



Pneumococcal Vaccine Schedules Study

Checklist of procedures to monitor in the Basse laboratory

Date of visit:/ Name of attendant/s:						
Evaluation Crit	eria A- EQI	ITDMF	NT/D	FVICE	<u> </u>	
	Maintenance/Technical control/Functionality		POOR	LVICE	FAIR	GOOD
Thermometer(s)	Calibration	YES	NO	Other	-,	
	Comment:					
Refrigerator(s)	Maintenance/Technical control/Functionality		D POOR		FAIR	GOOD
	Calibration	YES	NO NO	Other	,	
	Temperature control sheet(s)/recording available and been recorded appropriately Comment:			□ NO		
	Measurements - if temperature out of range YES NO					□ NO
	Comment:					





	Maintenance/Technical control/Functionality	POOR	FAIR	GOOD	
	Calibration	YES NO Other	r,		
-70/80°C Freezer(s)	Temperature control she available and been recor		YES	□ NO	
	Comment:				
	Measurements - if temper	erature out of range	e YES	□ NO	
	Comment:				
	Maintenance/Technical control/Functionality	POOR	FAIR	GOOD	
	Calibration	YES NO Other	r,		
	CO2 environment	Y	ES NO)	
Incubator(s)	Comment				
	Temperature control she available and been recor		YES	□ NO	
	Comment:				
	Measurements - if temperature out of range YES NO				
	Comment:				





	Maintenance/Technical control/Functionality	POOR	FAIR	GOOD		
	Calibration	YES NO Other	r,			
Blood culture	Comment:					
device BACTEC	Temperature control she available and been record Comment:		YES	□ NO		
	Measurements - if out of	f range	YES	□ NO		
	Comment:					
	Maintenance/Technical control/Functionality	POOR	FAIR	GOOD		
Autoclave(s)	Calibration	YES NO Other	r,			
	Comment:					
Microscope(s)	Maintenance/Technical control/Functionality	POOR	FAIR	GOOD		
	Calibration	YES O Other	r,			
	Comment:					
Safety cabinet(s)	Maintenance/Technical control/Functionality	POOR	FAIR	GOOD		





	Calibration	YES NO Other,	,			
	Comment:					
	Maintenance/Technical control/Functionality	POOR	FAIR	GOOD		
Pipette(s)	Calibration	YES NO Other	· ,			
	Comment:					
	Maintenance/Technical control/Functionality	POOR	FAIR	GOOD		
Centrifuge(s)	Calibration	YES NO Other,	/			
	Comment					
	Cleanliness	POOR	FAIR	GOOD		
Working Environment	Organization	POOR	FAIR	GOOD		
	Comment:					
B - REAGENTS/KITS/MEDIA/REFERENCE STRAINS						
	Storage conditions (as p		YES	∏ NO		
Reagents/ Test kits	Expiration date (as per r	•	YES	□ NO		
I COL NICO	Constant supply availabl	e	YES	NO NO		





	Comment:		
	QC documentation (as per SSP)	YES	□ NO
	Expiration date (as per manufacturer)	YES	☐ NO
Pneumococcal antisera	Constant supply available	YES	☐ NO
41100014	Comment:		
	Preparation date recorded	YES	□ NO
	Expiration date recorded	YES	☐ NO
Modia/	Preparation process: appropriate conditions	YES	☐ NO
Media/ Media	Control of functionality/growth	YES	☐ NO
preparation	Control of sterility	YES	☐ NO
	Comment:		
	Reference strains used/available	YES	☐ NO
	Comment:		
	Strains stored in appropriate medium (e.g. microbank, glycerol medium)	YES	☐ NO
Reference strains	Comment:		
	Storage temperature -20°C	YES	□ NO
	Strains handled for sub-cultivation according to SOP	YES	☐ NO
	Comment:		





C - HANDLING OF SPECIMEN						
	Transportation of specimen as per protocol/SOPs (i.e. temperature/time)	YES	□ NO			
	Specimen recorded in EMR first on reception	YES	□ NO			
Specimen transportation	Specimen recorded in EMR and log book	YES	□ NO			
transfer	Internal laboratory number given to specimen (if applicable)	YES	□ NO			
	Comment:					
	Culturing of NPS STGG as per protocol/SOPs	YES	NO			
	Identification of cultured microorganisms as per protocol/SOPs	YES	□ NO			
NPS processin		YES	□ NO			
	Comment:	1	1			
	Culture and identification according to protocol	YES	□ NO			
	Contamination rate of blood cultures calculated (ideally up to 5%)	YES	□ NO			
	Measures taken if high contamination rate	YES	□ NO			
	Comment					
Blood culture documentation interpretation	n/					
interpretation	Positivity rate of blood cultures (ideally 5%)	YES	□ NO			
	Measures taken if low positivity rate	YES	□ NO			
	Comment					
D – QUALITY CONTROL						
	Control of testing done regularly using the reference strains	YES	☐ NO			
Internal quality	Control of identification reactions according to SOPs	YES	☐ NO			
control	Comment:					





	Does the laboratory participate in an external quality control program for microbiology?	YES		□ NO		
External	If "Yes", state the external quality control program and result of most recent evaluation. Check documentation is filed in laboratory.					
quality control	Quarterly external QC of sweep serotyping done?	YES NO) [NOT DUE		
	If yes, result – agreement for x/y STGG specimens	, ,				
	Comment:					
	E – NPS & BC SAMPLE MO					
	Pick two sample ID from reception logb and follow through:	OOK				
NPS	Is sample recorded in lab EMR?	YI	ES	□ NO		
INFS	If sample is isolate is it stored?	YI	ES	☐ NO		
	Comment:					
	Pick two sample ID from reception logb and follow through:	ook				
D.C.	Is sample recorded in lab EMR?	YI	ES	☐ NO		
BC	If sample is isolate, is it stored?	YI	ES	☐ NO		
	Comment:					
	F - STANDARD OPERATING PROC	EDURES/TRAIN	ING			
	PVS SOPs available & used		YES	NO NO		
	Monthly/Quarterly staff proficiency doc	umented				
Standard Operating Procedures	Comment	Comment				
(SOPs)	Additional SOPs of collaborators availal used (If applicable)	ble &	YES	□ NO		
	Comment					
	Reference literature available & used		YES	NO		





	Comment		
Reference	Training of staff at site	YES	□ NO
literature/ Training	Comment/Trained on:		
	Comment/Frequency of training:		

Evaluation Outcome/comments: -

Recommendations/Action points:

Name of person conducting supervision: