



MicroPhase Clinical Laboratory

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 256-882-7275 T, 256-882-3853 F

SPECIMEN COLLECTION FORM

PATIENT DEMOGRAPHIC SHEET, COPY OF LICENSE AND INSURANCE CARD CAN BE ATTACHED INSTEAD OF COMPLETING INFORMATION BELOW

Patient Last Name	Patient First Name	Date of Birth	Sex
Address			
City		State	Zip

INSURANCE

Primary (Check One): <input type="checkbox"/> BCBS <input type="checkbox"/> MEDICARE <input type="checkbox"/> MEDICAID <input type="checkbox"/> UHC <input type="checkbox"/> Humana <input type="checkbox"/> Tricare <input type="checkbox"/> Other	ID#	Date Collected	Time
Secondary (Check One): <input type="checkbox"/> BCBS <input type="checkbox"/> MEDICARE <input type="checkbox"/> MEDICAID <input type="checkbox"/> UHC <input type="checkbox"/> Humana <input type="checkbox"/> Tricare <input type="checkbox"/> Other	ID#		
Referring Physician			
Source of Specimen			
Diagnosis			
PLEASE CALL 256-882-7275 OR TEXT 256-509-6543 FOR CULTURE PICKUP			

TESTS

<input type="checkbox"/>	ROUTINE BACTERIAL CULTURE - AEROBIC AND ANAEROBIC
<input type="checkbox"/>	URINE CULTURE
<input type="checkbox"/>	THROAT CULTURE
<input type="checkbox"/>	FUNGUS CULTURE AND SMEAR
<input type="checkbox"/>	AFB CULTURE
<input type="checkbox"/>	RT-PCR — SARS-COV-2 (SAME DAY RESULTS IF SWABBED AND ONSITE BY 1:30) \$150 TRAVEL
<input type="checkbox"/>	SARS-COV-2 ANIGEN (RAPID) RESULTS WITHIN THE HOUR
<input type="checkbox"/>	SARS-COV-2 ANTIBODY - IGM AND IGG RESULTS WITHIN THE HOUR
<input type="checkbox"/>	SARS-COV-2, FLU A, FLU B, RSV, RT-PCR COMBO TEST (SINGLE SWAB) RESULTS WITHIN THE HOUR
<input type="checkbox"/>	LYME DISEASE BLOOD 30 MICROLITERS, SERUM OR PLASMA
<input type="checkbox"/>	STREP COMPLETE Group A/C/G/Molecular Assay
<input type="checkbox"/>	OTHER
<input type="checkbox"/>	FLU A and B Molecular Assay
<input type="checkbox"/>	RSV / hMPV Molecular Assay

The SARS-CoV-2 has been authorized by the FDA under an Emergency Use Authorization (EUA) for the detection and/or diagnosis of the SARS-CoV-2 Virus and is only authorized for the duration of the declaration that circumstances exist justifying the authorization of emergency use of in vitro diagnostics for detection and / or diagnosis of Covid-19 under section 564(B)(1) of the Federal Food, Drug, and Cosmetic Act, 21 USC 360 BBB-3(B)(1) unless the authorization is terminated or revoked sooner. Tests performed at MicroPhase Clinical Laboratory CLIA #01D0923800

Signature _____ Fax Results to _____