

# Concomitant piperacillin-tazobactam and vancomycin use increases the risk of acute kidney injury

**Medical history** A male in his 40s was hospitalized for infection after artificial urinary sphincter implantation.

Day 1 *Escherichia coli* detected in urine culture on admission. Piperacillin-tazobactam (PIPC/TAZ) **4.5 grams three times per day started** on same day.

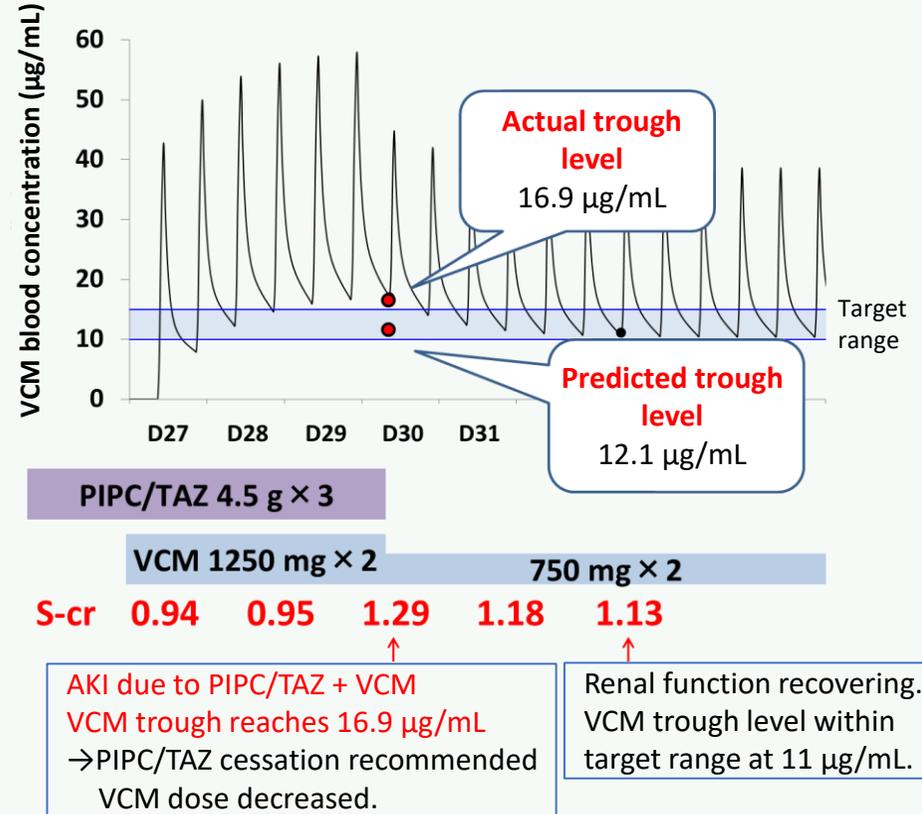
Day 27 The patient presented with fever of 38°C. **Methicillin-resistant *Staphylococcus aureus* (MRSA)** and ***Corynebacterium striatum*** detected in a urine culture. Treatment with vancomycin (VCM) **1250 mg two times per day started** (S-cr 0.94 mg/dL, Ccr 144 mL/min).

Day 30 **Acute kidney injury (AKI) detected** (S-cr 1.29 mg/dL, Ccr 105 mL/min). **Predicted VCM trough blood level was 12.1 µg/mL but actual level was 16.9 µg/mL** (Target VCM trough range: 10-15 µg/mL). Because concomitant PIPC/TAZ and VCM use frequently causes AKI, and because the PIPC/TAZ treatment period was long (30 days), **discontinuation of PIPC/TAZ therapy was recommended**.

**PIPC/TAZ therapy was stopped the same day and the VCM dose was decreased to 750 mg twice per day.**

Day 33 Renal function recovering. VCM trough level decreased to 11 µg/mL, within target range.

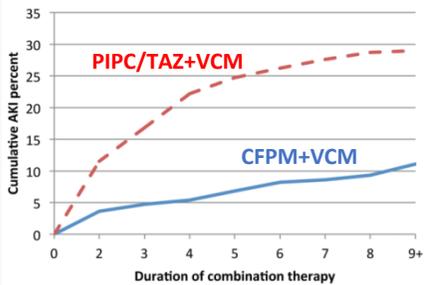
Day 48 Administration of VCM therapy completed without additional AKI.



## Beware of increased risk of side effects due to unnecessary antimicrobial use

- Concomitant use of PIPC/TAZ and VCM is known to increase the risk for AKI. When possible, it is important to change one to a different antimicrobial or to discontinue one if unnecessary.
- In this case, the addition of VCM to prolonged PIPC/TAZ therapy resulted in AKI.

## The risk of AKI due to concomitant PIPC/TAZ and VCM use



AKI risk is up to **THREE TIMES HIGHER** for PIPC/TAZ plus VCM compared to CFPM plus VCM.

Navalkele B et al. Clin Infect Dis. 64:116-23,2017.

PIPC/TAZ: Piperacillin-tazobactam, VCM: vancomycin, CFPM: Cefepime