2019 HAIC Invasive *Staphylococcus aureus* Supplemental Surveillance Officer Survey

Please answer the following questions for the year <u>2018</u>. The purpose of the survey is to verify and document current surveillance procedures, including cases ascertainment and auditing methods. Please return an electronically completed copy or scanned copy of the completed survey to <u>gqv8@cdc.gov</u>. If you have any questions, please contact Kelly Jackson (<u>gqv8@cdc.gov</u>).

Site: ____CA ___CT ___GA ___MD ___MN ___NY ___TN Person(s) Completing the Form: ____

Surveillance Area Characteristics

- 1. Is MRSA reportable at your site? _____ yes _____ no

 a. If yes:
 - a. If yes:
 - i. What is your reportable definition of MRSA?
 - _____ All invasive MRSA statewide
 - _____ Invasive MRSA in residents among defined catchment area
 - _____ Healthcare-associated invasive MRSA infection
 - _____ Other, please define: ______
 - ii. Is isolate submission to the State Health Department Laboratory required?
 - _____ yes _____ no
 - b. If no:
 - i. What mechanism do you have in place that allows for SOs to have access to case counts and medical records?
 - _____ Agent of the state
 - _____ State Health Department Regulation
 - _____ Other, please explain: ______
 - ii. Does your state/site plan to make MRSA reportable? _____yes _____no
- Did your site send MRSA isolates to CDC for characterization in 2017? _____yes _____no
 a. If yes, how were isolates selected? ______
- 3. How does your site complete SA case report forms (please select all that apply)?
 - ____On a computer or tablet
 - _____With paper and pen
 - ____Other, please explain: _____

4. Did your site participate in MSSA surveillance in 2017? _____yes _____no

- a. If yes, what mechanism did you have in place that allowed for SOs to have access to MSSA case counts and medical records?
- MSSA is a reportable condition
- _____ Agent of the state
- _____ State Health Department Regulation
- _____ Other, please explain: _____

Public reporting burden of this collection of information is estimated to average 10 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 current 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 Rd NE, MS D-74, Atlanta, Georgia 30329; ATTN: PRA (xxxx-xxxx)

b. If yes, please complete the date range for which MSSA surveillance was conducted in 2017 (month/year) as well as the catchment area:

	2017
Dates of MSSA surveillance	
Catchment area	

- 5. Are you able to directly access any National Healthcare Safety Network (NHSN) data?
 - a. If yes:
 - i. Please mark which NHSN data your site can access
 - _____ Hospital MRSA LabID event
 - _____ Hospital central line-associated bloodstream infection (CLABSI) data _____ Dialysis event

b. If no:

i. Does a public health partner have access to NHSN data? For example, if your site is not based at the state HD does the state HD have access to the data? If your site is part of the state HD and you do not have direct access to the NHSN data, does another department/section have access to the data?

_____ yes _____ no

- 1. If yes:
 - a. Please mark which NHSN data can be accessed _____ Hospital MRSA LabID event _____ Hospital CLABSI data
 - _____ Dialysis event
- 3. How often do you access/are you provided with NHSN data?
 - _____ Weekly
 - _____ Monthly
 - _____ Never
 - _____ Other, please specify ______
- 4. What do you use NHSN data for?_____

Lab Participation and Case Finding

Please answer the following questions for hospitals and labs under surveillance for 2017.

1. Please list the total number of each type of lab <u>serving</u> your MRSA surveillance catchment area (both inside and outside of the catchment area):

Inside catchment area	Outside catchment area	
		Hospital laboratories
		Dialysis referral laboratories
		Commercial/outpatient laboratories*
		Other; please specify:
		Total number (Add above together)
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*For the purpose of the survey, we are defining "Commercial/Outpatient Laboratories" as any for profit laboratory, not including dialysis referral laboratories, that serve health care facilities in a given surveillance catchment area. Examples include LabCorp and Quest.

2. Please list the total number of each type of lab <u>serving</u> your MSSA surveillance catchment area *if different catchment than MRSA* (both inside and outside of the catchment area):

Inside catchment area	Outside catchment area	
		Hospital laboratories
		Dialysis referral laboratories
		Commercial/outpatient laboratories*
		Other; please specify:
		Total number (Add above together)
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*For the purpose of the survey, we are defining "Commercial/Outpatient Laboratories" as any for profit laboratory, not including dialysis referral laboratories, that serve health care facilities in a given surveillance catchment area. Examples include LabCorp and Quest.

3. Please indicate the culture sources your site requests from participating labs for surveillance other than blood, CSF, pleural fluid, peritoneal fluid, pericardial fluid, joint/synovial fluid, bone, and muscle?

4. Indicate the percentage contribution of each case finding method to your site's total SA case counts (100%) in 2017.

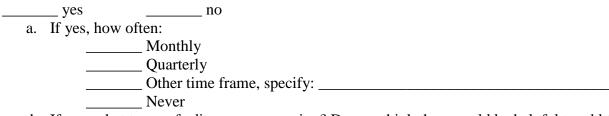
Case Finding	% MSSA	% MRSA	Method
Method Used?	Case Count	Case Count	
	Contribution	Contribution	
$\square Y \square N$			NETSS/NEDSS or other passive state reporting system
$\square Y \square N$			Retrospective review of received line lists from <i>hospital</i> labs
			Routinely received line lists from <i><u>commercial/outpatient</u></i> labs
			Routinely received line lists from <i>dialysis referral</i> labs
Y N			Regular lab visits; <i>frequency</i> :
Y N			ICPs submitting case report form
Y N			Isolates being received at state lab
Y N			NHSN
Y N			Other, please specify

	a. Do	 you expect this distribution and/or percentage values to change in 2017? yes no i. If yes, please explain why:
		eporting invasive SA, how many of the participating labs are providing case reports irect electronic messaging, such as HL7 messaging?
	a. If le	ess <100%, how else are you receiving reports?
	b. Wł	nat are the perceived barriers to use of direct electronic messaging?
		ow important is electronic messaging for your site? (1 not important at all; 5very high iority)
Γ	Did any la	abs drop out of participation in 2017? yesno
		a. If yes, how many?b. Why did these labs drop out of participation?
		lid you identify any additional labs, regardless of location, which identify invasive SA om persons who are residents of your catchment area?
	a.	
	b.	-
		If yes, how did you find these labs? If yes, how many labs did you find?

d. Approximately how many cases does this/these lab(s) identify each year among residents of your catchment area? _____

Data Edits

1. Does your site run a data edit program in addition to the CDC edit program that is sent out monthly?



b. If yes, what types of edits are you running? Do you think they would be helpful to add to CDC's edit process?

Ascertainment of Surveillance Area* and Case Audits*

*"Case ascertainment" should include ongoing attempts to identify new or additional laboratories inside and outside of your defined catchment area which may be processing MRSA specimens for surveillance area residents.

*Audits of all laboratories both within the ABCs MRSA surveillance area and those outside are required once a year. The purpose of the audit is to ensure that all cases of invasive MRSA are being captured. The primary data source at every reporting laboratory (e.g. laboratory log slips/log book, computer-generated electronic printouts, case reports, line lists) should be reviewed for invasive MRSA cases and compared to the list of cases that were reported to the surveillance personnel.

- 1. How did your site define an audit case in 2017?
- 2. Indicate the percentage contribution of each case finding method to your site's <u>audit counts</u> (100%) in 2017.

Audit Method	% MSSA	% MRSA	Method
Used?	Audit Count	Audit Count	
	Contribution	Contribution	
$\square Y \square N$			NETSS/NEDSS or other passive state reporting system
$\Box Y \Box N$			Retrospective review of received line lists from <i>hospital</i> labs
$\Box Y \Box N$			Routinely received line lists from <i><u>commercial/outpatient</u></i> labs
$\Box Y \Box N$			Routinely received line lists from <i>dialysis referral</i> labs
$\Box Y \Box N$			Regular lab visits; <i>frequency</i> :
$\Box Y \Box N$			ICPs submitting case report form
			Isolates being received at state lab
Y N			NHSN comparison
Y N			Other, please specify

3.	*Au to e slip	your site assess your individual laboratory case auditing* methods? dits of all laboratories both within the HAIC MRSA surveillance area and those outside are required once a year. The purpose of the audit is nsure that all cases of invasive MRSA are being captured. The primary data source at every reporting laboratory (e.g. laboratory log s/log book, computer-generated electronic printouts, case reports, line lists) should be reviewed for invasive MRSA cases and compared to list of cases that were reported to the surveillance personnel. Yesno
	a.	If no, please explain why:
	b.	If yes, how often is this performed? When was this last performed?
	c.	If yes, how does your site perform audits*? Remove negative restrictions from line list reports Review actual query codes Review selection criteria Other, please specify
		 i. If you picked "Review actual query codes", can you see pathogen resistance information? For example, in the query code can you see that the laboratory is including isolates resistant to oxacillin for MRSA? YesNo
1.	*"C	Your site perform routine ascertainment* of the surveillance area? <i>Tase ascertainment</i> " should include ongoing attempts to identify new or additional laboratories inside and outside of your defined catchment a which may be processing specimens for surveillance area residents. <u>Yes</u> <u>no</u>
	a.	If yes, how does your site assess case ascertainment* methods? (examples include: physician surveys, LTCF surveys, outreach to new dialysis centers, etc).
	b.	If yes, how often is this performed? When was this last performed?
5.	Are th	ere specific labs that you have difficulty obtaining line lists from?
	a.	If yes, what types of labs?

6. Does your site have checks in place to recognize decreasing/increasing case counts or rates of MRSA disease?

a. If yes, please describe the check(s) that you use_____

- _____
- b. If yes, how often are the check(s) used?

c. If yes, do you plan to use these for MSSA once more surveillance data are available? _____ yes _____ no

Geocoding

1. Is your site continuing to geocode SA cases? ______ yes ______ no

CDC Responsibilities

- 1. CDC staff are responsive to questions/concerns/emails (e.g., Valerie Albrecht, Kelly Jackson, Isaac See, and Shirley Zhang).
 - ______ Strongly agree

 ______ Agree

 ______ Neutral

 ______ Disagree

 ______ Strongly disagree
 - a. If you disagree or strongly disagree, please explain and provide improvement suggestions:

2. Monthly surveillance officer calls are a valuable use of my time.

Strongly agree
Agree
Neutral
Disagree
Strongly disagree

a. If you disagree or strongly disagree, please explain and provide improvement suggestions:

- 3. What parts of the SA SharePoint site do you use the least or find the least useful?
- 4. Can you suggest any future training topics that might be useful for SA surveillance officers?

5. What SA topics would you like to see covered at the HAIC Surveillance Officer's Meeting during the SA session next year?

END Thank you very much!