Department of Health & Human Services Centers for Disease Control and Prevention (CDC), Atlanta, Georgia, 30333 **NREVSS Laboratory Assessment** Form Approved OMB 0920-0004

Date of interview:		Inte	Interviewer Name:					
Laboratory Name:					Lab ID:			
City:				State:	Zip Code:			
Contact Person:			Position:					
Contact Number/En	mail:							
Type of institution:								
Hospital affiliated	Clinic or physician's office	Private commercial Refe	rence University	Military (Government			
		g and the contributing facto	rs in changes to test	ing practices f	or laboratories reporting to NREVSS			
Section I: Demogra	<u>phics</u>							
	how many respiratory speci specimens in peak w		outum, BALs, NP asp	irates) does yo	our laboratory test during the winte			
2. (For reference l	abs only?) What is the geog	raphic location from which	your specimens are	collected? For	example, only specific states?			
3. Approximately, Patient type	what percentage of the spe e	ecimens tested by your lab a	re from the followin	ng:				
Jublic roporting burdon	of this collection of information i	s astimated to average 20 minute	s nor rosnonso, includio	a tha tima far ra	wlowing instructions, soarching 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 information, including suggestions for reducing this burden to CDC,

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a.	Inpatient		% / Unknowr	า			
b. (Outpatient		% / Unknow	n			
Age	categories						
c	Pediatric (0 – 18	vears)	% / Unknowr	า			
	Adults (18 years	•					
	, ,		<u> </u>				
4. Do othe	r institutions sen	d specimens to	your laboratory	for testing? If so, pleas	e describe.		
F D		.1					
5. Does you	ur institution ser	id specimens to	other laboratori	es for testing? If so, ple	ease describe.		
Section II: Te	esting Procedure	<u>es</u>					
6. Which o	f the following m	nultiplex PCR res	piratory virus as	says are used in your l	ab? (circle all that appl	y)	
EraGen	GenMark	Seegene	Luminex	FilmArray			
Qiagen	Fastrack	Other (Pleas	o List			١	
Qiageii	Tastrack	Other (Fleas	C LIST			/	
data sources, g	athering and				onse, including the time for	-	
	e data needed, and formation unless	completing and rev	lewing the collectio	n or information. An agency	may not conduct or sponse	or, and a person is not	required to respond t

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7. For each of the following 12 viral agents, please indicate the methods of detection your lab uses (Y/N).

						Nucl	eic Acid Aı	mplification Test / PCR					
		mmunoass	ay	_	"Hom	e Brew"*	<u> </u>	Culture					
	Rapid	IFA	EIA		CDC	Other	CDC Kit	Commercial**		Conventional	Shell Vial	No	one
(Example)	N	Υ	N		N	N	Υ	6/2013-Switched from Luminex resp panel to Genmark resp panel		N	Υ	N	
Influenza							<u> </u>	Commany resp parter					
RSV													
Rotavirus													
Adenovirus													
hMPV													
Rhinovirus													
Enterovirus													
PIV1													
PIV2													
PIV3													
PIV4													
Coronavirus													

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*Produced In-house**Please indicate product name if commercial assay is used. Also indicate if a commercial assay was recently introduced or recently discontinued.	
Notes/explanations regarding testing practices:	
8. Do you use a standard protocol or physician order in selecting testing methods? If standard protocol, please describe.	
9. Do your testing practices for RSV and influenza change between the on- and off-seasons? Please describe.	
 10. Do you test for RSV all year? Yes / No / Unknown a. If not, at what time of year do you generally start testing for RSV? b. At what time of year do you generally stop testing for RSV? c. At which factors influence this decision? 	
11. Do you test for influenza all year? Yes / No / Unknown 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 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 reducing this burden to CDC, Do not send the completed form to this address. OMB No. 0920-0004	to a

----National Disease Surveillance Program – II. Disease Summaries, OMB 0920-0004----National Respiratory and Enteric Virus Surveillance System (NREVSS)

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- a. If not, at what time of year do you generally start testing for influenza?
- b. At what time of year do you generally stop testing for influenza?
- c. At which factors influence this decision?
- 12. Have your laboratory's routine testing practices changed in the past 5 years? If so, in which ways? What factors have influenced these changes?

Section III: Data Recording and Reporting Practices

- 13. How do you keep records of test results (e.g. MS Excel, Access, paper ledger)?
- 14. Do you have any issues or suggestions regarding reporting data to NREVSS?

15. Additional comments or suggestions:

Supplemental Questions

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Would you change anything about the NREVSS data entry or submission process?
How do you feel about NREVSS security (your lab ID, password, etc.)? Any problems to report?
How often do you view the public NREVSS website? Any suggestions regarding the content presented?
What is the best way to reach your lab? (Phone, Email, Fax, Mail, other)
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