



Rate of Fetal Macrosomia with Maternal and Early Neonatal Complications in Internally Moved People Affected by Violence

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ABSTRACT

Background: Fetal macrosomia is usually distressing to obstetricians and neonatologists. In the current study, involved mothers had poor social and medical circumstances, as they were migrated forcefully within the country borders due to war, from their original homeland to safer camps which had miserable situations. **Objectives:** To study rate, risk factors, and complications of macrosomia in people with low socio-economic living conditions and missed medical follow up. **Methods:** All internally displaced pregnant women who gave birth to neonates weighed ≥ 4000 g were involved in the study. All required history, examination, care, and investigations were practiced by the attending obstetrician and neonatologist. Cases of normal birth weight neonates from the same sample of internally moved mothers were considered as controls. **Results:** Fetal macrosomia rate was 15.77% (143 out of 907). Observed significant macrosomia risk factors were maternal age ≥ 30 years, multiparity, body mass index ≥ 30 , previous or family history of macrosomia, gestational age > 40 weeks, cesarean section, diabetes, and hypertension. Meconium-stained liquor, shoulder dystocia, uterine atony, and genital trauma, were major maternal complications, while main neonatal sequelae were Apgar score (> 7) at first minute, birth asphyxia, admission to NICU, hypoglycemia, polycythemia, and respiratory distress. **Conclusion:** Higher rate and more frequently encountered risk factors of macrosomia than national and international figures found in our sample (of forcefully moved mothers) were probably related to poor living circumstances, and absence of regular medical follow up with antenatal care.

Keywords: Macrosomia, Rate, Sequelae, Forcefully displaced mothers, Poor medical care, Socio-economic state

INTRODUCTION

Exaggerated intrauterine fetal growth or birth weight ≥ 4000 g could define macrosomia, but in fact, no neonatal birth weight parameter has been used to identify macrosomia, including > 4000 g to 4500 g, and $> 90^{\text{th}}$ - 95^{th} centile for gestational age and sex [1,2].

Delivering a giant baby is distressing to the mother, her baby, obstetrician, and neonatologist. It may lead to unfavorable outcomes during the whole process started from pregnancy through delivery and finally after giving birth [3].

Maternal risk factors with certain diseases are a major key that assists macrosomia occurrence, leading to different complications affecting both mother and her baby [2].

Little attention has been offered to high birth weight infants even though they have elevated perinatal morbidity and mortality rates. It is important to study fetal macrosomia in order to aid in better management of affected pregnancies, and anticipate their sequelae [4,5].

Objectives

To look deeply into the issue of fetal macrosomia regarding its rate, risk factors, and complications (maternal or neonatal), in mothers forcefully displaced because of violence, and lived in camps and habitations with bad circumstances and poor medical services.

PATIENTS AND METHODS

This prospective work took place at Medical City Hospital, Department of Obstetrics and Gynecology, and in neonatology unit, in Baghdad, Iraq, from 1st of December 2017 till the end of May 2018. During the above 6 months, only pregnant ladies who visited the hospital aiming to deliver their babies and forcefully left their own houses and land were involved.

These women came from internally moved or displaced families to safer places or camps within the country, because of the war that broke out in their areas. Although these camps were away from violence, they had low social, medical, and environmental living conditions. These pregnant ladies did not get regular antenatal care.

After identification of mother's background address (moved or native), and after delivery all neonates (of internally migrated mothers) were weighed. Only macrosomic term (≥ 37 weeks of gestation) newborns ≥ 4000 grams were recruited in the study with their mothers, twins, and triplets were excluded from the study [6].

Newborns delivered to single term mothers (≥ 37 weeks of gestation) with birth weights 2500-3999 grams (g), were randomly selected as a control group after matching of the sex of babies, in a ratio of 3 controls per one case (of macrosomia). An informed consent was obtained from all mothers and/or caregivers.

Relevant data were collected by the attending obstetrician with full history. In addition, all pregnant women were sent routinely for serum blood sugar (once or more as recommended by the available obstetrician), it should have been ≥ 92 mg/dl for fasting samples or ≥ 180 mg/dl for 60 minutes' postprandial, to diagnose diabetes when no history was given [7].

Also, blood pressure was measured for all pregnant ladies; the cut-line to diagnose hypertension was $\geq 140/90$ mmHg [8].

Postpartum hemorrhage was defined as an excessive blood loss of >500 milliliters after delivery [9]. Shoulder dystocia defined as "requirement of additional obstetric maneuvers when gentle downward traction has failed to affect the delivery of the shoulders" [10].

Body mass index (BMI) was evaluated for all involved women based on height and weight. Results were divided into two categories, either <30 (under, normal or over-weight) or ≥ 30 (obese) [11].

Apgar score was evaluated by the attending neonatologist within 1 minute and 5 minutes after birth. According to the hospital policy, a score of 7-10 needed no intervention; a score of 4-7 might necessitate some prompt and temporary resuscitation parameters, while the least score of 0-3 required immediate aggressive resuscitation including intubation and admission to intensive care unit of neonates [12].

Within the first hour after delivery, all early neonatal macrosomia-related complications were estimated through clinical examination and/or radiological assessment.

A heel pricks capillary blood sample was taken from all involved neonates to test hypoglycemia (<47 mg/dl), and polycythemia [13,14].

This study was approved by a scientific and ethical committee of the College of Medicine, and Al-Kindy College of Medicine, University of Baghdad, Iraq. All procedures performed in the current study were in accordance with the 1964 Helsinki declaration and its later amendments or comparable ethical standards.

Statistical Analysis

Statistical package for the social sciences (SPSS) version 22 was used to analyze the data; statistical tests were utilized as needed. A p-value ≤ 0.05 was considered as statistically significant. Univariate analysis of risk factors was performed by conditional logistic regression so that we could identify how powerful the association was with the outcome.

Analysis through multivariate conditional logistic regression was done on variables that appeared significantly in the previous univariate analysis. Binary logistic regression approach was helpful in analyzing the maternal or the fetal sequelae. Odds ratio (OR) with 95% confidence intervals (CIs), were evaluated for both the risk factors and the complications.

RESULTS

During the study period, the total number of women gave birth to a single newborn in our hospital was 3248. We found 907 (27.92%) women were internally displaced (moved) due to fighting. Of them, 143 (15.77%) women delivered macrosomic babies (≥ 4000 grams), which could express the prevalence rate of macrosomia in the current sample. For each woman with macrosomia, another 3 pregnant females without macrosomia were randomly selected from the same total number of internally moved mothers (907), which was named as the control group, to compare it with the case group (with macrosomia).

Case and control groups had 143, 429 pregnant ladies, respectively. Only 1 macrosomic baby within our sample died because of birth asphyxia. His mother was diabetic and obese. Maternal age of 30 years and more, multiparity, obese mothers, previous history or family history of macrosomia, postdate pregnancy, cesarean section delivery, maternal diabetes mellitus, and mother's hypertension, were statistically significant risk factors for fetal macrosomia (Table 1).

Table 1 General mother's risk factors

Risk factor	Case (macrosomia) group (143) n (%)	Control (normal birth weight) group (429) n (%)	Odds ratio	95% confidence interval (95% CI)	p-value
Maternal age					
<18 years	12 (8.39%)	43 (10.02%)	1.000	-	-
18->30 years	55 (38.46%)	270 (62.94)	0.3681	0.2493 to 0.5435	0.0001
≥ 30 years	76 (53.15%)	116 (27.04%)	3.0607	2.0692 to 4.5274	0.0001
Multiparity	122 (85.31%)	176 (41.03%)	8.3512	5.0580 to 13.7885	0.0001
BMI (kg/m²)					
<30 (under, normal or over-weight)	52 (36.36%)	301 (70.16%)	0.243	0.1631 to 0.3620	0.0001
≥ 30 (Obese)	91 (63.64%)	128 (29.84%)	4.1152	2.7624 to 6.1305	0.0001
Smoking	5 (3.50%)	21 (4.90%)	0.7039	0.2605 to 1.9025	0.4889
Previous mother's history of macrosomia	30 (20.98%)	5 (1.17%)	22.5133	8.5409 to 59.3438	0.0001
Family history of macrosomia	21 (14.69%)	4 (0.93%)	18.2889	6.1610 to 54.2905	0.0001
Gestational age >40 weeks	63 (44.06%)	15 (3.50%)	21.735	11.7877 to 40.0766	0.0001
Mode of Delivery					
Cesarean section	133 (93.01%)	295 (68.76%)	6.0414	3.0780 to 11.8576	0.0001
Normal vaginal	10 (6.99%)	134 (31.34%)	0.1655	0.0843 to 0.3249	0.0001
Diabetes mellitus	63 (44.05%)	18 (4.19)	17.9813	10.1081 to 31.9867	0.0001
Hypertension	36 (25.17%)	52 (12.12%)	2.4393	1.5153 to 3.9267	0.0002

Meconium-stained liquor, shoulder dystocia, uterine atony, and genital tract trauma were statistically significant obstetrical risk factors of macrosomia (Table 2). Regarding genital tract injury, only perineal tear caught significance level.

Table 2 Obstetrical maternal complications

Event	Macrosomia group (143) n (%)	Control group (429) n (%)	Odds ratio	95% confidence interval (95% CI)	p-value
Meconium-stained liquor	37 (25.87%)	30 (6.99%)	4.6425	2.7407 to 7.8637	0.0001
Shoulder dystocia	19 (13.29%)	3 (0.70%)	21.7581	6.3344 to 74.7366	0.0001
Post-partum hemorrhage	5 (3.50%)	4 (0.93%)	3.8496	1.0194 to 14.5376	0.0468
Atony of uterus	18 (12.59%)	6 (1.40%)	10.152	3.9449 to 26.1258	0.0001
Genital tract injury	9 (6.29%)	3 (0.70%)	9.5373	2.5452 to 35.7382	0.0008
Perineal tear	5 (3.50%)	2 (0.47%)	7.7355	1.4840 to 40.3208	0.0052
Vaginal lacerations	3 (2.10%)	1 (0.23%)	9.1714	0.9463 to 88.8844	0.0558
Cervical lacerations	1 (0.69%)	0 (0.00%)	9.0421	0.3663 to 223.2210	0.1783

Table 3 concentrated on neonates. Apgar score (<7) at first minute, birth asphyxia, admission to neonatal intensive care unit (NICU), hypoglycemia, polycythemia, and respiratory distress were significantly associated with macrosomia.

Table 3 Major neonatal sequelae due to macrosomia and general characteristics

Outcome	Macrosomia group (143) n (%)	Control group (429) n (%)	Odds ratio	95% confidence interval (95% CI)	P value
Brachial plexus injury	2 (1.41%)	0 (0.0%)	15.1767	0.7243 to 318.0207	0.0797
Skeletal fractures (clavicle or long bones)	1 (0.69%)	0 (0.0%)	9.0421	0.3663 to 223.2210	0.1783
Apgar score (<7) at					
One minute	10 (6.99%)	4 (0.93%)	7.9887	2.4651 to 25.8891	0.0005
Five minutes	1 (0.69%)	3 (0.70%)	1.0000	0.1032 to 9.6907	1.0000
Birth asphyxia	19 (13.29%)	9 (2.1%)	7.1505	3.1556 to 16.2032	0.0001
Admission to neonatal intensive care unit (NICU)	17 (11.89%)	22 (5.19%)	2.496	1.2853 to 4.8473	0.0069
Hypoglycemia	36 (25.17%)	22 (5.13%)	6.2243	3.5145 to 11.0233	0.0001
Polycythemia	8 (5.60%)	1 (0.23%)	25.363	3.1437 to 204.6226	0.0024
Respiratory distress	18 (12.59%)	19 (4.43%)	3.1074	1.5820 to 6.1034	0.0010
Sex of newborn					
Female	48 (33.57%)	188 (43.82%)	0.6477	0.4359 to 0.9624	0.0316
Male	95 (66.43%)	241 (56.18%)	1.5439	1.0390 to 2.2941	0.0316

DISCUSSION

Fetal macrosomia is an identifiable challenge in current practice due to associated fetal and maternal complications [2]. Nevertheless, while some of the known risk factors and complications of macrosomia are preventable, there is a worldwide increase in its incidence [4].

There are many publishes throughout the world talk about macrosomia, but herein this study we tried to involve a unique sample of forcefully displaced pregnant women (with their macrosomic babies) to find out if they had the same local or global rates, risk factors, and complications, as most of them had been settled in exceptional conditions of living which might have an impact on the outcome results.

We found that the prevalence rate of fetal macrosomia was (15.77%) which is considered higher than the others average rate. It was (1.4%) in a previous local Iraqi paper and in Turkey, it was (5.2%), while in Hawaii it was (10.9%) [15-17].

These differences in rates may be related to differences in race, ethnicity, climate, and genetic influences [17]. Based on our data, low socio-economic factors with poor medical care might have an influence on such higher rates.

Age of the pregnant mother (≥ 30 years) was a strong risk factor for fetal macrosomia. It was mentioned in another paper, this may be due to the fact that with increasing mother's age, there may be an effect on metabolism, thereby accelerating the velocity of growth in the baby [4,18].

Multiparity and increased maternal body mass index were noticed significantly with macrosomia, as said by other researchers with a higher incidence of occurrence [19]. This may be explained by lower socio-economic state and poorer nutritional background related to difficult living conditions seen in our pregnant women, they were apparently slimmer and with less parity than usual local pregnant ladies.

Previous mother's history or family history of macrosomia were found to have a significant relationship as a risk factor. This opinion was advocated by some workers [20]. Diabetes was historically one of the most famous risk factors of macrosomia and considered as an independent risk factor, it occurred in (44.05%) of our women, which is higher than what was reported by Chinese authors (26%) [10,21].

Our internally displaced women could not reach their obstetricians for antenatal care regularly, and they missed educational sessions about low glycemic index lifestyle during pregnancy, in addition to the virtual hardness of having highly-priced diet food, all these factors, added to racial, social, and environmental parameters, may be behind our elevated rates of diabetes in macrosomia cases.

Maternal complications significantly related to obstetrical events in our sample were meconium-stained liquor, shoulder dystocia, atony of uterus, and genital tract injury [22,23]. Meconium stained liquor occurred in (25.87%),

while shoulder dystocia took place in (13.29%). These results were compared to another local study that mentioned (29.8%) and (13.4%), respectively [15].

Neonatal brachial plexus injury and skeletal fractures were found mainly in macrosomic babies (1.41% versus zero, and 0.69% versus zero) respectively. Nearly similar results were mentioned by other papers [24,25]. Apgar score (<7) at one minute was significantly complicating macrosomic newborns, occurred in (6.99%), which is just close to what was found by reporters from Denmark [26].

Other significant neonatal complications were birth asphyxia, admission to neonatal intensive care unit (NICU), hypoglycemia, polycythemia, and respiratory distress [27,28]. We noticed that rates of maternal and neonatal complications found within the current study were in parallel with other published data, this may be related to optimum (or near optimum) medical care offered by our involved hospital during delivery, and shortly after that.

Some limitations were encountered that adversely affect this study, like its short duration (just 6 months), data collected in a single setting, non-availability of some laboratory investigations such as serum electrolytes (for neonates), and some cases of unnoticed gestational diabetes could be missed by absence of glucose tolerance test (in spite of a routine maternal blood sugar sampling on admission).

The prospective approach of this study, sample population type (which was studied for a first time, up to our knowledge), and a cooperative team (obstetricians and neonatologists) who dealt promptly with complications occurred in both sides (maternal and neonatal), represent strong aspects of this study.

CONCLUSION

After a detailed inspection of our results in contrast to others throughout the globe, the finding of higher rates of macrosomia, with more frequently seen risk factors in forcefully moved people with unhealthy living conditions, should draw the attention of obstetricians, neonatologists, medical staffs, and policymakers. We suggest setting regular visits of health care professionals to camps or interim places (where these mothers live) before and during pregnancy, to educate them about the most important prophylactic health parameters, encourage them to protect themselves and their future babies from macrosomia, and screening for any unnoticed relevant disease such as diabetes or hypertension.

DECLARATIONS

Conflict of Interest

The authors declared no potential conflicts of interest with respect to the research, authorship, and/or publication of this article.

REFERENCES

- [1] Malin, G. L., et al. "Antenatal magnetic resonance imaging versus ultrasound for predicting neonatal macrosomia: a systematic review and meta-analysis." *BJOG: An International Journal of Obstetrics and Gynaecology*, Vol. 123, No. 1, 2016, pp. 77-88.
- [2] Allen K and Wallace SV. "Fetal macrosomia." *Obstetrics, Gynaecology and Reproductive Medicine*, Vol. 23, 2013, pp. 185-88.
- [3] Levitt, Lorinne, et al. "720: Macrosomia-self-fulfilling prophecy." *American Journal of Obstetrics & Gynecology*, Vol. 216, No. 1, 2017, pp. 420-21.
- [4] Said, Aisha Salim, and Karim Premji Manji. "Risk factors and outcomes of fetal macrosomia in a tertiary center in Tanzania: a case-control study." *BMC Pregnancy and Childbirth*, Vol. 16, No. 1, 2016, p. 243.
- [5] Galvin, Daniel M., et al. "94: Accuracy of prenatal detection of macrosomia >4,000 g and outcomes in the absence of intervention: results of the prospective multicenter genesis study." *American Journal of Obstetrics and Gynecology*, Vol. 216, No. 1, 2017, p. 68.
- [6] Kamana, K. C., Sumisti Shakya, and Hua Zhang. "Gestational diabetes mellitus and macrosomia: a literature review." *Annals of Nutrition and Metabolism*, Vol. 66, No. 2, 2015, pp. 14-20.
- [7] Kerner, W., and J. Brückel. "Definition, classification and diagnosis of diabetes mellitus." *Experimental and Clinical Endocrinology and Diabetes*, Vol. 122, No. 7, 2014, pp. 384-86.

- [8] Watanabe, Kazushi, et al. "Outline of definition and classification of pregnancy-induced hypertension (PIH)." *Hypertension Research in Pregnancy*, Vol. 1, No. 1, 2013, pp. 3-4.
- [9] Kerr, Robbie, et al. "Postpartum hemorrhage: Case definition and guidelines for data collection, analysis, and presentation of immunization safety data." *Vaccine*, Vol. 34, No. 49, 2016, p. 6102.
- [10] Hansen, Alexandra, and Suneet P. Chauhan. "Shoulder dystocia: definitions and incidence." *Seminars in Perinatology*, Vol. 38. No. 4. WB Saunders, 2014.
- [11] Flegal, Katherine M., et al. "Association of all-cause mortality with overweight and obesity using standard body mass index categories: a systematic review and meta-analysis." *Jama*, Vol. 309, No. 1, 2013, pp. 71-82.
- [12] Iliodromiti, Stamatina, et al. "Apgar score and the risk of cause-specific infant mortality: a population-based cohort study." *The Lancet*, Vol. 384, No. 9956, 2014, pp. 1749-55.
- [13] Tin, Win. "Defining neonatal hypoglycemia: a continuing debate." *Seminars in Fetal and Neonatal Medicine*. Vol. 19. No. 1. WB Saunders, 2014.
- [14] Alsafadi, Tariq Rushdi Mohieldeen, et al. "Polycythemia in a neonatal intensive care unit, risk factors, symptoms, pattern, and management controversy." *Journal of Clinical Neonatology*, Vol. 3, No. 2, 2014, p. 93.
- [15] Alrubae, Methal-A., and Kloud Jafer. "Macrosomia; Risk factors and labor outcome." *The Medical Journal of Basrah University*, Vol. 28, No. 2, 2010, pp. 77-84.
- [16] Usta, Akin, et al. "Frequency of fetal macrosomia and the associated risk factors in pregnancies without gestational diabetes mellitus." *The Pan African Medical Journal*, Vol. 26, 2017.
- [17] Tsai, Pai-Jong Stacy, Emily Roberson, and Timothy Dye. "Gestational diabetes and macrosomia by race/ethnicity in Hawaii." *BMC Research Notes*, Vol. 6, No. 1, 2013, p. 395.
- [18] Li, Yi, et al. "Weight gain in pregnancy, maternal age and gestational age in relation to fetal macrosomia." *Clinical Nutrition Research*, Vol. 4, No. 2, 2015, pp. 104-09.
- [19] Alberico, Salvatore, et al. "The role of gestational diabetes, pre-pregnancy body mass index and gestational weight gain on the risk of newborn macrosomia: results from a prospective multicentre study." *BMC Pregnancy and Childbirth*, Vol. 14, No. 1, 2014, p. 23.
- [20] Mohammadbeigi, A., et al. "Fetal macrosomia: risk factors, maternal, and perinatal outcome." *Annals of Medical and Health Sciences Research*, Vol. 3, No. 3, 2013, pp. 546-50.
- [21] He, Xiu-Jie, et al. "Is gestational diabetes mellitus an independent risk factor for macrosomia: a meta-analysis?" *Archives of Gynecology and Obstetrics*, Vol. 291, No. 4, 2015, pp. 729-35.
- [22] Young, Brett C., and Jeffrey L. Ecker. "Fetal macrosomia and shoulder dystocia in women with gestational diabetes: risks amenable to treatment?" *Current Diabetes Reports*, Vol. 13, No. 1, 2013, pp. 12-18.
- [23] Fuchs, Florent, et al. "Adverse maternal outcomes associated with fetal macrosomia: what are the risk factors beyond birthweight?" *BMC Pregnancy and Childbirth*, Vol. 13, No. 1, 2013, p. 90.
- [24] Hammad, Ibrahim A., et al. "Neonatal brachial plexus palsy with vaginal birth after cesarean delivery: a case-control study." *American Journal of Obstetrics and Gynecology*, Vol. 208, No. 3, 2013, p. 229.
- [25] Cheng, Yvonne Kwun-Yue, and Terence T. Lao. "Fetal and maternal complications in macrosomic pregnancies." *Research and Reports in Neonatology*, Vol. 4, 2014, pp. 65-70.
- [26] Ovesen, Per Glud, et al. "Maternal and neonatal outcomes in pregnancies complicated by gestational diabetes. A nation-wide study." *The Journal of Maternal-Fetal & Neonatal Medicine*, Vol. 28, No. 14, 2015, pp. 1720-24.
- [27] Lipschuetz, Michal, et al. "A large head circumference is more strongly associated with unplanned cesarean or instrumental delivery and neonatal complications than high birthweight." *American Journal of Obstetrics and Gynecology*, Vol. 213, No. 6, 2015, p. 833.
- [28] Åberg, Katarina, et al. "Vacuum extraction in fetal macrosomia and risk of neonatal complications: a population based cohort study." *Acta Obstetrica et Gynecologica Scandinavica*, Vol. 95, No. 10, 2016, pp. 1089-96.