

Mitigating Long-Term Risks: Supporting Infants with Macrosomia to Promote Healthy Development

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DESCRIPTION

Macrosomia, a condition characterized by excessive birth weight, poses unique challenges for both infants and healthcare professionals. This article aims to provide a comprehensive overview of infants with macrosomia, exploring its causes, potential complications, and the essential steps to ensure optimal care for these new-borns.

Macrosomia refers to infants who are born with a birth weight significantly higher than the average for their gestational age. The condition is typically diagnosed when an infant weighs over 4,000 grams (8 pounds, 13 ounces) at birth. Macrosomia is relatively common, with an estimated prevalence of 10% to 15% of all births globally. Several factors contribute to the development of macrosomia, including maternal diabetes, excessive maternal weight gain, and genetic factors.

Maternal diabetes, particularly gestational diabetes, is one of the leading causes of macrosomia. When a pregnant woman's blood sugar levels are elevated, the fetus is exposed to excess glucose, leading to increased insulin production and subsequent overgrowth. Maternal obesity is another significant risk factor for macrosomia, as excess maternal weight gain can contribute to the accumulation of adipose tissue in the fetus.

Genetic factors also play a role in macrosomia. Certain ethnicities, such as individuals of African, Hispanic, or Native American descent, have a higher predisposition to delivering macrosomic infants. Additionally, family history of large birth weight or a previous macrosomic delivery increases the likelihood of having a macrosomic infant.

While macrosomia itself may not always result in immediate complications, it can increase the risks associated with childbirth for both the infant and the mother. Macrosomic infants are more likely to experience birth trauma, including shoulder dystocia, where the baby's shoulder becomes stuck behind the mother's pelvic bone. This complication can lead to nerve injuries, fractures, and other birth-related complications.

Furthermore, mothers delivering macrosomic infants are at a higher risk of experiencing postpartum haemorrhage and perineal trauma during delivery. The size disparity between the infant and the birth canal can necessitate the use of assisted delivery techniques, such as forceps or vacuum extraction.

Prenatal diagnosis of macrosomia can be challenging but is crucial for appropriate management. Ultrasound examinations during the third trimester are commonly used to estimate fatal weight and assess potential risks associated with macrosomia. Close monitoring of maternal blood sugar levels, particularly in cases of diabetes, is also necessary.

The management of infants with macrosomia involves a multidisciplinary approach. Obstetricians, pediatricians, and neonatal nurses collaborate to ensure the well-being of both the mother and the infant. A carefully planned delivery, considering factors such as the estimated fatal weight and maternal health, is crucial. In cases of suspected macrosomia, healthcare providers may opt for an elective caesarean section to reduce the risk of complications.

Infants with macrosomia may be at increased risk of developing metabolic syndrome, obesity, and type 2 diabetes later in life. Consequently, regular follow-up visits with pediatricians are essential to monitor the child's growth, development, and metabolic health. Encouraging a healthy lifestyle, including balanced nutrition and regular physical activity, is vital in mitigating these long-term risks.

In conclusion, infants with macrosomia require specialized care throughout the pregnancy, delivery, and postpartum periods. By understanding the causes, potential complications, and appropriate management strategies, healthcare professionals can ensure optimal care for these new-borns and support their longterm health and well-being.

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