

**LifeCare Medical Center
Main Laboratory
Laboratory Aide Competency Assessment**

Name: _____ Year: _____ Assessment: () Initial () 6 months () Annual () Follow- up

Employee will be assessed as (**C**) – Competent (**NC**) – Non-Competent. Evaluation will be discussed with employee to establish an action plan if needed.

Department: Outpatient/ Serology/Phlebotomy	DO 1	TR 2	ID 3	PM 4	TP 5	PS 6	C	NC **	Date
Test / Test System									
Nova BioMedical Stat Strip Glucometer (waived)									
Hemocue Hb801 (waived)									
Roche Coaguchek XS Plus (waived)									
Siemens Uristix (waived)									
Abbott ID Now (Group A Strep, COVID- 19) (waived)									
BD Veritor System (Rapid Influenza A/B) (waived)									
Sekisui Diagnostics OSOM Card Pregnancy Test (waived)									
Cardinal Health Mono II Rapid Test (waived)									

DO=Direct observation of routine patient performance; **TR**=Monitoring and recording of test results; **ID**=Review of intermediate test results/worksheets/quality control/preventive maintenance records; **PM**=Direct observation of performance of instrument maintenance and function checks; **TP**=Assessment of test performance/proficiency/blind testing samples/previously analyzed specimens; **PS**=Assessment of problem solving skills

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Non-Technical	DO 1	TR 2	ID 3	PM 4	TP 5	PS 6	C	NC **	Date
Process									
Venipuncture									
Capillary Puncture									
Blood Alcohol Collection									
Blood Culture Collection									
Isolation									

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***= Describe component requiring further action (attach corrective action documentation and use a new checklist for re-assessment)*

Laboratory Aide : I feel competent in the tests/test systems/tasks evaluated above : **YES** **NO**

If **NO**, describe test/test system/task in which you need additional training:

Director/Lead: I have reviewed this employee's competency in the above functions and determined that the employee is competent to perform the tests measured above.

Employee's Full signature

Director/Lead's Full Signature**

Date

*****=Director/Lead must be a qualified individual under CLIA Subpart M std. 493.1411 for moderate complexity and std. 493.1419 for high complexity tests.***