		ACTIVE BACTERIAL CORE SUR	VEILLANCE CASE	REPORT –		
Patient's Name:					Phone No.:()	
Address:		Patient Chart No.:				
(Number, Street, Apt. No.)		Hospital:				
(City, State) - Patient Identifier information is not transn	sitted to CDC	(Zip Code	,			
Patient identifier miormation is not transm DEPARTMENT OF HEALTH AND HUMAN SE CENTERS FOR DISEASE CONTROL AND PREVENTION ATLANTA, GA 30333	202	4 ACTIVE BA SILLANCE (AI NENT OF THE EM - DARK SHADED AREAS	3Cs) Ca erging i	SE REPORT NFECTIONS P		
1. STATE: 2. STATE I.D.: 3. PA	TIENT I.D.: 4. Date reported Mo. Day	to Eir Site.	RF Status:	Incomplete 3 E	6. COUNTY: (Patient Residence	6a. PLANNING REGION: (Patient Residence)
			Chart unavailat after 3 requests	ole 7 QA Review	Change	
7a. HOSPITAL/LAB I.D. 8. DATE OF BIRTH	l: 9a. AGE:		D.SEX: 11	a. ETHNIC ORIGIN:	11b. RACE: (Check all that	
WHERE PATIENT TREATED:: Mo. Day	Year 9b. Is age i		LI Male	Hispanic or Latino Not Hispanic or La		n ve Hawaiian
		s 2 Mos. 3 Yrs.		Unknown	or O	ther Pacific Islander rican Indian or Alaska Native
Lab Repeating Group Section T1-T10						
T1 T2	of Specimen Collection	T3 Test Method Hos	a spital/Lab I.D.	T4 Site from which	T5 h Bacterial Spec	T6 ies Test Result
Test Type Date			ere test identifi			—
1						1=Positive 0=Negative
2				_		1=Positive
3				=		0=Negative
				_		0=Negative
4				_		1=Positive 0=Negative
T7 T8 Isolate/Specimen If isolate/specime	T9 T10	#T1 - Test		Test Method (if non-cultur ofire Filmarray Meningitis/E		T5 - Bacterial Species Isolated
Isolate/Specimen If isolate/specime Available? If isolate/specime N/A, why not?	en Shipped to If shipp CDC? access	eu,	2=Oti 3=Bio	her ofire Filmarray Blood Cultu	re ID (BCID) Panel	1=Neisseria meningitidis 2=Haemophilus influenzae
1 =Yes	1=Yes 0=No	9=Unknov	5=Bri	rigene Gram + Blood Culti uker MALDI Biotyper CA S nknown		3=Group B Streptococcus 5=Group A Streptococcus 6=Streptococcus pneumoniae
2 = 1=Yes	1=Yes					* For other bacterial pathogens (i.e. non-ABCs), write in pathogen name
☐ 2=No	0=No	T4 - Site 1=Blood 2=Bone	8=Other 9=Unkno		Non Sterile Sites icardial Fluid 27=Wound itoneal Fluid	T8 - No Isolate, why not 1=N/A at Hospital Lab
3 \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \	□ 0=No	3=Brain 4=CSF	10=Liver 11=Lymp	17=Ple th Node 18=Spl	ural Fluid een	2=N/A at Hospital Lab 2=N/A at State Lab 3=Hospital Refuses
4 1=Yes 2=No	1=Yes 0=No	5=Heart 6=Joint 7=Kidney	12=Musc 13=Ovar 14=Panc	20-111	cular Tissue eous Fluid	4=Isolate Discrepancy (2x) 5=No DNA (non-viable) 6=Isolate Not Needed
		Date of discharge:	14=1 0.10	1	nospitalized, was this patient a	·
HOSPITALIZED? Mo. I	Day Year	Mo. Day Yea	ar	ICU during ho		
T La Yes 2 La No				1	□ No 9 □ Unknown	
18a. Where was the patient a resident at t	_	medical ward		nt of a facility, what name of the facility?	19a. Was patient transferred from another hospital?	19b. If YES, hospital I.D.:
- I made residence		r (specify):			1 ☐ Yes 2 ☐ No	
3 Long term acute care facility 6 Co		* * * * * * * * * * * * * * * * * * * *	Facility ID: _		9 🗆 Unknown	
			PE OF INSURANCE: (Check all that apply)			
1 Private						
		1 Medicare 1 Indian Health Service (IHS) 1 Uninsured				
20c. BMI: OR		1 ☐ Medicaid/state 1 ☐ Incarcerated 1 ☐ Unknown assistance program				
22. OUTCOME: 1 Survived 2 Died 9 Unknown 22a. If survi			ived, patient discharged to: 1 Home 2 LTC/SNF 3 LTACH 5 Left AMA 9 Unknown			
23. If patient died, was the culture obtain 1 Yes 2 No 9 Unkn	If discharged to LTC/SNF or LTACH, list Facility ID: 4 Other, Specify:					
24a.At time of first positive culture, 24b. If pregnant or postpartum, what w			s the outcome of fetus: 25. If patient <1 month of age, indicate gestational age and birth weight.			
1 Survived, no apparent illness 1 Pregnant 2 Postpartum 2 Survived, clinical infection 3 Live birth/neor			makal de sils	If pregnant, indicate gestational age of fetus, only.		
3 Neither 9 Unknown 4 Abortion/stillbirth 5 Induced aborti				Gestational ag	e: (wks) Birth weight	t: (gms)

- IMPORTANT - PLEASE COMPLETE THE BACK OF THIS FORM -

9 Unknown

6 Still pregnant

Public reporting burden to collect this information is estimated to average 15 minutes per response, including time for reviewing instructions, searching existing data sources, gathering/maintaining the data needed, and completing/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 information, including suggestions for reducing this burden to CDC, CDC/ATSDR Reports Clearance Officer, 1600 Clifton Rd. MS D-74, Atlanta, GA, 30333, ATTN: PRA(0920-0978) **Do not send the completed form to this address.**

1 ☐ Abscess (not skin) 1 ☐ Chorioamnionitis 1 ☐ Empyema 1 ☐ Necrotizing fasc							
1 Bacteremia 1 Endocarditis 1 Hemolytic uremic 1 Osteomyelitis	1 Pericarditis 1 Septic abortion 1 STSS						
without Focus 1 Epiglottitis syndrome (HUS) 1 Otitis media 1 Cellulitis 1 Fodometritis 1 Meningitis	1 ☐ Pneumonia 1 ☐ Septic arthritis 1 ☐ Other (specify):						
The Endometries							
27. UNDERLYING CAUSES OR PRIOR ILLNESSES: (Check all that apply OR if NONE or CHART							
1 AIDS or CD4 count <200 1 Connective Tissue Disease (Lupus, etc.)	1 ☐ Immunosuppressive Therapy (Steroids, etc.) 1 ☐ Peripheral Neuropathy 1 ☐ Eculizumab (Soliris) - N.men. only 1 ☐ Peripheral Vascular Disease						
1	1 Ravulizumab (Ultomiris) - N.men. only 1 Peripiretal vascular bisease						
1 Bone Marrow Transplant (BMT) 1 Dear/Prolound Hearing Loss	1 ☐ Leukemia 1 ☐ Premature Birth (specify gestational						
1 CVA/Stroke/TIA 1 Diabetes Mellitus,	1 Multiple Myeloma age at birth) (wks)						
1 Chronic Hepatitis C 1 HbA1C(%), Date//	1 Multiple Sclerosis 1 Seizure/Seizure Disorder						
1 Chronic Kidney Disease 1 Emphysema/COPD	1 Myocardial Infarction 1 Sickle Cell Anemia						
1 L Chronic Liver Disease/cirrhosis 1 L Heart Failure/CHF	1 ☐ Nephrotic Syndrome 1 ☐ Solid Organ Malignancy						
1 ☐ Current Chronic Dialysis 1 ☐ HIV Infection 1 ☐ Chronic Skin Breakdown 1 ☐ Hodgkin's Disease/Lymphoma	1 ☐ Neuromuscular Disorder 1 ☐ Solid Organ Transplant 1 ☐ Obesity 1 ☐ Splenectomy/Asplenia						
1 Cochlear Implant 1 Immunoglobulin Deficiency 1 Parkinson's Disease							
1 Complement Deficiency	1 Peptic Ulcer Disease						
SUBSTANCE USE, CURRENT							
27b. SMOKING: 1 ☐ None 1 ☐ Unknown 1 ☐ Tobacco 1 ☐ E-Nicotine Delivery System (check all that apply)	1 Marijuana 27c. ALCOHOL ABUSE: 1 Yes 0 No 9 Unknown						
_	e Disorder (DUD)/Abuse Mode of delivery: (check all that apply)						
1 Marijuana/cannabinoid (other than smoking) 1 DUD or Al	: — : — : p -p p : — : : — : :						
1 ☐ Opioid, DEA schedule I (e.g., heroin) 1 ☐ Opioid, DEA schedule II - IV (e.g., methadone, oxycodone) 1 ☐ DUD or Al							
1 ☐ Opioid, DEA schedule II - IV (e.g., methadone, oxycodone) 1 ☐ DUD or AI 1 ☐ DUD or AI 1 ☐ DUD or AI							
1 ☐ Cocaine 1 ☐ DUD or Al							
1 DUD or Al	use 1 IDU 1 Skin popping 1 non-IDU 1 Unknown						
1 U Other* (specify): 1 DUD or Al							
1 Unknown substance 1 UDUD or Al							
- IMPORTANT - PLEASE COMPLETE F	OR THE RELEVANT ORGANISM –						
HAEMOPHILUS INFLUENZAE 28a. What was the serotype? 1 b 2 Not Typeable 3 a 4 c 5 d 6	e 7 🗆 f 8 🗆 Other (specify):9 🗔 Not tested or Unknown						
28b. If <15 years of age and serotype 'b' or 'unknown' did 1 Yes 2 No 9 Unknown patient receive Haemophilus influenzae b vaccine? If YES, please complete the list below.							
	OSE DATE GIVEN VACCINE NAME/MANUFACTURER Mo. Day Year						
	Mo. Day Year						
1							
2	4						
	4						
2							
NEISSERIA MENINGITIDIS							
NEISSERIA MENINGITIDIS 29. What was the serogroup? 1 A 2 B 3 C 4 Y 5 W135 6 30. Is patient currently attending college? 1 Yes 2 No 9 Unknown	Not Groupable 8 Other: 9 Unknown						
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NEISSERIA MENINGITIDIS 29. What was the serogroup? 1	Not Groupable 8 Other: 9 Unknown Complete the table DOSE TYPE DATE GIVEN VACCINE NAME/ MANUFACTURER 4						
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NEISSERIA MENINGITIDIS 29. What was the serogroup? 1 A 2 B 3 C 4 Y 5 W135 6 30. Is patient currently attending college? 1 Yes 2 No 9 Unknown 31. Did patient receive meningococcal vaccine? 1 Yes 2 No 9 Unknown If YES, Type Codes: DOSE TYPE DATE GIVEN VACCINE NAME MANUFACTURE (Menactra, Menveo, Menribirix, MenQuadfi) 1 2= ACWY polysaccharide (Menomune) 2 3= B (Bexsero, Trumenba) 9= Unknown 3 32. If survived, did patient have any of the following sequelae evident upon discharge? (Chamber 1) The patient deliver a baby (vaginal or C-section) 33. Did the patient have surgery or any skin incision?	Not Groupable 8 Other: 9 Unknown Complete the table DOSE TYPE DATE GIVEN VACCINE NAME/ Mo. Day Year MANUFACTURER 4						
NEISSERIA MENINGITIDIS 29. What was the serogroup? 1	Not Groupable 8 Other: 9 Unknown complete the table DOSE TYPE						
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NEISSERIA MENINGITIDIS 29. What was the serogroup? 1	Not Groupable 8 Other: 9 Unknown Complete the table DOSE TYPE DATE GIVEN WANUFACTURER Mo. Day Year MANUFACTURER 4 Unknown Falysis or spasticity 1 None 1 Unknown Falysis or spasticity 1 Skin Scarring/necrosis 1 Other (specify): 35. Did patient have: 1 Varicella 1 Surgical wound (post operative) 1 Penetrating trauma (post operative) 1 Blunt trauma If YES to any of the above, record the number of days prior to the first positive culture Date:						
NEISSERIA MENINGITIDIS 29. What was the serogroup? 1 A 2 B 3 C 4 Y 5 W135 6 30. Is patient currently attending college? 1 Yes 2 No 9 Unknown 31. Did patient receive meningococcal vaccine? 1 Yes 2 No 9 Unknown If YES, Type Codes: DOSE TYPE DATE GIVEN WACCINE NAME 1= ACWY conjugate (Menactra, Menveo,	Not Groupable 8 Other:						
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