Approach to a Pregnant Woman with Anti D + Anti C Reactivity Pattern: A Diagnostic Conundrum

Pathology Section

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ABSTRACT

The Rhesus G antigen is present on all RBCs that are C+ and also on most D+ RBCs. Due to this co-distribution of G with either C or D antigen, it mimics a reactivity pattern of anti C + anti D on Indirect Antiglobulin Test (IAT), though the role of Anti G in causing Hemolytic Disease of Newborn (HDN) is controversial. The differentiation of anti D, anti C, and anti G is essential particularly in pregnant females. We hereby report a rare case of anti G with anti D and anti G in a pregnant woman with emphasis on approach to identify anti D+C+G and its implications.

Keywords: Aloimmunisation, Arti D, Arti C, Anti G, Haemolytic, Irregular erythrocyte antibodies, Pregnancy

CASE REPORT

A 32-year-old, G2P1L1 female with 31 weeks of gestation presented in Obstetric Outpatient Department with a positive IAT reported from outside laboratory. The patient was Rh negative & thus suspected of Rh isoimmunisation. However, patient had past history of receiving single dose of anti D prophylaxis, 4000U during first pregnancy. Previous pregnancy led to full term normal delivery of healthy child with no history of neonatal america or jaundice. There was no significant medical history or obstetric history of any still britts, abortions or medical termination of pregnancy. There was no past history of blood transfusion. Patient had not received anti D in this pregnancy so far.

In view of positive IAT. Doppler ultrasound was done which revealed Middle Cerebral Artery Peak Systolic Welcotty (MCA-PSV) > 1.5 Multiples of Median (MOM); suggestive of severe fetal anemia. The patient was planned for Intrauterine Transfusion (IUT), however multiple units of O Rh negative leukoreduced packed RBCs put up for cross-match were found to be incompetible with maternal serum. Hence, the patient was referred to our Regional Blood Transfusion Centre (RBTC) for immunohaematology work-up.

First of all, ABO blood grouping and D typing of patient and her husband were performed. Patient's forward and reverse blood grouping done at room temperature (22°C) showed discrepancy. Following which, the patient's blood sample was collected in EDTA vial under strict warm conditions. The RBCs and serum were separated immediately by certiflugation at 2000 rpm for 5 minutes. The cells were washed multiple times with warm normal saline. Extended forward and reverse blood grouping was done at 22°C.

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[Flatble/Fig. 1]: Extended Blood grouping results of the patient.

37°C and 4°C by tube method. The blood group was confirmed as AB negative at 4°C. [Table/Fig-1] Patient was further confirmed to be negative for weak D by IAT using tube method and subsequently by column agglutination technology (Diamed get card method, Diamed, Switzerland). Autocontrols were negative at 3 temperatures ruling out autoantibody. Husband's blood group was A Rh positive. Also, polyspecific Direct Antiglobulin Test (DAT- anti-lgG and C3d) of the patient was negative. A commercially available three-cell antigen panel (ID-DiaCell I-II-III Asia, Diamed, Switzerland) was used for antibody screening by IAT. The patient's serum was reacted with reagent RBCs using LISS/Coombs ID-cards, at 37°C in AHG (antihuman globulin) phase. The cards were incubated for 15 minutes. and then centrifuged in ID-centrifuge for 10 minutes. The antibody screening panel was positive showing pan-agglutination. However, IAT by 3 cell panel was negative at 4°C in saline phase. An extended 11-cell panel ID-DiaPanel, DiaMed [Table/Fig-2] was used for antibody identification by IAT using ID-cards at 37°C. The reactions were suggestive of anti C + anti D antibodies [Table/Fig-3].

RivKeli/extended antigen profile of patient and her husband was done by using column agglutination technology (DiaClon gel card, Diamed Switzerland). Husband was strongly positive for C antigen whereas the patient was negative for it. [Table/Fig-4] Therefore, possibility of anti D + anti C with or without anti G could not be excluded. Meanwhile, a unit of irradiated, O Rh negative, C negative leucoreduced packed RBCs (heemstocrit of 80%) was found to be compatible with maternal serum and was successfully transfused in-

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[Babbs/Fig-2]: Phenotype of reagent red cells of extended 1.5 cell panel (DaMed 1 cell ID-DaPlanel) used for antibody dentification by WZ.

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