2021 HAIC Invasive Staphylococcus aureus Supplemental Surveillance Officer Survey

Please answer the following questions for the year $\underline{2021}$. The purpose of the survey is to verify and document current surveillance procedures, including cases ascertainment and auditing methods. Please enter your responses into the corresponding REDCap database. If you have any questions, please contact Kelly Jackson ($\underline{gqv8@cdc.gov}$).

Site: CA CT GA MD MN NY TN Person(s) Completing the Form:
Surveillance Area Characteristics
 Is MRSA reportable at your site? yes no 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?
ii. Does your state/site plan to make MRSA reportable?
 ii. Is isolate submission to the State Health Department Laboratory required?

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)

Other, please explain:
ii. Does your state/site plan to make MSSA reportable?yesno
3. Did your site send MRSA/MSSA isolates to CDC for characterization in 2021?yesno a. If yes, how were isolates selected?
b. If yes, how many isolates did you expect to be able to collect from clinical labs? MRSA, MSSA c. If yes, what was the total number of isolates collected from clinical labs? MRSA, MSSA
 4. 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:
5. Are you able to directly access any National Healthcare Safety Network (NHSN) data? yes no
 a. If yes: 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: 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
If yes: a. Please mark which NHSN data can be accessed Hospital MRSA LabID event Hospital CLABSI data Dialysis event
6. How often do you access/are you provided with NHSN data? Weekly Monthly Never Other, please specify

7. What do you us	e NHSN data for			

Lab Participation and Case Finding

Please answer the following questions for hospitals and labs under surveillance for <u>2021</u>.

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) and the total number of each type of lab <u>participating</u> (i.e., submit test results when available) in surveillance (both inside and outside the catchment area):

Inside catchment area Outside ca Serve Participate Serve		Outside catchment area		
		Participate		
				Hospital laboratories
				Dialysis referral laboratories
				Commercial/outpatient laboratories*
				Other; please
				specify:
				Total number (Add above together)

^{*}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. *If different catchment than MRSA*, please list the total number of each type of lab <u>serving</u> your MSSA surveillance catchment area (both inside and outside of the catchment area) and the total number of each type of lab <u>participating</u> (i.e., submit test results when available) <u>in surveillance</u> (both inside and outside the catchment area)::

Inside catchment area		Outside catchment area		
Serve	Participate	Serve Participa		
				Hospital laboratories
				Dialysis referral laboratories
				Commercial/outpatient laboratories*
				Other; please
				specify:
				Total number (Add above together)

^{*}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.		e indicate the culture sources your site requests from participating labs for surveillance othe blood, CSF, pleural fluid, peritoneal fluid, pericardial fluid, joint/synovial fluid, bone, and le?			
	-				
	_				
	_				
	_				

4. Indicate the percentage contribution of each case finding method to your site's total SA case counts (100%) in 2021.

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 <u>hospital</u> labs
\square Y \square N			Routinely received line lists from <u>commercial/outpatient</u> labs
\square Y \square N			Routinely received line lists from <u>dialysis referral</u> labs
\square Y \square N			Regular lab visits; frequency:
\square Y \square N			ICPs submitting case report form
\square Y \square N			Isolates being received at state lab
\square Y \square N			NHSN
\square Y \square N			Other, please specify

	a. Do you expect this distribution and/or percentage values to change in 2022? yes no i. If yes, please explain why:
5.	For labs reporting invasive SA, how many of the participating labs are providing case reports through direct electronic messaging, such as HL7 messaging?
	a. If less <100%, how else are you receiving reports?
	b. What are the perceived barriers to use of direct electronic messaging?
	c. How important is electronic messaging for your site? (1 not important at all; 5very high priority)
6.	Did any labs drop out of participation in 2021? yes no
	a. If yes, how many?b. Why did these labs drop out of participation?

		id you identify any additional labs, regardless of location, which identify invasive SA om persons who are residents of your catchment area?
150	14105 11	yes no
	a.	If yes, how did you find these labs?
	b.	If yes, how many labs did you find?
	c.	If yes, how many of these labs were added?
		i. If not all found labs were added, why not?
	d.	Approximately how many cases does this/these lab(s) identify each year among residents of your catchment area?
Data l	Edits	
	a. b.	ly?yesno If yes, how often:MonthlyQuarterlyOther time frame, specify:Never If yes, what types of edits are you running? Do you think they would be helpful to add to sedit process?
2.	a. If y i. l	our site complete CRF re-abstractions during 2021? yes no es, did you have any challenges completing the CRF re-abstractions? yes no f yes, please describe: o, why not?

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, computergenerated 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 2021?

2. Indicate the percentage contribution of each case finding method to your site's <u>audit counts</u> (100%) in 2021.

Audit Method	% MSSA	% MRSA	Method
Used?	Audit Count	Audit 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
\square Y \square N			Routinely received line lists from <u>commercial/outpatient</u> labs
\square Y \square N			Routinely received line lists from <u>dialysis referral</u> labs
\square Y \square N			Regular lab visits; frequency:
□Y □N			ICPs submitting case report form
□Y □N			Isolates being received at state lab
Y N			NHSN comparison
$\square_{Y} \square_{N}$			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
	d. How many laboratories did you audit in 2021?
4.	In 2021, did your site update its inventory of facilities within the EIP catchment area? yes no a. If no, why not?
	a. If no, why not:
	b. If yes, how many facilities serve the catchment area?
	c. If yes, how many facilities have you identified a clinical laboratory for?
5.	Does your site perform routine ascertainment* of the surveillance area? *"Case ascertainment" should include ongoing attempts to identify new or additional laboratories inside and outside of your defined catchment area which may be processing specimens for surveillance area residents.
	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?
	Are there specific labs that you have difficulty obtaining line lists from? yes no
	a. If yes, what types of labs?
7.	Does your site have checks in place to recognize decreasing/increasing case counts or rates of MRSA disease? yes no a. If yes, please describe the check(s) that you use
	a. If yes, how often are the check(s) used?
	b. If yes, do you plan to use these for MSSA once more surveillance data are available? yes no

Geocoding

•	site geocode SA cases in 2021?
	If yes, what is the most recent year of surveillance data that was geocoded? If no, why not?
Vital Record	s Linkages
•	ite link SA cases to vital records (mortality matching) in 2021?
	If yes, what is the most recent year of surveillance data that was linked? If no, why not?
COVID-19 I	mpact
deadlines dur	ID-19 response activities delay 2021 iSA surveillance work (e.g., unable to meet iSA ing 2021)? yes no If no, how were you able to meet iSA deadlines?
b.	If yes, how did COVID-19 response activities delay your iSA work?
Jackson, I	f are responsive to questions/concerns/emails (e.g., Davina Campbell, Runa Gokhale, Kelly saac See, and Shirley Zhang). Strongly agreeAgreeNeutralDisagreeStrongly disagreeStrongly disagree you disagree or strongly disagree, please explain and provide improvement suggestions:
·	surveillance officer calls are a valuable use of my time. Strongly agreeAgreeNeutralDisagreeStrongly disagree 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!