

Original Research Article

**Assessing the Median Serum Placental Growth Factor (PIGF) Levels in the First Trimester of Pregnancy regarding gestational age and maternal influencing factors from a prospective observational trial**

UNDER PEER REVIEW

## ABSTRACT

Objective: To investigate the association of risk factors, including ethnicity, maternal weight, smoking status, and assisted reproduction technology, with serum placental growth factor (PIGF) levels in the early first trimester of pregnancy.

Methods: This was a multicenter observational study involving pregnant females in the first trimester between 11+0 and 13+6 weeks of gestation. Serum PIGF concentrations were analyzed in relation to ethnicity, maternal weight, smoking status, and conception through assisted reproduction technology.

Results: A total of 367 **REVIEW THE VALUE. IN MATERIALS AND METHODS, YOU PRESENT ANOTHER QUANTITY.** patients were included in the study and had their serum PIGF levels measured. The results showed that the serum PIGF level was 4.3% higher in the Caucasian group compared to the non-Caucasian group. Smokers had 1.78 times higher levels of serum PIGF compared to non-smokers. Assisted reproduction technology was associated with lower levels of serum PIGF compared to spontaneous pregnancies. There was a negative correlation between serum PIGF levels and maternal weight, but it was statistically insignificant.

Conclusion: **Low serum PIGF levels were associated with an increased risk of pre-eclampsia. Maternal weight was negatively correlated with serum PIGF levels** **THIS PART IS WRITTEN AS RESULTS INSTEAD AS A CONCLUSIONS,** but this association was not statistically significant. High serum PIGF levels were observed in smokers and in non-Caucasians. Further research is needed to establish a concrete analysis of these findings.

Keywords: Maternal impact factors; Serum placental growth factor (PIGF); Median values

## Introduction

Currently, pre-eclampsia (PE) is a significant risk factor during pregnancy, with an occurrence rate of 3-5% [1-8] and is associated with increased morbidity, including low birth weight [2,10], and HELLP syndrome [11,12], as well as mortality [1-5,9]. PE is a pregnancy-related hypertensive disorder that can occur during the second half of pregnancy and is characterized by high blood pressure and proteinuria [2,10]. It is divided into an early-onset and late-onset form, occurring between 20-34 weeks and after 34 weeks of pregnancy, respectively [3,9]. PE is defined as a systolic blood pressure  $\geq 140$  mmHg or diastolic blood pressure  $\geq 90$  mmHg [2,10] which firstly occurs after week 20 of pregnancy and is accompanied by proteinuria protein  $\geq 3\text{g} / 24$  hours [2,10].

There are several known risks factors for PE [1,2,7], including antiphospholipid-syndrome (relative risk 9.72), diabetes mellitus (relative risk 3.56), twin pregnancy (relative risk 2.93), nulliparity (relative risk 2.91), positive family history (relative risk 2.90), high BMI (relative risk 2.47), advanced maternal age (relative risk 1.96) and a history of PE in previous pregnancies (relative risk 7.19) [15].

Several markers have been proposed as potential predictors of PE. The most commonly used is Doppler sonography of the uterine artery [1,16,17], but biochemical parameters such as free beta hCG, PAPP-A, PP-13, sFlt-1, P-selectin, NGAL, and Placenta Growth Factor (PIGF) have also been studied [1,2,12,18-20]. Out of these laboratory parameters, PIGF has shown most promising as a predictor of PE, particularly when used in combination with sFlt-1 in the second trimester [21,22], or as a single parameter in the first trimester of pregnancy [1-3].

PIGF is a member of the vascular endothelial factor family and is produced by the trophoblast cells in the placenta [1]. In normal pregnancies, PIGF concentration increases throughout pregnancy [1,2] and is influenced by a number of maternal factors, such as age, body weight, assisted reproduction technology (ART), racial origin, cigarette smoking, caffeine intake [1,2,6], or diabetes mellitus Type 1 and Type 2 [2]. However, in cases of

pathological conditions such as fetal aneuploidy and/or impaired placentation, PIGF concentrations may be reduced, leading to increased risk of PE, HELLP syndrome, or small for gestational age (SGA) [2,12]. Conversely, higher levels of PIGF are associated with reduced risk of PE and SGA [10,23].

**Aim:**

The Brahms/Kryptor is a novel analysis technique that allows for the measurement of PIGF serum concentration in the first trimester, between 11+0 and 13+6 weeks of pregnancy. In order to establish a screening test for PE based on very early PIGF serum concentration, it is necessary to compare measured concentrations to standard values. However, currently, only unpublished data are available in this regard. The aim of this study is to determine the median PIGF serum concentration values according to pregnancy age and to examine the influence of various factors on these values. **ADD THE AGE OF GESTATION**

**Material and Methods:**

This is a prospective multicenter observational study of pregnant women in the first trimester, conducted at two clinics in Germany: Women's Practice Bahnhofstrasse in Wolfenbüttel and MVZ Dr. Markus Luetge in Salzgitter. The study took place between September 2018 and December 2020 and is registered in the German Clinical Trials Registry (DRKS00014630).

Approval from local ethics boards were obtained and all enrolled patients provided written informed consent. The study population consisted of pregnant women who visited these clinics for routine monitoring and were between 11+0 to 13+6 weeks of pregnancy. Patients with pre-existing hypertension, those taking acetylsalicylic acid, or those pregnant with multiple births were excluded from participation.

Patients who did not meet the required time window in pregnancy, had miscarriages, or had incomplete data sets were also excluded from analysis.

After including only those patients who fulfilled all inclusion and none of the exclusion criteria, the study population's serum PIGF values were analyzed according to maternal weight,

**PLEASE; INCLUDE INFORMATION SUCH AS: TIME OF THE DAY THE SAMPLE WAS**

TAKEN, HOW MANY SAMMPLES PER PATIENT, WERE SEVERAL SAMPLES PER PATIENT OR JUST ONE. smoking status (yes or no), method of conception (spontaneous /assisted reproduction) and ethnic origin (Caucasian, Afro-Caribbean, Asian and Oriental).

Age depending normal WHAT IT MEANS NORMAL? ARE THE USUAL CONCENTRATION LEVELS IN PREGNANT WOMEN? OR WHICH WERE STATISTICALLY NORMAL? values were computed using various computed regression analysis methods, including linear, quadratic, cubic, exponential, logarithmic, polynomial, power, Gaussian, and hyperbolic regression analysis, with the use of the the Regression App for Mac (by Kevin Silmore v1.1) and OpenOffice software package (The Apache Software Foundation, v4.1.9).

To compare the measured PIGF values in relation to unpublished data from the test kit manufacturer (Brahms/Kryptor, Thermo-Fisher Scientific, Heringsdorf, Germany), a Student's t-test for connected samples was performed using JASP, (JASP team, Amsterdam, v0.8.1.1).

Additionally, the PIGF concentrations were transformed from raw data to multiples of median (MoM) according to the previously calculated gestational age-dependent reference values.

The PIGF-MoMs were compared between different ethnic origins (Caucasian vs. Oriental, Caucasian vs. Asian, Caucasian vs. Afro-Caribbean), smokers vs. non-smokers, and assisted reproduction technique (ART) vs. non-ART. After confirming the existence of a normal distribution using the Kolmogorov-Smirnov test, consistency among samples was checked using the Student's t-test for unconnected samples (WinSTAT, R. Fitch Software, v 2005.1). Statistical significance was assumed for a p-value <0.05.

As maternal weight is a metric scale factor, the correlation between maternal weight and PIGF concentration was analyzed using the Pearson correlation coefficient and a regression analysis (WinSTAT, R. Fitch Software, v 2005.1) after normal distribution was demonstrated again by the Kolmogorov-Smirnov test.

### **Results:**

In total, 432 cases were recruited for this prospective study. After excluding cases with missing data, such as those with missing smoking status (n=1), no status on ART (n=3), no

maternal weight status (n=6), no PIGF status (n=13), missing pregnancy week (n=1), no delivery weight status, or those who had an abortion or miscarriage (n=32), 376 cases

REVISED IT VALUE, IT IS NOT COMPARABLE WITH THE ABSTRACT VALUE remained for further analysis. THE BLUE PART SHOULD BE MOVED TO MATERIAL AND METHODS

The age range of the women study population was between 19 and 41 years, with a median MEDIAN OR MEAN?? of 30.8 years. The weight range was between 52.3 kg and 118 kg, with a median MEDIAN OR MEAN?? weight of 74.6 kg.

The mean PIGF serum concentration in correlation to pregnancy age was given as  $f(x)=0.0278531048 x^{2.826054962}$  ( $R^2=0.12$ , refer to table 1 and figure 1). IF BOTH SUPPORTS ARE REFERED TO THE SAME THING, ONE MUST BE DELETED. IT IS NOT RECOMMENDED TO REPEAT INFORMATION.

Other regression analyses using linear, quadratic, cubic, exponential, logarithmic, polynomial, power, Gaussian, and hyperbolic regression yielded in worse R-values INSERT REFERENCES.

When comparing our data to the normal values considered as normal and provided by the manufacturer, we found a good correlation (POSITIVE OR NEGATIVE?????) between the data sets, with  $p<0.001$ .

In our study group 26 cases belonged to the Oriental ethnic origin, 3 cases to the Asian ethnic origin, and 3 cases to the Afro-Caribbean ethnic origin (THESE ARE FEW OBSERVATIOS). We observed that in the Oriental population, the serum PIGF was 4.34% higher ( $p=0.63$ ) and in the Asian group, the serum PIGF values were 29.91% higher than in the Caucasians. Afro-Caribbean women had 26.01% higher PIGF levels ( $p=0.32$  ADD EE) compared to Caucasian women. THE RESULTS HAVE LITTLE RELIABILITY DUE TO THE REDUCED NUMBER OF SAMPLES.

Sixteen of 398 WHY ARE YOU TALKING ABOUT THIS AMOUNT IF YOU SAMPLE WAS LESS THAN THE VALUE MENTIONED HIRE (367 OR 376), PLEAS REVIEWED IT cases were smokers, who had 1.78-fold higher levels of PIGF ( $p=0.0003$ ).

Thirteen of 398 THE SAME OBSERVATION THAT THE PREVIOUS ONE cases had ART.

No differences could be detected between both groups.

The details of the impact factors are listed in table 2.

In our study, we found a small but measurable impact of maternal weight on PIGF levels, as demonstrated by a Pearson correlation of  $r = -0.01$  ( $R^2 = 0.0009$ ).

Regression analysis revealed that the best fit was given by  $f(x) = x / (6.908462154 + 0.979081743 \cdot x)$ . Correction factors are listed in table 3.

### **Discussion:**

Our values were consistent with those of the test kit manufacturer.

We also observed an increase in serum PIGF levels with advancing gestational age, which is in line with published data [1,2]. However, the increase was not linear [1], but rather exponential.

Kasdaglis et al. did not find a significant correlation between serum PIGF values and ethnical origin [1], unlike previous studies, that have shown higher levels in certain ethnic groups [2].

In our study population, subjects of African-Caribbean, South Asian, and East Asian origin

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CORRESPOND made up 8.8% of the total study population and had higher PIGF levels, similar to the findings of Tsiakkas et al [2]. Furthermore, we found that non-Caucasian ethnic groups had higher levels of PIGF than Caucasians; however, these results were not statistically significant due to the small sample size.

Additionally, we found that smokers had a 1.78-fold higher PIGF serum concentration compared to non-smokers, in line with previous studies [2,6,24].

We also found that assisted reproduction had no statistically significant impact on PIGF values similar to the findings of Coté [18].

Furthermore, we found a slightly negative correlation between BMI and PIGF levels, in line with previous studies [1,2,7].

**Conclusion:** Overall, our study found that several factors, including ethnic origin, nicotine consumption, and maternal weight were related to PIGF levels, but the results were mostly not significant due to small sample sizes in these subgroups.

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**Table 1:**

Gestational age (week + day)	Median PIGF concentration [pg/ml] (-2 std. dev / +2 std. dev)
11+0	24.4 (13.2 – 35.7)
11+1	25.3 (14.1 – 36.6)
11+2	26.3 (15.0 – 37.5)
11+3	27.2 (16.0 – 38.4)

11+4	28.2 (17.0 – 39.4)
11+5	29.2 (18.0 – 40.4)
11+6	30.2 (19.0 – 41.4)
12+0	31.2 (20.0 – 42.5)
12+1	32.3 (21.1 – 43.5)
12+2	33.4 (22.1 – 44.6)
12+3	34.5 (23.3 – 45.7)
12+4	35.6 (24.4 – 46.8)
12+5	36.8 (25.6 – 48.0)
12+6	38.0 (26.7 – 49.2)
13+0	39.2 (28.0 – 50.4)
13+1	40.4 (29.2 – 51.6)
13+2	41.7 (30.4 – 52.9)
13+3	43.0 (31.7 – 54.1)
13+4	44.2 (33.0 – 55.4)
13+5	45.6 (34.3 – 56.8)
13+6	46.9 (35.7 – 58.1)

Table 1: Median? OR MEAN??? PIGF value in pg/ml depending on the week of pregnancy.

Double standard deviation in brackets. THE TITLE SHOULD BE IN THE TOP OF THE TABLE.

Table 2:

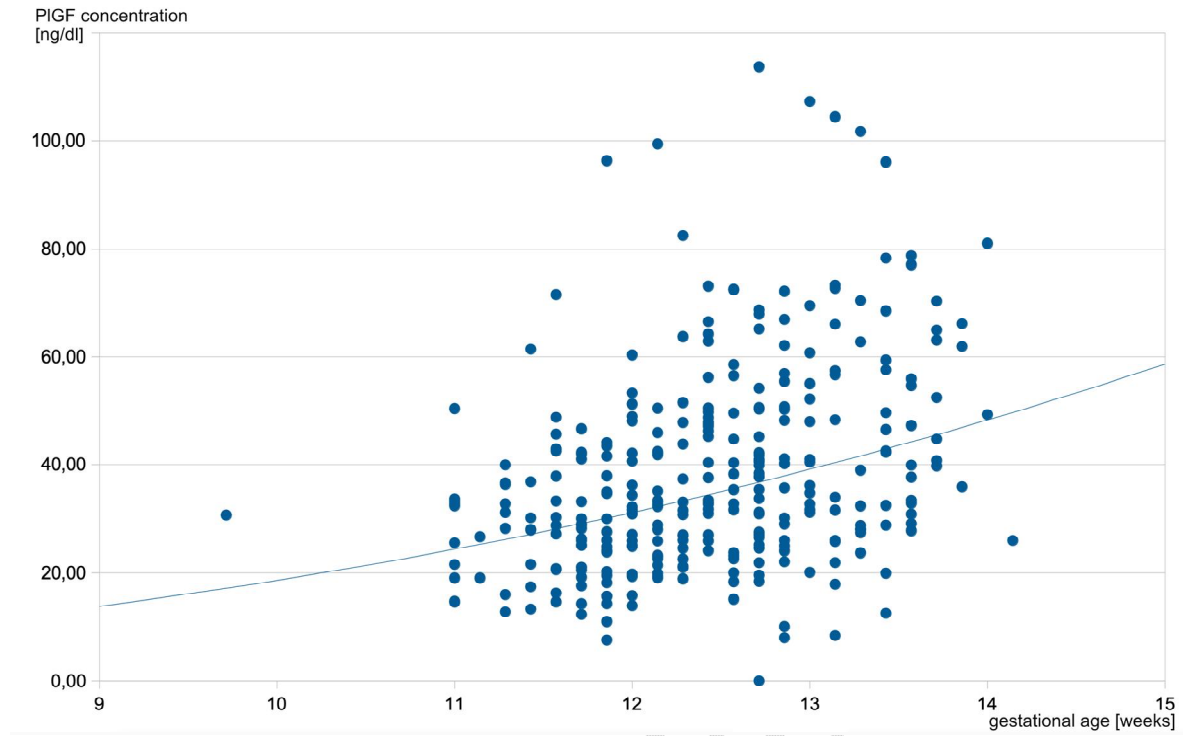
	Average MoM group 1	Average MoM group 2	Factor	p
Smoker/non-smoker	2.04 (n=16)	1.14 (n=398)	1.78	<0.0001
Asian/Caucasian	1.47 (n=3)	1.13 (n=350)	1.3	0.25
Oriental/Caucasian	1.18 (n=26)	1.13 (n=350)	1.04	0.63
Afro-Caribbean/Caucasian	1.43 (n=3)	1.13 (n=350)	1.26	0.32
ART/non-ART	0.99 (n=13)	1.18 (n=398)	0.83	0.20

Table 2: Comparison of the PIGF medians? OR MEAN regarding influencing factors with indication of the level of significance THE TITLE SHOULD BE IN THE TOP OF THE TABLE.

Table 3:

maternal weight [kg]	regressionanalysis	correction factor
50	0.895	1.042
60	0.914	1.020
70	0.928	1.005
74.12 *	0.933	1.000
80	0.939	0.994
90	0.947	0.985
100	0.954	0.978
110	0.960	0.972
120	0.967	0.967
130	0.963	0.963

Table 3: Specification of the correction factor for the maternal weight with the zero point for the correction being 74.1 kg (1.0 average correction factor). THE TITLE SHOULD BE IN THE TOP OF THE TABLE.



**Figure 1**

Figure 1: PGIF levels in ng/dl according to maternal gestational age in weeks, compared with **median???** **COULD BE MEAN???** values from Brams/Kryptor.