[P25-4] P25-4: Anti-infective drugs (4): Vancomycin

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[P25-4-5] Optimizing vancomycin dosage with a therapeutic drug

monitoring program in thrice-weekly hemodialysis patients

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Background

Poster

The treatment of patients with vancomycin and end-stage renal (ESR) function with regular hemodialysis schedule (thrice-weekly-HD) is complicated not only because therapeutic drug monitoring (TDM) is needed in order to achieve a target trough concentration (Cp), but also because of its irregular dosage regimen. Our objective was to evaluate a vancomycin TDM program in order to optimize the dosage in ESR patients treated with thrice-weekly hemodialysis to achieve stable vancomycin Cp the whole week.

Methods

Retrospective observational study. Patients over 18 years with ESR and thrice-weekly-HD, treated more than one week with vancomycin and at least one TDM report were included.

TDM protocol: a) an initial dose of 500 mg or 1000 mg iv upon clinician's criteria; b) TDM and dosage adjustment with Cp before the second dose of vancomycin.

TDM was based on Bayesian estimation of the pharmacokinetic parameters (PKSABBOTT v1.10) by Clinical Pharmacokinetics Unit.

Results

38 patients (36.8%/63.2% w/m) were included. The average age was 61 years (Cl95%: 57-65) and weight was 69 kg (Cl95%: 63-74). The indications were: tissue and skin infection (52.6%), bacteremia (26.3%), respiratory infection (10.5%) and unknown fever (10.5%). Median (p25-75) basal creatinine and CRP was: 6.2 mg/dL (5.9-7.3) and 6.2 (1.7-15.8), respectively. The median (p25-75) total and per kilogram initial dose was 1000 mg (1000-1000) and 15.8 mg/kg (11.0-18.2). Median basal pre-hemodialysis concentration after the initial dose was 12.1 mg/L (p25-75: 10.2-13.9). A new dosage regimen was initiated after the first TDM in 100% of patients. After dosage adjustment, the median (p25-75) total and per kilogram dose was: 500 mg (500-750) and 9.0 mg/kg (7.1-13-4) on Monday-Wednesday; and 1000 mg (750-1250) and 15.8 mg/kg (11.0-18.2) on Friday, with mean Cp pre-hemodialysis was 15.0 ng/mL (Cl95%: 14.1-16.0).

Conclusions

TDM of vancomycin is required in hemodialysis patients. In our population, to achieve target Cp after the initial dose it is necessary to administer a loading dose higher than 15 mg/kg (probably 20 mg/kg). To maintain target Cp pre-hemodialysis stable, it is necessary to increase the dose of vancomycin after last hemodialysis session of the week (ie. on Friday) in every patient by 50% (p25-75: 45.8-66.7).

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