

Patient Safety Indicators

Pre-analytic:

- **Phlebotomy associated adverse events:**
 - hematoma
 - multiple phlebotomy attempts >3
 - fainting
 - lapse in infection prevention/hand hygiene
 - skin reaction to tape/bandage/latex
 - sharps left in patient bed

- **Patient Identification:**
 - wrong patient drawn
 - failure to use 2 patient identifiers

- **Specimen Identification:**
 - unlabeled
 - mislabeled

- **Order Entry:**
 - incorrect or wrong patient demographics entered
 - test(s) ordered on the wrong patient
 - incorrect clinician entered
 - incorrect test or procedure ordered
 - wrong blood product ordered

- **Specimen integrity:**
 - contamination
 - hemolysis
 - fibrin clots
 - insufficient volume
 - inappropriate sample
 - lost or destroyed sample
 - specimen drawn above an intravenous line (diluted or contaminated)
 - improper temperature maintained

- **Effective use of Clinical Laboratory Services:**
 - inappropriate test(s) requested
 - inappropriate test frequency (too few or too many)
 - test requested at inappropriate time (e.g. therapeutic drug monitoring)
 - failure to order the appropriate test (failure to follow clinical practice guideline or clinical pathway)

Analytic:

- **Verify accuracy of abnormal test results:**
 - failure to verify abnormal or critical POCT result with clinical laboratory service
 - failure to recognize specimen integrity issues that affect test results
 - failure to recognize specimen process errors that affect test results

Post-analytic:

- **Wrong blood product dispensed/administered**
- **Communication of Test Results:**
 - test results not communicated to clinician
 - critical values not reported within defined timeframe to clinician
 - failure of timeliness in communication of results with clinician
- **Effective Utilization of Test Results:**
 - incorrect interpretation of test results
 - failure to order follow-up test(s)
 - continuing to re-order the same laboratory test
- **Outcomes of laboratory testing:**
 - failure to follow a best practice protocol (consider coagulation monitoring, therapeutic drug monitoring)
 - failure of Provider to notify patient of abnormal test results and required next steps