



# ARIZONA DEPARTMENT OF HEALTH SERVICES

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## PREPAREDNESS

### **Guide to Laboratory Services: Microbiology**

**Arizona Department of Health Services  
Bureau of State Laboratory Services  
250 North 17<sup>th</sup> Avenue  
Phoenix, Arizona 85007  
(602) 542-1188**

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## General Information

Bureau Chief, Laboratory Services	Victor Waddell, Ph.D.
Director, Laboratory Services	Daniel M. Lavine, M.D.
Assistant Bureau Chief	Kathryn Fitzpatrick

Hours of Operation: 8:00 AM to 5:00 PM Monday through Friday (Emergency services available outside business hours when required by public health needs.)

Receiving section only is open from 9:30 AM to 4:30 PM on Saturday.

Annual Holiday Schedule: Laboratory Services observes all state-recognized holidays.

Location: 250 North 17<sup>th</sup> Avenue, Phoenix, Arizona 85007

Telephone Number: (602) 542-1188

WATTS Line: (800) 525-8915

Fax Number: (602) 542-0760

Emergency Phone  
(Weekends/After Hours): (480) 955-0888

## Arizona State Public Health Laboratory Contact Information

Section	Supervisor	Contact Information
Chief, Microbiology	Drew Francis	(602) 542-6146 Drew.Francis@azdhs.gov
Virology / Serology	Alice Chow	(602) 542-6134 <a href="mailto:VirologySerologyLab@azdhs.gov">VirologySerologyLab@azdhs.gov</a>
Bacteriology / (Limited) Parasitology / / Food Testing	Katherine Nordell	(480) 298-3100 katherine.auernheimer@azdhs.gov
Mycobacteriology / Bioemergency Response and Detection for Select Agents	Rachel Wrobel	(480) 594-3780 rachel.schuester@azdhs.gov
Chief, Laboratory Support Services	Gail Weart	(480) 848-4395 gail.weart@azdhs.gov
Analytical Preparation Services	Michael Worsham	(602) 542-1147 <a href="mailto:Michael.Worsham@azdhs.gov">Michael.Worsham@azdhs.gov</a>
Receiving / Shipping	Adrian Fichter	(602) 542-1190 <a href="mailto:labreceiving@azdhs.gov">labreceiving@azdhs.gov</a>

Arizona Department of Health – Bureau of Epidemiology and Disease Control

Office of Infectious Disease Services	Main Number	(602) 364-3676
	After Hours	(480) 303-1191

State Epidemiologist	Ken Komatsu	(602) 364-3587
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## Core Functions and Capabilities of State Public Health Laboratories

State public health laboratories face the broad challenge of working towards prevention and control of disease and improvement of health. To function in this capacity, the public health labs provide testing for, and aid in the diagnosis of, unusual pathogens. The labs serve as the first line of defense in the rapid recognition and prevention of the spread of communicable diseases, while also serving as centers of expertise for the detection and identification of biologic agents of importance in human disease. The public health labs also perform testing to meet the specific program needs of the public health agencies.

### **Routine diagnostic testing for hospitals and private laboratories is provided through independent reference laboratories.**

The policy of the Arizona State Public Health Laboratory (ASPHL) is to provide microbiology and immunology diagnostic support to federal, state, county, and tribal agencies. In addition, the ASPHL serves as a reference microbiology laboratory to hospital and independent clinical laboratories in order to confirm their atypical results from cultures and clinical specimens. This information is also used as part of the Department of Health Services disease surveillance program. Selected diagnostic test procedures are available to private medical practitioners when a procedure is not available through independent reference laboratories or when intense surveillance is deemed necessary. The laboratory also accepts food and water from county and state agencies for outbreak investigations and surveillance. Upon specific request to the State Laboratory, the ordering provider (listed in the Ordering Provider Information section of the Microbiology Sample Submission form) will be given test results in addition to the submitting agency. **Otherwise, test results will be only given to the submitting agency.**

ASPHL reporting requirements can be found at <http://azdhs.gov/labreporting>. This report identifies agents which must be reported to the state and which isolates must be submitted to the laboratory. Please follow packing guidelines found in this manual, or on our website at <http://www.azdhs.gov/lab/>.

### **Sample Submission for Testing**

ASPHL requires all clinical samples to be submitted using either the electronic laboratory submission form or the Electronic Test Orders and Results (ETOR) portal. Please ensure to use the current submission form as samples submitted using the previous versions are subject to rejection.

Clinical microbiology submitters can use the Electronic Test Orders and Results (ETOR) portal. This platform will allow submitters to submit test orders and monitor the progress of the sample throughout the testing process. Notification of completion of testing and results reporting will be given to select users as specified in the portal. To access to this multifunctional portal, to go <https://lwp-web.aimsplatform.com/az/#/auth/login>. Click on the link labeled "New User Registration" to enter their account data and select a password.

The ETOR system sends an email to the submitter acknowledging receipt of the ETOR "New User Registration." Another email is sent when the registration has been completed and the

account is active. Registration should be completed within 72 hours. Food, water, and environmental samples may also be submitted for testing. The appropriate forms may be accessed at <https://www.azdhs.gov/preparedness/state-laboratory/shipping-receiving/index.php#forms-home>. Please do NOT use the clinical electronic submission form for environmental, food, or water testing.

## Collection Kits and Mailing Containers

ASPHL provides specimen collection materials and mailers free of charge. Further information regarding specimen collection materials, mailing containers and *Request for Materials Form* is located in **Section 10: Requesting Collection Kits and Mailing Containers**. Submission forms and supplies for specimen submission are available through the Receiving department at <https://www.azdhs.gov/preparedness/state-laboratory/shipping-receiving/index.php>.

## Courier Services

ASPHL uses Specialized Delivery Services for courier services. The schedule for routine pickup as well as will-call services for shipment and delivery can be found on the ADHS website at <https://www.azdhs.gov/documents/preparedness/state-laboratory/lcip/courier-schedule.pdf>. If your facility is not currently on the route for pickup, please contact Laboratory Support Services at (480) 848-4395.

## Reporting

Results will only be communicated to the submitting agencies and providers listed on the submission/requisition form. Laboratory results may be available to the local jurisdictional health departments. ASPHL has different mechanisms to report results. Reports can be emailed to a point of contact, faxed to a single number, or sent through mail. To have reports emailed or faxed to your facility, please contact Receiving at (602) 542-1190 or Laboratory Support Services at (480) 848-4395.

MyLIMS is a web interface application where submitting agencies can access their results/reports. ASPHL allows a limited number of representatives per site access to this application. To receive access, please contact Laboratory Support Services at (480) 848-4395

## Specimen Rejection Policy

The ASPHL currently has the following policy for rejection of laboratory specimens and/or requested examinations. The ASPHL will NOT examine clinical/reference specimens if the following circumstances exist:

- The test is routinely available at a hospital or a private independent laboratory.
- The identifier on the specimen did not match the identifier on the submission form, or there was no identification on the specimen.
- The quantity of specimen was not sufficient for examination.
- The specimen was too long in transit between the time of collection and receipt in the laboratory.
- The specimen was broken or leaked in transit.
- Clinical/epidemiological information submitted with the specimen was either insufficient or incomplete.
- Specimen was submitted in an improper or expired container, transport media or preservative.
- Blood specimens were hemolyzed or contaminated.
- Only acute blood specimen was submitted, no convalescent specimen (if applicable).
- Material for rabies examination was too decomposed or desiccated to test.
- Reference cultures were mixed or contaminated; only pure cultures are acceptable.
- Specimens were not submitted in individual containers.
- Test request deemed unnecessary by the Bureau of Epidemiology and Disease Control.
- Use of outdated submission forms.

Exceptions to this policy will be considered due to extenuating circumstances, however, final approval to make an exception can only be made by the Laboratory Director, Bureau Chief, Assistant Bureau Chief, or Chief of Microbiology and Molecular Diagnostics.

## Section 1: Sample Submission Guidelines

Submit all samples to the following location:

Arizona Department of Health Services  
State Public Health Laboratory  
250 N. 17<sup>th</sup> Ave  
Phoenix, AZ 85007  
(602) 542-1188

All infectious material must be classified as either Category A (UN2814) or B (UN3373) and must be transported to the Arizona State Public Health Laboratory according to appropriate IATA (International Air Transportation Association), USPS (United States Postal Service) and DOT (U.S. Department of Transportation) regulations. The list of Category A organisms, as outlined by IATA, is provided at the end of this section. All infectious material must be triple-packaged to protect against breakage and/or leakage during transportation. An ASPHL Submission Form must accompany every sample submitted for testing; the forms are available at:

<https://www.azdhs.gov/preparedness/state-laboratory/shipping-receiving/index.php#forms-home>

Category B shipments must follow Packaging Instruction (PI) 650. Category A shipments must follow PI 620 and shipments with dry ice must follow PI 954. All samples and their containers must be identified with the appropriate labels, client and patient information.

Any samples which are leaking and/or not properly identified will be rejected. The following are brief guidelines for properly triple-packaging and shipping specimens for infectious testing at the Arizona State Public Health Laboratory:

- **Primary Container**
  - Must be securely sealed; leak-proof for liquids and sift-proof for solids
    - NOTE: Screw caps and parafilm are recommended.
    - NOTE: Primary **OR** secondary container must be pressure and temperature capable (95 kPa) if air transportation is used.
  - Samples must be properly labeled with patient identifying information.
    - Specimen primary containers for Mycobacterial examination must be labeled with the patient's name, specimen type, date AND time of the collection.
  - For Category A, the maximum quantity for a cargo plane is 4 L or 4 kg. For a passenger plane, the maximum quantity is 50 mL or 50 g.
  - For Category B, the maximum quantity is 500 mL or 500 g.
  - Wrap with absorbent material sufficient for entire contents, and cushioning material.



- **Secondary Container**
  - Securely sealed and watertight/leak-proof
    - NOTE: Primary **OR** secondary container must be pressure and temperature capable (95 kPa) if air transportation is used.
    - NOTE: If you have the appropriate materials you can place multiple primary containers inside a secondary container.
  - A completed itemized list of contents must be placed outside of or surrounding the secondary container.
    - NOTE: An ASPHL Submission Form will satisfy the list of contents requirement.
  - Place absorbent and cushioning material between the primary and secondary containers.
  - Affix a biohazard symbol to the secondary container.
- **Tertiary/Outer Container for CATEGORY B shipments**
  - Outer package must be rigid and of good quality.
  - Affix UN3373 Biological Substance, Category B diamond shaped hazard label.
    - Do NOT affix biohazard symbol to outer package
  - Full name, complete address and phone number of shipper (responsible person).
  - Full name, complete address and phone number of recipient.
- **Tertiary/Outer Container for CATEGORY A shipments**
  - Outer package and inner containers must be UN certified, outer package must contain the UN symbol.
  - For Category A shipments containing infectious material affecting humans, affix **UN2814 Infectious Substances, Affecting Humans** diamond shaped hazard label.
  - For Category A shipments containing infectious material affecting animals, affix **UN2900 Infectious Substances, Affecting Animals** diamond shaped hazard label.
    - NOTE: A list of UN2814 and UN2900 organisms is contained at the end of this section.
    - NOTE: If the infectious material affects both humans and animals, then treat as UN2814.
  - Orientation marks (up arrows) must be present on two (2) sides of outer box.
  - Full name, complete address and 24-hour direct phone number of shipper (responsible person).
  - Full name, complete address and phone number of recipient.
  - NOTE: Make sure to write 6.2 above the UN2814 of the label. Leave the technical name off of the box.
  - NOTE: You must include the full technical name of the suspected unknown or Select Agent on the Dangerous Goods Form placed in with the Secondary Container.
  - For Infectious Substance and Dry Ice Label template examples, refer to the ASPHL Shipping/Receiving website.
    - <https://www.azdhs.gov/preparedness/state-laboratory/index.php#shipping-receiving-shipping>

- **Additional Documentation and Considerations**
  - **Temperature Considerations**
    - Consult appropriate sections within this *Guide to Laboratory Services* document for specific shipping temperatures based on the organism or laboratory section performing test.
    - If wet ice or ice packs are to be used for maintaining refrigerated shipping temperatures, ensure there is sufficient absorbent material contained within to absorb all moisture if ice melts during transit so integrity of box is not compromised.
      - NOTE: It is recommended to place wet ice and/or ice packs inside a zip-lock bag and surround this with absorbent material.
    - If dry ice is to be used to maintain sub-frozen temperatures ensure that the package conforms to PI 954 and that dry ice is not placed inside any tightly sealed container that will prevent the release of carbon dioxide gas during sublimation.
      - NOTE: Dry ice will degrade rapidly therefore it must be purchased, obtained and used as close to actual shipping as possible.
  - **Dangerous Goods Shipper's Declaration**
    - A Shipper's Declaration must accompany all Category A shipments.
      - NOTE: A minimum of 3 color and signed copies is needed.
    - A Shipper's Declaration is not needed for Category B shipments.
    - A Shipper's Declaration is not needed if only shipping dry ice, or dry ice with a Category B shipment.
  - **Select Agent and Toxin Transfers**
    - Shipping of any known Select Agent or Toxin must have prior approval and a completed CDC/APHIS Form 2.
    - Any "suspected" Select Agents must be shipped as either Category A or Category B as designated by its classification.
    - For additional information please visit the Select Agents website at: <http://www.selectagents.gov/form2.html> or contact the Arizona State Public Health Laboratory Bioemergency Response section at (602) 364-0999 for further assistance.
  - **Training**
    - Anyone who packages or ships infectious material must receive appropriate training. There are several "hands-on" and online courses and trainings available. For further information or the next scheduled course please contact the ASPHL Technical Trainer at (602) 542-6175.
  - **Supplies**
    - The Arizona State Public Health Laboratory offers several collection kits and materials for submitting samples. Please see Section 10 for further information.

○ **Regulations and Additional Guidance**

- Arizona State Reporting and Isolate Submission Requirements A.A.C. R9-6-204  
<http://azdhs.gov/labreporting>
- CDC/USDA Select Agent and Toxin list:  
<http://www.selectagents.gov/SelectAgentsandToxinsList.html>  
[https://www.asm.org/ASM/media/Policy-and-Advocacy/Biosafety\\_Sentinel\\_Guideline\\_October\\_2018\\_FINAL.pdf](https://www.asm.org/ASM/media/Policy-and-Advocacy/Biosafety_Sentinel_Guideline_October_2018_FINAL.pdf)
- DOT (Department of Transportation) 49 CFR Part 171-180, Hazardous Materials Regulations: [http://www.ecfr.gov/cgi-bin/text-idx?tpl=/ecfrbrowse/Title49/49cfrv2\\_02.tpl](http://www.ecfr.gov/cgi-bin/text-idx?tpl=/ecfrbrowse/Title49/49cfrv2_02.tpl)
- United States Postal Service (USPS) 39 CFR Part 20, International Postal Service (International Mail Manual), and Part 111, General Information on Postal Service (Domestic Mail Manual). Regulations on transporting infectious substances through the USPS are codified in Section 601.10.17 of the Domestic Mail Manual and Section 135 of the International Mail Manual. A copy of the Domestic and International Mail Manuals may be obtained from <https://about.usps.com/manuals/welcome.htm> or the Government Printing Office by calling Monday through Friday, 7:30 a.m. - 9:00 p.m. EST: (202) 512-1800, (866) 512-1800 (toll free), or from: <https://pe.usps.com/Downloads/DMMIMMQSG>
- Occupational Health and Safety Administration (OSHA). 29 CFR Part 1910.1030, Occupational Exposure to Bloodborne Pathogens: <https://www.osha.gov/laws-regs/regulations/standardnumber/1910/1910.1030>
- Technical Instructions for the Safe Transport of Dangerous Goods by Air (Technical Instructions). International Civil Aviation Organization (ICAO). A copy of these regulations may be obtained from the ICAO Document Sales Unit at (514) 954-8022, Fax: (514) 954-6769, E-Mail: [sales\\_unit@icao.int](mailto:sales_unit@icao.int), or from: <https://www.icao.int/safety/DangerousGoods/Pages/Doc9284-Technical-Instructions.aspx>
- Dangerous Goods Regulations International Air Transport Association (IATA). These regulations are issued by an airline association, are based on the ICAO Technical Instructions, and are followed by most airline carriers. A copy of these regulations can be obtained from: <https://www.iata.org/en/publications/dgr/> or [https://www.who.int/csr/resources/publications/biosafety/WHO\\_HSE\\_EP\\_R\\_2008\\_10.pdf](https://www.who.int/csr/resources/publications/biosafety/WHO_HSE_EP_R_2008_10.pdf)
- Please contact the Arizona State Public Health Laboratory (ASPHL) for appropriate specimen types and shipping instructions for specimen referral to the Centers for Disease Control and Prevention (CDC) for non-routine testing not offered at the ASPHL.

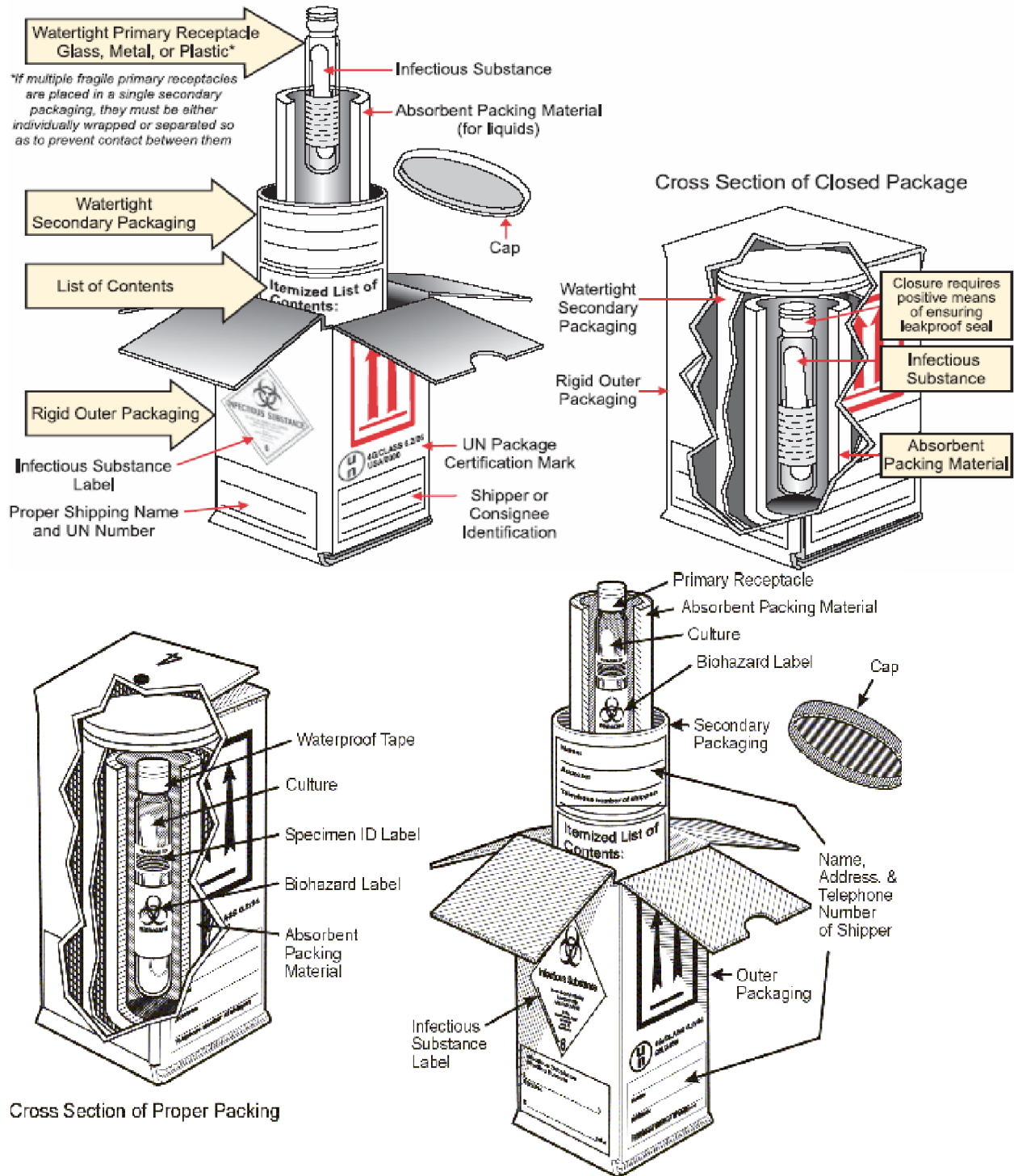
Section 1: Sample Submission Guidelines

The following list is not exhaustive. It is the list of Category A organisms as outlined by the IATA regulations. If there is any doubt as to whether the shipment should be sent as Category A or B, please contact the Arizona State Public Health Laboratory at (602) 542-1190 or (602) 364-0999 for assistance.

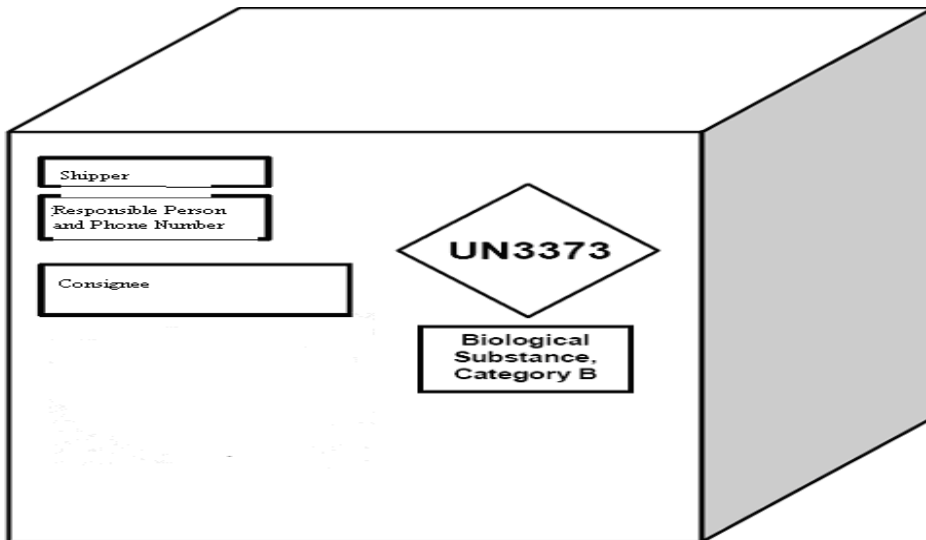
**IATA Category A Organisms**

Ref: IATA Dangerous Goods Regulation 60 <sup>th</sup> Edition, January 2019: 3.6.2.2 Classification of Infectious Substances Table 3.6 D UN number and proper shipping name	Micro-Organism
<p>UN 2814 Infectious substances, affecting humans</p>	<p><i>Bacillus anthracis</i> (cultures only)  <i>Brucella abortus</i> (cultures only)  <i>Brucella melitensis</i> (cultures only)  <i>Brucella suis</i> (cultures only)  <i>Burkholderia mallei</i>-<i>Pseudomonas mallei</i>-<i>Glanders</i> (cultures only)  <i>Burkholderia pseudomallei</i>-<i>Pseudomonas pseudomallei</i> (cultures only)  <i>Chlamydia psittaci</i> (avian) (cultures only)  <i>Clostridium botulinum</i> (cultures only)  <i>Coccidioides immitis</i> (cultures only)  <i>Coxiella burnetii</i> (cultures only)  Crimean-congo hemorrhagic fever virus  Dengue virus (cultures)  Eastern equine encephalitis virus (cultures only)  <i>Escherichia coli</i>, verotoxigenic (cultures only)  Ebola virus  Flexal virus  <i>Francisella tularensis</i> (cultures only)  Guanarito virus  Hantaan virus  Hantavirus causing hemorrhagic fever with renal syndrome  Hendra virus  Hepatitis B virus (cultures only)  Herpes B virus (cultures only)  Human immunodeficiency virus (cultures only)  Highly pathogenic avian influenza virus (culture only)  Japanese Encephalitis virus (culture only)  Junin virus  Kysanur Forest disease virus  Lassa virus  Machupo virus  Marburg virus  Monkeypox virus  <i>Mycobacterium tuberculosis</i> (cultures only)  Nipah virus  Omsk haemorrhagic fever virus  Poliovirus (cultures only)  Rabies virus  <i>Rickettsia prowazekii</i> (cultures only)  <i>Rickettsia rickettii</i> (cultures only)  Rift Valley fever virus (cultures only)  Russian spring-summer encephalitis virus (cultures only)  Sabia virus  <i>Shigella dysenteriae</i> type 1 (cultures only)  Tick-borne encephalitis virus (cultures only)  Variola virus  Venezuelan equine encephalitis virus  West Nile virus (cultures only)  Yellow fever virus (cultures only)  <i>Yersinia pestis</i> (cultures only)</p>
<p>UN 2900 Infectious substances, affecting animals</p>	<p>African swine fever virus (cultures only)  Avian paemyxovirus Type 1 – Velogenic Newcastle disease virus (cultures only)  Classical swine fever virus (cultures only)  Foot-and –mouth disease virus (cultures only)  Goatpox virus (cultures only)  Lumpy skin disease virus (cultures only)  <i>Mycoplasma mycoides</i> – Contagious bovine pleuropneumonia (cultures only)  Peste des petits ruminants virus (cultures only)  Rinderpest virus (cultures only)  Sheep-pox virus (cultures only)  Swine vesicular disease virus (cultures only)  Vesicular stomatitis virus (cultures only)</p>

# Category A Shipping Examples



# Category B Shipping Examples



A completely labeled outer package. The primary container inside the package contains a Biological Substance, Category B substance and is packed according to PI 650.

# Category B Checklist

Sample ID: \_\_\_\_\_  
Packager Name/Initial: \_\_\_\_\_  
Date: \_\_\_\_\_

## **CATEGORY B CHECKLIST**

-UN 3373 Biological Substances, Category B  
-IATA Packing Instruction (PI) 650  
-FedEx, UPS, USPS (Us Mail), private couriers

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### **Primary:**

- Specimen properly labeled with patient ID information
- 50mL or 50g maximum quantity
- Securely sealed & watertight/leakproof (screw cap receptacle and parafilm)  
Note: a Petri dish is not an acceptable primary container
- Wrapped in absorbent material sufficient for entire contents
- Wrapped in cushioning material (bubble wrap)
  - Primary OR secondary container pressure and temperature capable (95kPa)

### **Secondary:**

- Securely sealed and watertight/leakproof
  - Primary OR secondary container pressure and temperature capable (95kPa)
- A completed itemized list of contents (requisition or sample submission form) is placed between the secondary packaging and the outer packaging (NOT inside the secondary packaging)
- Absorbent material is placed between the primary and secondary packaging
- Biohazard symbol on secondary package required if shipping via US Mail (USPS)
- (Optional) Additional cushioning material placed between primary and secondary

### **Outer Package (Rigid):**

- Package is rigid and of good quality (acceptable to reuse Category B packages)
- UN 3373 Biological Substances, Category B diamond shaped label
- Quantity of infectious material is listed
- Quantity of sample –volume (mL) or weight (g)
- Must not contain more than 4 L, 4000mL or 4kg
- Do NOT put biohazard symbol on outer package
- Full name, complete address and phone number of person responsible for the shipment  
(This can be either the shipper or the recipient, but must be someone knowledgeable of the contents)
- Full name, complete address and telephone number of the shipper
- Full name, complete address and telephone number of the consignee/recipient

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NOTE: A Shipper's Declaration is not needed for Category B samples OR if dry ice is used.

If dry ice is used consult dry ice shipping checklist  
If overpack is used consult overpack shipping checklist

# Dry Ice Checklist

Sample ID: \_\_\_\_\_  
Packager Name/Initial: \_\_\_\_\_  
Date: \_\_\_\_\_

## **DRY ICE CHECKLIST**

Combined with Category A, B, or Exempt Shipments  
UN 1845 -Miscellaneous Hazard Class 9  
Packing Instruction (PI) 954 *formerly PI 904*

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- Properly ventilated package  
NOTE: Does not have to be a UN certified package
  - Overpack sticker/label needed for Category A shipments only  
(Category B packages will be packaged differently than A)
  - The net quantity of dry ice used in kg is listed
  - The quantity of the dry ice per package is less than 200kg
  - Irrelevant marks and labels removed from package
  - The UN number "UN 1845" label or sticker
  - Miscellaneous Hazard Class 9 label or sticker
  - Full name, complete address and phone number of person responsible for the shipment  
(This can be either the shipper or the recipient, but must be someone knowledgeable of the contents)
  - Full name, complete address and telephone number of the shipper
  - Full name, complete address and telephone number of the consignee/recipient
- 

## **Shipper's Declaration for Dry Ice**

Shippers Declaration must be completed for shipments containing Category A substances only, dry ice alone does not require a Shipper's Declaration.

- The words "Carbon dioxide, solid" or "Dry ice" is contained on Shipper's Declaration  
Recommended to use "Dry Ice"
- Packing group III
- Packing Instruction 954

Consult Checklist for Shipper's Declaration for complete list



## Section 2: Bacteriology

For testing assistance for organisms/diseases not offered by the ASPHL, please contact the Bacteriology section at (602) 542-6132. ASPHL partners with various public health laboratories for testing support, if necessary. Additional testing resources are available from the Centers for Disease Control and Prevention (CDC), Minnesota Public Health Laboratory (MPHL), Utah Public Health Laboratory (UPHL). Specimens should be forwarded to the CDC from the ASPHL and should not be sent directly from other facilities; please contact the Bacteriology section with any questions or for sample coordination with bacteriology or parasitology related samples.

Organism	Acceptable Specimen Source & Type	Test Method Performed by ASPHL	Comments	Turn Around Time (TAT) Business Days
<i>Bordetella pertussis</i> , <i>Bordetella parapertussis</i> , <i>Bordetella holmesii</i>	Nasopharyngeal swab in Regan-Lowe semi-solid transport medium	PCR	Swabs must be polyester, rayon, or nylon-flocked with a fine flexible wire or plastic shaft. Cotton-tipped or calcium alginate swabs, or swabs with a wooden shaft will be rejected.	7 days
Carbapenem Resistant <i>Enterobacteriales</i> , <i>Pseudomonas aeruginosa</i> , <i>Acinetobacter baumannii complex</i> (CRE, CRPA, CRAB)	Pure, viable isolate on non-selective agar plate or slant	Culture, Carbapenemase Screen, AST, PCR (CRE and CRPA only)	Attach complete susceptibility results that show carbapenem resistance. Carbapenem susceptible or intermediate only isolates will be rejected. <i>Pseudomonas aeruginosa</i> isolates susceptible to <u>both</u> ceftazidime and cefepime will be rejected. Fill out a separate submission form for each isolate submitted for testing.	7-14 days
<i>Clostridium botulinum</i>	Serum, Stool, Food - See Special Instructions for Botulism Testing	No testing performed - specimens are referred to the CDC	Approval for botulism testing must be obtained from the Bureau of Epidemiology and Disease Control prior to submission. Contact the Infectious Disease Section of the Bureau of Epidemiology and Disease Control at (602) 364-3676 (main number)/ (480) 303-1191 (after hours number).	12 weeks
<i>Corynebacterium diphtheriae</i>	Pure, viable isolate on non-selective agar plate or slant  Nasopharyngeal swabs, throat swabs, and/or wound swabs (cotton or polyester tip) in semi-solid transport medium	Culture	Approval for testing must be obtained from the Bureau of Epidemiology and Disease Control prior to submission. Contact the Infectious Disease Section of the Bureau of Epidemiology and Disease Control at (602) 364-3676 (main number)/ (480) 303-1191 (after hours number).  Isolates of <i>Corynebacterium diphtheriae</i> , <i>C. pseudotuberculosis</i> , and/or <i>C. ulcerans</i> will be referred to the CDC for toxin testing.  <b>Clinical specimens received after 24 hours of collection will be rejected.</b>	2-6 days

Section 2: Bacteriology

Organism	Acceptable Specimen Source & Type	Test Method Performed by ASPHL	Comments	Turn Around Time (TAT) Business Days
General Enteric Culture	Raw stool; preserved stool in ParaPak C&S, Modified Cary-Blair or equivalent  Aliquots of preserved stool are acceptable for testing.	Culture, PCR, WGS Serotyping	For optimal specimen recovery, preserved stool should be received by ASPHL within 3 days of collection.  <b>Raw stool received more than 2 hours after collection will be rejected.</b>  Stool samples submitted for General Enteric Culture will be screened for <i>Salmonella</i> , <i>Shigella</i> , <i>Campylobacter</i> , Shiga toxin-producing <i>E. coli</i> , <i>Aeromonas</i> , <i>Plesiomonas</i> , <i>Yersinia</i> , and <i>Vibrio</i> .	7-14 days
<i>Haemophilus influenzae</i>	Pure, viable isolate on chocolate agar plate or slant	Culture	Patients 5 years and younger  Isolates from patients older than 5 years must be approved by the Infectious Disease Section of the Bureau of Epidemiology and Disease Control. Contact at (602) 364-3676 (main number)/ (480) 303-1191 (after hours number).	7-14 days
<i>Legionella</i> spp.	Pure, viable isolate on BCYE or similar agar plate or slant	Culture, PCR		7-14 days
<i>Listeria</i> spp.	Pure, viable isolate on non-selective agar plate or slant	Culture, WGS		7-14 days
<i>Neisseria meningitidis</i>	Pure, viable isolate on chocolate or similar agar plate or slant	Culture	Referred to MDH for Serotyping	7-14 days
<i>Salmonella</i> spp.	Pure, viable isolate on non-selective agar plate or slant  Raw stool; preserved stool in ParaPak C&S, Modified Cary-Blair or equivalent  Aliquots of preserved stool are acceptable for testing.	Culture, WGS Serotyping	For optimal specimen recovery, preserved stool should be received by ASPHL within 3 days of collection.  <b>Raw stool received more than 2 hours after collection will be rejected.</b>	7-14 days

Section 2: Bacteriology

Organism	Acceptable Specimen Source & Type	Test Method Performed by ASPHL	Comments	Turn Around Time (TAT) Business Days
Shiga Toxin-Producing <i>Escherichia coli</i>	Pure, viable isolate on non-selective agar plate or slant  Raw stool; preserved stool in ParaPak C&S, Modified Cary-Blair or equivalent  Aliquots of preserved stool are acceptable for testing.	Culture, PCR, WGS Serotyping	For optimal specimen recovery, preserved stool should be received by ASPHL within 3 days of collection.  <b>Raw stool received more than 2 hours after collection will be rejected.</b>	7-14 days
<i>Vibrio spp.</i>	Pure, viable isolate on non-selective agar plate or slant  Raw stool; preserved stool in ParaPak C&S, Modified Cary-Blair or equivalent  Aliquots of preserved stool are acceptable for testing.	Culture	For optimal specimen recovery, preserved stool should be received by ASPHL within 3 days of collection.  <b>Raw stool received more than 2 hours after collection will be rejected.</b>  Isolates of <i>Vibrio cholerae</i> will be referred to the CDC for toxin testing.	7-14 days
VISA/VRSA	Pure, viable isolate on non-selective agar plate or slant	Culture, AST	Attach complete susceptibility results that show vancomycin intermediate or resistant results.  Isolates of <i>S. aureus</i> that have a vancomycin MIC of 8ug/mL or greater (when tested by ASPHL) will be referred to the CDC for further testing.	5-7 days
<i>Yersinia spp.</i>	Pure, viable isolate on non-selective agar plate or slant  Raw stool; preserved stool in ParaPak C&S, Modified Cary-Blair or equivalent  Aliquots of preserved stool are acceptable for testing.	Culture	For optimal specimen recovery, preserved stool should be received by ASPHL within 3 days of collection.  <b>Raw stool received more than 2 hours after collection will be rejected.</b>	7-14 days

**Shipping Conditions for Isolates:**

Isolates sent to ASPHL should be sealed with parafilm or an equivalent and shipped at room temperature (18-25°C).

### Specimen Collection Guidelines for Clinical Bacteriology Testing:

- **Swabs: Nasopharyngeal, Throat, Wound**
  - Collection Conditions:
    - Nasopharynx - Keep the swab in the nasopharynx for 10 - 30 seconds during the collection process. Collect from the inflamed areas of the nasopharynx, if possible.
    - Throat - If throat membranes are present and can be removed, swab from beneath the membrane.
    - Wound - Gently roll swab over the surface of the wound approximately five times, focusing on areas of pus and/or inflammation.
  - Transport Conditions, Regan-Lowe: Place the swab in Regan-Lowe semi-solid media, cutting the excess length of the swab shaft so that the tube can be closed securely. Do not coil, bend, or otherwise smash the swab into the collection tube - samples submitted in this manner will be rejected. Keep at room temperature (18-25°C) until shipment. Ship at room temperature (18-25°C).
  - Transport Conditions, Other: Refrigerate samples immediately after collection (2-8°C) and keep refrigerated until shipment. Ship in refrigerated (2-8°C) conditions.
- **Stool**
  - Collection Conditions:
    - Collect during the acute stage of infection (usually within 5-7 days of symptom onset), before the administration of antibiotics.
    - Stool should be passed into a clean, dry pan or toilet-mounted container. Do not collect using toilet paper, as toilet paper may be impregnated with barium salts which are inhibitory to some fecal pathogens.
    - Minimum quantity collected should be 1g of formed stool (approximately the size of a walnut) or 5mL of diarrheal stool. If sending an aliquot of preserved stool, please send a minimum of 5mL.
    - Please do not overfill the specimen container, as this may cause the container to leak. Leaking specimens or specimens with visible stool contamination on the outside of the container will be rejected.
    - Keep at room temperature (18-25°C) until shipment.
  - Transport Conditions: Ship at room temperature (18-25°C). If sending an aliquot of preserved stool, please indicate this clearly on the specimen container.

### Special Instructions for Botulism Testing

**IMPORTANT:** Collection and transport conditions are determined by the CDC and are subject to change. Always consult the most recent version of the CDC testing directory (<https://www.cdc.gov/laboratory/specimen-submission/list.html>) before collecting and/or sending samples for botulism testing.

- Specimen Source and Type:
  - Adult patients: Serum, Stool
  - Infant Patients: Stool
  
- Collection Conditions - Serum:
  - Minimum volume for serum is 1mL; at least 5mL serum is preferred
  - Serum samples must be collected before antitoxin treatment
  - Refrigerate samples immediately after collection (2-8°C) and keep refrigerated until shipment.
  
- Collection Conditions - Stool:
  - Minimum quantity for stool is 0.5 - 1g; at least 10g stool is preferred
  - Stool can be collected via enema, using sterile non-bacteriostatic water
  - Refrigerate samples immediately after collection (2-8°C) and keep refrigerated until shipment.
  
- Collection Conditions - Food:
  - Foods should be left in their original containers or placed in sterile unbreakable containers. Empty containers with remnants of foods are also acceptable for testing.
  - Refrigerate samples immediately after collection (2-8°C) and keep refrigerated until shipment.
  
- Transport Conditions:
  - For clinical samples, provide patient name, date of birth, history of present illness, and treatment history, including date of BabyBIG or BAT administration.
  - Neither ASPHL nor the CDC accept shipments on weekends or holidays. **Make sure specimens are received by ASPHL Monday - Thursday** so that they can be forwarded to the CDC immediately.
  - Ship stool specimens in refrigerated (2-8°C) conditions. **IMPORTANT: Stool specimens must be received by the CDC within 2 days of collection.** To prevent specimen rejection due to shipping delays, send stool specimens to ASPHL using a STAT courier (if available) as soon as possible after collection.
  - Ship serum specimens in refrigerated (2-8°C) conditions. Serum samples can be received by the CDC within 20 days of collection date.
  - Package must have proper labeling for biological hazards: UN3373 biological substance, Category B.

### Section 3: Mycobacteriology

For more information about specimen collection and shipping refer to the Specimen Collection & Shipment Guide found at <https://www.azdhs.gov/preparedness/state-laboratory/public-health-microbiology/index.php#tuberculosis>.

<b>Mycobacteria Sample Type</b>	<b>Specimens</b>	<b>Transport Conditions</b>	<b>Test Method Performed by ASPHL</b>	<b>Comments</b>	<b>Turn Around Time (TAT) Business Days</b>
Mycobacterium spp. – Clinical Samples	Acceptable clinical samples include: respiratory specimens, bodily fluids, tissues, urine, stool	Clinical Samples: Refrigerated (2-8°C)  Sample must be received by ASPHL within 7 days of collection.	AFB smear, Culture	Stool specimens will be subjected for microscopic analysis only.	6-8 Weeks
Mycobacterium tuberculosis complex – Direct Detection in Clinical Samples	NALC-NaOH digested/decontaminated respiratory specimens  Requires approval from the Arizona Department of Health Services, TB control section main telephone line at (602) 364-4750	Clinical Samples: Refrigerated (2-8°C)  Sample must be received by ASPHL within 7 days of collection.	NAAT PCR	May be sent to CDC for MDDR testing if Rifampin resistance is detected.	2-3 Days
Mycobacterium tuberculosis complex – Reference Drug Susceptibility Testing	Pure, viable isolate/culture in suitable medium or Growth in MGIT tube	Room temperature (18-25°C)	MGIT 960 DST and Agar Proportion DST methods	May be sent to CDC for additional testing.	3-4 weeks
Mycobacterium tuberculosis complex – Reference Identification & Confirmation of Isolate	Pure, viable isolate/culture in suitable medium or Growth in MGIT tube	Room temperature (18-25°C)	PCR, Culture	None	3-4 weeks
Mycobacterium spp. – Reference Identification & Confirmation of Isolate	Pure, viable isolate/culture in suitable medium or Growth in MGIT tube	Room temperature (18-25°C)	MALDI-TOF, Culture	None	3-4 weeks

### Section 3: Mycobacteriology

#### TB Specimen Collection Guidelines:

**AFB Clinical:** *Mycobacterium tuberculosis* complex, *Mycobacterium* spp.

Acceptable Clinical Sample Types: respiratory specimens (i.e. sputum, induced sputum, bronchial washings, NP swab), bodily fluids (i.e. gastric, pleural, ascitic, CSF), tissues (i.e. biopsy, autopsy, bone), skin lesions (i.e. abscess, wound swab), urine, stool

- The following sample types are acceptable but not optimal for testing, which may negatively impact the recovery of AFB: swabs, stool
- Specimens that have leaked in transit are subject to rejection

Unacceptable Clinical Sample Types: blood, serum, bone marrow

Collection Conditions: Collect from symptomatic individuals only, ideally within two weeks of symptom onset. Refrigerate samples immediately after collection (2-8°C) and keep refrigerated until shipment.

- Respiratory specimens
  - Minimum quantity collected should be of 5mL. Less than 5mL and greater than 1mL may be acceptable but is considered poor volume and may impact recovery of AFB.
  - Sputum samples should be collected in the morning before eating or drinking.
  - Use sterile 50mL conical tubes for sample collection.
- Bodily fluids
  - Minimum quantity collected should be of 5mL. Greater than 1mL and less than 5mL may be acceptable but is considered poor volume and may impact recovery of AFB.
  - Use sterile 50mL conical tubes for sample collection.
  - Indicate if the fluid collected is sterile or non-sterile
- Tissues & Skin lesions
  - Use sterile 50mL conical tubes for sample collection.
  - Indicate if the specimen collected is sterile or non-sterile.
- Stool
  - Minimum quantity for stool is 0.5 - 1g; at least 10g stool is preferred
  - Stool can be collected via enema, using sterile non-bacteriostatic water

Transport Conditions: Ship in refrigerated (2-8°C) conditions. Clinical samples must be received by ASPHL within 7 days of collection; samples received >7 days after collection will be rejected.

**AFB Reference:** *Mycobacterium tuberculosis* complex, *Mycobacterium* spp.

Acceptable Reference Sample Types: Pure, viable isolate/culture in suitable medium (i.e. MGIT broth tube, LJ slant, Middlebrook 7H11, Mitchison 7H11 Selective, Middlebrook 7H10, Middlebrook 7H9 broth, Chocolate agar, Versatrek)

Unacceptable Reference Sample Types: clinical samples

- Stool cultures will be rejected

### Section 3: Mycobacteriology

- Specimens that have leaked in transit are subject to rejection

Transport Conditions: Ship in room temperature (18-25°C) conditions, refrigerated (2-8°C) preferred. Reference cultures and isolates may be greater than 7 days after original specimen collection date.



## Section 4: Limited Parasitology

The ASPHL no longer accepts routine diagnostic samples for testing of parasites. The laboratory offers screening for *Giardia* and *Cryptosporidium* to assist in outbreak investigations with approval from the Bureau of Epidemiology and Disease Control. All other submissions are forwarded to CDC with the approval of the Bureau of Epidemiology and Disease Control.

For testing assistance for organisms/diseases not offered by the ASPHL please refer to the Centers for Disease Control and Prevention (CDC) Test Directory website (<http://www.cdc.gov/laboratory/specimen-submission/list.html>).

Organism	Acceptable Specimen Source & Type	Transport Conditions	Test Method Performed by ASPHL	Comments	Turn Around Time (TAT) Business Days
<i>Giardia</i> & <i>Cryptosporidium</i>	Preserved stool in liquid Cary-Blair	Refrigerated (2-8°C)	PCR	<b>Sample must be received by ASPHL within 4 days of collection.</b>	2-3 Days

### Collection Guidelines for Clinical Parasitology Testing

- Collection Conditions:
  - Collect during the acute stage of infection (usually within 5-7 days of symptom onset), before the administration of antibiotics.
  - Stool should be passed into a clean, dry pan or toilet-mounted container. Do not collect using toilet paper, as toilet paper may be impregnated with barium salts which are inhibitory to some fecal pathogens.
  - Minimum quantity collected should be 1g of formed stool (approximately the size of a walnut) or 5mL of diarrheal stool.
  - Please do not overfill the specimen container, as this may cause the container to leak. Leaking specimens or specimens with visible stool contamination on the outside of the container will be rejected.
  - Refrigerate samples immediately after collection (2-8°C) and keep refrigerated until shipment.
- Transport Conditions: Ship in refrigerated (2-8°C) conditions.

## Section 5: Food Product and Water Sample Testing

### Food Product Samples

#### Collection

At least 200 grams of a solid product or about 100 mL of a liquid.

Collection should be in a sterile whirl-pak plastic bag or sterile urine collection cup. The ASPHL does not provide sterile collection containers for food collection.

#### Shipment of Specimens

All samples must be kept cold (<10 °C) during transit to the laboratory.

See Section 1: Sample Submission Guidelines.

A properly completed *Microbiology Food Analysis Submittal Form* must accompany **each individual sample**. The form may be found at

<http://www.azdhs.gov/documents/preparedness/state-laboratory/public-health-microbiology/food-analysis-form.pdf>

Each sample must be identified by a unique number that corresponds to the identification number written on the submission form. More detailed information regarding how to obtain collection/submission supplies can be found in Section 10: Requesting Collection Kits and Mailing Containers.

#### Reporting and Interpretation of Results

Quality control samples are tested for aerobic plate count, total coliforms, fecal coliforms and *E. coli*.

Pathogen isolation and identification is available for foods implicated in foodborne illness outbreaks. Tests available include, but are not limited to, the following:

- *E. coli* O157:H7 detection and isolation
- *Salmonella* detection and isolation
- *Listeria* spp. detection and isolation
- Staphylococcal enterotoxins detection

Food samples are analyzed according to methods specified in the Bacteriological Analytical Manual (FDA BAM) Microbiology Laboratory Guide (USDA MLG) by methods specified by the Centers for Disease Control and Prevention (CDC), or the Food Emergency Response Network (FERN). When appropriate, rapid analytical test kits are used to screen samples for pathogens to provide quicker test results during food outbreak investigations or emergencies. The rapid test results usually take only 1 to 2 days. However, positive results of these tests are only presumptive and conventional tests need to be done to confirm these results.

Preliminary results are usually available within 48 to 72 hours after processing has begun. Confirmatory test results are usually available within 48 hours to ten days depending on the test organism. Please contact the Bacteriology Section at (602) 542-6130 at any time for updates on

the progress of the testing. Generally, final reports are mailed out 3 to 11 days after initial processing begins.

Interpretation of lab results is the responsibility of the submitter. The laboratory will consult with the submitter, if requested. No legal food standards are available on most products, so care and common sense are needed in the interpretation of lab data. Use your experience and comparisons to evaluate the results.

## **Water Samples**

The laboratory no longer accepts routine water samples for microbiological analysis. Samples are accepted from the county health departments with prior approval from management. The laboratory tests drinking water for the presence of coliforms and *E. coli* in compliance with the Safe Drinking Water Act. In addition, the laboratory tests surface or source waters, wastewater and runoff waters for indicator organisms. Please call Laboratory Support Services at (480) 848-4395 before submitting or shipping water samples for analysis.

### **Collection**

#### **Drinking Water Samples**

Drinking water samples should be collected in sterile four-ounce whirl-pak bags or sterile collection bottles with sodium thiosulfate added to neutralize any chlorine in the water. Aseptically collect water from the sample tap. If using sterile collection bottles fill to the 100 mL line and leave adequate air space. If using the whirl-pak bags, collect 125 mL of water. Be sure to whirl them closed tightly and tie the tabs together securely.

#### **Other Water Samples**

Surface water, source waters, runoff waters, etc., can be aseptically collected in any appropriate size sterile whirl-pak bag or bottle (sodium thiosulfate is not needed); however, at least 125 mL is needed to test.

### **Shipment of Specimens**

#### **Drinking Water Samples**

Drinking water samples must be received and tested within 30 hours of collection. It is recommended that samples arrive the first of the week whenever possible. Samples may be mailed or sent by courier to the Arizona State Public Health Laboratory to arrive the next day. While drinking water samples do not need to be iced during transit, it is recommended when feasible to cool samples. Each sample must be accompanied by a properly completed *Water Microbiology Sample Submission Form* (<https://azdhs.gov/documents/preparedness/state-laboratory/public-health-microbiology/water-microbiological-sample-submission-form.pdf>). Information regarding how to obtain collection/submission supplies can be found in Section 10: Requesting Collection Kits and Mailing Containers.

### Other Water Samples

These waters need to be received in the laboratory within six hours of collection, and must be iced during transit. Since the transit time is so short, it is usually best to send the water samples to the laboratory by courier. A properly completed *Water Microbiological Sample Submission Form* must accompany each sample. More detailed information regarding how to obtain collection/submission supplies can be found in Section 10: Requesting Collection Kits and Mailing Containers. Before submitting these water samples, please call the Analytical Preparation Services section at (602) 542-1147 to arrange for testing.

See Section 1: Sample Submission Guidelines.

### Reporting and Interpretation of Results

#### Drinking Water Samples

Drinking water samples are routinely tested for the presence of total coliforms and *E. coli* using the enzyme substrate coliform test. This method provides results in 18 to 24 hours. This is the EPA approved method SM 9223B.

Results of drinking water coliform tests are usually available within 18 to 24 hours after processing has begun. All positive results are called to the submitter, providing that a telephone number has been supplied. In addition, all compliance positive results and repeat samples are faxed to ADEQ (Arizona Department of Environmental Quality). Leaked in transit and too long in transit samples are also called to the submitter. Final reports will usually be mailed one to two days after initial processing. If the sample is checked as a compliance sample, a copy is sent to the submitter and ADEQ.

Normally, the maximum contaminant level for total coliforms in drinking water is based on the presence or absence of coliform organisms in a 100 mL sample. A single water sample can have 0 coliforms per 100 mL. Other rules apply when more routine samples are collected, as the ADEQ Compliance Department dictates. The number of samples required is based on the population served by a public water system. If a compliance sample is positive, repeat samples need to be collected. Please contact your ADEQ compliance officer to determine the number and location to collect these repeat samples.

Method Name	Units	Holding Time	Matrix	Temp (°C)
Enzyme Substrate Coliform Test (Colilert)	Presence or Absence per 100 mL	30 Hours	Drinking, Well or Ground Water	Ambient
Colilert MPN – Most Probable Number (QuantiTray – MPN)	MPN Index per 100 mL	8 Hours	Surface/Ambient and Wastewater	< 10 °C

- Holding time of 30 hours for drinking water is the time of collection to start of incubation.
- Holding time of 8 hours for surface/ambient and wastewater is time of collection to time of test start.

## Section 6: Serology

The Serology section is responsible for performing diagnostic testing for communicable diseases in support of outbreak investigations, and reference testing for private and public laboratories. The time required to process a serology specimen varies considerably, as indicated by the following table. Detailed information on the collection and submission of laboratory samples for any of the following tests can be obtained in the narrative guidelines that follow.

Organism/ Disease	Specimen Type	Minimum Sample Volume	Test Method	Comments	Turn Around Time (TAT) Business Days
Dengue virus* <sup>1, 4</sup>	Serum	1.0 mL	IgM EIA PCR PRNT	Date of onset needed, test method determined by case history.  PRNT testing performed on presumptive positive or equivocal samples.  Specimens may also be tested for West Nile virus and St. Louis Encephalitis virus.	2-7 days 2-7 days 14-28 days
Chikungunya virus* <sup>4</sup>	Serum	1.0 mL	IgM EIA PCR PRNT	Date of onset needed, test method determined by case history.  PRNT testing performed on presumptive positive or equivocal samples.	2-7 days 2-7 days 14-28 days
Hantavirus* <sup>2</sup>	Serum	1.0 mL	IgM EIA IgG EIA		1-7 days
HIV	Serum	0.5 mL	Ag/Ab EIA HIV-1, HIV-2 Ab Differentiation HIV-1/HIV-2 NAAT	ADHS HIV Prevention Program only.  Referred to Florida Department of Public Health	3-5 days
Measles	Serum	0.5 mL	IgM EIA IgG EIA	IgM and IgG are tested together. Sample will also be tested for Rubella IgM and IgG	3-5 days

Section 6: Serology

Organism/ Disease	Specimen Type	Minimum Sample Volume	Test Method	Comments	Turn Around Time (TAT) Business Days
Mumps	Serum	0.5 mL	IgM EIA IgG EIA		Referred to CDC
Q Fever ( <i>Coxiella</i> ), phase I & phase II	Serum	0.5 mL	IgG IFA		Referred to CDC
<i>Rickettsia</i> spp. Rocky Mountain Spotted Fever (RMSF) Group  Typhus Fever Group Other spp.	Paired Serum  Whole Blood	0.5 mL  2.0 mL	IgG IFA  PCR	Acute and convalescent serum preferred for testing.  Typhus Fever Group and other spp. referred to CDC.	IgG IFA 2-14 days  PCR 2-7 days
Rubella	Serum	0.5 mL	IgM EIA IgG EIA	IgM and IgG are tested together. Sample will also be tested for Measles IgM and IgG.	3-5 days
SARS CoV-2	Serum	0.5 mL	Total Ab EIA		3-5 days
St. Louis Encephalitis (SLE) virus <sup>1</sup>	Serum CSF	0.5 mL 0.5 mL	IgM EIA PRNT	Sample will also be tested for WNV  PRNT testing is performed on samples where the P/N ratio between WNV & SLE is <3x.	3-7 days 14-28 days
West Nile Virus (WNV) <sup>1</sup>	Serum CSF	0.5 mL 0.5 mL	IgM EIA PRNT	Sample will also be tested for SLE  PRNT testing is performed on samples where the P/N ratio between WNV & SLE is <3x.	3-7 days 14-28 days
Zika virus* <sup>1,3</sup>	<i>Serum</i> <sup>3</sup> CSF Whole Blood	2.0 mL 1.0 mL 1.0 mL	PCR		2-7 days

\* Prior notification and approval required

## Section 6: Serology

### Test abbreviations:

- EIA – Enzyme Immunoassay
- IFA – Indirect Fluorescent Antibody
- PCR – Polymerase Chain Reaction
- Ab – Antibody
- Ag – Antigen
- PRNT – Plaque Reduction Neutralization Test

1. Significant cross-reactivity has been observed within the viruses in the Flavivirus group including Dengue, SLE, WNV, and Zika. Confirmatory testing with Plaque Reduction Neutralization Testing (PRNT) will be performed on samples where cross-reactivity is suspected.
2. Specimens submitted for Hantavirus testing are tested for both IgG and IgM antibodies. Demonstration of the presence of IgM antibody is suggestive of recent exposure to Hantavirus (Sin Nombre virus). With prior notice and approval from the Arizona Department of Health Services Office of Infectious Disease Services, the turnaround time for test results can be shortened.
3. For Zika virus testing to proceed, serum submission is a requirement. Other specimen types such as urine, whole blood, CSF, and amniotic fluid can also be submitted and recommended but they must be accompanied by a serum specimen.
4. For Dengue virus and Chikungunya virus testing, patient history including travel, symptoms, and symptom onset is required. The test method utilized for detection of these viruses is dependent on clinical information.

### **Specimen Collection:**

#### **Blood**

Blood specimens should be collected aseptically in an appropriate collection tube and labeled with a patient identifier (e.g., patient name). Follow the manufacturer's instructions for volume to collect for each tube submitted. For pediatric patients, smaller volumes of blood may be collected utilizing pediatric tubes.

- **Serum:** red top, tiger top, gold top vacutainer tubes
- **Whole Blood/plasma:** lavender top vacutainer tube w/ EDTA anticoagulant

Acute blood samples should be drawn as soon as possible after appearance of symptoms. A convalescent sample should be drawn 10 – 14 days after the acute sample.

After collection, the tube may be transported directly to the Arizona State Public Health Laboratory (ASPHL) or the tube may be centrifuged and the serum/plasma poured off into a separate vial. Whole blood specimens for PCR testing should not be centrifuged prior to submission to the ASPHL.

#### **Other**

Other specimens may be sent to the ASPHL for serological testing. These include cerebrospinal fluid (CSF) and amniotic fluid which should be collected in a sterile container.

### **Transportation & Storage:**

Store samples refrigerated and ship on cold packs or wet ice. *Do not freeze whole blood specimens.* The specimen should be transported to the ASPHL as soon as possible. Due to the intense heat observed in the summertime, it is advisable to ship the specimen cold to prevent damage to the specimen during transit.

Samples must be transported with the appropriate paperwork, verifying that the information appearing on the specimen matches that on the submission form. Since the integrity of the sample must be maintained from the time of collection of the sample until testing is completed, **labeling errors will result in rejection of the specimen.**

Laboratory submission forms should be filled out completely with all pertinent demographic information. Successful tracking of positive cases is reliant on complete and accurate information being supplied on these forms, including patient name or identifier, date collected, date of onset of illness, submitter's name and address, and agency code.

For HIV serological testing, specimens are to be submitted with an *HIV Submission Form* only. All other serological specimens should be accompanied with a *Microbiology Submission Form*.

Specimens may be mailed or delivered by courier to the ASPHL. See Section 1: Sample Submission Guidelines.

#### **If sent by courier**

- Blood and blood products sent in vacutainer tubes should first be placed in a primary screw-cap leak proof container (such as a 50 mL plastic conical tube available from the ASPHL) to reduce the risk of shattering while in transit.
- The specimen should then be placed in a secondary container such as a plastic specimen bag with separate compartments for the submission form and specimen.
- All infectious material must be triple packaged and conform to U.S. Department of Transportation (DOT) requirements.
- Pack the specimen and its form in absorbent material to help prevent breakage.

Note: It is acceptable to send more than one specimen together, as long as they are properly secured and identified.

#### **If sent by mail**

- Blood sent in vacutainer tubes should first be placed in a leak proof primary container (such as a 50 mL conical tube available from the ASPHL) to reduce the risk of shattering while in transit.
- All infectious material must be triple-packaged and conform to current shipping regulations. Consult the Domestic Mail Manual published by the US Post Office (USPS) for current USPS requirements, and the Hazardous Material Regulations (HMR) for current DOT requirements.
- Wrap the submission form around the secondary container, and place inside the tertiary container or cardboard mailer. Package the specimen with enough absorbent material for entire contents and to help prevent breakage.



**Note: Do not put the submittal form around the primary container; it must be around the secondary container.**

- Place appropriate biohazard label on the outside of the secondary container before transportation to the ASPHL.

50 mL conical tubes and cardboard mailers are available from the ASPHL Receiving Section via request for materials form available at <http://www.azdhs.gov/preparedness/state-laboratory/shipping-receiving/index.php#forms-supply-order> or by emailing [labreceiving@azdhs.gov](mailto:labreceiving@azdhs.gov). Please submit your orders in advance to ensure prompt service and delivery. See Section 10: Requesting Collection Kits and Mailing Containers.

**Rejection Criteria:**

Samples may be considered unacceptable if they are:

- not properly identified or are improperly identified
- grossly hemolyzed (blood cells are lysed)
- contaminated with bacteria
- lipemic
- leaked in transit or broke in transit
- prior approval was not given
- no convalescent serum received.

The submitter will be notified of all rejected laboratory specimens by telephone and with a laboratory report mailed to the submitting agency confirming the reason for rejection.

## Section 7: Virology

The Virology section is responsible for performing diagnostic, reference, and surveillance testing for viruses. The following table provides the viruses the ASPHL Virology department can identify and the turnaround times to report results. Detailed information on the collection and submission of laboratory samples for any of the following tests can be obtained in the narrative guidelines that follow.

Note: Bacteriological collection swabs and transport medium are not acceptable for virus detection. Swabs should be placed into a liquid media such as viral transport media or universal transport media.

Organism	Specimen	Transport Medium & Volume	Comments	Turn Around Time (TAT) Business Days
Arbovirus Surveillance	Mosquito Pools <i>Culex spp.</i> <i>Aedes spp.</i>	None	Ship frozen	7-14 days
Enterovirus <sup>2</sup>	Respiratory Swabs <sup>1</sup> , Rectal Swab, Viral Isolate  Stool, CSF	Hanks, Viral Transport Media (VTM), Universal Transport Media (UTM), Sterile Saline.  Sterile collection container  Min. volume: 2 mL	Enterovirus D68 is referred to CDC. Must have prior approval.	Referred to CDC
Influenza virus <ul style="list-style-type: none"> <li>● Seasonal</li> <li>● Avian</li> <li>● Novel</li> <li>● Variant</li> <li>● A only</li> </ul>	Respiratory Swabs <sup>1</sup> , Viral Isolates  Respiratory washes/aspirates, Sputum, tissue	Hanks, Viral Transport Media (VTM), Universal Transport Media (UTM), Sterile Saline.  Sterile collection container  Min. volume: 2 mL	Surveillance only specimens– aliquot of sample can be submitted, minimum volume 0.5 mL	PCR: 1-5 days

Section 7: Virology

<b>Organism</b>	<b>Specimen</b>	<b>Transport Medium &amp; Volume</b>	<b>Comments</b>	<b>Turn Around Time (TAT) Business Days</b>
Measles virus	Respiratory Swabs <sup>1</sup> , viral isolates  Respiratory aspirates, urine	Hanks, Viral Transport Media (VTM), Universal Transport Media (UTM), Sterile Saline.  Sterile collection container  Min. volume: 2 mL  10 mL – Urine	Submission of both a throat swab and urine is preferred for testing.	PCR: 1-5 days
Mumps virus	Respiratory Swabs <sup>1</sup> , viral isolates  Respiratory aspirates, urine	Hanks, Viral Transport Media (VTM), Universal Transport Media (UTM), Sterile Saline.  Sterile collection container  Min. volume: 2 mL  10 mL – Urine	Submission of both a buccal swab and urine is preferred for testing.	PCR: 1-5 days
Norovirus PCR	Stool (raw)	Sterile collection container	Do not freeze.	2-10 days
Rabies	Small animal (bat), animal head, brain tissue	None	Refer to Section 8	1-2 days
Respiratory Virus Panel (PCR)	NP Swab	Viral Transport Media (VTM), Universal Transport Media (UTM)  Min. volume: 2 mL	Prior Approval Required	1-5 days

Organism	Specimen	Transport Medium & Volume	Comments	Turn Around Time (TAT) Business Days
SARS-CoV-2	Respiratory Swabs <sup>1</sup>  Respiratory washes/aspirates, Sputum	Hanks, Viral Transport Media (VTM), Universal Transport Media (UTM), Sterile Saline.  Sterile collection container		1-5 days
Zika virus <sup>3</sup>	Amniotic Fluid Urine	Sterile collection container  Min. volume: 2 mL	Testing determined by case history and/or travel history and follows the current guidelines from the CDC	2-7 days

1. Respiratory Swabs: Nasopharyngeal, nasal, throat, buccal
2. Enterovirus D68 requests need prior approval, specimens referred to the CDC.
3. For Zika virus testing, serum must be submitted in addition to other specimen types.

### Collection

In order to optimize the ability of the Virology section to identify viral agents from clinical specimens, it is very important that the specimens be collected, handled, and transported in a manner that minimizes deleterious effects on any viral agents present. In addition, sufficient information should be provided with a submitted specimen to guide the laboratory in the selection of proper inoculation techniques for the suspected viral agents.

### Transport Media

Transport media for viral detection should be a liquid medium free from serum. The following liquid media is approved for use for viral sample collections

- Hank’s Balanced Salt Solution
- Viral Transport Media (VTM)
- Universal Transport Media (UTM)
- Sterile Saline (only when other media is not available)

### Collection Swabs

Recovery of virus is also dependent upon the swab the specimen was collected. Acceptable swabs include:

- Polyester
- Synthetic
- Flocked swabs (FLOQ)
- Dacron
- Nylon

Unacceptable swab types include:

- Calcium alginate swabs
- Cotton tip swabs with wooden shafts

### Nasopharyngeal/Throat

Virus isolation is most successful if respiratory specimens are collected within 3 to 5 days of onset of illness. Swabs from both the throat and nasal passage should be collected. The pharynx is swabbed vigorously with a swab moistened with collection medium and then placed in a transport container containing transport medium. Break off the ends of the applicator sticks leaving the swab tips in the collection medium.

NP swabs are used to collect specimens from the nasal passage. Allow the swabs to remain in the nasal passages for a few seconds to absorb the nasal secretions laden with virus. Place the swabs in the Hanks or VTM and label vial.

Store specimens frozen at -70 °C if they cannot be tested within 72 hours. Transport to the ASPHL on wet ice. Do not freeze at -20 °C.

### Urine

Urine specimens can be submitted for further aid in diagnosis of a viral infection. Generally, Measles, Mumps, and Zika virus can be found in the urine. For measles virus and mumps virus testing, it is recommended that a respiratory specimen and a urine specimen be submitted for a suspect patient.

Collect the specimen as soon as possible after onset of illness. Clean voided specimens (10-20 mL) are collected in sterile containers and transported immediately to the laboratory on wet ice or cold packs.

### Throat Washings

Throat washings should be collected by gargling with Hanks Balanced Salt Solution (HBSS). Collect the specimen in a sterile container. Collect the specimen as soon as possible after onset of illness. Transport specimen immediately to the laboratory on wet ice or cold packs.

### Cerebrospinal Fluid (CSF)

For virus isolation, 3-4 mL of CSF should be collected no later than 7-10 days after onset of illness. Place in a sterile screw capped tube without a collection medium. If delays in transport, store frozen at -70 °C. Transport to the laboratory on wet ice or cold pack.

### Stools

Place three to four grams of stool into a sterile container and transport to the laboratory on wet ice or a cold pack.

Stool specimens collected to test for the presence of Norovirus must be refrigerated (not frozen) as soon as possible after collection.

### Autopsy or Biopsy Specimens

Autopsy specimens should be collected within 24 hours after death. Samples from probable sites of pathology are collected using separate, sterile instruments and separate sterile containers for each specimen. Tissues are transported to the laboratory on wet ice or cold pack. If they cannot be tested within 48 hours, they should be stored frozen at -70 °C.

### Shipment of Specimens

All infectious material must be triple-packaged. Place specimens in screw-cap leak proof primary container. Place the primary container in a leak proof secondary container and wrap submission form around the secondary container. Place the secondary container and submission form in an appropriate tertiary container. Ensure adequate ice or cool packs are used if required. Each specimen must be accompanied with a *Microbiology Submission Form*. Mail, ship or courier specimens to the ASPHL.

See Section 1: Sample Submission Guidelines.

### Reporting and Interpretation of Results

Influenza and COVID-19 diagnostic samples must be shipped at -70 °C if testing cannot be performed within 72 hours. A disclaimer will be added to all samples submitting for influenza testing that states: “Specimens should be shipped at -70 °C if testing cannot be performed within 72 hours of collection. If shipping conditions are not met, a negative test result does not rule out the presence of Influenza/COVID-19 virus.”

## Section 8: Special Virology – Rabies

### Collection

Updated collection information can be found at

<http://www.azdhs.gov/preparedness/epidemiology-disease-control/rabies/index.php>

Prior approval from the local health department and Bureau of Epidemiology and Disease Control Vector-Borne and Zoonotic Disease Section is needed on all submissions. Refer to the link above for information on approval.

To identify animals that are rabid, testing requires samplings of brain tissue. For animals that are extremely small such as bats, mice, rats, and other rodents, the animal should be sent intact. Larger animals but no larger than the size of a dog should have the head of the animal submitted. The head should be severed close to the shoulders allowing sufficient tissue of the throat to remain, to ensure inclusion of salivary glands. For larger animals, such as cows, horses, and horned animals, the brain should be removed by a veterinarian and sent to the laboratory. Arrangements can be made with ADHS for removal of brain tissue at the Arizona Veterinary Diagnostic Laboratory in Tucson.

- **Entire Animal:** Bats, mice, rats, rodents
- **Head only:** animals the size of skunks, foxes, wild cats, domestic cats, domestic dogs, javelina, etc.
- **Brain only:** horses, cows, horned animals

**Please Note: Rodents will be tested only by prior approval from the Vector-Borne and Zoonotic Disease Section of the Bureau of Epidemiology and Disease Control.** Rodents may carry other serious and deadly diseases, such as plague, tularemia, or Hantavirus, and should be handled with extreme caution.

Birds and reptiles *will not* be accepted for examination.

Specimens for rabies examination should be collected immediately after the death of the animal. Decomposed specimens or specimens infested with maggots may not be testable but will be determined by the Virology section.

### Shipment of Specimens

All infectious material including specimens submitted for rabies testing should be triple-packaged. Place the head in a leak proof primary container, and place the primary container in another leak proof secondary container. The secondary container should be placed in a tertiary container that is filled with wet ice or cold packs as necessary. An animal ID should be written on the primary, secondary, or tertiary container to ID each sample submitted. An approved *Rabies Submission Form* should be placed in a separate sealed plastic bag outside of the secondary container, or in a separate plastic bag or envelope taped to the outside of the box. Ship the specimens to the Arizona State Public Health Laboratory. Testing delays may be experienced on specimens that are received frozen.

See Section 1: Sample Submission Guidelines.

**Reporting and Interpretation of Results**

In all cases when exposure of a human is reported by a physician or veterinarian, laboratory examination will be made consisting of microscopic examination of smears prepared from brain material. The results of the microscopic examinations will be available 24 to 48 hours after receipt of the specimen. Positive results will be reported by telephone to the Vector-Borne and Zoonotic Disease Section of the Bureau of Epidemiology and Disease Control.



## **Section 9: Bioemergency Detection and Response (Select Agents)**

Please contact the State Bureau of Epidemiology and Disease Control at (602) 364-3676 (main number) or (480) 303-1191 (after hours number) and the Bioemergency Detection and Response Laboratory at (602) 364-0999 before submitting samples for potential outbreak. In the event that an intentional release of any biological agent is suspected, contact your local county health department, local law enforcement agencies, and the FBI Phoenix field office at (602) 279-5511 to inform them of the incident.

### **Specimen Collection and Shipping**

Refer to the [American Society for Microbiology \(ASM\) Sentinel Laboratory procedures](#) for acceptable specimen type and handling. Sample collection should be consistent with current medical practices for the disease/organism biology.

Environmental specimens should be of sufficient quantity and may consist of food, soil, smooth non-porous surfaces swab/wipe, powders, or packages (Table 1). Prior to submitting environmental samples, contact the Bioemergency Detection and Response testing laboratory section at (602) 364-0999 for submission guidance. For details regarding the collection and submission of powder samples/suspicious unknowns including those associated with a threat, please contact your local law enforcement and refer to the Arizona Department of Health Services Suspicious Substances guidelines:

<http://azdhs.gov/documents/preparedness/emergency-preparedness/response-plans/suspicious-substance-protocol.pdf>

All organisms (in culture form) listed in this section are considered Category A infectious substances and must be shipped accordingly. For information regarding the packaging and shipping of Category A infectious agents please refer to Section 1: Sample Submission Guidelines of this manual and/or the ASM Packaging and Shipping Guidelines for Sentinel Laboratories.

[https://www.asm.org/ASM/media/Policy-and-Advocacy/Biosafety\\_Sentinel\\_Guideline\\_October\\_2018\\_FINAL.pdf](https://www.asm.org/ASM/media/Policy-and-Advocacy/Biosafety_Sentinel_Guideline_October_2018_FINAL.pdf)

Section 9: Bioemergency Detection and Response (Select Agents)

**Table 1 – Environmental specimens**

Specimen Type	Amount	Notes
Smooth non-porous Surface (counter, instrument, etc.)	26 cm <sup>2</sup> (4 in <sup>2</sup> ) – Macrofoam swabs 645 cm <sup>2</sup> (100 in <sup>2</sup> ) – Cellulose sponge-wipe 929 cm <sup>2</sup> (144 in <sup>2</sup> ) – Gauze	Use sterile swab, Cellulose sponge-wipe or gauze. Synthetic fibers, synthetic or metal shafts strongly preferred.
Powder	Up to 5 g	Collect aseptically.
Food	25 – 100 g	If food is not available, submit empty containers. Food must be suspected of being intentionally contaminated with one of the agents listed below.
Isolate	Isolate plate or slant	Send in both plates or tubes.

**Table 2 – Clinical specimens**

**Approval is required for clinical samples.** Approval must be obtained from the Bureau of Epidemiology and Disease Control prior to submission. Contact the Infectious Disease Section of the Bureau of Epidemiology and Disease Control at (602) 364-3676 (main number)/ (480) 303-1191 (after hours number).

Organism	Specimen	Test Method	Comments	Turn Around Time (TAT) Business Days
<i>Bacillus anthracis</i>	<ul style="list-style-type: none"> <li>◦ Recovered isolates from clinical specimens</li> <li>◦ For cutaneous anthrax, sterile swabs are appropriate for collection of vesicular fluid and eschar material.</li> <li>◦ For intestinal or pulmonary anthrax, whole blood, serum, plasma, pleural fluid, transtracheal aspirates, sputum, fresh or frozen tissue, stool or rectal swab can be submitted.</li> </ul>	<p>PCR, Culture</p> <p>BioFire PCR (Blood only)</p>	<p>For optimal specimen recovery, use Dacron swabs for swab site collection</p>	7 days

Section 9: Bioemergency Detection and Response (Select Agents)

<i>Brucella spp.</i>	<ul style="list-style-type: none"> <li>◦ <i>Recovered isolates from clinical specimens</i></li> <li>◦ <i>Acceptable sample types: blood<sup>1</sup>, serum, bone marrow, abscess fluid, spleen and liver biopsies, cerebrospinal fluid (CSF), and joint fluid, spleen and liver biopsies, breast milk.</i></li> </ul>	PCR, Culture	<i>When brucellosis is suspected, multiple blood cultures should be obtained.</i>	14 days
<i>Burkholderia mallei/pseudomallei</i>	<ul style="list-style-type: none"> <li>◦ <i>Recovered isolates from clinical specimens</i></li> <li>◦ <i>Acceptable samples include: whole blood, serum, urine, sputum, abscesses and tissue aspirates.</i></li> </ul>	PCR, Culture	None	14 days
<i>Coxiella burnetii</i>	<ul style="list-style-type: none"> <li>◦ <i>Acceptable samples include: whole blood in EDTA</i></li> </ul>	PCR	None	3 days
<i>Ebola</i>	<ul style="list-style-type: none"> <li>◦ <i>Acceptable samples include: whole blood, serum, plasma, urine<sup>2</sup></i></li> </ul>	PCR  BioFire PCR (Blood only)	<i>Urine should not be the sole specimen tested. If a urine specimen from a patient is tested, it should be tested alongside a blood specimen.</i>	3 days
<i>Francisella tularensis</i>	<ul style="list-style-type: none"> <li>◦ <b><i>Recovered isolates from clinical specimens</i></b></li> <li>◦ <b><i>Acceptable samples include: whole blood, ulcer scrapings/swabs, lymph node biopsies, bronchial/tracheal washings, sputum, and pleural fluid, clinical swabs</i></b></li> <li><i>Autopsy specimens: abscess material, biopsy of lymph node, lung, liver, spleen.</i></li> <li>◦ <b><i>Acceptable animal samples include: aspirates, necropsy specimens: abscess material, biopsy of lymph node, lung, liver, spleen, bone marrow scrapings</i></b></li> </ul>	PCR, Culture  BioFire PCR (Blood and sputum only)	<i>Use polyester NP swabs</i>	7 days

Section 9: Bioemergency Detection and Response (Select Agents)

<p><i>Orthopoxvirus</i></p>	<p>◦ <i>Acceptable samples include:</i> <i>Synthetic swabs of lesion, vesicular material, crust or scab from the roof of a vesicle, or fresh tissue biopsy.</i></p>	<p>PCR</p>	<p><i>Submit dry swabs, biopsy tissue and scabs. A dry, synthetic swab is preferred. Do not use cotton swabs.</i></p>	<p>3 days</p>
<p><i>Yersinia pestis</i></p>	<p>◦ <i>Recovered isolates from clinical specimens</i></p> <p>◦ <i>Acceptable samples include:</i> <i>aspirates, biopsy of affected area, sputum, bronchial wash, transtracheal aspirate, blood, and sputum.</i> <i>Autopsy specimens: abscess material, biopsy of lymph node, lung, liver, spleen, bone marrow scrapings.</i></p> <p>◦ <i>Autopsy specimens include:</i> <i>abscess material or sections of lymph node, lung, liver, spleen.</i></p> <p>◦ <i>Acceptable animal samples include:</i> <i>aspirates, necropsy specimens: abscess material, biopsy of lymph node, lung, liver, spleen, bone marrow scrapings.</i></p>	<p>PCR, Culture</p> <p>BioFire PCR (Blood and sputum only)</p>	<p>None</p>	<p>7 days</p>

For questions regarding sample submission, turnaround times or reporting mechanisms please contact the Bioemergency Detection and Response laboratory section at (602) 364-0999.

When a select agent is identified by ASPHL, the submitting laboratory will be contacted by ASPHL with the information regarding the need to fill out and submit the sections C and D of the APHIS/CDC Form 4A to CDC. A complete list of Select Agents can be found on the following website: <http://www.selectagents.gov/SelectAgentsandToxinsList.html>

## LRN Specimen Collection and Transport Guidelines

### **Bacillus anthracis Specimens**

- Cutaneous - Vesicular (early) stage
  - Unroof vesicle and aspirate fluid or collect with two sterile swabs (dacron).
- Cutaneous - Eschar (late) stage
  - Insert swab (dacron) beneath the edge of the eschar, rotate swab or obtain an aspirate.
  - Transport specimens at room temperature for immediate processing.
- Gastrointestinal
  - Stool (> 5 grams), collect and transport in a leak proof sealed container.
  - Collect blood (late stage of infection) directly into an appropriate blood culture bottle (aerobic and anaerobic).
  - Transport specimens and bottles at room temperature for immediate processing.
- Inhalational
  - Collect expectorated sputum into sterile transport cup or collect during bronchoscopy procedure.
  - Collect blood directly into an appropriate blood culture bottle (aerobic and anaerobic).
  - Collect cerebrospinal fluid (CSF) only if signs of meningitis occur.
  - Transport specimens at room temperature for immediate processing.
- Postmortem tissue
  - Tissue pieces should be collected and kept moist.
  - Transport in sterile container at room temperature within 1 hour of collection.

### **Brucella spp. Specimens**

- Bone marrow or whole blood
  - Collect bone marrow in a sterile container.
  - Collect blood directly into an appropriate blood culture bottle.
  - Transport at room temperature for immediate processing.
  - Do not refrigerate.
- Serum
  - Collect at least 1 ml without anticoagulants for serologic diagnosis.
  - Store at 4°C until testing is performed.
  - Acute specimens are collected as soon as possible after onset of disease.
  - Convalescent-phase should be collected >14 days after the acute specimen.
- Joint or abdominal fluid
  - Collect directly into an appropriate blood culture bottle.
  - Transport bottles at room temperature for immediate processing.
- Spleen, liver abscesses

## Section 9: Bioemergency Detection and Response (Select Agents)

- Tissue pieces (at least the size of a pea) should be collected and kept moist.
- Transport in sterile container at room temperature within 1 hour of collection.

### **Burkholderia mallei and Burkholderia pseudomallei Specimens**

- Bone marrow or whole blood
  - Collect bone marrow in a sterile container.
  - Collect blood directly into an appropriate blood culture bottle.
  - Transport at room temperature for immediate processing.
  - Do not refrigerate.
- Sputum
  - Collect expectorated specimen into sterile transport cup or collect during bronchoscopy procedure.
  - Transport at room temperature up to 2 h.
  - If it is known that material will be transported from 2-24 h after collection, then store and transport at 2-8°C.
  - Saliva is not acceptable.
- Tissue specimens (biopsies, abscess aspirates)
  - Tissue pieces (at least the size of a pea) should be collected and kept moist.
  - Transport in sterile container at room temperature within 1 hour of collection.
- Urine
  - Collect at least 1 ml into a leak-proof container.
  - Transport at room temperature up to 2 h.
  - Refrigerate 2 up to 24 h until culture inoculation.

### **Coxiella Specimen Collection**

- Whole blood
  - Collect blood in EDTA (lavender) or sodium citrate (blue) top tube.
  - Maintain at 4 °C for storage and shipping for PCR or special cultures.
  - Collect specimens prior to antimicrobial therapy, if possible.

### **Franciscella tularensis Specimens**

- Tissue
  - For small tissue samples, add several drops of sterile normal saline into a sterile container to keep the tissue moist.
  - Transport at room temperature for immediate processing.
  - If processing of specimen is delayed beyond 2 hours, keep specimen chilled (2-8°C).
  - Amies transport media is an appropriate transport medium.
- Aspirate (Lymph node or lesion)

## Section 9: Bioemergency Detection and Response (Select Agents)

- Collect in a sterile container.
- Transport at room temperature for immediate processing.
- If processing of specimen is delayed beyond 2 hours, keep specimen chilled (2-8°C).
- Bone marrow
  - Collect in a sterile container.
  - Transport at room temperature for immediate processing.
  - If processing of specimen is delayed beyond 2 hours, keep specimen chilled (2-8°C).
- Blood
  - Collect directly into an appropriate blood culture bottle.
  - Transport bottles at room temperature for immediate processing.
  - Do not refrigerate.
- Serum
  - Collect at least 1 ml without anticoagulant for serologic diagnosis.
  - Store at 4°C until testing is performed.
  - Acute specimens are collected as soon as possible after onset of disease.
  - Convalescent-phase should be collected >14 days after the acute specimen.
- Respiratory specimens
  - Collect in a sterile container.
  - Transport at room temperature for immediate processing.
  - If processing of specimen is delayed beyond 2 hours, keep specimen chilled (2-8°C).

### **Orthopoxvirus Specimens**

- Swab of lesion material
  - Collect 2 swabs from each lesion, preferably from different location on the body or from lesions which differ in appearance.
    - Do not clean the lesion prior to swabbing.
  - Use sterile, dry synthetic swabs (i.e. polyester, nylon, Dacron)
    - Do not use cotton swabs.
  - Vigorously swab the surface of the lesion. It is not necessary to de-roof the lesion. If the lesion ruptures while swabbing, ensure that swab collects lesion fluid.
  - Put each swab into a separate container by either steps:
    - Break or cut-off the end of the swab applicator into a 1.5- or 2-ml screw-capped tubes with o-ring or other sterile leak-proof container.
    - Put the entire swab into a sterile container with a gasket seal.
  - Store refrigerated (2-8°C) or frozen (-20°C or lower) within 1 hour of collection.
- Vesicular material
  - Sanitize the patient's skin with an alcohol wipe and allow skin to dry.
  - Open the top of a vesicle or pustule with a scalpel.

## Section 9: Bioemergency Detection and Response (Select Agents)

- Collect the skin of the vesicle top in a dry, sterile 1.5- to 2-mL screw-capped tube. Label the tube.
- Scrape the base of the vesicle or pustule with the wooden end of an applicator stick or swab and smear the scrapings onto a sterile container.
- Swab the base of the lesion with a synthetic swab, place in screw-capped plastic vial, break off applicator handle, and seal.
- Repeat this procedure for 2 or more lesions.
- Lesion crust (scab) specimens
  - Sanitize the patient's skin with an alcohol wipe and allow skin to dry.
  - Use forceps to remove all or a piece of the crust at least 4mm x 4mm
  - Place each crust into dry, separate sterile screw-capped plastic tubes.
  - Wrap parafilm around the juncture of the cap and vial.
  - Label the tube.
- Biopsy lesions (At least 2 specimens obtained by using a 3.5- or 4-mm punch biopsy kit.)
  - Use sterile technique and appropriate anesthetic.
  - Place 1 sample in formalin for immunohistochemical or histopathologic evaluation and store at room temperature.
  - The second specimen should be placed dry (do not add transport medium) in a sterile 1.5- to 2-mL screw-capped container (do not add transport medium).
  - Refrigerate if shipment occurs within 24 hours; otherwise, the specimen should be frozen.

### ***Yersinia pestis* Specimens**

- Lower respiratory tract
  - Transport specimens in sterile, screw-capped containers at room temperature.
  - If it is known that material will be transported from 2-24 h after collection, then store and transport at 2-8°C.
- Blood
  - Collect directly into an appropriate blood culture bottle.
  - Transport bottles at room temperature for immediate processing.
  - Do not refrigerate.
- Aspirate, tissue or biopsy specimen
  - Submit tissue or aspirate in a sterile container.
    - A swab of tissue is not recommended.
  - For small samples, add 1–2 drops of sterile normal saline to keep the tissue moist.
  - Transport the sample at room temperature for immediate processing.
  - Keep the specimen chilled if processing will be delayed (> 2 h).



## Section 10: Requesting Collection Kits and Mailing Containers

Supplies ordered from the Arizona State Public Health Laboratory (ASPHL) are to be used **ONLY** to submit specimens to the ASPHL. There are two Requests for Materials forms currently in use: a *Newborn Screening Supplies Request Form* and a *Request Form* for all other supplies available from the ASPHL. Supplies can be requested by faxing, emailing or calling the Receiving section. All request forms are available as fillable or printable documents at:

<https://www.azdhs.gov/preparedness/state-laboratory/shipping-receiving/index.php#forms-supply-order>

Arizona Department of Health Services  
 Bureau of State Laboratory Services  
 ATTN: Receiving Section  
 250 North 17<sup>th</sup> Avenue  
 Phoenix, AZ 85007

Fax: (602) 364-0758  
 Phone: (602) 542-1190  
 Email: [labreceiving@azdhs.gov](mailto:labreceiving@azdhs.gov)

Please request materials before they are required as the expected turnaround time per order is **5 business days**. Most materials do have a limited shelf life; therefore, only order what will be used before the expiration date. Please do not use expired kits or any kits in which the medium has changed characteristics. Dispose of the media properly and order replacement supplies. The following table provides information regarding submission forms, kit contents and expiration period of each kit. Submitters may use the Request for Materials form, found at <https://www.azdhs.gov/preparedness/state-laboratory/shipping-receiving/index.php#forms-supply-order>, to order entire kits, as well as individual components.

KIT	CONTENTS	SHELF LIFE
Enteric Kit	Instruction Sheet Baggie Media: Cary Blair Store +20 to +25 °C	1-2 years
Influenza Kit	Instruction Sheet N/P Swab Media: Universal Transport Medium Store +2 to +25 °C	1-2 years

Section 10: Requesting Collection Kits and Mailing Containers

KIT	CONTENTS	SHELF LIFE
Pertussis Kit	Instruction Sheet N/P Swab Media: Regan Lowe. Store +2 to +8 °C	4-6 months
Tuberculosis Kit	Sputum Vial Metal Container Cardboard Mailer Store +20 to +25°C	n/a

All submission forms can be obtained at:

<https://www.azdhs.gov/preparedness/state-laboratory/shipping-receiving/index.php#forms-home>

The ASPHL requests that facilities submitting samples/specimens to the laboratory complete the submission forms electronically. The electronic completion of submission forms benefits everyone through ensuring that all information needed is completed, reducing interruptions to submitters, and improves the overall efficiency of the submission of samples/specimens.