Updated Guidance for Evaluation of Severe Respiratory Illness Associated with Middle East Respiratory Syndrome Coronavirus (MERS-CoV)

August 13, 2013

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Health Update: Updated Guidance for Evaluation of Severe Respiratory Illness Associated with Middle East Respiratory Syndrome Coronavirus (MERS-CoV)

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SUBJECT: Updated Guidance for Evaluation of Severe Respiratory Illness Associated with Middle East Respiratory Syndrome Coronavirus (MERS-CoV)

On June 10, 2013, the Missouri Department of Health and Senior Services (DHSS) issued a Health Advisory entitled “Updated Guidelines for Evaluation of Severe Respiratory Illness Associated with MERS-CoV.” On August 12, 2013, the Centers for Disease Control and Prevention (CDC) provided updated guidance on who should be tested for MERS-CoV infection, as well as information on changes to CDC’s “probable case” definition, and on what specimens should be obtained when testing for MERS-CoV. This DHSS Health Update contains the updated guidance and information from CDC, and also provides instructions for submitting clinical specimens to the Missouri State Public Health Laboratory (MSPHL) for MERS-CoV testing.

The following is taken from the CDC Health Update entitled “Notice to Healthcare Providers and Public Health Officials: Updated Guidance for the Evaluation of Severe Respiratory Illness Associated with Middle East Respiratory Syndrome Coronavirus (MERS-CoV),” distributed by the CDC Health Alert Network, August 12, 2013.

**Summary**

The Centers for Disease Control and Prevention (CDC) continues to work closely with the World Health Organization (WHO) and other partners to better understand the public health risks posed by Middle East Respiratory Syndrome Coronavirus (MERS-CoV). To date, no cases have been reported in the United States. The purpose of this health update is 1) to provide updated guidance to healthcare providers and state and local health departments regarding who should be tested for MERS-CoV infection, 2) to make them aware of changes to CDC’s “probable case” definition, and 3) to clarify what specimens should be obtained when testing for MERS-CoV. Please disseminate this information to infectious disease specialists, intensive care physicians, primary care physicians, and infection preventionists, as well as to emergency departments and microbiology laboratories.

**Background**

MERS-CoV, formerly called novel coronavirus, is a beta coronavirus that was first described in September 2012. As of August 12, 2013, 94 laboratory-confirmed cases have been reported to WHO. Of those cases, 49% (46) were fatal. All diagnosed cases were among people who resided in or traveled from four countries (Kingdom of Saudi Arabia, United Arab Emirates, Qatar, or Jordan) within 14 days of their symptom onset, or who had close contact with people who resided in or traveled from those countries. Cases with a history of travel from these countries or contact with travelers from these countries have been identified in residents of France, the United Kingdom, Tunisia, and Italy. To date, no cases have been reported in the United States. The most up-to-date details about the number of MERS-CoV cases and deaths by country of residence are on CDC’s MERS website at http://www.cdc.gov/coronavirus/mers/index.html.
Updates to Interim Guidance and Case Definitions

Interim Guidance for Health Professionals: Patients in the U.S. Who Should Be Evaluated

CDC has changed its criteria for who should be evaluated for MERS-CoV. In the previous guidance (HAN 348, June 7, 2013), CDC did not recommend MERS-CoV testing for people whose illness could be explained by another etiology. The new guidance, available at http://www.cdc.gov/coronavirus/mers/interim-guidance.html, states that, in patients who meet certain clinical and epidemiologic criteria, testing for MERS-CoV and other respiratory pathogens can be done simultaneously and that positive results for another respiratory pathogen should not necessarily preclude testing for MERS-CoV.

The new guidance also clarifies recommendations for investigating clusters of severe acute respiratory illness when there is not an apparent link to a MERS-CoV case. Clusters* of patients with severe acute respiratory illness (e.g., fever and pneumonia requiring hospitalization) should be evaluated for common respiratory pathogens and reported to local and state health departments. If the illnesses remain unexplained, testing for MERS-CoV should be considered, in consultation with state and local health departments.

For CDC’s updated interim guidance for healthcare professionals, see http://www.cdc.gov/coronavirus/mers/interim-guidance.html.

Case Definitions

CDC has not changed the case definition of a confirmed case, but the criteria for laboratory confirmation have been clarified. CDC has changed its definition of a probable case so that identification of another etiology does not exclude someone from being classified as a “probable case.”

For CDC’s updated case definitions, see http://www.cdc.gov/coronavirus/mers/case-def.html.

CDC may change its guidance about who should be evaluated and considered a case as we learn more about the epidemiology of MERS-CoV infection and risk of transmission.

Interim Guidance about Testing of Clinical Specimens

CDC recommends collecting multiple specimens from different sites at different times after symptom onset. Lower respiratory specimens are preferred, but collecting nasopharyngeal and oropharyngeal (NP/OP) specimens, as well as stool and serum, are strongly recommended. This will increase the likelihood of detecting MERS-CoV infection. For more information, see CDC’s Interim Guidelines for Collecting, Handling, and Testing Clinical Specimens at http://www.cdc.gov/coronavirus/mers/guidelines-clinical-specimens.html. Many state health department laboratories are approved for MERS-CoV testing using the CDC rRT-PCR assay. Contact your state health department to notify them of people who should be evaluated for MERS-CoV and to request MERS-CoV testing.

[If a Missouri medical provider has a patient that appears to meet the above-mentioned CDC criteria for who should be evaluated for MERS-CoV, that provider should immediately contact DHSS at 800/392-0272 (24/7) to discuss sending specimens for testing at the Missouri State Public Health Laboratory (MSPHL). Note that before any specimen is sent for testing, DHSS staff must first be consulted at 800/392-0272. After consultation with DHSS and determination that the patient meets the criteria for testing, the medical provider should then contact MSPHL at 573/751-3334 or 800/392-0272 for guidance on specimen collection and shipping prior to collecting the specimens. This will help ensure that proper specimens are obtained in the right quantity, and that they are packed and transported properly.]

*In accordance with the WHO’s guidance for MERS-CoV, a cluster is defined as two or more persons with onset of symptoms within the same 14-day period, and who are associated with a specific setting such as a classroom, workplace, household, extended family, hospital, other residential institution, military barracks, or recreational camp. See WHO’s “Interim Surveillance Recommendations for Human Infection with Middle East Respiratory Syndrome Coronavirus” at http://www.who.int/csr/disease/coronavirus_infections/InterimRevisedSurveillanceRecommendations_nCoVInfection_27Jun13.pdf.

Questions should be directed to DHSS’ Bureau of Communicable Disease Control and Prevention at 573/751-6113, or 800/392-0272 (24/7).