Fetal anaemia: two clinical cases with fetal blood transfusion

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Summary

Fetal anaemia in pregnancy, although less prevalent since the introduction of anti-D immunoglobulin, remains a clinical issue. In this article the authors describe two clinical cases of challenging fetal anaemia; the first following a parvovirus B19 infection during pregnancy and the second after Rh isoimmunization (due to Rh positive maternal transfusions in childhood), sooner and more severe in a second pregnancy. A brief review of the topic was also conducted.

Key words: Fetal anaemia; Parvovirus; Transfusion; Systolic peak middle cerebral artery; Fetal transfusion.

Introduction

When any blood group factor inherited via father is not possessed by the mother, antepartum or intrapartum hemorrhage may stimulate an immunological reaction [1-3]. This can also occur if the mother received unmatched blood transfusion [1-3]. The formation of maternal antibodies is called isoimmunization, which can lead to varying degrees of transplacental passage of these antibodies into the fetal circulation, causing a sufficient immune response to destroy fetal red blood cells [1-3]. More commonly, isoimmunization occurs in subsequent pregnancies, causing haemolytic disease of the fetus or newborn (NB), characterized by hemolysis, bilirubin release, and anaemia. The severity of the disease will depend on the degree of immune response (amount of antibodies produced and their affinity for the antigens), the gestational age at the diagnosis, and the fetal capacity to replace the destroyed erythrocytes, maintaining a sufficient hematocrit for their growth and development [1-3].

Fetal anaemia in pregnancy remains a diagnosis of suspicion. The etiology includes genetic diseases, metabolic deficits, infections, and thoracic abnormalities [2]. The most frequent causes of severe fetal anaemia are erythrocyte alloimmunization, parvovirus B19 infection, chronic fetomaternal haemorrhage, genetic erythrocytic diseases, and complications of twin fetal transfusion treatment [2]. Although relatively rare, the fetus is at risk of transient aplastic crisis, high-output heart failure, non-immune hydrops or even demise [1-3].

Case Report

Case 1

The patient was 38-years-old, 0 Rh negative, G2 P1, with first pregnancy at risk in 2010 due to positive Coombs and a steadily increase in irregular antibodies titer. Fetal anaemia and hydramnios developed at 33 weeks, conditioning a caesarean delivery two weeks later. The newborn needed an exchange transfusion in the second week of life. The investigation highlighted a history of maternal transfusion (0 Rh positive) in her childhood (at 6-7 years). At second pregnancy (2013), fetal blood group screening in maternal blood at 18 weeks revealed Rh positive fetus. After a weekly systolic peak of the middle cerebral artery (SP-MCA) surveillance beginning at 20 weeks, nine weeks later the ultrasound was suggestive of fetal anaemia (Figure 1). Three cordocentesis were then performed with blood cell transfusion (fetal hemoglobin before the first transfusion - 2.1 g/dL; after four transfusions - 9.5 g/dL). At 33+1 weeks a caesarean section was decided (NB 2,534 grams, 9/10 Apgar and 12 g/dL hemoglobin). A new NB blood transfusion was required at the third day of life due to a decrease in hemoglobin in the neonatal period (9 g/dL). Currently, at four-years-old, the child presents a normal psychomotor development.

Case 2

The patient was 32-years-old and was first surveilled with a pregnancy with normal development until week 24. In this period, fetal hydrops was observed (ascites and pericardial effusion), conditioning probable fetal anaemia - SP-MCA above 2 MoM (Figure 3). After investigation, the hypothesis of fetal infection by parvovirus B19 seemed probable, confirmed by amniocentesis. Then, managed in a tertiary hospital, a four-time cordocentesis with blood cell transfusion (fetal haemoglobin before the first transfusion - 2.1 g/dL; after four transfusions - 9.5 g/dL) was performed. Subsequently, in serial ultrasonography with SP-MCA, the fetus remained slightly above 1.5 MoM and with hydrops reversal until term (Figure 4). Eutocic delivery of NB with 2,592 grams, 9/10 Apgar occurred at 38 weeks. Blood NB samples highlighted 15 g/dL haemoglobin. Currently, at four-years-old the child presents a normal psychomotor development.

Discussion

More than 50 anti-erythrocyte antibodies have been associated with fetal haemolytic disease [2]. The most com-
mon antigens responsible for fetal erythroblastosis are anti-D, anti-K1 (Kell) and anti-c [1-4]. Despite the implementation of prophylactic anti-D isoimmunization, maternal Rh is still a cause of fetal hemolytic disease [1-3, 5].

The first step seems to be assessing paternal blood test to predict fetal Rh and determine the fetus risk of alloimmunization [2, 3, 5]. If this is negative, alloimmunization will be mainly due to past transfusions or injection sharing [2, 4, 5]. Next, if the paternal Rh is positive, free fetal DNA can be screened in maternal blood to determine the Rh type [1-3, 5]. Less commonly used option is PCR determination [2].

The first pregnancy with Rh positive alloimmunization is followed differently from subsequent ones, due to less severe fetal effects (worsening with each pregnancy) [1-5]. In the first one, the indirect Coombs titer is used not only for detection but also to characterize the degree of alloimmunization [2, 4, 5]. It is considered a screening test, with the critical titer (associated with fetal hydrops risk) being between 1/8 and 1/32 [2, 4, 5]. Until this threshold is reached, bi-monthly to monthly monitoring is acceptable; if this occurs, it becomes necessary to characterize the severity of fetal anaemia [2].

Administration of anti-D globulin after the primary immune response to the D antigen has occurred does not prevent a titer increase [2, 5]. Subsequent pregnancies are more serious due to the anamnestic response of maternal antibodies to the presence of fetal cells in the maternal cir-
The diagnosis of primary infection is essentially serological (IgM and IgG), although PCR techniques may be useful [1-6]. It is difficult to carry out its culture [6]. For fetal diagnosis, the method of choice, remains the isolation of small amounts of viral DNA through amniotic fluid PCR techniques [6]. If there is confirmation of infection, the clinical situation and risks should be explained, the necessary ultrasound surveillance implemented, and the possibility of intrauterine transfusion equated. This is mostly underlined in severe cases, since the fetus generally tolerates mild to moderate degrees of anaemia [1-6]. Doppler SP-MCA is effective in the non-invasive determination of the degree of fetal anaemia [1-6].

Fetal transfusion in clinical cases of fetal anaemia should be considered for hemoglobin levels below 7 g / dL (adjusted for gestational age) or two standard deviations below the mean, with erythrocytes submitted to the same quality control of the general population, preferably from Rh negative donor, irradiated, leuko-depleted, concentrated to a 75% to 85% hematocrit, CMV negative, and with less than five-day storage [2, 7]. Vascular access is mainly used, via a distal portion of the umbilical vein in the just-placental umbilical cord or intrahepatic umbilical vein, for a target of a 40-50% of final hematocrit [2, 7]. In some cases, intraperitoneal transfusion can be a choice [2, 7].

After the first transfusion, a decline in fetal hematocrit of approximately one percent per day is expected if the fetus has no hydrops (1.88 with hydrops) [2, 4, 7]. Therefore, a second transfusion will be necessary in about ten to 14 days after, depending on the hematocrit and the post-transfusion results; after the third, it is prudent to separate them three to four weeks by the suppression of fetal erythropoiesis [6, 7]. Subsequently, Doppler of the middle cerebral artery is not advised, by the presence of adult hemoglobin [2]. Ideally, they should be performed between 18 and 35 weeks (poor visualization of structures before this threshold and with risk of childbirth being lower after week 35 than transfusion) [6, 7].

Although data are still limited, fetal prognosis is good, with about 90% of children reaching a normal neurological development, even in the presence of hydrops [2, 3, 6, 7]. Delivery should be scheduled at a tertiary center [1-6].

Conclusion

After the onset of fetal hydrops, is important to rule out the diagnosis of anemia. The infectious aetiology should be investigated in fetal anaemia, namely parvovirus B19 infection and Rh isoimmunization, with even more severe and early recurrences. Surveillance is critical for fetal op-
timization. Anaemia can be severe and benefit from intrauterine transfusion, optimizing maternal-fetal prognosis.

References


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