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# Case Report: Autoimmune anemia and thrombocytopenia following hematopoietic stem cell transplant

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## Purpose

- **Autoimmune cytopenias (AIC)** post hematopoietic stem cell transplant (HSCT) are rare but exceptionally challenging complications.
- Here, we report **anti E, warm autoimmune hemolytic anemia (WAIHA), cold autoantibody, polyagglutination** and **immune thrombocytopenia** in a **5-year-old boy** with a history of **chronic granulomatous disease**, who underwent an **ABO incompatible, mismatched (9/10) unrelated HSCT**.

## Methods

- **Case Report**
- **4 months following HSCT** patient was admitted for **fever, fatigue, and decreased appetite**.
- **Patient's** original blood type was **O+** and **donor's** blood type was **B+**.
- **Hematopoietic chimera evaluation** demonstrated **100% donor-derived cells**.
- Initial laboratory work-up revealed:
  - ❑ White blood count **10.07x10<sup>3</sup>/uL**
  - ❑ Hemoglobin **4.2 gm/dL**
  - ❑ Platelets **137x10<sup>3</sup>/uL**
  - ❑ Absolute retic count **0.123x10<sup>6</sup>/mcl**
  - ❑ Total bilirubin **1.8 mg/dl** (mainly indirect)
  - ❑ Lactate dehydrogenase **1,037 unit/L**
- **Several months later**, the patient developed **thrombocytopenia with a platelet count of 45x10<sup>3</sup>/uL**.

## Results

Test	Results	Comments
ABO typing	Invalid	Spontaneous agglutination of patient's red cells
Antibody screen and work-up	Positive	Anti E
Direct antiglobulin test (DAT)	Invalid	Spontaneous agglutination of patient's red cells
Eluate	Positive at 22°C and 37°C	Cold antibody present and warm antibody confirmed
Cold antibody titer	Negative	Titer < 2
Thermal amplitude	Negative	No reactivity at 30°C and 37°C
Donath Landsteiner test	Negative	
DTT treated eluate 22°C PEG IAT	Decreased reactivity Weak reactivity	Supports the presence of IgM and IgG antibodies in the eluate
Polyagglutination testing (Patient red cells mixed with donor plasma)	Strongly positive	AB plasma: 9/10 reacted B plasma: 4/4 reacted
Lectin studies Arachis hypogea Glycine soja Salvia sclarea Salvia horminum	Strongly positive Negative Negative Negative	Spontaneous polyagglutination but not due to the classic T activation. Tx, Tk or Th variants of T activation were neither ruled out nor confirmed.
Platelet antibody studies SPRCA test ELISA test	Negative Positive	Anti HLA, not present Anti GPIIb/IIIa, present

## Discussion

- This is a report of a **5-year-old male post-HSCT** who developed **overt hemolytic anemia** with laboratory testing that demonstrated evidence of **WAIHA, anti-E antibody**, clinically insignificant **cold autoantibody**, and **polyagglutination**.
- **Classic T activation** was ruled out.
- Several months later, **hemolysis subsided** but **immune mediated thrombocytopenia ensued**.
- This case **demonstrates the complexity of AIC** that rarely develops in patients who underwent HSCT.

## Summary

- Autoimmune cytopenias are a rare but exceptionally challenging complications of HSCT.
- This report demonstrates a case of anti-E, WAIHA with cold antibodies and polyagglutination who later developed immune-mediated thrombocytopenia.