

TRALI – multiple suspects for a crime

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Introduction

- Transfusion-related acute lung injury (TRALI) is a life threatening complication of transfusion characterized by the development of acute respiratory distress associated with non-cardiogenic pulmonary edema, occurring within 6 h after receiving a blood transfusion.
- TRALI has been associated with transfusion of virtually all blood components. Donors involved in a TRALI event should be screened for the presence of HLA Class I and II and HNA antibodies and the transfusion recipient assessed for the presence of the corresponding antigens. Donors involved in a TRALI event should be deferred from donating blood.
- We report the case of a 56 year old woman admitted for acute myeloid leukemia and petechiae, who initiated symptoms of dyspnea, tachycardia and vomiting during the administration of a pool of platelets. She died a few hours later from respiratory failure. TRALI was diagnosed.

Material and Methods

- Donors anti-HLA antibodies specificity was performed with LABScreen™ Single Antigen (One Lambda), anti HNA antibodies were analyzed with LABScreen™ Multi (One Lambda) and recipient HLA typing was performed by PCR-SSO LIFECODES HLA Typing Kits (HLA-A,B,C, DRB1, DQB1 and DPB1 – Immucor).

Results

- HLA genotyping of the patient : A*01, 02; B*07,08; C*07; DRB1*04,17; DQB1*02,08; DPB1*06,13.

Patient HLA Genotyping

A*01,02
 B*07,08
 C*07
 DRB1*04,17
 DQB1*02,08
 DPB1*06:13

Anti HLA antibodies - Median Fluorescent Intensity

| | Donor 1 | Donor 2 | Donor 3 | Donor 4 |
|------|---------|---------|---------|---------|
| A1 | 2107 | | | |
| A36 | 1432 | | | |
| B8 | 3579 | | | 2897 |
| DR4 | | | 1067 | |
| DR7 | | | 1441 | |
| DR10 | 1023 | | | |
| DQ7 | | 1153 | | |
| DQ8 | | | 1823 | |
| DP1 | | | 3394 | 1066 |
| DP5 | | | 1307 | |
| DP6 | | | 1118 | |
| DP10 | | | 1043 | |
| DP14 | | | 1509 | |

- Anti-HLA antibodies were identified in all donors, with maximum Median Fluorescent Intensity ranging from 3579 to 1023.
- Patient specific antibodies were identified in 3 out of the 4 donors: donor 1 anti- HLA- A1 and HLA-B8, donor 3 anti- HLA- DR4, DQ-8 and DP-6, and donor 4 anti- HLA B8.
- None of the donors had anti HNA antibodies.

Conclusion

- It is noteworthy that anti-HLA antibodies against some of the patient's antigens were identified in 3 donors, thereby straining the identification of the donor or donors responsible for the TRALI event.
- The finding of multiple female donors with the theoretical ability to elicit TRALI might suggest reappraisal of the criteria for the selection of blood donors in order to prevent TRALI.