

#### **Zika Virus Disease Case Investigation Form**

Arboviral Diseases Branch Version 3.1

Form Approved OMB No. 0920-1011 Exp. Date 03/31/2017



FOR CDC USE ONLY	
CDC R-number	ZIKVID:
CDC staff initial:	Date form completed:/
CDC investigating group:	
Reporting Jurisdiction	
Jurisdiction (state/territory):	Agency:
Contact Name:	Contact Phone:
Contact Position:	Contact Email:
Alternate Contact Name:	Alternate Contact Phone:
Demographic Information	
State of residence:	State patient ID number:
Patient last name:	Patient first name:
Age:   Years   Months   Days	Sex: ☐ Male ☐ Female
Travel History	
Dates of travel:	
Country(s) visited:	
Vaccination History	
Previously vaccinated for:	apanese Encephalitis
Cases of Special Interest	
Please indicate if patient meets any of the following	criteria:
Local vector-borne transmission	☐ Yes ☐ No ☐ Suspect
Pregnant	☐ Yes ☐ No ☐ Unknown
	If yes: Current gestational week: Gestational week at illness onset (if applicable):
Fetal loss	☐ Yes ☐ No
	If yes: Gestational week at time of fetal loss:
Microcephaly	☐ Yes ☐ No ☐ Suspect
Guillain-Barre syndrome/acute flaccid paralysis	☐ Yes ☐ No ☐ Suspect
Sexual transmission	☐ Yes ☐ No ☐ Suspect
Blood/blood product transfusion transmission	☐ Yes ☐ No ☐ Suspect
Organ/tissue transplant transmission	☐ Yes ☐ No ☐ Suspect
Breastfeeding transmission	☐ Yes ☐ No ☐ Suspect



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Arboviral Diseases Branch Version 3.1



Illness Ir	Illness Information					
Illness on	set date:	/ 🗆 н	ospitalized			
Fever	□ Yes □ I	No				
	If yes: □ S	ubjective fever	ever (Maximum measured temperature:)			
Rash	□ Yes □ I	No				
	If yes:	Type: ☐ Maculopapular Pruritic: ☐ Yes ☐ No Distribution:				
☐ Arth	ralgia	☐ Myalgia	☐ Oral ulcers			
☐ Conj	unctivitis	☐ Vomiting	☐ Hematospermia (for males)			
☐ Head	dache	☐ Diarrhea	☐ Peripheral edema			
Specime	n Informati	on				
Specimer	n 1 collected:		Type: ☐ Serum ☐ CSF ☐ Amniotic fluid ☐ Tissue ☐ Saliva ☐ Urine ☐ Semen			
Specimer	n 2 collected:		Type: ☐ Serum ☐ CSF ☐ Amniotic fluid ☐ Tissue ☐ Saliva ☐ Urine ☐ Semen			
Specimen 3 collected:/			Type: ☐ Serum ☐ CSF ☐ Amniotic fluid ☐ Tissue ☐ Saliva ☐ Urine ☐ Semen			
Specimen 4 collected:/			Type: ☐ Serum ☐ CSF ☐ Amniotic fluid ☐ Tissue ☐ Saliva ☐ Urine ☐ Semen			
Specimer	າ 5 collected:		Type: ☐ Serum ☐ CSF ☐ Amniotic fluid ☐ Tissue ☐ Saliva ☐ Urine ☐ Semen			
Specimer	n 6 collected:	/	Type: ☐ Serum ☐ CSF ☐ Amniotic fluid ☐ Tissue ☐ Saliva ☐ Urine ☐ Semen			
Specimer	n 7 collected:	/	Type: ☐ Serum ☐ CSF ☐ Amniotic fluid ☐ Tissue ☐ Saliva ☐ Urine ☐ Semen			

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#### ANEXO H: EXTRAÇÃO DE PRONTUÁRIO PARA BEBÊS CASOS

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Equipe:	entificação do bebê ca	iso:	Data d	a extração:	Exp. Date 05/51/2017
Nome da unidade da saú	úde:				
Esta	ado:		Município	:	
HISTÓRICO DO BEB	Ê				
DT Nasc Bebe		Data de nascimento mãe		Peso ao n	nascer:
Comprimento:	PC:	Data das medições:		Hora das med	ições:
PROBLEMAS DE SAÚ	DE DURANTE O INT	ERNAMENTO			
Problemas de audiçã	ío 🔲 Cegueira	Convulsões Difi	culdade na deglutiçã	ăo <u>□</u> D∈	esconforto respiratório
Sepse	Nenhum prob	olema Out	ro: Especifique		
RESULTADOS DA IM	AGIOLOGIA E EXAM	IES PARA O BEBÊ			_
Tomografia computador	izada:	▼ Se sim, data:			
Normal Calcifica	ações 🔲 Lisencefalia	Atrofia cerebral V	entriculomegalia 🔲	Suturas calcif	ficadas Outras
Outras,especificar:					
Ultrassonografia transfo	ntanelar:	▼ Se sim, data:			
Normal Calcifica	ações 🔲 Lisencefalia	Atrofia cerebral V	entriculomegalia 🔲	Suturas calcif	ficadas 🔲 Outras
Outras, especificar:					
Ressonância magnética	a: 🔻 :	Se sim,data:			
Normal Calcifi	cações Lisencefali	a Atrofia cerebral	Ventriculomegalia	Suturas ca	alcificadas 🔲 Outras
Outras, especificar:					
Ecocardiograma:	▼ Se	sim, data:		٦	
Se sim:		alterado (resultado):			
USG abdominal:	▼ Se	sim, data:		 	
Se sim:		alterado (resultado):			
EXAMES E HISTÓRIO	O PRÉ-NATAL				
Ultrassonografia pré-na	atal:	Resultado	▼ Se for anor	mal, data:	
Se anormal, especifique:					
Amostragem vilo corial:					
Descrever					
Amniocente	ese 🔻	Resultado:	▼		
Descrever:					

Complicações dur	ante a gestação?	1	•					
–Se sim, quais: —								
ITU		Anemia		DM ges	tacional		HAS gestacional	
Pré eclampsia		Placenta	a prévia	Oligodr	amnio		Polidramnio	
Insuficiência o	olo uterino	Hiperen	nese gravidica	Anoma	lias anatômicas	no útero	Descolamento de pla	acenta
Crescimento i	ntraútero restrito	Incisura	à	Outras				
Se outras, espec	ificar:							
Medicamentos da	ı mãe durante a g	ravidez:						
_			Medicamento	1				
Medicamento 1			30 dias ant	eriores da g	gravidez 🔲 1	o Tri	2º Tri 🔲 3º Tri	
Medicamento 2:	Medicamento 2:							
Medicamento 2.			30 dias ant	eriores da g	gravidez 🔲 1	o Tri	2º Tri 3º Tri	
Medicamento 3:			Medicamento					
Tredicamento 5.			30 dias ant	eriores da g	gravidez 🔲 1	º Tri	2º Tri 3º Tri	
Medicamento 4:			Medicamento					
			30 dias ant	eriores da g	gravidez 🔲 1	Tri	2º Tri 3º Tri	
Medicamento 5:			Medicamento		=		=	
L			30 dias ant	eriores da g	gravidez 🔲 1	o Tri	2º Tri 3º Tri	
–Exames de doen	ças infecciosas p	ara o bebê	<u> </u>					
VDRL	Res	ultado	▼					
CMV	▼	IgM	_	IgG		PCF		
	•							
HSV 1	▼	IgM	▼	IgG	•	PCF	▼	
HSV 2	•	IgM	•	IgG	•	PCF	₹	
Rubéola	▼	IgM	▼	IgG	•	-		
Тохо		IgM		IgG				
	▼	Igin	•	igo	`			
Dengue	•	IgM	•	IgG	•	PCR	•	
Zika	_	IgM	_	IgG		PCR	_	
	<u> </u>	-5	•	-3-			•	
Chikungunya	•	IgM	•	IgG	•	PCR	•	
Outros1	▼ Bosult	adas Outras	.1.					
Outros 2:	Result	ados Outros ados Outros						
Outros Z.	Kesuit	auus Outi 05						

Ν

Exames de d	oenças infecci	osas durante a gravidez  —					
VDRL	_	Resultado	▼				
CMV	<b>—</b>	IgM	-	IgG	<b>•</b>	PCR	▼
HSV 1	<b>-</b>	IgM	<b>~</b>	IgG	·	PCR	▼
HSV 2	_	IgM	<b>~</b>	IgG	<b>-</b>	PCR	▼
Rubéola	_	IgM	~	IgG	•		
Тохо	-	IgM	<b>~</b>	IgG	_		
Dengue		IgM	•	IgG		PCR	▼
Zika		IgM	▼	IgG	•	PCR	•
Chikungunya		IgM	▼	IgG	•	PCR	▼
Outros1		Resultados Outros1:					
Outros 2	-	Resultados Outros2:					
O bebê fez exa	ame de vista?	•		Resulta	do de exame de vi	sta do bel	bê: ▼
Caso anormal,	descrever:						
Outras anom	alias/defeitos	do bebê ————					
Presença de m	alformações no	RN:					
Se sim, espe	ecificar:						
Aparelho o	circulatório 🔲	Aparelho digestivo 🔲 Aparel	lho respirató	rio 🔳	Órgãos genitais	Apare	lho osteomuscular
Outras							
Descreva a ma	lformação enco	ntrada:					
	,						
Descrever res	sultados de ex	xames ou defeitos específic	cos de forn	na mai	s detalhada		

#### Questionário da pesquisa - Microcefalia

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Equipe:								- :   <b>-</b> : <b>-</b> :		
	2 @ 3 @ 4	<b>5</b>	6 0	7		NU		e identificação		
							ı	Número de pare	amento:	
Entrevista	dor						Da	ta da entrevista	:	
Endereço				Latitud	de 🗀					٦
-								Gerar coord	lenadas	
				Longitud	de					
Data Visita 1	_T	urno				Situação 1				
		) Manhã	⊚ Tar	de   Noite	9	<ul><li>Participa</li></ul>	ante @	Recusou-se	Indispor	nível
Data Visita 2		urno —				Situação 2				
		) Manhã	⊚ Tar	de   Noite	Э	<ul><li>Participa</li></ul>	ante @	Recusou-se	Indispor	nível
Visita 3		urno —				Situação 3				
		) Manhã	Tar	de   Noite	Э	Participa	ante @	Recusou-se	Indispor	nível
A Introdu	ução									
melhor por	que alguns beb ual da mãe (and ual do bebê:	pês têm m		a e outros não	).	3. Sexo do		ue nos ajudem a		•
5. Localiza	ação da residên	cia:			•					
	cia e histórico nto tempo você Meses	mora na		s da mãe						
2. Há quar	nto tempo você	mora em	seu end	ereço atual?				▼		
3. Durante	e a gravidez, vo	cê morou						▼		
4. Durant	e a gravidez, vo	ocê passou	u 3 noites	s consecutivas	ou mais	fora de casa, oi	nde o tr	ajeto foi superio	or a 3 horas?	
									•	-
	nãe estiver mor para o estudo;					a idade do bel	bê), talv	ez não atenda a	os critérios d	е
_	atas e destinos									
Data		l	_ocais:				Dias:			
Data			ocais				Dias:			
Data			Locais:				Dias			

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 Reports Clearance Officer; 1600 Clifton Road NE, MS D-74 Atlanta, Georgia 30333; ATTN: PRA (0920-1011)

C. Informações sobre a gravidez						
1. Data da útima mestruação:		2. Qual f	oi a data prováve	el do parto inforr	nada pelo médico?	
	■ Não	sei				Não sei
3. Tipo de gravidez ▼	Se gemela	r, número de be	ebês:	4. Os outros be	ebês nasceram vivos?	•
Se sim, teve alguma malformação:	•	Especificar				
5. Tipo de parto:		•				
6. Quantas vezes você engravidou ante bebê nascido morto ou outros resultad		videz, incluindo	gestações que p	odem ter termin	ado em abortos natur	ais,
Número de gestaç	ões:		Número de nasc	idos vivos		
Número de ab	oortos:	Núi	mero de nascido:	s mortos:		
7. Há alguma (outra) criança ou adulto	da sua fam	nília que nasceu	com microcefalia	9? ▼		
Se sim, especifique (grau de parente	sco em relaç	ção ao bebê e o	momento do dia	agnóstico):		
8. Você e o paí da criança têm algum ç	grau de pare	entesco?	<b>▼</b> S	e sim, qual?		
9. Qual é a data de nascimento deste l	oebê?					
<ul><li>D. Doenças durante a gravidez</li><li>Agora, vou fazer algumas perguntas</li><li>1. No período de 30 DIAS ANTES d</li></ul>					os seguintes sintomas	:
			Semanas ou	meses		
Manchas vermelhas no corpo:	•	Quando	Semanas	Meses	Não sei	
Febre:	•	Quando	Semanas ou		Não sei	
Coceira:	•	Quando	Semanas ou		Não sei	
Dores articulações:	•	Quando	Semanas ou		Não sei	
Olhos vermelhos:	•	Quando	Semanas ou		Não sei	

2. Durante o PRIMEIRO TRIME sintomas?	STRE de gravidez (até 13 semanas), você t	eve alguma doença com algum dos seguintes
Manchas vermelhas no corpo:	→ Quando	Semanas ou meses
Febre:	<b>→</b> Quando	Semanas ou meses
Coceira:	<b>▼</b> Quando	Semanas ou meses
oocana.	• quanto	
Dores articulações:	<b>▼</b> Quando	Semanas ou meses
	· Camina	
Olhos vermelhos:	<b>▼</b> Quando	Semanas ou meses
Ollios vertileinos.	<b>↓</b> Quando	
3. Durante o SEGUNDO TRIME sintomas? Manchas vermelhas no corpo:	STRE de gravidez (14 a 26 semana), você te	eve alguma doença com algum dos seguintes Semanas ou meses
·		
Febre:	<b>▼</b> Quando	Semanas ou meses
	- Quanto	
Coceira:	<b>▼</b> Quando	Semanas ou meses
Cocena.	→ Quando	
Dores articulações:	<b>▼</b> Quando	Semanas ou meses
Bores ar clearagoes.	Qualita	
Ollege vermeellege		Semanas ou meses
Olhos vermelhos:	▼ Quando	

4. Durante o TERCEIRO TRIMES sintomas?	TRE de gravidez (27	a 42 semanas), você	teve alguma doença com algum dos seguintes
Manchas vermelhas no corpo:	<b>▼</b> Q	uando	Semanas ou meses
Febre:	<b>→</b> Q	uando	Semanas ou meses
Coceira:	<b>▼</b> Q	uando	Semanas ou meses
Dores articulações:		uando	Semanas ou meses
Dores articulações.	▼ 4	dando	
Olhos vermelhos:		uando	Semanas ou meses
omos vermemos.	▼ 4	dando	
afirmativo, registrar a seman	ividez e o final da gra a de gravidez, se pos	videz, você teve algu sível, e o(s) mês(es)	ıma das seguintes doenças ou infecções? [Em caso de gravidez, caso a semana seja desconhecida]
Infecção urinária			Semanas ou meses
-30 Infecção dos rins, bexiga ou t	rato urinário	<b>→</b> Quando	Semanas Meses Não sei
			Semanas ou meses
1º Tri Infecção dos rins, bexiga ou	ı trato urinário	<b>→</b> Quando	Semanas Meses Não sei
			Semanas ou meses
2º Infecção dos rins, bexiga ou tra	ato urinário	<b>→</b> Quando	Semanas Meses Não sei
3º Tri Infecção dos rins, bexiga ou	u trato urinário	<b>→</b> Quando	Semanas ou meses
			Semanas Meses Não sei
Infecções por fungos			
20 7-6		0	Semanas ou meses
-30 Infecção por fungos	•	Quando	
			Semanas ou meses
1º Tri Infecção por fungos	•	Quando	Semanas Meses Não sei
2º Tri Infecção por fungos		Quando	Semanas ou meses
2º 111 Illiecção por lungos	•	Qualido	
			Semanas ou meses
3º Tri Infecção por fungos	•	Quando	

Toxoplasmose  Quando  Semanas ou meses Rubéola  Quando  Semanas ou meses Depectifique: Quando Semanas ou meses Não sei Não sei Semanas ou meses Não sei Não sei Semanas ou meses Não sei Semanas ou meses Não sei Semanas ou meses				Semanas ou me	eses	
Quando Semanas o Meses Não sei  Semanas ou meses Semanas	Toxoplasmose	•	Quando	Semanas	Meses	Não sei
Rubéola  Quando  Semanas ou meses  Demças desconhecidal  Semanas ou meses  Não sei	Citomegalovírus (CMV)	•	Quando			Não sei
Augundo Semanas Meses Não sei  Semanas ou meses Semanas o	Rubéola	•	Quando			Não sei
Quando Semanas Meses Não sei  Quando Semanas ou meses Meses Não sei  Entre o mês anterior à gravidez e o final da gravidez, você teve alguma outra infecção que não citamos?  En caso afirmativo, registrar a semana de gravidez, se possível, e o(s) mês(es) de gravidez, caso a semana seja desconhecida] Semanas ou meses Despecifique: Diabetes (fora do período da gravidez) Diabetes durante a gravidez Doenças respiratórias Doenças neurológicas Doenças cardíacas Outro problema de saúde crônico Nenhum dos anteriores Não sei	Herpes	•	Quando			Não sei
Catapora  Quando  Quando  Semanas Meses  Não sei  LCMV (coriomeningite linfocitária)  Quando  Semanas ou meses  Semanas ou meses  Semanas Meses  Não sei  Semanas ou meses  Semanas ou meses  Semanas ou meses  Em caso afirmativo, registrar a semana de gravidez, se possível, e o(s) mês(es) de gravidez, caso a semana seja desconhecida]  Semanas ou meses  Doenças femanas ou meses  Semanas ou meses  Semanas ou meses  Doenças femanas ou meses  Semanas ou meses  Nemanas ou meses  Nemanas ou meses  Nemanas ou meses  Semanas ou meses  Semanas ou meses  Nemanas ou meses  Nem	Sífilis	•	Quando			⊚ Não sei
LCMV (coriomeningite linfocitária)  Semanas Meses Não sei  Entre o mês anterior à gravidez e o final da gravidez, você teve alguma outra infecção que não citamos?  Em caso afirmativo, registrar a semana de gravidez, se possível, e o(s) mês(es) de gravidez, caso a semana seja desconhecida]  Semanas ou meses  Semanas Meses  A. Você já foi diagnosticada com algum dos seguintes problemas de saúde?  Pressão alta  Diabetes (fora do período da gravidez)  Doenças respiratórias  Doenças neurológicas  Doenças cardíacas  Outro problema de saúde crônico  Nenhum dos anteriores  Se outra doença especificar?	Catapora	•	Quando			Não sei
Em caso afirmativo, registrar a semana de gravidez, se possível, e o(s) mês(es) de gravidez, caso a semana seja desconhecida]  Semanas ou meses  Semanas ou meses  Semanas ou meses  Meses  -4. Você já foi diagnosticada com algum dos seguintes problemas de saúde?  Pressão alta  Diabetes (fora do período da gravidez)  Doenças respiratórias  Doenças neurológicas  Doenças cardíacas  Outro problema de saúde crônico  Nenhum dos anteriores  Não sei	LCMV (coriomeningite	linfocitária) 🔻	Quando			⊚ Não sei
Especifique: Quando Semanas Meses  4. Você já foi diagnosticada com algum dos seguintes problemas de saúde?  Pressão alta Diabetes (fora do período da gravidez) Diabetes durante a gravidez  Doenças respiratórias Doenças neurológicas Doenças cardíacas  Outro problema de saúde crônico Nenhum dos anteriores Não sei  Se outra doença especificar?						
Pressão alta  Diabetes (fora do período da gravidez)  Doenças respiratórias  Doenças neurológicas  Doenças cardíacas  Nenhum dos anteriores  Se outra doença especificar?	Especifique:	Quando				
Doenças respiratórias  Doenças neurológicas  Outro problema de saúde crônico  Nenhum dos anteriores  Não sei  Se outra doença especificar?	4. Você já foi diagnost	 :icada com algum dos segui	intes problemas de sa	aúde? ———		
Outro problema de saúde crônico  Nenhum dos anteriores  Não sei  Se outra doença especificar?	Pressão alta		Diabetes (fora do p	período da gravid	ez) 🔲 Diabe	tes durante a gravidez
Se outra doença especificar?	Doenças respiratória	is [	Doenças neurológi	cas	Doen-	ças cardíacas
	Outro problema de	saúde crônico	Nenhum dos anter	iores	■ Não	sei
Se marcou alguma das doenças acima (respiratória, neurológica e cardíaca), especifique	Se outra doença espec	ificar?				
	Ge marcou alguma das o	doenças acima (respiratória	, neurológica e cardía	aca), especifique		

#### E. Medicamentos Agora, vou fazer perguntas sobre medicamentos que você pode ter tomado durante a gravidez. 1. Entre o mês anterior à gravidez e o final da gravidez, você tomou algum medicamento com ou sem prescrição? -Período -Medicamento 🔲 30 dias antes 📗 1º Tri Período -Medicamento 30 dias antes 1º Tri 2º Tri -Período -Medicamento 30 dias antes 1º Tri Período = Medicamento 30 dias antes 1º Tri 2º Tri Período – Medicamento 30 dias antes 2º Tri 1º Tri 2. Entre o mês anterior à gravidez e o final da gravidez, você tomou algum medicamento tradicional ou medicamento homeopático? Período -Medicamento 30 dias antes 1º Tri 2º Tri Período = Medicamento 30 dias antes 2º Tri 1º Tri =Período = Medicamento 30 dias antes 1º Tri 2º Tri ·Período -Medicamento 30 dias antes 1º Tri 2º Tri 3º Tri Período = Medicamento 30 dias antes 1º Tri 2º Tri 3. Entre o mês anterior à gravidez e o final da gravidez, você tomou alguma multivitamina, vitamina pré-natal ou suplemento de ácido fólico? Semanas ou meses Sulfato ferroso: Quando iniciou o uso Anterior Semanas Meses

Medicamento

30 dias antes 1º Tri 2º Tri 3º Tri

30 Tri

31. Entre o mês anterior à gravidez e o final da gravidez, você tomou alguma multivitamina, vitamina pré-natal ou suplemento de ácido fólico?

Semanas ou meses

Anterior Semanas ou meses

#### As próximas perguntas tratam do consumo de cigarros e álcool. 1. Entre o mês anterior à gravidez e o final da gravidez, você... Fumou cigarros? Por quanto tempo: Anterior Periodo Quantos por dia: 2. Entre o mês anterior à gravidez e o final da gravidez, você conviveu com alguém que ... Dentro de casa Fumou cigarros? Quanto: Anterior Periodo Quantos por dia: 2A. Fumou narguilê ou algo semelhante Dentro de casa: Quanto: Anterior Semanas ou meses Quantas horas por dia: 3. Entre o mês anterior à gravidez e o final da gravidez, você bebeu vinho, cerveja, bebidas destiladas, como cachaça, ou coquetéis de bebidas? Quanto: Anterior Semanas ou meses Frequência: G. Exposições ambientais Agora, vamos fazer perguntas sobre outras exposições que você pode ter tido durante a gravidez. Qual foi sua principal fonte de água para beber durante a gravidez? Torneira Aqueduto rural Agua mineral/agua filtrada Poço Rio ou lagoa Cisterna ou tanque Outra fonte Especificar Não sei O que? 2. Você faz alguma coisa para filtrar ou purificar a água que você bebe? 3. Você fez consumo de peixes e/ou frutos do mar durante a gestação? 4. Quanto tempo você ficou ao ar livre por dia durante a gravidez? 5. Você mantinha janelas e portas abertas durante o dia e noite quando estava grávida?

F. Exposições ao tabaco e álcool

6. Suas janelas e portas tinham telas protetoras?

7. Você usou repelente contra insetos quando estava ao ar livre durante a gravidez?

7. Durante a gravidez, voc de gravidez, caso a seman		rar a semana de gravidez, se possível, e o(s) mês(es)
Pesticidas 🔻	Especifique o(s) pesticida(s):	
Quando	Periodo  Semanas Meses Não sei	Frequência: ▼
Inseticida 🔻	Especifique o(s) inseticida(s):	
Quando	Periodo  Semanas Meses Não sei	Frequência: ▼
Raticidas 🔻	Especifique o(s) raticida(s):	
Quando	Semanas ou meses  Semanas Meses Não sei	Frequência:
Fertilizantes 🔻	Especifique o(s) fertilizante(s):	
Quando	Semanas ou meses  Semanas Meses  Não sei	Frequência:
Fumigação 🔻	Especifique o(s) produto(s):	
Quando	Semanas ou meses  Semanas Meses  Não sei	Frequência:
H. Avaliação do bebê		
Agora, vou fazer algumas	perguntas sobre a saúde do seu bebê.	
1. Em geral, como você cla	ssificaria a saúde do seu bebê?	▼
Caso :	seja regular ou ruim, explique:	
2. Desde que seu bebê nas	sceu, ele(a) apresentou algum dos seguintes pro	blemas?
Convulsões	▼ Febre ▼	Problemas de audição
Problemas de visão	→ Outro problema de saúde →	Especifique:
	demográficas e da residência	
Agora, gostaria apenas de	fazer as últimas perguntas sobre você e sua fam	
1. Como você classificaria	sua raça? ▼ Se out	tra, especifique:
2. Qual era a sua escolario	lade quando o bebê nasceu? (considerar o maior	nível completo)
3. Durante os 9 meses de	gravidez, qual era a renda mensal de sua família	9? ▼
4. Quantas pessoas eram	sustentadas por essa renda, inclusive adultos e c	crianças?

PC(cm) Estat	tura (cm)
Há alguma observação sobre o procedimen	to:
K. Observações finais e coleta das amost	
pouco de sangue para ver se você ou o seu b	to pela sua atenção em responder às nossas perguntas e nos fornecer um pebê foram infectados pelo vírus Zika. Sua contribuição para este estudo nos preender melhor a razão para tantos bebês estarem nascendo com microcefalia
🔳 1. Uma amostra de sangue foi colhida da	mãe?
🔲 2. Uma amostra de sangue foi colhida do	bebê?
3. O bebê foi fotografado?	
Observações finais	

J. Medidas antropométricas no momento da entrevista:

Form Approved OMB# 0929-1011 Expires 03/31/2017

Appendix 1: Case Investigation Form

Public reporting burden of this collection of information is estimated to average 75 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; ATTN: PRA (0920-1011).

This form is intended to interview patients with:					
<ul> <li>Isolates of any Elizabethkingia spp from a AND</li> </ul>	ny body site with PFGE matching outbreak pattern;				
The specimen was collected on or after N	lovember 1, 2015				
When initiating an interview, please use the script appropriate to a	participant. Please fill out completely, if patient or proxy does not				
know the information, then please check 'unknown' or note that qu	uestion was asked and information is not available.				
Was consent given: Yes No (DO NOT PROCEED)					
Contact Information					
atient contact information	Proxy contact information (if applicable):				
(gather at least State and Zip Code, even if proxy was interviewed):	Name:				
Name:	Relation to patient: Relative:				
Address:	☐ Clinician ☐ Other:				
City, State, Zip:	Address: Same as patient				
Phone: ( )					
Name of residence, if applicable (i.e. nursing home, assisted	City, State, Zip:				
living)	Phone: ( )				

CDCID\_\_\_\_\_

Date of interview://	
Zip code of residence:	
State Epi ID (state use only):	

Interview Information
Date first culture collected:/(MM/DD/YYYY)
First date of 30-day exposure period (date of culture collection – 30 days):/(MM/DD/YYYY)
First date of 7-day exposure period (date of culture collection – 7 days):/(MM/DD/YYYY)
Date interview completed://(MM/DD/YYYY)
The interview completed
Interviewer: Name:
Affiliation (state health dept. or CDC):
Linelist patient ID
For interviewer use only:
Information on this report was collected through (check all that apply): Patient/proxy interview (specify:)
Record Review  Review of health department notes  Other:
Must be filled BEFORE faxing to DPH:
Does this patient have laboratory-confirmation of Elizabethkingia spp infection?
Hollo Lam (name affiliation)
Hello, I am (name, affiliation).  Thank you for taking the time to talk to me today. Understanding healthcare and community exposures you had before you got sick with
Elizabethkingia is critical for identifying the source of these illnesses and stopping more people from getting sick. During this interview I will ask you
about your health, healthcare, and activities in the 30 days before you tested positive for Elizabethkingia. This is the period from (first date of
exposure period) to (date of culture collection) [if conducting in person interview, show a calendar]. To answer these questions, it might be helpful
for you to gather information that will help you remember what you did in the month before you became ill, such as an appointment diary, calendar,
statements from healthcare providers, and receipts from restaurants or travel. This is a standardized interview form we are using for all the patients
affected, to see if we can find some things in common that may have led to people becoming sick with this bacteria. We are still not sure the source of
this outbreak. This bacteria is very rare and relatively newly discovered, so there are a lot of things we don't know about it. What we do know is that
it likes to live in water and when it has infected people in the past that has typically been people who are already sick in the hospital. For that reason many of these questions will focus on prior healthcare exposures in the month before you became sick. I'll also be asking about home exposures,
including water and soil exposures. Then we'll also talk about food exposures.
The same of the second of the
$\underline{Instructions to the Interviewer-If the case is still only Possible and not Confirmed, besure to state:}$
"The state health department is automatically receiving any samples of this bacteria from hospital labs, and so we have been informed that you grew
this bacteria on(date) from(body location) We still need to check the DNA fingerprint of the sample we received and see if it matches the
same fingerprint of the other patients in this outbreak, and so we cannot confirm at this time that you are actually a part of the outbreak.  Nevertheless, we are trying to get ahead of things and start contacting anyone we think MIGHT be affected by the outbreak to ask some questions."
The verticless, we are a ying to get aread or a migs and start contacting anyone we a mink when his be affected by the outbreak to ask some questions.
Please remember that all of your responses are confidential. This interview will take up to an hour.
Are you ready to begin?
···- / · / · / · / · / · / · / /
Patient Provider (Patient interview or Medical Pacerd Paview)
Patient Provider (Patient interview or Medical Record Review)  1. Primary care provider name:
Location and phone number of Primary care provider:

Date of interview://
Zip code of residence:
State Epi ID (state use only):

ion (Medical Record Review and Patient Interview)	
e questions about your background and where you live.	
D Unk	on) did you stay at least one night in an institutional setting?  Assisted living facility Acute care facility Other, specify
☐ Male ☐ Female ☐ Unk	Unk
Patient Interview)	
· · · · · · · · · · · · · · · · · · ·	days before you became ill, the period from (first date of exposur
e collection). I will refer to this period as the month before	
you became ill, did you receive home health services (include each of these home health visits, starting with the most recese list:	ling wound checks, dressing changes, baths)? Yes No ent.
Agency Name and Contact Information	Reason for Visit
	with healthcare providers (this does not include outpatient visits for the such as cardiologists or oncologists, eye doctors, and dentists.
se list your appointments:	
Clinic Name (phone number and address, if known) and Specialty Type	Reason for Visit
you became ill, were you receiving outpatient dialysis?  t type of dialysis:  Hemodialysis  Peritoneal Dialysis contact information for dialysis facility:  lysis please specify access type:  fistula  graft  cent do you receive dialysis  MWF  T TH S Other  Us s session before symptoms onset. Date:	ral venous catheter  other  Unk
	e questions about your background and where you live.

Date of interview://
Zip code of residence:
State Epi ID (state use only):

Date	Hospital Na	ime and Contact Information		Reason for	ER Visit	How did you get to the hospital?
						EMS
						POV
						Other
						EMS
						POV
						Other
						EMS
						POV
						Other
						EMS
						POV
						Other
						☐ EMS ☐ POV
						Other
	L					1
-	een to an Urgent ( yes, please tell us	Care in the month prior to illno s more:	ess onset?	Yes	□No	□Unk
-	yes, please tell us			Yes	□ No  Reason for Urgent (	
a. If	yes, please tell us	s more:		Yes		
a. If	yes, please tell us	s more:		☐ Yes		
a. If y	yes, please tell us te Urg	s more:	nformation		Reason for Urgent (	Care Visit
a. If y Date  In the month  Yes  In the month  a. Ple	h before you because tell me all lo	gent Care Name and Contact II  gent Care Name and Contact II  ame ill, did you have an overni  Unk  ame ill, were you hospitalized ng term care facilities and hos	nformation  ight stay at a nu  overnight?	ursing home? Th	Reason for Urgent of the second secon	Care Visit  isted living facilities?
a. If y Date  In the month  Yes  In the month  a. Ple	yes, please tell us te Urg  h before you beca	gent Care Name and Contact II  gent Care Name and Contact II  ame ill, did you have an overni  Unk  ame ill, were you hospitalized ng term care facilities and hos	nformation  ight stay at a nu  overnight?	ursing home? Th	Reason for Urgent of the second secon	Care Visit  isted living facilities?
In the montl Yes  In the montl a. Ple	h before you because tell me all louding telescent tell me all louding type of Facility	gent Care Name and Contact II  ame ill, did you have an overni  Unk  ame ill, were you hospitalized ng term care facilities and hos dmissions).  Location (Address and	nformation  ight stay at a nu  overnight?	ursing home? Th	Reason for Urgent of the second secon	isted living facilities?  oth before you became ill (inc
In the montl Yes  In the montl a. Ple m	h before you because tell me all loudingle stays or according to the stays of the stays or according to the stays of the stay of t	ame ill, did you have an overni Unk ame ill, were you hospitalized ng term care facilities and hos dmissions).	ight stay at a nu overnight?	ursing home? Th	Reason for Urgent of the second secon	isted living facilities?
In the montl Yes  In the montl a. Ple	h before you because tell me all louding telescent tell me all louding type of Facility	gent Care Name and Contact II  ame ill, did you have an overni  Unk  ame ill, were you hospitalized ng term care facilities and hos dmissions).  Location (Address and	ight stay at a nu overnight?	ursing home? Th	Reason for Urgent of the second secon	isted living facilities?  oth before you became ill (inc
In the montl Yes  In the montl a. Ple	h before you because tell me all louding telescent tell me all louding type of Facility	gent Care Name and Contact II  ame ill, did you have an overni  Unk  ame ill, were you hospitalized ng term care facilities and hos dmissions).  Location (Address and	ight stay at a nu overnight?	ursing home? Th	Reason for Urgent of the second secon	isted living facilities?  oth before you became ill (inc

Date of interview://	
Zip code of residence:	
State Epi ID (state use only):	

16.	Have you received ca	re from any	of the following	in the month	prior to illness	onset?

Exposure	Yes	No	Unk	Location	Were any procedures beyond a routine examination performed?	Date(s) (MM/DD/YYYY)
Dentist					If yes, describe.	
Podiatrist						
Chiropractor						
Massage Therapist						
Naturopath						
In the month before y		Yes [	□No [		call that apply	

	a.		NO [								
	b.	Nebulizers Yes	No [	Unk							
	C.	Nasal sprays Yes	No [	Unk							
	d.	Eye drops	No [	Unk							
	e.	Oxygen Yes N	ol [	Unk							
	f. Over the counter supplements, including vitamins, probiotics, powders added to a drink or smoothie (e.g., protein powder)  Yes No Unk										
							orand	_			
	g.	Thickened juice, food, or sha i. If yes, which brand specify:		Unknown	ւԱր :k-I <sup>†</sup>	k *	ReadyCare 2.0 <sup>™</sup> Hori	me	l Thick & Easy®   Simply Thick   Oth		
		ii. If yes, specify type	of thi	ckener:					Unknown		
	h.	Proton pump inhibitors (PPI).							□ No □ Unk		
	i.	H2 blockers. Examples of H2	block	ers are Zantac and T	aga	me	et. 🗌 Yes 🗌 No 🔲 Ur	٦k			
	j.	Antibiotics Yes No		nk							
		i. If yes, check all tho	se tha	at were received:							
		Amikacin		Cefprozil			Doxycycline		Penicillin		
		Amoxicillin		Ceftazidime			Ertapenem		Piperacillin-Tazobactam		
		Amoxicillin/Clavulanic Acid		Ceftizoxime			Fosfomycin		Polymyxin B		
	/	Ampicillin/sulbactam		Ceftriaxone			Gentamicin		Rifampin		
	/	Azithromycin		Cefuroxime			Imipenem		Tetracycline		
		Aztreonam		Cephalexin			Levofloxacin		Ticarcillin/Clavulanic Acid		
		Cefaclor		Ciprofloxacin			Linezolid		Tigecycline		
		Cefazolin		Clarithromycin			Meropenem		Tobramycin		
		Cefdinir		Clindamycin			Metronidazole		Trimethoprim-Sulfamethoxazole		
	(	Cefepime		Colistin			Moxifloxacin		Vancomycin		
Ī		Cefotaxime		Daptomycin			Nitrofurantoin		Other (specify):		

Cefpodoxime

17.

Ofloxacin

Doripenem

Other (specify):

Date of interview://	
Zip code of residence:	
State Epi ID (state use only):	

Sign/Symptom	Present	If Yes, Date of Symptom Onset (MM/DD/YYYY) Write UNK if unknown	Notes (describe circumstances)	Treatment (include description of products used)
Open wounds, sores, or skin injury (i.e. ulcers, burns, cuts, or scrapes)	☐ Yes ☐ No ☐ Unk			
•	t through a catheter or during	•	on injections (i.e., intramuscular infusions)?	(IM), subcutaneous (SQ),
Medication	How	frequent are the injections?	D	ate of last injection?
				_
a. If yes, please to Medication/Vitami Substance (including s	in or	acility or Location (Address/F	Phone number)	Date(s) (MM/DD/YYYY)
heparin				
heparin				
In the month before you including a dialysis graft o	or fistula) 'ER: THIS LINE COULD HAVE E		atheters present? (for example DAYS, BUT MUST HAVE BEEN F	•
In the month before you including a dialysis graft of **NOTE FOR INTERVIEW TO ILLNESS.  Yes No Unk a. If yes, please tell us mo	or fistula) 'ER: THIS LINE COULD HAVE E			·

Date of interview://
Zip code of residence:
State Epi ID (state use only):

22.	2. In the month before you became ill, did you have any implanted medical devices (includes any device regardless of time placed)? (joint replacements, bone plates, cardiac defibrillator/pacer, heart valves, vascular stents, urinary catheter, etc.). Note: This does not include central or peripheral venous catheters which should be captured above  Yes No Unk  a. If yes, please tell us more:				
	Device Type	Device Location (note Left/Right if applicable)	Year Implanted		

Date of interview://
Zip code of residence:
State Epi ID (state use only):

	me Exposures (Patient Interview)				
Tha	Thank you for providing that information. Now I am going to ask you questions about potential exposures at home and in the community.				
23.	How long before you became ill did you live in your current home?months/years				
24.	In the month before you became ill, did you make any changes to your plumbing, heating, or cooling systems? Yes No Unk a. If yes, please explain:				
25.	In the <u>three</u> months before you became ill, were your plumbing, heating, or cooling systems serviced? Yes No Unk  a. If yes, please explain:				
26.	Describe the water supply used in the month prior to becoming ill? Private Well City or Municipal water (Specify municipality) Other, specify Unk				
27.	Does your home water use a de-chlorinator Yes				
	<ul><li>a. If yes, when was the filter last replaced prior to illness onset? Unk</li><li>b. Type of filter? Unk</li></ul>				
	Does your home water use a softener  Yes  No  Unk Where did you get your drinking water in the month before you became ill, check all that apply?				
	☐ Home Tap ☐ Point of Use Filter ☐ Bottled ☐ Other, specify ☐ ☐ Unk				
30.	In the month before you became ill, did you consume commercially bought ice? Yes No Unk  a. If yes, specify brand and location:				
31.	In the month before you became ill, did you use a humidifier in your home? Yes No Unk				
32.	In the month before you became ill, did you use a Neti-Pot or performed nasal rinsing?				
	a. If yes, what is the water source used? Unk				
33.	In the month before you became ill, did you have an aquarium in your home? Yes No Unk				
34.	In the month before you became ill, did you have any pets at home? Yes, specify No Unk				
35.	Did you have any exposure to animals in the 2 weeks before you became ill? Yes, specify No Unk				
36.	Did you have any plants in your home in the month before you became ill? Yes, specify No Unk				
	In the month before you became ill, did you have any contact with cut flowers?				
39.	How do you bathe, check all that apply?  Bath Shower Sponge bath Whirlpool Other, please specify Unk				
40.	Do you have dentures? Yes No Unk				
41.	What brand of toothpaste did you use in the month before you became ill? Unk				

Date of interview://
Zip code of residence:
State Epi ID (state use only):

	a. If yes, wh	nich brand? re you became ill, did	Unl you use any topica	al products (e.g., lotions	, creams, liniments, or ointments	
44.		•		•	edications? Yes No bu used during this period.	] Unk
45.		•		or bodywashes? \(\sigma\) ashes you used during t	Yes No Unk his period.	
46.		•		poos? Yes ed during this period.	□ No □ Unk	
47.	toothpaste, deodoi	rant)? Check all that	apply and specify I	ocation:	efore you became ill (e.g., soap, s	hampoo, creams and lotions,
	☐ Drug Store	Grocery Sto	ore 🔲	On-line	ier	
	Name			Location		
Ţ						
48.	In the month befor ☐ Yes ☐ I		I you use marijuana swer 🔲 Unk	a, also called cannabis,	mber that your responses are cor in any form? pical (oils)	nfidential.
49.		•	e or community tha	at has experienced a sir	milar illness? Yes No	] Unk
E0	a. If yes, wh		a manth hafara ya	u became ill? Yes		
50.	•	ch?	•	u became iii:		
51.	In the month befor	re you became ill, hav	e any pets in your	home had insect infesta	ations (i.e., fleas)? Yes	□ No □ Unk
52.		re you became ill, did he more about all the			our home? Yes No	
	Name	Phone Number	Relationship	Occupation	Place of Employment	Check if surveillance
						cultures were obtained

Date of interview://	
Zip code of residence:	
State Epi ID (state use only):	

Name	Phone Number	Relationship	County of Residence	Occupation	Place of Employment
					. L
Outside Exposure	(Patient Interview)				
'	,				
	efore you became ill did yo			that apply:	igious service Support Group [
	oup Club Social of				
5. If yes to any of	the above, please specify v	when and where _			
		ic i			
	<u>youwilibeconductingtheful</u>	<u>irooaexposureque</u>	stionnairewiththisp	atienttnenskipquest	tions56-58andproceedtoquestion
<u>9.</u> am now going to a	ck valusama allostians aha	out food you have	oaton which will fo	ocus only on the 7 da	ays before you became ill, which is the period
	date of culture) to (date	•		cus only on the <u>7 us</u>	before you became in, which is the period
rom (7 days prior to	date of cartal of to (date t	or curtare concern	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,		
66. In the 7 days be	fore you became ill, where	did your food con	ne from that was pr	epared at home	
•	-	S ☐ No ☐ Unk	·	•	
	i. If yes, specify 1			3	
b. F	armers market/food purch	ased directly from	farm \( \square\)	'es 🗌 No 🔲 Unk	
	i. If yes, specify 1	2		3	
c. H	lealth food store 🔲 Yes				
	<ol><li>If yes, specify 1</li></ol>			3	
d. E	thnic specialty market				
	i. If yes, specify 1			3	
e. F	ish or meat shop (e.g., but			_	
	i. If yes, specify 1			3	
f. H	lunting or fishing Yes			0	
	i. If yes, specify 1			3	
g. L	ocally grown fresh foods, e				
□ Not an	<ul><li>i. If yes, specify 1 olicable (my food is prepare</li></ul>	Z ad and controlled I	ny my rosidonco fac	ility)	
	fore you became ill, did far		, ,	J.	□ Yes □ No □ Unk
•	se describe:	•		iside the idenity.	
				ch as from a restaur	ant, meal delivery service, or at a school, or
•	cludes foods that you ate				•
	Restaurants (including deliv		-		
	i. If yes, specify 1	•			
	, , <u>,                                </u>		<u></u>		
b. N	Meal delivery service, such	as meals on wheel	s 🗌 Yes 🗌 No [	Unk	
	i. If yes, specify 1	2		3	
c. I	nstitution, such as hospital	or school 🔲 Yes	s □ No □ Unk		
	<ol> <li>If yes, specify 1.</li> </ol>	2		3.	

#### Elizabethkingia Spp.

				Elizabethkingia Spp. Interview Form:	Date of interview:// Zip code of residence: State Epi ID (state use only):
				use any consumables (lotions, balms, salves) prepared on onsume any foods from outside the U.S.?	
k you. Now I am goin posure period) to (dat	-	-		re questions about your activities. These questions w ).	ill refer to the full <u>30 day</u> period from (first d
n the month before ye	ou beca	ame ill,	did you	do any of the following activities?	
Exposure	Yes	No	Unk	Location	Date(s) (MM/DD/YYYY)
Swimming, hot tub					
Water aerobics					
Water park					
Fishing					
Steam room, or					
wet sauna					
Hot tub or					
whirlpool/spa					
Other:					
	Locat	ion		Date(s) (MM/DD/YYYY)	
Any additional comme	ents or I	notes (	e.g. trav	l details, additional visits to healthcare providers, othe	r diagnostic testing, and information)?

Date of interview:/ /
Zip code of residence:
State Epi ID (state use only):

 $\underline{Interviewer Instructions: If you will be conducting the full food exposure question naire proceed to it at this point. If you will not be conducting the full food exposure question naire then the interview is completed.}$ 

This is the end of the interview. Thank you very much for your time and willingness to provide this valuable information.

If you have any questions please feel free to contact Wisconsin Division of Public Health at 608-267-9003.

If necessary, would it be okay to contact you again in the future with any follow-up questions?

Thank you, and take care.

Interviewer: Please fax completed forms to 608-261-4976



Appendix 3: Case Series Form

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; ATTN: PRA (0920-1011).

Patient (	CDCID:	
Wiscon	sin Clinical Course Abstraction Form	
SECTIO	N A	
includes collecte culture	A Instructions: Abstract Section A using all medical records the medical record from the outpatient encounter, ER, or ed and follow-up care (hospitalization for EK, care at AL for ). If a patient was transferred during their EK hospitalization from both facilities.	r hospitalization where the positive culture was r EK, or outpatient treatment for the positive
Obtain	list of ICU locations from hospital prior to abstraction to e	ensure that all ICU stays are captured in item 21.
Patient	CDCID:	
Abstrac	ctor Name:	
Date of	Abstraction:	
Specify	name of facilities included in abstraction for Section A:	
	Age:	
	Sex: Male Female Unk	
3.	Race (check all that apply):  White Asian America  Native Hawaiian/Other Pacific Islander Unk	n Indian/Alaska Native 🗌 Black
4.	Ethnicity: Hispanic or Latino Not Hispanic or Latino [	☐ Unk
5.	County of residence:	
6.	Date of collection of first specimen positive for E. anophe	elis:/(MM/DD/YYYY)
7. 8.	Time of collection of first specimen positive for E. anophe Where was the first specimen positive for E. anophelis co	,
	☐ Inpatient	☐ LTCF/SNF
	☐ Emergency Room	LTACH
	Observational unit/Clinical Decision Unit	☐ Dialysis clinic
	Outpatient	Other (specify):
	☐ Assisted Living	Unknown

Patient CDCID:								
9.	9. Was patient hospitalized at the time of or during the 7 days after the first specimen positive for E. anophelis was collected? ☐ Yes ☐ No ☐ Unk							
	a. If yes, specify date of admission:/(MM/DD/YYYY) (If patient was hospitalized and transferred, specify first date of admission)   Unk							
	b. Was patient transported to the hospital by EMS/ambulance (this is only intended to capture emergency transports by EMS not planned transfers via private ambulance)? ☐ Yes ☐ No ☐							
	C.	Was patient transferred to a different short stay a specimen positive for E. anophelis was collected?						
		If yes, specify date(s) of transfer:/	(MM/DD/YYYY)					
	d.	List all admission diagnoses on H&P:						
10		nere was the patient admitted from? (select one)						
		Private residence	☐ LTCF/SNF					
		Acute care hospital inpatient	LTACH					
		Homeless	Other (specify):					
		☐ Assisted living	Unknown					
		etory of present illness at the time of hospital admis	ssion (please briefly summarize details from the					

□None □Unknown	
AIDS Atrial Fibrillation History of alcohol abuse Asplenia Asthma Autoimmune disease Cerebrovascular disease/stroke (except hemiplegia) Cerebral palsy Chronic cognitive deficit Chronic kidney disease (not on dialysis) Chronic liver disease without cirrhosis Cirrhosis Cirrhosis Chronic obstructive pulmonary disease (COPD)/emphysema Chronic lung disease (other than COPD/emphysema, asthma) Chronic steroid or other immunosuppressive therapy Chronic ventilation/tracheostomy Congenital heart disease	Failure to thrive
Congestive heart failure Connective tissue disease Cystic fibrosis Dementia Diabetes mellitus with complications Diabetes mellitus without complications Eczema	Solid tumor malignancy, not metastatic Spinal cord injury or paraplegia or quadriplegia Transplant, hematopoietic stem cell or bone marrow Transplant, solid organ Other Other Other
13. Did the patient have any of the following expos  a. Current smoking: ☐ Yes ☐ No ☐ Unk, if y	sures at time of hospital admission? yes, specify pack years: □ Unk if yes, specify drinks per week: □ Unk

nen was collected and icate the patient
If Yes, Date of Symptom Onset (MM/DD/YYYY) Write UNK if unknown
_
_
_

Yes No Unk

Other specify\_

Patient CDCID:						
16. Specify all physical exam findings	s documented by the clinical team on the date the first positive					
	No only if the records indicate the finding was not present (e.g., if team					
-	al then would check No for skin signs); otherwise specify unknown.					
Sign	Sign Present?					
Altered mental status	Yes No Unk					
Rash	Yes No Unk					
Skin redness	Yes No Unk					
Skin tenderness	☐ Yes ☐ No ☐ Unk					
Skin warmth	☐ Yes ☐ No ☐ Unk					
Skin wound (including decubitus u						
Cellulitis specifically documented	Yes No Unk					
Other specify	Yes No Unk					
Other specify	_ Yes No Unk					
Other specify	Yes No Unk					
Other specify	_ Yes □No □Unk □ Yes □No □Unk					
Other specifyOther specify	Yes No Unk					
Other specify	Yes No Unk					
отног оросну						
17. Vital signs documented closest to time of collection of first positive specimen  None Unknown  Date_/_/(MM/DD/YYYY) Unk  Time: (HOURS/MINUTES; 24 HOUR CLOCK) Unk						
Parameter (include units)	Result					
Systolic Blood pressure (mmHg)						
Diastolic Blood pressure (mmHg)						
Pulse (beats per minute)						
Respiratory rate (breaths per minute)						
Temperature (degrees F)						
Pulse Ox (percent)	Percent saturation					
	On O <sub>2</sub> Yes No Unk					
	If Yes, mode of delivery: Nasal cannula Intubated Other (specify):					
If yes, FiO2: or L/Min:(if FiO2 not documented)  Unk						

and the lowest.			
	Day 1 (Day culture was	Day 2	Day 3
	performed)	Date://	Date://
	Date:/		
Highest systolic blood pressure			
Lowest systolic blood pressure			
Highest heart rate			
Lowest heart rate			
Highest respiratory rate			
Lowest respiratory rate			
Highest WBC			
Lowest WBC			
Highest proportion bands			
Altered mental status present			
(Yes/No/Unknown)			

19. Complete supplementary Table 1. For each day of the patient's hospital admission, record vitals

collection date through the duration of their hospitalization.

documented closest to 6am and 6pm in the medical record. If the patient was already hospitalized at the time their first positive specimen was collected, document vitals starting 7 days prior to specimen

18. Record each for the 3 days beginning with the day of collection of first positive specimen. If only 1 value was documented (e.g., only 1 wbc value recorded for a given day) record the value as both the highest

Patient CDCID: \_\_\_\_

Patient CDCID:					
		nted as specified below			
Parameter (include units)	Results on day patient was admitted to hospital (if multiple results, select first collected)  Date: _/_/	Results on day first positive specimen was collected (select results closest to specimen collection)  Date: _/_/	Results of highest value obtained during hospitalizatio n  Specify date for each value	Results of lowest value obtained during hospitalization  Specify date for each value	Results on day discharged from hospital or died  Date:
WBC	<u> </u>				
Percent neutrophils (corresponds to WBC count above)					
Percent bands (corresponds to WBC count above)					
Platelets					
Hematocrit BUN					
Creatinine					
AST ALT					
INR					
Alkaline phosphatase					
Bilirubin					
Glucose					
CRP					
Anion Gap: Sodium – (Chloride + Bicarbonate)					

21. Was the patient admitted to an intensive care unit during his/her stay? $\ \ \ \ $	□No	Unk
If yes, specify dates of admission to ICU:		

	Date of ICU Admit			Date of ICU Discharge		
1	/_	/	Unk	/_	/	Unk
2	/_	/	Unk	/_	/	Unk
3	/_	/	Unk	/_	/	Unk
4	/_	/	Unk	/_	/	Unk

22. Specify if any of the following procedures were performed or provided during the patient's hospitalization:

Procedure	Performed?	Date of Procedure  Indicate Start Date for those procedures where number of days is documented	If yes, describe indication for procedure	Number of Days (do not fill if box is greyed)
Placement of Chest tube	☐ Yes ☐ No ☐ Unk	//		
Placement of other drain Specify	☐ Yes ☐ No ☐ Unk	_//_		
Acute hemodialysis	☐ Yes ☐ No ☐ Unk	_//_		
Mechanical ventilation	Yes No Unk	_//_		
Noninvasive ventilation (CPAP or BiPAP)	☐ Yes ☐ No ☐ Unk	//		
Placement of Central Venous Catheter	☐ Yes ☐ No ☐ Unk	_//_		
Bronchoscopy	Yes No Unk	//		
Endoscopy Specify	☐ Yes ☐ No ☐ Unk	_//_		
Surgery Specify	☐ Yes ☐ No ☐ Unk	//		
Other Specify	☐ Yes ☐ No ☐ Unk	//		
Other Specify	☐ Yes ☐ No ☐ Unk	//		
Other Specify	☐ Yes ☐ No ☐ Unk	_//_		

23. Did the pat stop dates:	ient require vasopressors? Ye	s No Unk If	yes, specify wh	nich ones and all	start and
List all vaso	pressors that were started at an r was stopped and then restarte		-		
Examples: (specify)	Dopamine, Dobutamine, Epinepl	nrine, Norepinephrin	e, Neosyneph	rine, Vasopressir	n, Other
Vasopressor	Start d	ate	Stop date	<del></del>	
•	/	/	/	/	
		<u> </u>		<del></del>	
		<u></u>	<u> </u>	<u>'</u>	
		<u></u>		<u></u>	
			/	_/	
			/		
	/_	/	/	_/	
Antimicrobial	Route (IV, IM, PO, Topical, Inhaled)	Start date		Stop date	
	IV IM PO Topical Inhaled	_//_	Unk	/	/_ Unl
	IV IM PO Topical Inhaled	//	Unk	/	/_ Unl
	☐ IV ☐ IM ☐ PO ☐ Topical ☐ Inhaled	//	Unk	/_	/_ Unl
	IV IM PO Topical Inhaled	_//_	Unk	/_	/_ Unl
	IV IM PO		Unk	/	/ Uni
	☐ IV ☐ IM ☐ PO☐ Topical ☐ Inhaled☐ IV ☐ IM ☐ PO☐ INHALED☐ IV ☐ IM ☐ PO☐ INHALED☐ IN		Unk		/ Un
	Topical Inhaled		Unk		/ Uni
	Topical Inhaled	/ /	Unk		/ Unl
	Topical Inhaled	/ /	Unk		
	Topical Inhaled	<del>''</del>		—' ——' ——	

Patient CDCID: \_\_\_\_\_

Patient CDCID:	_		
25. Were antibiotics p a. If yes, specify:	rescribed for patient at discha	arge? ☐ Yes ☐ No ☐ I	Jnk
Antimicrobial		Route (IV, IM, PO, Topical)	Prescribed Duration in Days (specify unknown it not documented)
		IV IM PO Topical Inhaled IV IM PO IV IM PO Topical Inhaled	
		IV IM PO Topical Inhaled	
Imaging  26. List all imaging stu  Note: Do not include r	dies and results results from X-Rays except Ch	nest X-Ray as described in ite	m 27
Performed	Location	Impression	Date
CT MRI Ultrasound Nuclear Medicine Other (specify):	Head Abdomen Chest Extremity (specify): Other (specify):		//
CT MRI Ultrasound Nuclear Medicine Other (specify):	Head Abdomen Chest Extremity (specify): Other (specify):		_//
CT MRI Ultrasound Nuclear Medicine Other (specify):	Head Abdomen Chest Extremity (specify): Other (specify):		_//
CT MRI Ultrasound Nuclear Medicine Other (specify):	Head Abdomen Chest Extremity (specify): Other (specify):		/

Patient CDCID:					
CT MRI Ultrasound Nuclear Medicine Other (specify):	Head Abdomen Chest Extremity (specify):  Other (specify):		//		
CT MRI Ultrasound Nuclear Medicine Other (specify):	Head Abdomen Chest Extremity (specify): Other (specify):		//		
CT MRI Ultrasound Nuclear Medicine Other (specify):	Head Abdomen Chest Extremity (specify): Other (specify):		//		
CT MRI Ultrasound Nuclear Medicine Other (specify):	Head Abdomen Chest Extremity (specify): Other (specify):		//		
27. Chest X-Ray (Document findings from the X-ray that was performed most proximal to the time the first positive specimen was collected; only Chest X-rays on day of positive culture or the 2 days following)    Yes					
Were any of the following n radiologist. Please also doc		nly check items that were expression above)?	licitly documented by the		
Pleural effusion Inf	iltrate Consolidation [	Bronchopneumonia/pneum	nonia		
☐ No evidence of pneumo	onia 🗌 Cannot rule out pne	monia			

Patient CDCID:			
Procedures			
28. Did patient have a lu	umbar puncture?	☐ No ☐ Unk	
If yes, record results for eac	h lumbar puncture performe	d (include units)	
Date://	Unk		
Parameter	First tube	Subsequent tube	
WBC count			List differential
RBC count			
Protein			
Glucose			
Opening pressure			
Gram stain			
Date:/	Unk		
Parameter	First tube	Subsequent tube	
WBC count		·	List differential
RBC count			
Protein			
Glucose			
Opening pressure			
Gram stain			
29. Did patient have pa	racentesis? Yes Nh paracentesis performed (ii		
Date:/	Unk		
Parameter	First tube		
WBC		Differential:	
RBC			
LDH			
Protein			
Gram stain			
			'
Date:/	Unk		
Parameter	First tube		
WBC		Differential:	
RBC			
LDH			
Protein			
Gram stain			

Patient CDCID:	<u></u>	
30. Did patient hav	re thoracentesis?  Yes	☐ No ☐ Unk
If yes, record results fo	r each thoracentesis perf	ormed (include units)
Date://	Unk	
Parameter	First tube	
WBC		Differential:
RBC		
LDH		
Protein		
рН		
Gram stain		
Date://	Unk	
Parameter	First tube	
WBC		Differential:
RBC		
LDH		
Protein		
рН		
Gram stain		
If yes, record results fo	e joint aspiration? Ye	
Date://	Offic	
Parameter	First tube	
WBC		Differential:
RBC		
LDH		
Protein		
Gram stain		
Date://	Unk	
Parameter	First tube	
WBC		Differential:
RBC		
LDH		
Protein		
Gram stain		

	Date			Result		If blood culture is a "set"
(specimen type)						of cultures specify total number of bottles and
						the number positive
_	_/_	_/	Unk	No gr		Unl
				Organ	nism specify	
_	_/_	_/	Unk	☐ No gr		Unl
				│	nism specify	
_	/	_/	Unk	☐ No gr		Un
				Orgar	nism specify	
	_/_		Unk	☐ No gr	owth	☐ Uni
				Organ	nism specify	
	/	/	Unk	│ No gr	owth	☐ Unl
				Organ	nism specify_	
	/	_/	Unk	☐ No gr	owth	☐ Un
					nism specify	_
33 For F anonhelis isola						
medical record for fi	-	-	e dian	neter and i	nterpretation (i	f available) listed in the
medical record for fi	-	-		neter and in	Zone	f available) listed in the  Interpretation
medical record for fi	-	ate				Interpretation
medical record for fi Isolate collection date Piperacillin/tazobactam	irst isol	ate			Zone	Interpretation  S I R Non
medical record for fi Isolate collection date Piperacillin/tazobactam	irst isol	Date  //_ Unk //			Zone	Interpretation  S I R Non
medical record for fi  Isolate collection date  Piperacillin/tazobactam  Trimethoprim/sulfamethox	irst isol	Date			Zone	Interpretation  S I R Non
medical record for fi Isolate collection date Piperacillin/tazobactam Trimethoprim/sulfamethox Levofloxacin	irst isol	Date  //_ Unk //			Zone	Interpretation  S I R None S I R None
medical record for fi Isolate collection date Piperacillin/tazobactam Trimethoprim/sulfamethox Levofloxacin	irst isol	Date //Unk// Unk// Unk//			Zone	Interpretation  S I R Non
medical record for fi Isolate collection date Piperacillin/tazobactam Trimethoprim/sulfamethox Levofloxacin Ciprofloxacin	irst isol	Date //Unk//Unk//Unk//			Zone	Interpretation  S I R Non  S I R Non  S I R Non  S I R Non
medical record for fi Isolate collection date Piperacillin/tazobactam Trimethoprim/sulfamethox Levofloxacin Ciprofloxacin Moxifloxacin	irst isol	Date //Unk// Unk// Unk//			Zone	Interpretation  S I R Non
medical record for fi  Isolate collection date  Piperacillin/tazobactam  Trimethoprim/sulfamethox  Levofloxacin  Ciprofloxacin  Moxifloxacin  Rifampin	irst isol	Date // Unk// Unk// Unk// Unk// Unk// Unk			Zone	Interpretation  S I R Non
medical record for fi  Isolate collection date  Piperacillin/tazobactam  Trimethoprim/sulfamethox  Levofloxacin  Ciprofloxacin  Moxifloxacin  Rifampin	irst isol	Date //Unk// Unk//Unk//Unk//Unk//Unk// Unk			Zone	Interpretation  S I R Non
medical record for fi Isolate collection date Piperacillin/tazobactam Trimethoprim/sulfamethox Levofloxacin Ciprofloxacin Moxifloxacin Rifampin Vancomycin	irst isol	Date // Unk// Unk// Unk// Unk// Unk// Unk			Zone	Interpretation  S I R Non
·	irst isol	Date //Unk// Unk//Unk//Unk//Unk//Unk// Unk			Zone	Interpretation  S I R Non  S I R Non

Patient CDCID: \_\_\_\_\_

Patient CDCID:			
Other (specify):			S I R None
Other (specify):	Unk//		S I R None
Other (specify):	Unk //		S I R None
Other (specify):	Unk //		S I R None
· · · · · ·	Unk		
b. How was organism report  Outcomes (at end of discharge for a second s	Died Discharged S  //_ discharged:// e 30 days after discharge from Yes No Unk	Species are hospital): still inpatient at time o Unk	• • • • • • • • • • • • • • • • • • • •
Date of death:	// Unk		
Cause(s) of dea	th listed on death certificate	in the order they are	listed:
	_		
	re facility Long-term acu	te care hospital 🔲 A	ssisted living Unk
Other, specify			
36. Specify all discharge dia	gnoses in the order listed in	the medical record:	
37. Was patient readmitted	l to acute care hospital withi	in 30 days from discha	rge from the hospital?
Yes No Unk			
·	on	_	
<ul> <li>b. If yes, indication for rea</li> </ul>	dmission		

Patient CDCID:							
SECTION B							
•	Compete Section B using all available records from the 30 days prior to the collection date of the first culture positive specimen to complete the following section.						
Did the patient rece specimen?	Did the patient receive any of the following medications in the 30 days prior to collection of the first positive specimen?						
38. H2 blocker:	Yes No Unk						
	If yes, were H2 blockers listed as outpatient medication in the HPI from the time of first positive culture collection?   Yes   No  Unk						
39. Proton pum	p inhibitor: Yes No Un	k					
	PI listed as outpatient medication in t	he HPI from the time of first	positive culture collection?				
40. Were antibi provider no Yes No		s prior to collection of positi	ve culture, based on				
Do not include t	hose already documented in item 24	above.					
If yes, specify:							
Antimicrobial	Route	Start date	Stop date				

	Route	Start date	Stop date
☐ Amikacin	□ IV □IM □PO □Topical □nhaled	/Unk	/Unk
☐Amoxicillin	□ IV □IM □PO □Topical □nhaled	/Unk	/Unk
	□ IV □IM □PO □Topical □nhaled	//Unk	/Unk
Amoxicillin/Clavulanic			
Acid			
	□ IV □IM □PO □Topical □nhaled	/Unk	/Unk
Ampicillin/sulbactam			
Azithromycin	□ IV □IM □PO □Topical □nhaled	/Unk	/Unk
Aztreonam	□ IV □IM □PO □Topical □nhaled	/Unk	/Unk
Cefaclor	□ IV □IM □PO □Topical □nhaled	/Unk	/Unk
Cefazolin	□ IV □IM □PO □Topical □nhaled	/Unk	/Unk
Cefdinir	□ IV □IM □PO □Topical □nhaled	/Unk	/Unk
Cefepime	□ IV □IM □PO □Topical □nhaled	/Unk	/Unk
☐ Cefotaxime	□ IV □IM □PO □Topical □nhaled	/Unk	/Unk
Cefpodoxime	□ IV □IM □PO □Topical □nhaled	/Unk	/Unk
☐ Cefprozil	□ IV □IM □PO □Topical □nhaled	/Unk	/Unk
Ceftazidime	□ IV □IM □PO □Topical □nhaled	/Unk	/Unk
Ceftizoxime	□ IV □IM □PO □Topical □nhaled	/Unk	/Unk
Ceftriaxone	□ IV □IM □PO □Topical □nhaled	/Unk	/Unk
Cefuroxime	□ IV □IM □PO □Topical □nhaled	/Unk	/Unk
Cephalexin	□ IV □IM □PO □Topical □nhaled	/Unk	/Unk
Ciprofloxacin	□ IV □IM □PO □Topical □nhaled	//Unk	/Unk
Clarithromycin	□ IV □IM □PO □Topical □nhaled	/Unk	/Unk
Clindamycin	□ IV □IM □PO □Topical □nhaled	/Unk	/Unk
Colistin	□ IV □IM □PO □Topical □nhaled	/Unk	/Unk
□ Daptomycin	□ IV □IM □PO □Topical □nhaled	/Unk	/Unk
Doripenem	□ IV □IM □PO □Topical □nhaled	/Unk	/Unk
Doxycycline	□ IV □IM □PO □Topical □nhaled	/Unk	/Unk
Ertapenem	□ IV □IM □PO □Topical □nhaled	/Unk	/Unk
Fosfomycin	IV IM PO Topical Inhaled	/Unk	/Unk

Patient CDCID:	
----------------	--

Gentamicin	□IV □IM □PO □	Topical Inhaled	//	Unk	//	Unk
☐ Imipenem	□IV □IM □PO □	Topical nhaled	//	_ Unk	/	Unk
Levofloxacin	□IV □IM □PO □	Topical nhaled	//	Unk	//	Unk
Linezolid	□IV □IM □PO □	Topical Inhaled	//	Unk	/	Unk
Meropenem	□IV □IM □PO □	Topical nhaled	//	_ Unk	/	Unk
Metronidazole	□IV □IM □PO □	Topical Inhaled	//	_ Unk	/	Unk
Moxifloxacin	□IV □IM □PO □	Topical Inhaled	//	_ Unk	//	Unk
Nitrofurantoin	□IV □IM □PO □	Topical Inhaled	//	_ 🗌 Unk	//	Unk
Ofloxacin		Topicalnhaled	//	Unk	/	Unk
Penicillin		Topical nhaled	//	_ 🗌 Unk	/	Unk
☐ Piperacillin-	□ IV □ IM □ PO □	Topical nhaled	//	_ 🗌 Unk	//	Unk
Tazobactam						
Polymyxin B		Topical Inhaled	//	Unk	/	Unk
Rifampin		Topical Inhaled	//	Unk	//	Unk
Tetracycline		Topical nhaled	//	_ Unk	/	Unk
	□IV □IM □PO □	Topical Inhaled	//	_ 🗌 Unk	//	Unk
Ticarcillin/Clavulanic						
Acid						
Tigecycline		Topical Inhaled	//	_ Unk	//	Unk
Tobramycin		Topical Inhaled	//	_ Unk	//	Unk
☐ Trimethoprim-	□IV □IM □PO □	Topical Inhaled	//	_ Unk	/	Unk
Sulfamethoxazole						
Vancomycin		Topical Inhaled	//	Unk	//	Unk
Other	□IV □IM □PO □	Topical Inhaled	//	Unk	/	Unk
(specify):						
		🗖	, ,		, ,	
Other	□IV □IM □PO □	Topicalnhaled	//	Unk	/	Unk
(specify):						
Other	□IV □IM □PO □	TopicalInhaled	/ /	Unk	/ /	Unk
(specify):		Topical Linnaled	//	_ LI OLIK	'	.⊔UIK
(apecity).						
Other		Topical Inhaled	/ /	□Unk	/ /	□Unk
(specify):		Topical Littlated	//		''	
(opcony)						
	l					

Form Approved
OMB# 0929-1011
Expires 03/31/2017

Appendix 2: Medical Abstraction Form

Public reporting burden of this collection of information is estimated to average 75 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; ATTN: PRA (0920-1011)

Line list patient l	e list patient ID (CDCID) Wisconsin State Laboratory of Hygiene ID Abs			
This form is in	ntended to be u	sed for abstraction of medical rec	ords for patients in \	Nisconsin with:
	AND	f <i>Elizabethkingia</i> spp cultured fror on or after November 1, 2015	n sterile sites,	
Patient NAMI	E:			
Patient DOB:	//	(MM/DD/YYYY)		
		<del>-</del>		
Abstraction I				
Date medical	record abstract	ion completed:/ (N	/IM/DD/YYYY)	
Abstractor:		te health dept. or CDC):		

Line list patient ID (CDCID) \_\_\_\_\_ Wisconsin State Laboratory of Hygiene ID \_\_\_\_\_ Abstractor Initials\_\_\_\_

Line list patient ID (CDCID) Wisconsin State Laboratory of Hygie	ne ID Abstractor Initials					
SECTION 1: Case Background Information. Complete this section using the Case Report Form  1. Date positive culture collected:/ (MM/DD/YYYY) Time positive culture collected (24 hour): HH:MM  2. First date of 30-day exposure period (date of culture collection – 30 days):/ (MM/DD/YYYY)  3. First date of 7-day exposure period (date of culture collection – 7 days):/ (MM/DD/YYYY)  4. Name of facility where first positive culture was collected:						
LOCATION OF CULTURE COLLECTION:    Inpatient						
<ul> <li>5. Was this collected more than 3 calendar days after admission: ☐ Yes ☐ No</li> <li>6. Where was patient <b>residing</b> at time of culture collection:</li> </ul>						
☐ Private residence ☐ Acute care hospital inpatient ☐ Homeless ☐ Assisted living	☐ LTCF/SNF ☐ LTACH ☐ Other ☐ Unknown					
<ol> <li>Was the patient hospitalized for <i>Elizabethkingia</i> infection?  Yes No</li> <li>If yes to 5: Admission date:  (MM/DD/YYYY)</li> <li>If culture was not collected in a hospital facility, what was the reason for collected in a hospital facility.</li> </ol>	Admission time(24 hour):HH:MM culture?					

Line list patient ID (CDCID)	Wisconsin State Laboratory of Hygiene ID	Abstractor Initials
. , ,		

10. Please list all known medical encounters in 30 days prior. Medical records should be requested from each of the listed facilities.

HC	Date of Health			
Encounter #	Care	Emagyintar lagation	Type of encounter	Record or interview included, Yes or No
	Encounter	Encounter location	<b>5.</b>	
1		outpatient clinic		
		home health		
		☐ EMS ☐ emergency room (no admy to ACH)		
		ACH (admission )		
		Assisted living		
		☐ LTCF ☐ LTACH		
		☐ Dialysis		
		☐ Dental		
		Other If admitted:		
		Dates//		
2		outpatient clinic		
		home health EMS		
		emergency room (no admy to ACH) ACH (admission)		
		ACH (admission )		
		☐ Assisted living ☐ LTCF		
		□LTACH		
		☐ Dialysis		
		☐ Dental ☐ Other		
		If admitted:		
		Dates//		
3		outpatient clinic		
		home health EMS		
		emergency room (no admy to ACH)		
		ACH (admission )		
		Assisted living		
		LTCF LTACH		
		☐ Dialysis		
		☐ Dental ☐ Other		
		If admitted:		
		Dates// outpatient clinic		
4		outpatient clinic home health		
		☐ EMS		
		emergency room (no admy to ACH)		
		ACH (admission )		
		Assisted living LTCF		
		□LTACH		
		Dialysis		
		☐ Dental ☐ Other		
		If admitted:		
		Dates/		
5		outpatient clinic home health		
		☐ EMS ☐ emergency room (no admy to ACH)		
		emergency room (no admy to ACH)		
		ACH (admission ) Assisted living		
		LTCF		
		□LTACH		
		☐ Dialysis ☐ Dental		
		Other		
		If admitted:		
		Dates//		

Line list patient ID (CDCID) Wisconsin State La	Abstractor Initials				
SECTION 2: Overall Medical History. Complete th	is section using <u>all available</u> medical records				
Medical History					
11. Females only: Were you pregnant or ≤6 weeks postpart	um at the time of first positive EK culture?				
Yes, pregnant (weeks pregnant at onset)	Yes, postpartum (delivery date)/ No Unk				
12. Height (use record closest to EK positive culture)	_ftincm				
13. Weight (use record closest to EK positive culture)	lbkg				
14. BMI					
15. Did the patient have any of the following medical condit	· · · · · · · · · · · · · · · · · · ·				
Coronary artery disease	Connective Tissue Disease				
Atrial Fibrillation	Malignant Lymphoma				
Congestive Heart Failure	Solid Tumor				
Peripheral Vascular Disease	Mild Liver Disease				
Cerebrovascular Disease	HIV without AIDS				
☐ Dementia	AIDS				
Chronic Obstructive Pulmonary Disease (COPD)	History of decubitus ulcers				
Pulmonary Hypertension	Cellulitis				
Peptic Ulcer Disease	Pancreatitis				
Diabetes Mellitus without complications	Current alcohol dependence				
☐ moderate or severe renal disease ☐ Inflammatory bowel disease(Ulcerative Colitis/Crohns)					
Hemiplegia	Smoking (previous year)				
Hematologic Malignancy	Solid organ transplant				
Moderate or severe liver disease	Asthma				
Diabetes mellitus with end organ damage	Other				
Dialysis					
16. Did the patient get dialysis in the <b>30 days</b> prior to positive	<del>_</del> _ <del>_</del>				
a. What type of dialysis was performed? 🗌 F					
b. Does the patient have permanent vascular					
c. What type of vascular access was used?					
d. Did the patient get this type of dialysis in the <b>7 days</b> prior to positive EKM culture? 🔲 Yes 🔲 No 🗍 Unk					
i. Date of most recent dialysis	<del></del>				
	ter (Name)				
_ • •	Department (Name)				
	e: peritoneal dialysis is usually done at home				
Other:					
17. Did the patient have CRRT in the <b>30 days</b> prior to positiv	vo EV culturo2 🗆 Voc 🗀 NO				
	<u> </u>				
e. Date f. Name of facility:					
i. Name of facility	<del></del>				

Line list patient ID (CDCID)	Wisconsin Sta	ite Laboratory o	f Hygiene ID	Abstrac	tor Initials
18. Did the patient receive any IV If yes, complete the table bel		the 30 days to	2 hours prior to pos	sitive culture? 🗌 Yes	□No
Medication Name	Facility Name	Route (IV, IM)	Most Recent Date Prior to Positive Cx (MM/DD?YYY)	Approximate start date	Comments
19. Did the patient have surgery	in the 30 days prior to	positive EK cult	rure?  Yes	] No	
Type of Surgery	Date (MM/DD	/YYY)	Facility	Name	
20. Did patient have a wound in a. Wound type (de	the thirty days prior to escription and location				
b. Topical treatme	nts received:				

Line list patient ID (CDCID) Wisconsin State Laboratory of Hygiene ID Abs  21. Did patient have any indwelling devices present in the 30 days prior  Yes  No					stractor Initials	
Device	Present at EK Cx		Details			Placed in last 30 days?
Cardiac Pacemaker/ICD	Yes No	Unk				Yes No Unk
Cardiac Defibrillator	Yes No	Unk				Yes No Unk
Prosthetic Cardiac Valve	Yes No	] Unk				Yes No Unk
Vascular Stent	Yes No	] Unk	cardiac other_	peripheral		Yes No Unk
Vascular grafts	Yes No	Unk	cardiac	aortic other_		Yes No Unk
Indwelling vascular catheter	Yes No	] Unk	Port other	Picc HD permca	ath	Yes No Unk
Urinary Catheter	Yes No	Unk	ourier			Yes No Unk
Prosthetic joint	Yes No	Unk	Location:			Yes No Unk
Orthopedic implants	Yes No	Unk				Yes No Unk
(plates/screws)						
Other Implant1	Yes No	] Unk				Yes No Unk
Other Implant2	Yes No	] Unk				Yes No Unk
22. Has patient used any immunosuppressant medications in the last 30 days: Yes No						
Immunosuppressant		In 30 days	prior to Cx?	Medication name	Date of mos	t recent administration
Corticosteroid (e.g. Predisone	>20mg daily)	Yes	☐ No			
Biologics		Yes	☐ No			
Chemotherapy		Yes	☐ No			
Radiation		Yes	☐ No			
Other1		Yes	☐ No			
Other2		Yes	□No			

<sup>\*</sup>examples of common biologics include Humira (adalimumab), Remicade (infliximab), Rituxan (rituximab), Enbrel (etanercept), or other medications ending in –mab or –cept

## Line list patient ID (CDCID) \_\_\_\_\_ Wisconsin State Laboratory of Hygiene ID \_\_\_\_\_ Abstractor Initials\_ 23.Culture Data: Complete for <u>all cultures</u> collected 7 days prior to positive EK culture (<u>EXCEPT EK POSTIVE CULTURES</u>)

Culture No.	Specimen		Collect date (mm/dd/yy) Time (HH:MM)	Pathogens identified
1	Blood Urine  BAL CSF Ascites	Pleural fluid Synovial fluid Wound Stool Sputum	//	Path1Path3
2	Blood Urine  BAL CSF Ascites	Pleural fluid Synovial fluid Wound Stool Sputum	//	Path1Path3
3	Blood Urine  BAL  CSF  Ascites	Pleural fluid Synovial fluid Wound Stool Sputum	//	Path1Path3
4	Blood Urine  BAL CSF Ascites	Pleural fluid Synovial fluid Wound Stool Sputum	//	Path1Path3
5	Blood Urine  BAL CSF Ascites	Pleural fluid Synovial fluid Wound Stool Sputum	//	Path1Path3
6	Blood Urine  BAL CSF Ascites	Pleural fluid Synovial fluid Wound Stool Sputum	//	Path1Path3
7	Blood Urine BAL CSF Ascites	Pleural fluid Synovial fluid Wound Stool Sputum	//	Path1Path3
8	Blood Urine BAL CSF Ascites	Pleural fluid Synovial fluid Wound Stool Sputum	//	Path1Path3
9	Blood Urine BAL CSF Ascites	Pleural fluid Synovial fluid Wound Stool Sputum	//	Path1Path2Path3
10	Blood Urine  BAL CSF Ascites	Pleural fluid Synovial fluid Wound Stool Sputum	//	Path1Path3

Line list	patient ID (	CDCID	)
Line in	puncin in ,	CDCL	<i>'</i>

Abstractor Initials\_

Line list patient ID (CDCID) \_\_\_\_\_ Wisconsin State Laboratory of Hygiene ID \_\_\_\_\_ 24.Culture Data: Complete for all *positive EK cultures* 

Culture No.	Specimen		Collect date (mm/dd/yy) Time (HH:MM)	Comments
1	☐Blood ☐Urine ☐BAL	☐ Pleural fluid ☐ Synovial fluid ☐ Wound	//	
	☐CSF ☐Ascites	☐Stool ☐Sputum	<u></u> :	
2	☐Blood ☐Urine ☐BAL	☐Pleural fluid ☐Synovial fluid ☐Wound	//	
۷	☐CSF ☐Ascites	Stool Sputum	<u></u> :	
3	☐Blood ☐Urine ☐BAL ☐CSF	☐ Pleural fluid ☐ Synovial fluid ☐ Wound ☐ Stool	//	
	Ascites	Sputum	:	
4	☐Blood ☐Urine ☐BAL ☐CSF	☐ Pleural fluid ☐ Synovial fluid ☐ Wound	//	
	☐ Ascites	☐Stool ☐ Sputum	:	
5	☐Blood ☐Urine ☐BAL	☐Pleural fluid ☐Synovial fluid ☐Wound	//	
	☐CSF ☐Ascites	Stool Sputum	:	

Line list patient ID (CDCID) Wiscons	Wisconsin State Laboratory of Hygiene ID Abstractor Initials					
Section III: Visit with the positive culture.  Location of visit: acute care hospital LTAG  Skilled Nursing Facility Urgent Ca  Date of symptom onset (for positive culture):	re Other	<del>-</del>				
25. Chief Complaint (i.e. what were the patie	nt's symptoms) at time of positive culture:					
	Approximate Start Date (MM/DI	D/YYY)				
Abdominal Pain						
Altered Mental Status						
Chest Pain						
Cough						
Diarrhea						
Dysuria						
Facial Droop						
Fever						
Hyperglycemia						
Rash/Redness						
Tachypnea/Dyspnea/Shortness of Breath						
Swelling						
Vomiting/Nausea						
Weakness						
None, Asymptomatic						
Other1						
Othor?	<del>                                     </del>					
Other2						
a. If patient asymptomatic did the	patient develop signs of infection later?	Yes No Nunk				
	mptom(s)					
ii What day did symptom	s develop?(MM/DD	 /VVV)				
ii. What day did symptoni	(IVIIVII DD	, , , , , ,				
26. Patient Labs & Vitals within 2 hours of cul	ture collection					
		/// Time o				
Value	Date:MMDD	YY Time::				
Temperature Heart Rate						
Blood Pressure						
Respiratory Rate						
Pulse Ox						
Lactate						
WBC						
******						
27. Diagnosis for the visit when the positive of	ulture was collected:					
Acute Respiratory Failure	Pleural Effusion					
Atrial Fibrillation	Pneumothorax					
Bacteremia	Renal Failure					
Cellulitis	Sepsis					
□DKA	Stroke					
Heart Failure						
	Hyperglycemia Other1:					
Myocardial Infarction	Other2:					
28. Was Patient Admitted in response to t	he positive culture? $\square$ Yes $\square$ I	Vo				
29. Did the patient have evidence of soft t	issue infection at time of positive cu	Iture: Yes or No				
a. If ves, describe:		<del></del>				

Line list patient ID (CDCID) Wisconsin State Laboratory of Hygiene ID _	Abstractor Initials					
30. Did the patient have diagnosed pulmonary infection at time of pos	itive culture? Yes or No					
a. If yes, describe:						
31. Did the patient have any other infections at time of positive culture? Yes or No?						
a. If yes, describe:						
32. What was patient's disposition from hospitalization in which EK po	32. What was patient's disposition from hospitalization in which EK positive culture was collected?					
33. Death Home Inpatient rehab LTCF/SNF	Hospice Other:					
34. If the patient Died: Date of death//(MM/DD/YYYY	)					
35. Location of death: home Inpatient rehab LTCF/SNF	hospice Other:					
36. Diagnosis at time of death?						

· ·				ited in the 30 days prior to positive Ek
culture				
Name of Facility/Clinic			_	_
Date(s) of visit to healthcare far Inpatient or outpatient:	ility	are Other (MM/DD/YYYY)	-	
If inpatient, date of admission:	//	(MM/DD/YYYY)		
If inpatient, date of discharge:				
Reason for visit or chief compla	int:			
Percutaneous Exposures		Date	Notes	Specify products used
		(MM/DD/YYYY)		
1. IV infusion	Yes No			<u>Infusate:</u>
				Antiseptic:
				Antisoptic.
2. IM injection	Yes No			<u>Infusate:</u>
				Antiseptic:
3. Thoracentesis	Yes No			Specify antiseptic
4. Paracentesis	Yes No			Specify antiseptic
5. Peripheral IV insertion	Yes No			Specify antiseptic
6. Central line placement.	Yes No			Specify antiseptic
7. Type 1:				
8. Central line placement.	Yes No			
9. Type 2:				
10. Interventional Radiology	Yes No			<u>Infusate:</u>
				Antiseptic:
				Апизерис.
11. Radiology with contrast	Yes No			Infusate:
				Antiseptic:
12. Labs Drawn	Yes No		Specify:	Specify antiseptic
13. Bedside tests (e.g. Blood	Yes No		Specify:	Specify antiseptic
glucose, lactate)				
14. Other	Yes No			
1(specify)				
15. Other 2	Yes No			
(specify)				
16. Other 3	Yes No			
(specify)				

Respiratory Exposures			
17. Oxygen Administered (e.g	Yes No		
face mask, nasal cannula)			
18. Intubation	Yes No		
19. Nebulizer	Yes No		Specify agent
20. Metered Dose Inhaler (MDI)	Yes No		Specify agent
21. Other(specify)	Yes No		
Topical Exposures			
22. Podiatry care	Yes No		Specify any topical
			<u>treatments</u>
23. Whirlpool therapy	Yes No		
24. Any topical treatments	Yes No		
Other exposures:			
25. Endoscopy	Yes No	Specify:	
26. Other exposure 1	Yes No		

Line list patient ID (CDCID) \_\_\_\_\_ Wisconsin State Laboratory of Hygiene ID \_\_\_\_\_ Abstractor Initials\_\_\_\_\_

Form Approved OMB No. 0920-1011 Exp. Date 03/31/2017

# 2016 Urgent Assessment of Blood Collection and Use in Puerto Rico in Response to the Zika Virus Outbreak

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; ATTN: PRA (0920-1011)

### ¡Hola!

Please fill out the following sections that pertain to your institution as completely as possible. The arrows will help you progress through the survey. In general, Section 2 corresponds to Blood Collections and Section 3 corresponds to Blood Transfusions.

Please complete and return to the below email by February 19, 2016

Amber Vasquez, MD, MPH Zika Blood Safety Team

email: amber.vasquez@salud.pr.gov

cell: 937-269-3169

Please do not hesitate to call or email with questions.

Primary person responsible for completing this section	
Prefix	
First name	
Last name	
Title/Position	
Name of Institution	
Address of Institution	
Telephone	
Email	

Facility included in the survey	
Facility name	
Address	

#### **General Information**

Which of the following best describes your institution?	Select one (with "X")
A local or regional blood center (non-hospital) that collects blood from donors and supplies blood and components to other institutions, but does not perform transfusion services	
A hospital-based blood bank and transfusion service that collects blood from donors (may be only autologous or directed) and provides blood and components for transfusion primarily to your own institution	
A transfusion service that provides blood and components for transfusion, but does not collect blood from donors	
A local or regional blood center that collects blood from donors and supplies blood, components, and cross matched blood products to participating facilities (e.g., centralized transfusion services). In this category, the service is not limited to reference laboratory work, but includes routine transfusion service work	

	Yes/No
Does your institution collect blood from donors? (Even if you collect autologous units only, enter "Yes.")	

#### **Section 2 Blood collections**

From Jan 1, 2015 through Dec 31, 2015, how many collections were successfully completed by your institution in each of the following categories? (*indicates required	Number of Collection Procedures*	Number of Units
fields)		
Whole Blood		
Allogeneic (non-directed donations)*		
Autologous*		
Directed*		
Total*		
Red Blood Cells		
Apheresis		
Allogeneic*		
Autologous*		
Directed*		
Concurrent red cells (from apheresis platelets)	_	
Total Apheresis Red Blood Cells*	-	
Whole-blood-derived Allogeneic*	_	
Allogeneic Autologous*	-	
Directed*	-	
Total WBD Red Blood Cells*	-	
Total Mad Hod Blood Colle		
Platelets		
Apheresis		
Single-donor		
Directed single-donor		
Single collection	_	
Double collection <sup>1</sup>	_	
Triple collection <sup>1</sup>	_	
Total Apheresis Platelets*  Total apheresis platelet units subjected to pathogen reduction technology	_	
Whole-blood-derived		
Individual* <sup>2</sup>	-	
Total whole blood-derived individual units subjected to pathogen reduction technology	-	
Diamon		
Plasma Apheresis		
FFP		
PF24		
PF24RT24		
Jumbo FFP (>400 mL)		
Total Apheresis Plasma*		
Total Apheresis plasma units subjected to pathogen reduction technology		
Whole-blood-derived FFP	-	
PF24		
Cryoprecipitate reduced		
Liquid		
Total WBD Plasma*		
Total WBD plasma units subjected to pathogen reduction technology		
Cryoprecipitate		
Individual* <sup>3</sup>		
Total Granulocytes*		

<sup>&</sup>lt;sup>1</sup> Count double collections as two units and triple collections as three units

<sup>&</sup>lt;sup>2</sup> Enter the number of individual platelet units prepared from whole blood collections

<sup>&</sup>lt;sup>3</sup> Enter the number of individual cryoprecipitate units prepared from whole blood collections

## 2.3 Blood collections

2.3. From Jan 1, 2015 through Dec 31, 2015, from how many of the following types of donors did your institution successfully collect blood?	Number of Donors
First-time allogeneic donors	
Repeat allogeneic donors (Count multiple donations from a single repeat donor only once)	
Directed donors	
Autologous donors	
Total number of donors	

#### 2.4 Blood collections

Whole Blood for distribution as Whole Blood  Allogeneic (non-directed donations)  Lutologous  Directed  Total*  Red Blood Cells  Apheresis  Allogeneic  Autologous  Directed  Concurrent ed cells (from Total Apheresis Red Blood Cells*  Whole Blood chrived  Autologous  Directed  Total WBD Red Blood Cells*  Total WBD Red Blood Cells*  Patients  Single-donor  Single-donor  Doubte cells cells (from Total Apheresis Patients  Single-donor  Single-donor  Total WBD Red Blood Cells*  Patients  Single-donor  Total Cells  Fire Fire Fire Fire Fire Fire Fire Fir	From Jan 1, 2015 through Dec 31, 2015, how many units of each product were imported, distributed, and outdated by your institution? (* indicate required fields)	Total Units Imported	(including imported	Total Units Outdated
Autologous Directed Total*  Red Blood Cells  Apheresis Aliogeneic Autologous Directed Concurrent ed cells (from Total Apheresis Red Blood Cells* Whole-blood-derived Autologous Directed Autologous Directed Total Apheresis Red Blood Cells* Total Apheresis Red Blood Cells*  Platelets Autologous  Directed Total WBD Red Blood Cells*  Platelets Apprecis Apprecis Total WBD Red Blood Cells*  Platelets Apprecis Total Cells (From Ce	Whole Blood for distribution as Whole Blood			
Directed	Allogeneic (non-directed donations)			
Directed	Autologous			
Total*  Red Bload Cells  Apheresis  Allogeneic Autologous Directed Concurrent red cells (from Total Apheresis Red Bload Cells* Whole-bload-derived Autogeneic Autologous Directed  Total WBD Red Bload Cells*  Total WBD Red Bload Cells*  Platelets Apheresis Single-donor Directed single-donor Single collection* Total Apheresis Platelets* Whole-bload-derived Individual* Pooled*  PERMENTAL STATES AND STATES AN				
Red Blood Cells  Apheresis  Allogensic  Autologous  Directed  Concurrent red cells (from  Total Apheresis Red Blood Cells'  Whole-blood-derived  Autologous  Directed  Total WBD Red Blood Cells'  Plateles  Apheresis  Single donor  Directed single-donor  Single collection  Double collection'  Triple collect				
Apheresis  Allogeneic Autologous Directed Concurrent red cells (from Total Apheresis Red Blood Cells* Whole blood-derived Allogeneic Autologous Directed  Total WBD Red Blood Cells*  Platelets Apheresis Single-donor Directed single-donor Direc	Total			
Autologous Directed Concurrent red cells (from Total Aphresis Red Blood Cells* Whole-blood-derived Autologous Directed Total WBD Red Blood Cells*  Plateles Aphresis Aphresis Aphresis Concurrent red cells (from Total WBD Red Blood Cells*  Plateles Aphresis Aphresis Aphresis Single collection Directed single-dronor Single collection Double collection Total Aphresis Plateles* Triple collection Total Aphresis Plateles* Pooled*  Pooled*  Plasma Aphresis Plateles  Aphresis Plateles Triple collection Total Aphresis Plateles* Triple collection Total Aphresis Plateles Triple collection Total Aphresis Plasma* Triple collection Triple collectio	Red Blood Cells			
Autologous Directed Concurrent red cells (from Total Aphreresis Red Blood Cells' Whole-blood-derived Allogeneic Autologous Directed  Total WBD Red Blood Cells'  Platelets Aphreresis Single-donor Directed single-donor Directed single-donor Directed single-donor Total Mpresis Platelets Aphreresis Single collection Single-donor Double collection Single collection Total Aphreresis Platelets Total Aphreresis Platelets Aphreresis FFP Plasma Aphreresis FFP PP24 PP24RT24 PP24RT25 PP24 Cryoprecipitate reduced Liquid Total MpB Plasma' Cryoprecipitate reduced Liquid Total WBD Plasma' Cryoprecipitate Individual' Total WBD Plasma' Cryoprecipitate Individual' Total WBD Plasma' Cryoprecipitate Individual' Pooled Single-colored Individual' Total WBD Plasma' Cryoprecipitate Individual' Total WBD Plasma' Cryoprecipitate Individual' Pooled Single-colored Individual' Pooled Single-colored Individual' Total WBD Plasma'	Apheresis			
Directed	Allogeneic			
Concurrent red cells (from Total Apheresis Red Blood Cells* Whole-blood-derived Allogeneic Autologus  Directed  Total WBD Red Blood Cells*  Platelets Apheresis Single-donor Directed single-donor Directed single-donor Single collection Total Apheresis Platelets  Apheresis Tiple collection Total Apheresis Platelets' Whole-blood-derived Individual* Pooted*  Plasma Apheresis FFP FP FP24 PP24T24 PP24T724 Jumbo FFP (<400 mL) Total Apheresis Plasma' Whole-blood-derived FFP FPP FPP FPP FPP FPP FPP FPP FPP FP	Autologous			
Total Apheresis Red Blood Cells*  Whole-blood-derived  Allogons  Directed  Total WBD Red Blood Cells*  Platelets  Apheresis  Single-donor  Directed single-donor  Single collection  Double collection <sup>2</sup> Total Apheresis Platelets*  Whole-blood-derived Individual*  Pooled*  Plasma  Apheresis  FFP  PF24  PF24  PF24  PF24  Jumbo FFP (-400 mL)  Total Apheresis Plasma*  Whele-blood-derived  United by the plasma*  Whele-blood derived  PF24  PF25  PF24  PF24  PF24  PF24  PF24  PF24  PF25  PF24  PF24  PF25  PF24  PF24  PF25  PF24  PF25  PF24  PF24  PF25  PF24  PF24  PF25  PF24  PF25  PF24  PF25  PF24  PF25  PF24  PF25  PF24  PF25  PF25  PF24  PF25  PF24  PF27  PF24  PF29  PF24  PF28  PF29  PF24  PF29				
## Allogonic ## Al	Concurrent red cells (from			
Allogeneic Autologous  Directed  Total WBD Red Blood Cells*  Platelets Apheresis Single-donor Directed single-donor Single collection* Obuble collection* Triple collection* Triple collection* Total Apheresis Platelets* Whole-blood-derived Individual* Pooled*  Plasma Apheresis FFP PF24 PF24T24 Jumbo FFP (1400 mL) Total Apheresis Plasma* Whole-blood-derived  FFP PF24 Jumbo FFP (2400 mL) Total Apheresis Plasma* Whole-blood-derived  FFP PF24 Jumbo FFP (2400 mL) Total Apheresis Plasma* Whole-blood-derived  FFP PF24 Jumbo FFP (2400 mL) Total Apheresis Plasma* Whole-blood-derived FFP PF24 Jumbo FFP (2400 mL) Total Apheresis Plasma* Whole-blood-derived FFP PF24 Jumbo FFP (2400 mL) Total Apheresis Plasma* Total WBD Plasma*  Whole-blood-derived FFP PF24 Cryoprecipitate reduced Liquid Total WBD Plasma*	Total Apheresis Red Blood Cells*			
Autologous  Directed  Total WBD Red Blood Cells*  Platelets Apheresis Single-donor Directed single-donor Single collection Double collection Total Apheresis Platelets* Whole-blood-derived Individual* Pooled  Plasma Apheresis FFP PF24 PF24RT24 Jumbo FFP (-400 mL) Total Apheresis Plasma* Whole-blood-derived FFP PF24 Cryoprecipitate reduced Liquid Total WBD Red Blood Cells*  Total Apheresis Plasma* Whole-blood-derived Cryoprecipitate reduced Liquid Total WBD Plasma*  Cryoprecipitate Individual*  Pooled*	Whole-blood-derived			
Directed  Total WBD Red Blood Cells*  Platelets Apheresis Single-donor Directed single-donor Single collection Double collection* Triple collection* Total Apheresis Platelets* Whole-blood-derived Individual* Pooled*  PEP24 PP24 PP244 PP24T24 Jumbo FFP (-400 mL) Total Apheresis Plasma* Whole-blood-derived  Individual* PFP PFP PFP PFP PFP PFP PFP PFP PFP PF				
Total WBD Red Blood Cells*  Platelets Apheresis Single-donor Directed single-donor Single collection* Double collection* Total Apheresis Platelets* Whole-blood-derived Individual* Pooled*  PP24 PP24 PP24 PP24 PP24 PP24 PP24 PP	Autologous			
Platelets Apheresis Single-donor Directed single-donor Single collection Double collection Total Apheresis Platelets  Hodividual' Pooled PF24 Jumbo FFP (>400 mL) Total Apheresis Plasma* Whole-blood-derived Jumbo FFP (>400 mL) Total Apheresis Plasma* Cryoprecipitate Individual'  Pooled  FOR Double Collection Double Co	Directed			
Apheresis   Single-donor   Single-collection   Single collection	Total WBD Red Blood Cells*			
Apheresis   Single-donor   Single-collection   Single collection				
Single-donor				
Directed single-donor   Single collection	Aprieresis Single-donor			
Double collection <sup>2</sup>				
Triple collection <sup>2</sup> Total Apheresis Platelets*  Whole-blood-derived Individual*  Pooled <sup>3</sup> Plasma  Apheresis  FFP  PF24  PF24A  Jumbo FFP (>400 mL)  Total Apheresis Plasma*  Whole-blood-derived  FFP  PF24  Cryoprecipitate reduced  Liquid  Total WBD Plasma*  Cryoprecipitate  Individual*  Pooled <sup>4</sup> Individual*  Pooled <sup>4</sup>				
Total Apheresis Platelets*  Whole-blood-derived  Individual*  Pooled³  Plasma  Apheresis  FFP  PF24  PF24  PF24RT24  Jumbo FFP (>400 mL)  Total Apheresis Plasma*  Whole-blood-derived  FFP  PF24  Cryoprecipitate reduced  Liquid  Total WBD Plasma*  Cryoprecipitate  Individual*  Pooled⁴  Whole-blood-derived  FFP  Precipitate  Individual*  Pooled⁴  Pooled⁴  Pooled⁴				
Whole-blood-derived	Triple collection <sup>2</sup>			
Individual* Pooled³  Plasma Apheresis  FFP PP24 PF24RT24 Jumbo FFP (>400 mL) Total Apheresis Plasma*  Whole-blood-derived FFP PF24 Cryoprecipitate reduced Liquid Total WBD Plasma*  Cryoprecipitate Individual* Pooled⁴  Pooled⁴  Pooled⁴  Pooled⁴  Pooled⁴  Individual* Pooled⁴  Pooled*	Total Apheresis Platelets*			
Plasma Apheresis  FFP PF24 PF24RT24 Jumbo FFP (>400 mL) Total Apheresis Plasma* Whole-blood-derived FFP PF24 Cryoprecipitate reduced Liquid Total WBD Plasma* Cryoprecipitate Individual* Pooled <sup>4</sup> Pooled <sup>4</sup> Plasma  Apheresis  Apheresis Apheresis Apheresis Plasma Apheresis Plasma Apheresis Plasma Apheresis Plasma Apheresis Plasma Apheresis Plasma Apheresis Apheres				
Apheresis   FFP	Pooled <sup>3</sup>			
Apheresis   FFP				
FFP				
PF24	FFP			
Jumbo FFP (>400 mL)  Total Apheresis Plasma*  Whole-blood-derived  FFP  PF24  Cryoprecipitate reduced  Liquid  Total WBD Plasma*  Cryoprecipitate  Individual*  Pooled*	PF24			
Total Apheresis Plasma* Whole-blood-derived FFP PF24 Cryoprecipitate reduced Liquid Total WBD Plasma*  Cryoprecipitate Individual* Pooled*	PF24RT24			
Whole-blood-derived         Image: Control of the	Jumbo FFP (>400 mL)			
FFP         Image: Control of the				
Cryoprecipitate reduced Liquid Total WBD Plasma*  Cryoprecipitate Individual* Pooled <sup>4</sup>	FFP			
Liquid Total WBD Plasma*  Cryoprecipitate Individual* Pooled <sup>4</sup> Individual* In				
Total WBD Plasma*  Cryoprecipitate Individual* Pooled*  Pooled*				
Cryoprecipitate Individual* Pooled <sup>4</sup> Individual*				
Individual*  Pooled <sup>4</sup> Individual *  Pooled *				
Pooled <sup>4</sup>				
Total Craw Joseph 1	Pooled*			
ODIA CITADUOCNES	Total Granulocytes*			

<sup>&</sup>lt;sup>1</sup> Units returned and distributed more than once should be counted only once

<sup>&</sup>lt;sup>2</sup> Count double collections as two units and triple collections as three units

<sup>&</sup>lt;sup>3</sup> Total number of platelet pools prepared from whole blood collections
<sup>4</sup> Total number of cryoprecipitate pools prepared from whole blood collections

#### 2.5-2.6 Blood collections

2.5 What was the average whole dollar amount your institution was reimbursed (by hospital or clinical facility) per unit in 2015 for the following components? (Include discounts in your calculations. If you do not use a particular component, select "Not Applicable". CPT/HCPCS codes are in in parenthesis.)	Average Amount Paid Per Unit (\$)
Plasma, single donor, frozen with 8 hours of phlebotomy (P9017)	
Plasma, frozen between 8 and 24 hours of phlebotomy (P9059)	
Red cells, leuko-reduced (P9016)	
Red cells, non-leuko-reduced (P9021)	
WBD platelets, each unit, not leuko-reduced, not irradiated (P9019)	
Apheresis platelets, leuko-reduced (P9035)	
Cryoprecipitate, each unit (P9012)	

2.6. If your facility does not use pathogen reduction technology for apheresis platelet or plasma collections	Cost
What is the estimated total cost of implementation (this includes equipment, capital investment, training, etc)?	
What is the estimated additional cost per each unit type below if your facility adopted pathogen reduction technology?	

## Section 3 - Blood utilization

	Yes/No
Is your institution directly involved in the transfusion of blood to patients?	

## 3.3 Blood utilization

3.3. From Jan 1, 2015 through Dec 31, 2015, how many units of allogeneic whole blood and red blood cells did your institution transfuse? (Leave the field blank if you do not know the answer).	Total Number of Units Transfused	 Total outdated units
Allogeneic Whole Blood		
Allogeneic Red Blood Cells (include all blood groups)		
Allogeneic Group O Positive RBCs		
Allogeneic Group O Negative RBCs		
Allogeneic Group A Positive RBCs		
Allogeneic Group A Negative RBCs		
Allogeneic Group B Positive RBCs		
Allogeneic Group B Negative RBCs		
Allogeneic Group AB Positive RBCs		
Allogeneic Group AB Negative RBCs		

## 3.4 Blood utilization

3.4. Indicate the disposition of directed and autologous units in 2015	Total Number of Units Transfused to Intended Recipient	Total Number of Recipients	Outdated Units
Directed Whole Blood Units			
Directed RBC Units			
Autologous Whole Blood Units			
Autologous RBC Units			

## 3.5 Blood utilization

3.5. From Jan 1, 2015 through Dec 31, 2015, how many units of each of the following components did your institution transfuse and how many units were outdated while on your shelf (include units transfused to pediatric patients)? (* indicates required fields)	Total Number of Units Transfused	Total Number of Units Outdated
WBD Platelets (individual concentrates and pools expressed as individual concentrate equivalents)*		
Apheresis Platelet units – Full dose*		
Directed Platelets to intended recipients		
Total Plasma*		
Fresh Frozen Plasma (FFP)		
FFP, pediatric size (≤100 mL)		
Plasma, Frozen within 24 hours (PF24)		
PF24RT24		
Jumbo FFP (>400 mL)		
Liquid plasma		
Directed plasma to intended recipients		
Thawed plasma		
Plasma, cryoprecipitate reduced		
Group AB plasma		
Granulocytes*		
Platelets with pathogen reduction technology		
Plasma with pathogen reduction technology		

## 3.6 Blood utilization

3.6. Indicate the total number of units transfused to pediatric populations in 2015	Number of Adult Equivalent Units in Whole or in Part for Pediatric Patients <sup>1</sup>	Total Number of Pediatric Recipients
Whole Blood		
RBCs		
Plasma		
Platelets		

<sup>&</sup>lt;sup>1</sup> This should be a subset of data reported in question 4 and 5 if your hospital transfuses non-pediatric patients.

### 3.7 Blood utilization

following adult e	ndicate how many irradiated, leuko-reduced, and leuko-filtered units for each of the ing components your institution transfused in 2015. For pediatrics, use the number of equivalent units used in whole or part. For components that are irradiated and leukoed, include these in the count for both columns.	Components Irradiated	Components Leuko- reduced Before or After Storage (not at bedside)	filtered at the
a.	Whole Blood			
b.	RBCs			
c.	Apheresis platelets (single donor platelets)			
d.	WBD platelets			
Total c	omponents (if the number for a-d is 'unknown', enter the total number of components for the eation)			

## 3.8-3.9 Blood utilization

	Yes/No
3.8. Does your institution have a policy to transfuse only leuko-reduced (LR)	
components?	

3.9a. In 2015, how many total units of RBCs transfused were	Number of Units
1 - 35 day(s) old	
36 – 42 days old	

3.9b. In 2015, how many total units of WBD platelets transfused were	Number of Units
1-3  day(s) old	
4 – 5 days old	

3.9c. In 2015, how many total units of Apheresis platelets transfused were	Number of Units
1 - 3 day(s) old	
4 – 5 days old	

## 3.10-3.11 Blood utilization

	Number of platelet units
3.10. In your institution, on average, how many individual platelet units were included in a pooled WBD platelet dose in 2015?	

3.11. Indicate the number of units that were transfused in inpatient or outpatient settings.	Number of RBC Units	Number of Platelet Units	Total	Don't Know
All Surgery (including transplant)				
Inpatient Medicine (including hematology/oncology)				
Emergency Department				
Obstetrics/Gynecology				
Pregnant females				
Pediatrics				
Neonates				
Outpatient and non-acute inpatient settings <sup>1</sup>				

<sup>&</sup>lt;sup>1</sup> E.g., outpatient dialysis, rehabilitation, long term care, etc.

### 3.12 Blood utilization

3.12. What was the average whole dollar amount your institution paid per unit in 2015 for the following components? (Include discounts in your calculations. If you do not use a particular component, select "Not Applicable". CPT/HCPCS codes are in in parenthesis.)	Average Amount Paid Per Unit (\$)
Plasma, single donor, frozen with 8 hours of phlebotomy (P9017)	
Plasma, frozen between 8 and 24 hours of phlebotomy (P9059)	
Red cells, leuko-reduced (P9016)	
Red cells, non-leuko-reduced (P9021)	
WBD platelets, each unit, not leuko-reduced, not irradiated (P9019)	
Apheresis platelets, leuko-reduced (P9035)	
Cryoprecipitate, each unit (P9012)	

## 3.13 Blood utilization

	Yes/No
3.13a. Were any elective surgeries postponed due to blood inventory shortages in 2015?	

	Number of days
3.13b. How many days were elective surgeries postponed?	

	Number of surgeries
3.13c. How many elective surgeries were postponed in 2015?	

### 3.16 - 3.17 Blood utilization

16. In 2015, how many days was your institution's order incomplete for the following components?	Number of days
Whole Blood	
RBCs	
Plasma	
Apheresis platelets	
WBD platelets	

	Number of days
17. In 2015, how many days were you unable to meet other non-surgical blood requests (e.g., red cells, platelets)?	

## 3.18-3.20 Blood utilization

	Number of units
18. At your institution, how many units of Group O red cells are on your shelf on	
an average weekday?	

	Number of units
19. At what number of Group O positive and Group O negative RBC units in uncrossmatched inventory do you consider your inventory to be "critically low"?	

	Yes/No
20. Does your facility have an electronic system for tracking transfusion-related adverse events (e.g., unplanned, unexpected, and undesired occurrences)?	

## 3.21 Blood utilization

	Number of units
3.21a. How many total red blood cell units did you buy from a non-American Red	
Cross blood center in 2015?	

	Number of units
3.21b. How many total red blood cell units did you buy from an American Red Cross	
blood center in 2015?	

## **Survey Completed!**

Thank you for taking the time to complete this survey.

Please return to the below email by February 19, 2016

Amber Vasquez, MD, MPH Zika Blood Safety Team

email: amber.vasquez@salud.pr.gov

cell: 937-269-3169

Please do not hesitate to call or email with questions.

### **Survey Glossary**

Autologous: Self-directed donations.

**Centralized transfusion service:** A hospital or blood center that collects blood from donors and supplies blood, components, medical services and/or crossmatched blood products to multiple transfusing facilities.

**Collected:** Successful whole blood or apheresis collections placed into production (not QNS, or other removals).

**Deferrals:** The number of donors deferred for specific reasons:

- a) Donors deferred for low hemographic do not meet the current FDA blood hemographic requirements for b)^^deferrais for other medical reasons may include the use of medications on the medication deferral list, growth hormone from human pituitary glands, insulin from cows (bovine, or beef, insulin), Hepatitis B Immune Globulin (HBIG), unlicensed vaccines, or presenting with physical conditions or symptoms that do not qualify a person to be a blood donor.
- c) High-risk behavior deferrals include deferrals intended to reduce the risk of transmission of infectious diseases including HIV and hepatitis viruses. Examples of questions intended to identify these risks are sexual contact (e.g., men who have sex with men (MSM)) and non-medical injection drug use questions.
- d) Travel deferrals are deferrals for travel to a specific region of the world.

**Directed:** Allogeneic donations intended for a specific patient.

**Donation:** The collection of a unit of blood or blood component from a volunteer donor.

Dose/Dosage: a quantity administered at one time, such as a specified volume of platelet concentrates.

First-time allogeneic donor: A donor who is donating for the first time at your center.

**Imported**: Units not collected by your institution, but obtained by your institution from another institution for distribution to a transfusion facility.

**Modify:** Procedures applied by a blood center, hospital blood bank, or transfusion service that may affect the quality or quantity of the final product (e.g., irradiation, leukofiltration, or production of aliquots of lesser volume).

Outdated: Units that expire on your shelf.

#### Plasma:

- a) **Plasma, frozen within 24 hours of phlebotomy (PF24):** plasma separated from the blood of an individual donor and placed at -18 C or colder within 24 hours of collection from the donor.
- b) Fresh frozen plasma (FFP): Plasma frozen within 8 hours of collection.
- c) Plasma, Jumbo: FFP having a volume greater than 400 mL.
- d) Plasma frozen within 24 hours of phlebotomy and held at room temperature up to 24 hours after phlebotomy (PF24RT24): Plasma held at room temperature for up to 24 hours after collection and then frozen at -18 C or colder.

Recipient: A unique individual patient receiving a transfusion one or more times in a calendar year.

Distributed: units that have runnied an processing requirements and have been made available for transfer to

Repeat allogeneic donor: A donor who has previously donated a blood component.

**Severe Donor-Related Adverse Events:** adverse events occurring in donors attributed to the donation process that include, for example, major allergic reaction, arterial puncture, loss of consciousness of a minute or more, loss of consciousness with injury, nerve irritation, etc.

<u>Transfusion Related Adverse Reactions: An undesirable response or effect in a patient temporally associated with the administration of blood or blood components. For a list of adverse reaction types and case definitions, visit <a href="http://www.cdc.gov/nhsn/PDFs/Biovigilance/BV-HV-protocol-current.pdf">http://www.cdc.gov/nhsn/PDFs/Biovigilance/BV-HV-protocol-current.pdf</a>.</u>

**Transfusion Service:** a facility that performs, or is responsible for the performance of, the storage, selection, and issuance of blood and blood components to intended recipients.

## Appendix 1. Invasive GAS in Long Term Care Facility 2016 <a href="EmployeeSurvey"><u>Employee Survey</u></a>

**Date Completed:** \_\_\_\_/\_\_\_\_

Form Approved; OMB No. 0920-1011 Exp. Date 03/31/2017

☐ Check box if documented case

A. Employee Backgroun	1. Name:				2	2. Age:			
3. Sex: ÿ Male ÿ	Female	4. Employ	4. Employed at Facility since:/						
5. List occupation: ÿ Activi	ty aid j	ÿ Adminis	trative	ÿ CNA		ÿ Diet	ary	ÿ Food service	
ÿ Housel	keeping j	ÿ Laundry ÿ PT/OT ÿ			ÿ Pha	harmacist ÿ Physician			
ÿ Mainte	enance	ÿ RNA		ÿ RN/LF	PN	ÿ Soc	cial service ÿ Van driver		
ÿ Wour	nd care team	ÿ(	Other				_		
6. Since Thanksgiving to pre	esent, have you	worked in a	ny other pa	ntient-care fa	acility?		ÿ Yes	ÿ No (If no, skip to Section B)	
Name & city of facility	Dates of empl	oyment	byment Have you been in contact with a patient infected with group A strep?				What was t	he patient's diagnosis?	
	Start:		ÿ Yes				ÿ Strep th	roat ÿ Impetigo	
	/	/	ÿ No				ÿ Celluliti	s ÿ Bacteremia/Sepsis	
	End:/	/		te of contact	t: 		ÿ Other, specify:		
	Start:			ÿ Yes			ÿ Strep throat ÿ Impetigo		
	ÿ Nо			ÿ Cellulitis ÿ Bacteremia/Sepsis					
	End:			If yes, date of contact://			ÿ Other, specify:		
	Start:			ÿ Yes			ÿ Strep th	roat ÿ Impetigo	
	/   ÿ No			ÿ Celluliti	s ÿ Bacteremia/Sepsis				
	If yes, date of contact:/			ÿ Other, specify:					
7. a. Since the outbreak						ococcu	s? ÿ Yes	ÿ No (If no, skip to # 8)	
b. If yes, when?	//_								
c. Where was the cultur	re obtained from	n? ÿ Thro	oat ÿ Re	ectal ÿ	Vaginal	ÿ Ski	n/wound	ÿ Other	
d. What were the result	ts? ÿ Positive	e ÿ Neg	gative						
B. Job Description at		8. As part	of your job	o, do you ha	ve physica	l contac	ct with patien		
•	Facility A (If no, skip to Section D)								
9. Areas usually worked:		-		-	a y Keha	b floor	y Other		
10. Shifts usually worked: ÿ Day ÿ Evening ÿ Night ÿ Other									
11. Patient units usually worked: ÿ 1 ÿ 2 ÿ 3 ÿ 4 ÿ 5 ÿ 6 ÿ 7 ÿ 8 ÿ Do not work in patient units ÿ All patient units									
12. Which days do you usual		•		_					
Sunday Monday	7 Tue	esday	Wedne	esday	Thursday	y	Friday	Saturday	

Public reporting burden of this collection of information is estimated to average 15 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; ATTN: PRA (0920-1011)

13.	What	t kind of patient conta	act do you have? (check ALL th	hat appl	y)								
ÿ(	$\ddot{y}$ Give oral medications $\ddot{y}$ Feeding resident $\ddot{y}$ Respiratory therapy							ÿ Tracheostomy care					
ÿ(	ÿ Change dressings/wound care ÿ Gastrostomy care ÿ Handle urinary catheter						•	ÿ Bathe resident					
ÿ Assist with patient transfer ÿ Clean room ÿ Handle soiled linens/bedding ÿ Handle								le soil	ed diar	pers/be	edpans		
ÿΙ	ÿ Deliver meal trays ÿ Take vital signs ÿ Bedside incision and debridement aspiration/drainage												
ÿΡ	ÿ Provide PT/OT ÿ Other beside surgical procedures												
7	C. Work Practice   14. Do you use soap and water to clean your hands?   ÿ Yes   ÿ No												
C.	Wor	rk Practice	15. Do you use alcohol-base		-		ur hands?	ÿΥ		ÿ No			
16.	Pleas	se answer the followin						Never				Always	
	a.	Do you perform han	nd hygiene BEFORE physical	contact	with patie	nts?		1	2	3	4	5	N/A
	b.		nd hygiene BEFORE physical ongings (e.g. bedside table, refi				;.)?	1	2	3	4	5	N/A
	c.	Do you perform han	nd hygiene AFTER physical co	ontact w	ith patient	is?		1	2	3	4	5	N/A
	d.		nd hygiene AFTER physical cobedside table, refrigerator, rolli			atient's en	vironment	1	2	3	4	5	N/A
	e.	Do you perform han	nd hygiene BETWEEN contact	t with p	atients?			1	2	3	4	5	N/A
	f.	Do you use the sink room?	or alcohol-based sanitizer in t	he patie	nt's room	or outside	patient's	1	2	3	4	5	N/A
	g.	Do you use the sink	or alcohol-based sanitizer at the	he nurse	e's station'	?		1	2	3	4	5	N/A
	h.		when changing bandages/dress					1	2	3	4	5	N/A
		i. If yes, do you	ou change gloves between patie	ents/pati	ient rooms			1	2	3	4	5	N/A
			ou perform hand hygiene befor ou perform hand hygiene after i					1 1	2 2	3	4 4	5 5	N/A N/A
	1.		when cleaning soiled patients of					1	2	3	4	5	N/A
		m. If yes, do you	ou change gloves between patie	ents/pati	ient rooms			1	2	3	4	5	N/A
			ou perform hand hygiene before					1	2	3	4	5	N/A
		o. If yes, do you perform hand hygiene after removing gloves?  1 2 3 4 5 N/A											
	p.		protective equipment (PPE) whe specify type of PPE:			ts?		1	2	3	4	5	N/A
D.	You	ur Health	17. Do you have paid "Sick 18. Did you receive prophyla		•	•		n? ÿ Ye	s ÿ i	No Wł	nen? _	/	/
19.	a.	Since Thanksgiving	g, have you had a sore throat?			ÿ Yes	ÿ No	(If no, s	kip to	#20)			
	b.	When? /	/			•							ļ
	c.		for testing collected from you?		ÿ Yes	•	d. If yes	s, specify	montł	a:			
	e.	Was a rapid strep the	nroat test done (you would have	e been g	-		•						ĺ
			specify month:			, was the	result positi	-	ÿ Yes	•			
	h.	Were you diagnosed	•	ÿ Yes	ÿ No		i. If yes,	-					
	j.	Did you miss work f		ÿ Yes	ÿ No		k. How n	nany day	s did y	you mis	ss?		
	1.		ere you ill?										
	m.	· · · · · · · · · · · · · · · · · · ·		ÿ Yes	ÿ No		n. If yes,						
20.			g, did you have a rash, open wo			-				_			
		d. Did you miss work f f. How many days were	for this illness? re you ill?		ÿ Yes	ÿ No	How man	ny days d	id you	a miss?	<u>'</u>		
			ibiotics for this condition?		ÿ Yes	ÿ No	If yes, an	ntibiotic n	iame _				

Study ID #: \_CHŸŸŸ

## Appendix 1. Invasive GAS in Long Term Care Facility 2016 <a href="mailto:EmployeeSurvey"><u>EmployeeSurvey</u></a>

Form Approved; OMB No. 0920-1011 Exp. Date 03/31/2017

21.	a. Since Thanksgiving, did you have fever, cough, and/or other respiratory infects	ection? ÿ Yes ÿ No (If no, skip to #22))
		How many days did you miss?
	d. How many days were you ill?  e. Did you receive antibiotics for this condition?	If yes, antibiotic name
22. If	you're feeling sick before a work shift, how do you notify Warren Barr Gold Coa	ast?
	<del></del>	
23.	a. How many people are in your household? (If none, END)	
	b. How many children under 18 years of age are in your household?  c. Since Thanksgiving, did anyone in your household have a sore throat?  d. When? / / e. Who (relationship)? _	ÿ Yes ÿ No
	e. Was he/she diagnosed with strep throat?	ÿ Yes ÿ No
	g. Were they treated? $\ddot{y}$ Yes $\ddot{y}$ No If so, with what?	
	h. During the past 3 months, did anyone in your household have impetigo or cel i. When? / /	ellulitis (skin infections)? ÿ Yes ÿ No

END - Thank you!

Study ID #: _CH
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Form Approved; OMB No. OMB No. 0920-1011 Exp. Date 03/31/2017

Person completing form							
Resident (check one): ÿ Case ÿ Control							
If case, indicate disease classification: □invasive	□noninvasive □colonized						
If CONTROL, date of matched case's GAS culture://							
Date 14 days prior to positive culture of case://							
Why was the culture obtained? ÿ Screening	g ÿ Illness						
A. GAS TESTING RESULTS							
1. Has the resident had any cultures/tests for GA	S from July 17, 2015 to present?						
Ÿ Yes ÿ No							

#	Date obtained	Site cultured	Culture	If nonsterile site,	Result
			obtained for	was culture	
			Screening	associated with	
				illness	
a.	, ,	ÿ Blood	ÿ Yes	ÿ Yes	ÿ Positive
	//	ÿ Skin/Wound:			
		ÿ Other	ÿ No	ÿ No	ÿ Negative
		ÿ Throat			
		ÿ Central line/TPN			
		ÿ Catheter ÿ Diabetic ulcer			
		ÿ Post-surgical wound ÿ G-			
		tube			
b.	, ,	ÿ Blood	ÿ Yes	ÿ Yes	ÿ Positive
	//	ÿ Skin/Wound:			
		ÿ Other	ÿ No	ÿ No	ÿ Negative
		ÿ Throat			
		ÿ Central line/TPN			
		ÿ Catheter ÿ Diabetic ulcer			
		ÿ Post-surgical wound ÿ G-			
		tube			

C.		ÿ Blood	ÿ Yes	ÿ Yes	ÿ Positive
	//	ÿ Skin/Wound:			
		ÿ Other	ÿ No	ÿ No	ÿ Negative
		ÿ Throat			
		ÿ Central line/TPN			
		ÿ Catheter ÿ Diabetic ulcer			
		ÿ Post-surgical wound ÿ G-			
		tube			
d.	, ,	ÿ Blood	ÿ Yes	ÿ Yes	ÿ Positive
		ÿ Skin/Wound:			
		ÿ Other	ÿ No	ÿ No	ÿ Negative
		ÿ Throat			
		ÿ Central line/TPN			
		ÿ Catheter ÿ Diabetic ulcer			
		ÿ Post-surgical wound ÿ G-			
		tube			
e.	/ /	ÿ Blood	ÿ Yes	ÿ Yes	ÿ Positive
		ÿ Skin/Wound:			
		ÿ Other	ÿ No	ÿ No	ÿ Negative
		ÿ Throat			
		ÿ Central line/TPN			
		ÿ Catheter ÿ Diabetic ulcer			
		ÿ Post-surgical wound ÿ G-			
f.		tube	" *7	" **	" " "
'-	/ /	ÿ Blood	ÿ Yes	ÿ Yes	ÿ Positive
		ÿ Skin/Wound:	∴ NI.	C. NI.	C. Nie auties
		ÿ Other	ÿ No	ÿ No	ÿ Negative
		ÿ Throat			
		ÿ Central line/TPN			
		ÿ Catheter ÿ Diabetic ulcer			
		ÿ Post-surgical wound ÿ G-			
		tube			

#### **B. RESIDENT BACKGROUND**

2. Sex:	ÿ Male	ÿ Female	3. Age:	_ 4. Date of birth:	/	/
---------	--------	----------	---------	---------------------	---	---

5a. Room history within 14 days prior to GAS culture:

Room # (floor/wing)	Dates in room	Roommate Yes/No	Roommate (dates)
	/to//	ÿ Yes ÿ No	/to//
	/to//	ÿ Yes ÿ No	/to//
	/to//	ÿ Yes ÿ No	/to//

/to//	ÿ Yes ÿ No	/to//
/to/	ÿ Yes ÿ No	/to//
/_ /to/_/_	ÿ Yes ÿ No	/ / to / /

5b. Did the resident have a re	commate with GAS infection	on or colonization within 30 days	prior to GAS culture?
ÿ Yes ÿ No ÿ Uı	nknown		
If yes: initials of GAS+	roommate	Dates room shared:/_	
5c. Number of visitors the res	sident had within 14 days ¡	orior to GAS culture?	
6. Total length of stay at facil months and_		at time of GAS culture (mark only	y one):
b. If resident died, de	eath was: ÿ Related to G ÿ Not related	If yes, date of death:/_ GAS infection	
8. Resident's physicians with	in 14 days prior to GAS cu	ılture'?	
Physician's name	Spec	cialty (e.g., wound care, etc.)	
			_
			_
	<u></u>		_
C. MEDICAL HISTORY 9. Which medical condition(s	) does the resident have?	(mark ALL that apply):	
ÿ Diabetes	ÿ CHF/history of MI	ÿ Peripheral vascular disease	ÿ Stroke
ÿ Asthma/COPD	ÿ Hypertension	ÿ Chronic leg edema	ÿ Recent herpes zoster
ÿ Dialysis	ÿ Renal insufficiency	ÿ Dementia	□糎久のCメ畑 Cヤ屋DC8 IDC
ÿ Cancer, specify typ	oe:	ÿ Immunosuppressed/immuno	suppression ÿ None
ÿ Cirrhosis □Malnutrition	ÿ Recent IV Drug Use	ÿ Prosthetic ÿ Other:	

Study ID #: CH	Study	ID #:	CH	
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(**Note**: immunosuppression includes: HIV/AIDS, chemo, radiation, immunosuppressive meds, including tacrolimus [Prograf], sirolimus [Rapamune], mycophenolate mofetil [Cellcept], high-dose or chronic steroids [prednisone, methylprednisone, hydrocortisone, dexamethasone] methotrexate.)

10a. Weight:	lbs or kg	(circle unit of measure)	10b. l	Height:	
•	, ,	nds, pressure ulcers, or	other wounds (de	fined as skin breakdow	vn) at the time
of admission to	•				
-	If yes, how many	•			
If Yes, s	ize of largest wound:		(e.g.	, largest width in inche	s or cm)
12. Did patient h	nave any surgical wou	nds, pressure ulcers, or	other wounds wit	hin 14 days prior to GA	S culture?
ÿ Yes	ÿ No				
If yes, p	lease specify site and	number of wounds.			
	ÿ Right/Left upper ex	tremity	ÿ Back	ÿ Pe	erineal
	ÿ Right/Left lower ext	remity	ÿ Abdomen	<del></del>	
	ÿ Right/Left Hand		ÿ Sacrum	_	
	ÿ Right/Left Foot	_	ÿ Chest		
	□ <b>癰</b> her				
13. Did the patie	ent receive <u>Wound Ca</u> ÿ No	re Team consultation se	ervices within 14 d	ays prior to GAS cultur	e?
Dates		Initial(s) of doctors or	nurses	Team	
14. Did the pation	ent receive wound car	e <u>WITHOUT Wound Ca</u>	re Team consultat	<u>ion</u> within 14 days prio	r to GAS
ÿ Yes	ÿ No	ÿ Unkown			
15. Products us	ed for wound care (su	rgical and nonsurgical)	within 14 days prid	or to GAS culture ( <i>ched</i>	:k all):
ÿ Calciu	ım Alginate ÿ Dakir	ns ÿ Dry Gauze ÿ F	oam: type	ÿ Hibicleanse	ÿ lodosorb
ÿ Medih	oney ÿ Santyl ÿ	Saf-gel ÿ Sterile Sali	ne ÿ Antimicrol	oial cleanser/cream	
ÿ Woun	d vac ÿ None ÿ Ot	her:			

16a. Did resident have a wound vac at any	time from July 17, 2015 – current?	ÿ Yes ÿ No
b. Date applied?//	Date removed?//	-
ÿ Medela ÿ Pico	ÿ Pressure:	
c. Date applied?//	Date removed?//	
ÿ Medela ÿ Pico	ÿ Pressure:	
d. Date applied?//	Date removed?	
ÿ Medela ÿ Pico	ÿ Pressure:	
17. Has the patient had a surgical procedu	re within 14 days prior to GAS culture	?
ÿ Yes ÿ No		
Procedure	Date	Incision Site
	///	
	//	
b. Wound infection ÿ c. Pharyngitis ÿ d. Bacteremia ÿ e. Pneumonia f. Joint Infection g. Necrotizing fasciitis h. Septic Shock	Yes ÿ No Date of onset	
19. Within 14 days of GAS culture, did the <i>apply</i> )	resident have any of the following sig	ns or symptoms? ( <i>mark ALL that</i>
	Date of onset (dd/mm/yy)	
a. ÿ Fever (≥100.5°F or 38°C)	//	Max temp recorded:

		Date of onset (dd/mm/yy)	
a.	ÿ Fever (≥100.5°F or 38°C)	//	Max temp recorded:
b.	ÿ Sore throat	//	
d.	ÿ Purulent discharge from wound	//	Site:
e.	ÿ Wound – warm on touch	//	Site:
f.	ÿ Wound – redness	//	Site:

g.	ÿ Edema at the site	///	Site:
h.	ÿ Increased pain at the site	///	Site:
i.	ÿ Joint – warm on touch	//	Site:
j.	ÿ Joint – redness	//	Site:

I Wh	a. Hospitalization date	resident taking within 14	1 days prior to GAS guil	turo?
1. VVII	at medications was the	Start Date	Finish Date	Indication
ntibio	otics			
'h a m	oth or on oution			
nem	otherapeutics			
teroi	ds			
nmur	nosuppressives			

23. Within 14 days prior to GAS culture was the resident ambulatory?

ÿ Yes ÿ No

hin 14 da	ays prior to (	GAS cultu	re, was the	e resident incontinent	of: (mark ALL that apply)	
ÿ Not Ir	ncontinent	ÿ Stool	ÿ Urine	ÿ Urinary catheter	ÿ Colostomy/Ileostomy	ÿ Unknowr
25. Did the resident participate in the following within 14 days prior to GAS culture (mark ALL that apply):						
a.	ў РТ/ОТ			Times in 14	l day period:	
b.	ÿ Speech p	athology		Times in 14	l day period:	
C.	ÿ Podiatry			Times in 14	l day period:	
d.	ÿ Other:			Times in 14	l day period:	
	ÿ Not Ir the resid a. b. c.	ÿ Not Incontinent  the resident particip  a. ÿ PT/OT  b. ÿ Speech p c. ÿ Podiatry	ÿ Not Incontinent ÿ Stool  the resident participate in the  a. ÿ PT/OT  b. ÿ Speech pathology c. ÿ Podiatry	ÿ Not Incontinent ÿ Stool ÿ Urine the resident participate in the following v a. ÿ PT/OT b. ÿ Speech pathology c. ÿ Podiatry	ÿ Not Incontinent ÿ Stool ÿ Urine ÿ Urinary catheter  the resident participate in the following within 14 days prior to  a. ÿ PT/OT Times in 14  b. ÿ Speech pathology Times in 14  c. ÿ Podiatry Times in 14	a. ÿ PT/OT Times in 14 day period: b. ÿ Speech pathology Times in 14 day period: c. ÿ Podiatry Times in 14 day period:

Study ID #: \_CHŸŸŸ

### Appendix 3. Invasive GAS in Long Term Care Facility 2016 Wound Care Survey

Form Approved; OMB No. 0920-1011 Exp. Date 03/31/2017

A. Employee Background	1. Name:	2. Age:				
3. Sex: ÿ Male ÿ Female	4. Employed at Facility since:	/				
5. What is your level of professional tra	nining on the wound care team?	ÿ RN ÿ MD ÿ LPN ÿ LVN ÿ Other				
6. a. Have you received training in	infection control practices?	ÿ Yes ÿ No ÿ Unknown				
b. If yes, when was your most red	cent training?	$\ddot{y} \leq 1 month \   \ddot{y}  2\text{-}6  months  \ddot{y}  6\text{-}12 months  \ddot{y}  > 1 year$				
B. Wound care	sults do you see per day? more					
8. On average, how many patients with	wounds do you see per day? ÿ (	0-10 ÿ 10-20 ÿ 20-30 ÿ 30 or more				
9. a. When evaluating a new consult	or reassessing an old patient, d	o you perform a full skin examination? ÿ Yes ÿ No				
b. If so, how do you report new w ÿ Medical Chart ÿ Report	younds found on your exam? to Nurse ÿ Report to Doctor ÿ	Other				
10. Is there a standardized risk assessm Scale) ÿ Yes ÿ No ÿ Unknown	ent tool used to document skin	breakdown/ pressure ulcer formation (e.g. Braden				
11. How often do you reassess wounds						
ÿ Daily ÿ 3-7 days ÿ 8-14 days ÿ Monthly ÿ Quarterly ÿ Other:						
12. What types of care do you perform on the wound care team?						
ÿ Incision and Drainage ÿ Undressing/Redressing ÿ Cleaning wound ÿ Wound vac care ÿ Other:						
13. Have you ever discovered pieces of foam/cotton gauze present in the wound from previous dressing changes? ÿ Yes ÿ No ÿ Unknown						

# Appendix 3. Invasive GAS in Long Term Care Facility 2016 <u>Wound Care Survey</u>

C. Negative-pressure wound therapy	14. Have you been specifically trained in the use of negative-pressure wound therapy? ÿ Yes ÿ No				
15. If so, when was your most recent tra	aining? $\ddot{y} \le 1$ month $\ddot{y}$ 2-6 months $\ddot{y}$ 6-12months $\ddot{y} > 1$ year				
16. How many residents require negative	re-pressure wound therapy/wound vac?				
17. What type of wound vac is used at y	our facility?				
18. Who is responsible for the original p	placement and replacement of the wound vac?				
ÿ Patient RN ÿ CNA ÿ MD ÿ	Only wound care team ÿ Other				
19. Who is allowed to change the woun	d vac cartridges and settings? (select more than 1 if applicable)?				
ÿ Patient RN ÿ CNA ÿ MD ÿ	Only wound care team  ÿ Other				
20. How often is a patient with a wound	I vac reassessed?				
ÿ Daily ÿ 2-3xweek ÿ Weekly ÿ Monthly ÿ Other					
21. Are their patients per week are foun	d to have full drainage cartridges or fluid backing up into the drainage tubing?				
22. If yes, how would this issue be repo	rted?				
ÿ Medical Chart ÿ Report to Nurse ÿ R	eport to Doctor ÿ Other				
23. When replacing the wound vac on the	he same patient, are any of the following re-used?				
(select more than 1 if applies)					
ÿ foam/gauze ÿ adhesive dressing	ÿ drainage tubing ÿ other				
24. If worsening wound is observed, is the wound vac replaced before a physician consult?					
ÿ Yes ÿ No ÿ Symptoms specific					
25. If symptoms specific please specify what symptoms would prompt you to replace the wound vac <i>before</i> a physician consult?					
26. What symptoms for a "worsening wound" prompts a physician consult?					
ÿ change in character of drained fluid ÿ	ÿ increase in fluid drainage ÿ increasing erythema ÿ pain ÿ increase in size				

						FU
Parish	Village	GPS Coordinates: Lat	Long	Elevation (m)	DD/MM/YY	
						г,

Form Approved OMB
No. 0920-1011
Exp. Date 03/31/2017

No	Animal	Animal ID	Owner	Age	Species	Gender	Breed	Current	Past Year	Comments:
	Sample	(tagged)		I = Infant	C=Cattle				Health:	· Symptoms
	ID .	. 55 /	Sample ID		G=Goats			(Vet)	(Owner)	<ul> <li>Abortion/stillbirth history</li> </ul>
		Or	-	M=Middle	S=Sheep	F=Female		S=Sick	S=Sick	<ul> <li>location of origin for slaughterhouse</li> </ul>
			Or	Age				H=Healthy	H=Healthy	animals)
		Name/Color		A=adult		C=Castrate		A=aborted	A=Aborted	*Common RVF symptoms: decreased appetite, decreased
		(not tagged)	Name	A=auun						milk production, nose/eye discharge, diarrhea, jaundice,
										prostration, lymph node swelling
1										
2										
3										
4										
5										
6										
7										
8										
9										
10										
11										
12										
13										
14										
15										
16										
17										
18										
19										
20										
21										
22										

Public reporting burden of this collection of information is estimated to average 1 minute 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; ATTN: PRA (0920-1011)

								Form Approved OMB
Parish	Village	GPS Coordinates: Lat	Long	Elevation (m)	DD	/MM	/YY	No. 0920-1011
								Exp. Date 03/31/2017

		T =		Ι _	T		1_	L	
No		Animal ID	Owner	Age	Species	Gender			Comments:
	Sample	(tagged)			C=Cattle	M=Male	Health	Health	· Symptoms
	ID		Sample ID		G=Goats		Status:	Status:	<ul> <li>Abortion/stillbirth history</li> </ul>
		Or		M=Middle	S=Sheep	F=Female	(Vet)	(Owner)	· location of origin for slaughterhouse
			Or	Age			S=Sick	S=Sick	animals)
		Name/Color				C=Castrate	H=Healthy	H-Healthy	*Common RVF symptoms: decreased appetite, decreased
		(not tagged)	Name	A=adult			Δ-aborted	$\Lambda = \Lambda$	milk production, nose/eye discharge, diarrhea, jaundice,
		(not taggea)	rearrie				rt-abortea	A-Aborted	prostration, lymph node swelling
-									prostation, tymph hode swelling
-									
	l	ı		1	1	1		1	

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Sample ID	Date/

Form Approved OMB No. 0920-1011 Exp. Date 03/31/2017

<u>Human</u>	Questior	nnaire

Participant classification (A/B/C/D)	Team (1/2)	Site (1/2/3/etc)

Form Completed by						
Name:	Position:	District:				
		<del>-</del>				
Section 1.	Assessment Participant	Information				
ID Number:	Family Name:	English Name	Age:	Gender:		
Village/Town:	Parish:	Sub-County:				
District:	Nationality:	_ Marital status:	ngle			
GPS Coordinates: Lat	Long	Elevation		_		
Section 2.	Epidemiological Risk Fa	ctors and Exposures				
1. Education level:	None	dary Post-Secondary Unkn    Housewife Student Come   Great Comestic animals Yes No   No ontact with (tick all that apply)   Poultry Dogs others speci-   During milking During graing or handling raw meat Step Secondary Yes No   On handling or preparing meat     No on handling or preparing meat     No nandling or	Child Animal Head PLETE LIVESTOCK FO Unkn (If NO/I  fy razing Grooming eeping near animals Unkn Unkn Yes No Yes No We birth Yes No Unkn No Unkn Others (specify) Unkn (If NO/Unk Others specify Other specify Jnkn, skip to #21)  O Unk Porons/ovals Other NO/Unknown, skip Yes No Unkr Unkn (If NO/Unk Unkn, skip to #21)  O Unk Others Specify Unk Others Specify Unk Others Specify Unk Other Specify Other S	alth worker  Butcher  DRM ON LAST PAGE) Unkn, skip to #7)  G		



		Form Approved OME
Sample ID	Date//	No. 0920-1011
		Exp. Date 03/31/2017

32. Have you had <i>el nino</i> (have you had more rainfall than usual) rains in the last one year?  Yes  No Unkn  33. Have you had flooding in this sub-county in past one year?  Yes  No Unkn (If NO/Unkn, skip to #35)  34. If yes, which months do you get flooding  35. In the past year, have you seen more mosquitoes than usual in this village?  No Unkn  36. In the past year, have you been bitten by more mosquitoes than usual?  Yes  No Unkn  37. In the past year, have you been using a mosquito net?  Yes  No Unkn (If NO/Unkn, skip to #39)  38. If no, why  39. In past year, have you sprayed animals against external parasites?  Yes  No Unkn  40. Have you done indoor spraying against mosquitoes in the last one year?  Yes  No Unkn  41. Have you been the forest/bush in the past one year Yes  No Unkn
Knowledge & Attitude Questions  42. Have you heard about Rift Valley Fever Disease?   Yes
Thank you for your Time End of Interview
Section 3. Specimen Information
Specimen identification number:
Specimen collection date:/ (DD/MM/YYYY)
Laboratory testing date:/ (DD/MM/YYYY)
Results/Titer level: IgM IgG



Sample ID	Date/

Form Approved OMB No. 0920-1011 Exp. Date 03/31/2017

#### **Livestock Assessment Form**

Sec	ction 1.	Herd Demographics
1)	What is your relationship to the livestock?   Owner	☐ Hardsman ☐ Other(specify)
2)		P Yes No Unkn Other (specify)
3)		area Grazing ground Other, specify
4)	Number of animals currently owned? Cattle	
,	Poultry	_ Dogs Cats
5)	What is the herd's typical grazing pattern? ☐ Shared	Enclosed Non-grazing Other (specify)
6)	In the past year, has the herd left the village? Yes	□No □Unknown
	If yes, why? Nomadic grazing Trade Gift	:/dowry
	If yes, to where? If yes, how many months ago? <1 month 1-3 m	II
	If yes, how many months ago? $\  \  \  \  \  \  \  \  \  \  \  \  \ $	nonths 3-6 months 6-12 months
Sec	ction 2.	Herd Health Status
_\		
7)	In the past year, has your <u>cattle</u> had unusual:	
		If yes, how many?
	• Stillbirths?	If yes, how many?
	Deaths in adults?    Yes   No   No   No   No   No   No   No   N	If yes, how many?
		If yes, how many? If yes, what?
8)	In the past year, has your <b>goats</b> had unusual:	ii yes, what?
0)		If yes, how many?
	, , , <u> </u>	If yes, what?
9)	In the past year, has your sheep had unusual:	
	· Abortions?	If yes, how many?
		If yes, how many?
		If yes, how many?
		If yes, how many?
	<ul> <li>Other health problems? ☐Yes ☐No</li> </ul>	If yes, what?
*C0	ommon DVE symptoms: docroased apportite, docroased milk	production, nose/eye discharge, diarrhea, jaundice, prostration, lymph node swelling
		Herd Treatment
JCC	Stion 3.	nord incamont
10)	) Have your <u>cattle</u> received:	
		If yes, what vaccines?
	<ul> <li>Insecticide treatment?  Yes No</li> </ul>	If yes, how often?
	Other treatments? ☐ Yes ☐ No	If yes, what treatment?
11)	Have your <b>goats</b> received:	
		If yes, what vaccines?
		If yes, when?
		If yes, what treatment?
12)	) Have your <u>sheep</u> received:	
		If yes, what vaccines?
		If yes, when?
	<ul> <li>Other treatments? ☐ Yes ☐ No</li> </ul>	If yes, what treatment?

#### **Telephone Interview Form**

SECTION A: General	exposure and	demographics.	Circle response.

Did you (your child) visit the Oak Leaf Dairy farm?
 Yes No

#### (If YES) ► PROCEED TO QUESTION 2

- (If NO) ► "Did anyone else in your household go to the farm? IF yes, may we speak to them? Go to question 2. If No, "Thank you for your time and participation, I have no further questions."
  - Since your visit to the farm have you been ill (defined as diarrhea (3 or more loose stools per day), vomiting, or abdominal cramps)?
     Yes No

#### (If NO) ► PROCEED TO QUESTION 3

(If YES) ► "Are there other members of your household that went to the farm and have not been ill as defined above?"

If yes, "May we ask to interview and proceed to question 3.

If no, "Thank you for your time someone else from the health department may call you back to ask additional questions about your illness."

3.	Are th	nere others ir	n your house	ehold who	visited	the farm	and hav	e also ı	not be	en ill?
	Yes	No								
	If yes	, How many f	family memb	oers visite	d the fa	rm and a	re not ill'	?		

4. What is your (your child's) birthdate?	//	(mm/dd/yyyy)
---	----	--------------

- What is your (your child's) gender?
   Male Female Prefer not to answer
- 6. What is your (your child's) town of residence?

## SECTION B. Hand-to-mouth habits "Let's talk about hand-to-mouth habits"

7. In general, do you (does your child) chew on or bite your (their) fingernails?

Yes No Don't Know

8. In general, do you (does your child), suck your (their) thumb or fingers? Yes No Don't Know

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# SECTION C. Prior Animal exposure "Let's talk about prior animal exposures"

9.	Do you (does your child) live on a property where farm animals such as cattle, sheep, or goats are kept?
	Yes No Don't Know
IF I	NO, SKIP TO SECTION D
10.	Which of these animals are kept on the property where you live (where your child lives)? Please answer yes or no.
	Cattle Yes No
	Sheep Yes No
	Goats Yes No
	Other Yes No
	Specify other animals kept on property:
	(If yes) How long have you owned: (months)
SECTIO	N D: Oak Leaf Dairy farm visit to tour.
"Let's t	alk about your / your child's visit to the Oak Leaf Dairy farm."
11.	Did you (your child) visit the farm more than once? Yes No
12.	On what date did you (your child) visit the farm?/ (mm/dd/yyyy)
13.	Approximately how much time did you (your child) spend at the farm? Less than 1 hour Between 1 and 2 hours Between 2 and 3 hours More than 3 hours
14.	(skip if only one visit) On what date did you (your child) visit the farm?//(mm/dd/yyyy)
15.	(skip if only one visit) Approximately how much time did you (your child) spend at the farm?
	Less than 1 hour Between 1 and 2 hours Between 2 and 3 hours More than 3 hours
16.	(skip if only one visit) On what date did you (your child) visit the farm?/(mm/dd/yyyy)
17.	(skip if only one visit) Approximately how much time did you (your child) spend at the farm?
	Less than 1 hour Between 1 and 2 hours Between 2 and 3 hours More than 3 hours
18.	Did you (your child) attend the Goat Keeping 101 class before Open House on March 12 <sup>th</sup> ? Yes No Don't Know

"Now, let's talk about areas on the farm. In this first section we will discuss is the baby goat barn."

19.	Did you (your child) enter the baby goat barn? Yes No IF NO SKIP TO QUESTION 34	Don'	t Know		
20.	Did you (your child) use hand sanitizer <u>BEFORE</u> visiting the baby Yes No Don't Know	goat	barn?		
21.	Did you (your child) sit on the ground in the baby goat barn?	Yes	No	Don't k	Cnow
22.	Did you (your child) touch/pet the <i>adult</i> goats in the baby goat	barn?	Yes	No D	on't Know
23.	Did you (your child) feed the <i>adult</i> goats in the baby goat barn?	Yes	No [	Oon't Kno	)W
24.	Did you (your child) enter a pen with the baby goats? (If yes) Did you (your child) sit on the ground in the pen? (If yes) Did you (your child) sit on a hay bale in the pen?	Yes Yes Yes	No No No	Don't k Don't k Don't k	(now
25.	Did you (your child) touch/pet the baby goats?	Yes	No	Don't k	
26.	Did you (your child) hold/snuggle the baby goats?	Yes	No	Don't	Know
27.	Did you (your child) kiss the baby goats?	Yes	No	Don't	Know
28.	Did the baby goats lick you (your child)?	Yes	No	Don't	Know
29.	Did you (your child) feed the baby goats?	Yes	No	Don't	Know
30.	Did you (your child) touch a railing while at the baby goat barn?	Yes	No	Don't	Know
31.	Did you (your child) use your cell phone in the baby goat barn?  If Yes:		Yes	No	N/A
	Did you (your child) talk on your phone?	Yes	No	Don't	Know
	Did you (your child) text on your phone?	Yes	No	Don't	Know
	Did you (your child) take pictures with your phone?	Yes	No	Don't	Know
	(If yes) Did you (your child) take pictures with goats?	Yes	No	Don't	Know
	Did you (your child) place your phone down (on hay bales/o Yes No Don't Know	n raili	ngs/on	floor)?	
32.	Did you (your child) use hand sanitizer <u>AFTER</u> visiting the baby g No Don't Know	oat ba	arn?	Ye	S
33.	Did you (your child) use baby wipes to clean your (their) hands barn?  Yes No Don't Know	<u>AFTER</u>	visiting	the bab	y goat

### "Now, let's talk about the adult goat barn."

,	g			
34.	Did you (your child) visit the adult goat barn? Yes No Don't IF NO, SKIP TO QUESTION 43	t Know		
35.	Did you (your child) use hand sanitizer <u>BEFORE</u> visiting the add Yes No Don't Know	ult goat	barn?	•
36.	Did you (your child) touch/pet the adult goats at the adult goa	at barn?	Yes	No Don't Know
37.	Did you (your child) feed the adult goats at the adult goat bar	n? Yes	No D	on't Know
38.	Did you (your child) touch a railing while at the adult goat bar	n? Yes	No I	Don't Know
39.	Did you (your child) sit on a hay bale while at the adult goat b	arn? Ye	s No	Don't Know
40.	Did you (your child) use your cell phone while at the adult goa	nt barn?	Yes	No N/A
	If yes:			
	Did you (your child) talk on your phone?	Yes	No	Don't Know
	Did you (your child) text on your phone?	Yes	No	Don't Know
	Did you (your child) take pictures with your phone?	Yes	No	Don't Know
	If yes, Did you (your child) take pictures with	goats?		
	Did you (your child) place your phone down (on hay b	ales/on	railin	gs/on floor)?
	Yes No Don't Know			
41.	Did you (your child) use hand sanitizer <u>AFTER</u> visiting the adul Yes No Don't Know	t goat b	arn?	
42.	Did you (your child) use baby wipes to clean your (their) hand	s <u>AFTER</u>	visitii	ng the adult goat
	barn? Yes No Don't Know			
"Now,	let's talk about other things you may have done at the farm."	,		
43.	Did your child have a pacifier at the farm?  Yes No N/A			
	If yes, did your child take it into the baby goat barn?	Yes	No	Don't Know
	If yes, did your child take it near the adult goat barn?	Yes	No	Don't Know
	<b>3</b>			
44.	Did your child have a sippy cup at the farm?  Yes No N/A			
	If yes, did your child take it into the baby goat barn?	Yes	No	Don't Know
	If yes, did your child take it near the adult goat barn?	Yes	No	Don't Know
	•			
45.	Did you (your child) chew gum while at the farm? Yes	No	Don	t Know
46.	Did you (your child) eat candy while at the farm? Yes	No	Don	t Know
47.	Did you bring a stroller on the farm?	Yes	No	N/A
	If yes, did you bring it in the baby goat barn?	Yes	No	Don't Know
	If yes, did you bring it near the adult goat barn?	Yes	No	Don't Know

"Now, let's talk about other animal contact	ct you may have had at the farm."
---	-----------------------------------

- 48. Did you (your child) use hand sanitizer <u>BEFORE</u> touching any animals besides goats?
  - Yes No Don't Know
- 49. Did you (your child) touch/pet the rabbits? Yes No Don't Know
- 50. Did you (your child) touch/pet the dogs?

  Yes No Don't Know
- 51. Did you (your child) use hand sanitizer <u>AFTER</u> touching any animals besides goats? Yes No Don't Know
- 52. Did you (your child) use baby wipes to clean your (their) hands <u>AFTER</u> touching any animals besides goats? Yes No Don't Know

#### "Now I'm going to ask you some questions about eating and drinking at the farm."

53. Did you (your child) eat any food products you may have purchased while at the farm?

Yes No Don't Know

#### (IF NO SKIP TO QUESTION 58)

If yes, did you (your child) use hand sanitizer **BEFORE** eating?

Yes No Don't Know

54. Did you (your child) eat cheese bought from the farm while at the farm?

Yes No Don't Know

If yes, where did you (your child) eat the cheese bought from the farm (circle all that apply)?

Farm store Picnic table Adult goat barn Baby goat barn

Milking parlor Other Don't Know

55. Did you (your child) drink milk (pasteurized) bought from the farm while at the farm?

Yes No Don't Know

If yes, where did you (your child) drink milk (pasteurized) bought from the farm (circle all that apply)?

Farm store Picnic table Adult goat barn Baby goat barn

Milking parlor Other Don't Know

56. Did you (your child) drink raw milk (unpasteurized) bought from the farm while at the farm?

Yes No Don't Know

If yes, where did you (your child) drink raw milk (unpasteurized) bought from the farm (circle all that apply)?

Farm store Picnic table Adult goat barn Baby goat barn

Milking parlor Other Don't Know

57. Did you (your child) eat caramels bought from the farm while at the farm? Yes No Don't Know If yes, where did you (your child) eat caramels bought from the farm (circle all that apply)? Farm store Picnic table Adult goat barn Baby goat barn Don't Know Milking parlor Other 58. ► Did you (your child) taste any samples at farm? Yes No Don't Know (IF NO SKIP TO QUESTION 63) Did you (your child) use hand sanitizer BEFORE tasting the sample? Yes No Don't Know 59. Did you (your child) taste cheese samples from the farm while at the farm? No Don't Know If yes, where did you (your child) eat the cheese sample from the farm (circle all that apply)? Farm store Picnic table Adult goat barn Baby goat barn Milking parlor Other Don't Know 60. Did you (your child) drink milk (pasteurized) samples from the farm while at the farm? No Don't Know If yes, where did you (your child) drink milk (pasteurized) sample from the farm (circle all that apply)? Farm store Picnic table Adult goat barn Baby goat barn Milking parlor Don't Know Other Did you (your child) drink raw milk (unpasteurized) sample from the farm while at the 61. farm? Yes No Don't Know If yes, where did you (your child) drink raw milk (unpasteurized) sample from the farm (circle all that apply)? Farm store Picnic table Adult goat barn Baby goat barn Milking parlor Other Don't Know Did you (your child) eat caramels samples from the farm while at the farm? 62. No Don't Know If yes, where did you (your child) eat caramels samples from the farm (circle all that apply)? Farm store Picnic table Adult goat barn Baby goat barn Milking parlor Other Don't Know 63. ▶ Did you (your child) bring food to the farm and eat it on the farm (for example, to have a picnic)? Yes No Don't Know (IF NO SKIP TO QUESTION 64) Did you (your child) use hand sanitizer BEFORE eating? Yes Don't Know No Where on the farm did you (your child) eat the food (circle all that apply)?

Farm store Picnic table Adult goat barn Baby goat barn Milking parlor Other Don't Know

64. Did you (your child) drink any beverages that you brought with you at the farm?

Yes No Don't Know

#### (IF NO SKIP TO QUESTION 65)

Where on the farm did you (your child) drink it (circle all that apply)?

Farm store Picnic table Adult goat barn Baby goat barn

Milking parlor Other Don't Know

65. Did you (your child) drink any water from a faucet at the farm? Yes No Don't Know

"Now I'm going to ask you some questions about activities after leaving the farm."

66. After visiting the farm, did you (your child) stop to eat? Yes No Don't Know If yes, did you (your child) wash your hands before eating? Yes No Don't Know If yes, did you (your child) use hand sanitizer before eating? Yes No Don't Know

67. After visiting the farm, did you (your child) come home with any of the following?

Dirty or stained clothing? Yes No Don't Know Dirty shoes? Yes No Don't Know

- 68. Did you (your child) remove shoes before walking in the home? Yes No Don't Know
- 69. Did you (your child) change your clothes immediately when you returned home?

Yes No Don't Know

#### **SECTION E: General knowledge and awareness**

"Now I would like to ask you some questions about general knowledge on interaction with animals."

- 70. In general, were you (was your child) aware that some diseases can be spread by having contact with farm animals?

  Yes No Don't Know
- 71. In general, were you (was your child) aware that some diseases can be spread by having contact with surfaces at a farm, such as the ground, railings?

Yes No Don't Know

#### SECTION H: Pre-existing medical conditions and medication use

"Now I would like to ask you a few questions about your (your child's) health in March, 2016. We would like to know about long-standing medical conditions or other specific medical conditions in the month of March. You do not need to answer the questions if you don't want to."

72. During the month of March did you (your child) have any of the following medical conditions?

PLEASE READ EACH CONDITION AND CHECK YES NO DK

	Yes	No	DK
Diabetes			
Kidney Disease			
If YES ► Are you/your child on dialysis?			
Organ or Bone Marrow Transplant			
Leukemia or Cancer			
If YES ► Treatment with radiation or chemotherapy in previous			
month?			

"I would now like to ask some questions about medications that you (your child) may have been taking in the month of March."

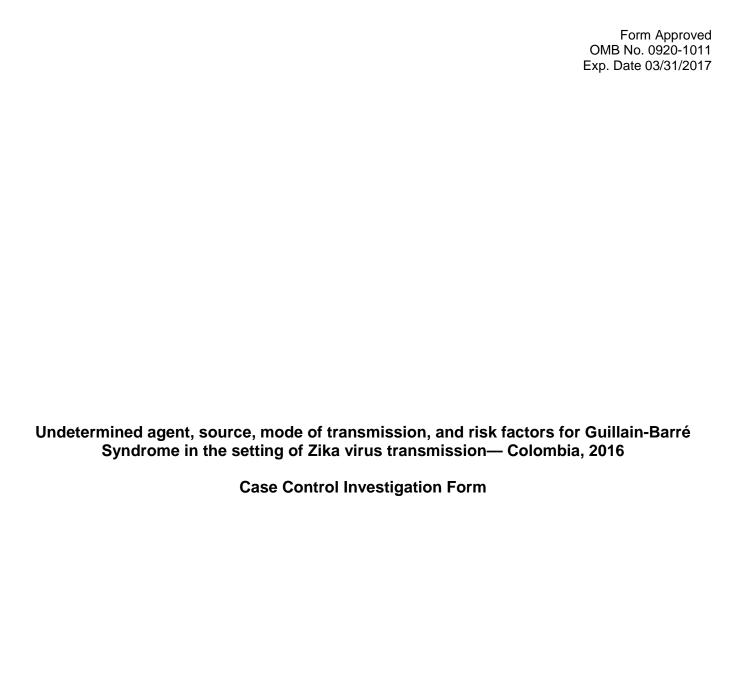
73. In the month of March, did you (your child) take any of the following types of medications?

PLE	ASE READ EACH MEDICATION AND CHECK YES NO DK	Yes	No	DK
	Any antibiotics			
	Any oral steroid, such as Prednisone?			
	Any immune-suppressing medication, such as to treat juvenile arthritis?			

"Now, I would like to gather some additional information."

74.	Did you hear or see in the media that the Department of	Public Health	was requesting	ill
	and non-ill people that visited the farm to contact them?	? Yes	No	

	and non-iii people that visited the farm to contact them?
75.	What is your (your child's) race? White Black/African American American Indian/Alaskan Native Native Hawaiian or Pacific Islander Asian Other, specify: Unknown Prefer not to answer
74.	Do you consider yourself (your child) to be of Hispanic ethnicity? Yes No Prefer not to answer
75.	If you (your child) visited the farm more than once, please describe any differences between your visits:



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Investi	gation ID Num	ber COL		_	•	$\Box$ Control	
"B" fo	r the first contr		econd control, an	ed a"D" for the thi		an "A" for the case p For example, the se	
Intervi	ewer:			Date of Interview:		_///	
Neuro	Symptom Onse	et Date for Case		YYY	DD	MM YYYY	
The fo	llowing questio	ns are to be asked	of cases AND co	ontrols during the	interview:		
1. Curi	rent Address:	(Street)		// (Province		<u></u>	
		(Street)	(Town)	(Province	e)	(District)	
2. Ons	et Address:		1	1		1	
(for ca	ses only if diffe	rent from above; wi	here cases spent	most nights in the	2 months p	orior to neuro onset)	
a CDC	. C 1: (6	Daniel Communication			C		Е
3. GPS 4. Sex:		Onset for cases; curl  ☐ Female	rent for controls)	):	S,	·	E
			_		_	_	
			-			ellow   Other:	
		veloped first neuro			·		
7. Wha	nt is your occup	ation?					
8. Wha	at is the highest	level of education	you completed?				
□ Prir	mary   Secon	ndary   Technic	al Universi	ty   None			
9. Did	you travel anyv	vhere two weeks pr	ior to onset of sy	mptoms?			
	□ Ye	es $\square$ No	□ Unkı	nown			
Where	:						
10. Ha	ve you ever bee	en told by a clinicia	n that you have a	any of the followin	g medical	conditions?	
	☐ Diabetes	☐ High blood p	·	☐ Heart disease		☐ High colester	ol
	☐ Stroke	☐ Kidney disea	se	☐ Liver disease		☐ Rheumatologi	c disease
	☐ Asthma	□ COPD	1	☐ Cancer		☐ Surgery (within symptom onset)	2 months of
	☐ Other neur	ologic illness:					
	☐ Take any reprednisone):	medication or have	any condition th	at might impact yo	our ability t	o fight infections (e.ş	).
11.		nths prior to		neuro onset date fo	r case), ha	ve YOU been sick at	all?
	□ Ye		□ Unkı	nown			
	b. If so, when	did you first feel si	ck?		/		

☐ Fevers	☐ Chills	☐ Nausea or Vomiting	☐ Diarrhea						
☐ Muscle pains	☐ Joint pains	☐ Skin rash	☐ Abnormally red eye						
☐ Headache	☐ Pain behind eye	es	☐ Confusion						
☐ Abdominal pain	☐ Coughing	☐ Runny nose ☐ Sore	throat   Calf pain						
☐ Pruritus									
d. If so, did you see a	doctor or go to the ho	spital for this illness?							
	No □ Unknown	Which hospital? _							
e. If so, did they draw	any blood for testing	?	Unknown						
a. In the 2 months price	or to //	(neuro onset date for case)	), has anyone in your						
	HOUSEHOLD been sick at all?  ☐ Yes ☐ No ☐ Unknown								
b. If so, when did the		per become sick?	/						
		members have (check all that a							
☐ Fevers	☐ Chills	☐ Nausea or Vomiting	☐ Diarrhea						
☐ Muscle pains	☐ Joint pains	☐ Skin rash	☐ Abnormally red eye						
☐ Headache	☐ Pain behind eye	es   Stiff neck	☐ Confusion						
☐ Abdominal pain	☐ Coughing [	☐ Runny nose ☐ Sore	e throat $\Box$ Calf pain						
☐ Pruritus		·	•						
Which vaccinations h	ave you received and	when?							
☐ Information verified on vaccine card ☐ Information provided verbally  Vaccine DD MM YYYY Additional doses									
a. MMR		/	4000						
b. Polio			<del></del>						
	/								
c. Yellow fever	<del></del> /_								
c. Yellow fever d. BCG	/	/							
	/ /	/							
d. BCG	/	/							
d. BCG e. DPT	/	/							
d. BCG e. DPT f. HiB	/	/							
d. BCG e. DPT f. HiB g. Pneumococcal	/								
d. BCG e. DPT f. HiB g. Pneumococcal h. Meningitis i. Hepatitis B	/	apanese encephalitis, etc.):							

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							CO	OL
	he two months bor on your proper			oms, wha	at pets, i	farm, or o	other animals	have lived in your
	□ Dogs	□ Cats	☐ Mice/rats		□ Pet	t birds	□ P	Pet lizards /turtles
	☐ Goats	☐ Sheep	□ Cows	□ Chi	ckens	□ Pigs		Other
15. In t tap?	he two months b	pefore onset of n	eurologic sympto	oms, hov	v often l	have you	gotten your d	rinking water from the
	□Almost alway	ys (>75%)	□Often (25-75	5%)		□Rarel	y (<25%)	□Never (0%)
	If ever, was the	water boiled or	treated?	□Yes		No	□Unknow	n
	he two months b		eurologic sympto	oms, hov	v often l	have you	gotten your d	rinking water from a
	□Almost alway	ys (>75%)	□Often (25-75	5%)		□Rarel	y (<25%)	□Never (0%)
	If ever, was the	water boiled or	treated?   Yes			□No		□Unknown
17. In t	he two months b	efore onset of n	eurologic sympto	oms, hov	v often	do you wa	alk around ba	refoot outside?
	□Almost alway	ys (>75%)	□Often (25-75	5%)		□Rarel	y (<25%)	□Never (0%)
18. In t pond?	he two months b	pefore onset of n	eurologic sympto	oms, hav	e you s	wam or w	aded in a fres	shwater river, stream, or
	□Daily	□Weekly	$\square$ Monthly	□Rare	ly ( <on< td=""><td>ce per mo</td><td>onth)</td><td>□Never</td></on<>	ce per mo	onth)	□Never
19. Ho	w much time do	you spend outdo	oors each day?					
	□<1 h	our	□1-4 hours		□5-8	hours	□>8	3 hours
20. Ho	w often do you w	vear long sleeve	s and pants?					
	□Almost alway	ys (>75%)	☐Often (25-75%) ☐Rare		rely (<25%) □Nev		ever (0%)	
22. Do	you normally we	ear insect repell	ant?					
	□Almost alway	ys (>75%)	□Often (25-75	5%)	□Rare	ely (<25%	5) □No	ever (0%)
23. Do	you leave the wi	indows open at	your house?					
	☐Yes, during t	he day  \sum Yes	, at night □Ye	es, all tin	nes	□Wind	lows are not l	eft open at this house
24. Ho	w many of your	windows or doo	rs have intact sci	reens?				
	□All of them		□Some of the	m		□None	of them	
25. Do	es your home use	e any of the follo	owing for air con	nditioning	g (check	k all that a	apply)?	
	•	nditioning (at lea	•	∃Fans		□None		
	w often do you h /cistern, septic ta		standing water ar	ound the	outside	e of your l	house (e.g. bu	ickets, water
		2-3 times/week	□Once/weel	k □E	Every of	her week	□Never	
27. Are	these containers	s covered?						
	□ Yes □ N	No 🗆 Unkno	own					

28. In the two months before onset of neurologic symptoms, have you handled any dead animals?

	□Yes   Which?		known					
29. In 2	2016, have yo	u eaten or drunk an	y of the following f	foods at least on	ice per week (check all that apply)?			
	$\square$ Beef	☐ Lamb	☐ Chicken	☐ Fish	☐ Shellfish			
	☐ Milk	☐ Cheese	☐ Yogurt	☐ Fresh salad	/ uncooked greens			
30. <u>Hu</u>	ghes Disabilit	y Score: (Date re	corded/	_/)				
	Hughes Disability Score (0 to 6):							
$[0 = Complete\ recovery;\ no\ sequelae,\ 1 = Minor\ symptoms\ and\ capable\ of\ running,\ 2 = Able\ to\ walk\ 10\ metres\ or\ more\ without\ assistance\ but\ unable\ to\ run,\ 3 = Able\ to\ walk\ 10\ metres\ with\ help,\ 4 = Bedridden\ or\ chairbound$								

(unable to walk 10 meters with help), 5 = Requiring assisted ventilation for at least part of the day, 6 = Dead]

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Formulario Aprobado OMB No. 0920-1011

Fecha de vencimiento: 03/31/2017

Cuestionario de Caracterización Síndrome de Guillain-Barré – Colombia, 2016



	ro de Identificación COL Caso
"A" pa	ara el paciente que sea un caso, una "B" para el primer control, una "C" para el segundo control Por o, el segundo individuo control emparejado con el caso número 8 sería marcado como "COL-008-C".
Entrev	istador: Fecha de Entrevista:/ DD MM AAAA
Fecha	de Inicio de Síntomas Neurológicos:  DD MM AAAA  DD MM AAAA
Las sig	guientes preguntas son para ser realizadas durante la entrevista:
1.	Dirección actual:
	Dirección Ciudad o Municipio Distrito o Departamento
2.	Dirección donde se presentaron los síntomas:
	(solamente si es diferente de la dirección actual; donde los casos pasaron el mayor número de noches en los dos meses previos al inicio del cuadro neurológico)
3.	Coordenadas GPS (Inicio de síntomas): N, O
4.	Sexo:   Masculino   Femenino
5.	Pertenencia étinica: ☐ Indígena ☐ ROM/Gitano ☐ Raizal ☐ Palenquero
	□ Negro/mulato/Afrocolombiano □ Otro
6.	Edad cuando presentó los primeros síntomas neurológicos: años
7.	¿Cuál es su ocupación?
8.	Cuál es su nivel educativo (marque si fue cursado completo):   Primaria   Secundaria   Técnica   Universitaria   Ninguno
9.	Dos semanas antes del inicio de los síntomas neurológicos viajó a otro lugar?
	A dónde:
10	. ¿Ha sido informado por algún médico que usted padece alguna de las siguientes condiciones médicas?
	☐ Diabetes ☐ Presión Arterial Alta ☐ Enfermedad del Corazón ☐ Colesterol Elevado
	☐ Accidente Cerebrovascular (Derrame cerebral) ☐ Enfermedad Renal ☐ Enfermedad Hepática
	☐ Enfermedad Reumatológica
	☐ Asma ☐ Enfermedad Obstructiva Pulmonar Crónica ☐ Cáncer
	☐ Cirugía (dentro de los meses de inicio de síntomas)
	☐ Otra enfermedad neurológica:



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			a condición que pueda afectar su	-				
11.	a. ¿En los dos meses ar enfermo (a)?  □ Sí □ No	nteriores a// DD MM  □ No sabe	(fecha de inicio de cuadro r AAAA	neurológico), estuvo				
	<b>b.</b> Si estuvo enfermo (a	.), ¿cuándo se sintió enfe	ermo(a) por primera vez?/	//				
	c. Si estuvo enfermo (a)	) ; Oué síntomas tuvo?	DD (Marque todos los que aplican)	MM AAAA				
	☐ Fiebre	☐ Escalofrío	☐ Nausea o Vómito	☐ Diarrea				
	☐ Dolor muscular	☐ Dolor articular	☐ Rash cutáneo	☐ Ojos anormalmente rojos				
	☐ Dolor de cabeza	☐ Dolor retro ocular	☐ Rigidez nucal	☐ Confusión				
	☐ Dolor abdominal	□ Tos □ Sec	creción nasal	☐ Dolor de pantorrillas				
	<b>d.</b> Si estuvo enfermo (a ☐ Sí ☐ No		fue al hospital por esta enfermed	lad?				
	□ Sí □ No □ No sabe     ¿Cuál médico? ¿Qué hospital?							
	e. Si estuvo enfermo (a)	), ¿le tomaron muestra d	e sangre? □ Sí □ No □	No sabe				
12.		DD MM A e haya estado enfermo (a		s neurológicos), ¿hubo				
	<b>b.</b> Si la respuesta es afin	rmativa, ¿en qué fecha s	e enfermó la primera persona de	su hogar?				
	DD MM AAAA  c. Si alguien en su hoga	nr estuvo enfermo (a) ¿Q	ué síntomas tuvo? (Marque tod	os los que aplican)				
	☐ Fiebre	☐ Escalofrío	☐ Nausea o Vómito	☐ Diarrea				
	☐ Dolor muscular	☐ Dolor articular	☐ Rash cutáneo	☐ Ojos anormalmente rojos				
	☐ Dolor de cabeza	☐ Dolor retro ocular	☐ Rigidez nucal	☐ Confusión				
	☐ Dolor abdominal	□ Tos □ Sec	creción nasal	☐ Dolor de pantorrillas				
13.	¿Qué vacunas ha recibi	do y cuándo?						
	☐ Información verifica	nda en el carné de vacuna	as   Información proveída ven	rbalmente				
	<ul><li>a. Triple viral (SRP o N</li><li>b. Polio</li><li>c. Fiebre Amarilla</li></ul>	MMR) DD MM A	AAAA  Dosis adicionales:					



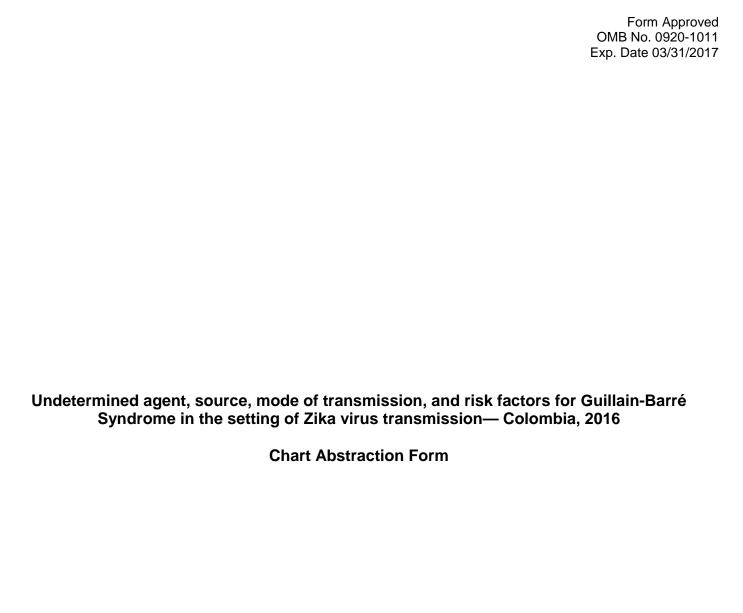
SAI	LUD	COL			
	d. BCG e. DPT f. Haemophilus Influenza B g. Neumococo h. Meningitis i. Hepatitis B j. Otras vacunas (ejemplo: rabia, encefalitis Japone ¿Cuál?				
14.	En los dos meses antes del inicio de los síntomas no animales vivieron en su casa o su propiedad? (Marc			les de granja u otro	os
	☐ Perros ☐ Gatos ☐ Ratone	es/ratas	☐ Pájaros o	lomésticos	
	☐ Lagartijas /tortugas ☐ Cabras ☐ Ovejas	□ Vacas	☐ Gallinas		
	☐ Cerdos ☐ Otros				
15.	En los dos meses antes del inicio de los síntomas no estaba hervida o tratada?	eurológicos, ¿qué ta	n frecuentemei	nte tomó agua que i	no
	$\Box$ Casi siempre (>75%) $\Box$ A veces (25-75%)	%) □Ra	ra vez (<25%)	□Nunca (0%)	Į.
16.	En los dos meses antes del inicio de los síntomas no quebrada o lago?	eurológicos, ¿qué ta	n a menudo toi	nó agua de un pozo	o, rio
	□Casi siempre (>75%) □A veces (25-75%)	□Rara vez (<	<25%) □N	(unca (0%)	
	Si tomó alguna vez, ¿hirvieron o trataron el agua?	□Sí □	]No □	No sabe	
17.	En los dos meses antes del inicio de los síntomas no descalzo(a)?	eurológicos, ¿qué ta	n a menudo sal	ió de su casa a can	ninar
	□Casi siempre (>75%) □A veces (25-75%)	□Rara vez (<	<25%) □N	□Nunca (0%)	
18.	En los dos meses antes del inicio de los síntomas no arroyo, o lago?  □Diariamente □Semanalmente □Mensua		·	propios medios un de una vez al mes)	rio,
	□Nunca				
19.	¿Cuántas horas al día está al aire libre?				
	□Menos de 1 hora □1-4 hora □5-8 ho	ras □>8 horas			
20.	Qué tan frecuente es el uso de ropa con mangas y p	antalón largos?			
	□Casi siempre (>75%) □A menudo (25-75%)	□Rara vez (<	<25%) □N	unca (0%)	
21.	¿Usa repelente de insectos?				
	$\Box$ Casi siempre (>75%) $\Box$ A menudo (25-75%)	□Rara vez (<	<25%) □N	unca (0%)	
22.	¿Fumiga dentro su vivienda?				
,	☐ Casi siempre (>75%) ☐ A menudo (25-75%)	□Rara vez (<	<25%) □N	unca (0%)	



23.	¿Deja	las ventanas	de su casa abiert	as?				
	□Sí, c	lurante el día	□Sí, en la nocl	ne □Sí, todo	o el tiempo	□Las venta	anas no se deja	n abiertas en
24.	¿Cuán	itas de sus vei	ntanas o puertas	tienen angeos i	intactos?	esta casa	•	
	□Tod	as [	☐Algunas de ellas	□Ninguna				
25.	¿Tiene	en en su casa	alguno de los sig	guientes tipos d	le aire acondi	cionado?		
	□Aire	acondicionad	o central	Aire acondicion	ado por habita	ción	□Ninguno	
26.			tiene recipientes ernas, tanques, inse		su casa donde	puede hab	er agua estano	eada? (por
	□Dian	riamente	l2-3 veces/semana	□Una vez	a la semana	□Semana	a de por medio	□Nunca
27.	Estos	recipientes se	encuentran tapa	dos				
	□Sí	□No □N	No sabe					
28.	muert	o? □Sí	ntes del inicio de □No □No	sabe			ulado algún ar	nimal
29.			consumido algun odos los que apli		entes alimento	s o bebidas	s por lo menos	s una vez a la
	a.	☐ Carne	☐ Cordero	□ Pollo	☐ Pescad	lo 🗆 Ma	ariscos	
	b.	☐ Leche	☐ Queso	☐ Yogurt	□ Ensala	da / verdura	as crudas	
30.	Puntaj	e de Discapa	cidad de Hughes	: (Fecha de re	gistro:	//	)	
		Puntaje de I	Discapacidad de	Hughes (0 a 6)	: □	] Descono	cido	
	[0= R	ecuperación (	completa: sin sec	cuelas. 1= Sínt	omas menore	s v capaz a	le correr. 2= 1	Puede

COL-\_\_\_\_ - \_\_

[0= Recuperación completa; sin secuelas, 1= Síntomas menores y capaz de correr, 2= Puede caminar 10 metros o más sin asistencia pero no puede correr, 3=Puede caminar 10 metros con ayuda, 4= Postrado en cama o en silla de ruedas (no puede caminar 10 metros con ayuda), 5= Requiere ventilación asistida por lo menos una parte del día, 6=Muerto]



Study 1	ID Number COL	Encount	er level (Brighton 1-5) or not neuro (6):
The ID physici	-	ase number (for example CO	L-01). Information as documented by attending
The fo	llowing pages are to be abstracte	ed from the medical records	exam:
Chart A MRN:		Ab	straction Date: / / MM DD YYYY
1.	First name:	Mio	ldle name:
2.	Paternal name:		ternal name:
3.	Age (years):		e of birth://
			MM DD YYYY
4.	Sex: ☐ Male ☐ Fem	ale	
5.	Patient address:		
6.	Patient phone number:		
7.	Date of neuro symptom onset:	/ / Dar MM DD YYYY	e first sought care:/ / MM DD YYYY
	Date of admission:/ MM DD		e of discharge/death: / / / MM DD YYYY
8.	Discharged to:		
	☐ Home ☐ Rehab/skilled n	ursing facility     Transferr	ed Died Died Other (specify)
		CURRENT I	LLNESS
9.	How long from onset until hosp	oital admission?	minutes/hours/days/weeks
10	• What were the initial neurologi from PE, symptoms from HPI)	c symptoms (i.e. within the th	ree days of illness onset)? (check all that apply, signs
	☐ Leg weakness	☐ Arm weakness	☐ Diplopia/Ophthalmoplegia
	☐ Leg numbness/paresthesias	☐ Arm numbness/paresthe	sias
	☐ SOB / respiratory distress	☐ Gait imbalance (not wea	kness)/ataxia

☐ Face weakness ☐ Dysarthria

☐ Arm weakness

 $\square$  Leg numbness/paresthesias  $\square$  Arm numbness/paresthesias

11. What neurologic symptoms occurred AT ANY TIME during the neuro illness? (check all that apply, signs from

☐ Dysphagia

☐ Hyporeflexia/areflexia

PE, symptoms from HPI)

☐ Leg weakness

 $\square$  Dysautonomia

 $\square$  Diplopia/Ophthalmoplegia

☐ Face numbness/paresthesias

tudy ID Number	COL		I	Encounter 1	level (Br	righton 1-	-5) or not neu	ıro (6):	
□ SOB / res	piratory distress	☐ Gait	imbalance (	not weakn	ess)/atax	ia	☐ Hand cl	umsiness/ataxia	
☐ Hyporefle	exia/areflexia	☐ Face	weakness	☐ Dysai	thria	□ Dys	sphagia	☐ Dysautonon	nia
•	om onset until max point during this n		•	•			minutes/hour	s/days/weeks	
☐ Unable to	walk without assi	stance (e.	g. cane, wal	ker)			☐ Unable	to walk at all	
☐ Admitted	to the hospital		☐ Admitted	d to the IC	U/CCU		☐ Intubate	ed	
<b>14.</b> If any blood	was taken for this	neurologi	c illness, ple	ase fill out	t the foll	owing for	r the INITIA	L blood draw:	
	DD YYYY	WBC _	HgF	3	Plts _		Na	K	
BUN	Cr	Glucose	TBi	li	AST _		ALT	AlkPhos	-
	cumented hyporef documentation of				s	□ No		Unknown	
☐ Yes b. If yes, whi	□ No ch:	□ Unkr	nown						
☐ Hyperrefl	exia 🗆 Inc	reased to	ne/spasticity	□ B	abinski/I	Hoffman		Sustained clonus	
17. Was there an	y sensory level do	cumented	?	□ Ye	s	□ No		Unknown	
	LABORA	TORY, II	MAGING, A	ND ELEC	CTROPH	IYSIOLO	OGIC STUDI	IES	
	r puncture (LP) do			Yes			☐ Unknow		
	/ RBCS DD YYYY		WBCS	Prot	tein (mg/	/dL)	Glucose	e (mg/dL)	_
			_IgG index_	Oli	goclonal	bands	IgG syı	nthesis	
	/ RBCS		WBCS	Prot	tein (mg/	/dL)	Glucose	e (mg/dL)	_
Differential_			_IgG index_	Oli	goclonal	bands	IgG syı	nthesis	
19. Did they rece	eive any targeted tr	eatment (	(IVIG/steroid	ds/plasma	exchange	e) for this	s neuro illnes	ss?	
IVIG	☐ Yes	□ No	□ Unkno	wn	Start d	late	/_ MM DD	/	
Plasma excha	ange	□ No	□ Unkno	wn	Start d	late		/	

Study I	D Number COL		Er	counter le	evel (Brighton	1-5) or no	t neuro	(6):
	Steroids	□ Yes □ No	□ Unknow	n	Start date	/ MM		YYYY
	Mechanical ventilation	□ Yes □ No	□ Unknow	n	Start date	/		YYYY
	Other	☐ Yes ☐ No	☐ Unknow	n	Start date			YYYY
20.	Did the patient receive	blood transfusion	/blood produc	ts? (other	than IVIG)	IVIIVI	DD	1111
	□ Yes □ No □ U	Jnknown which	one		Start			/
21.	Were any of the follow	ing diseases tested	d for? If so, w	hat was th	ne result? (inclu	MN uding spec		
	a. Campylobacter jejun	i	□ Y€	es 🗆 No	Result:			<del></del>
	b. Mycoplasma pneumo	oniae	□ Y€	es 🗆 No	Result:			
	c. Haemophilus influen	zae	□ Y€	es 🗆 No	Result:			
	d. Salmonella spp.		□ Y€	es 🗆 No	Result:			
	e. Cytomegalovirus (Cl	MV)	□ Y€	es 🗆 No	Result:			<del></del>
	f. Epstein-Barr virus (E	(BV)	□ Y€	es 🗆 No	Result:			<del></del>
	g. Varicella-zoster viru	s (VZV)	□ Y€	es 🗆 No	Result:			<del></del>
	h. Human immunodefic	ciency virus (HIV	) \( \sum \text{Y}\epsilon	es 🗆 No	Result:			
	i. Enterovirus / Rhinovi	irus	□ Y€	es 🗆 No	Result:			
	j. Arboviruses		□ Y€	es 🗆 No	Result:			
	k. Other		□ Y€	es 🗆 No	Result:			
22.	. Was neuro imaging dor	ne? If so, what wa	s the result? (	Transcrib	e the impression	on)		
	☐ Yes ☐ No Result		•		•	,		
								/
23.	Were electro-diagnostic	cs done (e.g. EMC	G)? If so, what	were the	results? (Trans			OD YYYY sion)
	☐ Yes ☐ No Result							
						Date _		/
24.	What was the GBS Brig	ghton level?	1	2	3 4	5 N	MM ]	OD YYYY
	of Diagnostic Certain					T _		
Level 1	ee of an alternative diagnosis f	Level 2 For weakness			Level 3	Level 4*		NOT a case
Acute of	onset of bilateral and relatively sed or absent deep tendon refl	y symmetric flaccid w		mbs		* Lacking	_	1101 a cusc

Monophasic illness pattern with weakn	ed by clinical	fulfill minimal		
plateau			case criteria	
Albuminocytologic dissociation	CSF with a total white cell count < 50			
(elevation of CSF protein level above	cells/mm <sup>3</sup> (with or without CSF protein			
laboratory normal value and CSF	elevation above laboratory normal value) or			
total white cell count < 50	if CSF not collected or results not available,			
cells/mm <sup>3</sup> )	and electrodiagnostic studies consistent			
	with GBS			
Electrophysiologic findings				
consistent with GBS				

	ANTECEDENT ILLNESS						
25.	<b>a.</b> ) In the 2 months prior	or to neuro onset date, did	I the individual experience	ce an acu	ite illness? (	other than their neur	
	illness)? ☐ Yes	□No □ Unknown					
26	minutes/hours/days/wee	or acute illness onset until eks they report having or wh					
20.	☐ Fevers	☐ Chills	☐ Nausea or Vomiting		☐ Diarrhe		
				3			
☐ Muscle pains ☐ Joint pains ☐ Skin rash ☐ Conjunctivitis							
	☐ Headache	☐ Pain behind eyes	☐ Stiff neck		☐ Confusi	ion	
	☐ Back pain	☐ Abdominal pain	☐ Coughing		□ Runny	nose	
	☐ Sore throat	☐ Calf pain	☐ Pruritis				
	<b>b.</b> ) If any blood was take Date//	ken for this acute illness, WBC _	please fill out the follow HgB	-		blood draw: K	
	DD MM YY BUN Cr		TBili AST _		ALT	AlkPhos	
	<b>c.</b> ) Were they hospitaliz	zed for this acute illness?		□ Yes	s □ No	☐ Unknown	
	<b>d.</b> ) Did they receive any	y blood products / IVIG f	for this illness?	□ Yes	s 🗆 No	☐ Unknown	
	What product? Date?// MM _DD _YYYY						
	e.) Did they receive pla	smapheresis / plasma exc If yes, date?/ MM	_	□ Yes	s 🗆 No	☐ Unknown	
27.		ilable for dengue from th		☐ Yes	s 🗆 No	□ Unknown	
				_	_	_	
28.		ilable for chikungunya fi		☐ Yes	s 🗆 No	☐ Unknown	
	If yes, please specify:						

udy I	D Number	COL	<del></del>	Encounter le	evel (Bri	ghton 1-5)	or not net	ıro (6):
			lable for Zika from this			□ Yes	□ No	□ Unknown
			PAST MEDICAI	L, SOCIAL ANI	) FAMII	Y HISTO	RY	
30.	What medical c	condition	s are listed in the admiss	sion history and	physical	(H&P)?		
	☐ Hypertensio	n	☐ Diabetes	☐ HIV	☐ Aut	oimmune o	lisorder	<del></del>
	☐ Prior GBS		☐ Hemoglobinopathy	☐ B12 deficie	ency	☐ Cance	r	
31.	What social cor	nditions a	are listed in admission H	I&P?	•			
	☐ Alcohol use	;	☐ Drug use	☐ Tobacco		☐ Other		
32.	What condition	s are list	ed in family history of H	1&P?				
	☐ Autoimmun	e disorde	er (specify)		□ Can	cer (specif	y)	
	☐ Hemoglobin	nopathy (	(specify)		□ Neu	ro (specif	y)	



Formulario Aprobado OMB No. 0920-1011 Fecha de vencimiento: 03/31/2017

Instrumento para la recolección de datos de historias clínicas. Caracterización de casos con Síndrome de Guillain-Barré – Colombia, 2016

NACIONAL DE SALED		
NACIONAL DE SALED NT	INSTITUTO	
SALED NT	NACIONAL DE	
	SALED	Nıí

Número de Identificación

COL-		

Nivel (Brighton 1-5) o no neurológico (6): \_\_\_\_

El número de identificación comienza con los 3 dígitos del número de caso (por ejemplo COL-001). Información según lo documentado por el médico tratante

	evisor de Historia Clínica:	
INL	ímero de Historia Clínica:	
		Segundo Nombre:
		Segundo Apellido:
3.	Edad (años):	Fecha de Nacimiento: / / MM DD YYYY
4.	Sexo: ☐ Masculino ☐ Femenino	
		ección completa, ciudad o municipio y departamento):
6.	Número de teléfono del paciente:	
7.	a.) Fecha de ingreso hospitalario:/ MM DD Y	
	<b>b.</b> ) Fecha en la que buscó atención por primera v	vez:// MM
	<b>c.)</b> Fecha de egreso hospitalario/muerte: /_ MM DI	/ OYYYY
8.	Egresó hacia:	
	☐ Hogar ☐ Centro de Rehabilitación ☐	☐ Remitido a otra institución hospitalaria
	☐ Muerte ☐ Otro (Especifique)	
		ENFERMEDAD ACTUAL
9.	¿Cuánto tiempo transcurrió desde el inicio de los	s síntomas hasta el ingreso al hospital?minutos/horas/días/semanas
10	· ·	les dentro de los tres días previos al inicio de la enfermedad? (Marque nen físico y síntomas de historia de enfermedad actual)
	$\square$ Debilidad en extremidades inferiores $\square$	Debilidad en extremidades superiores   Diplopia/Oftalmoplejía
	☐ Adormecimiento de extremidades inferio	pres/parestesias
	☐ Adormecimiento de extremidades superi	ores/parestesias
	☐ Adormecimiento de la cara/parestesias	
	☐ Dificultad para respirar/distress respirato	orio   Trastornos de la marcha (sin debilidad)/ataxia

	Número de Identificación COL Nivel (Bri	ghton 1-5) o no neurológico (6):							
	☐ Trastornos de la motricidad manual/ataxia								
11.	☐ Hiporeflexia/areflexia ☐ Debilidad en la cara ☐ Disartria ☐ 11. ¿Qué síntomas neurológicos ocurrieron en CUALQUIER MOMENTO durante l todas las opciones que apliquen, signos del examen físico y síntomas de historia	•							
	☐ Debilidad en extremidades inferiores ☐ Debilidad en extremidades sup	eriores   Diplopia/Oftalmoplejía							
	☐ Adormecimiento de extremidades inferiores /parestesias								
	☐ Adormecimiento de extremidades superiores /parestesias	☐ Adormecimiento de extremidades superiores /parestesias							
	☐ Adormecimiento de la cara /parestesias	☐ Adormecimiento de la cara /parestesias							
	☐ Dificultad para respirar / distress respiratorio ☐ Trastornos de la march	a (sin debilidad)/ataxia							
	☐ Trastornos de la motricidad manual/ataxia								
	☐ Hiporeflexia/areflexia ☐ Debilidad en la cara ☐ Disartria ☐	Disfagia							
12.	12. ¿Cuánto tiempo transcurrió desde el inicio hasta la presentación de los síntomas minutos/horas/días/semanas	neurológicos más severos?							
13.	<b>13.</b> Marque todas las opciones que apliquen al paciente que se presentaron al momer neurológico:	nto de mayor severidad del cuadro							
	☐ Incapacidad para caminar sin asistencia (por ejemplo: bastón, caminador)	)   Incapacidad total para caminar							
	☐ Ingreso al hospital ☐ Ingreso a Unidad de Cuidado Intensivo (UC	CI)   Intubación							
14.	<b>14.</b> Si se extrajo muestra de sangre como parte de los análisis de laboratorio para el complete las siguiente información de la muestra de sangre obtenida INICIALM	9 1							
	Fecha/ / Recuento de blancos Hemoglo MM DD YYYY	bina Plaquetas							
	Sodio Potasio BUN Creatinina Gluco	sa Bilirrubina Total							
	AST ALT Fosfatasa Alcalina								
15.	<b>15.</b> ¿Se documentó hiporeflexia/areflexia? ☐ Sí ☐ No ☐	Desconocido							
16.	<b>16. a.</b> ) ¿Hubo evidencia de signos de motoneurona superior?								
	□ Sí □ No □ Desconocido								
	<b>b.</b> ) En caso afirmativo, ¿Se documentó algunos de los siguientes hallazgos	s?							
	☐ Hiperreflexia ☐ Aumento en el tono/espasticidad ☐ Babins	ki/Hoffman ☐ Clonus sostenido							

Número de Identificación  Número de Identificación	COL		I	Nivel (Brighton 1-5) o	no neui	ológico	o (6):
17. ¿Se documentó algún nivel sensit	tivo?	□ Sí	□ No	☐ Desconocido	)		
☐ Guante y Bota ☐ C	Cervical		☐ Dorsal	☐ Lumbar			
LABORATORIO	, IMÁGENI	ES DIA	GNOSTICAS Y E	STUDIOS ELECTRO	FISIOL	.ÓGIC	OS
18. ¿Se llevó a cabo una punción lun	nbar?	□ Sí	□ No	☐ Desconocide	0		
Fecha punción lumbar/_			ento de eritrocitos _	Recuento d	de leuc	ocitos _	
	DD YYYY		sa (mg/dL)	_ Diferencial			
Fecha punción lumbar/_ MM	/ DD YYYY		ento de eritrocitos _	Recuento o	de leuc	ocitos _	
Proteínas (mg/dL) Diferencial		Glucos					
	<b>19.</b> ¿Recibieron algún tratamiento específico para manejar esta enfermedad neurológica (Inmunoglobulina intravenosa/esteroides/recambio plasmático)?						
a. Inmunoglobulina intraveno	osa □ Sí □	□ No	☐ Desconocido	Fecha de inicio			/
b. Recambio plasmático	□ Sí □	□ No	☐ Desconocido	Fecha de inicio	/	/	/
					MM	DD	YYYY
c. Esteroides	□ Sí □	□ No	☐ Desconocido	Fecha de inicio			
1 37 (1 12 2 1		¬		F 1 1	MM	DD	YYYY
d. Ventilación mecánica	□ S1 L	□ No	☐ Desconocido	Fecha de inicio			YYYY
e. Otro	□ Sí □	□ No	☐ Desconocido	Fecha de inicio			
					MM	DD	YYYY
20. ¿Recibió el paciente transfusión o intravenosa)	de sangre o	algún o	otro hemoproducto?	otros diferentes a Inn	nunoglo	bulina	
□ Sí □ No □ Descond	ocido ¿Cu	ál?		Fecha de inicio			_/ YYYY
21. ¿Fueron algunos de los siguientes espécimen y el tipo de prueba)	s patógenos	estudia	ndos? En caso afirm	nativo, ¿cuál fue el resu	ıltado?	(incluy	endo el
a. Campylobacter jejuni			□ Sí □ No	Resultado:			

□ Sí

**b.** Mycoplasma pneumoniae

☐ No Resultado: \_\_\_\_\_

=	Instruction
•	24
	NACIONAL DE
_	SALUD

Número de Identificación COL		_	Nivel (Brighton 1-5) o no neurológico (6):
<b>c.</b> Haemophilus influenzae	□ Sí	□ No	Resultado:
<b>d.</b> Salmonella spp.	□ Sí	□ No	Resultado:
e. Citomegalovirus (CMV)	□ Sí	□ No	Resultado:
f. Virus Epstein-Barr (EBV)	□ Sí	□ No	Resultado:
g. Virus Varicella-zoster (VZV)	□ Sí	□ No	Resultado:
h. Virus de Inmunodeficiencia Humana (VIH)	□ Sí	□ No	Resultado:
i. Enterovirus / Rhinovirus	□ Sí	□ No	Resultado:
<b>j.</b> Arbovirus	□ Sí	□ No	Resultado:
k. Otro. ¿Cuál?	□ Sí	□ No	Resultado:
23. ¿Se llevaron a cabo pruebas electrodiagnósticas? (peresultado? (Transcriba el resultado reportado)  □ Sí □ No  Resultado			
Fecha // MM DD YYYY		_	
24. ¿Cuál fue el nivel de SGB en la escala de Brighton?	' 1	2	3 4 5
Niveles de Certeza Diagnóstica			

Nivel 1	Nivel 2	Nivel 3	Nivel 4*	Level 5	
Ausencia de un diagnóstico alternativo para debilidad					
Inicio agudo de debilidad flácida bilitaral y relativamente simétrica de las extremidades * Al carecer de					
Reflejos tendinosos profundos dismi	documentación				
Patrón de enfermedad monofásica con nadir de debilidad entre 12 horas y 28 días, seguido de para cumplir con					
meseta clínica					



Número de Identificación	COL-	Nivel (Brighton 1-5) o no neurológico (6):
vuillelo de lacitificación	COL	Triver (Brighton 1-3) ono neurologico (o).

Disociación albuminocitológica	LCR con un total de recuento de glóbulos	los criterios	
(elevación del nivel de proteínas en	blancos<50 células / mm3 (con o sin	mínimos de caso	
el LCR por encima del valor	elevación de proteínas en LCR sobre el		
normal de laboratorio y recuento	valor normal de laboratorio) o si el LCR		
total de glóbulos blancos en LCR	no fue recolectado o los resultados no		
<50 células / mm3)	están disponibles y los estudios de		
	electrodiagnóstico son consistentes con		
	SGB		
Hallazgos electrofisiológicos			
consistentes con SGB			

## ANTECEDENTES DE LA ENFERMEDAD

25.	aguda? (diferente a su enf	ermedad neurológica)	de síntomas neurológicos, tuvo  nta 30, sección Antecedentes Cl	•	
	□ Sí □ No □ Desconocido				
	<b>b.</b> ) ¿Cuánto tiempo se pre neurológica?1			eso hospitalario por la condición	
26.	a.) ¿Qué síntomas reporta apliquen)	ron haber tenido o qué sig	gnos fueron evidenciados? (Marq	ue todas las opciones que	
	☐ Fiebre	☐ Escalofrío	☐ Nausea o Vómito	☐ Diarrea	
	☐ Dolor muscular	☐ Dolor articular	☐ Rash cutáneo	☐ Conjuntivitis	
	☐ Cefalea	☐ Dolor retro ocular	☐ Rigidez nucal	☐ Confusión	
	☐ Dolor de espalda	☐ Dolor abdominal	□ Tos	☐ Secreción nasal	
	☐ Odinofagia	☐ Dolor de pantorrilla	a		
			urte de los análisis de la enfern nuestra de sangre obtenida INI		
	Fecha / / / MM		lancos Hemoglobina_	Plaquetas	
	Sodio Potas	io BUN	Creatinina Gluco	sa Bilirrubina Total	
	AST ALT_	Fosfatasa Alca	lina		
	c.) ¿Hubo hospitalizad	ción por esta enfermedad a	aguda? □ Sí □ No □ I	Desconocido	
	<b>d.</b> ) ¿Recibió alguna tr	ansfusión de cualquier he	moproducto/administración de In	nmunoglobulina intravenosa para	
	esta enfermedad agud	a? □ Sí □ No □	Desconocido		

Número de Identificación COL Nivel (Brighton)	n 1-5) o	no neu	rológi	co (6):
En caso afirmativo, ¿Qué producto?	Fecha:		/	/
		MM	DD	YYYY
e.) ¿Recibió plasmaféresis / recambio plasmático para esta enfermedad aguda?	□ Sí	□ No		Desconocido
En caso afirmativo, ¿qué fecha? // MM DD YYYY				
27. ¿Hay algún resultado de laboratorio para dengue en esta visita médica?	□ No	☐ De	sconoc	cido
Resultado □ Positivo □ Ne				
<b>28.</b> ¿Hay algún resultado de laboratorio para chikungunya en esta visita médica? ☐ Sí	□ No	□ De	sconoc	cido
<b>Resultado</b> □ Positivo □ Ne	gativo	□ Des	conoc	ido
29. ¿Hay algún resultado de laboratorio para zika en esta visita médica?	□ No	□ De	sconoc	cido
<b>Resultado</b> □ Positivo □ Ne	egativo	□ De	sconoc	eido
ANTECEDENTES CLINICOS, SOCIALES Y FAM	ILIARE	S		
<b>30.</b> ¿Qué antecedentes clínicos están registrados en la historia clínica de ingreso?				
☐ Hipertensión ☐ Diabetes ☐ VIH				
☐ Trastorno autoimmune. En caso afirmativo, ¿cuál?				
☐ SGB previo ☐ Hemoglobinopatía ☐ Deficiencia de Vitamina B12	2			
☐ Cancer. En caso afirmativo, ¿cuál?				
21 . On form the design of the configuration and the little in 1/2 in the income 0				
31. ¿Qué antecedentes sociales están registrados en la historia clínica de ingreso?				
☐ Uso de alcohol ☐ Uso de drogas ☐ Tabaquismo				
☐ Otros. En caso afirmativo, ¿cuáles?				
<b>32.</b> ¿Qué antecedentes familiares están registrados en la historia clínica de ingreso?				
☐ Trastornos autoinmunes (Especifique)				
☐ Cáncer (Especifique)				
☐ Hemoglobinopatías (Especifique)				
☐ Neurológicos (Especifique)				

### **Case Abstraction Form**

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Question	Code	Variable
RVCT number		RVCT
Last Name		Lname
First Name		Fname
Alternate Names/Nicknames/Aliases:		Alias
Date of Birth (MM/DD/YY)		DOB
Age (years)		Age
Gender (1=Male, 2=Female, 3=Other, 99=missing)		Sex
Race/Ethnicity (1=Black, 2=White, 3=Hispanic/Latino, 4= American Indian/Alaskan Native, 5=Native Hawaiian/Pacific Islander, 6=Asian, 7=Other, 99=Missing) [Mark all that apply]		Race
Tribe If American Indian, then specify tribe:		Tribe
<b>Tribe A residence</b> If lives on Tribe A reservation, specify which area: 1=northwest of Yuma, 2=southwest of Yuma, 3=south of Yuma		Residence
If lives elsewhere, specify		Other
Locating Information, if available:		
Addresses: Phone	S:	
How long at this address?		
Be sure to list any other known addresses during last 3 years.		
Country of Birth (1=United States, 2=Other [foreign-born], 99=missing)		Birth
If foreign-born, then specify country:		Country
Date of arrival (MM/DD/YY) For patients born outside the		Arrival

Public reporting burden of this collection of information is estimated to average 4 hours 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; ATTN: PRA (0920-1011)

United States,	
enter the date of arrival in the United States.	

### **TB Risk Factors**

HIV infection (0=No, 1=Yes, 99=Unknown)  Diabetes (0=No, 1=Yes, 99=Unknown)  If diabetic, most recent HbA1C  Chronic Renal Failure (0=No, 1=Yes, 99=Unknown)  Immunosuppression other than HIV (e.g. organ transplant, chemotherapy, medications such as steroids, TNF blockers. 0=No, 1=Yes, 99=Unknown)  Mental illness (0=No, 1=Yes, 99=Unknown) (Axis I diagnosis not related to substance abuse, e.g. mood disorders, schizophrenia, anxiety disorders) Injection drug use-recent (Within 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Injection drug use-ever (Prior to 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Injection drug use-recent (Within 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Non-injection drug use-recent (Within 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Non-injection drug use-ever (Prior to 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Non-injection drug use-ever (Prior to 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Non-injection drug use-ever (Prior to 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Excess alcohol use-recent (Within 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Excess alcohol use-recent (Within 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Smoking commercial tobacco regularly (i.e., most days) for at least 1 year at time of diagnosis (0=No, 1=Yes, 99=Unknown)  Homeless/unstable housing within 1 year of diagnosis? (0=No, 1=Yes, 99=Unknown)  Homeless/unstable housing >1 year before diagnosis? (0=No, 1=Yes, 99=Unknown)	Question	Code	Variable
Diabetes (0=No, 1=Yes, 99=Unknown)  If diabetic, most recent HbA1C  Chronic Renal Failure (0=No, 1=Yes, 99=Unknown)  Immunosuppression other than HIV (e.g. organ transplant, chemotherapy, medications such as steroids, TNF blockers. 0=No, 1=Yes, 99=Unknown)  Mental illness (0=No, 1=Yes, 99=Unknown) (Axis I diagnosis not related to substance abuse, e.g. mood disorders, schizophrenia, anxiety disorders)  Injection drug use-recent (Within 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Injection drug use-ever (Prior to 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Injection drug use-recent (Within 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Non-injection drug use-ever (Within 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Non-injection drug use-ever (Prior to 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Non-injection drug use-ever (Prior to 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Non-injection drug use-ever (Within 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Non-injection drug names (open ended) list all non-injection drugs used ever Excess alcohol use-recent (Within 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Excess alcohol use-ever (Prior to 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Smoking commercial tobacco regularly (i.e., most days) for at least 1 year at time of diagnosis (0=No, 1=Yes, 99=Unknown)  Homeless/unstable housing within 1 year of diagnosis? (0=No, 1=Yes, 99=Unknown)  Homeless/unstable housing >1 year before diagnosis? Home2			
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Immunosuppression other than HIV (e.g. organ transplant, chemotherapy, medications such as steroids, TNF blockers. 0=No, 1=Yes, 99=Unknown)   Mental illness (0=No, 1=Yes, 99=Unknown) (Axis I diagnosis not related to substance abuse, e.g. mood disorders, schizophrenia, anxiety disorders)   Injection drug use-recent (Within 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)   IDU_rvct (Within 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)   Injection drug use-ever (Prior to 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)   IDU_list (open ended) list all drugs injected ever   NIDU_rvct (Within 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)   Non-injection drug use-ever (Prior to 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)   Non-Injection drug use-ever (Prior to 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)   Non-Injection drug names (open ended) list all non-injection drugs used ever   EtOH_rvct (Within 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)   Excess alcohol use-ever (Prior to 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)   EtOH_ever (Prior to 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)   Tobacco for at least 1 year at time of diagnosis (0=No, 1=Yes, 99=Unknown)   Homeless/unstable housing within 1 year of diagnosis? (Deno, 1=Yes, 99=Unknown)   Homeless/unstable housing within 1 year of diagnosis? (Deno, 1=Yes, 99=Unknown)   Homeless/unstable housing variety (Deno, 1=Yes, 99=			
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schizophrenia, anxiety disorders)  Injection drug use-recent (Within 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Injection drug use-ever (Prior to 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Injection drug names (open ended) list all drugs injected ever  Non-injection drug use-recent (Within 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Non-injection drug use-ever (Prior to 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Non-Injection drug names (open ended) list all non-injection drugs used ever  Excess alcohol use-recent (Within 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Excess alcohol use-ever (Prior to 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Excess alcohol use-ever (Prior to 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Excess alcohol use-ever (Prior to 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Excess alcohol use-ever (Prior to 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Excess alcohol use-ever (Prior to 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Homeless/unstable housing within 1 year of diagnosis?  Home1 (0=No, 1=Yes 99=Unknown)  Homeless/unstable housing >1 year before diagnosis?	, , , , , , , , , , , , , , , , , , , ,		Mental
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Non-injection drug use-ever (Prior to 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Non-Injection drug names (open ended) list all non-injection drugs used ever  Excess alcohol use-recent (Within 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Excess alcohol use-ever (Prior to 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Smoking commercial tobacco regularly (i.e., most days) for at least 1 year at time of diagnosis (0=No, 1=Yes, 99=Unknown)  Homeless/unstable housing within 1 year of diagnosis? (0=No, 1=Yes 99=Unknown)  Homeless/unstable housing >1 year before diagnosis?  Home2	(Within 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)		
Prior to 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)   Non-Injection drug names (open ended) list all non-injection drugs used ever			NIDU ever
NIDU_list (open ended) list all non-injection drugs used ever  Excess alcohol use-recent (Within 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Excess alcohol use-ever (Prior to 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Smoking commercial tobacco regularly (i.e., most days) for at least 1 year at time of diagnosis (0=No, 1=Yes, 99=Unknown)  Homeless/unstable housing within 1 year of diagnosis? (0=No, 1=Yes 99=Unknown)  Homeless/unstable housing >1 year before diagnosis?  Home2			
(open ended) list all non-injection drugs used ever  Excess alcohol use-recent (Within 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Excess alcohol use-ever (Prior to 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Smoking commercial tobacco regularly (i.e., most days) for at least 1 year at time of diagnosis (0=No, 1=Yes, 99=Unknown)  Homeless/unstable housing within 1 year of diagnosis? (0=No, 1=Yes 99=Unknown)  Homeless/unstable housing >1 year before diagnosis?  Home2			NIDU list
Excess alcohol use-recent (Within 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Excess alcohol use-ever (Prior to 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Smoking commercial tobacco regularly (i.e., most days) for at least 1 year at time of diagnosis (0=No, 1=Yes, 99=Unknown)  Homeless/unstable housing within 1 year of diagnosis? (0=No, 1=Yes 99=Unknown)  Homeless/unstable housing >1 year before diagnosis?  Home2	•		
(Within 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Excess alcohol use-ever (Prior to 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Smoking commercial tobacco regularly (i.e., most days) for at least 1 year at time of diagnosis (0=No, 1=Yes, 99=Unknown)  Homeless/unstable housing within 1 year of diagnosis? (0=No, 1=Yes 99=Unknown)  Homeless/unstable housing >1 year before diagnosis?  Home2			EtOH rvct
Excess alcohol use-ever (Prior to 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Smoking commercial tobacco regularly (i.e., most days) for at least 1 year at time of diagnosis (0=No, 1=Yes, 99=Unknown)  Homeless/unstable housing within 1 year of diagnosis? (0=No, 1=Yes 99=Unknown)  Homeless/unstable housing >1 year before diagnosis?  Home2			
(Prior to 1 year of TB diagnosis. 0=No, 1=Yes, 99=Unknown)  Smoking commercial tobacco regularly (i.e., most days) for at least 1 year at time of diagnosis (0=No, 1=Yes, 99=Unknown)  Homeless/unstable housing within 1 year of diagnosis? (0=No, 1=Yes 99=Unknown)  Homeless/unstable housing >1 year before diagnosis?  Home2			EtOH ever
Smoking commercial tobacco regularly (i.e., most days) for at least 1 year at time of diagnosis (0=No, 1=Yes, 99=Unknown) Homeless/unstable housing within 1 year of diagnosis? (0=No, 1=Yes 99=Unknown) Homeless/unstable housing >1 year before diagnosis? Home2			
for at least 1 year at time of diagnosis (0=No, 1=Yes, 99=Unknown)  Homeless/unstable housing within 1 year of diagnosis? (0=No, 1=Yes 99=Unknown)  Homeless/unstable housing >1 year before diagnosis?  Home2			Tobacco
99=Unknown)  Homeless/unstable housing within 1 year of diagnosis?  (0=No, 1=Yes 99=Unknown)  Homeless/unstable housing >1 year before diagnosis?  Home2			
Homeless/unstable housing within 1 year of diagnosis?  (0=No, 1=Yes 99=Unknown)  Homeless/unstable housing >1 year before diagnosis?  Home2			
(0=No, 1=Yes 99=Unknown)  Homeless/unstable housing >1 year before diagnosis?  Home2			Home1
Homeless/unstable housing >1 year before diagnosis? Home2			
			Home2
\- ··-, · · •• •·····	(0=No, 1=Yes 99=Unknown)		
Use of homeless shelter within 1 year of diagnosis? 0=No,  Shelter1			Shelter1
1=Yes 99=Unknown			
Use of homeless shelter >1 year before diagnosis? 0=No, Shelter2			Shelter2
1=Yes 99=Unknown			
			Shelter_list
(open ended) list all homeless shelters used			
At least 1 night in correctional/detention facility within 1 Incarc1			Incarc1

year of diagnosis? (0=No, 1=Yes 99=Unknown)	
At least 1 night in correctional/detention facility >1 year	Incarc2
before diagnosis? (0=No, 1=Yes 99=Unknown)	
Incarceration facility names	Incarc_list
(open ended) list all correctional/detention facilities where	
stayed at least 1 night	
Residence in long term care facility within 1 year of	LTCF1
diagnosis?	
(0=No, 1=Yes 99=Unknown)	
Residence in long term care facility>1 year before	LTCF2
diagnosis?	
(0=No, 1=Yes 99=Unknown)	
If known exposure to TB case, exposure type:	TBexp
(1=own household, 2=homeless shelter, 3=jail, 4=other	
household, 5=bar, 6= hotel, 7=Other:)	ExpOth
List name of site if known:	ExpSite

# TB Case Characteristics

Question	Code	Variable
How was case recognized or detected?		Caserec
(1=symptoms, 2=contact investigation, 3=routine TB		
screening by healthcare provider, 4=incidental finding by		
healthcare provider, 5=other, 99=unknown)		
Cough (0=not present 1= present, 99=unknown)		Cough
Fever (0=not present 1= present, 99=unknown)		Fever
Night Sweats (0=not present 1= present, 99=unknown)		Sweats
Weight Loss (0=not present 1= present, 99=unknown)		Weight
Other TB Symptoms (list)		OthSx
Date of first symptom onset (Enter the first date the patient		DateSx
began experiencing symptoms in the format MM/DD/YY)		
Site of disease		TBSite
(1=pulmonary, 2=extrapulmonary, 3=both pulmonary and		
extrapulmonary)		CXRrslt
<b>Diagnostic CXR result</b> (1=Negative, 2=Abnormal, possibly TB, 3=Abnormal, not consistent with TB, 4=Unknown [not completed or not available])		CARISIL
Diagnostic chest radiograph (CXR) result date (Enter the		CXRdate
date of the patient's most recent CXR completed as part of		
current diagnostic workup leading to patient's current		
diagnosis of TB. MM/DD/YY)		
Cavitary disease on CXR? (0=No, 1=Yes, 99=Unknown)		CavCXR
Cavitary disease on CT? (0=No, 1=Yes, 99=Unknown)		CavCT

Sputum AFB smear positive disease? (0=No, 1=Yes,	Sputum
2=Sputum never submitted)	
Sputum smear converted to negative (0=No, 1=Yes ≤2	Smearconv
months of treatment, 2=Yes >2 months of treatment,	
3=Unknown/NA)	
Other site AFB smear positive? (0=No, 1=Yes,	OthSmear
99=Unknown)	OthSite
Specify Site:	
Culture-confirmed disease? (0=No, 1=sputum only,	Culture
2=non-sputum specimen, 3=both sputum and non-sputum	
specimens, 4=specimens never submitted, 99=Unknown)	
If culture confirmed, list GENType	GENType
Culture converted to negative (0=No, 1=Yes ≤2 months of	Cxconv
treatment, 2=Yes >2 months of treatment, 3=Unknown/NA)	C/(00/14
Diagnosis date (MM/DD/YY) (the earliest date of the	Dxdate
following: positive smear, positive culture, positive PCR test,	
or abnormal chest x-ray/CT scan)	
Drug susceptibility based on molecular testing	Suscept_Mol
(1=Pan-susceptible, 2=INH resistance, 3=rifampin	. –
resistance, 4=multiple resistance, including MDR TB,	
88=pending, 99=unknown)	
Drug susceptibility based on culture	Suscept_DST
(1=Pan-susceptible, 2=INH resistance, 3=rifampin	
resistance, 4=multiple resistance, including MDR TB,	
88=pending, 99=unknown)	
INH resistance level (highest concentration at which isolate	INHR
is resistant)	
RIF resistance level (highest concentration at which isolate	RIFR
is resistant)	
	Ott. D
Specify any other detected resistance	Oth_R
Diagnostic TST result (Enter the patient's TST result, if	TST
completed as part of the diagnostic workup leading to the	
patient's current diagnosis of TB. 1=negative, 2=positive,	
3=positive with conversion [≥10mm increase in last 2 years],	
4=not done due to prior positive TST, 5=not done for other	
reason, 99=result unknown)	
Diagnostic TST reading (mm reading)	TSTmm
Diagnostic TST date (MM/DD/YY)	TSTdate
, ,	
Diagnostic QFT result (Enter the patient's qualitative QFT	QFT
result, if completed as part of the diagnostic workup leading	
to the patient's current diagnosis of TB. 1=negative,	
2=positive, 3=indeterminate, 4=not done, 99=unknown)	
Diagnostic QFT value (result-nil). (Enter the quantitative	QFTvalue

result of the patient's current QFT result, 99=Unknown.	
Leave blank if not performed.)	
Diagnostic QFT date (MM/DD/YY)	QFTdate
<b>Diagnostic T.Spot result</b> (Enter the patient's qualitative result, if completed as part of the diagnostic workup leading to the patient's current diagnosis of TB. 1=negative, 2=positive, 3=indeterminate, 4=borderline, 5=not done, 99=unknown)	TSpot
<b>Diagnostic T.Spot value (</b> Enter the quantitative result of the patient's current result, 99=Unknown. Leave blank if not performed.)	TSpotvalue
Diagnostic T.Spot date (MM/DD/YY)	TSpotdate
<b>Treatment</b> (1=On treatment, 2=Completed full treatment, 3=Completed partial treatment, 4=Died during treatment, 5= Died before treatment, 6=died after treatment, 7=awaiting treatment initiation, 8=refused treatment, 99=Unknown)	TBrx
Start date of initial TB treatment (Enter the date of antituberculosis medication in the format MM/DD/YY.)	TBRxdate
If applicable, date of change to MDR TB regimen (Enter the date of antituberculosis medication in the format MM/DD/YY.)	MDRRxdate
List MDR TB regimen	MDRregimen
<b>Date of treatment completion if done</b> (Enter the date of antituberculosis medication in the format MM/DD/YY.)	Rxcomp
History of loss to follow-up or non-compliance during this TB treatment course (0= No, 1= Yes, 99=Unknown)	TBfu
If died, then enter date of death (MM/DD/YY)	Deathdate
If died, then enter cause of death	Deathcause

Previous TB episodes and LTBI history

Question	Code	Variable
Prior TB disease? (0=No, 1=Yes, 99=Unknown)		PrevTB
Year of previous diagnosis (YYYY)		Prevyr
If prior TB, exposure type (1=own household, 2=homeless		PrevTBexp
shelter, 3=jail, 4=other household, 5=bar, 6= hotel, 7=Other:		
		PrevTBexp oth
If prior TB, drug susceptibility (1=Pan-susceptible, 2=INH		Prevresist
resistance, 3=rifampin resistance, 4=multiple resistance, incl.		
MDR TB, 88=pending, 99=unknown)		
If prior TB, Genotype (GENType)		PrevGENty

	pe
<b>TB treatment completed (</b> 0= No, 1= Yes, 2=In progress,	PrevTBRx
99=Unknown)	
<u>'</u>	
History of loss to follow-up or non-compliance during TB	PrevTBfu
treatment (0= No, 1= Yes, 99=Unknown)	
Previous positive test for LTBI	HxLTBI
0= No, 1= Pos TST, 2=Pos IGRA, 99=Unknown)	TIXETBI
Previous TST result date (Enter the date of the patient's most	PrevTSTdat
recent TST before any test conducted as part of current	
	е
diagnostic workup leading to patient's current diagnosis of TB.	
MM/DD/YY)	Dray TOT-
Previous TST result (MM) (Enter the mm reading of the	PrevTSTm
patient's previous TST result. 99=Unknown)	m
Previous TST interpretation (1=Negative, 2=Positive,	PrevTSTrslt
3=Unknown)	
Previous QFT result date (Enter the date of the patient's most	PrevQFTdat
recent QFT before any a test conducted as part of current	е
diagnostic workup leading to patient's current diagnosis of TB.	
MM/DD/YY)	
Previous QFT result (Enter value [result-nil]. 99= unknown)	PrevQFTnu
, , ,	m
Previous QFT interpretation (1=Negative, 2=Convertor,	PrevQFTrslt
3=Unknown)	
Diagnostic T.Spot result (Enter the patient's qualitative result, if	PrevTSpot
completed as part of the diagnostic workup leading to the	
patient's current diagnosis of TB. 1=negative, 2=positive,	
3=indeterminate, 4=not done, 99=unknown)	
Diagnostic T.Spot value (Enter the quantitative result of the	PrevTSpotv
patient's current result, 99=Unknown. Leave blank if not	alue
performed.)	alue
Diagnostic T.Spot date (MM/DD/YY)	PrevTSpotd
Diagnostic 1.5pot date (WilVI/DD/11)	ate
Previous chest radiograph (CXR) result date (Enter the date	DateprevCX
<u> </u>	R
of the patient's most recent CXR <i>before</i> any CXR conducted as	K
part of current diagnostic workup leading to patient's current	
diagnosis of TB. MM/DD/YY)	D OVD
Previous CXR result (1=Negative, 2=Abnormal, possibly TB,	PrevCXRrsl
3=Abnormal, not consistent with TB, 99=Unknown [not	t
completed or not available])	ļ
Initiated treatment for LTBI?	LTBIRxStart
0=offered but refused, 1=never offered by provider, 2=yes,	
initiated, 99=unknown	
Prior LTBI treatment completed 0= No, 1= Yes,	HxLTBIRx
99=Unknown	

### Infectious Period Determination

Question	Code	Variable
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Date of infectious period beginning (format MM/DD/YY) -For symptomatic patients, start the infectious period 3 months before "Date of symptom onset" recorded on page 2For asymptomatic patients who have sputum smear-positive or cavitary disease, start the infectious period 3 months before the "Diagnosis date" recorded on page 2For asymptomatic patients without sputum smear-positive or cavitary disease, start the infectious period 1 month before the "Diagnosis date" recorded on page 2	IPopen
<ul> <li>Date of infectious period end (format MM/DD/YY)</li> <li>For patients who are not isolated, the infectious period can be closed when the following three conditions are met: <ol> <li>Treatment with an adequate regimen (based on drug susceptibility results) for ≥2 weeks, AND</li> <li>The patient shows clinical improvement, AND</li> <li>Three consecutive sputum smears are negative (which have been obtained at least 8 hours apart)</li> </ol> </li> <li>For patients who are isolated (e.g. in a hospital) until these three conditions are met, then use date of isolation as the end of the infectious period.</li> </ul>	IPend

#### **Case Interview Form**

Question	Response	Variable Name
Case Last Name		Lname
Case First Name		Fname
Alternate Names/Nicknames/Aliases:		Alias
Age		Age
Date of Birth		DOB
Check the database for the patient's estin	mated infectious period	'. -
Start of infectious period:		
End of infectious period:		
Explain to the patient that you have been why there have been more cases of tuber series of questions to try to identify where people who have TB, as well as to figure Acknowledge that the patient has already providers. Reassure the patient that all ar of the interview is to learn information that people from getting sick (emphasize prote time and for speaking with us.	rculosis, or TB. Explain the health department out where the patient no participated in many in Inswers will be kept con to to the spreas	n that you will be asking a t might be able to find other might have gotten sick. Interviews with health care fidential, and that the purpose ead of TB and prevent other
Note that throughout the interview, the 2 years before the start of the infectious	is period to the end o	f the infectious period.
Ask patient whether they are from the If not, ask where patient came from an		to the area.

Public reporting burden of this collection of information is estimated to average 2 hours 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; ATTN: PRA (0920-1011)

Discuss symptom onset date. Confirm based on chart data.
We are interested in learning where you could have been exposed to TB in the 2 years
perfore you got sick with TB. People sick with TB often have a bad cough, or might lose
lot of weight. TB is spread through the air when a person who is sick coughs or speaks and does anything that brings up air from the lungs. How do you think that you
not TB?" Mention household exposure (i.e. people you visited or people who visited you).
Attempt to elicit names of sick contacts who might have been source patients. Note when and
where the exposure occurred. Emphasize that these people are not in trouble, and we are not
rying to blame anyone. We are trying to make sure we can find all sick people and treat them.
TD is commonly appead among poople staying in the same boundhald. Wa're were ad-
TB is commonly spread among people staying in the same household. We're worried

"TB is commonly spread among people staying in the same household. We're worried about people who may have been staying with you or people you may have stayed with when you were coughing a lot or started feeling sick. I know it might be hard to remember, but please try your best. During [infectious period], where did you live, and who was staying with you?" *Emphasize protecting family.* 

Time period(s)	Last time visited	Location	People in household	

T		

"TB can also be spread to people you spend a lot of time around, even if you don't stay in the same household. During [infectious period], could you tell us where you worked, where you hung out, and who else was usually there?" Emphasize protecting friends and family. Mention work sites, bars, friends' homes.

Location	Dates of first attendance	Dates of most recent attendance	Frequency of attendance	Contacts present

Ask patient how else he/she passes time. As examples, you could mention cards, bingo, video lottery. Record locations and contacts present.

Activity	Location	Dates of first attendance	Dates of most recent attendance	Contacts present

of the reservati		or whether fri	ends/family fro	tends social events ON a om on a reservation visite d contacts.	_
reservations in		ether friends/fa	amily from off	tends social events OFF the reservations visited to ontacts.	
infection, and r	ent that certain a make a person n following TB ris	nore likely to b	ecome sick.	able to fight off a TB	
<b>0</b> =None <b>1</b> =Less than Da	nercial tobacco	during the yea	r before diagn	osis?	
2=Daily 3=Does not reca	all or refuses				
Smoking traditi 0=None 1=Less than Da	ional tobacco di	uring the year	before diagnos	sis?	
<b>2</b> =Daily <b>3</b> =Does not reca	•				

If so: What substance:
Participates in "sweats" (traditional sweat lodge purification ceremony): Y N
Location:
Alcohol use ("drinking") within 1 year before diagnosis?  0=Never  1=Rarely (1-2 times ever)  2=Occasionally (more than 1 or 2 times, but less than most days or nights)  3=Frequently (most days or nights of the week)  4=Does not recall or refuses
Note the locations where patient drank alcohol? Smoked?
With whom would the patient usually drink? Smoke?
Among the group that the patient drank with/smoked with, did anyone possibly have TB?

Non-injection drug ("taking anything for recreation, e.g. marijuana") use within 1 year before diagnosis 0=Never 1=Rarely (1-2 times ever) 2=Occasionally (more than 1 or 2 times, but less than most days or nights) 3=Frequently (most days or nights of the week) 4=Does not recall or refuses					
What kinds of drugs were used before diagnosis? Circle all that apply.					
Marijuana Crack or cocaine Methamphetamine Heroin Prescription drugs					
Other drugs:					
lote the locations where non-injection drugs were used:					
Orug use with anyone with possible TB?					
njection drug use ("shooting up") within 1 year before diagnosis					
<ul> <li>0=Never</li> <li>1=Rarely (1-2 times ever)</li> <li>2=Occasionally (more than 1 or 2 times, but less than most days or nights)</li> <li>3=Frequently (most days or nights of the week)</li> <li>4=Does not recall or refuses</li> </ul>					
What kinds of drugs were used before diagnosis?					
Note the locations where injection drugs were used or obtained:					
Drug use with anyone with possible TB?					
ANY drug use prior to the year before diagnosis					
What kinds of drugs were used?  Note the locations where drugs were used or obtained:					
Drug use with anyone with possible TB?					

(i.e., where people were coughing a lot) or people we should contact?				

Any other contacts not yet discussed:

Name of Contact (and contact info if available)	Where and when had contact	How often had contact? (1=daily, 2=few times/week, 3=weekly or less, 99=unk)	Activities Together	Smoked together? (0=no, 1=yes, 99=unk)	Drank together? (0=no, 1=yes, 99=unk)	Drugs together? (0=no, 1=yes, 99=unk)	Comments

**Tuberculosis Contact Screening Form** q Male DOB: Contact Name: Age: a Female **Current Location:** Contact Exposure History (During the Infectious Period) **Date of Last Exposure:** Contact's Relationship to Index: Location of Exposure: 1. How much time did you spend in the same room or house as Number of days per week: the index while he/she was contagious (during the infectious Number of hours per day: 2. How much time did you spend in a bar or drug-using location Number of days per week: as the index while he/she was contagious (during the infectious Number of hours per day: 3. How much time did you spend in the same room in the hospital Number of days per week: while he/she was contagious (during the infectious period)? Number of hours per day: 4. If you are a healthcare worker, did you perform any procedures q Yes (If Yes, person is automatically a on the index patient that may have caused them to cough (such close contact) as suctioning, collecting sputum, performing CPR, using a bag mask, or intubation) q No IF YES, specify type of procedure(s) and date(s) 5. Specify other contact setting and any related details Ø Based upon the answers above, is this a "close" contact? A "close" contact is a person who spent ≥4 hours multiple times *or* spent ≥8 hours qYes qNo at least one time inside the same room as the index patient (during the infectious period)? TB Symptom Screening (Current Symptoms) **Start Date and Duration** Instructions: Screen to see if the contact currently has TB symptoms. Consider the contact "symptomatic for TB" if they have: (1) A cough for ≥2 weeks duration OR (2) Two "yes" responses to symptoms #2-8 that cannot be explained by another medical condition

Public reporting burden of this collection of information is estimated to average 15 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; ATTN: PRA (0920-1011)

qNo

qNo

qYes

qYes

1. Have you been coughing for ≥2 weeks?

2. Have you been coughing up blood?

3. Have you had difficulty breathing?	qYes qNo				
4. Have you had fevers or chills?	qYes qNo				
5. Have you had night sweats? (completely soaking your clothes at night)	qYes qNo				
Have you been tired or feeling weak lately?	qYes qNo				
7. Have you lost your appetite?	qYes qNo				
8. Have you had unplanned weight loss?	qYes qNo qUnknown	yes, how much?			
$\varnothing$ Is this contact symptomatic for TB?	qYes qNo  If yes, specify symptom star	t date:/			
TB Risk Factor Screening		Notes			
Instructions: Screen to see if the contact has risk factors	s that could increase their risk for progre	ssion to active TB disease.			
1. Is this contact >50 years old?	qYes qNo				
Was this contact <5 years old during the exposure period?	2. Was this contact <5 years old during the exposure period?  qYes qNo				
3. Do you have diabetes?	qYes qNo or Unknown				
4. Do you have HIV?	qYes qNo or Unknown				
5. Do you have cancer?	qYes qNo or Unknown				
6. Do you take prednisone every day?	qYes qNo				
7. Do you smoke tobacco?	qYes qNo				
3. Do you drink alcohol? qYes qNo		If yes, specify amount/frequency			
9. Do you use any other substances?	qYes qNo	If yes, include types/routes, frequency, and locations where substances acquired and used			
Ø Does this contact have a high-risk cond If the contact answers "yes" to questions 1-6 above then the contact has a high-risk condition.	qYes qNo				
Additional Questions					
Have you ever been diagnosed with active TB disease?     If so, please provide details including treatment if any.					
Have you ever been diagnosed with latent TB infection?     If so, please provide details including treatment if any.					
3. Have you ever known anybody with TB?					
If yes, what was/is the nature of your relationship and contact? What did/does this person do during the day? How did/does he/she spend his/her time? Who spent/spends a lot of time with that person?					

<ul> <li>4. Do you know anybody now who might have TB symptoms?</li> <li>(e.g., cough ≥2 weeks, fevers, chills, unintended weight loss)</li> </ul>					
			E	ND QUEST	IONS
W		Date TST Placed	Date TST Read MM		Chest X-Ray
Test Results	TST 1: TST 2: TST Inter q Negation		If pos, Convers	ion? q	CXR Date://  CXR Result: q Not Suggestive of TB q Suggestive of TB
lts	IGRA 1:	Date of IGRA	IGRA Result		Chest X-Ray
Test Results	IGRA 2:	erpretation: ve q Positive	If pos, Conversion? q		CXR Date:/  CXR Result: q Not Suggestive of TB q Suggestive of TB
q Adverse q Lost to fe			Treatment Outcome	ted LTBI treatment q Provider decision to stop q Moved q Died d treatment q Other (specify):	
TB Status  q LTBI q TB Disease q Not infected (test negative 8 weeks after last exposure) q Lost to follow-up					
Interviewer Name: Date:/					