

Centre hospitalier universitaire vaudois

# Infectious Complications after Treatment of Antibody-Mediated Kidney Allograft Rejection: A National Cohort Study



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# Background

Multimodal therapeutic strategies used to treat acute antibody-mediated rejection (AMR) could enhance the risk of infection.

# Objectives

• To describe the occurrence of infectious complications

# Conclusion

- Infectious complications were common after acute AMR treatment.
- Infection-associated mortality was low (3.1%).
- Plasmapheresis was associated with an increased risk of infection.
- IVIg may reduce the incidence of bacterial infection.
- To analyze the impact of the different therapeutic strategies on the incidence of infection after AMR treatment

# Methods

## **Study population**

- All kidney transplant (KT) recipients from the Swiss Transplant Cohort Study (STCS)<sup>1</sup>
- Who received a treatment for an acute AMR episode occurring in the first year post-transplantation (Tx) (2008-2014).

### Data

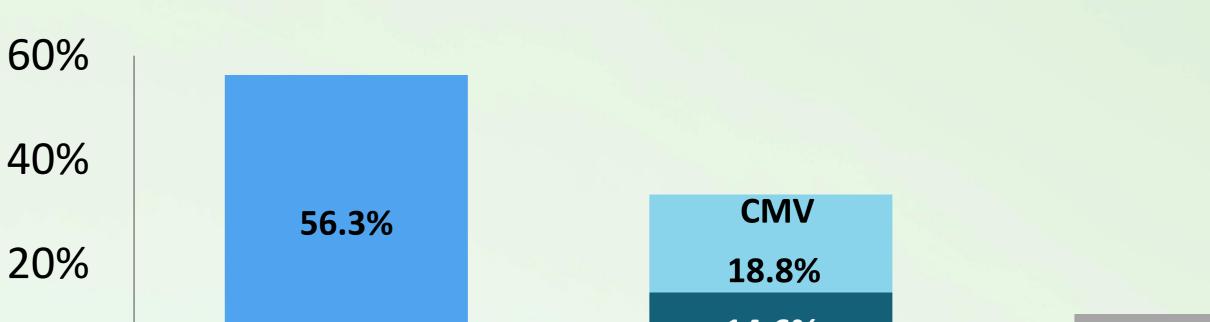
- Acute AMR treatment used
- Infectious complications occurring in the 6 months following acute AMR treatment

### 1 year survival outcomes:

- Graft survival (death-censored) : 90.9%
- Patient survival : 93.8% (2/4 patients died of severe infection)

## Infectious complications within 6 months:

- 63.6% (42/66) of transplants
- 2.3 episodes/patients



#### **Episodes of infection (n=96)**

Patient and graft survival

#### Analysis

- Risk factors of infection after AMR treatment: uni- and multivariate Cox regression models
- Time-to-event curves: Kaplan-Meier method (log-rank test for inter-group differences)

### Results

66/1669 (3.9%) KT recipients were treated for an acute AMR episode in the first year post-Tx.

Value
65 (66 Tx)
56.1
$46.1 \pm 18.5$
27.3
53.0
34.8
33.3
16.7
15.2
98.5
6.6
6.1
3.0
56.1
98.5

0%10.4%BacterialViralFungalFigure 1: Type of infectious complications within the first 6 months after<br/>acute AMR treatment. Main bacterial infections: urinary tract inf. 37.0%,<br/>respiratory tract inf. 20.4%, blood stream inf. 18.5%.

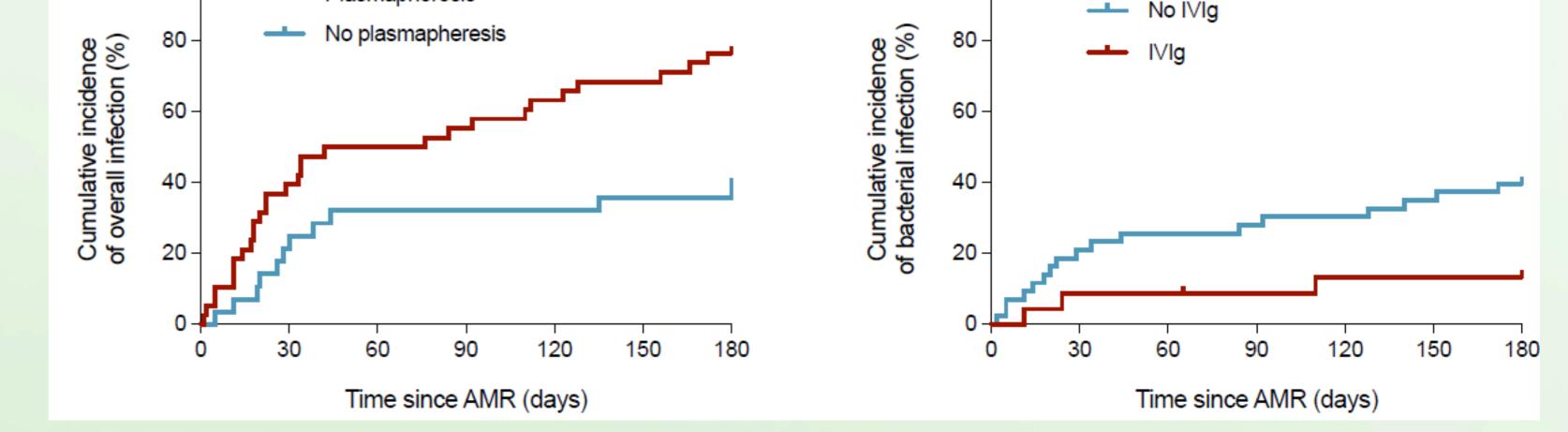
**Table 2:** Multivariate analysis of risk factors predicting the occurrence of infection

Risk factor	Hazard Ratio (HR)
For overall infection:	
Plasmapheresis	HR: 2.9 (95%CI: 1.5-5.7), P = 0.002
For bacterial infection:	
Induction with Rituximab	HR: 6.6 (95%CI: 2.1-20.7), P = 0.001
IVIg	HR: 0.3 (95%CI: 0.1-1.0), P = 0.053
For opportunistic infection:	
Plasmapheresis	HR: 5.3 (95%CI:1.2-27.7), P = 0.033

100 <sub>7</sub>

a)

۲ 100



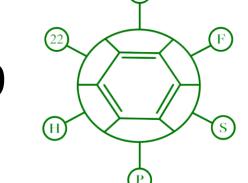
b)

**Figure 2: a.** Overall infection according to the use of plasmapheresis (log-rank test *P*-value = 0.002), **b.** Bacterial infection according to the use of IVIg (log rank test *P*-value = 0.035). IVIg: intravenous immunoglobulins

#### Reference

1. Koller M.T., et al. Eur J Epidemiol, 2013. 28(4): p. 347.

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