Form Approved OMB No. 0920-0009 Exp Date: 04/30/2016

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PART I. Acute Neurological Illness with Limb Weakness in Children: Patient Summary Form

Form to be completed by, or in conjunction with, a physician who provided care to the patient during the neurological illness.

Confirmation of case:	Yes	No	Unknown
a. Neurological findings (upon examination by clinician) include focal limb weakness			
b. MRI of spinal cord demonstrates spinal lesion(s) largely restricted to or predominantly affecting the			
gray matter.			
(Terms in the spinal cord MRI report such as "affecting mostly gray matter," "affecting the anterior horn			
or anterior horn cells," "affecting the central cord," "anterior myelitis," or "poliomyelitis" would all be			
consistent with this. If still unsure if this criterion is met, consider asking the radiologist directly.)			
c. Age at onset of limb weakness is 21 years or less			
d. Onset of limb weakness was August 1, 2014 or later			

Answer to <u>ALL 4 criteria must be YES</u>. If yes, continue to Part II on pages 2 - 5. If you have any questions about whether your patient meets all 4 criteria, please e-mail us at limbweakness@cdc.gov

Public reporting burden of this collection of information is estimated to average 60 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 Reports Clearance Officer; 1600 Clifton Road NE, MS D-74 Atlanta, Georgia 30333;

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PART II. Acute Neurological Illness with Limb Weakness in Children: Patient Summary Form

Form to be completed by, or in conjunction with, a physician who provided care to the patient during the neurological illness. Once completed, submit to Health Department (HD). HD can also facilitate specimen testing.

1.Today's dat	:e/	/	(mm/dd/yyyy) 2	.Name of person cor	mple	ting fo	rm:									
3. Affiliation_			[Phone:				En	nail:							
4. Name of p	hysician who	can provide a	ndditional clinical/	lab information, if n	eede	ed										
5. Affiliation_				Phone:				E	mail:							
6. Name of m	nain hospital t	hat provided:	patient's care:					7.	.State	:	8 .0	County:				
9. Patient ID:			10. 9	State ID:							_ 11.	Patient	t's sex	x: □	M [⊒F
12. Patient's	age:y	ears AND	months	Patient's resid	lence	e: 13 . S	tate		14. (County_						
15 . Race:	□Asian □White	□Black or a (check all ti	African American hat apply)				Pacific Is			⊐Ameri	can	Indian c	or Alas	ka N	ative	
			//	(<i>mm/dd/yyyy</i>) 20. Date of disc												known alized)
21 . Current c	linical status:	□recovered	□not recover	red, but improved	□no	t impro	oved l	□Dec	ease	d: 22 .Da	te o	f death_	/		_/	
Signs/symp	toms/condi	tion at <u>ANY</u>	time during the	e illness:										\equiv		
						Right A	Arm		Left A	rm		Right L	eg	L	Left L	eg
[indicate yes	(y), no (n), uni	known (u) for	h limbs have beer each limb] at worst weaknes	·	Υ	N	U	Υ	N	U /	/ Y		U	Υ	N	U
(mm/dd/yyyy	<i>')</i>									/			-			
26. Any senso	ory loss/numb	ness in the a	corded at worst w ffected limb(s), at considered here)	,		Arefle	xic/hypo	orefle	xic (0	-1) □ N Y N		nal (2) [J	⊒ Нур	erre	flexic (3-4+)
`'			•	me during illness)	Υ	N	U	Υ	N	U	Υ	N	U	Υ	N	U
,,			.,,,,,	<u> </u>								Yes	No		Unkn	own
28. Sensory l	evel on the to	rso (ie, reduc	ed sensation belo	ow a certain level of	the t	orso)?	(at any	time	durir	ng illnes	s)					
29. At any tin	ne during the	illness, pleas	e check if the pati	ent had any of the f	ollov	ving cr	anial ne	rve si	gns:							
□Dip	lopia/double	vision (If yes,	circle the cranial	nerve involved if kn	own	: 3 / 4	4 / 6)									
	s of sensation					sphag			ysart	hria						
30. Any pain	or burning in	neck or back	? (at any time dui	ring illness)												
			any time during i													
				alternating tachy/bra	adyca	ardia)?	(at any	time	durir	ng illnes	s)					
				encephalopathic)? (
	? (at any tim					-										
			•	n? (at any time duri	ng ill	ness)								\neg		
				tracheostomy) hec			ırologic	al con	ditio	n?						

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Other patient information:

Neuroradiographic findings: (Indicate based on <u>most abnormal</u> study)

enhance with GAD?

Within the 4-week period BEFORE onset of				
limb weakness, did patient:	Yes	No	Unk	
37. Have a respiratory illness?				38 . If yes, date of onset//
39 . Have a fever, measured by parent				40 . If yes, date of onset / /
or provider and ≥ 38.0°C/100.4°F?				40. If yes, date of offset
41. Receive oral, IM or IV steroids?				
42. Receive any other systemic				43. If yes, list:
Immunosuppressant(s)?				
44. Travel outside the US?				45. If yes, list country
46 . Does patient have any underlying illnesses?				47. If yes, list
48. On the day of onset of limb weakness, did				
patient have a fever? (see definition above)				

Polio vaccination history:			
49. How many doses of inactivated polio vaccine (IPV) are documented to have been received by			
the patient before the onset of weakness?	_doses	□unknown	
50. How many doses of oral polio vaccine (OPV) are documented to have been received by the			
patient before the onset of weakness?	doses	□unknown	
51. If you do not have documentation of the <i>type</i> of polio vaccine received:			
a. What is total number of documented polio vaccine doses received before onset of weakness?	doses	□unknown	

53. Levels imaged:	52 . Date of study// □cervical □thoracic □lumbosacral □yes □no □unknown		
55. Location of lesions:	□cervical cord □thoracic cord □conus □cauda equina □unknown	Levels of cord affected (if applicab 56. Cervical:	le): 62. Thoracic:
For cervical and thoracic	57. What areas of spinal cord	□predominantly gray matter	□predominantly white matter
cord lesions	were affected?	□both equally affected	□unknown
	58 . Was there cord edema?	□yes □no □unknown	
For cervical, thoracic cord or conus lesions	59 . Did any lesions enhance with GAD?	□yes □no □unknown	
For cauda equina lesions	60 . Did the ventral nerve roots enhance with GAD?	□yes □no □unknown	
	61. Did the dorsal nerve roots	□yes □no □unknown	

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MRI of brain 63. Gadoliniu		Date of study _ ves □no	<i>//</i> unknown		_ (mm	n/dd/yyyy)				
	atentorial (i.e, lobe ortical, basal gangl esions	I	□no □unk	nown						
		65 .If yes	, indicate locat	tion(s)	□cort	tex □subo	cortex 🗆 ba	sal ganglia	□thalamus	□unknown
			s, did any lesion		□yes	□no	□unknown			
67. Any brain	stem lesions?	□yes	□no □unk	nown						
		68 . If yes	, indicate loca	tion:	□mid	lbrain □p	ons □m	edulla	□unknown	
		1	s, did any lesion nce with GAD?		□yes	□no	□unknown			
70. Any crani	al nerve lesions?	□yes								
		71. If yes CN(s	s, indicate which):	ch	CN	Dunilater	ral □bilatera	l CN	□unilateral	□bilateral
					CN	🗆 unilater	ral □bilatera	l CN_	□unilateral	□bilateral
			s, did any lesion nce with GAD		□yes	□no	□unknown			
73. Any lesion	ns affecting the m ?	□yes	□no □unk	nown	-					
CSF examin	MG done?	f acute motor n umbar puncture		otor neu			erve or anterio			
	Date of lumbar puncture	WBC/mm3	% neutrophils	% lympho	ocytes	% monocytes	% eosinophils	RBC/mm3	Glucose mg/dl	Protein mg/dl
77a. CSF from LP1	panetare	WBC/IIIIIS	псинорииз	Tymphe	reytes	monocytes	СОЗПОРППЗ	NDC/IIIII3	Glucose mg/ui	Trotein ing/ui
77b. CSF										
from LP2										
Pathogen to	esting performed	:								
78. Was CSF to pathogens?	tested for the follo	wing	Date of spec	cimen co	ollectio	n/	/		☐ Not done	
				s PCR: □ tive: typ		ve □ Negat	ive □ Not o			
						Positive \square	Negative \square			
							PCR	□ N-+ '		
							□ Negative		one	
			Cytomegalo Varicella Zo			□ Positive □	Negative C		<u> </u>	
			Other patho				☐ Negative	□ Not do	ліс	
			1 - 3 - 10 - 1110	5		-1				

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Type of test:

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79 . Was a respiratory tract specimen tester for the following pathogens?	ed [Date o	f specimen coll	ection//	☐ Not done
	E		virus/rhinoviru positive: type:	s PCR: ☐ Positive ☐ Negative ☐ Not d ☐ Not typed	lone
	1	Adeno	virus PCR: DP	ositive	
			positive: type:	☐ Not typed☐ Positive☐ Negative☐ Not done☐	
	'		positive: type:		
		Other p	oathogen ident	ified: specify:	
	1	ype o	f test:		
80. Was a stool specimen tested for the following pathogens?	[Date of	f specimen coll	ection//	☐ Not done
			ovirus PCR: positive: type:	Positive ☐ Negative ☐ Not done ☐ Not typed	
	F			sitive □ Negative □ Not done	
	F	Poliovi	rus culture: 🏻	Positive ☐ Negative ☐ Not done	
			oathogen ident	ified: specify:	
		ype o	t test:		
81. Was serum tested for the following pathogens?				ection/	☐ Not done
	\			Positive □ Negative □ Not done ype: □ IgM □ PCR	
			oathogen ident	ified: specify:	
82. Describe any other laboratory finding(s		ype o			
52. Describe any other laboratory infamily	,, consid	ici cu t	o be significan		
83. Was/Is a specific etiology considered to 84. If yes, please list etiology and reason(s)			•		
-					
Treatment: 85. Were any of these the	rapies a	admin	istered for th	e acute neurologic illness? (as of time o	of form completion)
	Yes	No	Unknown		
a. Antibiotics				If yes, date first administered:/	
b. Antivirals					te first administered://
c. Corticosteroids				If yes, date first administered:/	
d. Intravenous immune globulin (IVIG)				If yes, date first administered:/	
e. Plasma exchange or Plasmapheresis				If yes, date first administered:/	
f. Interferon				If yes, specify; dat	te first administered://
g. Other immunosuppressive therapy				If yes, specify; dat	te first administered://
86. Other information you would like us to	know _				
87. Indicate which type(s) of specimens from			_		-
☐ CSF ☐ Nasal wash/aspirate ☐ BAL sp	oec □	Trach	eal aspirate	□NP/OP swab □Stool □Serum □	Other, list
☐ No specimens stored					

[Type text	[7	Гу	pe	te	ext
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This section below for CDC use	
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Patient Number (assigned by CDC PPLB Lab)	
Patient Number (assigned by CDC PPLB Lab) State Specimen ID	State Specimen ID