

#### **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:	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	
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
1 Ctal 1033	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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Illness Information								
Illness on	set date:	/ 🗆 н	ospitalized					
Fever	□ Yes □ I	□ Yes □ 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					
Specimer	Specimen 3 collected:/ Type: □ Serum □ CSF □ Amniotic fluid □ Tissue □ Saliva □ Urine □ Semen							
Specimer	n 4 collected:	/	Type: ☐ Serum ☐ CSF ☐ Amniotic fluid ☐ Tissue ☐ Saliva ☐ Urine ☐ Semen					
Specimer	Specimen 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 medições:							
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:		$\neg$			
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 HISTÓRICO PRÉ-NATAL							
Ultrassonografia pré-na	atal:	Resultado	▼ Se for anor	mal, data:			
Se anormal, especifique:							
Amostragem vilo cori	al: ▼	Resultado:	•				
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	Oligodramnio			Polidramnio	
Insuficiência o	olo uterino	Hiperen	nese gravidica	Anomalias anatômicas no útero Descolamento de			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 4:							
	30 dias anteriores da gravidez 1 Tri 2º Tri 3º Tri							
Medicamento 5:	Medicamento 5: ———————————————————————————————————							
L			30 dias ant	eriores da g	gravidez 🔲 1	o Tri	2º Tri 3º Tri	
–Exames de doen	ças infecciosas p	ara o bebê	ė: ————					
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	iosas durante a gravidez —					
VDRL	_	Resultado	▼				
CMV	<b>—</b>	IgM	<b>*</b>	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	<b>▼</b>	IgM	<b>~</b>	IgG	•	PCR	▼
Outros1	<b>—</b>	Resultados Outros1:					
Outros 2	<b>-</b>	Resultados Outros2:					
O bebê fez exa	ame de vista?	•		Resulta	ido de exame de vi	sta do be	bê: ▼
Caso anormal,	descrever:						
Outras anom	alias/defeitos	s do bebê —————					
Presença de m	alformações no	RN:					
Se sim, espe	ecificar:						
Aparelho o	circulatório 🔲	Aparelho digestivo 🔲 Aparell	ho respirató	rio 🔳	Órgãos genitais	Apare	elho osteomuscular
Outras							
Descreva a ma	Descreva a malformação encontrada:						
2 000. 014 4	o.mayao enee						
Descrever re	sultados de e	xames ou defeitos específic	os de forn	na mai	s detalhada		
	Descrever resultados de exames ou defeitos específicos de forma mais detalhada						

#### Questionário da pesquisa - Microcefalia

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Equipe:								- :   <b>-</b> : <b>-</b> :		
	□ 1							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 gravid	ez					
1. Data da útima mestruação:		2. Qual foi	a data prováve	el do parto inforn	nada pelo médico?	
	■ Nã	o sei				Não sei
3. Tipo de gravidez	▼ Se gemela	ar, número de beb	ês:	4. Os outros be	ebês nasceram vivos?	•
Se sim, teve alguma malformação	: 🔻	Especificar				
5. Tipo de parto:		▼				
6. Quantas vezes você engravidou a bebê nascido morto ou outros resul		avidez, incluindo g	estações que p	odem ter termin	ado em abortos natur	ais,
Número de gest	tações:	N	úmero de nasci	dos vivos		
Número de	abortos:	Núm	ero de nascidos	s mortos:		
7. Há alguma (outra) criança ou adu	ılto da sua fan	nília que nasceu c	om microcefalia	n? <b>▼</b>		
Se sim, especifique (grau de parer	ntesco em rela	ção ao bebê e o n	nomento do dia	gnóstico):		
8. Você e o paí da criança têm algur	m grau de par	entesco?	▼ Se	e sim, qual?		
9. Qual é a data de nascimento dest	e bebê?					
D. Doenças durante a gravidez  Agora, vou fazer algumas pergunt  1. No período de 30 DIAS ANTES	as sobre doen				os seguintes sintomas	:
Manchas vermelhas no corpo:		Ouando .	Semanas ou r	meses		
Marichas Vermenias no corpo.	•	Quando	<ul><li>Semanas</li></ul>	Meses	Não sei	
Febre:	•	Quando	Semanas ou r		Não sei	
Coceira:	•	Quando	Semanas ou r		Não sei	
Dores articulações:	•	Quando	Semanas ou r		Não sei	
Olhos vermelhos:	•	Quando	Semanas ou r		Não sei	

2. Durante o PRIMEIRO TRIME sintomas?	ESTRE 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
0000	<b>↓</b> Qualita	
Dores articulações:	<b>▼</b> Quando	Semanas ou meses
	· Camara	
Olhos vermelhos:	<b>▼</b> Quando	Semanas ou meses
Ollios verniellos.	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
, 33, 3,	Quanto	
Coceira:	<b>→</b> Quando	Semanas ou meses
Cocena.	<b>→</b> Quando	
Dores articulações:	<b>▼</b> Quando	Semanas ou meses
bores ar cicarações.	Quanto	
Olhos vermelhesi	Ouendo [	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.	▼	dando	
Olhos vermelhos:		uando	Semanas ou meses
omos remenos.	¥ 4		
<ol> <li>Entre o mês anterior à gra afirmativo, registrar a seman</li> <li>Infecção urinária</li> </ol>	videz e o final da gra a de gravidez, se pos	videz, você teve algu sível, e o(s) mês(es)	ima das seguintes doenças ou infecções? [Em caso de gravidez, caso a semana seja desconhecida]
iniceção dimaria			Semanas ou meses
-30 Infecção dos rins, bexiga ou tr	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	→ Quando	Semanas Meses Não sei
3º Tri Infecção dos rins, bexiga ou	ı trato urinário	<b>→</b> Quando	Semanas ou meses
			Semanas Meses Não sei
Infecções por fungos			
20 1-6		Outrada 🗔	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 poi lungos	•	Quando	Semanas
			Semanas ou meses
3º Tri Infecção por fungos	•	Quando	

Citomegalovírus (CMV)  Quando  Semanas ou meses	Meses Não sei  Meses Não sei
Citomenalovírus (CMV) Quando	
Rubéola   ✓ Quando  Semanas ou meses  ⑤ Semanas ⑥ I	Meses Não sei
Herpes   ✓ Quando   Semanas ou meses  ⑤ Semanas	Meses    Não sei
Semanas ou meses  Quando  Semanas ou meses	Meses Não sei
Catapora Quando Quando Semanas ou meses	Meses Não sei
LCMV (coriomeningite linfocitária)	Meses    Não sei
. Entre o mês anterior à gravidez e o final da gravidez, você teve alguma outra infecção que Em caso afirmativo, registrar a semana de gravidez, se possível, e o(s) mês(es) de gravidez,	
Especifique: Quando Semanas ou meses Semanas ou Mes	ses
-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?	
Se marcou alguma das doenças acima (respiratória, neurológica e cardíaca), 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		egistrar 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 se	Frequência: ▼
Fertilizantes 🔻	Especifique o(s) fertilizante(s):	
Quando	Semanas ou meses  Semanas Meses  Não s	Frequência:   ▼
Fumigação 🔻	Especifique o(s) produto(s):	
Quando	Semanas ou meses  Semanas Meses  Não s	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 p	problemas?
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 f	
1. Como você classificaria	sua raça?   ▼ Se	e outra, especifique:
2. Qual era a sua escolario	dade quando o bebê nasceu? (considerar o ma	naior nível completo)
3. Durante os 9 meses de	gravidez, qual era a renda mensal de sua fam	mília?
4. Quantas pessoas eram	sustentadas por essa renda, inclusive adultos	s e 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)
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:) Medical 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?  Yes No (STOP interview)
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 get sick with
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.
InstructionstotheInterviewer-IfthecaseisstillonlyPossibleandnotConfirmed,besuretostate:
"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 (data), from (back loss tien). We still peed to sheek the DNA fingerprint of the sample we received and see if it matches the
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."
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 Record Review)
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):

Dem	nographic Infor	mation (Medical Record Review and F	Patient Interview)	
l will	start by asking	ome questions about your background ar	nd where you live.	
	In the time period	No Unk		on) did you stay at least one night in an institutional setting?  Assisted living facility Acute care facility Other, specify:
6. 7.	☐ Black	:	☐ Unk Not Hispanic or Latino ☐ Unk	
Ноз	Ithcara Evnosu	re (Patient Interview)		
		,	at occurred in the 30	days before you became ill, the period from (first date of exposure
		Iture collection). I will refer to this period		
9.	Unk	out each of these home health visits, start		ing wound checks, dressing changes, baths)? Yes No ent.
	Date	Agency Name and Contact Info	ormation	Reason for Visit
10.	dialysis)? Exam	oles of healthcare providers are primary can		with healthcare providers (this does not include outpatient visits for such as cardiologists or oncologists, eye doctors, and dentists.
	a. If yes,	Clinic Name (phone number and Specialty T		Reason for Visit
11.	<ul><li>a. If yes,</li><li>b. Name</li></ul>	ore you became ill, were you receiving out what type of dialysis:   Hemodialysis and contact information for dialysis facility	Peritoneal Dialysis	Unk
		odialysis please specify access type: fist		
		lays do you receive dialysis 🏻 MWF 🔲 T alysis session before symptoms onset. Date		

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

Date	Hospital Na	ime and Contact Information		Reason for	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  h before you beca	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 or according to the stay of	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 receive	ed care from ar	ny of the follow	ina in the mo	nth prior to illnes	s onset?

Exposure	Yes	No	Unk	Location	Were any procedures beyond a routine examination performed?	Date(s) (MM/DD/YYYY)
					If yes, describe.	
Dentist						
Podiatrist						
Chiropractor						
Massage Therapist						
Naturopath						
In the month before a. Inhalers			did you	use any of the following medications, check	call that apply	

		6.1. 6.11					
the month before you became ill o	-	•	wing r	nedications, check	all that ap	ply	
a. Inhalers Yes		<del></del>					
b. Nebulizers Yes		_					
c. Nasal sprays 🗌 Yes 🗌	No [	Unk					
d. Eye drops  Yes	No [	Unk					
e. Oxygen 🗌 Yes 🗀	]No [	Unk					
f. Over the counter suppleme	ents, inc	luding vitamins, pro	biotic	s, powders added t	to a drink	or s	moothie (e.g., protein powder)
☐ Yes	☐ No	Unk					
		If yes, specify type	e and	brand		_	
g. Thickened juice, food, or st	nakes?	∏Yes  No □	Unk	·		_	
i. If yes, which brar	nd of th	ickener? Thic	ck-It®	ReadyCare 2.0 <sup>™</sup>	Horm	nel T	Thick & Easy®   Simply Thick   Othe
specify:		Unknown	l	•	ı		, , , ,
ii. If yes, specify typ	e of thi	ckener:					Unknown
h. Proton pump inhibitors (PF							∐No ∐Unk
i. H2 blockers. Examples of H			agam	et. 🔲 Yes 🔲 N	o 🔲 Unl	K	
j. Antibiotics Yes No							
i. If yes, check all th	nose 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
Cefotaxime		Daptomycin		Nitrofurantoin			Other (specify):
Cefpodoxime		Doripenem		Ofloxacin			Other (specify):

17.

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):

. 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):

	Home 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				
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, what is a second of the month before	nich brand? re you became ill, did	Un I you use any topic	ık al products (e.g., lotior	s, creams, liniments, or ointm	ents)? Yes No Unk
	-				Unk
	•		-		
toothpaste, deodo	rant)? Check all that	apply and specify	location:		ap, shampoo, creams and lotions,
☐ Drug Store	Grocery Sto	ore $\square$	On-line Ot	her	
Name			Location		
In the month befo ☐ Yes ☐	ore you became ill, did No	d you use marijuan nswer 🔲 Unk	a, also called cannabis	in any form?	
Do you know of any	one else in your hom	e or community th	nat has experienced a s	milar illness? Yes No	Unk
		-	ou became III? Yes	∐ NO ∐ UNK	
In the month before	re you became ill, ha	ve any pets in your	home had insect infes	tations (i.e., fleas)? Yes	☐ No ☐ Unk
	•		0 .	your home? Yes No	
Name	Phone Number		·	Place of Employment	Check if surveillance
				1, 1, 1	cultures were obtained
	a. If yes, will in the month before a. If yes: Plant in the month before a. If yes, specific in the month before a. If yes, will were you been bitted a. If so whill in the month before in the month	a. If yes, which brand? In the month before you became ill, did a. If yes: Please tell me all of the a. If yes: Please tell me all of the a. If yes: Please tell me all of the a. If yes: Please tell me all of the a. If yes: Please tell me all of the a. If yes: Please tell me all of the a. If yes: Please tell me all of the b.  Where did you purchase the personal c toothpaste, deodorant)? Check all that b.  Drug Store Grocery Store  Name  Now I would like to ask about a topic the lin the month before you became ill, did a. If yes, specify route (check all the b.)  Do you know of anyone else in your hom a. If yes, who:  Were you been bitten by any insect in the a. If so which?  In the month before you became ill, did If yes: Please tell me more about all the ling the ling the line in the month before you became ill, did ling yes: Please tell me more about all the ling the line in the month before you became ill, did ling yes: Please tell me more about all the line in the month before you became ill, did ling yes: Please tell me more about all the line in	a. If yes, which brand? Ur In the month before you became ill, did you use any topic a. If yes: Please tell me all of the products you use	In the month before you became ill, did you use any topical products (e.g., lotion a. If yes: Please tell me all of the products you used during this period	a. If yes, which brand?

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

	Name	Phone Number	Relationship	County of Residence	Occupation	Place of Employment
Оп	tside Exposure (Pati	ent Interview)				
	risido Exposar o (r dir					
54.	In the month before	you became ill did you	attend any of the	following check all	that apply: Rel	igious service Support Group [
		Club Social g				
55.	If yes to any of the al	pove, please specify w	hen and where _			
Mot	tatalatarviawar:lfvauw	villboconductingthofull	foodovnosuroguos	etionnairowiththisn	ationtthonskinguost	tions56-58andproceedtoquestion
59.		<u>mibeconductingmenui</u>	<u>noodexposureques</u>	<u>stiorii aii ewiti iti ii sp</u>	atterittileriskipquesi	nonsso-soandproceedtoquestion
		u some questions abo	ut food you have	eaten which will fo	cus only on the 7 da	ays before you became ill, which is the period
	m (7 days prior to date	•	•		, <u>—</u>	<del></del>
56.	In the 7 days before			e from that was pr	epared at home	
		ry store Yes			2	
	h Farmo	i. If yes, specify 1 ers market/food purch	Z asod directly from	farm D V	3 es	
		i. If yes, specify 1			3	
		n food store  Yes			o. <u></u>	
		i. If yes, specify 1			3	
		specialty market				
		i. If yes, specify 1			3	
		r meat shop (e.g., buto i. If yes, specify 1			2	
		ng or fishing Yes			3	
		i. If yes, specify 1			3	
		y grown fresh foods, e				
	_	i. If yes, specify 1			3	
		ole (my food is prepare				
57.			-		tside the facility? [	Yes No Unk
- 0		scribe:			ab as from a rostour	rant, meal delivery service, or at a school, or
00.	,	es foods that you ate c				•
	•	urants (including delive		· ·		CS NO DIIK
		i. If yes, specify 1	•		3	
		, , <u> </u>				<del></del>
		delivery service, such a				
		i. If yes, specify 1	2		3	
	- L- 00	ution or oh as besself of	or cohool 🗆 V	□No □U		
		ition, such as hospital		□ INO □ UNK	3.	
		i. If yes, specify 1			4	

#### 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):

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

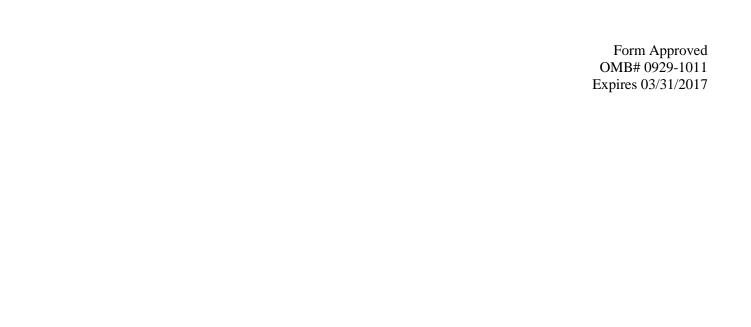
This is the end of the interview. Thank you very much for your time and willingness to provide this valuable information.

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 CDC	CID:	
Wisconsin	Clinical Course Abstraction Form	
SECTION A		
includes the collected a culture). If	nstructions: Abstract Section A using all medical records e medical record from the outpatient encounter, ER, or nd follow-up care (hospitalization for EK, care at AL for a patient was transferred during their EK hospitalization from both facilities.	hospitalization where the positive culture was EK, or outpatient treatment for the positive
Obtain list	of ICU locations from hospital prior to abstraction to en	nsure that all ICU stays are captured in item 21.
Patient CD	CID:	
Abstractor	Name:	
Date of Ab	straction:	
Specify nar	me of facilities included in abstraction for Section A:	
	e:	
	x: Male Female Unk	
	ce (check all that apply):	n Indian/Alaska Native 🗌 Black
4. Eth	nnicity: Hispanic or Latino Not Hispanic or Latino	] Unk
5. Co	unty of residence:	
6. Da	te of collection of first specimen positive for E. anophel	is:/(MM/DD/YYYY)
	me of collection of first specimen positive for E. anophel nere 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						
	M/DD/YYYY) (If patient was hospitalized and						
	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)?						
	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 kidney disease (on dialysis) Chronic liver disease without cirrhosis Cirrhosis Chronic obstructive pulmonary disease (COPD)/emphysema Chronic lung disease (other than COPD/emphysema, asthma)	Failure to thrive
asthma)  Chronic steroid or other immunosuppressive therapy  Chronic ventilation/tracheostomy  Congenital heart disease  Congestive heart failure  Connective tissue disease  Cystic fibrosis  Dementia  Diabetes mellitus with complications  Diabetes mellitus without complications  Eczema	Recurrent cystitis or urinary tract infection  Sickle cell disease  Solid tumor malignancy, metastatic  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

I. Did the patient have any medication a	llergies?: □Yes □No □Unk	
,	norgioca i El 100 El 110 El olim	
If yes, specify all:	<u></u>	
. Specify all symptoms reported by pat onset date reported by the patient (if the symptom was present; otherwise s	known). Check No only if the records	
Symptom	Symptom Present?	If Yes, Date of Symptom Onset (MM/DD/YYYY) Write UNK if unknown
Weakness	Yes No Unk	
Headache	Yes No Unk	
Lightheadedness, Dizziness	Yes No Unk	
Blurry vision	Yes No Unk	
Documented Fever (T >100.3)	Yes No Unk	
Subjective fever	Yes No Unk	
Night sweats	Yes No Unk	
Chills	Yes No Unk	
Cough	Yes No Unk	
Wheezing	Yes No Unk	
Sore throat / difficulty swallowing	Yes No Unk	
Muscle aches	Yes No Unk	
Chest pain	Yes No Unk	
Shortness of breath	☐ Yes ☐ No ☐ Unk	
Vomiting	Yes No Unk	
Nausea	Yes No Unk	
Diarrhea	Yes No Unk	
Abdominal pain	Yes No Unk	
Joint pain	☐ Yes ☐ No ☐ Unk	
Bleeding gums	☐ Yes ☐ No ☐ Unk	
Mouth sores	☐ Yes ☐ No ☐ Unk	
Skin wound	Yes No Unk	
Rashes	Yes No Unk	
Skin warmth	☐ Yes ☐ No ☐ Unk	
Skin redness	Yes No Unk	
Skin pain	Yes No Unk	
Altered mental status	Yes No Unk	
Other specify	Yes No Unk	
Other specify	Yes No Unk	

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:						
_						
20. Record la	aboratories documen	ted as specified below.				
Parameter (include units)	Results on day patient was admitted to hospital (if	Results on day first positive specimen was collected (select results	Results of highest value obtained during	Results of lowest value obtained during	Results on day discharged from hospital	
	multiple results, select first collected)  Date: _/_/	closest to specimen collection)  Date: _/_/	hospitalizatio n Specify date for each value	hospitalization Specify date for each value	or died  Date: _/_/	
WBC						
Percent neutrophils (corresponds to WBC count above)						
Percent bands (corresponds to WBC count above)						
Platelets						
Hematocrit BUN						
Creatinine						
Lactate						
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	/	
2	//	Unk	/	
3	//	Unk	/	
4	//	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 pati	ient require vasopressors?	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 ntry.				
Examples: (specify)	Dopamine, Dobutamine, Epinepl	nrine, Norepinephrir	ne, Neosyneph	rine, Vasopressir	n, Other
Vasopressor	Start da	ate	Stop date	<del></del>	
•	/	/		/	
	<u> </u>	<u> </u>		<u> </u>	
		<u></u>	<u> </u>	<u>-'</u>	
		<u></u>			
	<u> </u>	<u></u>	<del></del>		
			/	_/	
		/	/	_/	
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 Topical Inhaled	_//_	Unk	/	/ Unl
	☐ IV ☐ IM ☐ PO☐ Topical ☐ Inhaled☐ IV ☐ IM ☐ PO☐ INHALED☐ INHALED		Unk	/	/ Unl
	Topical Inhaled		Unk		/_ Unl
	Topical Inhaled	/ /	Unk	/	/ Unl
	Topical Inhaled	/ /	Unk		/ 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						
•	oted (check all that apply; o ument their full written impi	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:		
30. Did patient have th	oracentesis?  Yes  No	o 🗌 Unk
If yes, record results for ear	ch thoracentesis performed (	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 for ear	int aspiration? Yes Ch joint aspiration performed	_
Date:/	Unk	
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
				∐  Orgar	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					
	ockers listed as outpatient medications \( \subseteq \mathbb{No} \subseteq \mathbb{Unk} \)	on in the HPI from the time o	of first positive culture			
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		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	ine list patient ID (CDCID) Wisconsin State Laboratory of Hygiene ID Abstr						
This form is in	ntended to be us	ed for abstraction of medi	cal records for patien	ts in Wisconsin with:			
	AND	Elizabethkingia spp cultur on or after November 1, 20					
Patient NAM	E:						
Patient DOB:	//	MM/DD/YYYY)					
Abstraction I	Information						
Date medical	l record abstracti	on completed://_	(MM/DD/YYYY)				
Abstractor:		e 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   Emergency Room   Observational unit/Clinical Decision Unit   Outpatient   Assisted Living	☐ LTCF/SNF ☐ LTACH ☐ Dialysis clinic ☐ Unknown						
<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 the collected in th</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?	
	he <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)			•	•	Ab	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 imm	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.	Spec	imen	Collect date (mm/dd/yy) Time (HH:MM)	Pathogens identified
1	□Blood □Urine □BAL □CSF	☐ Pleural fluid☐ Synovial fluid☐ Wound☐ Stool	/	Path1Path2
	Ascites Blood Urine	Sputum Pleural fluid Synovial fluid	//	Path3CX is Neg Path1
2	☐BAL ☐CSF ☐Ascites ☐Blood	☐ Wound ☐ Stool ☐ Sputum ☐ Pleural fluid	:	Path2 Path3
3	Urine BAL CSF Ascites	Synovial fluid Synovial fluid Wound Stool Sputum	//	Path1Path2Path3
4	Blood Urine BAL CSF	Pleural fluid Synovial fluid Wound Stool	//	CX is Neg Path1 Path2 Path3
5	☐Ascites ☐Blood ☐Urine ☐BAL ☐CSF	Sputum Pleural fluid Synovial fluid Wound Stool	//	CX is Neg Path1 Path2
6	Ascites Blood Urine BAL CSF Ascites	Sputum Pleural fluid Synovial fluid Wound Stool Sputum	//	Path3CX is Neg Path1 Path2 Path3
7	Blood Urine BAL CSF Ascites	Pleural fluid Synovial fluid Wound Stool Sputum	//	Path1Path2Path3
8	Blood Urine BAL CSF Ascites	Pleural fluid Synovial fluid Wound Stool Sputum	//	Path1Path2Path3
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)	)

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       □Pleural fluid         □Urine       □Synovial fluid         □BAL       □Wound         □CSF       □Stool	//	
	Ascites Sputum	:	
2	□ Blood       □ Pleural fluid         □ Urine       □ Synovial fluid         □ BAL       □ Wound	//	
	☐CSF ☐Stool ☐Sputum	<u>:</u>	
3	□Blood       □Pleural fluid         □Urine       □Synovial fluid         □BAL       □Wound         □CSF       □Stool         □Ascites       □Sputum	//	
4	□Blood       □Pleural fluid         □Urine       □Synovial fluid         □BAL       □Wound         □CSF       □Stool         □Ascites       □Sputum	//	
5	□Blood       □Pleural fluid         □Urine       □Synovial fluid         □BAL       □Wound         □CSF       □Stool         □Ascites       □Sputum	//	

Line list patient ID (CDCID) Wisconsin State Laboratory of Hygiene ID Abstractor Initials					
Section III: Visit with the positive culture.  Location of visit: acute care hospital Drach Skilled Nursing Facility Urgent Care Date of symptom onset (for positive culture):	e Other	<del>-</del>			
25. Chief Complaint (i.e. what were the patient	t's symptoms) at time of positive culture:				
	Approximate Start Date (MM/DD/)	(YY)			
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					
Other2					
i. If yes, what were the sym	atient develop signs of infection later? Yenptom(s)(MM/DD/Y	?			
26. Patient Labs & Vitals within 2 hours of cultu	ure collection				
Value	Date:MMDDYY	Time::			
Temperature					
Heart Rate					
Blood Pressure					
Respiratory Rate					
Pulse Ox Lactate					
WBC					
WBO					
27. Diagnosis for the visit when the positive cu	lture was collected:				
Acute Respiratory Failure	Pleural Effusion				
Atrial Fibrillation	Pneumothorax				
Bacteremia	Renal Failure				
Cellulitis	Sepsis				
DKA	Stroke				
Heart Failure	UTI				
<ul><li>☐ Hyperglycemia</li><li>☐ Myocardial Infarction</li></ul>	☐Other1: ☐Other2:				
28. Was Patient Admitted in response to th	e positive culture? Yes No				
·	· — —				
29. Did the patient have evidence of soft tis a. If yes, describe:	ssue infection at time of positive cult	ure: Yes or INO			
a. ii ves. describe:					

Line list patient ID (CDCID)	actor Initials						
30. Did the patient have diagnosed pulmonary infection at time of positive 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 positive culture was coll	ected?						
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 ItCF/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	-	
Diama		
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*  Proteic direction Total Aphresis Plateles*  Pooled*  Pooled*  Plasma Aphresis Plateles  Aphresis Plateles Triple collection Total Aphresis Plateles* Total Aphresis Plateles  Whole-blood-derived Individual* Pooled*  Proteid FP24 PF24 PF24 PF24 PF24 PF24 PF24 PF24	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 MpD Plasma' Cryoprecipitate Individual' Pooled Single-colored Individual' Total MpD Plasma' Cryoprecipitate Individual' Total MpD Plasma' Cryoprecipitate Individual' Pooled Single-colored Individual' Pooled Single-colored Individual' Total MpD 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  Whole-blood-derived Individual* Pooted*  Plasma Apheresis FFP 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  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* I	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 PP24  PF24  PF24  Cryoprecipitate Individual FOR POOLED FOR PP24  Cryoprecipitate Individual FOR POOLED FOR PP35  FFP FFP FFP FFP FFP FFP FFP FFP FFP F	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="mailto:EmployeeSurvey"><u>EmployeeSurvey</u></a>

Form Approved; OMB No. 0920-1011 Exp. Date 03/31/2017 □ **Check box if documented case** 

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

A. Employee Background	1. Name:			2. Age:				
3. Sex: ÿ Male ÿ Female	4. Employed at F	Facility since:	/					
5. List occupation: ÿ Activity aid	ÿ Administrative	e ÿ CNA	ÿ Die	etary	ÿ Food service			
ÿ Housekeeping	ÿ Laundry	ÿ PT/OT	ÿ Pha	armacist	ÿ Physician			
ÿ Maintenance	ÿ RNA	ÿ RN/LP	ÿ RN/LPN ÿ Social service ÿ Van driv					
ÿ Wound care team	ÿ Other_							
6. Since <u>Thanksgiving to present</u> , have you	worked in any oth	her patient-care fac	cility?	ÿ Yes	ÿ No (If no, skip to Section B)			
Name & city of facility Dates of empl	-	ve you been in con ent infected with g		What was the	e patient's diagnosis?			
Start:	ÿΥϵ	es		ÿ Strep thro	oat ÿ Impetigo			
/ End:	/ ÿ No			ÿ Cellulitis	ÿ Bacteremia/Sepsis			
Enu.	/   If ye	es, date of contact: _//	: 	ÿ Other, specify:				
Start:	ÿΥϵ	es		ÿ Strep thro	oat ÿ Impetigo			
/	/ ÿ No			ÿ Cellulitis	ÿ Bacteremia/Sepsis			
End:/	/ If ye	es, date of contact:	: —	ÿ Other, sp	ecify:			
Start:	ў Үе	es		ÿ Strep thro	oat ÿ Impetigo			
/ End:	/ ÿ No			ÿ Cellulitis	ÿ Bacteremia/Sepsis			
/	/ If ye	es, date of contact: _//	: 	ÿ Other, sp	ecify:			
7. a. Since the outbreak, have you ha	ad a screening cu		A Streptococci	ıs? ÿ Yes	ÿ No (If no, skip to #8)			
b. If yes, when?//		5 D 4 1 - 5 3	, , 1 <u>0</u> 01	• /1	ë oa			
c. Where was the culture obtained from d. What were the results? ÿ Positiv	•	-	Vaginal ÿ Sk	cin/wound	ÿ Other			
B. Job Description at		ur job, do you hav	e physical conta	ect with nationt	s? ÿ Yes ÿ No			
Facility A	o. As part of you	ui joo, uo you nav	e physical coma	et with patient	(If no, skip to Section D)			
9. Areas usually worked: ÿ Patient room	9. Areas usually worked: ÿ Patient rooms ÿ Nurses' station ÿ Cafeteria ÿ Rehab floor ÿ 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 usually work (circle	ALL that apply):							
Sunday Monday Tu	esday W	Wednesday	Thursday	Friday	Saturday			

13.	What	t kind of patient conta	act do you have? (check ALL th	hat appl	y)								
ÿ Give oral medications ÿ Feeding resident ÿ 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							dding	ÿ Hand	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 1	2 2	3	4	5	N/A
			ou perform hand hygiene after					_			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. When? / /	ction? ÿ	Yes ÿ N	lo (If no,		2))
	c. Did you miss work for this illness?	How m	any days d	iss?		
	e. Did you receive antibiotics for this condition?	o If yes, antibiotic name				
22. If	you're feeling sick before a work shift, how do you notify Warren Barr Gold Coa	ıst?				
23.	a. How many people are in your household? (If none, END)					
	<ul><li>b. How many children under 18 years of age are in your household?</li><li>c. Since Thanksgiving, did anyone in your household have a sore throat?</li></ul>	-	ÿ Yes	ÿ No		
	d. When?/ e. Who (relationship)? _		-	-		
	e. Was he/she diagnosed with strep throat?	ÿ Yes	ÿ No			
	g. Were they treated? ÿ Yes ÿ No If so, with what?					
	h. During the past 3 months, did anyone in your household have impetigo or cel	Ilulitis (ski	n infection	s)?	ÿ Yes	ÿ No
	i. When?//					

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GDCメ뭬 C根で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:	

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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

24. Wit	hin 14	days prior to	GAS cultu	re, was the	e resident incontinent	t of: (mark ALL that apply)	
	ÿ Not	Incontinent	ÿ Stool	ÿ Urine	ÿ Urinary catheter	ÿ Colostomy/Ileostomy	ÿ Unknowr
25. Did	the res	sident particip	oate in the	following	within 14 days prior to	o GAS culture (mark ALL th	nat apply):
	a.	ÿ PT/OT			Times in 14	4 day period:	
	b.	ÿ Speech ı	pathology		Times in 14	4 day period:	
	C.	ÿ Podiatry			Times in 14	4 day period:	
	d.	ÿ Other: _			Times in 14	4 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	7. How many new wound con ÿ 0-4 ÿ 5-9 ÿ 10 or ı	* *
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 Mon	thly ÿ Quarterly y Other:	
12. What types of care do you perform		
ÿ Incision and Drainage ÿ Undressing	g/Redressing ÿ Cleaning wo	und ÿ Wound vac care ÿ Other:
13. Have you ever discovered pieces of ÿ Unknown	foam/cotton gauze present in the	he wound from previous dressing changes? ÿ Yes ÿ No

# 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 consult?	what symptoms would prompt you to replace the wound vac before a physician
26. What symptoms for a "worsening w	yound" 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,

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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	I .		1	1	1		1	

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Sample ID	Date//

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

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_				

Participant classification (	(A/B/C/D)	Team (1/2)	Site (1/	2/3/etc)
Form Completed by				
Name:	Position:	District:		
Phone Number/email: _				
Section 1.	Assessment Participa	ant Information		
ID Number:	Family Name:	English Name	Age:	Gender:
Village/Town:	Parish:	Sub-County:		
District:	Nationality:	Marital status:	ngle	

ון טו	mber: Family Name: English Name Age: Gender: [] Male [] Female
Villa	e/Town: Parish: Sub-County:
Dist	t:Nationality: Marital status: Married Single Widowed
GPS	oordinates: LatLong Elevation
Sect	n 2. Epidemiological Risk Factors and Exposures
1. 2. 3. 4. 5. 6. 7. 8. 9. 10. 11. 12. 13. 14. 15. 16. 17. 18. 19.	Contact   Carrier   Carr
21.	Which animals are usually huntedn past one year, have you had contact with dead wild animals Yes No Unk To you use PPE when handling animals Yes No Unkn Not applicable (If NO/Unkn/NA, skip to #24)
23. 24.	f yes, which ones Gloves Gumboots Mask Eye protection Aprons/ovals Others (specify) Have you eaten wild meat in the past one year Yes No Unk (If NO/Unknown, skip to #26)  f Yes, which species
26.	Have you traveled outside your home or village/town in the past one year  Yes  No Unkn (If NO/Unkn, skip to #28) If yes, specify location and date:
28.	old you ever suffer from undiagnosed fever or illness in the past one year  Yes  No Unkn (If NO/Unkn, skip to #31)  f Yes, when? Month Year

Public reporting burden of this collection of information is estimated to average 20 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)

31. Did someone you know in the last one year have unexplained fever or diagnosis? Yes \( \subseteq \text{No} \subseteq \text{Unkn} \)

30. If yes, did you seek medical attention: Yes No



Form  Sample ID Date//  Exp. [
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33. 34. 35. 36. 37. 38. 39. 40.	Have you had <i>el nino</i> (have you had more rainfall than usual) rains in the last one year?  Yes  No Unkn Have you had flooding in this sub-county in past one year?  Yes  No Unkn (If NO/Unkn, skip to #35) If yes, which months do you get flooding
42. 43. 44. 45. 46. 47.	Knowledge & Attitude Questions  Have you heard about Rift Valley Fever Disease?
51. 52.	If no, why
<ul><li>55.</li><li>56.</li><li>57.</li></ul>	If no, why?
59.	How do you think you can protect yourself from acquiring RVF disease?  vaccination avoiding contact with animals traditional medicine avoiding sick people sleeping in a mosquito net Unkn Others (specify)  How do you think RVF disease can best be healed or treated?  Traditional medicine Spiritual healing Modern medicine Herbal medicine Unkn  Others (specify)
	Do you think you are at risk of contracting RVF virus disease?  Yes No Unk  If yes/no, why
	nk you for your Time End of Interview
Sect	ion 3. Specimen Information
Spec	cimen identification number:
Spec	cimen collection date:/ (DD/MM/YYYY)
Labo	pratory testing date:/ (DD/MM/YYYY)
Resi	ults/Titer level: IaM IaG



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

#### **Livestock Assessment Form**

Sect	Section 1. Herd Demographics	
1)	1) What is your relationship to the livestock?  Owner Herdsman Other(specify)	
2)		
3)	3) Herd location at time of survey? Central collection area Grazing ground Other, specify	
4)		
_,	Poultry Dogs Cats	
5)	5) What is the herd's typical grazing pattern? Shared Enclosed Non-grazing Other (specify)	
6)		
	If yes, why? Nomadic grazing Trade Gift/dowry Other (specify)  If yes, to where?	
	If yes, to where? If yes, how many months ago? <1 month 1-3 months 3-6 months 6-12 months	
Sect	Section 2. Herd Health Status	
7)	· · · · · · · · · · · · · · · · · · ·	
	<ul> <li>Abortions?</li> <li>Yes \( \sum_{No} \) If yes, how many? \( \sum_{No} \)</li> </ul>	
	• Stillbirths?	
	Deaths in adults?	
	Deaths in young?	
0)	<ul> <li>Other health problems?  Yes No If yes, what?</li> <li>In the past year, has your <u>qoats</u> had unusual:</li> </ul>	
8)	Abortions?	
	Stillbirths?	
	Deaths in adults?	
	<ul> <li>Deaths in young?</li> <li>Yes \( \subseteq No \) If yes, how many?</li> </ul>	
	Other health problems? Yes No If yes, what?	
9)	9) In the past year, has your <u>sheep</u> had unusual:	
	<ul> <li>Abortions?</li> <li>Yes \( \sum_{No} \) If yes, how many?</li> </ul>	
	<ul> <li>Stillbirths?</li></ul>	
	Deaths in adults? Yes No If yes, how many?	
	Death in young?	
	Other health problems?	
*Co	*Common RVF symptoms: decreased appetite, decreased milk production, nose/eye discharge, diarrhea, jaundice, prostrati	on, lymph node swelling
	Section 3. Herd Treatment	
10)	10) Hous your acttle resolved.	
10)	10) Have your <u>cattle</u> received:  Vaccines?	
	Insecticide treatment?	
	Other treatments?	
11)	11) Have your <b>goats</b> received:	
,	Vaccines? Yes No If yes, what vaccines?	
	Insecticide treatment? Yes No If yes, when?	
	<ul> <li>Other treatments?</li></ul>	
12)	12) Have your <u>sheep</u> received:	
	<ul> <li>Vaccines?</li> </ul>	
	Insecticide treatment?	
	Other treatments? Yes No 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 i	n your hou	usehold	who '	visited	the fa	rm and	l have	also ı	not I	oeen	ill?
	Yes	No											
	If yes	, How many	family me	mbers v	isited	I the fa	rm an	d are n	ot 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
goats are kept? Yes No Don't Know  IF NO, SKIP TO SECTION D  10. Which of these animals are kept on the property where you live (where your of 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:  "Let's talk 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? Less than 1 hour Between 1 and 2 hours Between 2 and 3 hours More than (mm/dd/yyyy)  15. (skip if only one visit) Approximately how much time did you (your child) spend farm? Less than 1 hour Between 1 and 2 hours Between 2 and 3 hours More than farm? Less than 1 hour Between 1 and 2 hours Between 2 and 3 hours More than farm? Less than 1 hour Between 1 and 2 hours Between 2 and 3 hours More than farm? Less than 1 hour Between 1 and 2 hours Between 2 and 3 hours More than farm? Less than 1 hour Between 1 and 2 hours Between 2 and 3 hours More than farm? Less than 1 hour Between 1 and 2 hours Between 2 and 3 hours More than farm? Less than 1 hour Between 1 and 2 hours Between 2 and 3 hours More than farm?	NO, SKIP TO SECTION D
10.	
	· ·
	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."

34.	Did you (your child) visit the adult goat barn? Yes No Don	t Know		
	IF NO, SKIP TO QUESTION 43			
35.	Did you (your child) use hand sanitizer <u>BEFORE</u> visiting the ad 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	at 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	_		
	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	t goat b	arn?	
40	Yes No Don't Know	- ACTCD		
42.	Did you (your child) use baby wipes to clean your (their) hand barn? Yes No Don't Know	S <u>AFTER</u>	VISILII	ig the adult goat
	barn? Yes No Don't Know			
"Now	let's talk about other things you may have done at the farm.'	,		
14011,	ict's taik about other tillings 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
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
45.	bld you (your child) thew guilt wrille at the fairth?	INO	DON	LKIIOW
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."

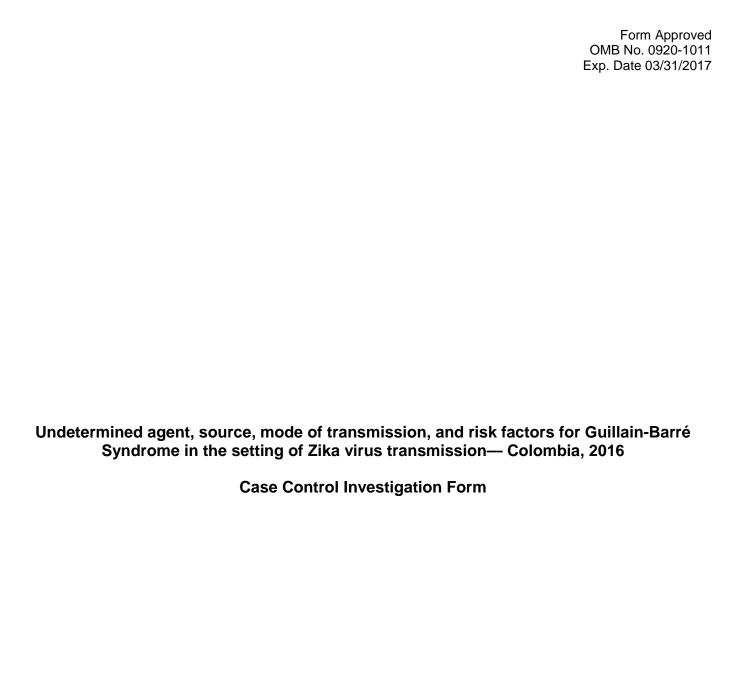
73. In the month of March, did you (your child) take any of the following types of medications?

PLE	PLEASE READ EACH MEDICATION AND CHECK YES NO DK			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 Puk	olic Health	was requesting	g 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:



COL	
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Investi	gation ID Num	ber COL		_	•	$\square$ 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 HOUSEHOLD been s	or to //	(neuro onset date for case)	), has anyone in your					
	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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							C	OL
	he two months bor on your proper			oms, wha	at pets, i	farm, or c	other animals	have lived in your
	□ Dogs	□ Cats	☐ Mice/rats		□ Pet	birds	□ F	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	lrinking water from the
	□Almost alwa	ys (>75%)	□Often (25-75	5%)		□Rarel	ly (<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	lrinking water from a
	□Almost alwa	ys (>75%)	□Often (25-75	5%)		□Rarel	ly (<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 alwa	ys (>75%)	□Often (25-75	5%)		□Rarel	ly (<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	vaded in a free	shwater river, stream, or
	□Daily	$\square$ 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	8 hours
20. Ho	w often do you v	vear long sleeve	s and pants?					
	□Almost alwa	ys (>75%)	☐Often (25-75	5%)	□Rare	ely (<25%	(a) □N	ever (0%)
22. Do	you normally we	ear insect repell	ant?					
	□Almost alwa	ys (>75%)	□Often (25-75	5%)	□Rare	ely (<25%	6) □N	ever (0%)
23. Do	you leave the wi	indows open at	your house?					
	☐Yes, during t	the day   \text{Yes}	, at night $\Box Y \epsilon$	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	c all that a	apply)?	
	□Local air con	nditioning (at lea	ast 1 room)	∃Fans		□None	,	
	w often do you h /cistern, septic ta		standing water ar	ound the	outside	e of your l	house (e.g. bı	ickets, water
		2-3 times/week	□Once/weel	k □E	Every of	her week	□Never	
27. Are	these containers	s covered?						
	□ Yes □ N		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 you	ı eaten or drunk an	y of the following f	foods at least on	ce per week (check all that apply)?			
	$\square$ Beef	☐ Lamb	☐ Chicken	☐ Fish	☐ Shellfish			
	□ Milk	☐ Cheese	☐ Yogurt	☐ Fresh salad	/ uncooked greens			
30. <u>Hu</u>	ghes Disability	y Score: (Date re	corded/	_/)				
	Hughes Disability Score (0 to 6):							
	-	-			f running, 2 = Able to walk 10 metres or h 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



	DD MM AAAA le Inicio de Síntomas Neurológicos:  DD MM AAAA  DD MM AAAA
s sig	DD WIN AAAA
	uientes preguntas son para ser realizadas durante la entrevista:
1.	Dirección actual:
	Dirección Ciudad o Municipio Distrito o Departamen
2	Dirección donde se presentaron los síntomas:
4.	/
	(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
	Sexo:   Masculino   Femenino
5.	Pertenencia étinica: ☐ Indígena ☐ ROM/Gitano ☐ Raizal ☐ Palenquero
٠.	□ 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?
	□ Sí □No □ No sabe
	A dónde:
10	
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 Pulmanar Cránica ☐ Cánacr
	<ul> <li>☐ Asma</li> <li>☐ Enfermedad Obstructiva Pulmonar Crónica</li> <li>☐ Cáncer</li> <li>☐ Cirugía (dentro de los meses de inicio de síntomas)</li> </ul>



COL			
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	•	•	na condiction que pueda afectar su	• •						
11.	enfermo (a)?	DD MM	/(fecha de inicio de cuadro l AAAA	neurológico), estuvo						
	□ Sí □No	☐ No sabe								
			ermo(a) por primera vez? 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 □ Se	creción nasal	☐ Dolor de pantorrillas						
	<b>d.</b> Si estuvo enfermo (a ☐ Sí ☐ No		o fue al hospital por esta enfermed	dad?						
	¿Cuál médico?		¿Qué hospital?							
				☐ No sabe						
12.		DD MM a e haya estado enfermo (a		s neurológicos), ¿hubo						
	<b>b.</b> Si la respuesta es afi	rmativa, ¿en qué fecha s	se enfermó la primera persona de	su hogar?						
	DD MM AAAA  c. Si alguien en su hoga	ar estuvo enfermo (a) ¿(	Qué síntomas tuvo? (Marque tod	los 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 □ Se	creción nasal	☐ Dolor de pantorrillas						
13.	¿Qué vacunas ha recibi	do y cuándo?								
	☐ Información verifica	ada en el carné de vacun	as   Información proveída ve	rbalmente						
	<ul><li>a. Triple viral (SRP o N</li><li>b. Polio</li><li>c. Fiebre Amarilla</li></ul>	<b>a.</b> Triple viral (SRP o MMR) <b>b.</b> Polio  DD MM AAAA  Dosis adicionales:								



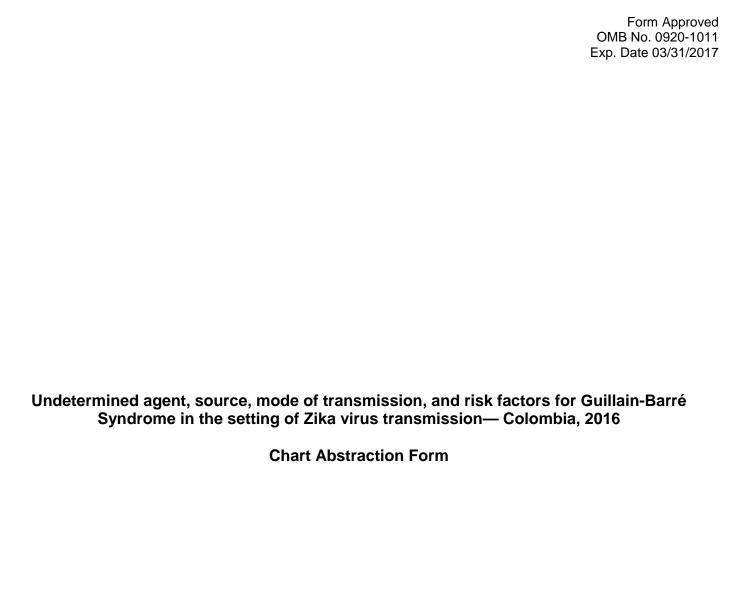
SAI	LUD	COL			
	d. BCG e. DPT f. Haemophilus Influenza B g. Neumococo h. Meningitis i. Hepatitis B j. Otras vacunas (ejemplo: rabia, encefalitis Japone ¿Cuál?				
14.	En los dos meses antes del inicio de los síntomas no animales vivieron en su casa o su propiedad? (Marc			les de granja u otro	os
	☐ Perros ☐ 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	unca (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 d	e su casa abierta	s?			
	□Sí, d	urante el día	□Sí, en la noch	e □Sí, todo el	tiempo [	□Las ventanas no se de esta casa	jan abiertas en
24.	¿Cuán	tas de sus ven	tanas o puertas ti	ienen angeos inta	actos?		
	□Toda	as $\square$	Algunas de ellas	□Ninguna			
25.	¿Tiene	en en su casa a	lguno de los sig	uientes tipos de a	ire acondici	onado?	
	□Aire	acondicionado	central □A	Aire acondicionado	por habitaci	ón □Ninguno	
26.			tiene recipientes mas, tanques, inse		casa donde p	ouede haber agua esta	ncada? (por
	□Diar	iamente $\Box$ 2	2-3 veces/semana	□Una vez a la	a semana	☐Semana de por medi	o □Nunca
27.	Estos 1	recipientes se	encuentran tapac	los			
	□Sí	□No □No	o sabe				
28.	muerto	o? □Sí	□No □No			ha manipulado algún	animal
29.	En el a	ño 2016, ¿ha c		o de los siguiente		o bebidas por lo men	os una vez a l
	a.	☐ Carne	☐ Cordero	□ Pollo	☐ Pescado	☐ Mariscos	
	b.	☐ Leche	☐ Queso	☐ Yogurt	☐ Ensalada	a / verduras crudas	
30.	Puntaj	e de Discapac	idad de Hughes:	(Fecha de regis	tro:/_	/)	
		Puntaje de D	iscapacidad de H	Hughes (0 a 6): _		Desconocido	
	[()- R	ecuneración c	omnleta: sin sec	uelas 1— Síntom	as menores	v capaz de correr 2-	- Puede

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[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		F	Encounter l	level (Bı	righton 1	-5) or not neu	nro (6):
□ SOB / res	piratory distress	☐ Gait	imbalance (	not weakn	ess)/atax	aia	☐ Hand cl	umsiness/ataxia
☐ Hyporefle	exia/areflexia	☐ Face	weakness	□ Dysar	thria	□ Dys	sphagia	☐ Dysautonomia
•	om onset until max point during this n		•	•			minutes/hour	s/days/weeks
☐ Unable to	walk without assi	stance (e.	g. cane, wall	ker)			☐ Unable	to walk at all
☐ Admitted	to the hospital		☐ Admitted	d to the IC	U/CCU		☐ Intubate	d
<b>14.</b> If any blood	was taken for this	neurologi	c illness, ple	ase fill out	t the foll	owing fo	r the INITIA	L blood draw:
	DD YYYY	WBC _	HgE	<b>3</b>	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	□ Ba	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	TROPH	IYSIOLO	OGIC STUDI	ES
18. Was a lumba	r puncture (LP) do	ne?		Yes	□ No	)	☐ Unknow	'n
	/ RBCS DD YYYY		WBCS	Prot	tein (mg/	/dL)	Glucose	e (mg/dL)
			_IgG index	Olig	goclonal	bands	IgG syı	nthesis
	/ RBCS		WBCS	Prot	tein (mg/	/dL)	Glucose	e (mg/dL)
Differential_			_IgG index_	Olig	goclonal	bands	IgG syı	nthesis
<b>19.</b> Did they rece	eive any targeted tr	eatment (	IVIG/steroic	ls/plasma e	exchang	e) for this	s neuro illnes	s?
IVIG	☐ Yes	□ No	□ Unknov	wn	Start c	late	/_ MM DD	_/ <u></u>
Plasma excha	ange	□ No	□ Unknow	wn	Start c	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				
total white cell count < 50	tal 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				

		AN	NTECEDENT 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	<ul> <li>b.) How long from prior acute illness onset until admission for neuro illness? minutes/hours/days/weeks</li> <li>26. a.) What symptoms did they report having or what signs were noticed? (check all that apply)</li> </ul>						
20.	☐ Fevers	☐ Chills	☐ Nausea or Vomiting		☐ Diarrhe		
				3			
	☐ Muscle pains	☐ Joint pains	☐ Skin rash		☐ Conjun	ctivitis	
	☐ 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	uro (6):
			lable for Zika from this			□ Yes	□ No	□ Unknown
			PAST MEDICAL	L, SOCIAL ANI	O FAMIL	Y HISTO	RY	
30.	What medical c	condition	s are listed in the admiss	sion history and	physical	(H&P)?		
	☐ Hypertensio	n	☐ Diabetes	$\square$ HIV	□ Aut	oimmune d	lisorder	<u>-</u>
	☐ 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

<b>.</b>	
NACIONAL DE	
SALED	Nú

Número	4.	140.	tifi.	ممنذم
Niimero	ae	Ider	TTTTC.	acıon

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

		echa de revisión: //
Nú	Número de Historia Clínica:	MM DD YYYY
1.	1. Primer Nombre: Segundo Nombre	:
	2. Primer Apellido: Segundo Apellido	
3.	3. Edad (años): Fecha de Nacimie	
		MM DD YYYY
4.	<b>4.</b> Sexo: ☐ Masculino ☐ Femenino	
5.	5. Dirección de residencia del paciente (Incluir dirección completa, ciudad	d o municipio y departamento):
6.	6. Número de teléfono del paciente:	
7.	7. <b>a.</b> ) Fecha de ingreso hospitalario:/	
	MM DD YYYY	
	<b>b.</b> ) Fecha en la que buscó atención por primera vez:// MM DD YYYY	
	c.) Fecha de egreso hospitalario/muerte: / /	
	MM DD YYYY	
8.	8. Egresó hacia:	
	☐ Hogar ☐ Centro de Rehabilitación ☐ Remitido a otra insti	tución hospitalaria
	☐ Muerte ☐ Otro (Especifique)	
	ENFERMEDAD ACT	TUAL
9.	9. ¿Cuánto tiempo transcurrió desde el inicio de los síntomas hasta el ingr	reso al hospital? minutos/horas/días/semanas
		•
10.	10. ¿Cuáles fueron los síntomas neurológicos iniciales dentro de los tres dí todas las opciones que apliquen, signos del examen físico y síntomas de	
	☐ Debilidad en extremidades inferiores ☐ Debilidad en extremidades	dades superiores   Diplopia/Oftalmoplejía
	☐ Adormecimiento de extremidades inferiores/parestesias	
	☐ Adormecimiento de extremidades superiores/parestesias	
	☐ Adormecimiento de la cara/parestesias	
	☐ Difficulted para respirar/distress respiratorio ☐ Trastornos de	la marcha (sin dahilidad)/atavia

	Número de Identificación COL Nivel (Brighton 1-5) o no neurológico	(6):
	☐ Trastornos de la motricidad manual/ataxia	
11.	☐ Hiporeflexia/areflexia ☐ Debilidad en la cara ☐ Disartria ☐ Disfagia ☐ Disauto  11. ¿Qué síntomas neurológicos ocurrieron en CUALQUIER MOMENTO durante la enfermedad neurológica? ( todas las opciones que apliquen, signos del examen físico y síntomas de historia de enfermedad actual)	
	☐ Debilidad en extremidades inferiores ☐ Debilidad en extremidades superiores ☐ Diplopia/Oftalm	noplejía
	☐ Adormecimiento de extremidades inferiores /parestesias	
	☐ Adormecimiento de extremidades superiores /parestesias	
	☐ Adormecimiento de la cara /parestesias	
	☐ Dificultad para respirar / distress respiratorio ☐ Trastornos de la marcha (sin debilidad)/ataxia	
	☐ Trastornos de la motricidad manual/ataxia	
	☐ Hiporeflexia/areflexia ☐ Debilidad en la cara ☐ Disartria ☐ Disfagia ☐ Disautonoi	mía
	12. ¿Cuánto tiempo transcurrió desde el inicio hasta la presentación de los síntomas neurológicos más severos? minutos/horas/días/semanas	
13.	13. Marque todas las opciones que apliquen al paciente que se presentaron al momento de mayor severidad del c neurológico:	cuadro
	☐ Incapacidad para caminar sin asistencia (por ejemplo: bastón, caminador) ☐ Incapacidad total par	ra caminar
	☐ Ingreso al hospital ☐ Ingreso a Unidad de Cuidado Intensivo (UCI) ☐ Intubación	
14.	14. Si se extrajo muestra de sangre como parte de los análisis de laboratorio para el cuadro neurológico, por favo complete las siguiente información de la muestra de sangre obtenida INICIALMENTE:	or
	Fecha/ / Recuento de blancos Hemoglobina Plaquetas	
	Sodio Potasio BUN Creatinina Glucosa Bilirrubina To	tal
	AST ALT Fosfatasa Alcalina	
15.	<b>15.</b> ¿Se documentó hiporeflexia/areflexia? ☐ Sí ☐ No ☐ Desconocido	
16.	16. a.) ¿Hubo evidencia de signos de motoneurona superior?	
	□ Sí □ No □ Desconocido	
	b.) En caso afirmativo, ¿Se documentó algunos de los siguientes hallazgos?	
	☐ Hiperreflexia ☐ Aumento en el tono/espasticidad ☐ Babinski/Hoffman ☐ Clonus sos	tenido

Número de Identificación	COL		1	Nivel (Brighton 1-5) o	no neui	ológico	o (6):
17. ¿Se documentó algún nivel sensi	tivo?	□ Sí	□ No	☐ Desconocido	)		
☐ Guante y Bota ☐ C	Cervical		$\square$ Dorsal	☐ Lumbar			
LABORATORIO	, IMÁGENE	S DIA	.GNOSTICAS Y E	STUDIOS ELECTRO	FISIOL	LÓGIC	OS
18. ¿Se llevó a cabo una punción lun	nbar? [	□ Sí	□ No	☐ Desconocide	0		
Fecha punción lumbar/_		Recue	ento de eritrocitos _	Recuento d	de leuc	ocitos _	
	DD YYYY (	Glucos	a (mg/dL)	_ Diferencial			
Fecha punción lumbar/_ MM	/ DD YYYY	Recue	ento de eritrocitos _	Recuento o	de leuc	ocitos _	
Proteínas (mg/dL) Diferencial	(						
19. ¿Recibieron algún tratamiento es intravenosa/esteroides/recambio			jar esta enfermedad	d neurológica (Inmuno	globuli	na	
a. Inmunoglobulina intraven	osa □ Sí □	l No	☐ Desconocido	Fecha de inicio			/
b. Recambio plasmático	□ Sí □	l No	☐ Desconocido	Fecha de inicio		/	/
					MM	DD	YYYY
c. Esteroides	□ Sí □	l No	☐ Desconocido	Fecha de inicio			
1 17 (1 17 7 7		1 37		T 1 1	MM	DD	YYYY
d. Ventilación mecánica	□ 51 □	l No	☐ Desconocido	Fecha de inicio			YYYY
e. Otro	□ Sí □	l No	☐ Desconocido	Fecha de inicio			
					MM	DD	YYYY
20. ¿Recibió el paciente transfusión e intravenosa)	de sangre o a	lgún o	tro hemoproducto?	(otros diferentes a Inn	nunoglo	obulina	
□ Sí □ No □ Descond	cido ¿Cuá	1?		Fecha de inicio			_/ YYYY
21. ¿Fueron algunos de los siguiente espécimen y el tipo de prueba)	s patógenos e	estudia	dos? 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
•	**
	NACIONAL DE
•	SALED

SALED	Número de Identificación COL		_	Nivel (Brighton 1-5) o no neurológico (6):
	c. 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:
resu	llevaron a cabo pruebas electrodiagnósticas? (politado? (Transcriba el resultado reportado)  Sí No  Resultado			
	Fecha / / MM DD YYYY			
Ü	nál 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			NO es un caso	
Inicio agudo de debilidad flácida bilitaral y relativamente simétrica de las extremidades * Al carecer de				
Reflejos tendinosos profundos disminuidos o ausentes en las extremidades afectadas		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	•
		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**

$\Box$	Δn	$\sim$	a	c	nk	٦i	CC	
$\boldsymbol{L}$	en	IU	yι	а	РΙ	Ш	C5	

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:	1	1
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**

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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)		
<b>Diagnostic CXR result</b> (1=Negative, 2=Abnormal, possibly		CXRrslt
TB, 3=Abnormal, not consistent with TB, 4=Unknown [not		
completed or not available])		_
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)		0.00/5
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)	
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)	
DIF and a factor and the second of the secon	DIED
RIF resistance level (highest concentration at which isolate	RIFR
is resistant)	
Specify any other detected resistance	Oth_R
oposity any other detected resistance	Oun_rc
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, it completed as part of the diagnostic workup leading	
result, if completed as part of the diagnostic workup leading to the patient's current diagnosis of TB. 1=negative.	
to the patient's current diagnosis of TB. 1=negative, 2=positive, 3=indeterminate, 4=not done, 99=unknown)	

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)	
, ,	
History of loss to follow-up or non-compliance during TB	PrevTBfu
treatment (0= No, 1= Yes, 99=Unknown)	1
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 (WIWI/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])	
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**

Case Last Name Case First Name Alternate Names/Nicknames/Aliases:  Age Date of Birth If proxy interviewed, name and relationship to case patient:  Check the database for the patient's estimated infectious period.  Start of infectious period:  Explain to the patient that you have been asked by the health department why there have been more cases of tuberculosis, or TB. Explain that you series of questions to try to identify where the health department might health people who have TB, as well as to figure out where the patient might has Acknowledge that the patient has already participated in many interview providers. Reassure the patient that all answers will be kept confidentia of the interview is to learn information that can help stop the spread of The people from getting sick (emphasize protection of families). Thank the people from getting sick (emphasize protection of families).	Lname Fname Alias  Age DOB
Age Date of Birth  If proxy interviewed, name and relationship to case patient:  Check the database for the patient's estimated infectious period.  Start of infectious period:  End of infectious period:  Explain to the patient that you have been asked by the health department why there have been more cases of tuberculosis, or TB. Explain that you series of questions to try to identify where the health department might be people who have TB, as well as to figure out where the patient might have providers. Reassure the patient that all answers will be kept confidentiat of the interview is to learn information that can help stop the spread of Talenta in the patient of the interview is to learn information that can help stop the spread of Talenta in the patient in the patient information that can help stop the spread of Talenta in the patient information that can help stop the spread of Talenta in the patient information that can help stop the spread of Talenta in the patient information that can help stop the spread of Talenta in the patient information that can help stop the spread of Talenta information that can help stop the spread of Talenta information that can help stop the spread of Talenta information that can help stop the spread of Talenta information that can help stop the spread of Talenta information that can help stop the spread of Talenta information that can help stop the spread of Talenta information that can help stop the spread of Talenta information that can help stop the spread of Talenta information that can help stop the spread of Talenta information that can help stop the spread of Talenta information that can help stop the spread of Talenta information that can help stop the spread of Talenta information that can help stop the spread of Talenta information that can help stop the spread of Talenta information that can help stop the spread of Talenta information that can help stop the spread of Talenta information that can help stop the spread of Talenta information that can help stop the sp	Alias Age
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End of infectious period:  Explain to the patient that you have been asked by the health department why there have been more cases of tuberculosis, or TB. Explain that you series of questions to try to identify where the health department might be people who have TB, as well as to figure out where the patient might have Acknowledge that the patient has already participated in many interview providers. Reassure the patient that all answers will be kept confidential of the interview is to learn information that can help stop the spread of Tallows	
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time and for speaking with us.	ou will be asking a be able to find other ave gotten sick. vs with health care I, and that the purpose TB and prevent other
Note that throughout the interview, the period of interest is 2 years before the start of the infectious period to the end of the in	fectious period.
Ask patient whether they are from the Tribe A Reservation. If not, ask where patient came from and when he/she came to the a	

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 before you got sick with TB. People sick with TB often have a bad cough, or might lose a 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 got 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 trained to blome anything the people are not in trouble, and treat them
trying to blame anyone. We are trying to make sure we can find all sick people and treat them.
'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?

lon-injection drug ("taking anything for recreation, e.g. marijuana") se within 1 year before diagnosis =Never =Rarely (1-2 times ever) =Occasionally (more than 1 or 2 times, but less than most days or nights) =Frequently (most days or nights of the week) =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:						
Drug 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: **q** 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 qYes (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?	<b>q</b>	res <b>q</b> N	lo				
4. Have you had fevers or chills?	<b>q</b> \	res <b>q</b> N	lo				
Have you had night sweats?     (completely soaking your clothes at night)	<b>q</b> /	es <b>q</b> N	lo				
Have you been tired or feeling weak lately?	<b>q</b> /	res <b>q</b> N	lo				
7. Have you lost your appetite?		res <b>q</b> N					
8. Have you had unplanned weight loss?		res <b>q</b> N Jnknown		If yes,	yes, how much?		
Ø Is this contact symptomatic for TB?	qYes <b>q</b> No ? If yes, specify symptom start date://						
TB Risk Factor Screening					Notes		
Instructions: Screen to see if the contact has risk factors	that	could incre	ase their risk for pro	gression t	to active TB disease.		
1. Is this contact >50 years old?		<b>q</b> Yes	qNo				
2. Was this contact <5 years old during the exposure period?		<b>q</b> Yes	<b>q</b> No				
3. Do you have diabetes?		<b>q</b> Yes	<b>q</b> No or Unknow	wn			
4. Do you have HIV?		<b>q</b> Yes	<b>q</b> No or Unknow	wn			
5. Do you have cancer?		qYes	<b>q</b> No or Unknow	wn			
6. Do you take prednisone every day?		<b>q</b> Yes	<b>q</b> No				
7. Do you smoke tobacco?		<b>q</b> Yes	qNo				
8. Do you drink alcohol?		<b>q</b> Yes	<b>q</b> No	If y	es, specify amount/frequency		
9. Do you use any other substances?		<b>q</b> Yes	<b>q</b> No	and	ves, include types/routes, freque d locations where substances quired and used	∍ncy,	
Ø Does this contact have a high-risk condition?  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>						
			END	QUEST	TIONS		
ts		Date TST Placed	Date TST Read	ММ	Chest X-Ray		
Test Results	TST 1: TST 2: TST Inter  Regative	pretation: ve <b>q</b> Positive	If pos, Conversion	n? <b>q</b>	CXR Date:/  CXR Result: <b>q</b> Not Suggestive of TB <b>q</b> Suggestive of TB		
Test Results		Date of IGRA erpretation: ve <b>q</b> Positive	IGRA Resu		Chest X-Ray  CXR Date:/  CXR Result: <b>q</b> Not Suggestive of TB <b>q</b> Suggestive of TB		
Rx Start Date:    Rx End Date:							
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:/							