIGF and fetal flow).

Considering the PIGF test results for the 28 women who had both a term birth and an infant of normal birth weight, 1/1 woman with preeclampsia, 2/3 women with SIPE, 9/11 women with GHT and 5/13 women with CHT had a normal PIGF result. All of these women had a normal fetal flow.

Table 3 shows PIGF test results in women with normal and abnormal fetal flow. Of those who delivered preterm, 20/20 women with abnormal fetal flow and 36/41 women with normal fetal flow had low or very low PIGF. Five women who had a preterm delivery had both a normal fetal flow and a normal PIGF test; of these five, 2 women had an early delivery because of premature rupture of membranes. All 5 women had infants with a normal birth weight. Of the IUGR neonates, 22/22 had low or very low maternal PIGF and 10/22 had abnormal fetal flow. In the subset of 69 women who had normal fetal flow, a positive PIGF test result was significantly associated with adverse fetal outcomes (p-value < 0.0001 for preterm delivery and p-value=0.0069 for IUGR).

Twenty-four of our study participants had pathological CTG, 21 of whom had low or very low PIGF. Oligohydramnios occurred in 27 cases, 20 of them had low or very low PIGF. Of those who delivered preterm, 19/20 women with pathological CTG and 37/41 women with normal CTG had low or very low PIGF. In the preterm delivery group, 19/21 women with oligohydramnios and 37/40 women with normal amniotic fluid volume had low or very low PIGF. In the subset of 41 women with preterm delivery and normal fetal flow, 10/11 women with pathological CTG and 11/13 women with oligohydramnios had low or very low PIGF. Of the 22 IUGR neonates, 10 had pathological CTG and 10 had oligohydramnios.

Table 1 Demographics and clinical characteristics (N=89)

Diagnosis	All (N=89)	GHT (n=17)	CHT (n=24)	PE (n=19)	HELLP (n=12)	SIPE (n=17)
Maternal age (years)	33 (30–36)	34 (33–35)	33 (30–37)	33 (27–37)	30 (29–33)	34 (32–36)
Pre-gestational BMI (kg/m ²)	26.2 (23.1–33.1)	26.2 (23.1–30.8)	30 (25.9–36.2)	25.6 (22.4–29.7)	29.8 (23.7–35.5)	24.4 (21.9–30)
Primiparous women (%)	51 (57.3)	11 (64.7)	12 (50)	11 (57.9)	9 (75)	8 (47.1)
Current smoking (%)	7 (7.9)	0 (0)	2 (8.3)	1 (5.3)	2 (16.7)	2 (11.8)
Previous history of PE (%)	12 (13.5)	1 (5.9)	3 (12.5)	3 (15.8)	2 (16.7)	3 (17.6)
GA at blood draw (weeks)	32 (28–33)	33 (31–34)	33 (31–34)	30 (28–33)	29 (27–32)	31 (28–32)
GA at delivery (weeks)	34 (30–37)	38 (36–39)	37 (34–38)	31 (28–34)	29 (27–32)	32 (30–36)
Blood draw to delivery time interval (days)	8 (1–29)	29 (24–51)	20 (4–43)	1 (0–3)	1 (0–2)	8 (3–25)
Fetal birth weight (grams)	1900 (1150–3100)	3330 (2690–3860)	2700 (2105–3360)	1220 (990–1770)	1125 (845–1420)	1750 (1100–2560)
Fetal birth length (cms)	45 (39–52)	51 (48–55)	51 (45–53)	40 (37–45)	39 (36–44)	45 (40–51)
Preterm birth (%)	61 (68.5)	6 (35.3)	11 (45.8)	18 (94.7)	12 (100)	14 (82.4)
IUGR (%)	22 (24.7)	3 (17.6)	2 (8.3)	8 (42.1)	4 (33.3)	5 (29.4)
Early-onset PE (%)	44 (49.4)	0 (0)	0 (0)	17 (89.5)	11 (91.7)	16 (94.1)
Severe PE (%)	41 (46.1)	0 (0)	0 (0)	18 (94.7)	12 (100)	11 (64.7)
C-Section (%)	78 (87.6)	12 (70.6)	19 (79.2)	19 (100)	12 (100)	16 (94.1)

Continuous variables are reported as Median (IQR) and categorical variables are reported as n (%), where percent is relative to the column total. Gestational hypertension (GHT); chronic hypertension (CHT); preeclampsia (PE); hemolysis, elevated liver enzymes and low platelet count syndrome (HELLP); superimposed preeclampsia (SIPE); body mass index (BMI); gestational age (GA); intrauterine growth restriction (IUGR); Cesarean section (C-section).

Discussion

This study aimed to assess the performance of the Alere Triage® PIGF test, a rapid test for measuring PIGF levels in the maternal peripheral blood. Our study population consisted of women with chronic hypertension, gestational hypertension, HELLP syndrome, preeclampsia and superimposed preeclampsia. We earlier evaluated the diagnostic accuracy of this test in the study groups and healthy pregnant control subjects [25] and found that using a gestational age-dependent threshold of 5% of a normal population, a positive PIGF test predicted delivery before 37 weeks in over 90% of hypertensive women.

In this study we compared the diagnostic value of PIGF measured before 35 weeks using two absolute thresholds with the last Doppler ultrasonography before delivery in identifying preterm birth and IUGR neonates.

We found that PIGF concentration below 100 pg/ml identified all women with hypertensive disorders of pregnancy who required urgent delivery following an abnormal fetal flow result, and predicted all IUGR neonates, and 56/61 women who had a preterm delivery. Nearly 60% of women with normal fetal flow results had a preterm delivery. Our protocol for delivery dictated birth by Cesarean section, as clinically required, for women at

Table 2 Results of PIGF and fetal flow tests, by outcome and diagnosis (N=89)

Fetal flow	Normal PIGF		Low PIGF		Very Low PIGF	
	Normal	Abnormal	Normal	Abnormal	Normal	Abnormal
n	22	0	23	3	24	17
Percent of N	(24.7)	0.0	(25.8)	(3.4)	(27.0)	(19.1)
Preterm	5 (22.7)		14 (60.9)	3 (100.0)	22 (91.7)	17 (100.0)
IUGR	0 (0.0)		3 (13.0)	3 (100.0)	9 (37.5)	7 (41.2)
CHT	8 (36.4)		11 (47.8)	0 (0.0)	2 (8.3)	3 (17.6)
GHT	10 (45.5)		3 (13.0)	1 (33.3)	2 (8.3)	1 (5.9)
HELLP	0 (0.0)		2 (8.7)	0 (0.0)	6 (25.0)	4 (23.5)
PE	1 (4.5)		2 (8.7)	1 (33.3)	10 (41.7)	5 (29.4)
SIPE	3 (13.6)		5 (21.7)	1 (33.3)	4 (16.7)	4 (23.5)

Breakout of N=89 subjects by three placental growth factor test result groups (normal PIGF \geq 100, low 12<PIGF<100, and very low PIGF \leq 12) and two fetal flow groups (abnormal, normal). The breakout by outcomes (preterm birth, IUGR infant) and diagnosis shows the number and percent (relative to n) for each category. Intrauterine growth restriction (IUGR); chronic hypertension (CHT); gestational hypertension (GHT); hemolysis, elevated liver enzymes and low platelet count syndrome (HELLP); preeclampsia (PE); superimposed preeclampsia (SIPE).

Table 3 Adverse fetal outcomes by fetal flow (abnormal, normal) and PIGF (normal, low, very low) results (N=89)

PIGF	Abnormal fetal flow			Normal fetal flow		
	Very Low	Low	Normal	Very Low	Low	Normal
n	17	3	0	24	23	22
Percent of N	19.1	3.4	0.0	27.0	25.8	24.7
Preterm	17 (100.0)	3 (100.0)		22 (91.7)	14 (60.9)	5 (22.7)
p-value					<0.0001	
IUGR	7 (41.2)	3 (100.0)		9 (37.5)	3 (13.0)	0 (0.0)
p-value					0.0069	

Breakout of N=89 subjects by three placental growth factor groups (normal is $PIGF \geq 100$, low is $12 < PIGF < 100$, and very low is $PIGF \leq 12$) and fetal flow (abnormal vs normal). The breakout by preterm delivery and IUGR shows the number and percent (relative to n) for each category. The p-values are from a two-tailed Fisher Exact test of the 2x2 contingency table (with PIGF Low and Very Low grouped together) for each outcome (preterm and IUGR) in the subset with normal fetal flow.

any gestational age with evidence of fetal compromise including abnormal fetal flow, pathological CTG and severe oligohydramnios. The Triage PIGF test identified the majority of women with normal fetal flow who needed to be delivered preterm due to pathological CTG or oligohydramnios. Considering both test results together, no woman with a normal fetal flow and normal PIGF had an IUGR infant, and only 5 had a preterm delivery, 2 of whom were due to premature rupture of membranes.

In a small study Benton et al. [28] found that Triage PIGF differentiated placental IUGR from constitutionally small fetuses with a sensitivity of 100% and a specificity of 86%. Consistently, all of our SGA neonates were asymmetrically small with no fetal structural abnormalities or genetic diseases suggesting placental IUGR and had low or very low PIGF. In 2007 Schlembach et al. [29] correlated levels of angiogenic growth factors, including PIGF, with Doppler ultrasound parameters in women with preeclampsia and intrauterine growth restriction (suspected by ultrasound measurements and confirmed by birth weight). Maternal levels of PIGF were inversely correlated with PI values in both the umbilical and uterine arteries. PIGF levels in the umbilical vein were below the detection limit in nearly all samples of IUGR fetuses and lower than in those with preeclampsia ($p < 0.001$). In our study, we observed low or very low PIGF even in hypertensive women with normal fetal flow and an IUGR neonate.

Taylor et al. reported in 2003 that maternal circulating PIGF levels in preeclampsia are lower if accompanied by IUGR [11]. Furthermore they showed that in normotensive subjects with IUGR, PIGF concentrations are also decreased compared to controls. Our knowledge about IUGR in chronic and gestational hypertension in this regard is deficient. We found a positive PIGF test in all women with an abnormal fetal flow, all women with an IUGR neonate, the majority of women with pathological CTG or oligohydramnios, as well as in a high proportion of women carrying normal-size fetuses, many of whom

had preterm delivery. Thus, the sensitivity of the PIGF test for fetal risk (IUGR, abnormal fetal flow, pathological CTG, oligohydramnios, preterm delivery) in the hypertension group with and without proteinuria was excellent.

The potential clinical impact of these findings is that PIGF may provide useful information before 35th gestational week to identify fetuses requiring urgent delivery, and those at risk of later adverse outcomes not identified by fetal flow Doppler ultrasonography. Similarly, for all women with hypertensive disorders of pregnancy, a combination of a normal fetal flow and normal PIGF test may identify women at lower risk of adverse outcomes.

However, the weakness of our study is its retrospective observational design following normal clinical practice and the low case number. Additionally, we did not measure circulating levels of anti-angiogenic factors (sFlt-1, soluble endoglin). In this retrospective observational study of a specific target population we are unable to calculate the specificity of either test with respect to preterm delivery or IUGR infant, or to calculate negative or positive predictive values. There is a need for prospective studies to prove the safety and efficiency of the test in the clinical management of hypertension in pregnancy.

Conclusions

The Triage test is a quick reliable method for measuring circulating levels of free PIGF. The test may provide useful information before 35th gestational week to identify fetuses requiring urgent delivery, and those at risk of later adverse outcomes not identified by fetal flow Doppler ultrasonography.

Ethical approval

The study protocol was approved by the Regional and Institutional Committee of Science and Research Ethics of the Semmelweis University, Budapest, Hungary (TUKÉB 52/2008).

Appendix

Personal communication from the following researchers

Lucy C Chappell, PhD; Suzy Duckworth, MBBS; Paul T Seed, CStat; Melanie Griffin, MBChB; Lucy Mackillop, MA; Nigel Simpson, MBBS; Jason Waugh, MBBS; Dilly Anumba, MD; Louise C Kenny, PhD; Christopher W Redman, MBChB; Andrew H Shennan, MD.

Abbreviations

(NHBPEP): National High Blood Pressure Education Program Working Group on High Blood Pressure in Pregnancy; (ACOG): American Congress of Obstetricians and Gynecologists; (PIGF): Placental growth factor; (VEGF): Vascular endothelial growth factor; (sFlt-1): Soluble fms-like tyrosine kinase-1; (HELLP): Hemolysis, elevated liver enzymes and low platelet count syndrome; (SGA): Small for gestational age; (IUGR): Intrauterine growth restriction; (SIPE): Superimposed preeclampsia; (SGOT): Serum glutamic oxaloacetic transaminase; (SGPT): Serum glutamic pyruvic transaminase; (CTG): Cardiotocography; (EDTA): Ethylenediaminetetraacetic acid.

Competing interests

The authors declare that they have no competing interests.

Authors' contributions

AM conceived of the study, participated in its design and coordination, performed statistical analyses and drafted the manuscript. NG, BS and GF collected biological samples, acquired clinical data and performed PIGF measurements. BN and JR participated in the design and coordination of the study. All authors read and approved the final manuscript.

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References

1. National High Blood Pressure Education Program Working Group on high blood pressure in pregnancy: Report of the National High Blood Pressure Education Program Working Group on high blood pressure in pregnancy. *Am J Obstet Gynecol* 2000, **183**:S1–S22.
2. Walker JJ: Pre-eclampsia. *Lancet* 2000, **356**:1260–1265.
3. Gardosi J, Madurasinghe V, Williams M, Malik A, Francis A: Maternal and fetal risk factors for stillbirth: population based study. *Brit Med J* 2013, **346**:f108.
4. Knudsen UB, Kronborg CS, von-Dadelszen P, Kupfer K, Lee S-W, Vittinghus E, Allen JG, Redman CW: A single rapid point of care placental growth factor determination as an aid in the diagnosis of preeclampsia. *Pregnancy Hypertens* 2012, **2**:8–15.
5. Levine RJ, Maynard SE, Qian C, Lim KH, England LJ, Yu KF, Schisterman EF, Thadhani R, Sachs BP, Epstein FH, Sibai BM, Sukhatme VP, Karumanchi SA: Circulating angiogenic factors and the risk of preeclampsia. *N Engl J Med* 2004, **350**:672–683.
6. Hertig A, Berkane N, Lefevre G, Toumi K, Marti HP, Capeau J, Uzan S, Rondeau E: Maternal serum sFlt1 concentration is an early and reliable predictive marker of preeclampsia. *Clin Chem* 2004, **50**:1702–1703.
7. McKeeman GC, Ardill JE, Caldwell CM, Hunter AJ, McClure N: Soluble vascular endothelial growth factor receptor-1 (sFlt-1) is increased throughout gestation in patients who have preeclampsia develop. *Am J Obstet Gynecol* 2004, **191**:1240–1246.
8. Shibata E, Rajakumar A, Powers RW, Larkin RW, Gilmour C, Bodnar LM, Crombleholme WR, Ness RB, Roberts JM, Hubel CA: Soluble fms-like tyrosine kinase 1 is increased in preeclampsia but not in normotensive pregnancies with small-for-gestational-age neonates: relationship to circulating placental growth factor. *J Clin Endocrinol Metab* 2005, **90**:4895–4903.
9. Kim SY, Ryu HM, Yang JH, Kim MY, Han JY, Kim JO, Chung JH, Park SY, Lee MH, Kim DJ: Increased sFlt-1 to PIGF ratio in women who subsequently develop preeclampsia. *J Korean Med Sci* 2007, **22**:873–877.
10. Torry DS, Wang HS, Wang TH, Caudle MR, Torry RJ: Preeclampsia is associated with reduced serum levels of placenta growth factor. *Am J Obstet Gynecol* 1998, **179**:1539–1544.
11. Taylor RN, Grimwood J, Taylor RS, McMaster MT, Fisher SJ, North RA: Longitudinal serum concentrations of placental growth factor: evidence for abnormal placental angiogenesis in pathologic pregnancies. *Am J Obstet Gynecol* 2003, **188**:177–182.
12. Reuvekamp A, Velsing-Aarts FV, Poulina IE, Capello JJ, Duits AJ: Selective deficit of angiogenic growth factors characterises pregnancies complicated by pre-eclampsia. *Br J Obstet Gynaecol* 1999, **106**:1019–1022.
13. Livingston JC, Chin R, Haddad B, McKinney ET, Ahokas R, Sibai BM: Reductions of vascular endothelial growth factor and placental growth factor concentrations in severe preeclampsia. *Am J Obstet Gynecol* 2000, **183**:1554–1557.
14. Sunderji S, Gaziano E, Wothe D, Rogers LC, Sibai B, Karumanchi SA, Hodges-Savola C: Automated assays for sVEGF R1 and PIGF as an aid in the diagnosis of preterm preeclampsia: a prospective clinical study. *Am J Obstet Gynecol* 2010, **202**:40:e1–7.
15. De-Vivo A, Baviera G, Giordano D, Todarello G, Corrado F, D'Anna R: Endoglin, PIGF and sFlt-1 as markers for predicting pre-eclampsia. *Acta Obstet Gynecol Scand* 2008, **87**:837–842.
16. Ohkuchi A, Hirashima C, Suzuki H, Takahashi K, Yoshida M, Matsubara S, Suzuki M: Evaluation of a new and automated electrochemiluminescence immunoassay for plasma sFlt-1 and PIGF levels in women with preeclampsia. *Hypertens Res* 2010, **33**:422–427.
17. Kusanovic JP, Romero R, Chaiworapongsa T, Erez O, Mittal P, Vaisbuch E, Mazaki-Tovi S, Gotsch F, Edwin SS, Gomez R, Yeo L, Conde-Agudelo A, Hassan SS: A prospective cohort study of the value of maternal plasma concentrations of angiogenic and anti-angiogenic factors in early pregnancy and midtrimester in the identification of patients destined to develop preeclampsia. *J Matern Fetal Neonatal Med* 2009, **22**:1021–1038.
18. Lim JH, Kim SY, Park SY, Yang JH, Kim MY, Ryu HM: Effective prediction of preeclampsia by a combined ratio of angiogenesis-related factors. *Obstet Gynecol* 2008, **111**:1403–1409.
19. Molvarec A, Szarka A, Walentin S, Szucs E, Nagy B, Rigo J Jr: Circulating angiogenic factors determined by electrochemiluminescence immunoassay in relation to the clinical features and laboratory parameters in women with pre-eclampsia. *Hypertens Res* 2010, **33**:892–898.
20. Verlohren S, Galindo A, Schlembach D, Zeisler H, Herraiz I, Moertl MG, Pape J, Dudenhausen JW, Denk B, Stepan H: An automated method for the determination of the sFlt-1/PIGF ratio in the assessment of preeclampsia. *Am J Obstet Gynecol* 2010, **202**:161:e1–11.
21. Powers RW, Jeyabalan A, Clifton RG, Van-Dorsten P, Hauth JC, Klebanoff MA, Lindheimer MD, Sibai B, Landon M, Miodovnik M, Eunice Kennedy Shriver National Institute of Child Health Human Development Maternal-Fetal Medicine Units Network: Soluble fms-like tyrosine kinase 1 (sFlt1), endoglin and placental growth factor (PIGF) in preeclampsia among high risk pregnancies. *PLoS One* 2010, **5**:e13263.
22. Nadarajah VD, Min RG, Judson JP, Jegasothy R, Ling EH: Maternal plasma soluble fms-like tyrosine kinase-1 and placental growth factor levels as biochemical markers of gestational hypertension for Malaysian mothers. *J Obstet Gynaecol Res* 2009, **35**:855–863.
23. Khalil A, Muttukrishna S, Harrington K, Jauniaux E: Effect of antihypertensive therapy with alpha methyl dopa on levels of angiogenic factors in pregnancies with hypertensive disorders. *PLoS One* 2008, **3**:e2766.
24. Verlohren S, Herraiz I, Lapaire O, Schlembach D, Moertl M, Zeisler H, Calda P, Holzgreve W, Galindo A, Engels T, Denk B, Stepan H: The sFlt-1/PIGF ratio in different types of hypertensive pregnancy disorders and its prognostic potential in preeclamptic patients. *Am J Obstet Gynecol* 2012, **206**:58:e1–8.
25. Gullai N, Stenczer B, Molvarec A, Fugedi G, Veresh Z, Nagy B, Rigo J Jr: Evaluation of a rapid and simple placental growth factor test in hypertensive disorders of pregnancy. *Hypertens Res* 2013, **36**:457–462.
26. Chaiworapongsa T, Romero R, Korzeniewski SJ, Cortez JM, Pappas A, Tarca AL, Chaemsaihong P, Dong Z, Yeo L, Hassan SS: Plasma concentrations of angiogenic/anti-angiogenic factors have prognostic value in women

presenting with suspected preeclampsia to the obstetrical triage area: a prospective study. *J Matern Fetal Neonatal Med.* in press. <http://dx.doi.org/10.3109/14767058.2013.806905>.

27. Saffer C, Olson G, Boggess KA, Beyerlein R, Eubank C, Sibai BM, the NORMALS Study Group: Determination of placental growth factor (PIGF) levels in healthy pregnant women without signs or symptoms of preeclampsia. *Pregnancy Hypertens* 2013, **3**:124–132.
28. Benton SJ, Hu Y, Xie F, Kupfer K, Lee SW, Magee LA, von-Dadelszen P: Can placental growth factor in maternal circulation identify fetuses with placental intrauterine growth restriction? *Am J Obstet Gynecol* 2012, **206**:163:e1–7.
29. Schlembach D, Wallner W, Sengenberger R, Stiegler E, Mortl M, Beckmann MW, Lang U: Angiogenic growth factor levels in maternal and fetal blood: correlation with Doppler ultrasound parameters in pregnancies complicated by pre-eclampsia and intrauterine growth restriction. *Ultrasound Obstet Gynecol* 2007, **29**:407–413.

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