LISTERIOSIS

IMMEDIATE NOTIFICATION

EPIDEMIOLOGY PROGRAM

Event Name:	LIST									
Event Time Period:	1 year									
Clinical Description (CDC 2019):	 Invasive listeriosis: Systemic illness caused by <i>Listeria monocytogenes</i> manifests most commonly as bacteremia or central nervous system infection. Other manifestations can include pneumonia, peritonitis, endocarditis, and focal injections of joints and bones. Pregnancy-associated listeriosis has generally been classified as illness occurring in a pregnant woman or in an infant age ≤ 28 days. Listeriosis may result in pregnancy loss (fetal loss before 20 weeks gestation), intrauterine fetal demise (≥ 20 weeks gestation), pre-term labor, or neonatal infection, while causing minimal or no systemic symptoms in the mother. Pregnancy loss and intrauterine fetal demise are considered to be maternal outcomes. Neonatal listeriosis commonly manifests as bacteremia, central nervous system infection, and pneumonia, and is associated with high fatality rates. Transmission of Listeria from mother to baby transplacentally or during delivery is almost always the source of early-onset neonatal infections (diagnosed between birth and 6 days), and the most likely source of late-onset neonatal listeriosis (diagnosed between 7–28 days). Non-invasive Listeria infections: Listeria infection manifesting as an isolate from a non-invasive clinical specimen suggestive of a non-invasive infection; includes febrile gastroenteritis, urinary tract infection, and wound infection. 									
CDC Event Classification (2019) 18-ID-06:	 Isolation of <i>L. monocytogenes</i> from a clinical specimen collected from a normally sterile site reflective of an invasive infection (e.g., blood or cerebrospinal fluid [CSF] or, less commonly: pleural, peritoneal, pericardial, hepatobiliary, or vitreous fluid; orthopedic site such as bone, bone marrow, or joint; or other sterile sites including organs such as spleen, liver, and heart, but not sources such as urine, stool, or external wounds). For maternal isolates: In the setting of pregnancy, pregnancy loss, intrauterine fetal demise, or birth, isolation of <i>L. monocytogenes</i> from products of conception (e.g., chorionic villi, placenta, fetal tissue, umbilical cord blood, amniotic fluid) collected at the time of delivery. For neonatal isolates: In the setting of live birth, isolation of <i>L. monocytogenes</i> from a non-sterile neonatal specimen (e.g., meconium, tracheal aspirate, but not products of conception) collected within 48 hours of delivery. 									

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	Probable	 Detection of <i>L. monocytogenes</i> by culture-independent diagnostic test (CIDT) in a specimen collected from a normally sterile site (e.g., blood or cerebrospinal fluid or, less commonly: pleural, peritoneal, pericardial, hepatobiliary, or vitreous fluid; orthopedic site such as bone, bone marrow, or joint; or other sterile sites including organs such as spleen, liver, and heart, but not sources such as urine, stool, or external wounds). For maternal isolates: In the setting of pregnancy, pregnancy loss, intrauterine fetal demise, or birth, detection of <i>L. monocytogenes</i> by CIDT from products of conception (e.g., chorionic villi, placenta, fetal tissue, umbilical cord blood, amniotic fluid) collected at the time of delivery. For neonatal isolates: In the setting of live birth, detection of <i>L. monocytogenes</i> by CIDT from a non-sterile
		neonatal specimen (e.g., meconium, tracheal aspirate, but not products of conception) collected within 48 hours of delivery. Epidemiologic linkage
		4. For probable maternal cases: A mother who does not meet the confirmed case criteria BUT who gave birth to a neonate who meets confirmed or probable laboratory criteria for diagnosis AND neonatal specimen was collected up to 28 days of birth.
		5. For probable neonatal cases: Neonate(s) who does not meet the confirmed case criteria AND whose mother had <i>L. monocytogenes</i> isolated or detected by CIDT for diagnosis from products of conception OR a clinically compatible neonate whose mother had <i>L. monocytogenes</i> isolated or detected by CIDT for diagnosis from a normally sterile site.
	Suspect	Isolation of <i>L. monocytogenes</i> from a non-invasive clinical specimen source, e.g., stool, urine, wound other than those specified under maternal and neonatal specimens in confirmed section above.
Massachusetts Event Classification:	Follows CD	C event classification

Report type	Test Type	Source	Result	New event or beyond report period?	Data Entry			
Laboratory Report	Culture	Normally sterile site OR	Listeria monocytogenes	Yes	New event CONFIRMED			
		Products of conception collected at time of delivery OR non-sterile neonatal specimen collected within 48 hours of		No	Same event			
Select:	Migroorga	delivery nism: PrId: Pt: xx	v. Nom. Cultura					
Laboratory Report	Culture	Normally	Listeria spp. other than	Yes	New event			
		sterile site OR Products of conception collected at time of delivery OR non-sterile neonatal specimen collected within 48 hours of delivery	Listeria monocytogenes	No	SUSPECT Same event			
Select:	Microorga	Microorganism: PrId: Pt: xxx: Nom: Culture						
Laboratory Report	Culture	Non-sterile site	Listeria monocytogenes	Yes	New event SUSPECT			
	3.6	· DIID	N. C.1	No	Same event			
Select:	Select: Microorganism: PrId: Pt: xxx: Nom: Culture							

Laboratory Report	PCR	Normally sterile site OR	Positive	Yes	New event PROBABLE		
				NT -			
		Products of		No	Same event		
		conception					
		collected at					
		time of					
		delivery OR					
		non-sterile					
		neonatal					
		specimen					
		collected					
		within 48					
		hours of					
		delivery					
Select:	Listeria monocytogenes rRNA: ACnc: Pt: xxx: Ord: Probe						
Select:	Listeria monocytogenes DNA: ACnc: Pt: xxx: Ord: Probe.Amp.Tar						
Select:	Meningitis+Encephalitis pathogens DNA and RNA panel - Cerebral spinal fluid by NAA with non-probe detection						

Data entry notes:

- Cases in neonates and mothers should be reported separately when each meets the case definition. A case in a neonate is counted if live-born.
- Pregnancy loss and intrauterine fetal demise are considered maternal outcomes and would be counted as a single case in the mother.