Saddleback Memorial Medical Center
24451 Health Center Drive
Laguna Hills, CA 92653
(949) 452-3554
LABORATORY SERVICES

ACCREDITATIONS

College of American Pathologists (CAP)  AU-ID# 1187862/AP# 2337601
Health Care Financing Administration (CLIA)  # 05D0578029
State of California Department of Health Services License  # CIF 539
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INTRODUCTION TO

SADDLEBACK MEMORIAL MEDICAL CENTER

LABORATORY SERVICES

SMMC MISSION:

To improve the health and well being of individuals, families and our communities through innovation and the pursuit of excellence.

SMMC VISION:

To become Southern California’s preferred, operationally excellent, fiscally sound provider of comprehensive, high-quality health services.

DESCRIPTION

At Saddleback Memorial Medical Center, we are dedicated to providing quality services to our patients with the most advanced laboratory testing and uncompromising customer service. We offer:

- Laboratory services 24 hours a day, 365 days a year
- State-of-the-art testing methodologies
- A wide range of in-house testing
- Quick and accurate STAT testing
- Reliable specimen handling and processing

DEDICATED & PROFESSIONAL STAFF

Our highly trained and dedicated staff is committed to providing you with the highest level of service:

- Courteous and knowledgeable staff
- Experienced phlebotomists (neonatal and pediatric proficient)
- Portocath nurse available at main laboratory location
- High ratio of licensed clinical laboratory scientists (CLS) to technicians
- Board-certified pathologists available for consultation 24 hours a day

For questions regarding test results, please call (949) 452-3554.
COMPREHENSIVE LAB TESTING

Following is sampling of our comprehensive laboratory testing services:

- Blood Bank, Definitive
- Antibody Testing
- Bone Marrow Examinations
- Chemistry, Automated & Esoteric
- Coagulation Assays
- Consultative Pathology
- Endocrinology
- Fetal Lung Maturity Studies
- Hematopathology
- HIV Antibody Testing
- Microbiology
- Obstetrical/Prenatal Studies
- Therapeutic Drug Monitoring
- Urine Chemistry

PATHOLOGIST-SUPPORTED SPECIALTY TESTING AVAILABLE

Our pathologists are available to assist physicians with their decisions in a variety of medical specialties, including:

- Surgical and Cytopathology
- Fine Needle Aspiration
- Non-gynecological Cytology
- Clinical Pathology
- Breast Pathology
- Immunopathology
- Blood Bank and Transfusion Medicine
- Hematopathology
- Coagulation and Cellular Immunology
- Pediatric and Placental Pathology
- Bone Marrow Examination
- Cytochemical Stain
DIRECTORY

Laboratory: Saddleback Memorial Medical Center

Locations: 24451 Health Center Drive
Laguna Hills, CA 92653

Phone/FAX:

- Clinical Laboratory: (949) 452-3445
- Clinical Lab (Fax): (949) 452-3563
- Anatomic Pathology: (949) 452-3562
- Anatomic Pathology (Fax): (949) 452-3066

Facility CEO: Marcia Manker (949) 452-3622
Facility COO: Cheryl Jacob (949) 452-3853
Vice President, Ancillary Services: Hugo Folli (949) 452-7109
Laboratory Medical Director: Vivian Mendoza, M.D. (949) 452-3562
Transfusion Medicine Director: Thomas Hirose, M.D. (949) 587-3056

Laboratory Leadership:

- Executive Director: Cheryl Dilbeck, CLS (949) 452-3613
- Technical Manager: Linda Stricklin, CLS (949) 452-3047
- Phleb/Point of Care Supervisor: Joanne Johansen, CLS (949) 452-7430
- Operations Manager/Specimen Processing/Blood Donor Center: Susan Robertson, CLS (949) 452-3055

Technical Specialists:

- Blood Bank Coordinator: Cheryl Louie, CLS (949) 452-3559
- Hematology: Sabrina Uy, CLS (949) 452-3557
- Microbiology/Chemistry Coordinator: Dolores Wertemberger, CLS (949) 452-3560
- Quality: Barbara Purdy, CLS (949) 498-4581

Laboratory Information Systems:

- .......................................................... (949) 452-3668

Blood Donor Center:

- Blood Donor Center Coordinator: Elisita Healey, RN (949) 452-3161
- Blood Donor Center Program Support Representative: Carol Carranza (949) 452-3159
TESTING GUIDELINES

ACCREDITATION/LICENSURE

SMMC Laboratory maintains a current CLIA number with the U.S. Department of Health and Human Services Health Care Financing Administration, is accredited by the College of American Pathologists (CAP), and holds all other necessary state licenses. For additional information or copies of our certificates, please call (949) 452-3445.

ALPHABETICAL TEST LIST

Information about specific tests can be found in the alphabetical test list section of this Compendium. This section contains test numbers, mnemonics, methodologies, performed/reported schedules, specimen requirements, CPT codes, and notes.

MEDICARE COVERAGE OF LABORATORY TESTING

When ordering laboratory tests that are billed to Medicare/Medicaid or other federally-funded programs, the following requirements may apply:

1. Only tests that are medically necessary for the diagnosis or treatment of the patient should be ordered. Medicare does not pay for screening tests, except for certain specifically approved procedures, and may not pay for non-FDA approved tests or those tests considered experimental.

2. If there is reason to believe that Medicare will not pay for a test, the patient must be informed. The patient must sign an Advance Beneficiary Notice (ABN) to indicate that he or she is responsible for the cost of the test if Medicare denies payment.

3. The ordering physician must provide an ICD-9 diagnosis code, not a narrative description, if required by the fiscal intermediary or carrier.

4. Organ or disease oriented panels should be billed only when all components of the panel are medically necessary.

5. Client customized panels should be billed to Medicare only when every component of the customized panel is medically necessary.

6. Medicare National Limitation Amounts for CPT codes are available through HCFA or its intermediaries. Medicaid reimbursement will be equal to or less than the amount of Medicare reimbursement.

Questions concerning utilization of CPT codes should be addressed with your local carrier, as local medical review policies may vary from one third-party payer to another. Call Admitting at 949-52-3546 or Outpatient Express Registration at 949-452-7420 for more information.

MINIMUM ACCEPTABLE VOLUMES

This Compendium lists minimum acceptable volumes. If there is insufficient volume for testing, attempts will be made to locate any additional sample at SMMC Laboratory which was collected at the same time. In this case, there may be delays.

This Compendium also lists pediatric minimum acceptable volumes. This volume is defined as the amount sufficient to perform one test, with no repeat or confirmatory testing.
PANELS
SMMC offers groups of standard HCFA Panels approved for Medicare patients; customized profiles require medical staff approval or physician signature at time of request. Individual components of Custom Panels may also be ordered individually, unless otherwise indicated.

REFLEX TESTING
Approved parameters for reflex testing and mandatory confirmation testing, which will occur when initial test results are positive or outside normal parameters and indicate that a second related test is medically indicated are defined in policy PL-403, APPROVAL OF LABORATORY PANELS FOR USE ON SMMC HOSPITAL PATIENTS, Attachment F-3.

REFERRAL TESTING
One of SMMC Laboratory’s service goals is to support clients by providing comprehensive service for all reference laboratory testing. To accomplish this goal, SMMC has enhanced its in-house test menu and test combinations by selecting primary vendors to perform additional tests not performed at SMMC. Primary vendors are selected based upon aspects of service, quality, reliability, turnaround time, and price.

SMMC’s established referral test vendors fax or transmit test results expediently to SMMC so that clients receive results by the fastest, most error-free method possible. If a client requests that a test performed by SMMC, or by one of our primary vendors, be performed by another laboratory, SMMC will honor that request if it is clearly indicated on the test request form submitted at the time the test is ordered. A handling fee will be added to reference testing.

SMMC reserves the right to change vendors, test parameters, and costs for referral testing at any time. For questions, please contact staff at (949) 452-3554.

SPECIMEN CONTAINERS AND TUBES
SMMC Laboratory requests that clients use the following guidelines to ensure safe handling procedures, non-compromised specimens, and fast and accurate test results.

ACCEPTABLE CONTAINERS

• Sterile specimen containers
• Swabs (sterile)
• Transport media

UNACCEPTABLE CONTAINERS/CONDITIONS

• Syringes with needles attached
• Specimens received in expired transport containers, media, or tubes
• Transfer tubes secured with parafilm
• Specimens that are mislabeled or unlabeled or don’t have at least two (2) identifiers
• Specimens that are leaking from the container
SPECIMEN REJECTION/TEST CANCELLATION

All specimens must be collected, labeled, minimum last and first name with date of birth, transported, and processed according to procedure. Selecting the container type, volume, and special handling requirements needed for analysis before the specimen is collected is essential. If the criteria for these processes are not met, the specimen may be rejected or the test may be canceled. The following represent some reasons for specimen rejection or test cancellation:

• Inappropriate specimen type or difficulty in obtaining the specimen (e.g. hemolysis, clotted specimens)
• Insufficient volume for analysis
• Improperly labeled specimen
• Inappropriate specimen container
• Improper specimen transport
• Specimen that has leaked in transit
• Specimen that has been sent in expired transport media
• Incomplete or incorrect test request form; for example, testing not marked
• Test request without a specimen
• Specimen without a test request

SPECIMEN STABILITY

Specimen stability for many tests is described in the alphabetical test list, for client convenience. Stability information may be useful in determining whether a previously processed patient sample is still appropriate for testing at SMMC Laboratory. The temperatures and times listed under stability do not indicate preferred transport conditions or temperatures. If a particular specimen fits the stability criteria for an assay, please refer to the Transport section on how to send the specimen to SMMC Laboratory.

STAT TESTING

Many procedures are offered by SMMC Laboratory on a STAT (NOW) basis for emergency purposes 24 hours per day with an expected turn-around-time (TAT) of approximately one hour from the time the laboratory receives the specimen. ASAP orders are defined with an expected TAT of two to four hours from the time the laboratory receives the specimen.

Routine testing is generally completed within four hours and in all cases within the shift received.

All tests ordered on infants as STAT will be performed STAT.

The list of those tests which will be performed on a STAT basis for all other age populations are defined in policy PL-400, STAT PROCEDURES.
SPECIMEN COLLECTION

GENERAL INSTRUCTIONS FOR SPECIMEN COLLECTION
Collection instructions for individual tests appear in the alphabetical test list of this Compendium under Specimen Required. Please refer to this information to ensure that the appropriate sample has been collected for the test ordered.

HEMOSTASIS/THROMBOSIS SPECIMENS

I. ANTICOAGULATION LABORATORY MONITORING

To: SMMC Medical Staff

From: Thomas G. Hirose, M.D.
Director, Transfusion Medicine

Date: February 25, 2013

Subject: Anticoagulation Laboratory Monitoring

Anticoagulation therapy is expanding with the addition of several new agents and newer medication on the near horizon. The following is a summary of laboratory monitoring for these agents. When a laboratory assay is not used, weight adjusted administration provides the therapeutic dose. While factor Xa levels have been used in the past for the monitoring of the therapeutic effects of several anticoagulants, their utility has been limited and in many cases, does not correlate to the degree of therapeutic anticoagulation (e.g. LOVENOX). Currently, one of the only indications for Xa level monitoring is when a “lupus anticoagulant” (aka anti-phospholipid antibody) is present which is interfering with the aPTT and/or Prot ime monitoring.

Anticoagulation should be used with caution for the following patients: obese, newborns, patients with renal failure (unless the medication is metabolized by the liver).

Unfractionated heparin should be administered by the SMMC dosing protocol (bolus and units/kg/hour). A copy of the protocol may be found in all clinical treatment areas.

Recommendation for laboratory monitoring of patient anticoagulation:

<table>
<thead>
<tr>
<th>Medication</th>
<th>Laboratory Monitoring</th>
<th>Metabolized</th>
</tr>
</thead>
<tbody>
<tr>
<td>Coumadin</td>
<td>Prot im e</td>
<td>Liver</td>
</tr>
<tr>
<td>Unfractionated Heparin</td>
<td>aPTT</td>
<td>Liver</td>
</tr>
<tr>
<td>Low Molecular Weight Heparin</td>
<td>Weight &amp; Creatinine adjusted</td>
<td>Kidney</td>
</tr>
<tr>
<td>(Enoxaparin)</td>
<td>No laboratory monitoring</td>
<td></td>
</tr>
<tr>
<td>(Rivaroxaban)</td>
<td>Weight &amp; Creatinine adjusted</td>
<td>Kidney</td>
</tr>
<tr>
<td></td>
<td>No laboratory monitoring</td>
<td></td>
</tr>
<tr>
<td>Direct Thrombin Inhibitors</td>
<td>aPTT</td>
<td>Liver</td>
</tr>
<tr>
<td>(Argatroban)</td>
<td>Weight &amp; Creatinine adjusted</td>
<td>Kidney</td>
</tr>
<tr>
<td>(Dabigatran)</td>
<td>No laboratory monitoring</td>
<td></td>
</tr>
<tr>
<td>Pentasaccharide Anticoagulants</td>
<td>Weight &amp; Creatinine adjusted</td>
<td>Kidney</td>
</tr>
<tr>
<td>(Fondaparidux)</td>
<td>No laboratory monitoring</td>
<td></td>
</tr>
</tbody>
</table>

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Notes:

1. **Warfarin** is a vitamin K inhibitor. Vitamin K is required for the terminal carboxylation of Factors II, VII, IX, X, and Protein C & Protein S. Coumadin is given as an oral medication. Initiation of therapy should not include “bolus” dosing. The average dose is approx. 4 mg p.o. daily. A therapeutic dose usually requires 4-6 days. The target INR for Coumadin in most cases is 2-3. Mechanical valves have a target INR of 2.5 - 3.5. Coumadin is contraindicated for the therapy of H.I.T. Type II until the platelet is above 150,000.

2. **Unfractionated Heparin** is monitored by aPTT. The target degree is based on Laboratory Heparin Therapeutic Range.

3. **Enoxaparin** prophylactic dose is 40 mg. daily administered subcutaneously. Full course anticoagulation is 1mg/kg twice daily or 1.5 mg/kg daily. Anti-Factor Xa Level Assays may be considered when an overdose of LMWH needs to be ruled out. Enoxaparin is only partially reversed by Protamine Sulfate (30%) and has no reversing agents. FFP may be considered for hemorrhagic complications but may have a paradoxical effect due to AT-III content of the plasma. This may be associated with increased bleeding.

4. **Ultra LMWH-Pentasaccharide Anti-coagulant** has no reversing agents. The prophylactic dose of Fondaparinux is 2.5 mg. subcutaneous daily. Full course anticoagulation is 7.5 mg. subcutaneous daily for patients < 100 kg. Studies have shown that there is no correlation with Anti-Factor Xa Level and should not be used for monitoring of the therapeutic effect. FFP may be considered for hemorrhagic complications. Fondaparinux may be considered for the treatment of H.I.T. Type II. Full course anticoagulation would be used.

5. **Argatroban and Lepirudin** is a direct thrombin anticoagulant used for the treatment of H.I.T. Type II. They are initially dosed at 0.5 ug/kg/minute as an intravenous infusion. The dose is adjusted until the aPTT is 1.5 to 2x the mean of the normal aPTT. There is no reversing agent. FFP may be considered for hemorrhagic complications. They should not be used as an alternative for UFH while on cardio-pulmonary bypass machine as it can be associated with life threatening hemorrhagic complication in as many as 50% of cases.

6. **Dabigatran** is prophylactic treatment for non-valvular atrial fibrillation. The dose is 150 mg./12 hours.

7. **Rivaroxaban** is prophylactic treatment for non-valvular atrial fibrillation, DVT, PE and for knee & hip replacement.
   - Knee replacement: 10 mg. PO qDay for 12 days; may take with or without food.
   - Hip Replacement: 10 mg. PO qDay for 35 days; may take with or without food.
   - DVT and PE: 15 mg PO BID for 21 days with food, THEN 20 mg. PO qDay for 6 months.
   - Nonvalvular Atrial Fibrillation: 20 mg./day PO with the evening meal.
II. HEMOSTASIS/THROMBOSIS SPECIMEN COLLECTION

To produce valid results for hemostasis/thrombosis tests and factor assays, specimen integrity is crucial and must be maintained. All specimens sent for testing must be collected and shipped in the following manner:

1. Obtain venous blood by clean venipuncture. Avoid slow flowing draws and/or traumatic venipunctures as either of these may result in an activated or clotted sample. Do not use needles smaller than 23 gauge.

2. Always draw a pilot tube (blue top tube – marked with X) before drawing coagulation specimens in light blue.
   
   Note: Reference ranges have been established using 3.2% buffered sodium citrate.

3. Fill light blue top tubes as far as vacuum will allow, and mix by gentle inversion. Exact ratio of 9 parts blood to 1 part anticoagulant must be maintained. Underfilled specimens will be rejected. Send samples within 4 hours after draw.

4. If transporting plasma or freezing coagulation specimens, the specimen must be processed as follows:
   
   a. Centrifuge the capped tube at 2500xg for 10 minutes or at 5600 rpm for 5 minutes. Inspect for hemolysis or clot, which is not acceptable for coagulation testing and requires that specimen needs to be redrawn.
   
   b. Immediately remove the top two-thirds of the platelet-poor plasma from the specimen using a plastic transfer pipet (use of glass pipets can result in activation and/or coagulation of the plasma).
   
   c. Place the plasma in a properly labeled polypropylene tube. Seal the tube. Make sure to label that the specimen is PLASMA. (Glass vials will be rejected).
   
   d. Freeze the specimen immediately at -70°Celsius. Specimens should not be frozen in an ordinary household or self defrosting freezer for storage.
   
   e. A separate tube must be submitted for each assay requested. (Those being sent for special studies such as Factor V, XII ETC.)
   
   f. Send specimen on dry ice – specimens must remain frozen during transport.

5. Some assays are performed on a priority basis if a medical emergency exists. Contact Client Services to make arrangements.

6. All requests for coagulation assays must include a brief patient history and other pertinent clinical information (e.g., medications, blood products, etc.).

   Note: Samples containing heparin should not be used for coagulation studies. If possible, stop heparin therapy before the draw to avoid contamination. Heparin interferes with most clotting assays.
GENERAL INSTRUCTIONS

INFECTIOUS DISEASE SPECIMENS

I. SPECIMENS - GENERAL REQUIREMENTS

A. SPECIMEN CONTAINERS

1. SHIPPING CONTAINERS, STERILE SPECIMEN CONTAINERS, TRANSPORT MEDIA, AND SWABS ARE AVAILABLE FROM SMMC. REFER TO INFECTIOUS DISEASES: COLLECTION AND TRANSPORT MEDIA AT THE END OF THIS SECTION FOR GUIDELINES. SPECIMENS ARE ACCEPTABLE FOR PROCESSING ONLY WHEN COLLECTED AND SUBMITTED IN THE APPROPRIATE CONTAINER. SPECIMEN CONTAINERS MUST BE SECURELY TIGHTENED TO ELIMINATE ANY LEAKAGE. USE OF COLLECTION AND TRANSPORT CONTAINERS THAT ARE PAST THE EXPIRATION DATE IS UNACCEPTABLE.

B. RESULT REPORTING

1. PRELIMINARY RESULTS ARE ISSUED AS SOON AS ACCURATE DATA IS AVAILABLE. FINAL RESULTS ARE GENERATED AT THE COMPLETION OF THE CULTURE. IF REQUESTED, PRELIMINARY RESULTS WILL BE CALLED TO THE PHYSICIAN OR REQUESTING LAB ON ANY POSITIVE, SIGNIFICANT STAIN OR CULTURE (E.G. BLOOD, CSF, STERILE BODY FLUID). THE REQUEST FOR PHONE NOTIFICATION SHOULD BE WRITTEN ON THE TEST REQUEST FORM. INFORMATION MUST INCLUDE THE NAME OF THE PERSON OR LAB TO CONTACT, AND THE TELEPHONE NUMBER. RESULTS OR INTERPRETATIONS OF RESULTS MAY BE OBTAINED BY CALLING THE LABORATORY DURING THE HOURS OF OPERATION OR BY CALLING (949) 452-3554.

C. REPORTABLE DISEASES

1. SMMC LABORATORY FOLLOWS APPLICABLE LOCAL AND STATE REQUIREMENTS FOR REPORTING.

II. SPECIMEN - SPECIFIC COLLECTION GUIDELINES

The proper collection of a specimen for culture is the most important step in the recovery of pathogenic organisms responsible for infectious disease. A poorly collected specimen may lead to failure in isolating the causative organism(s) and result in the recovery and subsequent treatment of contaminating organisms.

Basic Concepts for Collection

1. Collect the specimen from the actual site of infection, avoiding contamination from adjacent tissues or secretions.
2. Collect the specimen at optimal times (for example, early morning sputum for AFB culture).
3. Collect a sufficient quantity of material.
4. Use appropriate collection devices: sterile, leak-proof specimen containers. Use appropriate transport media (Amies Gel Swab for bacterial culture, Cary-Blair for stool culture, UTM for viral and chlamydia cultures).
5. Whenever possible, collect specimens prior to administration of antibiotics or antivirals.
6. Properly label the specimen and complete the test request form. The source of specimen, date and time of collection, and initials of collector is required.
7. Minimize transport time. Maintain an appropriate environment between collection of specimens and delivery to the laboratory.
8. If appropriate, decontaminate the skin surface. Use 70-95% alcohol (ALC) and 1-2% tincture of iodine (TIO) to prepare the site. Allow a contact time of two minutes to maximize the antiseptic effect.
SPECIAL GUIDELINES FOR SPECIMEN COLLECTION AND TRANSPORT

I. Principle

All diagnostic information from the Microbiology Laboratory is contingent on the quality of the specimen received. Consequences of a poorly collected and/or poorly transported specimen include failure to isolate the causative microorganism and recovery of contaminants or normal microbiota. This can lead to improper treatment of the patient. Often, direct specimen smears are utilized to determine the quality of the specimen, to provide rapid information for diagnosis and therapy, and to allow the physician to determine if additional, better quality specimens should be collected.

II. Specimen

A. General Considerations

This procedure addresses instructions that must be communicated to physicians, nurses, and phlebotomy teams.

1. Safety considerations
   a) Follow universal precaution guidelines. Treating all specimens as potentially hazardous eliminates the need for warning labels.
      i. laboratory coat or gown when collecting or handling specimens. If splashing may occur, protective eyewear, face masks and aprons may be necessary.
   b) Do not contaminate the external surface of the collection container and/or its accompanying paperwork.
   c) Minimize direct handling of specimens in transit from the patient to the laboratory. See Section B – Transport of Diagnostic Specimens

2. General guidelines for proper specimen collection
   a) Collect specimen before administering antimicrobial agents when possible.
   b) Collect specimen with as little contamination from indigenous microbiota as possible to ensure that the sample will be representative of the infected site.
   c) Utilize appropriate collection devices. Use sterile equipment and aseptic technique to collect specimens to prevent introduction of microorganisms during invasive procedures
   d) Clearly label the specimen container with the patient's full name and date of birth, and with the date and time of collection with initials or ID# of collector and RN initials.
   e) Collect an adequate amount of specimen. Inadequate amounts of specimen may yield false-negative results.
   f) Develop an understanding of the Microbiology Laboratory’s source identification schemes. Know when to include "rule-out" requests. For example, the laboratory routinely screens for Shigella, Salmonella, Plesiomona, Aeromonas, Vibrio and Campylobacter species in stool cultures but not for Yersinia.
   g) Identify the specimen source and/or specific site fully and correctly on the specimen so that proper culture media will be selected during processing in the laboratory. For example, use Abscess, right leg.
   h) If a specimen is to be collected through intact skin, cleanse the skin first. For example, use 70% alcohol followed by iodine solution (1 - 2% tincture of iodine or 10% solution of povidone-iodine.) Prevent burn by tincture of iodine by removing excess after specimen has been collected.
   i) Before collecting the specimen, consider the risk/benefit ratio of the collection procedure to the patient.
      Collect specimens in sturdy, sterile, screw-cap, leakproof containers with lids that do not create an aerosol when opened.
3. General guidelines for proper specimen transport
   a) Transport all specimens to the laboratory promptly.
      i. To ensure the survival and isolation of fastidious organisms and to prevent overgrowth by more hardy bacteria
      ii. To shorten the duration of specimen contact with some local anesthetics used in collection procedures that may have antibacterial activity
      iii. To provide a more accurate diagnosis of the infectious disease process.
   b) Alternative to prompt delivery
      i. Refrigerate most specimens at 2 - 8° Celsius. The following are exceptions:
         a) Specimens that may harbor temperature-sensitive organisms such as Neisseria species should be left at room temperature.
         b) For anaerobic specimens, use Amies Gel Swab. (See Table 1.)
         c) Stool specimens
            i. For bacterial culture, mix stool with a transport medium (Carey-Blair medium).
            ii. For parasitology examination, mix stool with preservative (PVA and formalin)
         d) Hold CSF specimens at room temperature unless they are to be cultured for viruses.
         e) Wound and body fluid cultures should be left at room temperatures.

4. Use of specimen transport systems
   a) Aerobic transport methods are listed in Table 2. Certain types of swabs should be used for the collection and transport of certain cultures, as described in Table 2.
   b) Specific transport containers for specimens are listed in Table 3.

Table 1: Transport Systems for Anaerobic Specimens

<table>
<thead>
<tr>
<th>System &amp; Supplier</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Syringe or needle aspirates</td>
<td>Express excess air from syringe, remove needle, and use syringe cover to cap syringe. If fairly large volume is collected (2 ml or more), transfer specimen to a sterile container. Anaerobic bacteria can survive for 24 hours at room temperature. a</td>
</tr>
</tbody>
</table>
| BD BD, Div. Of Becton Dickinson and Co., Rutherford, NJ | AMIES Gel without charcoal
Single swab
For abscess wound specimens |
| BD BD, Div. Of Becton Dickinson and Co., Rutherford, NJ | AMIES Gel without charcoal
Soft Aluminum wire
For male urethral sampling, as well as sinus and eye specimens |
<table>
<thead>
<tr>
<th>System &amp; Supplier (references)</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>BD</strong></td>
<td>AMIES Gel without charcoal</td>
</tr>
<tr>
<td>For throat, vaginal, skin and wound specimens (aerobic/anaerobic)</td>
<td></td>
</tr>
<tr>
<td><strong>BD</strong></td>
<td>AMIES Gel without charcoal</td>
</tr>
<tr>
<td>For male urethral sampling, as well as ear, nose, throat, and eye specimens</td>
<td></td>
</tr>
<tr>
<td>Calcium alginate swabs</td>
<td>Can be toxic for some strains of <em>N. gonorrhoeae</em>, HSV, and <em>Ureaplasma urealyticum</em>, and may be toxic for some cell cultures. Useful for collection of <em>Chlamydia</em> cultures.</td>
</tr>
<tr>
<td>Cotton swabs</td>
<td>Residual fatty acids may inhibit some bacteria and <em>Chlamydia</em> spp. If cotton is glued or spun to wooden applicator stick, wooden stick may inactivate HSV and interfere with some <em>Ureaplasma</em> identification tests.</td>
</tr>
<tr>
<td>Dacron swabs</td>
<td>Useful in collection of viral and group A streptococcus specimens</td>
</tr>
<tr>
<td>Nasopharyngeal-urethrogenital swabs (example: Calgiswab type IV; Spectrum Diagnostics, Glenwood, IL)</td>
<td>Flexible wire shafts and small tips provide easier specimen collection, especially for collection of nasopharyngeal specimens, <em>(UTM)</em> <em>B. perussis</em>, and male urethral specimens of <em>N. gonorrhoeae</em>. Male urogenital = Amies w/o charcoal soft aluminum wire.</td>
</tr>
<tr>
<td>Sterile screw-cap cups</td>
<td>Useful for collection of urine, sputum, stool, bronchoaveolar lavage, and biopsy specimens. Useful for hair or skin-scraping specimens. If biopsy specimen is small, add small amount of sterile nonbacteriostatic 0.85% NaCl to cup. <strong>Never place biopsy specimen in formalin or wrap in gauze.</strong></td>
</tr>
<tr>
<td>Sterile tubes (screw-cap plastic tubes, sterile Vacutainer tubes without additives)</td>
<td>Useful for collection of sterile fluids, bronchoaveolar lavage, drainage or brush specimens.</td>
</tr>
<tr>
<td>Source and Type of Specimen</td>
<td>Transport Method</td>
</tr>
<tr>
<td>---------------------------------------</td>
<td>----------------------------------------------------------------------------------</td>
</tr>
<tr>
<td>Blood</td>
<td>Bactec bottles plus Ferobic/F and Lytic Anaerobic or Peds Plus/F</td>
</tr>
<tr>
<td>CNS</td>
<td>Sterile screw-cap tube</td>
</tr>
<tr>
<td>CNS</td>
<td>Sterile screw-cap tube</td>
</tr>
<tr>
<td>CNS</td>
<td>Sterile screw-cap cup or tube.</td>
</tr>
<tr>
<td>Gastrointestinal System</td>
<td>Sterile screw-cap cup or tube.</td>
</tr>
<tr>
<td>Feces</td>
<td>Sterile screw-cap cup or tube.</td>
</tr>
<tr>
<td>Rectal swab</td>
<td>Sterile screw-cap or Lukens trap</td>
</tr>
<tr>
<td>Gastric lavage or washings</td>
<td>Sterile screw-cap or Lukens trap</td>
</tr>
<tr>
<td>Duodenal aspirate</td>
<td>Sterile screw-cap cup or tube.</td>
</tr>
<tr>
<td>Rectal biopsy</td>
<td>Sterile screw-cap cup or tube.</td>
</tr>
<tr>
<td>Signoidoscopy specimen</td>
<td>Sterile screw-cap cup or tube.</td>
</tr>
<tr>
<td>Eye</td>
<td>Send prepared smears and directly inoculated media.</td>
</tr>
<tr>
<td>Conjunctival scrapings</td>
<td>Send prepared smears and directly inoculated media.</td>
</tr>
<tr>
<td>Corneal scrapings</td>
<td>Send prepared smears and directly inoculated media or</td>
</tr>
<tr>
<td>Intraocular fluid</td>
<td>Sterile screw-cap tube/cup, or capped syringe without needle with air expelled.</td>
</tr>
<tr>
<td>Genital tract, female</td>
<td>Sterile screw cap cup of 1 - 2 ml sample</td>
</tr>
<tr>
<td>Amniotic fluid</td>
<td>Sterile screw cap cup of 1 - 2 ml sample</td>
</tr>
<tr>
<td>Bartholin fluid</td>
<td>Gel swab transport system 1- 2 ml of sample</td>
</tr>
<tr>
<td>Fallopian tube</td>
<td>Gel Swab transport, viral or chlamydial transport, Aptima Probe (GC/Chlamydia)</td>
</tr>
<tr>
<td>Cervical</td>
<td>Gel Swab transport, viral or chlamydial transport, Aptima Probe</td>
</tr>
<tr>
<td>Urethral</td>
<td>Gel Swab transport, viral or chlamydial transport, Aptima Probe</td>
</tr>
<tr>
<td>Vaginal/Rectal</td>
<td>Gel Swab transport, viral or chlamydial transport, Aptima Probe</td>
</tr>
<tr>
<td>Endometrial</td>
<td>Sterile screw-cap cup or tube</td>
</tr>
<tr>
<td>Vulva</td>
<td>Capped syringe without needle; swab transport, viral or chlamydial transport, Aptima Probe</td>
</tr>
</tbody>
</table>
Table 3: Specimen Transport Guide

<table>
<thead>
<tr>
<th>Source and Type of Specimen</th>
<th>Transport Method</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Genital tract, male</strong></td>
<td></td>
</tr>
<tr>
<td>Anal swab</td>
<td>Gel swab transport, viral or chlamydial transport, Aptima Probe</td>
</tr>
<tr>
<td>Urethral</td>
<td>Gel Swab transport, viral or chlamydial transport, Aptima Probe</td>
</tr>
<tr>
<td>Epididymis</td>
<td>Gel swab transport system; sterile screw-cap cup, Aptima Probe</td>
</tr>
<tr>
<td>Prostatic massage</td>
<td>Sterile screw-cap cup, tube or swab transport system, Aptima Probe</td>
</tr>
<tr>
<td>Semen</td>
<td>Sterile screw-cap cup, tube or swab transport system, Aptima Probe</td>
</tr>
<tr>
<td>Penile lesion</td>
<td>Capped syringe <strong>without needle</strong>; Gel swab transport, viral or chlamydial transport system, Aptima Probe</td>
</tr>
<tr>
<td><strong>Specimens for <em>N. gonorrhoeae</em></strong></td>
<td></td>
</tr>
<tr>
<td>Anal, cervical, urethral, vaginal</td>
<td>Gel swab transport</td>
</tr>
<tr>
<td><strong>Lower respiratory tract</strong></td>
<td></td>
</tr>
<tr>
<td>Lung biopsy</td>
<td>Sterile screw-cap cup; if specimen is small, place it in a small amount of sterile 0.85% NaCl (<strong>Never place in formalin-for Microbiology</strong>).</td>
</tr>
<tr>
<td>Expectorated sputum</td>
<td>Sterile screw-cap cup</td>
</tr>
<tr>
<td>Induced sputum</td>
<td>Sterile screw-cap cup</td>
</tr>
<tr>
<td>Tracheal or endotracheal aspirate</td>
<td>Lukens trap or sterile screw-cap cup or tube</td>
</tr>
<tr>
<td>Bronchoalveolar lavage fluid</td>
<td>Lukens trap or sterile screw-cap cup or tube</td>
</tr>
<tr>
<td>Bronchial washings</td>
<td>Lukens trap or sterile screw-cap cup or tube</td>
</tr>
<tr>
<td>Transbronchial biopsy</td>
<td>Sterile screw-cap tube with 1-2 ml of sterile 0.85% NaCl</td>
</tr>
<tr>
<td>Bronchial brush</td>
<td>1 – 2 unfixed slides labeled</td>
</tr>
<tr>
<td>Transtracheal aspirate</td>
<td>Sterile screw-cap cup or tube</td>
</tr>
<tr>
<td>Lung aspirate</td>
<td>Sterile screw-cap cup or tube</td>
</tr>
<tr>
<td><strong>Upper respiratory tract</strong></td>
<td></td>
</tr>
<tr>
<td>Throat swab</td>
<td>Gel swab transport or viral transport system</td>
</tr>
<tr>
<td>Nasal swab</td>
<td>Gel swab transport or viral transport system</td>
</tr>
<tr>
<td>Oral culture</td>
<td>Gel swab transport or viral transport system</td>
</tr>
<tr>
<td>Nasopharyngeal swab</td>
<td>Gel swab transport or viral transport system</td>
</tr>
<tr>
<td>Tympanocentesis fluid</td>
<td>Capped syringe <strong>without needle or sterile screw-cap cup</strong></td>
</tr>
<tr>
<td>Sinus aspirate</td>
<td>Capped syringe <strong>without needle or sterile screw-cap cup</strong></td>
</tr>
<tr>
<td>Nasopharyngeal suction/ Nasal washings</td>
<td>Sterile screw-cap cup or viral transport system</td>
</tr>
<tr>
<td><strong>Sterile body fluids (excluding CSF, urine, blood) Pleural, peritoneal, ascites, joint and synovial fluid</strong></td>
<td>Sterile container, capped syringe <strong>without needle</strong></td>
</tr>
<tr>
<td><strong>Deep wounds, aspirates, tissues</strong></td>
<td></td>
</tr>
<tr>
<td>Site wound</td>
<td>Gel swab transport</td>
</tr>
<tr>
<td>Deep wounds or abscesses</td>
<td>Sterile screw-cap container or gel swab transport system</td>
</tr>
<tr>
<td>Soft tissue aspirates</td>
<td>Capped syringe <strong>without needle or sterile screw-cap cup</strong></td>
</tr>
<tr>
<td>Bone</td>
<td>Sterile screw-cap cup; if specimen is small, place it in a small amount of sterile 0.85% NaCl to prevent drying.</td>
</tr>
<tr>
<td>Punch skin biopsy</td>
<td>Sterile screw-cap cup; if specimen is small, place it in a small amount of sterile 0.85% NaCl to prevent drying.</td>
</tr>
<tr>
<td><strong>Urine</strong></td>
<td></td>
</tr>
<tr>
<td>Clean catch</td>
<td>Sterile screw-cap cup or tube, boric acid tube (gray top) for culture</td>
</tr>
<tr>
<td>Catheter (In/Out or Indwelling)</td>
<td>Sterile screw-cap cup or tube, boric acid tube (gray top) for culture</td>
</tr>
<tr>
<td>Suprapubic aspirate</td>
<td>Capped syringe <strong>without needle or sterile screw-cap cup or tube</strong></td>
</tr>
<tr>
<td>Bladder washout</td>
<td>Sterile screw-cap cup or tube (be careful to label specimens with correct <strong>times and sites</strong>)</td>
</tr>
<tr>
<td>Bilateral ureteral catheterization</td>
<td>Sterile screw-cap cup or tube (be careful to label specimens with correct <strong>times and sites</strong>)</td>
</tr>
</tbody>
</table>
Specimens obtained by a physician using needle aspiration should be transferred to a sterile tube or container prior to transport of the specimen to the laboratory. If there is little material in the syringe, the physician should draw a small amount of sterile nonbacteriostatic 0.85% NaCl or sterile broth through the syringe and then transfer the specimen to a sterile tube. Alternatively, and only if the specimen will be compromised by transferring it from the syringe, a small amount of sterile 0.85% NaCl or broth may be drawn into the syringe prior to removal of the needle. The physician should use a protective device while removing the needle to avoid injury and should cap the syringe with a sterile cap prior to transporting it to the laboratory. **SYRINGES WITH NEEDLES WILL BE REJECTED.**

B. Transport of Diagnostic Specimens

1. SMMC Laboratory provides special transport bags to assist with handling of diagnostic specimens. All specimens must be in a leak-resistant primary container labeled “Biohazard” and must be placed in leak resistant secondary Biohazard plastic bags.

2. The requisition must be placed in the pouch on the back of the sample bag.

3. Couriers are not allowed to pick up specimens that are leaking and not in secondary containers such as the transport bag.

4. **Make sure specimens are sent at the appropriate temperature.** Most specimens should be sent at 2-8 degrees Celcius or refrigerated. Remember to NEVER FREEZE whole blood, except in very rare instances that instructions specify. Call SMMC staff at (949) 452-3554 if unsure about temperature requirements.

III. Collection Instructions for Different Anatomic Sites

Any collection method requiring an invasive technique should be performed by a physician. Some specimen collection techniques should be performed only by a physician specialist with advanced training and skills. The specimen collection guidelines in this section are brief summaries of the procedures used for specimen collection and are not intended to be used as a step-by-step guide for obtaining specimens.

A. Blood Cultures Collections are defined in policy **PL 417 - BLOOD CULTURE COLLECTION PROCEDURE.**

B. Central Nervous System (CNS) Specimens

1. CSF
   Suggested volumes are 1ml, 2ml, and 3 ml for routine, fungal, and mycobacterial cultures, respectively.

   a) Lumbar puncture
      i. Clean the puncture site with antiseptic solution and alcohol before needle insertion to prevent introduction of infection.
      ii. Insert a needle with stylet at the L3-L4, L4-L5, or L5-S1 interspace. When the subarachnoid space is reached, remove the stylet and spinal fluid will appear in the needle hub.
      iii. Slowly drain the CSF into the sterile leakproof tubes. Three tubes are generally required for microbiology, hematology, and chemistry testing. The second tube drawn will generally go to microbiology, and the last tube drawn will generally go to hematology. (In traumatic taps, the CSF will often clear as the later tubes are collected.)
b) Ommaya reservoir fluid
   i. Clean the Ommaya reservoir site with antiseptic solution and alcohol prior to removal of
      Ommaya fluid to prevent introduction of infection.
   ii. Remove Ommaya fluid via the Ommaya reservoir unit, and place it in a sterile tube.

2. Other CNS specimens
   a) Brain abscess
      i. Ninety percent of brain abscesses will grow anaerobic bacteria. Transport the specimen without
         delay in a sterile screw-cap cup to the microbiology laboratory for IMMEDIATE processing.
      ii. A physician aspirates material from a lesion and sends it to the microbiology laboratory in a
          sterile screw-cap cup or syringe. See Table 1 for transport of specimens in a syringe.
   b) CNS biopsy samples
      i. Obtain a biopsy sample from the lesion at surgery and send it to the microbiology laboratory in
         a sterile screw-cap cup for immediate processing. DO NOT ADD FORMALIN.

3. CNS specimen collection considerations are outlined in Table 4.

Table 4: Collection Considerations for CNS Specimens

<table>
<thead>
<tr>
<th>Culture</th>
<th>Vol (ml)</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bacteria</td>
<td>1</td>
<td>Send cloudiest CSF specimen to microbiology laboratory immediately; (Tube no. 2 is preferred).</td>
</tr>
<tr>
<td>Fungi</td>
<td>2</td>
<td>Rule out Cryptococcus spp., Coccidioides immitis Mycobacteria: Mycobacterium tuberculosis, Mycobacterium Avium, Mycobacterium intracellulare</td>
</tr>
<tr>
<td>Mycobacteria</td>
<td>2</td>
<td>Brain abscess or CNS biopsy specimens</td>
</tr>
<tr>
<td>Anaerobes</td>
<td>NA</td>
<td>Brain abscess or CNS biopsy specimens</td>
</tr>
<tr>
<td>Parasites</td>
<td>NA</td>
<td>Brain abscess or CNS biopsy specimens for Entamoeba histolytica, Toxoplasma gondii, Naegleria species(^b), Acanthamoeba spp(^b)</td>
</tr>
<tr>
<td>Virus</td>
<td>1-2</td>
<td>Send to laboratory on ice.</td>
</tr>
</tbody>
</table>

\(^a\) Amounts are guidelines applicable. Greater volumes increase the chance of organism recovery. NA, not applicable.\(^b\) CSF can be submitted.

C. Stool Specimens

1. GI PCR
   GI PCR will include bacteria, virus, and parasites. 5 ml of a diarrhea specimen is placed into a sterile
   container. Per Centers for Disease Control guidelines, the specimen must take the shape of the
   container for it to be acceptable.

2. Stool cultures
   a) Fecal specimens for culture are submitted primarily for the detection of Campylobacter, Shigella,
      Salmonella, Plesiomonas, E. coli-O157, Aeromonas species and Vibrio.

   b) General considerations
      i. Keep stool specimen cool (not frozen); do not incubate them.
      ii. The specimen should be placed in a sterile container.
      iii. Do not use toilet paper to collect stool. Toilet paper may be impregnated with barium salts
           which are inhibitory for some fecal pathogens.
c) Have patient obtain stool specimen by one of the following methods:
   i. Pass stool directly into a sterile, wide-mouth container. Transfer a portion of the stool into the sterile container.
   ii. Pass stool into a clean, dry bedpan, and transfer portion of the stool into a sterile container.

D. Gastrointestinal Tract
The gastrointestinal tract includes the esophagus, stomach, duodenum, small intestine, and colon.

1. Rectal swabs
   a) Submitted primarily for the detection of *Neisseria gonorrhoeae*, *Shigella* species, Herpes simplex virus (HSV), and anal carriage Vancomycin Resistant Enterococcus.
   b) Pass the tip of a sterile swab approximately 1 inch beyond the anal sphincter. Carefully rotate the swab to sample the anal crypts, and withdraw the swab. Send the swab in a gel swab transport (Table 2).

2. Gastric aspirates
   a) The patient should fast prior to each of the following procedures.
   i. Gastric lavage
      (a) Submitted primarily for the detection of *Mycobacterium tuberculosis* in patients (most frequently children) unable to produce quality sputum. Should be performed after the patient wakes in the morning so that sputum swallowed during sleep is still in the stomach.
      (b) Pass a well-lubricated tube orally or nasally through to the stomach of the patient, and perform the lavage. Before removing the tube, release the suction and clamp to prevent mucosal trauma and/or aspiration.
   ii. Duodenal aspiration
      (a) Submitted primarily for the detection of *Giardia* species and larvae of *Strongyloides stercoralis* and *Ascaris lumbricoides*.
      (i) Pass a tube orally through to the duodenum of the patient.
      (ii) To aspirate a sample for giardiasis, the tube should be at least in the third portion of the duodenum.

3. Gastric biopsies and washings
   a) The patient should fast prior to each of the following procedures.
   b) Esophageal, stomach, or duodenal specimens
      i. Esophageal specimens are primarily used to detect Candida species, cytomegalovirus (CMV), and HSV infections. Stomach and duodenal specimens are primarily used for the detection of *Helicobacter pylori*.
      Duodenal specimens can also be used for the detection of *Giardia* species and the larvae of *S. stercoralis* and *A. lumbricoides*.
      (a) Pass an endoscope orally.
      (b) Obtain specimens through a channel in the endoscope by using one of the following procedures.
      (i) Using biopsy forceps, obtain samples from the esophagus, stomach, or duodenum.
      (ii) Using a sheathed brush, brush suspicious areas several times to obtain adequate cellular material.
   c) Gastric Lavage
      i. Perform a wash by injecting approximately 25 to 30 ml of sterile nonbacteriostatic isotonic 0.85% NaCl through the biopsy channel onto the lesion.
      ii. Collect the specimen by aspirating the fluid through the scope into a sterile trap, which is connected to the suction tubing. NOTE: If a gastric ulcer is seen, obtain biopsy samples from the base, the surrounding gastric mucosa, and each of the four quadrants of the margin.
d) Rectal biopsy
i. Submitted primarily for the detection of *Entamoeba histolytica*, *Balantidium coli*, and HSV.
ii. If lesions are not evident, biopsy the posterior rectal mucosa below the peritoneal reflection (within 7 to 10 cm of the anal verge).

e) Small bowel biopsy
i. Submitted primarily for the detection of *Giardia*, *Cryptosporidium*, and *Microsporidium* species.
ii. Biopsies of the small intestine provide the highest diagnostic yield for *Microsporidia* species. Biopsies from other gastrointestinal sites (stomach, colon, rectum) have a much lower yield in comparison. Obtain biopsy sample of lesion at surgery.

f) Sigmoidoscopy
i. Useful in the detection of *E. histolytica* and *Mycobacterium* species and the diagnosis of pseudomembranous colitis associated with *C. difficile* and possibly *Staphylococcus aureus*.
   (a) Perform flexible or rigid sigmoidoscopy.
   (b) Obtain endoscopic pinch biopsy samples of any lesions seen. Additionally, aspirate liquid from the inflammed bowel with a pipette passed through the sigmoidoscope. Transport specimens in a sterile screw-cap container. If biopsy samples are small, add a small amount of sterile nonbacteriostatic 0.85% NaCl to prevent the specimen from drying.

g) Gastrointestinal specimen collection considerations are summarized in Table 5.

Table 5: Collection Considerations for Gastrointestinal Specimens

<table>
<thead>
<tr>
<th>Culture</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bacteria</td>
<td>Stools: One stool specimens recommended.</td>
</tr>
<tr>
<td></td>
<td>Gastric biopsy: Rule of <em>Helicobacter pylori</em>.</td>
</tr>
<tr>
<td></td>
<td>Rectal swab: Rule out enteric pathogens (especially <em>Shigella</em> spp.) (<em>Neisseria gonorrhoeae</em> or <em>VRE</em>).</td>
</tr>
<tr>
<td>Fungi</td>
<td>Gastric aspirate, gastric biopsy, esophageal brush, esophageal biopsy.</td>
</tr>
<tr>
<td>Pinworm</td>
<td>For pinworm collection kit, contact the laboratory.</td>
</tr>
<tr>
<td></td>
<td>Swab perianal area when patient gets up in the morning before patient bathes or defecates.</td>
</tr>
<tr>
<td>Mycobacteria</td>
<td>Gastric aspirate or gastric biopsy; feces.</td>
</tr>
<tr>
<td>Parasites</td>
<td>If transport to laboratory is delayed, place specimen in para-pak.</td>
</tr>
<tr>
<td></td>
<td>Duodenal aspirates are useful for detecting <em>Giardia</em> spp. and larvae of <em>S. stercoralis</em> and <em>A. lumbricoides</em>.</td>
</tr>
<tr>
<td></td>
<td>Use rectal biopsy specimens for <em>E. histolytica</em> and <em>B. coli</em>.</td>
</tr>
<tr>
<td></td>
<td>Use small-bowel biopsy specimens for <em>Giardia</em>, <em>Cryptosporidium</em>, and <em>Microsporidium</em> spp.</td>
</tr>
<tr>
<td>Virus</td>
<td>Use esophageal specimens for CMV and HSV and rectal biopsy specimens for HSV. Send to laboratory in Viral Transport Media. Do not freeze.</td>
</tr>
</tbody>
</table>

E. Genital Tract Specimens

1. Female
Genital tract specimens are submitted primarily for the detection of sexually transmitted pathogens (such as *N. gonorrhoeae*, *Chlamydia trachomatis*, lumphogranuloma venereum, HSV, human papillomavirus (HPV) with histological confirmation), trichomonads, group B streptococci, and *Candida* infections) (Table 1). See Table 2 for *N. gonorrhoeae* transport and swab transport systems. For chlamydial and viral transport systems, see Table 13.

   a) Amniotic fluid
      i. Aspirate fluid by catheter, at cesarean section, or at amniocentesis.
   b) Bartholin gland
      i. Decontaminate the skin with povidone-iodine, and aspirate material from the ducts.
c) Cervix
   i. Do not use lubricant during procedure.
   ii. Wipe the cervix clean of vaginal secretion and mucus.
   iii. Rotate a sterile swab, and obtain exudate from the endocervical glands.
   iv. If no exudate is seen, insert a sterile swab into the endocervical canal, and rotate the swab

d) Endometrium
   i. Collect the endometrium specimens by transcervical aspiration through a telescoping catheter.

e) Fallopian tubes
   i. Obtain aspirates (preferably) or swab specimens during surgery. Bronchoscopy cytology brushes may be used if exudate is not expressed.

f) Rectal swabs:
   i. Used primarily to detect N. gonorrhoeae, Shigella species, HSV, and anal carriage of VRE.
   ii. Pass the tip of a sterile swab approximately 1 inch beyond the anal sphincter. Carefully rotate the swab to sample the anal crypts, and withdraw it. Send the swab in a swab transport, viral transport (for HSV), or N. gonorrhoeae transport system (Table 2).

g) Urethra
   i. Collect specimens 1 hour or more after patient has urinated.
   ii. Stimulate discharge by gently massaging the urethra against the pubic symphysis through the vagina.
   iii. Collect the discharge with a sterile swab.
   iv. If discharge cannot be obtained, wash external urethra with betadine soap and rinse with water. Insert a urethrogenital swab 2 to 4 cm into the endourethra, gently rotate the swab, and leave it in place for 1-2 seconds. Withdraw the swab, and submit it in the appropriate transport system for culture.

h) Vagina
   i. Use a speculum without lubricant. Collect secretions from the mucosa high in the vaginal canal with sterile pipette or swab.

i) Vulva
   i. Clean the surface of the lesion with 0.85% NaCl. If there is a crust on the lesion, remove it.
   ii. Scrape the lesion until serous fluid emerges.
   iii. Wipe away fluid and debris with sterile gauze. (Try to avoid bleeding.)
   iv. Press the base of lesion until clear fluid is expressed.
   v. Aspirate vesicular fluid with a 26 to 27-gauge needle.
   vi. Unroof the vesicle, and collect fluid with a sterile swab (for HSV Detection.) OR
   vii. Scrape the base of an open vesicle with a sterile scalpel blade, and then rub the base vigorously with a sterile swab (for HSV and H. ducreyi detection).

j) Vaginal/pectal
   i. Screening for Group B Streptococcus at 35-37 weeks gestation.
   ii. Swab to lower vagina followed by the rectum using the same swab.
   iii. Submit swab to the laboratory.

2. Male
   See Table 2 for N. gonorrhoeae transport and swab transport systems.

a) Anal swab
   i. Submitted primarily for the detection of N. gonorrhoeae, Shigella species, HSV, and anal carriage of S. pyogenes.
      Pass the tip of a sterile swab approximately 1 inch beyond the anal sphincter. Carefully rotate the swab to sample the anal crypts, and withdraw it. Send the swab in a swab transport, viral transport (for HSV), or N. gonorrhoeae transport system (Table 2).
b) Epididymis
   i. Used primarily to diagnose nonspecific bacterial epididymitis and *M. tuberculosis* infections generally occur after involvement of the prostate or seminal vesicles. Sexually transmitted epididymitis is most commonly due to *C. trachomatis* and *N. gonorrhoeae*.
   ii. Use a needle and syringe to aspirate material from the epididymis.

c) Penile lesion
   i. Used primarily to detect sexually transmitted pathogens such as *N. gonorrhoeae*, *C. trachomatis*, lymphogranuloma venereum, HSV, and *H. ducreyi*.
   ii. Clean the surface of the lesion with 0.85% NaCl. If there is a crust on the lesion, remove it.
   iii. Scrape the lesion until serous fluid emerges.
   iv. Wipe away fluid and debris with sterile gauze. (Try to avoid bleeding.)
   v. Press the base of the lesion until clear fluid is expressed.
   vi. Aspirate vesicular fluid with a 26 to 27-gauge needle. Refer to Table 1 for instructions on properly transporting specimens collected in a syringe.
   vii. Unroof the vesicle, and collect fluid with a sterile swab (for HSV detection) OR
   viii. Scrape the base of an open vesicle with a sterile scalpel blade, and rub the base vigorously with a sterile swab (for HSV and *H. ducreyi* detection).

d) Prostatic massage
   i. Used primarily to diagnose acute or chronic prostatitis. For both diseases, gram-negative enteric organisms are the most frequently isolated pathogens. *N. gonorrhoeae* is found infrequently, but is sometimes implicated in acute protatitis.
   ii. Perform a digital massage through the rectum.
   iii. Collect the specimen in a sterile tube or on a sterile swab.

e) Urethra
   i. Used primarily to detect *N. gonorrhoeae* and *C. trachomatis*.
   ii. Collect specimens at least 2 hours after the patient has urinated.
   iii. Insert a thin urethrogenital swab 2-4 cm into the endourethra, gently rotate it, leave it in place for 1-2 seconds, and withdraw it.

f) Specimen collection considerations for genital tract specimens are summarized in Table 6.

<table>
<thead>
<tr>
<th>Table 6: Collection Considerations for Genital Tract Specimens</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Culture</strong></td>
</tr>
<tr>
<td>----------------</td>
</tr>
<tr>
<td><em>N. gonorrhoeae</em></td>
</tr>
<tr>
<td>Bacteria</td>
</tr>
<tr>
<td><em>Trichomonas vaginalis</em></td>
</tr>
<tr>
<td>Fungi</td>
</tr>
<tr>
<td>Anaerobes</td>
</tr>
<tr>
<td>HSV</td>
</tr>
<tr>
<td><em>C. trachomatis</em></td>
</tr>
<tr>
<td><em>U. urealyticum</em></td>
</tr>
<tr>
<td>LGV</td>
</tr>
<tr>
<td><em>H. ducreyi</em></td>
</tr>
</tbody>
</table>
F. Ocular Specimens

1. General considerations
   a) Obtain viral and chlamydia samples before topical anesthetics are instilled.
   b) Obtain samples for chlamydial cultures with calcium alginate swabs and for viral cultures with Dacron swabs or cotton swabs with non-wood shafts (also refer to Table 2).
   c) Send prepared smears and inoculated media to the laboratory immediately.
   d) For appropriate media, call the Microbiology Department at (949) 452-3560.

2. Conjunctival scrapings
   a) One or two drops of topical anesthetic are generally instilled.
   b) Scrape the lower tarsal conjunctiva with a sterilized kimura spatula.
   c) Inoculate the appropriate media directly.
   d) Prepare smears by applying the scraping in a circular manner to a clean glass slide or by compressing material between two glass slides and pulling the slides apart.
   e) Alternatively, use a calcium alginate swab or a cotton-tipped applicator to swab the inferior tarsal conjunctiva (inside surface of eyelid) and the fornix of the eye. However, organisms are more readily detected in scrapings than from a swab.

3. Corneal scrapings
   a) Obtain conjunctival samples prior to corneal scrapings. Sometimes conjunctival cultures are helpful in assessing the possibility of contamination of corneal cultures.
   b) One or two drops of topical anesthetic are generally instilled.
   c) Using short, firm strokes in one direction, scrape multiple areas of ulceration and suppuration with a sterilized kimura spatula. (Keep the eyelid open, and be careful not to touch the eyelashes.)
   d) Inoculate each scraping directly to appropriate media. (Multiple scrapings are recommended because the depth and extent of viable organisms may vary.)
   e) Prepare smears by applying the scrapings in a gentle circular motion over a clean glass slide or by compressing material between two clean glass slides and pulling the slides apart.

4. Intraocular fluid
   a) Use a needle aspiration technique to collect intraocular fluid.
   b) Inoculate appropriate media directly, and/or immediately transport the samples to the laboratory in an aerobic transport system or a capped syringe with air bubbles expelled. Refer to Table 1 for instructions on properly transporting specimens collected in a syringe. Prepare smears by spreading a drop of material over the surface of a cleaned glass slide with a sterile kimura spatula or by compressing the material between two glass slides and pulling the slides apart.

5. Specimen collection considerations for ocular specimens are summarized in Table 7.

<table>
<thead>
<tr>
<th>Culture</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bacteria</td>
<td>Inoculate media directly with ocular scrapings. (If <em>N. gonorrhoeae</em> is suspected, inoculate a Thayer-Martin agar plate also.</td>
</tr>
<tr>
<td>Fungi</td>
<td>Inoculate media directly with ocular scrapings.</td>
</tr>
<tr>
<td>Anaerobes</td>
<td>Use anaerobic transport medium, or inoculate medium directly. If inflammation occurs after extracapsular cataract extraction, rule out <em>Propionibacterium</em> spp.</td>
</tr>
<tr>
<td>Parasites</td>
<td>Use to detect <em>Acanthamoeba</em> spp.</td>
</tr>
<tr>
<td>Chlamydia</td>
<td>Do not use cotton swabs for specimen collection.</td>
</tr>
<tr>
<td>Virus</td>
<td>Do not use calcium alginate swabs for specimen collection.</td>
</tr>
<tr>
<td>Mycobacteria</td>
<td>Ocular scrapings: use sterile screw-cap cup or tube. If specimen is small, add a small amount of non-bacteriostatic 0.85% NaCl.</td>
</tr>
</tbody>
</table>
G. Respiratory Specimens

1. General considerations
   a) Twenty-four-hour sputum collections are not acceptable for culture.
   b) If *Corynebacterium diphtheriae*, *Arcanobacterium haemolyticum*, *Bordetella pertussis*, *N. gonorrhoeae*, legionellae, chlamydiae, or mycoplasmas are suspected, the physician should contact the clinical microbiology laboratory prior to specimen collection because special techniques and/or media are required for the isolation of these agents.

2. Lower respiratory tract
   a) Expectorated sputum
      i. If possible, have the patient rinse mouth and gargle with water prior to sputum collection.
      ii. Instruct the patient not to expectorate saliva or postnasal discharge into the container.
      iii. Collect specimen resulting from deep cough in sterile screw-cap cup or other suitable sterile collection assembly.
   b) Induced sputum
      i. Using a wet toothbrush, brush the buccal mucose, tongue, and gums prior to the procedure.
      ii. Rinse the patient's mouth thoroughly with water.
      iii. Using an ultrasonic nebulizer, have the patient inhale approximately 20 - 30 ml of 3 - 10% 0.85% NaCl.
      iv. Collect the induced sputum in a sterile screw-cap cup or other suitable sterile collection assembly.
   c) Tracheostomy and endotracheal aspirations
      i. Tracheostomy is followed by colonization within 24 hours of insertion of the tube. Results must be correlated with clinical findings such as fever or infiltrate on chest X-ray. Aspirate the specimen into a sterile sputum trap.
   d) Bronchoscopy specimens
      i. Bronchoscopy specimens include bronchoalveolar lavage, bronchial washing, bronchial brushing, and transbronchial biopsy specimens.
      ii. Pass the bronchoscope transnasally or transorally in non-intubated patients or via the endotrachael tube in intubated patients.
      iii. Wedge the tip of the bronchoscope in a segmental (for bronchial wash) or subsegmental (for bronchoalveolar lavage) bronchus.
      iv. To obtain specimens
         (a) Bronchial wash or bronchoalveolar lavage. Bronchial wash and bronchoalveolar lavage specimens are generally obtained before brushing or biopsy specimens to avoid excess blood in the recovered fluid, because blood may alter the concentration of cellular and noncellular components.
            (i) Inject sterile nonbacteriostatic 0.85% NaCl (generally 5 - 20 ml aliquots) from a syringe through a biopsy channel of the bronchoscope.
            (ii) Gently suction the 0.85% NaCl into a sterile container before administering the next aliquot. (In general, 50 - 75% of the 0.85% NaCl instilled is recovered in the lavage effluent.)
            (iii) Keep aliquots separate during collection. Combine aliquots from the same site for microbiology cultures and smears, but aliquots from separate sites (for example, right upper lobe and right lower lobe) should be combined only after consultation with the physician of record.
         (b) Bronchial brush specimens
            (i) Insert a telescoping double catheter plugged with polyethylene glycol at the distal end (to prevent contamination of the bronchial brush) through the biopsy channel of the bronchoscope.
(c) Transbronchial biopsies
   (i) Obtain the biopsy sample through the biopsy channel of the bronchoscope, and transport it in a sterile container with a small amount of nonbacteriostatic sterile 0.85% NaCl.

(d) Lung aspirations
   (i) Use a computed tomography scan to obtain lung aspirates by inserting a needle through the chest wall into a pulmonary infiltrate. Aspirate material from the lesion. If the lesion is large or if there are multiple lesions, collect multiple specimens from representative sites. Refer to Table 1 for instructions on properly transporting specimens collected in a syringe.

(e) Lung biopsies
   (i) Obtain a 1-3 cm square piece of tissue is possible. If the lesion is large or if there are multiple lesions, collect multiple specimens from representative sites. Submit in a sterile container(s) without formalin.

3. Upper respiratory tract infections
   (a) Throat (pharyngeal specimens)
      (i) Submitted primarily for the detection of group A streptococci (can also be used to detect *N. gonorrhoeae*, *Haemophilus influenzae* [for epiglottitis].
      (ii) Do not obtain throat samples if epiglottis is inflamed, as sampling may cause serious respiratory obstruction.
      (iii) Depress tongue gently with tongue depressor.
      (iv) Extend sterile swab between the tonsillar pillars and behind the uvula. (avoid touching the cheeks, tongue, uvula, or lips).
      (v) Sweep the swab back and forth across the posterior pharynx, tonsillar areas, and any inflamed or ulcerated areas to obtain sample.

(b) Nasal swabs
   (i) Submitted primarily for the detection of MRSA carriers.
   (ii) Insert a sterile swab into the nose until resistance is met at the level of the turbinates (approximately 1 inch into the nose).
   (iii) Rotate the swab against the nasal mucosa.
   (iv) Repeat the process on the other side.

(c) Nasopharyngeal swabs
   (i) Submitted primarily for the detection of carriers of *N. meningitidis*.
   (ii) Carefully insert a flexible-wire calcium alginate-tipped swab through the nose into the posterior nasopharynx, and rotate the swab. (Keep the swab near the septum and floor of the nose.)

(d) Nasal washings
   (i) Submitted primarily for Rapid Influenzae or Respiratory Panel by PCR.
   (ii) Instruct the patient not to swallow during the procedure.
   (iii) With the patient’s head hyperextended (approximately 70° angle), instill approximately 3 - 7 ml of sterile 0.85% NaCl into each nostril.
   (iv) To collect material, aspirate the fluid by inserting a rubber bulb syringe into each nostril.
   (v) Place the saline wash in an equal volume of viral transport medium, or transport it in a sterile container.

(e) Sinus aspirates
   (i) Using a syringe aspiration technique, a specially trained physician or an otolaryngologist will obtain material from maxillary, frontal, or other sinuses.
   (ii) Place the contents of the syringe into an anaerobic transport system, or send the specimen in the syringe.
(f) Typanocentesis fluid
(i) Submitted primarily to diagnose middle ear infections only if previous therapy has failed.
(ii) Clean the external canal with mild detergent.
(iii) Using a syringe aspiration technique, the physician will obtain the fluid from the eardrum. Send the specimen in a sterile container, or send it in the syringe.
(iv) If the eardrum is ruptured, collect exudate by inserting a sterile swab through an auditory speculum.

(g) Oral cultures
(i) Used to prepare smears for the detection of yeast or fusospirochetal disease.
(ii) Rinse mouth with sterile saline.
(iii) Wipe the lesion with dry sterile gauze.
(iv) Swab or scrape areas of exudation or ulceration.

4. Respiratory specimen collection considerations are summarized in Table 8.

Table 8: Collection Considerations for Respiratory Specimens

<table>
<thead>
<tr>
<th>Culture</th>
<th>Vol (ml)</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bacteria</td>
<td>NA</td>
<td>Contact laboratory if <em>Legionella</em> is suspected. Submit sputum only; saliva is unacceptable.</td>
</tr>
<tr>
<td>Fungi</td>
<td>3 - 5</td>
<td>Collect early morning fresh specimen resulting from deep cough or sputum induction on three consecutive days. Lung biopsy specimens or lung aspirates are also appropriate.</td>
</tr>
<tr>
<td>Anaerobes</td>
<td>1</td>
<td>Sinus aspirate, typanocentesis fluid, transtracheal aspirate, and lung aspirates or biopsy specimens are appropriate.</td>
</tr>
<tr>
<td>Mycobacteria</td>
<td>5 - 10</td>
<td>Collect three early morning fresh specimens resulting from deep cough or sputum induction. Lung biopsy specimens or lung aspirates are also appropriate.</td>
</tr>
<tr>
<td><em>Pneumocystis</em> spp.</td>
<td>2</td>
<td>Use induced sputum, bronchoaveolar lavage fluid, or lung biopsy specimen.</td>
</tr>
<tr>
<td>Parasites</td>
<td>3 - 5</td>
<td>Can be examined for amoebae, helminth eggs (<em>Paragonimus westermani</em>), hooklets of <em>Echinococcus</em> spp., larvae of hookworm, and <em>Ascaris</em> and <em>Strongyloides</em> spp.</td>
</tr>
</tbody>
</table>

a Amount are guidelines. NA, not applicable.

H. Sterile Body Fluids (excluding CSF, urine, and blood)

1. Clean the needle puncture site with alcohol, and disinfect it with an iodine solution (1-2% tincture of iodine or a 10% solution of povidone-iodine [1% free iodine]) to prevent introduction of infection. (If tincture of iodine is used, remove with 70% ethanol after the procedure to avoid burn.
2. The physician will aseptically perform percutaneous aspiration to obtain pleural, pericardial, peritoneal, or synovial fluids.
3. Expel any air bubbles from the syringe, and immediately inject the specimen into an anaerobic transport system or send the specimen in the syringe. Transport additional fluid or pus in a sterile screw-cap container.
4. Sterile body fluid collection considerations are summarized in Table 9.
Table 9: Collection Considerations for Sterile Body Fluids

<table>
<thead>
<tr>
<th>Culture</th>
<th>Vol (ml)</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bacteria</td>
<td>1 - 5</td>
<td>If gonococcal arthritis is suspected, notify laboratory to add modified Thayer-Martin plate.</td>
</tr>
<tr>
<td>Fungi</td>
<td>&gt; 10</td>
<td>Blood for <em>Histoplasma capsulatum</em> (AIDS), <em>Cryptococcus</em> spp., <em>Candida albicans</em>(^b), and <em>Candida tropicalis</em>(^b).</td>
</tr>
<tr>
<td>Anaerobes</td>
<td>1 - 5</td>
<td>Use sterile screw-cap cup or tube.</td>
</tr>
<tr>
<td>Mycobacteria</td>
<td>&gt; 10</td>
<td>Use sterile screw-cap cup or tube.</td>
</tr>
</tbody>
</table>

\(^a\) Amount are guidelines. Greater volumes will increase the chance of organism recovery.

\(^b\) Can also be recovered in most bacterial cultures.

I. Subcutaneous Tissue and Skin Specimens

1. Burn specimens
   The surfaces of burn wounds will become colonized by the patient's microbiota or by environmental organisms. When the organism load is large, infection of underlying tissue may occur, and bacteria may ensue. Cultures of the surface alone are misleading; therefore, biopsies of deeper tissue are often indicated. Additionally, organisms may not be distributed evenly in the burn wound, so sampling of different areas of the bum is recommended.
   a) Disinfect the surface of the burn with 70% alcohol and then with an iodine solution (1 - 2% tincture of iodine or a 10% solution of povidone-iodine [1% free iodine]). Allow the disinfectant to dry prior to collecting the specimen. *Note: Blood cultures should be used to monitor patient status.* If tincture of iodine is used, it must be removed with 70% alcohol after the procedure to prevent burns.
      b) Collect a punch biopsy sample (3 - 4 mm) for quantitative culture.

2. Superficial wound, bacterial
   a) Syringe aspiration is preferable to swab collection.
   b) Disinfect the surface of the wound with 70% alcohol and then with an iodine solution (1 - 2% tincture of iodine or a 10% solution of povidone-iodine [1% free iodine]). Allow the disinfectant to dry prior to collecting the specimen.
   c) Using a 3 -5 ml syringe with a 22 - 23 gauge needle, a physician will aspirate the deepest portion of the lesion. If a vesicle is present, collect both fluid and cells from the base of the lesion.
   d) If the initial aspiration fails to obtain material, inject sterile, nonbacteriostatic 0.85% NaCl subcutaneously.
   e) Repeat the aspiration attempt.
   f) If no material is obtained, rinse the needle and syringe with broth by drawing the culture medium through the needle into the syringe.

3. Superficial lesions, fungal
   a) Clean the surface with sterile water.
   b) Using a scalpel blade, scrape the periphery of the lesion border. Samples from scalp lesions should include hair that is selectively collected for examination. If there is nail involvement, obtain scrapings of debris or material beneath the nail plate. Transport in a sterile container or sterile petri dish.

4. Ulcers and nodules
   a) Clean the area with 70% alcohol and then with an iodine solution (1-2% tincture of iodine or a 10% solution of povidone-iodine [1% free iodine]). Tincture of iodine must be removed with alcohol after the procedure to prevent burns.
   b) Remove overlying debris.
   c) Curette the base of the ulcer or nodule.
   d) If exudate is present from ulcer or nodule, collect it with a syringe or sterile swab.

5. Subcutaneous tissue and skin specimen collection considerations are summarized in Table 10.
Table 10: Collection Considerations for Subcutaneous Tissue and Skin Specimens

<table>
<thead>
<tr>
<th>Culture</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bacteria</td>
<td>Syringe aspirates or biopsy specimens are preferable to swab specimens.</td>
</tr>
<tr>
<td>Fungi</td>
<td>Useful in diagnosing dermatophytes, yeast, filamentous fungi, and dimorphic fungi.</td>
</tr>
<tr>
<td>Anaerobes</td>
<td>Uncommon in burn, ulcer, nodules, or superficial skin infections; useful following bites and trauma.</td>
</tr>
<tr>
<td>Virus</td>
<td>Useful in diagnosing HSV and varicella-zoster virus.</td>
</tr>
<tr>
<td>Mycobacteria</td>
<td>Useful in diagnosing <em>Mycobacterium marinum</em>, <em>Mycobacterium fortuitum</em>, and <em>Mycobacterium chelonei</em>.</td>
</tr>
</tbody>
</table>

* Rate of recovery of HSV and varicella-zoster virus is highest from the youngest lesions (vesicles), then from pustules, ulcers, and crusted lesions, in that order.

J. Deep Wounds, Aspirates, and Tissue Specimens

1. Bite wounds
   a) Aspirate pus from the wound, or obtain it at the time of incision, drainage, or debridement of the infected wound. (Do not culture fresh bite wounds, as infectious agents will likely not be recovered.)
2. Bone
   a) Obtain bone specimen at surgery.
   b) Submit in sterile container without formalin. Specimen may be kept moist with sterile 0.85% NaCl.
3. Deep wounds or abscesses
   a) Disinfect the surface with 70% alcohol and then with an iodine solution (1 - 2% tincture of iodine or a 10% solution of povidone-iodine (1% free iodine). Tincture of iodine must be removed with 70% alcohol to prevent burns.
   b) Aspirate the deepest portion of the lesion, avoiding contamination by the wound surface. If collection is done at surgery, a portion of the abscess wall should also be sent for culture.
4. Punch skin biopsies
   a) Disinfect the skin surface with 70% alcohol and then with an iodine solution (1 - 2% tincture of iodine or a 10% solution of povidone-iodine (1% free iodine). Tincture of iodine must be removed at completion of procedure to prevent burns.
   b) Collect 3 - 4 mm sample with dermal punch.
   c) Submit for microbiological analysis in sterile container without formalin.
5. Soft tissue aspirate
   a) Disinfect the skin surface with 70% alcohol and then with an iodine solution (1 - 2% tincture of iodine or a 10% solution of povidone-iodine (1% free iodine). Remove tincture of iodine with alcohol after procedure to prevent burns.
   b) Aspirate the deepest portion of the lesion or sinus tract. Be careful to avoid contamination by the wound surface.
6. Deep wound, aspirate, and tissue specimen collection considerations are summarized in Table 11.

Table 11: Collection Considerations for Deep Wound, Aspirate, and Tissue Specimens

<table>
<thead>
<tr>
<th>Culture</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bacteria</td>
<td>Biopsy specimens or aspirates are better than swab specimens.</td>
</tr>
<tr>
<td>Fungi</td>
<td>Useful in diagnosing <em>Pseudallescheria boydii</em>, <em>Bipolaris</em> spp., <em>Exophiala</em> spp., and <em>Fusarium</em> spp..</td>
</tr>
<tr>
<td>Anaerobes</td>
<td>Useful in diagnosing actinomycosis; send in anaerobic transport system.</td>
</tr>
<tr>
<td>Mycobacteria</td>
<td>Useful in diagnosing <em>Mycobacterium tuberculosis</em>, <em>Mycobacterium bovis</em>, and <em>Mycobacterium kansasii</em>.</td>
</tr>
</tbody>
</table>
K. Urine

1. General considerations
   a) **Never** collect urine from a bedpan or urinal.
   b) Thoroughly clean the urethral opening (and vaginal vestibule in females) prior to collection procedures to ensure that the specimen obtained is not contaminated with colonizing microorganisms in this area.
   c) Soap rather than disinfectants is recommended for cleaning the urethral area. If disinfectants are introduced into the urine during collection, they may be inhibitory to the growth of microorganisms.
   d) Submit BD Collection container as follows:
      i. C&S preservative tube (grey top) for C&S,
      ii. UA preservative tube (cherry red/yellow top) for UA,
      iii. Plain red tube for urine chemistry tests.
   e) If not collected using BD Urine Collection system, transport specimen to laboratory **WITHIN 2 HOURS** of collection. If it cannot be transported **within 2 hours** of collection, the urine specimen should be refrigerated. (Bacterial counts remain stable for at least 24 hours at 4°Celsius.) **Do not freeze.**
   f) Use sterile cups or tubes to transport urine.
   g) Transport suprapubic bladder aspirate (SPA) specimens for anaerobic culture in a sterile screw-cap cup or tube.
   h) Any urine collection procedure involving catheterization must be done with scrupulous aseptic technique to avoid introducing microorganisms.
   i) Send the first morning voided urine. Three consecutive first morning urine specimens are recommended for mycobacterial culture.
   j) Do not submit 24 hour urine collections for culture.

2. Collection Techniques for random urine collection
   a) For collection procedures using BD Urine Collection System see **PL 406 – Urine Specimen Collection**.
   b) If indicated, refer to **Urine Collection via Foley Catheter PL 406.1**.
   c) See Patient Instructions Using BD Vacutainer Urine Collection Cup (English, Spanish).
   d) See Mid-Stream Clean Catch Instructions (English, Spanish).

3. Urine Collection (24 Hour) – see Patient Instructions.
Urinalysis Instructions for Patients

Routine Urinalysis

1. **Caution:** DO NOT REMOVE the label from the top of the BD Vacutainer® Urine Collection Cup. There is a needle under the label.
2. Wash hands thoroughly with soap and water.
3. Unscrew the blue cap.
4. Place the blue cap on counter with "straw" facing upwards. Do not touch inside of cap or straw. (Follow mid-stream clean catch directions below if instructed.)
5. Urinate into the BD Vacutainer® Urine Collection Cup without touching the inside or lip of the cup.
6. Replace the blue cap onto the BD Vacutainer® Urine Collection Cup.
7. Return the sample to the healthcare worker or place it in the designated area in the bathroom.

Mid-Stream Clean Catch Instructions

FEMALE CLEANSING INSTRUCTIONS

1. Stand in a squatting position over the toilet. Separate the folds of skin around the urinary opening.
2. Cleanse the area around the opening with the first castile soap towelette.
3. Repeat using a second clean towelette.
4. Urinate the first portion of urine in the toilet.
5. As you continue to urinate, bring the collection cup into the mid-stream to collect the urine sample.
6. Do not touch the inside or lip of the cup.
7. Urinate the remainder of urine into the toilet.
8. Replace the cap onto the urine collection cup.
9. Return the sample to the healthcare worker or place it in the designated area in the bathroom.

Mid-Stream Clean Catch Instructions

MALE CLEANSING INSTRUCTIONS

1. Cleanse the end of the penis with the first castile soap towelette beginning at the urethral opening and working away from it. (The foreskin of an uncircumcised male must be retracted.)
2. Repeat using a second clean towelette.
3. Urinate the first portion of urine in the toilet.
4. As you continue to urinate, bring the collection cup into the mid-stream to collect the urine sample.
5. Do not touch the inside or lip of the cup.
6. Urinate the remainder of urine into the toilet.
7. Replace the cap onto the urine collection cup.
8. Return the sample to the healthcare worker or place it in the designated area in the bathroom.
Uroanálisis Instrucciones para Paciente

Procedimiento de obtención de muestra de orina para prueba de uroanálisis

1. **Precaución: Aguja situada debajo de la etiqueta amarilla.** No remueva la etiqueta amarilla de la parte superior del vaso para recolección de orina BD Vacutainer®.

2. Lávese bien las manos con agua y jabón

3. Remueva la tapa azul. Asegúrese de no tocar el interior o el borde del vaso.

4. Asegúrese de colocar la tapa en el mostrador con la “pajita” hacia arriba. No toque el interior de la tapa ni la pajita. (Siga las instrucciones para obtención de muestra de orina detalladas en la siguiente sección solo si el profesional de la salud así lo indica.)

5. No remueva la etiqueta amarilla de la parte superior de la tapa.

6. Asegúrese de no tocar el interior o el borde del vaso

7. Entregue la muestra al personal del laboratorio o en el bano en el compartimento designado para la muestra.

**Instrucciones para las Mujeres:**

1. Párese sobre el inodoro en cuclillas. Separe los pliegues del piel alrededor de la abertura uniraria. Limpie el área alrededor de la abertura con la primera toallita impregnada con jabón de Castilla (castile soap).

2. Limpie el área alrededor de la abertura con la primera toallita impregnada con jabón de Castilla (castile soap)

3. Repita el procedimiento usando una segunda toallita limpia

4. Evacue la primera parte de la orina en el inodoro.

5. Mientras sigue evacuando la orina, coloque el vaso para recolección en mitad de la micción para obtener la muestra de orina.

6. NO toque el interior o el borde del vaso con las manos ni ninguna otra parte del cuerpo.

7. Evacue el resto de la orina en el inodoro.

8. Cierre el vaso tocando solo las superficies exteriores de la tapa y del vaso.

9. Entregue la muestra al personal del laboratorio o en el bano en el compartimento designado para la muestra.

**Instrucciones para los Hombres:**

a. Limpie la punta del pene con la primera toallita impregnada con jabón de Castilla (castile soap). Comenzando por la abertura uretral y hacia fuera (el prepucio de un hombre no circuncidado se debe retraer primero).

b. Repita el procedimiento usando una segunda toallita limpia

c. Evacue la primera parte de la orina en el inodoro.

d. Mientras sigue evacuando la orina, coloque el vaso para recolección en mitad de la micción para obtener la muestra de orina.

e. NO toque el interior o el borde del vaso con las manos ni ninguna otra parte del cuerpo.

f. Evacue el resto de la orina en el inodoro.

g. Cierre el vaso tocando solo las superficies exteriores de la tapa y del vaso.

h. Entregue la muestra al personal del laboratorio o en el bano en el compartimento designado para la muestra.
Mid-Stream Clean Catch Instructions

FEMALE CLEANSING INSTRUCTIONS

1. Stand in a squatting position over the toilet. Separate the folds of skin around the urinary opening.
2. Cleanse the area around the opening with the first castile soap towelette.
3. Repeat using a second clean towelette.
4. Urinate the first portion of urine in the toilet.
5. As you continue to urinate, bring the collection cup into the mid-stream to collect the urine sample.
6. Do not touch the inside or lip of the cup.
7. Urinate the remainder of urine into the toilet.
8. Replace the cap onto the urine collection cup.
9. Return the sample to the healthcare worker or place it in the designated area in the bathroom.

-----------------------------------------------------------------------------------

Mid-Stream Clean Catch Instructions

MALE CLEANSING INSTRUCTIONS

1. Cleanse the end of the penis with the first castile soap towelette beginning at the urethral opening and working away from it. (The foreskin of an uncircumcised male must be retracted.)
2. Repeat using a second clean towelette.
3. Urinate the first portion of urine in the toilet.
4. As you continue to urinate, bring the collection cup into the mid-stream to collect the urine sample.
5. Do not touch the inside or lip of the cup.
6. Urinate the remainder of urine into the toilet.
7. Replace the cap onto the urine collection cup.
8. Return the sample to the healthcare worker or place it in the designated area in the bathroom.
Procedimiento de obtención de muestra
de orina para prueba de uroanálisis

Instrucciones para las Mujeres:
1. Párese sobre el inodoro en cuclillas. Separe los pliegues del piel alrededor de la abertura uniraria. Limpie el área alrededor de la abertura con la primera toallita impregnada con jabón de Castilla (castile soap).
2. Limpie el área alrededor de la abertura con la primera toallita impregnada con jabón de Castilla (castile soap)
3. Repita el procedimiento usando una segunda toallita limpia
4. Evacue la primera parte de la orina en el inodoro.
5. Mientras sigue evacuando la orina, coloque el vaso para recolección en mitad de la micción para obtener la muestra de orina.
6. NO toque el interior o el borde del vaso con las manos ni ninguna otra parte del cuerpo.
7. Evacue el resto de la orina en el inodoro.
8. Cierre el vaso tocando solo las superficies exteriores de la tapa y del vaso.
9. Entregue la muestra al personal del laboratorio o en el baño en el compartimento designado para la muestra.

Instrucciones para los Hombres:
1. Limpie la punta del pene con la primera toallita impregnada con jabón de Castilla (castile soap). Comenzando por la abertura uretral y hacia fuera (el prepucio de un hombre no circuncidado se debe retraer primero).
2. Repita el procedimiento usando una segunda toallita limpia
3. Evacue la primera parte de la orina en el inodoro.
4. Mientras sigue evacuando la orina, coloque el vaso para recolección en mitad de la micción para obtener la muestra de orina.
5. NO toque el interior o el borde del vaso con las manos ni ninguna otra parte del cuerpo.
6. Evacue el resto de la orina en el inodoro.
7. Cierre el vaso tocando solo las superficies exteriores de la tapa y del vaso.
8. Entregue la muestra al personal del laboratorio o en el baño en el compartimento designado para la muestra.
PATIENT INSTRUCTIONS FOR 24 HOUR URINE COLLECTION

1. Obtain a container for the specimen collection from the laboratory.
2. Check and make sure the container is labeled with identification (last and first name and date of birth).
3. Patient voids completely and discard the specimen. **Record this start time and date.**
4. For the next 24 hours, collect urine specimens in a separate container and pour them into the large collection container.
5. **Keep refrigerated or on ice.**
6. At 24 hours, void and add this last specimen to the 24-hour collection container. **Record this end time and date.**
7. Record accurately on the container the time and the date the patient started collecting and the time and date of the last collection specimen.
8. If you have any questions, please call the laboratory at (949) 452-3554.

**Note:** 24 HR CREATININE CLEARANCE TEST REQUIRES A BLOOD SPECIMEN UPON DELIVERY OF THE 24 HR URINE. BLOOD SPECIMEN MAY BE DRAWN 24 HRS PRIOR TO OR AFTER THE 24 HR URINE COLLECTION.
4. Urine Collection - Other
   a) Ileal conduit urine
      i. Remove the external urinary appliance, and discard the urine within the appliance.
      ii. Gently swab and clean the stomal opening with a 70% alcohol pad and then with an iodine solution (1 -2% tincture of iodine or a 10% solution of povidone-iodine (1% free iodine). Remove excess tincture of iodine with 70% alcohol after procedure to avoid burn.
      iii. Using sterile technique, insert a double catheter into the stoma. (A double catheter helps to minimize contamination of the specimen with skin flora.)
      iv. Catheterize the ileal conduit to a depth beyond the fascial level.
      v. Collect the urine drained into a sterile container.
   b) Straight catheter urine (in/out catheter urine specimens)
      i. In/out catheter urine specimens are useful when clean-catch urine cannot be obtained or when results from clean-catch urine specimens are equivocal and a diagnosis is critical.
      ii. Prior to catheterization, the patient should force fluids until the bladder is full. (Forcing fluids may reduce organism number.)
      iii. Clean the patient’s urethral opening (and in females, the vaginal vestibule) with soap, and carefully rinse the area with water.
      iv. Using sterile technique, pass a catheter into the bladder.
      v. Collect the initial 15 - 30 ml of urine, and discard it from the mouth of the catheter.
      vi. Collect a sample from the mid or later flow of urine in a sterile container.
   c) Indwelling catheter urine -- Indwelling catheters are placed in patients who are unable to pass urine.
      i. Clean the catheter collection port with a 70% alcohol wipe.
      ii. Using sterile technique, puncture the collection port with a needle attached to a syringe. *(NOTE: Do not collect urine from collection bag.)*
      iii. Aspirate the urine, and place it in a sterile container.
   d) Suprapubic skin aspiration (SPA) of the urinary bladder
      i. SPA is useful in determining urinary infection in adults in whom infection is suspected and for whom results from routine procedures have been equivocal and diagnosis is critical. SPA is also useful in pediatric patients when clean-catch urine specimens are difficult to obtain.
      ii. Before SPA, the patient should force fluids until the bladder is full. (Forcing fluids may reduce the organism number.)
      iii. Shave and disinfect the suprapubic skin overlying the urinary bladder.
      iv. The physician will make a small lance wound through the epidermis, just above the symphysis pubis.
      v. Aspirate urine from the bladder by using a needle aspiration technique.
   e) Bladder washout test (Fairly)
      i. The bladder washout test is useful in determining the site of infection in the urinary tract. Results are equivocal in about 10 - 20% of patients.
      ii. Prior to test, have the patient force liquids until the bladder is full. (Forcing liquids may reduce organism number.)
      iii. Clean the urethral area with soapy water, and rinse the area well with water.
      iv. Insert an indwelling catheter into the bladder through the urethra.
      v. Collect an initial urine specimen into a sterile container, and refrigerate it.
      vi. Empty the bladder through the urethral catheter, and then irrigate it. (Use a sterile nonbacteriostatic 0.85% NaCl solution to irrigate the bladder.)
      vii. Collect three additional specimens (5 - 10 ml each) at 10-minute intervals into separately labeled containers after irrigation of the bladder is performed.
      viii. Submit the initial and timed collection samples to the clinical microbiology laboratory for culture. *(NOTE: It is imperative that each specimen container be clearly labeled with the time of specimen collection.)*
f) Cystoscopy: bilateral ureteral catheterization
   i. Bilateral ureteral catheterization is useful in determining the site of infection in the urinary tract.
   ii. Prior to cystoscopy, have the patient force liquids until the bladder is full. (Forcing liquids may reduce the organism number.)
   iii. Clean the urethral area (and vaginal vestibule in females) with soapy water, and rinse the area well with water. Insert a cystoscope (obturator in place) into the bladder.
   iv. With sterile technique, collect approximately 5-10 ml of urine from open stopcock into a sterile container.
   v. Label this sample CB, for catheterized bladder urine, and refrigerate it. Then irrigate the bladder. (Use sterile nonbacteriostatic 0.85% NaCl to irrigate the bladder.)
   vi. After irrigation of the bladder and insertion of the ureteral catheters, collect irrigating fluid passing from the bladder through the ureteral catheters by holding the ends of both catheters over an opened sterile container.
   vii. Label this sample WB, for washed bladder urine, and refrigerate it.
   viii. Pass the ureteral catheters to each midureter or renal pelvis without introducing additional irrigating fluid. Open both stopcocks of the cystoscope to empty the bladder.
   ix. Discard the first 5-10 ml of urine from each ureteral catheter.
   x. Collect four consecutive paired cultures (5-10 ml each) directly into opened sterile containers.
   xi. Label these specimens LK-1, RK-1, LK-2, and RK-2 (LK for left kidney and RK for right kidney). Submit all samples to the clinical microbiology laboratory for culture.

5. Urine specimen collection considerations are summarized in Table 12.

Table 12: Collection Considerations for Urine Specimens

<table>
<thead>
<tr>
<th>Culture</th>
<th>Vol (ml)$^a$</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bacteria</td>
<td>0.5 - 1.0</td>
<td>Do not collect 24-hour specimen. After proper cleansing of patient, use first morning midstream void.</td>
</tr>
<tr>
<td>Fungi</td>
<td>&gt; 20</td>
<td>Do not collect 24-hour specimen. First morning void is recommended.</td>
</tr>
<tr>
<td>Mycobacteria</td>
<td>&gt; 20</td>
<td>Do not collect 24-hour specimen. First morning three consecutive voided urine specimens are recommended.</td>
</tr>
<tr>
<td>Anaerobes</td>
<td>1</td>
<td>Use suprapubic aspirate. Send in sterile screw-cap cup or tube.</td>
</tr>
<tr>
<td>Virus</td>
<td>10 - 50</td>
<td>Do not collect 24-hour specimen. First morning void is recommended. Useful for adenovirus, mumps, and CMV detection. Send on ice, and transport to laboratory immediately.</td>
</tr>
<tr>
<td>Parasites</td>
<td>24-hour collection</td>
<td>Use for detecting <em>Schistosoma haematobium</em> eggs, <em>Trichomonas vaginalis</em> trophozoites in males, and <em>Onchocerca volvulus</em> microfilariae.</td>
</tr>
</tbody>
</table>

$^a$Amount are guidelines. Greater volumes will increase the chance of organism recovery.

L. Stool Specimens for Occult Blood (Fecal Immunoassay Test)
   1. Inpatient – 5 gm random stool (minimum 1 ml) in a sterile container
   2. Outpatient – Sample device distributed by laboratory. Instructions are included with sample device or as follows:
      a) Package Insert for Personal Use Kit – Read all package insert directions carefully before sample collection. Test results may be invalid if test is not performed properly.
      b) Sample Deposit
         i. Place supplied collection paper inside toilet bowl on top of water.
         ii. Deposit stool sample on top of collection paper.
         iii. Collect sample from stool before paper sinks and stool sample touches water.
         iv. Flush. Collection paper is biodegradable and will not harm septic systems.
c) Sample Collection
   i. Fill in all required information on the sampling bottle.
   ii. Open green cap by twisting and lifting.
   iii. Scrape the surface of the fecal sample with the sample probe.
   iv. Cover the grooved portion of the sample probe completely with stool sample.
   v. Close sampling bottle by inserting the sample probe and twist green cap on tightly. Do not reopen.
   vi. Return the sampling bottle to your doctor or laboratory in envelope provided. Please do not mail to Polymedco.

d) Transport Instructions:
   i. Collect stool per sample collection instructions.
   ii. Remove plastic bag and absorbent pad from envelope.
   iii. Wrap sampling bottle in absorbent pad and insert into plastic bag.
   iv. Insert plastic bag with sampling bottle and absorbent pad into envelope.
   v. Peel tape from flap.
   vi. Fold flap at prefold line.
   vii. Press firmly to seal.
   viii. Return to the laboratory.

M. Bordetella Pertussis (B. pertussis)

1. Bordetella pertussis/parapertussis by PCR – Collect respiratory specimen: Aspirate, broncoalveolar (BAL), swab or sputum. Transfer to a sterile container. It is also acceptable to transfer to ARUP’s lab Viral Transport Media.

2. Bordetella pertussis is part of the Respiratory Panel by PCR. A nasopharyngeal swab in UTM (universal transport medium).

For nursing, refer to collection procedure PC 095 – Nasal Wash/Swab Procedure. Respiratory Therapy is notified to collect outpatient nasopharyngeal washings in the laboratory. (Nasopharyngeal Washings RTD – 221).

N. Glucose Tolerance Testing

When collecting specimens for glucose tolerance tests (GTT’s), it is imperative that all specimens be collected at appropriately timed intervals.

1. Patients referred for tests requiring a single timed collection, such as the 1-hour post 50 gm. glucose load, should be in a fasting state.

2. Please feel free to contact the laboratory at (949) 452-3554 with any questions.

O. Glucose Tolerance Testing (3 Hour)

1. In an effort to standardize oral glucose tolerance testing, the Committee on Statistics of the American Diabetes Association (ADA) has recommended a set of conditions under which the test should be performed.

2. For three days prior to the GTT, a diet containing at least 300 grams per day of carbohydrate is required.
3. Two additional days of this diet are essential if the patient previously has not been on a diet sufficient in carbohydrates. The presence of anorexia or any other condition precluding adequate food intake automatically invalidates the test. Inactivity, such as bed rest, has been reported to reduce glucose tolerance; thus a GTT should not be performed in non-ambulatory patients.
4. During the 12 hours prior to a test, the patient must fast and avoid even black coffee. In addition, smoking and even mild exercise are not permitted.
5. The test should not be performed on those patients who have had an illness during the prior two weeks.
6. For non-pregnant or patient weighing 95 pounds or more, give 75 gm glucose.

**P. OB Glucose Tolerance Testing for gestational diabetes mellitus**

1. Screening 50-gram, 1-hour GTT at 24 - 28 weeks of gestation in women not previously diagnosed with diabetes (non-fasting) of glucose > 140 mg/dL, proceed to GTT when fasting.
2. 2- or 3-hour OB-GTT at 24 - 48 weeks of gestation in women previously not diagnosed with diabetes (overnight fasting of at least 8 hours).
   - 75-gram, 2-hr GTT
   - 100-gram, 3-hr GTT

**Q. Collection of Cytology Specimens (NON-GYN):** refer to policy PL-436, CYTOLOGY SPECIMENS (NON-GYN PROCESSING).

**R. Gynecological Specimen Collection - Thin Prep PAP Smear:** refer to policy PL-440 GYNECOLOGICAL SPECIMEN COLLECTION – THIN PREP PAP SMEAR.

**S. Surgical Pathology Guidelines:** refer to policy PL-435 Surgical Pathology Guidelines.

**T. General Venipuncture Procedure:** refer to policies PL-415, ADULT VENIPUNCTURE AND CAPILLARY PUNCTURE and PL-416, INFANT CAPILLARY AND VENIPUNCTURE.
SPECIMEN LABELING

Specimen Labeling - outpatient

To assure positive identification and optimum integrity of patient specimens from the time of collection until testing is completed and results reported, the staff must label all specimens submitted to SMMC for testing with the patient’s first and last name, correctly spelled, and date of birth (D.O.B.). Samples should be labeled with the date and time of collection and the collector’s initials.

Test request forms for outpatient can be requested by calling SMMC Laboratory at (949) 452-3554.

REQUISITION REQUIREMENTS:

1. Complete a separate test request form for each patient, as outlined here.

2. Be sure to record the following: PLEASE PRINT LEGIBLY.
   • Patient name, birth date, sex, address, phone number
   • COLLECTION TIME AND DATE
   • Order date
   • Physician
   • Type of specimen submitted
   • Source of specimen for infectious disease test(s) and other comments in the “Comments” box
   • Special instructions to the Laboratory (if any)

3. If STAT, be sure to mark the STAT box and enter the FAX and phone numbers of physician.

4. Mark box(es) [x] indicating the test(s) requested. ICD-9 Diagnosis Code MUST be entered.

5. If there is no box for a test you are requesting, print the test name on the blank line.

6. For an example of a requisition (front and back) see following pages.

SPECIMEN LABELING:

1. PRINT patient’s first and last name, legibly and spelled correctly, on the specimen container.

2. PRINT patient’s date of birth (D.O.B.).

3. PRINT date and time collected.

4. For multiple tests, extra tubes or containers may be needed. Call SMMC Laboratory with any questions at (949) 452-3554.
LABORATORY REQUEST FORM (FRONT)

ORDER DATE: ___________ TIME: ___________

DID YOU REMEMBER... TO INCLUDE DIAGNOSIS CODE(S)?

ALL MD ORDERS: FAX TO 949-452-3563

COMPLETE FOR ALL BILLING TYPES

PATIENT NAME (LAST, FIRST, MIDDLE) ____________

DATE OF BIRTH M M D D YEAR: ____________

PATIENT PHONE: ( ) ____________

STREET ADDRESS OF INSURED/RESPONSIBLE PARTY

CITY ____________ STATE ____________ ZIP ____________

ORDERING PHYSICIAN* ____________

STAT ____________ DURING OFFICE HOURS ONLY TO:

PHONE # ____________ FAX # ____________

INSURANCE

FIRST NAME OF INSURED/RESPONSIBLE PARTY (LAST, FIRST, MIDDLE) ____________

RELATIONSHIP TO INSURED: ____________

SPouse ____________ DEPENDENT ____________

DATE OF BIRTH M D YEAR ____________

PATIENT/INSURED/SPouse/CONSERVATOR

DATE ____________

Diagnosis(es) or Signs/Symptoms for each test:

REQUERED

ICD-10 / DIAGNOSIS **

1. ____________
2. ____________
3. ____________

Please provide all diagnosis information.

Saddleback Memorial Laboratory Services Compendium - 42 -
Saddleback Memorial Medical Center
24451 Health Center Drive
Laguna Hills, CA 92653
(949) 452-3554

LABORATORY REQUEST FORM (BACK)

Saddleback Memorial Laboratory Services Compendium

PATHOLOGISTS
Dr. Vivian Mendoza
Dr. Banafsheh Rashidi
Dr. Thomas Hirose
Dr. Michelle Fajardo
Dr. Sonja Matthias

Calif. License No. 206426
Medicare Provider No. M05003
Federal I.D. No. 95-2585792
CLIA No. 05D0578028

ADVANCE BENEFICIARY NOTICE

Medicare will only pay for services that it determines to be medically reasonable and necessary under section 1862(a)(1) of the Medicare Law. If Medicare determines that a particular test, although it would otherwise be covered, "is not reasonable and necessary", under the Medicare Program Standards, Medicare will deny payment.

Tests ordered by your physician which are likely to be denied for payment should be identified by the * symbol. By signing the separate acknowledgement form you are agreeing to be financially responsible for payment.