

Inspecting Point of Care

<p>Processes/Areas for Observation</p>	<ul style="list-style-type: none"> • Specimen collection including patient ID/specimen labeling • Storage condition of reagents and QC • Reagent labeling • Personnel access to procedure manual • Result reporting process • Infection control practices • System to detect and correct unusual, clerical, analytical errors(move to All Common)
<p>Key Documents to Review</p>	<ul style="list-style-type: none"> • Organizational chart for line of authority • Procedure manual including specimen collection • Patient Reports <ul style="list-style-type: none"> ○ ID of testing personnel ○ Reference range, if appropriate ○ Drugs of abuse reporting elements • Personnel Records <ul style="list-style-type: none"> ○ Training ○ Competency ○ Color discrimination testing ○ List of POCT personnel with tests authorized to perform ○ Non-waived testing – academic degree or transcript • Quality Control – Non-Waived <ul style="list-style-type: none"> ○ Daily QC/Monthly QC with review and corrective action ○ Reagent lot verification ○ Validation of internal QC process ○ Comparison of non-waived methods/instruments ○ Calibration/calibration verification/AMR procedure and records • Quality Control – Waived <ul style="list-style-type: none"> ○ Daily QC/Monthly QC with review and corrective action ○ Calibration and calibration verification per manufacturer's instructions • Arterial Blood Gases <ul style="list-style-type: none"> ○ Arterial puncture training ○ Collateral circulation test records ○ Calibration/QC records ○ AMR verification records • Physician Performed Testing <ul style="list-style-type: none"> ○ Physician credentialing records ○ Training/competency records and patient reports if not credentialed