Influenza-Associated Pediatric Mortality Case Report Form

Form Approved OMB No. 0920-0004

STATE USE ONLY – DO NOT SEND INFORMATION IN THIS SECTION TO CDC					
Last Name:	First Name:		County:		
Address:	City:		State, Zip:		
Patient Demographics					
1. State:	2. County:	3. State ID:	4. CDC ID:		
			7a. Is sex known? ☐ Yes ☐ No		
5. Age: O Days O Months O Years	6. Date of birth:// MM DD	YYYY	7b. Sex: O Male O Female		
8a. Is ethnicity known? ☐ Yes ☐	□ No				
8b. Ethnicity: O Hispanic or Latino					
oo. Eminerty: O mispanic of Eatino	O 100 Thispanic of Eatino				
9a. Is race known? ☐ Yes ☐ No)				
9b. Race: ☐ White ☐ Black	☐ Asian ☐ Native Hawaiian or C	Other Pacific Islander	☐ American Indian or Alaska Native	e	
Death Information 12. Was an autopsy performed?					
10. Date of illness onset:// _DD / YYYY					
13 a. Did cardiac/respiratory arrest occur outside the hospital? O Yes O No O Unknown					
13 b. Location of death: O Outside the Hospital (e.g. home or in transit to hospital) O Emergency Dept (ED) O Inpatient ward O ICU O Other (specify):					
13 c. If the death occurred in the hospital, what was the date of admission?/					
CDC Laboratory Specimens	1				
14 a. Were pathology specimens sent to CDC's Infectious Diseases Pathology Branch? O Yes O No O Unknown Please provide the lab ID No. if known					
14 b. Were influenza isolates or original clinical material sent to CDC's Influenza Division? O Yes O No O Unknown Please provide the lab ID No. if known					

Public reporting burden of this collection of information is estimated to average 30 minutes per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. An agency may not conduct or sponsor, and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to CDC/ATSDR Information Collection Review Office, 1600 Clifton Road NE, MS D-74, Atlanta, Georgia 30333; ATTN: PRA (0920-0004).

Influenza Testing (check all that were used)				
Test Type	Result	Specimen Collection Date		
15. ☐ Commercial rapid diagnostic test	O Influenza A O Influenza B O Negative O Influenza A/B (Not Distinguished) O 2009 Influenza A (H1N1) O Influenza virus co-infection (specify)	/		
☐ Viral culture	O Influenza A (Subtyping Not Done) O Influenza A (H3N2v) O Influenza A (Unable To Subtype) O Influenza B/Victoria lineage O Influenza B/Victoria lineage O Influenza Virus co-infection (specify) O Negative	/		
☐ Fluorescent antibody (IFA or DFA)	O Influenza A (Subtyping Not Done) O Influenza B O Negative O Influenza A (Unable To Subtype) O Influenza A (H3) O 2009 Influenza A (H1N1) O Influenza virus co-infection (specify)	/		
☐ Enzyme immunoassay (EIA)	O Influenza A (Subtyping Not Done) O Influenza B O Negative O Influenza A (Unable To Subtype) O Influenza A (H3) O 2009 Influenza A (H1N1) O Influenza virus co-infection (specify)	/		
□ RT-PCR	O Influenza A (Subtyping Not Done) O Influenza A (H1N1) O Influenza A (H3N2v) O Influenza A (H3N2v) O Influenza A (Unable To Subtype) O Influenza B (Lineage Not Determined) O Influenza B/Victoria lineage O Influenza Virus co-infection (specify) O Negative	/		
☐ Immunohistochemistry (IHC)	O Influenza A O Influenza B O Negative O Influenza virus co-infection (specify)	//		
Culture confirmation of bacter	rial nathagans from STEDII F (Invasiva) SITES			
Culture confirmation of bacterial pathogens from STERILE (Invasive) SITES 16 a. Was a specimen collected for bacterial culture from a normally sterile site (e.g., blood, cerebrospinal fluid [CSF], tissue, or pleural fluid? Specimens collected greater than 24 hours after death are not sterile. O Yes O No O Unknown				
16 b. If yes, please indicate the site from which the specimen was obtained and the result. If more than one specimen type is positive and more than one organism is identified please indicate the organism cultured from each specimen type in the comments section. Specimen Type				
☐ CSF ☐ Lung Tissue	Date _ / _ / O Positive O Negative O Unknown Date _ / _ / O Positive O Negative O Unknown Date _ / _ / O Positive O Negative O Unknown			
16 c. If positive, please check the organ	nism cultured.			
□ Streptococcus pneumoniae	☐ Staphylococcus aureus, methicillin sensitive ☐ Haemop (MSSA)	hilus influenzae not-type b		
☐ Group A Streptococcus	☐ Staphylococcus aureus, methicillin resistant ☐ Haemop (MRSA)	hilus influenzae type b		
☐ Other bacteria:(If reporting another viral co-infection section 18 Clinical Diagnosis and C	n please do so in	nonas aeruginosa		

Culture confirmation of bacterial pathogens from NON-STERILE SITES					
16 d. Were other <u>respiratory</u> specimens collected for bacterial culture (e.g., sputum, ET tube aspirate)? O Yes O No O Unknown					
	om which the specimen was obtained and the result.	If more than one specimen type is positive and more than in the comments section.			
Specimen Type	Collection Date Result				
□ Sputum □ ET tube □ Other □ Unknown	Date/ O Positive O Negative O Unl Date/_/ O Positive O Negative O Unl Date/_/ O Positive O Negative O Unl	known			
16 f. If positive, please check the orga	anism cultured.				
☐ Streptococcus pneumoniae	☐ Staphylococcus aureus, methicillin sensitive (MSSA)	☐ Haemophilus influenzae not-type b			
☐ Group A Streptococcus	☐ Staphylococcus aureus, methicillin resistant (MRSA)	☐ Haemophilus influenzae type b			
☐ Other bacteria:	☐ Staphylococcus aureus, sensitivity not done	☐ Pseudomonas aeruginosa			
(If reporting another viral co- infection please do so in section 18 Clinical Diagnosis and Complications)					
Pathology confirmation of ba	atorial nathogons				
16 g. Was a specimen (e.g., fixed lung or state pathologist? (If pathology res	stissue) collected from an autopsy for testing of bacter ults are available from CDC it is not necessary to impose esection 14 "CDC Laboratory Specimens")				
If yes please indicate the results of the	se tests in the comments section at the end of the form	n.			
W II 16					
Medical Care					
17. Was the patient placed on mechanical ventilation? O Yes O No O Unknown					

Clinical Diagnoses an	d Complications					
18 a. Did complications occ	18 a. Did complications occur during the acute illness? O Yes O No O Unknown					
18 b. If yes, check all comp	lications that occurred dur	ing the acut	te illness:			
☐ Pneumonia (Chest X	-Ray confirmed) □	d) ☐ Acute Respiratory Disease Syndrome (ARDS) ☐ Croup ☐ Seizures			☐ Seizures	
☐ Bronchiolitis		☐ Encephalopathy/encephalitis		☐ Reye syndrome	☐ Shock	
☐ Sepsis		☐ Hemorrhagic pneumonia/pneumonitis		☐ Cardiomyopathy/myocarditis		
☐ Another viral co-info	ection:	☐ Other:				
19 a. Did the child have an	y medical conditions that	existed befo	ore the start of the acute illness?	O Yes O No O Unkno	own	
19 b. If yes, check all med	ical conditions that existed	before the	start of the acute illness:			
☐ Moderate to severe developmental delay ☐ Hemoglobinopathy (e.g. sickle cell disease) ☐ Asthma/ reactive airway disease						
☐ Diabetes mellitus	☐ History seizures	of febrile	☐ Seizure disorder	☐ Cystic fibrosis	s	
☐ Cardiac disease/congenit	al heart disease (specify)		☐ Renal disease (specify)	Skin or soft ti	ssue infection (SSTI)	
☐ Chromosomal Abnormal	lity/Genetic Syndrome (sp	ecify)	☐ Mitochondrial Disorder (specif	ŷ)		
☐ Chronic pulmonary disea	ase (specify)		☐ Immunosuppressive condition	(specify)		
☐ Cancer (diagnosis and/or treatment began in previous 12 months) ———————————————————————————————————						
□ Neuromuscular disorder (e.g. muscular dystrophy) (specify) □ Other Neurological disorder (specify)						
☐ Pregnant (specify gestational age) weeks ☐ Other (specify)						
Medication and Ther	any History					
20 a. Was the patient receiving any of the following therapies <i>prior</i> to illness onset? (if yes, check all that apply)						
□Yes	□ No	□ Unknown				
□Antiviral Prophylaxis	☐ Chronic aspirin therapy	☐ Chemotherapy or radiation therapy		☐ Steroids by mouth or injection		
☐ Other immunosuppressive therapy:						
20 b. Did the patient receive any of the following <i>after</i> illness onset? (if yes, check all that apply)						
□ Yes □ No □ Unknown						
☐ Antibiotic therapy specify ☐ Antiviral therapy specify						

Influenza Vaccine History					
21. Did the patient receive any influenza vaccine during the current season (before illness) O Yes O No O Unknown					
22. If YES*, please specify the influenza vaccine received before illness onset: □ Inactivated influenza vaccine (IIV3) [injected] □ Quadrivalent inactivated influenza vaccine (IIV4) [injected] □ Live-attenuated influenza vaccine (LAIV4) [nasal spray] □ Unknown	Inactivated influenza vaccine (IIV3) [injected] Quadrivalent inactivated influenza vaccine (IIV4) [injected] Live-attenuated influenza vaccine (LAIV4) [nasal spray]				
23. If YES*, how many doses did the patient receive and what was the timing of each dose? (Enter vaccination dates if available)					
O 1 dose □ <14 days prior to illness onset □ ≥14 days prior to illness onset □ Date dose given:					
O 2 doses onset onset onset 2^{nd} dose given <14 days prior to onset 2^{nd} dose given \ge 14 days prior to onset 2^{nd} dose given \ge 14 days prior to onset 2^{nd} dose 2^{n	YY				
23b. IF the patient received two doses of influenza vaccine during the current season, please specify the SECOND influenza vaccine received before illness onset: □ Inactivated influenza vaccine (IIV3) [injected] □ Quadrivalent inactivated influenza vaccine (IIV4) [injected] □ Live-attenuated influenza vaccine (LAIV4) [nasal spray] □ Unknown					
24 . Did the patient receive any influenza vaccine in previous seasons? O Yes O No O Unknown					
24 a. If YES , and patient was ≤8 years of age at the time of death, did they receive 2 doses of vaccine during a previous season? O Yes O No O Unknown					
Submitted By: Date: / Phone No.: () MM DD YYYY E-mail Address: Case Investigation Closed: □ Yes □ No					