

# CASE REPORT: LIFE -SAVING INCOMPATIBLE TRANSFUSION

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## Background

Since clinical significant antibodies can cause hemolytic transfusion reactions, it is essential to consider them when transfusion is required. Our report is based on an 81-year old Caucasian male patient with anti-e, -Ce and -Fy<sup>a</sup> who required mitral valve surgery.

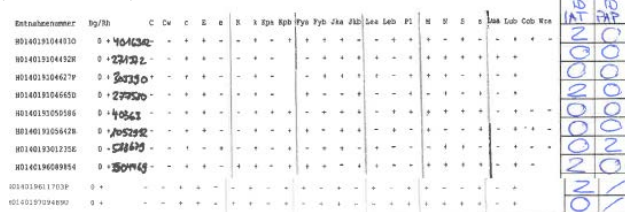
In consultation with the surgeon, 4 fully compatible red blood cell units were provided for the intervention.

Due to intraoperative complications, the patient required heart-lung machine support. As another 12 units were requested the challenge was to select the least incompatible units.

## Methods

Standard serological methods were used for antibody specification and direct antiglobulin test (DAT) (Grifols, CH, BioRad, CH and in-house).

Compatibility testing was performed using indirect antiglobulin test (IAT) at 37°C (gel-card, BioRad, CH) and molecular typing of the patient's so called rare blood group antigens by PCR-SSP (inno-train, D).



Entnahmesummer	Sp/Bl	C	Cw	E	e	B	k	Kpa	Kpb	Kya	Kyb	Jsa	Jsb	Fl	M	N	E	e	Jma	Lob	Cob	Wra			
H01401930449310	D +																								
H0140193044932K	D +																								
H0140193044937P	D +																								
H0140193044940D	D +																								
H014019305050586	D +																								
H014019305054126	D +																								
H01401930121216	D +																								
H01401960808954	D +																								
H0140196111039	D +																								
H0140197094890	D +																								

Figure 1. antibody specification showing anti-Fy<sup>a</sup> reactive in IAT and anti-e only reactive with papain-treated test cells

## Results

The patient's phenotype was B, ccD.EE (R<sub>2</sub>R<sub>2</sub>), K- and Fy(a-b+). Pretransfusion testing revealed anti-Ce and anti-Fy<sup>a</sup> reactive in IAT and anti-e only reactive with papain-treated test cells. In absence of compatible products we decided not to consider the anti-e and selected O ccDEe (R<sub>2</sub>r), K-, Fy<sup>a</sup>- RBC units for compatibility testing.

All of the incompatible units (n=12) showed negative cross matches. Finally, 7 of these incompatible products were transfused during the night.

A few days after incompatible transfusion the patient's serum showed a positive XM with a supposedly compatible red blood cell unit (R<sub>2</sub>R<sub>2</sub>, K-, Fy<sup>a</sup>-).

Subsequent analyses revealed an anti-Yt<sup>b</sup> (Cartwright<sup>b</sup>) and patients genotyping confirmed Yt<sup>b</sup> negativity. Consistently, the cross match positive red blood cell product as well as at least one of the cross match negative units transfused in emergency were determined Yt<sup>b</sup> positive.

The patient's DAT, autocontrol and eluate remained negative during the entire follow up and no clinical or laboratory signs of major intra- or extravascular hemolysis was observed.

## Transfusionsmerkblatt

Vor Bluttransfusionen dem Arzt vorweisen

**1938**

Blutgruppe **B** RhD **positiv** **ccD.EE, CW-, K-**  
Weitere Blutgruppenantigene: **Fy(a-b+), Yt(a+b-)**

Irreguläre Blutgruppenantikörper:

**Anti-e** (Rhesusantikörper)  
**Anti-Ce** (Compound Rhesusantikörper)  
**Anti-Fy<sup>a</sup>** (Anti-Duffy<sup>a</sup>)  
**Anti-Yt<sup>b</sup>** (Anti-Cartwright<sup>b</sup>)

Um hämolytische Reaktionen bei Transfusionen möglichst zu vermeiden müssen **e negative, C negative, Fy<sup>a</sup> negative und Yt<sup>b</sup> negative** Erythrozytenkonzentrate mit **einwandfrei** negativen Verträglichkeitsprüfungen verabreicht werden.  
Bei Blutbedarf ist eine frühzeitige Bestellung und Abklärung erforderlich.

Figure 2. transfusion document including recommendations for the selection of compatible blood to avoid transfusion reactions

## Conclusion

The fact that the anti-e was only enzyme-reactive was crucial for our decision to neglect this ab; thereby considering the other two antibodies (anti-Ce and anti-Fy<sup>a</sup>) as well as being able to issue units with negative cross matches.

After the incompatible transfusion, follow-up showed no signs of hemolysis. However, a newly formed or boosted low-prevalence antibody, anti-Yt<sup>b</sup>, was identified shortly thereafter, when a fresh patient sample showed positive cross match with a previously cross match negative red blood cell product.