

Vancomycin

Synonyms



Accredited to
ISO 15189:2022

Clinical Indication

Pre-dose monitoring is sufficient and post dose level is NOT required in most clinical cases.

Blood should be collected:

- a) Pre-dose: immediately prior to the drug being given
- b) Post-dose: 2 hours after completion of infusion

Ensure trough sample is collected within 2 hours prior to the next dose.

Do not withhold next dose unless patient has renal impairment or previous level was high.

Part of Profile / See Also

Request Form

Combined Pathology manual request form or ICE request. Ensure details of the date/time last dose are provided.

Availability / Frequency of Analysis

On request.

Turnaround Time

Performed on the same day within 2 hours of receipt in the laboratory.

Patient Preparation

Dosage should be 12 hourly.

Sample Requirements

Specimen Type

Serum and plasma

Volume

2 ml

Acceptable Containers



Yellow top (SST) tube



Green top (lithium-heparin) tube



paediatric orange top (lithium-heparin)



paediatric green top (lithium-heparin)

Plain serum samples may also be used.

Reference Range & Units

10.0-15.0mg/L for mild-moderate infections.
15-20mg/L for severe, deep seated infections.
(Except for ITU patients on continuous infusion – please refer to ITU protocol)
The suggested ranges are for guidance only, these may vary with disease process and the infecting organism, please refer to the appropriate trust protocol

Interferences

Interpretation & Clinical

Decision Value (if applicable)

References

Beckman kit insert.

Test code

VANC

Lab Handling

Analysed from primary tube and stored at 4°C.
Serum and plasma stable for 7 days at 2-8°C.