

PLACENTAL WEIGHT AND ITS ASSOCIATION WITH MATERNAL AND NEONATAL CHARACTERISTICS

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Abstract- Placenta plays a vital role in normal fetal development and failure of placenta to gain weight and insufficiency of its function can result in fetal disorders. We performed this study to determine placental weight and factors associated with low weight placentas. In a longitudinal cross-sectional study, women with single pregnancy, and gestational age between 37-42 weeks were studied. The subjects were categorized in high (> 750 g), normal (330-750 g), and low placental weights (< 330 g). The placental weight, birth weight, maternal age, gestational age, parity, pre-eclampsia, history of maternal diabetes, delivery approaches, infants' gender; and Apgar score in 5th minutes after delivery were examined. One thousand-eighty eight pregnant women were included in the study. The mean and standard deviation for maternal ages and gestational ages at deliveries were 25.35 ± 5.6 and 247.51 ± 9.56 days, respectively. The mean and standard deviation of neonates' weights at birth and placental weights were 3214.28 ± 529 and 529.72 ± 113 g, respectively. The prevalences of low and high placental weights were 2% and 2.8%, respectively. There were statistically significant relationships between placental weight and birth weight, fetal distress, Apgar score, maternal diabetes, pre-eclampsia and approaches of deliveries ($\alpha = 0.05$). Our findings indicate that placental weight can be associated with important variables influencing some maternal and neonatal outcomes and placental weight lower than 330 g can be a warning sign. Careful attention to placenta growth during pregnancy, for example by ultrasonography, can guide physicians to assess neonatal health.

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Key words: Birth weight, labor, placenta, pregnancy

INTRODUCTION

Placenta is the most important organ for maintaining and continuing healthy pregnancy. It transfers and exchanges oxygen and nutrition needed for fetus. The examination of placenta would demonstrate important information about whatever has happened on fetus. As fetus grows, many changes happen in placental shape and function that reflects changes in needs of fetus in different growth stages. To achieve

this, metabolic, immunologic and endocrine changes should happen in placental trophoblast.

Placental weight reflects placental development and functions and is correlated with maternal age, gestational age, history of maternal diabetes, preeclampsia, birth weight, parity, route of delivery, infants' gender and Apgar score and fetal distress. Other factors influencing placental weight include parity, maternal height and weight, and serum ferritin concentration (but not serum haemoglobin concentration) (1).

Increase in placental size is significantly associated with maternal weight, and it is an independent predictor of birth weight. Large placental size and low birth weight have been implicated as factors predicting high blood pressure

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in adulthood (1). It has been shown that maternal or fetal diseases (gestational diabetes, severe anemia, hypertension, hydrops fetalis) influence fetal and placental weight (2-5). Barker *et al.* showed the higher blood pressures have occurred in later life for men and women who had been small babies with large placentas (6). It has been shown that placental weight has a significant role in fetal growth in terms of weight, body length, and cord length but it has no significant role in the presence of meconium-stained fluid (7). While some other studies have shown less correlation between mentioned factors and placental weights, Little *et al.* showed that absolute measures of infant size and placental weight had mutual positive correlation (8).

We evaluate the prevalence of low placental weight; and determine the impact of placental weight on infants' characteristics. The results of this study would have an important implication for infants care, and decision-making for obstetrics.

MATERIALS AND METHODS

In a longitudinal cross-sectional study, from October 2000 to March 2000, in Al-Zahra Teaching Hospital, Rasht, Iran, we examined all pregnant women referred to our maternity center. Ethics committee of our institution approved the study protocol. We obtained informed consent from all participants.

Inclusion criteria were as follows: women with singleton pregnancy, and known gestational age ≥ 37 weeks. Exclusion criteria were as follows: women with unknown gestational age, intrauterine growth retardation (IUGR), and multiple pregnancies. In a pilot study after placentas expulsion, every placenta was dried, then its membrane was cut and the

placental weight measured. In addition, infants were weighted in neonatal care unit. We found mean and standard deviation (mean \pm 2 SD) as normal values 537 ± 105 g (328-744 g). Accordingly, the women were categorized as high placental weight (> 750 g), normal placental weight (330-750 g) and low placental weight (< 330 g) groups.

We used Statistical Package for Social Sciences (SPSS) for windows, version 9 for statistical analysis. Results presented in appropriate tables, and analyzes done with chi square statistical tests.

RESULTS

One thousand-eighty eight pregnant women were included in this study. Findings are represented in Tables 1, 2, and 3.

There were no significant difference between placenta weight and neonates genders; as well as between placenta weight and gestational ages ($P = 0.34$). Also, there was no significant difference between placenta weight and presence of major anomalies.

There were more low-weight placentas in under 19 years and more high-weight placentas in 30-34 years age groups. There were more low-weight placentas in preeclamptic mothers (8.1% vs. 1.6%) and more (20% vs. 2.6%) high-weight placentas in diabetic mothers. More (17.4% vs. 0.8%) low-weight placentas were found in under 2500 grams neonates while no low-weight placenta was found in over 3500 grams neonates weight group. There were lower Apgar scores in neonates with low-weight placenta and more fetal distresses and more Cesarean section in placentas with abnormal weight.

Table 1. Mean and standard deviation for some important variables

Variables	Mean	Max.	Min.	SD
Birth weight (g)	3214.3	4800	1200	529.74
Mother age (years)	25.35	45	15	5.64
Gestational age (days)	274.5	304	259	9.56
Placenta weight (g)	529.7	1200	250	113.52
Apgar score (in 5 th minute)	9.28	10	0	1.2
Parity (No.)	0.66	6	0	0.98

Abbreviation: SD, standard deviation.

Table 2. Placental weights distribution

Placenta weight group	Number	Percent
< 330 grams	22	2
330-750 grams	1036	95.2
> 750 grams	30	208
Total	1088	100

DISCUSSION

Placental hypertrophy and reduced fetal growth have been postulated to be an adaptation to maintain placental function in pregnant women with conditions such as malnutrition. As such, the pregnancy with impaired fetal growth, resulting in

small for gestational age (SGA) neonates, should have an increased placental weight to birth weight ratio (placental ratio) compared to those with appropriate for gestational age (AGA) or large for gestational age (LGA) infants; but studies have shown that this is not true (9). Heinonen *et al.* showed that placental actual weight was lower in SGA infants than in AGA infants with the same birth weight (2). It seems that low birth weight should be related to low functional tissue mass of placenta; and this is accompanied by diminution of the area for exchange between mother and fetus, both at the villi and at fetal capillary surface area. Thus, the ability of exchanging oxygen and nutrition from mother to fetus is curtailed.

Table 3. Summary of findings analyses (n=1088)*

Variables categories	Placental weight group Delivery time				P values
	< 330 g	330- 750 g	>750 g	Total	
Mothers parities	0	14	604	12	0.95 > P > 0.90
	1	6	278	12	
	2	2	92	2	
	3+	0	62	4	
	total	22	1036	30	
Mothers age groups	< 19	6	164	2	0.95 > P > 0.90
	20-24	8	334	4	
	25-29	4	280	12	
	30-34	2	184	10	
	35-39+	2	74	2	
	Total	22	1036	30	
Mothers preeclampsia (PIH)	Yes	6	64	4	0.0002
	No	16	968	26	
	Total (100%)	22	1032	30	
Mothers diabetes	Yes	0	8	2	0.00035
	No	22	1028	28	
	Total	22	1036	30	
Neonates weight groups	<2500 grams	16	76	0	< 0.0001
	2500-3500 grams	6	702	10	
	>3500 grams	0	258	20	
	Total	22	1036	30	
Apgar scores (5 th minutes)	0-8	8	66	2	< 0.00035
	8-10	14	967	28	
	Total (100%)	22	1033	30	
Fetal distress	Yes	8	188	8	0.05
	No	14	848	22	
	Total	22	1036	30	
Delivery approaches	Normal vaginal	4	610	10	0.00001
	Cesarean section	18	420	20	
	Total	22	1030	30	

* Data are given as number.

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Our findings showed that the prevalence of low weight placenta in SGA infants was 17.4%; but this is not true in infants with weight more than 3500 g ($P < 0.0001$). Furthermore, fetal growth capacity is determined by placental weight (7). The placental ratio tended to increase from the LGA group to the SGA group both for women with uncomplicated pregnancies and with pregnancies complicated by intrauterine growth retardation or for women with pregnancy induced hypertensions (3). Perry *et al.* showed there was no significant difference between the mean placental weight and preeclampsia and gestational hypertension (4). The pathophysiology of hypertension in pregnancy is unknown, and current therapies for preeclampsia aimed preventing the maternal syndrome, not preventing the primary pathophysiology but genetic and environment factors may caused this complication (10). Bortolus *et al.* have shown that pregnancies with induced hypertension result high weight placentas (11). Our findings indicate that not only the prevalence of high placental weight is higher in preeclampsia but low placental weight prevalence also in preeclampsia is higher than uncomplicated pregnancies. As if, the mean placental weight showed a significant increase from the SGA to the LGA. Our results were similar to this research.

One of the principal reasons for this problem in our study is because of the more nulliparous women; and nulliparity is a risk factor for high prevalence of this disease. Preeclampsia incidence according to our results and Perry's study on the healthy nulliparous pregnant women was 6.8% and 7.3%, respectively (4).

We found the prevalence of gestational diabetes mellitus (GDM) 0.9%; much lower than other studies; perhaps because we had more younger subjects than other investigators (with 75% < 29 years), as GDM is more common after 30 years old (12); also perhaps because insufficient screening test, and early detections (13).

One of the most common complications of GDM is macrosomic babies (> 4 Kg. and more babies weights) (14). Our results showed the adaptation of placenta with baby weight; because low placental weight was not seen in GDM women.

One of the influencing factors on placental weight is gestational age. Gestational age is known as a principal and determinant factor of placental weight. Kinare *et al.* stated that the capacity of fetal weight growth is determined by placental growth (15), and between placental volume in 15 to 18 weeks gestation ages to placental weight at birth and birth weight found significantly correlation. Molteni *et al.* have shown the average placental weight is related to gestational age (16). Placental weight increases in infants proportionately with gestational age. While in infants with SGA were not seen any change from 36 weeks gestation (12). Pardi *et al.* entitled there is a significant difference between placental weight in the second trimester gestation with placental weight at birth (7); but Lo *et al.* have proved there is no significant difference between placental weight and gestational age (7). Our data were similar to Lo's research perhaps because more admission pregnant women in term that placental growth was the same for all of them.

We found indirect correlation between Cesarean section and placental weight; perhaps because one of Cesarean section indications is fetal distress that consists of meconium defecation. Meconium defecation is affected on Apgar score. The result of Lao *et al.* study was vice versa (18). In their study, there were no significant differences in placental weight or birth weight between neonates with presence or absence of meconium stained amniotic fluid. In other research, although low Apgar score wasn't observed in high placental weight there were no significant differences between low Apgar score and high placental weight (19).

In different studies it has been demonstrated that male fetuses gain significantly greater weight as compared to female fetuses from 34th to 42nd week of gestation. Because placental weight has a relationship with infant weight, there should be a significant correlation between infant sex and placental weight (7, 19, 20). However, we found no correlation, perhaps due to differences in ethnic and/or genetic factors. In our research the prevalence of low- and high- weight placentas in fetuses under distress were 3.9% and 2.9%, respectively. In addition, the ratio of women with low placental weight who had Cesarean section to women with

natural vaginal delivery was 3.9/0.6. This ratio was 4.4/1.6 for high placental weight. Burkhardt *et al.* found mean placental weight from vaginal deliveries to be 76 g lighter than from Cesarean sections (21). The prevalence of low and high weight placentas from Cesarean section labors were more than from normal vaginal labors. This is corresponded with the prevalence of fetus distress, that consequently causes Apgar score decline especially in cases of low placental weight. It should be noted that we could not gather some variables such as CBC and hemoglobin during pregnancy; as well as body mass index in early pregnancy because most women have had prenatal cares in different health centers in other counties and villages. Godfrey *et al.* showed maternal hemoglobin levels influences placental weight, as if anemia and iron deficiency during pregnancy are associated with large placental weight (5).

Placenta is essential for normal fetal development and failure of the placenta can result in fetal problems. Because of growing evidence for a correlation of placental weight with chronic diseases in later life, we suggest attention and correct examination of placenta and recording all of the observations in patients' files as an important evidence for future. With evaluation and follow up of placenta growth in early pregnancy, we can prevent the risks for fetal life and improves infant health.

Conflict of interests

The authors declare that they have no competing interests.

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