Successful Identification of Anti-f Alloantibody in a Non-transfused Male Patient Employing ID-Micro Typing System Gel Cards®

Yuki Nakamura1, Yoshihisa Ohsawa1, Yoshihisa Furuta1, Miho Tokida1, Kayoko Ichikawa1 and Akimichi Ohsaka1,2,*

1Department of Transfusion Service, Juntendo University Hospital, Tokyo, Japan
2Department of Transfusion Medicine and Stem Cell Regulation, Juntendo University School of Medicine, Tokyo, Japan

Corresponding author: Akimichi Ohsaka, M.D., Ph.D., Department of Transfusion Medicine and Stem Cell Regulation, Juntendo University School of Medicine, 2-1-1 Hongo, Bunkyo-ku, Tokyo 113-8421, Japan. Tel: +81 3 5902 1108; Fax: +81 3 3811 2724; E-mail: ohsaka@juntendo.ac.jp

Received date: Sep 13, 2015, Accepted date: Nov 04, 2015, Publication date: Nov 09, 2015

Abstract

Anti-f alloantibody is produced by exposure to the f (ce) antigen on red blood cells (RBCs), which is a compound antigen of the Rhesus (Rh) blood group, expressed when both c and e antigens are present in the same haplotype (in cis). Although anti-f alloantibody was first discovered in 1953 and it has rarely been detected, the results of its detection were not reported as images in previous studies. We report a case of anti-f alloantibody in a Japanese male patient without a history of blood transfusion, whose anti-f alloantibody was successfully identified using a gel microcolumn assay (GMA). Furthermore, we reviewed the transfusion records between January 2002 and December 2010 and found that a total of 799 (1.1%) among 73,636 blood specimens were positive for irregular RBC antibody, of which anti-f alloantibody was only observed in one patient. The present case report focuses on the usefulness of GMA to preserve the results of detecting alloantibodies as images.

Keywords: f (ce) antigen; Anti-f alloantibody; cis; Rh blood group; Gel microcolumn assay

Introduction

The Rhesus (Rh) system is one of the most immunogenic and polymorphic human blood group systems. The RHCE gene encodes both C/c and E/e antigens on a single protein. The f (ce) antigen is a compound antigen of the Rh blood group and expressed on red blood cells (RBCs) when both c and e alleles of the RHCE gene are in the cis position on the Rh chromosome ([in cis]), e.g., DCCeEe (R1R2) and ccee (rr), but not in trans, e.g., DCCeEe (R1R2). Alloantibodies directed at cis product antigens have been encountered infrequently [1]. Anti-f alloantibody was first discovered in a multiply transfused 30-year-old Caucasian male who suffered from hemophilia in 1952 [2]. In general, alloantibodies in the Rh blood group can cause destruction of transfused RBCs and hemolytic disease of the newborn (HDN) [3]. Anti-f alloantibody has been implicated as the cause of HDN [4,5] and delayed hemolytic transfusion reaction [6]. In addition, an in vitro RBC survival test of a patient with anti-f antibody demonstrated the accelerated destruction of f antigen-positive RBCs [7]. Cases with anti-f alloantibody have rarely been reported in the gel column era [8], and the results of its detection were not reported as images in previous studies.

A gel microcolumn assay (GMA) is a modified serological technique and its principle is based on the use of gel to trap agglutinates. It offers advantages in terms of result stability, versatility, and the potential for automation [9]. It is noteworthy that the reactions are stable and allow for a secondary review of results for up to 24 hours after testing has been completed [9]. In this paper, we report a rare case of anti-f alloantibody in a Japanese male patient without a history of blood transfusion, whose anti-f alloantibody was successfully identified using GMA, and its detection was reported as color images.

Case Report

A 40-year-old Japanese male suffering from mitral regurgitation was hospitalized for mitral valveplasty, and he had denied previous blood transfusions. For pre-operative examination, blood samples were presented to the transfusion service in the hospital. The standard hemagglutination method was used employing ID-Micro Typing System Gel Cards® (Bio-Rad Laboratories, Tokyo, Japan). The blood type of the patient was reported to be B, D+C-E+c+c- (R1R2), K-, Le(a-b+), Fy(a+b-), Jk(a-b+), M+N+S-s+, and Di(a-). However, the patient’s plasma showed pan-reactivity on reverse blood grouping by GMA, resulting in an ABO discrepancy (data not shown). An indirect antiglobulin test (IAT) with the two-stage papain method was performed using a three-cell panel (ID-DiaScreen) and an 11-cell panel (ID-DiaPanel, each Bio-Rad Laboratories), showing reactivity with an anti-f antibody (Figure 1). To further investigate the patient’s plasma reactivity, adsorption/elution studies were carried out on RBCs with ccee (rr), DCCeEe (R1R2), and DCCeEe (R1R2) phenotypes. Adsorption of antibody in the patient’s plasma was only observed in the case of the ccee (rr) phenotype (Figure 2A), and the eluates of the adsorption study showed strong (3+) agglutinates with the ccee (rr) phenotype and weaker agglutinates with the ccee (‘r’ or ccEe ‘rr’) phenotype (Figure 2B), suggesting the presence of anti-f alloantibody. Next, a comparative titration study of the patient’s plasma was performed by a low ionic strength saline (LISS)-IAT, revealing that agglutinates before 2-mercaptoethanol (2-ME) treatment showed 8 titer (Figure 3A), whereas those after 2-ME treatment showed 2 titer (Figure 3B), suggesting that the patient’s anti-f alloantibody was a naturally acquired anti-f alloantibody.
Figure 1: Identification of an alloantibody of the patient's plasma. An indirect antiglobulin test (IAT) with the two-stage papain method was performed using a three-cell panel (a) and 3 of 11-cell panel (b), showing reactivity with an anti-f antibody.

Figure 2: Adsorption/elution studies with the two-stage papain-IAT. (A) Adsorption studies were performed on RBCs with ccee (rr), DCCee (R₁R₁), and Dccee (R₂R₂) phenotypes, and then the reactivity with RBCs with DCCee (R₁R₁), Dccee (R₂R₂), and ccee (rr) phenotypes was investigated. (B) The eluates of the adsorption studies were applied to GMA, and the reactivity with RBCs with DCCee (R₁R₁), Dccee (R₂R₂), Dccee (R₁R₂), Ccee (r'r), ccEe (r''r), and ccee (rr) phenotypes is shown.
Figure 3: Comparative titration studies with a low ionic strength saline (LISS)-IAT. The patient's plasma was applied to GMA before (A) and after (B) treatment with 2-ME, showing an anti-f antibody in the patient as a mixture of IgM and IgG.

Since anti-f alloantibody has been implicated as the cause of a delayed hemolytic transfusion reaction [6], patients with anti-f alloantibody require compatible blood units, being safely transfused with units of blood lacking both c and e antigens. The patient received mitral valveplasty with 800 mL of pre-operative autologous blood donation (PABD) and 100 mL of salvage blood in the operation. Although 1,000 mL of ce antigen-negative allogeneic blood units was reserved, the operation was completed without the need to use allogeneic blood.

Discussion

To elucidate the incidence of anti-f alloantibody, we reviewed the database of transfused patients in the hospital between January 2002 and December 2010, and found that a total of 73,636 blood specimens were screened for irregular RBC antibodies, of which 799 (1.1%) were positive for irregular antibodies in 774 patients. In this study, an antibody being identified from the same patient during the study period was counted as one. When one (or more) different antibody was additionally detected during the study period, they were counted as two (or more).

The number and rate (%) of the identified irregular RBC antibodies were as follows: anti-E 253 (31.7), anti-Lea 217 (27.2), anti-M 49 (6.1), anti-E+e 42 (5.3), anti-Cw 28 (3.5), anti-C+e 24 (3.0), anti-D 20 (2.5), anti-Lea+Leb 16 (2.0), anti-Fyb 17 (2.1), anti-P1 15 (1.9), anti-Dia 14 (1.8), anti-C 14 (1.8), anti-Leb 13 (1.6), anti-c 11 (1.4), anti-Jka 11 (1.4), anti-S 9 (1.1), anti-e 8 (1.0), anti-Jra 4 (0.5), anti-K 3 (0.4), anti-Bga 3 (0.4), anti-Jkb 2 (0.3), anti-Fy^a 2 (0.3), anti-E+Lea 2 (0.3), anti-E+Leb 2 (0.3), anti-E+Fyb 2 (0.3), anti-D+C 2 (0.3), anti-C+E 2 (0.3), 'anti-f 1(0.1)', anti-N 1 (0.1), anti-LW 1 (0.1), anti-Lu 1 (0.1), anti-Xg^a 1 (0.1), anti-S+P1 1 (0.1), anti-Lea+P1 1 (0.1), anti-Jka+Di 1 (0.1), anti-E+P1 1 (0.1), anti-E+Leb 1 (0.1), and anti-E+S+Fyb 1 (0.1). Among the irregular RBC antibodies identified during the study period, anti-f alloantibody was detected in only one patient who was presented in this paper. The rate of irregular RBC antibodies in our institution was consistent with that of a recent nationwide surveillance in Japan [10], where anti-f alloantibody was not reported. The prevalence and specificity of irregular RBC antibodies may vary among different geographic areas and races, being reported from 0.49 to 2.4% in hospitalized patients [11-16]. These studies also reported no case with anti-f alloantibody.

GMA has been used for ABO and Rh typing, the direct antiglobulin test (DAT), detecting alloantibodies, RBC phenotyping, and other applications. Although the role of GMA for DAT is controversial, GMA has been more sensitive than the conventional tube test (CTT) for detecting potentially significant antibodies coating RBCs in vivo [17]. Compared to CTT, an operator-dependent assay, GMA is less susceptible to handling errors, generates clear, more objective results, and may perform comparably to the CTT in titrating alloantibodies to Rh and Kell antigens [18]. In this study, we used ID-Micro Typing System Gel Cards® for detecting anti-f alloantibody in the patient, demonstrating detection as color images. Limitations of the present study are that it was a single case study and a single institutional investigation for detecting irregular RBC antibodies in hospitalized patients. Further studies are needed to establish the usefulness of GMA for preserving the results of detecting alloantibodies as images. To our knowledge, this study is the first to report the detection of anti-f antibody as color images.

Conclusion

We reported a rare case of anti-f alloantibody in a Japanese male patient without a history of blood transfusion, whose anti-f alloantibody was successfully identified by GMA. GMA may be useful for preserving the results of detecting alloantibodies as images.
Acknowledgements

We thank Dr. Makoto Uchikawa (Tokyo Red Cross Blood Center, Tokyo, Japan) for performing a confirmation study and Masahiro Ogasawara (Bio-Rad Laboratories, Tokyo, Japan) for helpful discussion.

Conflict of Interests

The authors declare no conflict of interests.

References