

## Reportable Diseases and Laboratory Reportable Significant Findings Changes for 2011

As required by Connecticut General Statutes Section 19a-2a and Section 19a-36-A2 of the Public Health Code, the lists of Reportable Diseases and Laboratory Reportable Significant Findings are revised annually by the Department of Public Health (DPH). An advisory committee, consisting of public health officials, clinicians, and laboratorians, contributes to the process. There are 2 additions, 2 deletions and 3 modifications to the lists effective January 1, 2011.

### Changes to the List of Reportable Diseases

#### **Chickenpox-hospitalizations and adult cases**

Chickenpox-related hospitalizations, and chickenpox in adults  $\geq 18$  years of age are deleted from the list of reportable diseases. This surveillance was originally initiated to detect smallpox. Currently, two syndromic surveillance systems monitor for syndromes consistent with smallpox. Chickenpox and chickenpox-related death remain on the List of Reportable Diseases as part of routine surveillance for vaccine-preventable diseases. Chickenpox also remains on the List of Laboratory Reportable Significant Findings. Smallpox remains a Category 1 reportable disease.

#### **Influenza-related hospitalization and deaths**

The reporting of influenza-related hospitalization and influenza-related death has been modified. These reportable findings have been moved from Category 1 diseases to Category 2 diseases to facilitate reporting. In 2009, these findings were added to the list of reportable diseases as Category 1 diseases to monitor the severity of pandemic and seasonal influenza. The 2010-2011 influenza season is expected to be characterized by currently circulating strains included in the vaccines.

#### **HIV infection and pregnancy**

"HIV-1 exposure in infants born 1/1/2001 or later" is modified to "HIV infection and pregnancy" on the list of reportable diseases and the Reportable Disease Confidential Case Report Form PD-23. Reliable data are necessary to determine needed resources for perinatal HIV prevention and to evaluate existing programs. Complete and timely reporting of HIV infection during pregnancy will

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facilitate case management, partner notification and referral, and monitoring adherence to prevention recommendations. DPH staff will conduct follow-up with cases to provide education, referrals, ensure appointments are kept, and conduct partner notification, testing, and referral services. Maternal and infant medical records will be abstracted to obtain standardized information for the perinatal HIV surveillance project funded by the Centers for Disease Control and Prevention (CDC).

### Changes to the List of Laboratory Reportable Significant Findings

#### **Antibiotic susceptibility results for invasive pneumococcal disease (IPD)**

Antibiotic susceptibility results for IPD are deleted from the laboratory significant findings list and the Laboratory Report of Significant Findings Form OL-15C. In 2010, IPD was nationally notifiable regardless of antibiotic susceptibility. This change was made after new breakpoints for defining susceptibility to penicillin were published in 2008. All *S. pneumoniae* isolates from patients with IPD are routinely submitted to the DPH Laboratory and forwarded to the CDC for antibiotic susceptibility testing. The DPH uses the CDC test results to monitor antibiotic resistance as part of the Connecticut Emerging Infections Program.

### Changes to both the Lists of Reportable Diseases and Laboratory Reportable Significant Findings

#### **Dengue**

Dengue is added to both lists under arboviral disease. In 2009, an outbreak in Key West, Florida was identified by the report from New York of a resident diagnosed with dengue infection who traveled to

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## REPORTABLE DISEASES 2011

The Commissioner of the Department of Public Health (DPH) is required to declare an annual list of reportable diseases. Each report (by mail, fax, or telephone) should include the full name and address of the person reporting, attending physician, disease being reported, and full name, address, date of birth, race/ethnicity, sex and occupation of the person affected. The PD-23 can be found on the DPH [website](#). Please see page 4 for a list of persons required to report reportable diseases. The reports should be sent in envelopes marked "CONFIDENTIAL." Changes for 2011 are noted in **bold** and with an asterisk (\*).

**Category 1 Diseases:** Report immediately by telephone on the day of recognition or strong suspicion of disease for those diseases marked with a telephone (☎). Also mail a report within 12 hours.

**Category 2 Diseases:** All other diseases not marked with a telephone are Category 2 diseases. Report by mail within 12 hours of recognition or strong suspicion of disease.

<p>Acquired Immunodeficiency Syndrome (1,2)</p> <p>☎ Anthrax</p> <p>Arboviral disease (California group, <b>Dengue*</b>, EEE, SLE, WNV)</p> <p>Babesiosis</p> <p>☎ Botulism</p> <p>☎ Brucellosis</p> <p>Campylobacteriosis</p> <p>Carbon monoxide poisoning (3)</p> <p>Central-line associated blood stream infections (Do not use this form to report) (4)</p> <p>Chancroid</p> <p>Chickenpox</p> <p>Chickenpox-related death</p> <p>Chlamydia (<i>C. trachomatis</i>) (all sites)</p> <p>☎ Cholera</p> <p><i>Clostridium difficile</i>, community-onset (5)</p> <p>Creutzfeldt-Jakob disease (age &lt; 55 years)</p> <p>Cryptosporidiosis</p> <p>Cyclosporiasis</p> <p>☎ Diphtheria</p> <p>Ehrlichiosis/Anaplasmosis</p> <p><i>Escherichia coli</i> O157:H7 gastroenteritis</p> <p>Gonorrhea</p> <p>Group A Streptococcal disease, invasive (6)</p> <p>Group B Streptococcal disease, invasive (6)</p> <p><i>Haemophilus influenzae</i> disease, invasive all serotypes (6)</p> <p>Hansen's disease (Leprosy)</p> <p>Hemolytic-uremic syndrome</p> <p>Hepatitis A</p> <p>Hepatitis B</p> <ul style="list-style-type: none"> <li>▪ acute infection (2)</li> <li>▪ HBsAg positive pregnant women</li> </ul> <p>Hepatitis C - acute infection (2)</p>	<p>HIV-1 infection in (1)</p> <ul style="list-style-type: none"> <li>▪ persons with active tuberculosis disease</li> <li>▪ persons with a latent tuberculous infection (history or tuberculin skin test <math>\geq 5</math>mm induration by Mantoux technique)</li> <li>▪ persons of any age</li> <li>▪ pregnant women</li> </ul> <p>HPV: biopsy proven CIN 2, CIN 3 or AIS or their equivalent (1)</p> <p><b>Influenza-associated death *</b></p> <p><b>Influenza-associated hospitalization (7) *</b></p> <p>Lead toxicity (blood level <math>\geq 15</math> <math>\mu</math>g/dL)</p> <p>Legionellosis</p> <p>Listeriosis</p> <p>Lyme disease</p> <p>Malaria</p> <p>☎ Measles</p> <p>☎ Meningococcal disease</p> <p>Mercury poisoning</p> <p>Mumps</p> <p>Neonatal herpes (<math>\leq 60</math> days of age)</p> <p>Neonatal bacterial sepsis (8*)</p> <p>Occupational asthma</p> <p>☎ Outbreaks:</p> <ul style="list-style-type: none"> <li>▪ Foodborne (involving <math>\geq 2</math> persons)</li> <li>▪ Institutional</li> <li>▪ Unusual disease or illness (9)</li> </ul> <p>☎ Pertussis</p> <p>☎ Plague</p> <p>Pneumococcal disease, invasive (6)</p> <p>☎ Poliomyelitis</p> <p>☎ Q fever</p> <p>☎ Rabies (human and animal)</p> <p>Reye syndrome</p> <p>Rheumatic fever</p> <p>☎ Ricin poisoning</p>	<p>Rocky Mountain spotted fever</p> <p><b>Rotavirus*</b></p> <p>☎ Rubella (including congenital)</p> <p>Salmonellosis</p> <p>☎ SARS-CoV</p> <p>☎ Septicemia or meningitis with growth of gram positive rods within 32 hours of inoculation</p> <p>Shiga toxin-related disease (gastroenteritis)</p> <p>Shigellosis</p> <p>Silicosis</p> <p>☎ Smallpox</p> <p>☎ Staphylococcal enterotoxin B pulmonary poisoning</p> <p>☎ <i>Staphylococcus aureus</i> disease, reduced or resistant susceptibility to vancomycin (1)</p> <p><i>Staphylococcus aureus</i> methicillin-resistant disease, invasive, community acquired (6,10)</p> <p><i>Staphylococcus epidermidis</i> disease, reduced or resistant susceptibility to vancomycin (1)</p> <p>Syphilis</p> <p>Tetanus</p> <p>Trichinosis</p> <p>☎ Tuberculosis</p> <p>☎ Tularemia</p> <p>Typhoid fever</p> <p>Vaccinia disease</p> <p>☎ Venezuelan equine encephalitis</p> <p><i>Vibrio</i> infection (<i>parahaemolyticus</i>, <i>vulnificus</i>, other)</p> <p>☎ Viral hemorrhagic fever</p> <p>☎ Yellow fever</p>
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### FOOTNOTES:

1. Report only to State.
2. CDC case definition.
3. Includes persons being treated in hyperbaric chambers for suspect CO poisoning.
4. Applies only to licensed hospitals (as defined by CGS. Ch368V). Hospitals report central-line associated blood stream infections associated with designated intensive care unites (ICUs): any pediatric ICU in the hospital (not including neonatal ICU) and the medical ICU, or, if no medical ICU, the medical-surgical ICU. Make reports to the DPH via the National Healthcare Safety Network (NHSN) using NHSN definitions, criteria, and protocols.
5. Community-onset: illness in a person living in the community at the time of illness onset and no known hospitalizations in preceding 3 months; if hospitalized, a positive test taken within 48 hours of admission.
6. Invasive disease: confirmed by isolation from sterile fluid (blood, CSF, pericardial, pleural, peritoneal, joint, or vitreous) bone, internal body sites, or other normally sterile sites. Includes muscle for group A *streptococcus*.
7. Reporting requirements are satisfied by submitting the Hospitalized and Fatal Cases of Influenza—Case Report Form to the DPH (or in Hartford, Middlesex, and New Haven counties to Yale Emerging Infections Program at 203-764-4357); in a manner specified by the DPH.
8. Clinical sepsis and blood or CSF isolate obtained from an infant  $\leq 72$  hours of age.
9. Individual cases of "significant unusual illness" are also reportable.
10. Community-acquired: infection present on admission to hospital, and person has no previous hospitalizations or regular contact with the health-care setting.

**How to report:** The PD-23 is the general disease reporting form and should be used if other specialized forms are not available. Specialized reporting forms from the following programs are available: HIV/AIDS Surveillance (860-509-7900), Sexually Transmitted Disease Program (860-509-7920), Tuberculosis Control Program (860-509-7722), Occupational Health Surveillance Program (860-509-7740), or Epidemiology and Emerging Infections Program for the PD-23 or Hospitalized and Fatal Cases of Influenza—Case Report Form (860-509-7994). The PD-23 can be found on the DPH [website](#) or by writing the Department of Public Health, 410 Capitol Ave., MS#11EPI, P.O. Box 340308, Hartford, CT 06134-0308 (860-509-7994); or by calling the individual program.

**Telephone reports** of Category 1 disease should be made to the local director of health for the town in which the patient resides and to the Epidemiology and Emerging Infections Program (860-509-7994). Tuberculosis cases should be directly reported to the Tuberculosis Control Program (860-509-7722). For the name, address, or telephone number of the local Director of Health for a specific town contact the Office of Local Health Administration (860-509-7660). **For public health emergencies, an epidemiologist can be reached evenings, weekends, and holidays through the DPH emergency number (860-509-8000).**

## LABORATORY REPORTABLE SIGNIFICANT FINDINGS 2011

The director of a clinical laboratory must report laboratory evidence suggestive of reportable diseases. The Laboratory Report of Significant Findings (OL-15C) can be obtained from the Connecticut Department of Public Health (DPH), 410 Capitol Ave., MS#11EPI, P.O. Box 340308, Hartford, CT 06134-0308; telephone: (860-509-7994) or on the DPH [website](#). The OL-15Cs are not substitutes for physician reports; they are supplements to physician reports which allow verification of diagnosis. A listing of possible bioterrorism diseases is highlighted at the end of this list. Changes for 2011 are noted in **bold** and with an asterisk (\*).

### AIDS (report only to the State)

- CD4+ T-lymphocyte counts <200 cells/μL: \_\_\_\_\_ cells/μL
- CD4+ count < 14% of total lymphocytes: \_\_\_\_\_%

### Arboviral infection (replaces "encephalitis")

California group virus (species): \_\_\_\_\_

### Dengue\*

Eastern equine encephalitis virus

St. Louis encephalitis virus

West Nile virus infection

Babesiosis:  IFA  IgM (titer): \_\_\_\_\_  IgG (titer): \_\_\_\_\_  
 Blood smear (1)  PCR  Other: \_\_\_\_\_

Campylobacteriosis (species): \_\_\_\_\_

Carboxyhemoglobin ≥ 9%: \_\_\_\_\_% COHb

### Chancroid

Chickenpox, acute:  IgM  Culture  PCR  
 DFA  Other: \_\_\_\_\_

Chlamydia (*C. trachomatis*) (test type: \_\_\_\_\_)

Creutzfeldt-Jakob disease, age < 55 years (biopsy)

Cryptosporidiosis (method of ID): \_\_\_\_\_

Cyclosporiasis (method of ID): \_\_\_\_\_

### Diphtheria (1)

Ehrlichiosis/Anaplasmosis (2)  *A. phagocytophilum*  *E. chaffeensis*

Unspecified  IFA (titers): IgM \_\_\_\_\_ IgG \_\_\_\_\_

Blood smear  PCR  Other: \_\_\_\_\_

Enterococcal infection, vancomycin-resistant (2,3): \_\_\_\_\_

*Escherichia coli* O157 infection (1)

### Giardiasis

Gonorrhea (test type: \_\_\_\_\_)

Group A streptococcal disease, invasive (3)

Group B streptococcal disease, invasive (3)

*Haemophilus influenzae* disease, invasive, all serotypes (1,3)

Hansen's disease (Leprosy)

Hepatitis A  IgM anti-HAV (1)

Hepatitis B  HBsAg  IgM anti-HBc (1)

Hepatitis C (anti-HCV) Ratio: \_\_\_\_\_  RIBA  PCR (4)

Herpes simplex virus, infant ≤ 60 days of age (specify type)

Culture  PCR  IFA  Ag detection

HIV genotype (electronic file) (report only to State) (5)

HIV Infection (report only to the State) (1,5)

HPV (report only to state): (6)

Biopsy proven  CIN 2  CIN 3  AIS

or their equivalent (specify): \_\_\_\_\_

Influenza:  A  B  Unk.  Subtype: \_\_\_\_\_

RT-PCR  Culture  Rapid test

Lead Poisoning (blood lead ≥10 μg/dL) (7)

Finger Stick: \_\_\_\_\_ μg/dL  Venous: \_\_\_\_\_ μg/dL

### Legionellosis

Culture  DFA  Ag positive

Four-fold serologic change (titers): \_\_\_\_\_

### Listeriosis (1)

Lyme disease (8)

Malaria/blood parasites (1,2): \_\_\_\_\_

Measles (Rubeola) (titer) (9): \_\_\_\_\_

Meningococcal disease, invasive (1,3)

### Mercury poisoning

Urine ≥ 35 μg/g creatinine: \_\_\_\_\_ μg/g

Blood ≥ 15 μg/L: \_\_\_\_\_ μg/L

Mumps (titer): \_\_\_\_\_

Neonatal bacterial sepsis (10\*) spp: \_\_\_\_\_

Pertussis (titer): \_\_\_\_\_

DFA Smear:  Positive  Negative

Culture:  Positive  Negative

### Pneumococcal disease, invasive (1,3) \*

Poliomyelitis

Rabies

Rocky Mountain spotted fever

### Rotavirus\*

Rubella (titer): \_\_\_\_\_

Salmonellosis (1,2) (serogroup/serotype): \_\_\_\_\_

SARS-CoV infection (1)  IgM/IgG

PCR: \_\_\_\_\_ (specimen)  Other: \_\_\_\_\_

Shiga toxin-related disease (1)

Shigellosis (1,2) (serogroup/species): \_\_\_\_\_

*Staphylococcus aureus* infection with MIC to

vancomycin ≥ 4 μg/mL (1)

MIC to vancomycin: \_\_\_\_\_ μg/mL

*Staphylococcus aureus* disease, invasive (3)

methicillin-resistant Date pt. Admitted: \_\_\_\_/\_\_\_\_/\_\_\_\_

*Staphylococcus epidermidis* infection with MIC to vancomycin

≥ 32 μg/mL (1)

MIC to vancomycin: \_\_\_\_\_ μg/mL

Syphilis  RPR (titer): \_\_\_\_\_  FTA (titer): \_\_\_\_\_

VDRL (titer): \_\_\_\_\_  MHA (titer): \_\_\_\_\_

Trichinosis

Tuberculosis (1)

AFB Smear:  Positive  Negative

If positive:  Rare  Few  Numerous

NAAT:  Positive  Negative  Indeterminate

Culture:  *Mycobacterium tuberculosis*

Non-tuberculosis mycobact. (specify: *M.* \_\_\_\_\_)

*Vibrio* infection (1) (species): \_\_\_\_\_

Yellow fever

Yersiniosis (species): \_\_\_\_\_

### Diseases that are possible indicators of bioterrorism

Anthrax (1,11)

Botulism (11)

Brucellosis (1,11)

Glanders (1,11)

*Bacillus* species, non-hemolytic, non-motile, from blood or CSF,

growth within 32 hours of inoculation (1,11)

Melioidosis (1,11)

Plague (1,11)

Q fever (11)

Ricin poisoning (11)

Smallpox (1,11)

Staphylococcal enterotoxin B pulmonary poisoning (11)

Tularemia (11)

Venezuelan equine encephalitis (11)

Viral hemorrhagic fever (11)

1. Send isolate, culture, or slide to the State Laboratory for confirmation. For Shiga-toxin, send positive broth. For positive HIV and IgM anti-HAV, send ≥ 0.5mL residual serum. For positive IgM anti-HBc, send ≥ 0.5mL residual serum within 6 months.
2. Specify species/serogroup.
3. Sterile site isolates: defined as sterile fluids (blood, CSF, pericardial, pleural, peritoneal, joint, or vitreous), bone, internal body site (lymph node, brain, heart, liver, spleen, kidney, pancreas, or ovary), or other normally sterile site; includes muscle for group A *streptococcus*.
4. Report all positive anti-HCV with signal to cutoff ratio, all positive RIBA, but only confirmatory PCR tests.
5. Laboratories conducting HIV genotype tests should report the HIV DNA sequence file electronically. Report all positive HIV antibody and antigen tests, and all viral load tests (including those with no virus detectable).
6. On request from the DPH and if adequate tissue is available, send fixed tissue from the specimen used to diagnose CIN2, 3 or cervical AIS or their equivalent for HPV typing according to instructions from the DPH.
7. Report lead results ≥10μg/dL within 48 hours to the Local Health Director and the DPH; submit ALL lead results at least monthly to the DPH.
8. Only laboratories with automated electronic reporting to the DPH are required to report positive results.
9. Report all IgM titers, but only IgG titers that are considered significant by the laboratory performing the test.
10. Report all bacterial isolates from blood or CSF obtained from an infant ≤72 hours of age.
11. Report by telephone to the DPH, weekdays 860-509-7994; evenings, weekends, and holidays 860-509-8000.

Florida. Surveillance will contribute to national surveillance for dengue infections acquired in the continental U.S. The mosquito species responsible for transmission of dengue are not established in Connecticut.

**Rotavirus**

Rotavirus is added to both lists. Before the advent of rotavirus vaccine, rotavirus caused approximately 600 hospitalizations annually in Connecticut and 1 death. Most of the children were <3 years of age, and 80% were infected by age 5 years. Currently, two rotavirus

vaccines are available, RotaTeq® and Rotarix®. Each protects against approximately 90% of strains. Connecticut has received funding from the CDC for a case-control study to monitor the effectiveness of Rotarix®. This surveillance is part of that initiative.

**Age criterion for neonatal bacterial sepsis**

The age criterion for neonatal bacterial sepsis is modified on both lists to match the national criterion. The revised age requirement for neonatal bacterial sepsis is ≤72 hours of age.

**Persons Required to Report Reportable Diseases**

1. Every health care provider who treats or examines any person who has or is suspected to have a reportable disease shall report the case to the local director of health or other health authority within whose jurisdiction the patient resides and to the Department of Public Health.
2. If the case or suspected case of reportable disease is in a health care facility, the person in charge of such facility shall ensure that reports are made to the local director of health and Department of Public Health. The person in charge shall designate appropriate infection control or record keeping personnel for this purpose.
3. If the case or suspected case of reportable disease is not in a health care facility, and if a health care provider is not in attendance or is not known to have made a report within the appropriate time, such report of reportable diseases shall be made to the local director of health or other health authority within whose jurisdiction the patient lives and the Department of Public Health by:
  - A. the administrator serving a public or private school or day care center attended by any person affected or apparently affected with such disease;
  - B. The person in charge of any camp;
  - C. The master or any other person in charge of any vessel lying within the jurisdiction of the state;
  - D. The master or any other person in charge of any aircraft landing within the jurisdiction of the state;
  - E. The owner or person in charge of any establishment producing, handling, or processing dairy products, other food or non-alcoholic beverages for sale or distribution;
  - F. Morticians and funeral directors.

**Persons Required to Report Laboratory Significant Findings**

The director of a laboratory that receives a primary specimen or sample, which yields a reportable laboratory finding, shall be responsible for reporting such findings within 48 hours to the local director of health of the town in which the affected person normally resides. In the absence of such information, the reports should go to the town from which the specimen originated and to the Department of Public Health.

**IMPORTANT NOTICE**

Reporting forms are available electronically on the DPH website. Persons required to report reportable diseases must use the Reportable Disease Confidential Case Report Form PD-23 to report any diseases found on the current Reportable Diseases List unless there is a specialized reporting form available. The director of a clinical laboratory must report laboratory evidence suggestive of reportable diseases using the Laboratory Report of Significant Findings Form OL-15C or other method specified by the DPH. Reporting forms can be obtained by writing or calling the Connecticut Department of Public Health, 410 Capitol Ave., MS#11EPI, P.O. Box 340308, Hartford, CT 06134-0308; telephone: (860-509-7994), or from the DPH website. Please follow these guidelines when submitting reports:

- Complete all required information (at minimum: full name and address of the person reporting, attending physician, disease/test result being reported, onset of illness date, and full name, address, date of birth, race/ethnicity, sex and occupation of the person affected if known).
- Make 2 copies of the report
  - ▶ Send one copy to the DPH via fax (860-509-7910), or mail to the State of Connecticut, Department of Public Health, 410 Capitol Ave., MS#11EPI, P.O. Box 340308, Hartford, CT 06134-0308. Any mailed documents should have “CONFIDENTIAL” marked on the envelope.
  - ▶ Send a copy of the report to the local health department of the town in which the patient resides.
  - ▶ Keep a copy for the patient’s medical record.

<p>J. Robert Galvin, MD, MPH, MBA Commissioner of Public Health</p> <p>Matthew L. Cartter, MD, MPH State Epidemiologist</p> <p>Lynn Sosa, MD Deputy State Epidemiologist</p>	<p>HIV/AIDS Surveillance 860-509-7900</p> <p>Epidemiology and Emerging Infections 860-509-7994</p> <p>Immunizations 860-509-7929</p> <p>Tuberculosis Control 860-509-7722</p> <p>Sexually Transmitted Diseases (STD) 860-509-7920</p>	<p><b>Connecticut Epidemiologist</b></p> <p>Editor: Matthew L. Cartter, MD, MPH</p> <p>Assistant Editor &amp; Producer: Starr-Hope Ertel</p>
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