Background: Diphtheria is very rare in the United States, with only two cases of respiratory diphtheria reported since 2003. Diphtheria is caused by Gram-positive *Corynebacterium diphtheriae* bacteria that produce diphtheria toxin (toxigenic strains). Important sites of infection are the respiratory mucosa (respiratory diphtheria) and the skin (cutaneous diphtheria). Rarely, other mucosa — the eye, ear, or genitals — may be infected. Both respiratory and non-respiratory disease caused by diphtheria toxin-producing *Corynebacteria* require public health follow-up. Non-toxigenic infections are much more common than toxigenic infections. Non-toxigenic infections are typically less severe and are not vaccine-preventable, as vaccines target diphtheria toxin rather than the bacteria.

Mode of transmission: Spread by respiratory droplets and/or by contact with discharges from skin lesions.

Incubation period: 2-5 days (range 1-10).

Communicability: In untreated people, organisms can be present in nose, throat, eye, and skin lesion discharges for 2-6 weeks. Chronic carriers may shed organisms for 6 months or more. Effective antibiotic therapy typically terminates shedding within 48 hours.

**Suspected Respiratory Diphtheria**


Alternate Diagnoses: Health departments may receive inquiries about patients with exudative pharyngitis with what appears to be an adherent pharyngeal membrane. Given the rarity of respiratory diphtheria, alternate diagnoses should be considered, especially in patients who have been immunized and who have not recently traveled internationally to diphtheria-endemic countries. More common causes of membranous pharyngitis include: Group A β-hemolytic *Streptococcus*, *Staphylococcus aureus*, *Arcanobacter hemolyticum*, *Candida albicans*, *Borellia vincenti* (Vincent’s angina), *H. influenzae* (acute epiglottitis), Epstein Barr virus, cytomegalovirus, adenovirus, herpes simplex, and Toxoplasma. In addition, some anti-neoplastic agents such as methotrexate may induce a pharyngeal membrane. Use of corticosteroids can cause oral candidiasis.

Symptoms: Respiratory diphtheria usually presents as membranous nasopharyngitis or obstructive laryngotracheitis. Initial symptoms include a sore throat, difficulty swallowing, malaise, and low-grade fever. The hallmark of respiratory diphtheria is the presence of a tough, grayish-white pseudomembrane over the tonsils, the pharynx, or larynx. The pseudomembrane is strongly adherent, and attempts to dislodge it usually result in bleeding. The membrane may progressively extend into the larynx and trachea and cause airway obstruction, which, if left untreated, can be fatal.

Swelling of the cervical lymph nodes and soft-tissue can lead to a “bull-neck” appearance in moderate to severe disease. Absorption of diphtheria toxin from the site of infection can cause systemic complications, including damage to the myocardium, nervous system and kidneys.

To obtain and administer diphtheria antitoxin (DAT) stocked at the SFO or LAX Quarantine Stations:
- During office hours contact the Immunization Branch at 510-620-3737.
- Outside of office hours contact the CDPH Duty Officer.
- Allergic reactions may occur in 5-20% of patients. Instructions for sensitivity testing accompany the antitoxin.
- For more information, please see CDC’s DAT webpage: https://www.cdc.gov/diphtheria/dat.html

Antibiotic treatment for 14 days is also indicated:
- Until the patient can swallow comfortably, administer
  - IM procaine penicillin G (units/dose q 12 hours: 300,000 if weight ≤10 kg, 600,000 units if >10 kg), or
  - IV erythromycin (10 mg/kg/dose, q 6 hours, up to maximum of 500 mg/dose)
- Once swallowing comfortably, may substitute oral doses every 6 hours:
  - erythromycin (10 mg/kg/dose, up to maximum of 500 mg/dose), or
  - penicillin V (250 mg per dose)

Elimination of the organism should be documented 24 hours after completion of treatment by two consecutive negative nose and throat cultures taken 24 hours apart. If cultures remain positive after treatment, a second antibiotic course should be given and follow-up cultures performed.

Because disease does not necessarily confer immunity, DTaP or Tdap should be administered during convalescence per ACIP recommendations for age.

Areas and countries with endemic diphtheria

<table>
<thead>
<tr>
<th>COUNTRIES</th>
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<tbody>
<tr>
<td>Africa</td>
</tr>
<tr>
<td>Americas</td>
</tr>
<tr>
<td>Asia/South Pacific</td>
</tr>
<tr>
<td>Middle East</td>
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<tr>
<td>Eastern Europe</td>
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</table>

SUSPECTED NON-RESPIRATORY DIPHTHERIA

Health departments may also receive inquiries about non-respiratory (usually wound) cultures growing *C. diphtheriae*. These specimens are very unlikely to be toxigenic; however, testing should be performed to rule out toxin production (see pages 3 and 6 of this document). Toxin-producing *C. diphtheriae* in non-respiratory sites may serve as reservoirs for respiratory and non-respiratory diphtheria in susceptible contacts, but are unlikely to cause systemic disease. Diphtheria in non-respiratory sites became reportable in 2019. **Antitoxin in not generally indicated for non-respiratory diphtheria.**

Symptoms:
- Cutaneous diphtheria, is usually mild, typically consisting of sores or shallow ulcers. [See www.nejm.org/doi/full/10.1056/NEJMicm1701825.] CDC is aware of a few recent cases of toxigenic cutaneous diphtheria in persons with a history of travel to endemic countries Antibiotic treatment as above is recommended.

Conjunctival infection may be mild, resembling a viral conjunctivitis, but the more typical infection is that of membranous conjunctivitis. Infiltration of the conjunctival surface leads to extreme edema and erythema of the eyelids with increasing stiffness of the lids. Membrane formation occurs over the conjunctivae, which ultimately becomes necrotic.
LABORATORY TESTING: If respiratory diphtheria is strongly suspected, treatment with antitoxin should occur whether or not laboratory testing has been completed, but the diagnosis should be confirmed with lab testing.

**Specimen collection:** For respiratory diphtheria, swab the underside of the pharyngeal membrane with a cotton or synthetic swab, or submit a portion of the membrane. For cutaneous diphtheria, swab the lesion.

**Clinical lab testing:** \( C. \ diphtheriae \) can be isolated from culture of nasal or throat swabs, membrane tissue, and/or swabs or tissue from other sites. Isolation of \( C. \ diphtheriae \) requires culture media containing tellurite. Clinical lab testing cannot determine whether \( C. \ diphtheriae \) is toxigenic.

**Public health laboratory testing:** Please send samples from all isolates of \( C. \ diphtheriae \) (regardless of suspected or known toxigenicity) to CDPH MDL for potential additional toxigenicity testing at CDC or CDPH.

Place all swabs in transport media such as Amies and ship overnight with ice packs. Dry swabs submitted in silica gel sachets are also acceptable. Store pieces of membrane or other tissue in sterile saline (not formalin) and ship overnight with ice packs. Alert the receiving laboratory to the suspicion of diphtheria so that tellurite-containing culture media is used.

Please see [https://www.cdph.ca.gov/Programs/CID/DCDC/Pages/MDLSpecimenSubmissionInstructions.aspx](https://www.cdph.ca.gov/Programs/CID/DCDC/Pages/MDLSpecimenSubmissionInstructions.aspx) for specimen submission information, or contact the CDPH Microbial Diseases Lab at 510-412-3903.

INFECTION CONTROL FOR CONFIRMED OR SUSPECTED DIPHTHERIA CASES: For respiratory diphtheria, droplet precautions, in addition to standard precautions, are indicated until elimination of the organism is documented 24 hours after the completion of antimicrobial therapy by 2 consecutive negative culture sets of both nose and throat collected 24 hours apart. For cutaneous diphtheria, contact precautions are indicated.

CLOSE CONTACT MANAGEMENT: Contact tracing and post-exposure prophylaxis (PEP) for confirmed toxigenic diphtheria cases is usually limited to household members and other people with a history of direct, habitual close contact (including kissing or sexual contacts), healthcare workers exposed to nasopharyngeal secretions, people sharing utensils or kitchen facilities, and people taking care of children.

Decisions regarding whether to undertake contact tracing and PEP for suspected diphtheria cases while confirmatory lab testing is pending should be made on a case-by-case basis. For patients in whom \( C. \ diphtheriae \) has been detected in a blood or skin culture, if the patient has:
- No exudative pharyngitis, and
- No history of travel to countries endemic for diphtheria or contact with travelers from those countries, public health personnel may defer contact tracing and management, including chemoprophylaxis. Additional reassurance is provided if the patient received one or more doses of diphtheria toxoid (e.g., DTaP, DT, Td, Tdap) in the last 10 years.

**Management of close contacts includes:**
- Surveillance for 7 days for evidence of disease
- Nose and throat cultures for \( C. \ diphtheriae \)
- Immunization, as appropriate (complete primary series if <3 doses; booster if last dose >5 years ago)
- Antibiotics: Options include:

<table>
<thead>
<tr>
<th>Antibiotic</th>
<th>Dosage</th>
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<tbody>
<tr>
<td>Benzathine penicillin G, IM, one dose</td>
<td>&lt;6 years of age 600,000 units</td>
</tr>
<tr>
<td></td>
<td>&gt;6 years of age 1.2 million units</td>
</tr>
<tr>
<td>Erythromycin, oral, 7 -10 days</td>
<td>Children: 10 mg/kg q 6 hours, up to 250 mg/dose</td>
</tr>
<tr>
<td></td>
<td>Adults: 250 mg q 6 hours</td>
</tr>
<tr>
<td>Azithromycin, oral, 7 -10 days</td>
<td>Children: 10-12 mg/kg once daily, up to 500mg/day</td>
</tr>
<tr>
<td></td>
<td>Adults: 500mg once daily</td>
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</tbody>
</table>

Persons who cannot be relied upon to complete a multiday course or who cannot be kept under surveillance should receive benzathine penicillin G and a dose of DTaP or Tdap per their age.
DETECTION OF NONTOXIGENIC C. DIPHTHERIAE

With the advent of technologies such as MALDI-TOF mass spectrometry, C. diphtheriae strains are being detected more frequently. Thus far, in California, recent C. diphtheriae isolates have been determined to be non-toxigenic upon further testing. Nontoxigenic strains of C. diphtheriae may be detected in:

• Healthy individuals
• Pharyngitis (unclear if causal or colonizing)
• Skin and soft tissue infections, including in persons experiencing homelessness; infections may be polymicrobial
• Invasive disease, including endocarditis

All specimens positive for C. diphtheriae should be submitted to CDPH MDL per laboratory testing guidance above.
CSTE DIPHTHERIA CASE DEFINITION (2019)

Clinical Criteria
• Upper respiratory tract illness with an adherent membrane of the nose, pharynx, tonsils, or larynx OR
• Infection of a non-respiratory anatomical site (e.g., skin, wound, conjunctiva, ear, genital mucosa)

Laboratory Criteria
*Confirmatory laboratory evidence:*
• Isolation of *C. diphtheriae* from any site AND
• Confirmation of toxin-production by Elek test or by another validated test capable of confirming toxin production

Supportive laboratory evidence: Histopathologic diagnosis

Epidemiologic linkage requires direct contact with a laboratory-confirmed case of diphtheria.

Case Classifications
*Confirmed:*
• An upper respiratory tract illness with an adherent membrane of the nose, pharynx, tonsils, or larynx AND any of the following:
  o isolation of toxin-producing *Corynebacterium diphtheriae* from the nose or throat OR
  o epidemiologic linkage to a laboratory-confirmed case of diphtheria.

OR
• An infection at a non-respiratory anatomical site (e.g., skin, wound, conjunctiva, ear, genital mucosa) WITH
  o isolation of toxin-producing *C. diphtheriae* from that site

*Suspect:*
• In the absence of a more likely diagnosis, an upper respiratory tract illness with each of the following:
  o an adherent membrane of the nose, pharynx, tonsils, or larynx AND
  o absence of laboratory confirmation AND
  o lack of epidemiologic linkage to a laboratory-confirmed case of diphtheria.

OR
• Histopathologic diagnosis