EMPLOYEE NAME (Printed):

ZUCKERBERG SAN FRANCISCO GENERAL HOSPITAL & TRAUMA CENTER Initial ORIENTATION & TRAINING

PROCEDURE: Chembio Rapid HIV 1/2 STAT PAK

This form is to be used for trainer-supervised initial orientation and training. The newly trained employee must demonstrate competency at the time of initial orientation & training. All other employees must demonstrate competency on an annual basis.

TRAINER(s) NAME (Printed)				
LICENSE NUMBER:	LOCA	ATION:		
I (initials)HAVE READ AND UNDERSTAN POLICY & PROCEDURE.	ID THE	E CHEMBIO RAPID I	HIV 1/2 STAT PAK	
STEF	<u>P 1</u>			
HAVE EMPLOYEE READ THE CHEMBIO RAPID HIV1/2 STAT POLICY & PROCEDURE	PAK	Date:	Trainer's Initials:	
STEP 2				
ORIENT TRAINEE TO POINT OF CARE TESTING ENVIRONME	ENT	Date:	Trainer's Initials:	
Location of Policy and Procedure, SFGH-POCT website for reference				
 Location of supplies and Reagents, and 	d wher	e to dispose ofused ma	terials	
STEF	P <u>3</u>			
INTRODUCTION OF THE CHEMBIO RAPID HIV 1/2 STAT PA	K	Date:	Trainer's Initials:	
Purpose and Principle				
Qualified Testing Personnel (Requirements: Initial Training and Annual Competency)				
 Equipment (Pouched test devices, sample loops, running buffer, Chembio HIV 1 & 2 reactive controls, nonreactive control, timer. For fingerstick: alcohol wipes, lancet, sterile gauze) 				
STEF	P 4			
SPECIMEN AND REAGENTS		Date:	Trainer's Initials:	
Specimen collection and stability (fing	erstick	sample must analyze ir	nmediately)	
Storage and Stability of Chembio Brand HIV 1/2 test devices (room temp)				
Storage and Stability of Chembio HIV 1/2 Reactive and Nonreactive Controls (refrigerated)				
Daily Temperature Log				

	STEP 5			
QUALITY CONTROL		Date:	Trainer's Initials:	
pos con ligh	ernal Quality Controls (included in the testitive internal quality control. A clear back trol. If the test is working properly, the bt pink. Ernal Quality Controls: HIV 1 REACTIVE, H	ground is an internal nackground in the result	egative background t area should be white to	
	trols are run WEEKLY, and on all new lot	•	•	
	STEP 6			
PATIENT TESTING PROCE	EDURE	Date:	Trainer's Initials:	
 Gather equipment needed. Cover workspace with a clean, disposable absorbent cover. Put on disposable gloves. 				
• Check expiration date of the test device. Label device with patient's medical record number.				
• Obt	tain a fingerstick blood sample: wipe awa	y 1st drop, then sampl	e with loop from 2nd drop.	
• Wit	th sample loop down, touch it to the pad	in the sample (S) well,	blood will flow onto pad.	
	d the Running Buffer bottle vertically (nong drops into (S) well. Three full drops ar		S) well and squeeze 3 free	
• Rea	d the test result after 15 minutes, do no	t read after 20 minute	S.	
	STEP 7			
INTERPRETATION OF RE	SULTS	Date:	Trainer's Initials:	
 <u>REACTIVE</u>: two pink/purple lines appear. One line should be in the control region (C) and another line (which can be faint) should be in the test region (T). 				
	GATIVE: one pink/purple line appears in the lears in the test region (T).	ne control region (C). N	o apparent line	
	• <u>INAVLID</u> : Control line fails to appear, is other than pink/purple color, or appears misplaced. First repeat with a new test device. If Invalid again, repeat with new unopened bottles of QC.			
	STEP 8			
LIMITATIONS OF METHO	OD	Date:	Trainer's Initials:	
CONFIRMATORY TESTIN	G	Date:	Trainer's Initials:	
MATERIAL SAFETY DATA	A SHEETS (MSDS)	Date:	Trainer's Initials:	

EMPLOYEE SIGNATURE:	DATE TRAINING COMPLETED:	
TRAINER(s) SIGNATURE:	DATE TRAINING COMPLETED:	

Upon Completion please scan and email to Point of Care Testing staff: francis.corteza@sfdph.org and cassiusmicho.santiag@sfdph.org

OR fax to POCT office: 415.206.3451

Unit Manager is encouraged to retain a copy in employee or unit file.