I. OBJECTIVES

Blood cultures are intended to provide information for the diagnosis and management of patients with suspected bacteremia or fungemia. The objective of this policy is to standardize optimum procedures for the ordering, procurement, and transport of blood cultures in a manner designed to provide accurate information.

II. INDICATIONS FOR USE

Used in the ordering, procurement and transportation of blood cultures.

III. DEFINITIONS

<table>
<thead>
<tr>
<th>Term</th>
<th>Definition</th>
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<tbody>
<tr>
<td>Blood Culture</td>
<td>A specimen of blood that is submitted for bacterial or fungal culture. NOTE: This is irrespective of the number of bottles or tubes into which the specimen is divided or distributed</td>
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<tr>
<td>Blood Culture Series</td>
<td>A group (usually two sets) of temporally related blood cultures that are collected to determine whether a patient has bacteremia or fungemia, and to help determine if a recovered organism is significant, or a likely contaminant.</td>
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<tr>
<td>Blood Culture Set</td>
<td>The combination of blood culture bottles or tubes into which a single blood specimen is inoculated</td>
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<tr>
<td>Contaminant</td>
<td>A microorganism isolated from a blood culture that was introduced into the culture during specimen collection or processing and that was not pathogenic for the patient from whom the blood was collected (i.e., the isolates were not present in the patient’s blood when the blood was sampled for culture).</td>
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<tr>
<td>CR-BSI</td>
<td>Catheter-related bloodstream infection</td>
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<tr>
<td>Aerobic Blood Culture Bottle</td>
<td>Used primarily for isolation of organisms that require oxygen as a hydrogen acceptor in their metabolic pathways. Optimum volume 8-10 ml of blood.</td>
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</table>
IV. RESPONSIBILITY

A. Authorized Prescribers
1. Order blood cultures according to this policy.
2. Indicate the time frame in which the cultures are to be drawn (or obtained per unit protocols/pathways).
3. Indicate any special media requirements.
4. Order no more than 4 sets of blood cultures per 24 hour period.
5. Draw the blood culture(s), if required. (See Appendix A)

B. Registered Nurses (RN)/ Licensed Practical Nurses (LPN)
1. With demonstrated/documented competency, may draw blood cultures according to Appendix A.
2. RN/LPN, do not perform femoral or lower extremity venous or arterial sticks.
3. Exception: Lower extremities can be used to obtain specimens for pediatric patients under 10 years old.

C. Microbiology Laboratory
1. Perform the analytical portion of blood culture processing.
2. Communicate positive results according to the critical action values policy.
3. Monitor blood culture contamination rates.
4. Provide education and training for proper blood culture procurement and transportation procedures

V. PROCEDURE

A. Standard Procedures
1. General Considerations:
   a. Ordering a Blood Culture:
      i. The prescriber must specify the site(s) of cultures to be obtained, i.e., peripheral, central line, other.
      ii. The prescriber must specify the type(s) of cultures needed: i.e., bacterial, fungal, isolator, mycobacterial, other.
      iii. The prescriber must specify number required and timing preferred, especially if antibiotics are to be held prior to obtaining the initial diagnostic cultures.
   b. Preparing the phlebotomy site:
i. Chlorhexidine Frepp preparation must be used unless, in a given patient, the product is contraindicated. This product has been proven to provide the most rapid and complete sterility of the skin site, and the lowest incidence of skin contamination and resulting false positive cultures.

ii. If the patient is allergic to Chlorhexidine, tincture of iodine may be substituted. Povidone iodine products must be avoided, except in the rare patient who cannot tolerate an alcohol-based product.

iii. The water-based, povidone products fail to dry in a timely manner, and release their iodine content too slowly to predictably de-fat skin, and kill underlying flora. These deficits greatly increase contamination rates.

2. Timing and Site Source:
   a. Except when not medically feasible, a minimum of two sets of blood cultures, from two separate sites, shall be drawn peripherally.
      i. If a different peripheral site is possible, the second set may be drawn immediately.
      ii. If using the same site (same vein), wait at least 10 minutes prior to obtaining the second set. A new sterile field must be created and site must be re-prepped.
      iii. If unsuccessful with second peripheral attempt, obtain one peripheral set and one set from a central venous access device, if present.
      iv. Up to two additional sets may be drawn within 24 hours.
   b. The practice of drawing blood from a catheter shall be avoided when a peripheral (non-catheterized) site is available.
      i. Blood cultures may not be drawn through an existing peripheral IV catheter.
      ii. A central catheter site is much more likely to give false positive results and may result in unnecessary interventions (antibiotics, line changes, etc.). Whenever possible, unless ruling out catheter sepsis, or unless the sample is drawn when the line is initially inserted, blood cultures shall not be drawn from an indwelling catheter site.
   c. Blood cultures for diagnostic purposes must be obtained prior to implementing or changing antibiotics, unless the authorized prescriber ordered otherwise. Cultures must be drawn promptly to prevent delay in initiation of antibiotic therapy. Blood cultures are most likely to be positive if the blood sample is collected as soon as possible after the development of fever.
   d. Surveillance or "test of cure" blood cultures are rarely indicated, and shall not be performed as a routine.

B. Specialized Blood Culture Procedures
   1. Suspected catheter sepsis:
      a. Draw two culture sets simultaneously.
         i. One set is to be obtained aseptically from the catheter hub or through the implanted venous access port (VAP).
         ii. The second set must be drawn from a peripheral site.
         iii. If a medical reason precludes drawing from a peripheral site, obtain a second culture from a different catheter lumen.
         iv. If the catheter is removed and culture is desired, a 5 cm section including the tip is to be cut aseptically, and sent to Microbiology in a dry sterile container. Catheter hardware cultures must always have an accompanying peripheral or line blood culture specimen sent within 24 hours to help validate the significance of any recoveries made from the hardware.
   2. Suspected acute endocarditis and endovascular infection or source
      a. Draw 2-3 culture sets from two separate sites, within 30 minutes of each other and before beginning antimicrobial therapy.
      b. At least one set shall be from a peripheral site.
      c. Begin therapy after cultures are obtained.
   3. Suspected subacute endocarditis:
Blood Cultures: Ordering, Procurement and Transport

4. Follow-up blood cultures
   a. Most patients with bacteremia or fungemia do not require follow-up blood cultures to document clearance. There are two exceptions for which at least two sets of blood cultures are to be drawn at 48 - 96 hours:
      i. Patients with infective endocarditis.
      ii. Patients with Staphylococcus aureus bacteremia not related to endocarditis.

5. Mycobacterial blood cultures (AFB):
   a. Use a mycobacterial blood culture bottle (BD BACTEC Myco F Lytic bottle). Because the media is unstable, it must be obtained from the Mycobacteriology (AFB) Lab, Meyer B1-124, 5-6470 on dayshift, or the main Microbiology Lab, 5-6510, during evening hours. The media is kept in a brown paper bag to protect it from light. The pneumatic tube can be used to obtain the culture bottles.
   b. Prepare the skin as for routine aerobic blood cultures.
   c. Inoculate 5 ml blood into a BACTEC Myco F Lytic bottle.
   d. Deliver the blood bottle in the brown bag promptly to the Microbiology Lab. The pneumatic tube can be used as long as sample is sent directly to the Microbiology Lab, and is sent within the brown paper bag. Shipment must be prompt to permit the timely inoculation of the additives that may be required.
   e. An authorized prescriber must have specifically placed an order for a Mycobacterial blood culture.

6. Fungal Cultures:
   a. Follow the routine procedure described above for bacterial cultures. If the order is specific for Histoplasma or Cryptococcus, an isolater tube culture must be inoculated. Obtain an isolator tube from the Microbiology Lab (call 5-6510). Because Isolator media is light sensitive, it also is stored in brown paper bags.
      i. Inoculate the isolator tube with 5cc of blood (absolute minimum is 3cc).
      ii. Send the culture in its brown paper bag directly to the Microbiology Lab. The pneumatic tube can be used as long as sample is sent directly to the Microbiology Lab.
   b. Submit an extra 3 ml purple top for the detection of viruses other than HSV by molecular testing if multiple viruses are in the differential.
      i. Molecular testing is available for detection of cytomegalovirus (CMV), Epstein Barr virus (EBV), enteroviruses, parvovirus B19 and adenoviruses.
      ii. Obtain one 3 ml purple top tube or plasma partitioning tube for detection of CMV, enteroviruses, parvovirus B19 and adenoviruses by molecular methods.
      iii. Obtain two 3 ml purple top tubes for EBV detection/quantification.

8. When rare organisms such as Brucella, Campylobacter or Bartonella are suspected:
   a. An ID physician shall be consulted.
   b. The Microbiology Lab shall be consulted to advise which type of specimen would be most likely to support the suspected organism.
9. Pediatric Blood Cultures:
   a. Pediatrics shall follow the same policies and procedures as described in this policy. See Appendix B for guidelines regarding blood culture volumes.

VI. REPORTABLE CONDITIONS
   A. Report to the authorized prescriber:
      1. When unable to obtain the required blood specimens for any reason.
      2. Any complications that occur during the procurement or transport procedures.

VII. DOCUMENTATION
   A. The date and collection time of a blood culture is automatically recorded in the laboratory if a blood culture specimen was scanned at the time of procurement. This is important for result interpretation, especially if the culture is negative. It permits correlation with the time of initiation of antibiotics.
   B. If a manual process is used, document the blood culture order in the medical record, noting the date and procurement time.
   C. Document any complications which may have occurred while obtaining a blood culture such as sub-optimal specimen amount or inability to obtain specimen from a peripheral site.

VIII. EDUCATION AND COMMUNICATION
   A. This policy will be communicated through Johns Hopkins Hospital publications.
   B. Placement of policy on-line at www.Hopkinsmedicine.org/heic
   C. This policy will be placed in the Interdisciplinary Clinical Practice Manual on the JHH Intranet site http://www.insidehopkinsmedicine.org/hpo . Paper distributions will be made to the Functional Unit Nursing offices in the event of web access difficulty.

IX. SUPPORTIVE INFORMATION
   See Also:
   The Johns Hopkins Hospital, Interdisciplinary Clinical Practice Manual
   • Adult Vascular Access Device Policy, IFC035
   • Mislabeled or Unidentified Patient Specimens, PAT027
   Microbiology Department. Guidelines for Drawing Blood Cultures.
   Children's Center Pediatric Policies, Procedures and Protocols
   • Pediatric VAD Central Maintenance Protocol

References:


Sponsor:

- Medical Care Evaluation Committee

Developers:

- Laboratory Advisory Committee
- VAD Committee
- Microbiology Lab
- Nursing Standards of Care
- HEIC

Review Cycle - Three (3) years  Medical Board - Approval: 11/30/10; Effective: 12/1/10

Vice President for Nursing and Patient Services  Vice President for Medical Affairs

Date:  Date: