CHAPTER 41 - EPIDEMIOLOGY HEALTH

SUBCHAPTER 41A - COMMUNICABLE DISEASE CONTROL

SECTION .0100 - COMMUNICABLE DISEASE CONTROL

10A NCAC 41A .0101 REPORTABLE DISEASES AND CONDITIONS

- (a) The following named diseases and conditions are declared to be dangerous to the public health and are hereby made reportable within the time period specified after the disease or condition is reasonably suspected to exist:
 - (1) acquired immune deficiency syndrome (AIDS) 24 hours;
 - (2) anthrax immediately;
 - (3) botulism immediately;
 - (4) brucellosis 7 days;
 - (5) campylobacter infection 24 hours;
 - (6) Candida auris 24 hours;
 - (7) Carbapenem-Resistant Enterobacteriaceae (CRE) 24 hours;
 - (8) chancroid 24 hours;
 - (9) chikungunya virus infection 24 hours;
 - (10) chlamydial infection (laboratory confirmed) 7 days;
 - (11) cholera 24 hours;
 - (12) Creutzfeldt-Jakob disease 7 days;
 - (13) cryptosporidiosis 24 hours;
 - (14) cyclosporiasis 24 hours;
 - (15) dengue 7 days;
 - (16) diphtheria 24 hours;
 - (17) Escherichia coli, shiga toxin-producing 24 hours;
 - (18) ehrlichiosis 7 days;
 - (19) encephalitis, arboviral 7 days;
 - (20) foodborne disease, including Clostridium perfringens, staphylococcal, Bacillus cereus, and other and unknown causes 24 hours;
 - (21) gonorrhea 24 hours;
 - (22) granuloma inguinale 24 hours;
 - (23) Haemophilus influenzae, invasive disease 24 hours;
 - (24) Hantavirus infection 7 days;
 - (25) Hemolytic-uremic syndrome 24 hours;
 - (26) Hemorrhagic fever virus infection immediately;
 - (27) hepatitis A 24 hours;
 - (28) hepatitis B 24 hours;
 - (29) hepatitis B carriage 7 days;
 - (30) hepatitis C, acute 7 days;
 - (31) human immunodeficiency virus (HIV) infection confirmed 24 hours;
 - influenza virus infection causing death 24 hours;
 - (33) legionellosis 7 days;
 - (34) leprosy 7 days;
 - (35) leptospirosis 7 days;
 - (36) listeriosis 24 hours;
 - (37) Lyme disease 7 days;
 - (38) Lymphogranuloma venereum 7 days;
 - (39) malaria 7 days;
 - (40) measles (rubeola) 24 hours;
 - (41) meningitis, pneumococcal 7 days;
 - (42) meningococcal disease 24 hours;

- (43) Middle East respiratory syndrome (MERS) 24 hours;
- (44) monkeypox 24 hours;
- (45) mumps 7 days;
- (46) nongonococcal urethritis 7 days;
- (47) novel influenza virus infection immediately;
- (48) plague immediately;
- (49) paralytic poliomyelitis 24 hours;
- (50) pelvic inflammatory disease 7 days;
- (51) psittacosis 7 days;
- (52) Q fever 7 days;
- (53) rabies, human 24 hours;
- (54) Rocky Mountain spotted fever 7 days;
- (55) rubella 24 hours;
- (56) rubella congenital syndrome 7 days;
- (57) salmonellosis 24 hours;
- (58) severe acute respiratory syndrome (SARS) 24 hours;
- (59) shigellosis 24 hours;
- (60) smallpox immediately;
- (61) Staphylococcus aureus with reduced susceptibility to vancomycin 24 hours;
- (62) streptococcal infection, Group A, invasive disease 7 days;
- (63) syphilis 24 hours;
- (64) tetanus 7 days;
- (65) toxic shock syndrome 7 days;
- (66) trichinosis 7 days;
- (67) tuberculosis 24 hours:
- (68) tularemia immediately;
- (69) typhoid 24 hours;
- (70) typhoid carriage (Salmonella typhi) 7 days;
- (71) typhus, epidemic (louse-borne) 7 days;
- (72) vaccinia 24 hours;
- (73) vibrio infection (other than cholera) -24 hours;
- (74) whooping cough 24 hours; and
- (75) yellow fever 7 days.
- (b) For purposes of reporting, "confirmed human immunodeficiency virus (HIV) infection" is defined as a positive virus culture, repeatedly reactive EIA antibody test confirmed by western blot or indirect immunofluorescent antibody test, positive nucleic acid detection (NAT) test, or other confirmed testing method approved by the Director of the State Public Health Laboratory conducted on or after February 1, 1990. In selecting additional tests for approval, the Director of the State Public Health Laboratory shall consider whether such tests have been approved by the federal Food and Drug Administration, recommended by the federal Centers for Disease Control and Prevention, and endorsed by the Association of Public Health Laboratories.
- (c) In addition to the laboratory reports for Mycobacterium tuberculosis, Neisseria gonorrhoeae, and syphilis specified in G.S. 130A-139, laboratories shall report using electronic laboratory reporting (ELR), secure telecommunication, or paper reports.
 - (1) Isolation or other specific identification of the following organisms or their products from human clinical specimens:
 - (A) Any hantavirus or hemorrhagic fever virus.
 - (B) Arthropod-borne virus (any type).
 - (C) Bacillus anthracis, the cause of anthrax.
 - (D) Bordetella pertussis, the cause of whooping cough (pertussis).
 - (E) Borrelia burgdorferi, the cause of Lyme disease (confirmed tests).
 - (F) Brucella spp., the causes of brucellosis.
 - (G) Campylobacter spp., the causes of campylobacteriosis.
 - (H) Candida auris.

- (I) Carbapenem-Resistant Enterobacteriaceae (CRE).
- (J) Chlamydia trachomatis, the cause of genital chlamydial infection, conjunctivitis (adult and newborn) and pneumonia of newborns.
- (K) Clostridium botulinum, a cause of botulism.
- (L) Clostridium tetani, the cause of tetanus.
- (M) Corynebacterium diphtheriae, the cause of diphtheria.
- (N) Coxiella burnetii, the cause of Q fever.
- (O) Cryptosporidium parvum, the cause of human cryptosporidiosis.
- (P) Cyclospora cayetanesis, the cause of cyclosporiasis.
- (Q) Ehrlichia spp., the causes of ehrlichiosis.
- (R) Shiga toxin-producing Escherichia coli, a cause of hemorrhagic colitis, hemolytic uremic syndrome, and thrombotic thrombocytopenic purpura.
- (S) Francisella tularensis, the cause of tularemia.
- (T) Hepatitis B virus or any component thereof, such as hepatitis B surface antigen.
- (U) Human Immunodeficiency Virus, the cause of AIDS.
- (V) Legionella spp., the causes of legionellosis.
- (W) Leptospira spp., the causes of leptospirosis.
- (X) Listeria monocytogenes, the cause of listeriosis.
- (Y) Middle East respiratory syndrome virus.
- (Z) Monkeypox.
- (AA) Mycobacterium leprae, the cause of leprosy.
- (BB) Plasmodium falciparum, P. malariae, P. ovale, and P. vivax, the causes of malaria in humans.
- (CC) Poliovirus (any), the cause of poliomyelitis.
- (DD) Rabies virus.
- (EE) Rickettsia rickettsii, the cause of Rocky Mountain spotted fever.
- (FF) Rubella virus.
- (GG) Salmonella spp., the causes of salmonellosis.
- (HH) Shigella spp., the causes of shigellosis.
- (II) Smallpox virus, the cause of smallpox.
- (JJ) Staphylococcus aureus with reduced susceptibility to vanomycin.
- (KK) Trichinella spiralis, the cause of trichinosis.
- (LL) Vaccinia virus.
- (MM) Vibrio spp., the causes of cholera and other vibrioses.
- (NN) Yellow fever virus.
- (OO) Yersinia pestis, the cause of plague.
- (2) Isolation or other specific identification of the following organisms from normally sterile human body sites:
 - (A) Group A Streptococcus pyogenes (group A streptococci).
 - (B) Haemophilus influenzae, serotype b.
 - (C) Neisseria meningitidis, the cause of meningococcal disease.
- (3) Positive serologic test results, as specified, for the following infections:
 - (A) Fourfold or greater changes or equivalent changes in serum antibody titers to:
 - (i) Any arthropod-borne viruses associated with meningitis or encephalitis in a human.
 - (ii) Any hantavirus or hemorrhagic fever virus.
 - (iii) Chlamydia psittaci, the cause of psittacosis.
 - (iv) Coxiella burnetii, the cause of Q fever.
 - (v) Dengue virus.
 - (vi) Ehrlichia spp., the causes of ehrlichiosis.
 - (vii) Measles (rubeola) virus.
 - (viii) Mumps virus.
 - (ix) Rickettsia rickettsii, the cause of Rocky Mountain spotted fever.
 - (x) Rubella virus.
 - (xi) Yellow fever virus.
 - (B) The presence of IgM serum antibodies to:

- (i) Chlamydia psittaci.
- (ii) Hepatitis A virus.
- (iii) Hepatitis B virus core antigen.
- (iv) Rubella virus.
- (v) Rubeola (measles) virus.
- (vi) Yellow fever virus.
- (4) Laboratory results from tests to determine the absolute and relative counts for the T-helper (CD4) subset of lymphocytes and all results from tests to determine HIV viral load.
- (5) Identification of CRE from a clinical specimen associated with either infection or colonization, including all susceptibility results and all phenotypic or molecular test results.
- (d) Laboratories utilizing electronic laboratory reporting (ELR) shall report in addition to those listed under Paragraph (c) of this Rule:
 - (1) All positive laboratory results from tests used to diagnosis chronic Hepatitis C Infection, including the following:
 - (A) Hepatitis C virus antibody tests (including the test specific signal to cut-off (s/c) ratio);
 - (B) Hepatitis C nucleic acid tests;
 - (C) Hepatitis C antigen(s) tests; and
 - (D) Hepatitis C genotypic tests.
 - (2) All HIV genotypic test results, including when available:
 - (A) The entire nucleotide sequence; or
 - (B) The pol region sequence (including all regions: protease (PR)/reverse transcriptase (RT) and integrase (INI) genes, if available).
- (e) For the purposes of reporting, Carbapenem-Resistant Enterobacteriaceae (CRE) are defined as:
 - (1) Enterobacter spp, E.coli or Klebsiella spp positive for a known carbapenemase resistance mechanism or positive on a phenotypic test for carbapenemase production; or
 - (2) Enterobacter spp, E.coli or Klebsiella spp resistant to any carbapenem in the absence of carbapenemase resistance mechanism testing or phenotypic testing for carbapenemase production.

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