

A Subsidiary of Laboratory Sciences of Arizona

2021 Laboratory & Testing Updates

(January 2021 to date)

If you would like to receive Client Grams via email, please contact your Account Manager or email us at <u>ClientGram@SonoraQuest.com</u>.

You may also visit <u>sonoraquest.com/test-directory</u> to use our Searchable Test Directory for the most accurate information.



UPDATE: Temporary Patient Service Center Closures and Appointment Only Conversions

Sonora Quest Laboratories is committed to ensuring the safety of our guests and communities along with our valued team members. As we manage through this pandemic, we have made a difficult decision to continue the **temporary** closures of the following Patient Service Centers until Monday, February 1, 2021.

City	Address
Glendale	3800 W. Happy Valley Rd. (Inside Safeway), Glendale, AZ 85310
Phoenix	4524 N. Maryvale Pkwy., Suite 120, Phoenix, AZ 85031
Phoenix	550 E. Bell Rd. (Inside Safeway), Phoenix, AZ 85022

Additionally, we have made the decision to **temporarily** close the following Patient Service Center beginning Monday, January 11, which will remain closed until further notice. Patients with appointments at this location will be contacted and rescheduled at alternative locations.

City	Address
Scottsdale	7281 E. Earll Drive, Suite #2, Scottsdale, AZ 85851

The following Patient Service Centers have been **temporarily** converted to appointment only as of Thursday, December 31, and will remain so until further notice.

City	Address	
Goodyear	9890 S. Estrella Pkwy., Goodyear, AZ 85338	
Fountain Hills	13620 N. Saguaro Blvd., #150, Fountain Hills, AZ 85268 (appointment only after 1 p.m.)	
Peoria	7757 W. Deer Valley Rd. Suite 265, Peoria, AZ 85382	
Sun City	10503 W. Thunderbird Blvd. Suite 105, Sun City, AZ 85351	
Surprise	15331 W. Bell Rd., Suite 110, Surprise, AZ 85374	

Patients can visit SonoraQuest.com for a complete listing of our Patient Service Centers and to schedule appointments.

UPDATE: Sonora Quest Patient Service Center Carside COVID-19 Collections

Please see the updated locations below that are providing carside collection for symptomatic/recently exposed patients. Carside testing is available **by appointment only**. Patients are encouraged to visit SonoraQuest.com/OrderCovid for up-to-date locations offering carside collection.

City	Address	Hours
Bullhead City	3003 Hwy 95., Suite H81	8 a.m. – 3:30 p.m.
	Bullhead City, AZ 86442	
Casa Grande	1860 E. Salk Dr., Suite A1	11 a.m. – 3:30 p.m.
	Casa Grande, AZ 85122	
Flagstaff	1100 N. San Francisco St., Suite C Flagstaff, AZ 86001	11 a.m. – 3:30 p.m.
Appointments Available by Jan 11:	13620 N. Saguaro Blvd., Suite 150	1:30 p.m. – 3:30 p.m.
Fountain Hills	Fountain Hills, AZ 85268	· ·
Glendale	6320 W. Union Hills Dr., Suite 160	Noon – 3:30 p.m.
	Glendale, AZ 85308	
Glendale	9980 W Glendale Ave., Suite 120	11 a.m. – 3 p.m.
	Glendale, AZ 85307	
Kingman	2505 Hualapai Mountain Rd., Suite A	1 p.m. – 4 p.m.
	Kingman, AZ 86401	
Maricopa	21300 N. John Wayne Pkwy., Suite 106	1 p.m. – 3:30 p.m.
	Maricopa, AZ 85139	
Mesa	6344 E. Broadway Rd., Suite 118	Noon – 5 p.m.
	Mesa, AZ 85206	
Phoenix	2640 W. Baseline Rd., Suite 115	Noon – 3:30 p.m.
	Phoenix, AZ 85041	
Show Low	2450 Show Low Lake Rd., Suite 3B	8 a.m. – 3:45 p.m.
	Show Low, AZ 85901	
Sierra Vista	1150 S. Highway 92, Suite E	1 p.m. – 3:30 p.m.
	Sierra Vista, AZ 85635	
Sun Lakes	10450 E. Riggs Rd., Suite 109	11:30 a.m. – 2: 30 p.m.
	Chandler, AZ 85248	
Surprise	13856 W. Waddell Rd., Suite 107	Noon – 3:30 p.m.
	Surprise, AZ 85379	
Tempe	1275 W. Washington St., Suite 109	1 p.m. – 3:30 p.m.
	Tempe, AZ 85281	
Tucson	630 N. Alvernon Way, Suite 200	1 p.m. – 3:30 p.m.
	Tucson, AZ 85711	



ANNOUNCEMENT: Sonora Quest Laboratories COVID-19 Webinar

Beginning Friday, January 8, we invite you to join Dr. Brian Koeneman, scientific medical director, molecular diagnostics, and Dr. Brian Mochon, system scientific medical director – infectious diseases division and clinical research program, as they navigate the pros and cons of the different types of COVID-19 testing in the market and their recommended approaches.

View the 17-minute recorded webinar now or at a time that is convenient for you by visiting SonoraQuest.com/COVIDWebinar.

ANNOUNCEMENT: eSwab - White-Cap Routine Culture Swabs Backorder

Due to manufacturer backorder, supply item #25784 - eSwab – White-cap Routine Culture Swab will be temporarily replaced with supply item #25785 - eSwab – Blue-cap Mini-Tip Culture Swab. Blue-cap eSwabs can be used as substitute collection devices for testing in which white-cap eSwabs are typically submitted.

We anticipate this to be a temporary substitution until white-cap eSwabs become available.

Supply orders can be made through our Provider Portal at SonoraQuest.com, our Quanum™ system, or by faxing a client supply requisition to our warehouse. For updated client supply requisitions, please call 602.685.5141.

For questions regarding supplies please contact our warehouse at the following numbers:

Phoenix: Phone (602) 685.5264; Fax (602) 685.5402

Toll-free (800) 766.6721, ext. 5264

Tucson: Phone (520) 784.8004; Fax (520) 296.5607

Toll-free (800) 266.8101



IMPORTANT COVID-19 TESTING REMINDERS: SARS-CoV-2 RNA (COVID-19), Qualitative, NAAT – Test 907080

SONORA QUEST LABORATORIES COVID-19 WEBINAR

Beginning Friday, January 8, we invite you to join Dr. Brian Koeneman, scientific medical director, molecular diagnostics, and Dr. Brian Mochon, system scientific medical director – infectious diseases division and clinical research program, as they navigate the pros and cons of the different types of COVID-19 testing in the market and their recommended approaches.

View the 17-minute recorded webinar now or at a time that is convenient for you by visiting SonoraQuest.com/COVIDWebinar.

OVERVIEW OF SERVICES

Sonora Quest currently offers gold standard molecular testing for diagnosing COVID-19 on nasopharyngeal, nasal, and oropharyngeal samples. Currently, >80% of tests are being reported within 1 day of sample collection and >95% are being reported within 2 days. Up to date turnaround time information is always available at SonoraQuest.com.

SUPPLIES

To request specimen transport media and swab supplies for SARS-CoV-2 RNA (COVID-19), Qualitative, NAAT (test code 907080), please order from the COVID-19 supplies listed below:

- COVID-19 Media with Nasopharyngeal Swab SQL Supply #44921
- COVID-19 Media with Anterior Nares (Nasal) / Oropharyngeal Swab SQL Supply #44947

Specimen collection guidelines and a complete list of acceptable collection devices are available at https://www.sonoraquest.com/covid-19-information-for-healthcare-providers/. Please note that comparable supply items may be substituted as needed.

Order supplies through our SonoraQuest.com Provider Portal, Quanum, or by faxing a Client Supply Request Form to 602.685.5402 in Phoenix (central and northern AZ), or 520.296.5607 in Tucson (southern AZ).

SAMPLE SUBMISSION

- To ensure timely processing of samples for testing:
 - The source MUST be written on the specimen container and included on the order.
 - Collection date and time MUST be included on the order. Accuracy of this information will be vital for reimbursement beginning January 1, 2021.
 - All patient demographics MUST be included on the order to ensure patients are able to access their results in the SonoraQuest.com Patient Results Portal.
 - If you do not have regularly scheduled SQL courier pick-ups, or if samples are collected on weekends or holidays, please contact our Logistics Department at 602.685.5052 or 520.886.8101 as soon as possible after sample collection.
- Orders for COVID-19 diagnostic testing MUST be submitted on a separate requisition and packaged separate from other tests.

SAMPLE COLLECTION

- Sonora Quest's in-office phlebotomists do not currently collect respiratory specimens, including those from patients suspected of having COVID-19.
- Select Sonora Quest Patient Service Centers now offer carside collection for active infection testing and we are working to make this service available at additional locations. Please see Client Gram vol. 1 sent on Thursday, January 7.
- We have partnered with select Walmart locations in Phoenix, Tucson, and Yuma for drive-thru observed collections.
 Note that you MUST submit an electronic order for testing to Sonora Quest for this sample collection option.
- Patients can schedule Sonora Quest Patient Service Center carside and Walmart drive-thru appointments at https://www.sonoraquest.com/appointments/.

PATIENT RESULTS PORTAL

In order to ensure patients get their COVID-19 test results as soon as they are available, we encourage you to let your patients know they can access their results through our Patient Results Portal by visiting https://www.sonoraquest.com/results. Registered patients can sign up for email and text notifications for new results.

BILLING

To enable Sonora Quest Laboratories to bill uninsured patient COVID-19 claims to the Department of Health and Human Services (HHS) Uninsured COVID-19 program administrator, Optum, we must receive either the Social Security, driver's license or state ID number on their COVID-19 order requisitions for all uninsured patients. If a patient does not wish to share this information, please indicate the patient's refusal to provide this information on the order form.

Things to know:

- Payor code 3588 "Uninsured COVID-19" has been added to Quanum for these patients.
- The Department of Health and Human Services (HHS) Uninsured COVID-19 program administrator, Optum, is requiring that Sonora Quest Laboratories collect this information.
- Sonora Quest Laboratories secures all patient information in accordance with regulatory requirements (e.g., HIPAA) and industry best practice.
- Patient Social Security, driver's license or state ID number will be transmitted only to HHS's COVID-19 Uninsured Program claims administrator, Optum, and solely for the purpose of billing uninsured laboratory claims.
- Sonora Quest Laboratories will seek reimbursement for uninsured laboratory claims from HHS by billing their plan
 administrator, Optum. If for any reason HHS does not pay as anticipated, Sonora Quest will not pursue the collection of
 any unpaid COVID-19 claim balances from patients.

Please visit https://www.sonoraquest.com/latest-covid-19-news/ for updates and additional resources and information.



ANNOUNCEMENT: Temporary Patient Service Center Closure

Sonora Quest Laboratories is committed to ensuring the safety of our guests and communities along with our valued team members. As we manage through this pandemic, we have made a difficult decision to **temporarily** close the following Patient Service Center until Monday, January 18, 2021.

City	Address
Sedona	2300 W. Highway 89A. (Inside Safeway), Sedona, AZ 86336

ASSAY CHANGES:

Test 102405	Old Name: Cryoglobulin Evaluation New Name: Cryoglobulin Screen
Effective:	1/18/2021

Test 1021	Albumin
Effective:	1/19/2021
Reference Range:	0-14 Days: 3.4-4.7 g/dL
_	15-364 Days: 3.0-5.3 g/dL
	≥1 Years: 3.8-5.1 g/dL

Test 1024	Chloride
Effective:	1/25/2021
Reference Range:	0-364 Days: 98-110 mmol/L
	1-59 Years: 98-108 mmol/L
	≥60 Years: 95-109 mmol/L

Test 2041	Lipase
Effective:	1/26/2021
Reference Range:	0-30 Days: 6-55 U/L
	31-364 Days: ≤29 U/L
	1-17 Years: ≤32 U/L
	≥18 Years: 13-60 U/L

Test 1020	Protein, Total
Effective:	1/18/2021
Reference Range:	0-14 Days: 5.5-8.3 g/dL
	15-364 Days: 4.6-7.2 g/dL
	1-3 Years: 5.9-7.7 g/dL
	4-49 Years: 6.3-8.0 g/dL
	≥50 Years: 6.0-7.7 g/dL

Test 1022	Urea Nitrogen (BUN)			
Effective:	1/25/2021			
Reference Range:		Male:	Female:	
	0-14 Days:	3-24	3-24 mg/dL	
	15-364 Days:	3-18	3-18 mg/dL	
	1-49 Years:	7-21	6-19 mg/dL	
	50-69 Years:	7-28	7-28 mg/dL	
	≥70 Years:	8-36	8-36 mg/dL	

Test 1025	Carbon Dioxide (CO2)	
Effective:	1/26/2021	
Reference Range:	0-364 Days: 13-29 mmol/L	
	≥1 Years: 20-31 mmol/L	

Test 1026	Sodium
Effective:	1/19/2021
Reference Range:	135-145 mmol/L

Test 1019	Bilirubin, Total
Effective:	1/18/2021
Reference Range:	31-364 Days: ≤0.6 mg/dL 1-3 Years: ≤0.6 mg/dL 4-12 Years: ≤1.0 mg/dL ≥13 Years: ≤1.3 mg/dL
Comment:	For testing on ages 0-30 days, please use test code 701487 – Bilirubin, Neonatal.

Test 701487	Bilirubin, Neonatal
Effective:	1/18/2021
Reference Range:	0-14 Days: ≤14.6 mg/dL
	15-30 Days: ≤0.6 mg/dL



ANNOUNCEMENT: CPT Coding Effective January 1, 2021

The American Medical Association (AMA) has made Current Procedural Terminology (CPT) code changes to the 2021 edition of the CPT coding manual. In addition, the CMS has also made changes to Healthcare Common Procedure Coding System (HCPCS) codes.

Sonora Quest Laboratories has implemented these changes effective January 1, 2021. The changes for 2021 affect the way we bill some of our tests. Please note these changes will not impact our service offerings or how you order them, but simply how we will bill third party payers.

The summary below outlines the 2021 CPT code changes that affect Sonora Quest Laboratories published test offerings. These tests may also be included in panels or profiles.

Test Code	Test Name	2020 CPT*	2021 CPT*
15039	Estradiol, Free	82670 (x2)	82681, 82670
10330	Felbamate	80299	80167
9058	Flecainide	80299	80181
904701	JAK2, Exon 12 Mutation Analysis	81403	81279
906964	JAK V617F Cascading to CALR, JAK2 EXON 12, MPL and	81479, 81403	81479, 81339,
	CSF3R	(x2), 81219,	81279, 81219,
		81270	81270
2250	Methotrexate Level	80299	80204
904702	MPL Mutation Analysis	81403	81339

ASSAY CHANGES:

Test 906366	BRCAvantage (™) Ashkenazi Jewish Screen
Effective:	1/25/2021
Specimen:	5 mL room temperature whole blood in a lavender-top (EDTA) tube (2 mL min).
Method:	DNA Bait Capture; Long Range Polymerase Chain Reaction; Next Generation Sequencing
Setup:	Days: Tuesday, Thursday, and Saturday
Reports:	15-22 days from completed prior authorization

Test 906474	BRCAvantage(™) Ashkenazi Jewish Screen w Rflx Comprehensive	
Effective:	1/25/2021	
Specimen:	5 mL room temperature whole blood in a lavender-top (EDTA) tube (2 mL min).	
Method:	DNA Bait Capture; Long	Range Polymerase Chain Reaction; Next Generation Sequencing
Setup:	Days: Tuesday, Thursda	y, and Saturday
Reports:	15-22 days from comple	ted prior authorization
Interface Mapping:	Result Code	Result Code
	10906474	Result
	Possible reflexes:	
	Result Code	Result Code
	19063691	Gene
	29063691	Variant
	39063691	Classification
	27906541	Gene 2
	28906541	Variant 2
	29906541	Classification 2
	19036393	Gene 3
	29036393	Variant 3
	39036393	Classification 3
	19063694	Gene 4
	29063694	Variant 4
	39063694	Classification 4
	19063695	Gene 5
	29063695	Variant 5
	39063695	Classification 5
	70906474	VUS(s)
	71906474	Gene List
	19063697	Clinical Interpretation
	29063697	Variant Information
	39906541	Reviewer
	29063698	Resources
	39063698	Methods and Limitations
	42906541	Additional Information
	72906474	AJ Screen
	73906474	Reflex Performed

Test 906367	BRCA 1 and BRCA 2 Deletion and Duplication		
Effective:	1/25/2021		
Specimen:	5 mL room temperature whole blood in a lavender-top (EDTA) tube (2 mL min).		
Method:	DNA Bait Capture; Long Range Polymerase Chain Reaction; Next Generation Sequencing		
Setup:	Days: Tuesday, Thursday, and Saturday		
Reports:	15-22 days from completed prior authorization		

Test 906369	BRCA Panel (BRCA1, BRCA2)			
Effective:	1/25/2021			
Specimen:	5 mL room temperature whole blood in a lavender-top (EDTA) tube (2 mL min).			
Method:		DNA Bait Capture; Long Range Polymerase Chain Reaction; Next Generation Sequencing		
Setup:	Days: Tuesday, Thursd		·	
Reports:	15-22 days from comple	eted prior authorization		
Interface Mapping:	Result Code	Result Name		
	10906369	Result		
	19063691	Gene		
	29063691	Variant		
	39063691	Classification		
	19063692	Gene 2		
	29063692	Variant 2		
	39063692	Classification 2		
	19036393	Gene 3		
	29036393	Variant 3		
	39036393	Classification 3		
	19063694	Gene 4		
	29063694	9063694 Variant 4		
	39063694	Classification 4		
	19063695	Gene 5		
	29063695	Variant 5		
	39063695	Classification 5		
	19063696	VUS(s)		
	29063696	Gene List		
	19063697	Clinical Interpretation		
	29063697	Variant Information		
	19063698	Reviewer		
	29063698	Resources		
	39063698	Methods and Limitations		
	24906369	Additional Information		
	Ask at order entry question			
	Result Code	Result Name	Response Options	
	11906369	Known Family BRCA Mutation:	Free Text	
	12906369	Spec. Familial Mutation:	Free Text	
	13906369	Spec. Familial Del/Dup:	Free Text	
	99811887	Pre-Auth?	Free Text	
	99911887	Pre-Authorization Code	Free Text	

Test 906541	Lynch Syndrome Panel		
Effective:	1/25/2021		
Specimen:	5 mL room temperatu	5 mL room temperature whole blood in a lavender-top (EDTA) tube (2 mL min).	
Method:	DNA Bait Capture; Lo	ong Range Polymerase Chain Reaction; Next Generation Sequencing	
Setup:	Days: Tuesday, Thurs	sday, and Saturday	
Reports:	15-22 days from com	pleted prior authorization	
Interface Mapping:	Result Code	Result Code	
	50906541	Lynch Syndrome Panel	
	Possible reflexes:		
	Result Code	Result Code	
	22906541	Gene 1	
	23906541	Variant 1	
	24906541	Classification 1	
	27906541	Gene 2	
	28906541	Variant 2	
	29906541	Classification 2	
	32906541	Gene 3	
	33906541	Variant 3	
	34906541	Classification 3	
	19063694	Gene 4	
	29063694	Variant 4	
	39063694	Classification 4	
	19063695	Gene 5	
	29063695	Variant 5	
	39063695	Classification 5	
	70906474	VUS(s)	
	71906474	Gene List	
	19063697	Clinical Interpretation	
	29063697	Variant Information	
	39906541	Reviewer	
	29063698	Resources	
	39063698	Methods and Limitations	
	42906541	Additional Information	



ANNOUNCEMENT: Scheduled Application Server Maintenance

On Sunday, January 24, 2021, Sonora Quest Laboratories will be performing application server maintenance. As a result, all computer systems will be unavailable beginning at 6:00 p.m. until approximately 11:00 p.m.

During this period, our Client Services Representatives will not have access to computerized information or patient results. All reporting will resume by normal processes once our systems are operational. Quanum users will have uninterrupted access to patient results that were completed prior to 6:00 p.m. Sunday. Quanum will be updated with results completed during the downtime once the server maintenance is completed.

STAT courier services and STAT testing will remain available during this time and will be reported manually by phone via our established down-time processes.

During the downtime, please contact us as follows:

- Client Services: 602.685.5050
- Logistics (for pick-ups including Mobile Diagnostic Services STAT phlebotomy requests): 602.685.5052



ANNOUNCEMENT: Testing Delays - Drug Screens of Abuse Meconium (DSAM)

The published turnaround time for test 907180 – Drug Screens of Abuse Meconium (DSAM) is 1-2 days. Due to instrumentation issues at the performing lab, Banner University Medical Center Phoenix (BUMCP), testing will be sent out to ARUP until further notice. This will add an additional 2 days to testing turnaround time. Additionally, if your facility receives electronic reporting, reports from ARUP will be transmitted as a hard copy report.

An update will be sent once the testing resumes at BUMCP.



UPDATE: Drug Screens of Abuse Meconium (DSAM) Testing Has Resumed

In Client Gram volume 7, it was announced that due to instrumentation issues at the performing lab, Banner University Medical Center Phoenix, test 907180 – Drug Screens of Abuse Meconium (DSAM) would be temporarily sent out to ARUP for testing and that clients would see slight delays in reporting. The instrumentation issues have been resolved and testing has resumed at the performing lab.



DISCONTINUED TEST:

Test 905601	Hepatitis C Virus RNA, Quant PCR w/Rflx to HCV Qual TMA
Effective:	2/10/2021
Comment:	Please note that this test contains discontinued Test 901153 - Hepatitis C Viral RNA, Qualitative, TMA and is also being discontinued. Please see below for the recommended alternative testing.

RECOMMENDED ALTERNATIVE:

Test 905541	Hepatitis C Virus RNA, Quantitative, Real-Time PCR	
Specimen:	2 mL frozen plasma	from a lavender-top (EDTA) tube (1 mL min).
Reference Ranges:	<15 Not Detected IU	J/mL
_	<1.18 Not Detected	Log IU/mL
Setup:	Days: Monday – Frid	day
Stability:	Ambient: Unacceptable	
-	Refrigerated: 6 Days	
	Frozen: 6 Weeks	
Reports:	2-4 Days	
Interface Mapping:	Result Code	Result Name
	10905541	Hepatitis C Virus RNA (IU/mL)
	11905541 Hepatitis C Virus RNA (Log IU/mL)	

ASSAY CHANGES:

Test 901873	StoneRisk Diagnostic Profile	
Effective:	2/22/2021	
Comment:	Please remove the below Interface Mapping code.	
Interface Mapping:	Result Code Result Name	
	99901837	StoneRisk Graphical Report

Test 901830	UroRisk® Diagnostic Profile	
Effective:	2/22/2021	
Comment:	Please remove the below Interface Mapping code.	
Interface Mapping:	Result Code Result Name	
	99901830	UroRisk Graphical Report

Test 2075	Protein Electropho	Protein Electrophoresis, Serum	
Effective:	2/22/2021		
Interface Mapping:	Result Code	Result Name	
	10001020	Protein, Total	
	10002426	Serum Proteins	
	10002583	A/G Ratio	
	10002427	Albumin	
	10002428	Alpha-1-Globulin	
	11302425	M-Spike (Alpha-1-Globulin)	
	11322425	M-Spike 2 (Alpha-1-Globulin)	
	11332425	M-Spike 3 (Alpha-1-Globulin)	
	10002429	Alpha-2-Globulin	
	11202425	M-spike (Alpha-2-Globulin)	
	11222425	M-Spike 2 (Alpha-2-Globulin)	
	12232425	M-Spike 3 (Alpha-2-Globulin)	
	10002430	Beta Globulin	
	11102425	M-spike (Beta Globulin)	
	15102425	M-Spike 2 (Beta Globulin)	
	14102425	M-Spike 3 (Beta Globulin)	
	10002425	Gamma Globulin	
	11002425	M-spike (Gamma Globulin)	
	81002425	M-Spike 2 (Gamma Globulin)	
	71002425	M-Spike 3 (Gamma Globulin)	
	13008469	PE Interpretation	

Test 709197	Protein Electropho	Protein Electrophoresis, Urine, 24 Hour	
Effective:	2/22/2021	2/22/2021	
Interface Mapping:	Result Code	Result Name	
	10001005	Volume (mL):	
	12005812	Duration (Hr):	
	10002533	Protein, Urine, 24 Hour	
	11002482	Protein, Urine, Timed	
	11021732	Urine Proteins, 24 Hour	
	10009652	Albumin	
	11021733	Globulin	
	19021734	% Albumin	
	19221733	M-spike (Alpha-1-Globulin)	
	19241733	M-Spike 2 (Alpha-1-Globulin)	
	19251733	M-Spike 3 (Alpha-1-Globulin)	
	19321733	M-spike (Alpha-2-Globulin)	
	19331733	M-Spike 2 (Alpha-2-Globulin)	
	19341733	M-Spike 3 (Alpha-2-Globulin)	
	19121733	M-spike (Beta Globulin)	
	19123733	M-Spike 2 (Beta Globulin)	
	19124733	M-Spike 3 (Beta Globulin)	
	19021733	M-spike (Gamma Globulin)	
	19422733	M-Spike 2 (Gamma Globulin)	
	19423733	M-Spike 3 (Gamma Globulin)	
	19008469	Interpretation	

Test 209198	Protein Electroph	noresis, Urine, Random
Effective:	2/22/2021	,
Interface Mapping:	Result Code	Result Name
	10002498	Creatinine, Urine, Random
	10002482	Protein, Urine, Random
	19002533	Protein, Urine, Normalized
	11021734	% Albumin
	11022371	Random Urine Proteins
	11022372	Albumin
	11022373	Globulin
	15008469	Interpretation
	19022373	M-spike (Gamma Globulin)
	19042373	M-Spike 2 (Gamma Globulin)
	19052373	M-Spike 3 (Gamma Globulin)
	19122373	M-spike (Beta Globulin)
	19133373	M-Spike 2 (Beta Globulin)
	19134373	M-Spike 3 (Beta Globulin)
	19222373	M-spike (Alpha-2-Globulin)
	19272373	M-Spike 2 (Alpha-2-Globulin)
	19282373	M-Spike 3 (Alpha-2-Globulin)
	19322373	M-Spike (Alpha-1-Globulin)
	19352373	M-Spike 2 (Alpha-1-Globulin)
	19362373	M-Spike 3 (Alpha-1-Globulin)



ANNOUNCEMENT: Testing Unavailable - Platelet Aggregation

Due to instrumentation issues at the performing lab, Banner University Medical Center Phoenix (BUMCP), testing for Platelet Aggregation (test code 710439) is currently unavailable. Since the specimen stability for this test is only 4 hours, testing should not be scheduled or collected. An update will be sent once the testing resumes at BUMCP, which is expected to be next week.

Please Note: This does not affect testing performed at Banner University Medical Center Tucson (BUMCT).

ANNOUNCEMENT: Scheduled Application Server Maintenance

On Sunday, February 28, 2021, Sonora Quest Laboratories will be performing application server maintenance. As a result, all computer systems will be unavailable beginning at 6:00 p.m. until approximately 11:00 p.m.

During this period, our Client Services Representatives will not have access to computerized information or patient results. All reporting will resume by normal processes once our systems are operational. Quanum users will have uninterrupted access to patient results that were completed prior to 6:00 p.m. on Sunday. Quanum will be updated with results completed during the downtime once the server maintenance is completed.

STAT courier services and STAT testing will remain available during this time and will be reported manually by phone via our established downtime processes.

During the downtime, please contact us as follows:

- Client Services: 602.685.5050
- Logistics (for pick-ups including Mobile Diagnostic Services STAT phlebotomy requests): 602.685.5052



DISCONTINUED TEST:

Test 708177	Culture, Yeast
Effective:	3/1/2021
Comment:	Please note that this test will be discontinued. Please see below for the recommended alternative testing.

RECOMMENDED ALTERNATIVE:

Test 907203	Culture, Yeast			
Effective:	3/1/2021			
Specimen:	Submit appropriate spe	ecimens in sterile container or swa	ab in transport media.	
Reference Ranges:	See Report			
Stability:	Room temperature: 24	Hours		
	Refrigerated: 72 Hours			
	Frozen: Unacceptable			
Method:	Conventional Yeast Cu	lture		
Setup:	Days: Monday – Sunda	ау		
Reports:	5 Days (growth depend	lent)		
CPT*:	87102			
Price:	Client: \$32.76 Patient: \$59.01			
Interface Mapping:	Result Code Result Name			
	10907203 Culture, Yeast			
	Ask at order entry questions			
	Result Code Result Name Result Options			
	99708177 Source: Free Text			
	99990000 Other: Free Text			
Comment:	Testing will be performed at our main laboratory in Phoenix. Antibiotic susceptibilities will be performed on			
	pathogens as necessar	ry at an additional charge. Additio	nal charge for organism identification.	

DISCONTINUED TEST:

Test 708086	Dermatophyte (Fungal) Screen
Effective:	3/1/2021
Comment:	Please note that this test will be discontinued. Please see below for the recommended alternative testing.

RECOMMENDED ALTERNATIVE:

Test 907206	Dermatophyte (Fungal) Screen			
Effective:	3/1/2021			
Specimen:	Cleanse skin with alcoh	nol before scraping. Remove hair	s with forceps, scrape skin or scalp scales, clip	
			apings, or hair in a leak-proof container, or the	
	sample can be submitted	ed on folded black paper inside a	n envelope or between two slides. Keep at room	
	temperature.			
Reference Ranges:	See Report			
Stability:	Room temperature: 24	Hours		
	Refrigerated: 7 Days			
	Frozen: Unacceptable			
Method:	Dermatophyte Screen	Culture		
Setup:	Days: Monday – Sunda	ау		
Reports:	3 Weeks (growth dependent)			
CPT*:	87101			
Price:	Client: \$32.76 Patient: \$59.01			
Interface Mapping:	Result Code	Result Name		
	10907206	Dermatophyte (Fungal) Screen		
	Ask at order entry questions			
	Result Code Result Name Result Options			
	99708086	Source: Free Text		
	99990000	9990000 Other: Free Text		
Comment:	Testing will be performed at our main laboratory in Phoenix. Antibiotic susceptibilities will be performed on			
	pathogens as necessary at an additional charge. Additional charge for organism identification.			

DISCONTINUED TEST:

Test 8321	Fungus Culture, Blood
Effective:	3/1/2021
Comment:	Please note that this test will be discontinued. Please see below for the recommended alternative testing.

RECOMMENDED ALTERNATIVE:

Test 907207	Fungus Culture, Bl	Fungus Culture, Blood		
Effective:	3/1/2021	3/1/2021		
Specimen:	Whole blood in an is container.	solator tube (supply order #6074). Ind	licate source on the request form and specimen	
Reference Ranges:	See Report			
Stability:		Room temperature: 24 Hours Refrigerated: Do Not Refrigerate Frozen: Unacceptable		
Method:	Conventional Fungu	s Culture		
Setup:	Days: Monday – Su	nday		
Reports:	3 Weeks (growth de	3 Weeks (growth dependent)		
CPT*:	87103	87103		
Price:	Client: 45.38 Patier	Client: 45.38 Patient: \$59.01		
Interface Mapping:	Result Code 10907207			
	Ask at order entry questions			
	Result Code 99908321	Result Name Result Options Source: Free Text		
Comment:	Testing will be performed at our main laboratory in Phoenix. Antibiotic susceptibilities will be performed on pathogens as necessary at an additional charge. Additional charge for organism identification.			

DISCONTINUED TEST:

Test 8176	Fungus Culture, Miscellaneous
Effective:	3/1/2021
Comment:	Please note that this test will be discontinued. Please see below for the recommended alternative testing.

RECOMMENDED ALTERNATIVE:

Test 907205	Fungus Culture, Misc	ellaneous		
Effective:	3/1/2021			
Specimen:	Bone Marrow: Submit r	refrigerated in an Isolator Tube (su	pply order #6074).	
	Sterile Body Fluids: Su	bmit refrigerated in an Isolator Tub	pe (supply order #6074).	
	Other specimens: Steri	ile container or swab in transport m	nedia.	
Reference Ranges:	See Report			
Stability:	Room temperature: 24			
	Refrigerated: 72 Hours			
	Frozen: Unacceptable			
Method:	Conventional Fungus (Culture		
Setup:	Days: Monday – Sunda	ау		
Reports:	3 Weeks (growth dependent)			
CPT*:	87102			
Price:	Client: \$35.51 Patient: \$59.01			
Interface Mapping:	Result Code	Result Name		
	10907205	10907205 Fungus Culture, Miscellaneous		
	Ask at order entry questions			
	Result Code Result Name Result Options			
	99908176	Source:	Free Text	
	99990000 Other: Free Text			
Comment:	Testing will be performed at our main laboratory in Phoenix. Antibiotic susceptibilities will be performed on			
	pathogens as necessar	ry at an additional charge. Addition	nal charge for organism identification.	



ANNOUNCEMENT: Testing Delays Due to Severe Weather

Reference Laboratory Testing

A number of referral laboratories that Sonora Quest sends samples to for testing are routed through the Memphis International Airport. As severe weather continues to result in flight cancelations, you may experience delays in reporting or potential cancellations due to sample stability. **We encourage pausing collection of these samples until the weather conditions stabilize.**

Below is a list of the impacted referral laboratories and the most frequently ordered tests performed at these locations:

- Advanced Diagnostics Lab at National Jewish Health
- Allermetrix
- Cleveland HeartLab
- Diatherix Laboratories
- Kansas State Rabies Laboratory
- Mayo Clinic (Rochester, MN)
- NMS Laboratories
- Quest Diagnostics Nichols Institute (Chantilly, VA)
- Quest Diagnostics TB (Memphis, TN)
- UT Department of Pathology Fungus Testing Laboratory
- Versiti (Wisconsin)
- Viracor

Test Name	Test Code
Bile Acids, Total	901268
Cardio IQ Myeloperoxidase (MPO)	906896
Cardio IQ Oxidized LDL	906898
Creatine	2300
Diatherix Bacterial Vaginosis Panel	906452
Diatherix Candidiasis Panel	906450
Diatherix Gastrointestinal Panel	906706
Diatherix Health Panel (15:1)	906142
Diatherix Sexually Transmitted Disease Panel	906451
Diatherix Viral Respiratory Panel	906704
Diptheria Antitoxoid	3171
Everolimus, LC/MS/MS, Blood	905453
Hemoglobin, Free, Plasma	703020
HIV-1 DNA, Qualitative PCR	3753
Oxacarbazepine Metabolite	79104
T-SPOT.TB	906927
Valproic Acid, Free	10286

Virus Cultures

The national weather conditions are also impacting our ability to obtain supplies necessary to perform virus cultures. Samples received for the following tests will be frozen until supplies are received. Test results may be delayed during this time.

Test Name	Test Code
Herpes Simplex Virus Culture with Typing	8181
Herpes Simplex Virus Culture without Typing	7158
Virus Culture, CMV	708144
Virus Culture, General	708169
Virus Culture, Sepsis	900039
Virus Culture, Varicella Zoster/Herpes Simplex with DFA	8158



ANNOUNCEMENT: Testing Resumed - Platelet Aggregation

In client gram volume 10, it was announced that due to instrumentation issues at the performing lab, Banner University Medical Center Phoenix (BUMCP), testing for Platelet Aggregation (test code 710439) is currently unavailable. Platelet Aggregation testing has resumed at BUMCP. Testing is available Monday through Friday.



UPDATE: Testing Delays Due to Severe Weather

As national weather conditions have improved, we are no longer experiencing testing delays or sample stability issues as referenced in a Client Gram sent on February 18.

Referral Laboratories

You may resume collection of samples sent to the following referral laboratories. The most commonly ordered tests sent to these locations are also listed below.

- Advanced Diagnostics Lab at National Jewish Health
- Allermetrix
- Cleveland HeartLab
- Diatherix Laboratories
- Kansas State Rabies Laboratory
- Mayo Clinic (Rochester, MN)
- NMS Laboratories
- Quest Diagnostics Nichols Institute (Chantilly, VA)
- Quest Diagnostics TB (Memphis, TN)
- UT Department of Pathology Fungus Testing Laboratory
- Versiti (Wisconsin)
- Viracor

Test Name	Test Code
Bile Acids, Total	901268
Cardio IQ Myeloperoxidase (MPO)	906896
Cardio IQ Oxidized LDL	906898
Creatine	2300
Diatherix Bacterial Vaginosis Panel	906452
Diatherix Candidiasis Panel	906450
Diatherix Gastrointestinal Panel	906706
Diatherix Health Panel (15:1)	906142
Diatherix Sexually Transmitted Disease Panel	906451
Diatherix Viral Respiratory Panel	906704
Diptheria Antitoxoid	3171
Everolimus, LC/MS/MS, Blood	905453
Hemoglobin, Free, Plasma	703020
HIV-1 DNA, Qualitative PCR	3753
Oxacarbazepine Metabolite	79104
T-SPOT.TB	906927
Valproic Acid, Free	10286

Virus Cultures

Supplies have been received for the following tests and we are no longer experiencing delays in testing.

Test Name	Test Code
Herpes Simplex Virus Culture with Typing	8181
Herpes Simplex Virus Culture without Typing	7158
Virus Culture, CMV	708144
Virus Culture, General	708169
Virus Culture, Sepsis	900039
Virus Culture, Varicella Zoster/Herpes Simplex with DFA	8158

DISCONTINUED TEST:

Test 9380	17-Ketogenic Steroids with Creatinine, Urine 24Hr
Effective:	Immediately
Comment:	Please note that this test has been discontinued by the reference laboratory due to the reagent is no longer being manufactured. There is no alternative testing at this time.
	no longer being manufactured. There is no alternative testing at this time.

DISCONTINUED TEST:

Test 903001	Aspirin Resistance (11-Dehydrothromboxane B2)
Effective:	3/8/2021
Comment:	Please note that this test will be discontinued. Please see below for the recommended alternative testing.

RECOMMENDED ALTERNATIVE:

Test 907223	Aspirin Works® (11-	-dhTXB2/Creatinine)	
Effective:	3/1/2021		
Specimen:		a C and S gray urine vacutainer tube (3 mL min). If urine is not collected in	
	C and S grey urine va	acutainer tube, it must be poured off into this tube within 4 hours of	
	collection. Store refrig	gerated and ship same day as collected if possible	
Reference Ranges:	See Report		
Stability:	Room temperature: 7 Days		
		Refrigerated: 14 Days	
	Frozen: 90 Days		
Method:	Enzyme Linked Immunoassay (ELISA), Calculation		
Setup:	Days: Monday – Friday		
Reports:	3-7 Days		
CPT*:	84431, 82570		
Price:	Client: \$125.00 Patient: \$158.70		
Interface Mapping:	Result Code	Result Name	
	11907223	Thromboxane (Aspirinworks)	
	12907223	11DHTXB2/Creatinine Ratio	
	20906899	Creatinine, Urine, Random	
Comment:	Testing will be performed at Cleveland Heart Labs.		



ASSAY CHANGE: SARS-CoV-2 Antibody (IgG), Spike, Semi-Quantitative

Detection of IgG antibodies may indicate prior exposure to a SARS-CoV-2 (COVID-19) infection or an immune response to a COVID-19 vaccination.

Testing is recommended at least 10 days after potential exposure to Coronavirus or the onset of symptoms, to allow for the development of IgG antibodies. The IgG antibody test provides insight into an individual's immune response after exposure to the SARS-CoV-2 virus (Coronavirus) but is not intended for diagnosis of active infection. Follow-up may be appropriate with a physician to determine if testing with a molecular diagnostic test should be considered to rule out an active infection.

For those vaccinated, the measurement of IgG levels can provide insight to an individual's immune response to the COVID-19 vaccine. The evaluation of an immune response should occur at least 10 days after the completion of the vaccine series. Although this test is designed to assess the level of an individual's immune response, studies are still needed to determine the level that indicates protective immunity as well as how long the immune response may last.

Interpretation of results:

- A **Nonreactive** result indicates that you have not developed a detectable level of SARS-CoV-2 IgG. It does not rule out a SARS-CoV-2 infection, particularly if you have been in contact with the virus. In addition, your immune response may be depressed if elderly, immunocompromised, or immunosuppressed.
- A Reactive result indicates that you have recovered from SARS-CoV-2 infection or have developed an immune
 response to a COVID-19 vaccine. Although uncommon, false-positive results may occur due to cross-reactivity from
 pre-existing antibodies to other human coronaviruses or other possible causes. A positive result does not mean you
 are immune to SARS-CoV-2 (Coronavirus) as additional studies are required to understand the immune status in
 relation to COVID-19.

Test 907097	SARS-CoV-2 Antibo	ody (IgG), Spike, Semi-Quantitative
Effective:	3/15/2021	
Specimen:	1 mL refrigerated serum from a serum separator tube (SST) (0.5 mL min).	
Reference Ranges:	Nonreactive: <1.00	
	Reactive: ≥1.00	
Stability:	Room temperature: 24 Hours	
	Refrigerated: 7 Days	
	Frozen: 30 Days	
Method:	Chemiluminescent Ir	
Setup:	Evenings: Monday –	
	Nights: Monday – Su	ınday
Reports:	1-3 Days	
CPT*:	86769	
Price:	Client: \$65.00 Patie	· ·
Interface Mapping:	Result Code	Result Name
	10907097	SARS-CoV-2 lgG, Result
	20907097	SARS-CoV-2 IgG, Interpretation
Comment:	The SARS-CoV-2 IgG assay is intended for qualitative and semi-quantitative detection of IgG antibodies to the S1 receptor binding domain (RBD) of the SARS-CoV-2 spike protein. The measurement of IgG levels can provide insight to an individual's adaptive immune response to a SARS-CoV-2 infection or vaccination. Although the assay is designed to assess the level of an individual's immune response, studies are still needed to determine the index level threshold that confers protective immunity as well as how long the adaptive immune response may last post-infection or via vaccination.	
	This test should not be used to diagnose or exclude an acute SARS-CoV-2 infection. If acute infection is suspected, direct testing by molecular methods for SARS-CoV-2 is necessary. False positive results for the test may occur due to cross-reactivity from pre-existing antibodies or other possible causes.	
	following websites: https://www.sonorag	act Sheets" available for health care providers and patients using the uest.com/covid-19-information-for-healthcare-providers/ uest.com/covid-19-information-for-patients/
		uthorized by the FDA under an Emergency Use Authorization (EUA) for boratories. The FDA authorized labeling is available at om/Antibody.



IMPORTANT BILLING UPDATE:

When submitting a COVID-19 laboratory order to Sonora Quest Laboratories or responding to requests for missing/invalid ICD-10 codes (changes effective 1/1/2021):

The Centers for Disease Control and Prevention's National Center for Health Statistics (CDC/NCHS), as a result of the ongoing COVID-19 public health emergency, are implementing additional codes into the International Classification of Diseases, Tenth Revision, Clinical Modifications (ICD-10-CM) for reporting to include:

- Encounter for screening for COVID-19 (Z11.52)
- Contact with and (suspected) exposure to COVID-19 (Z20.822)
- Personal history of COVID-19 (Z86.16)
- Multisystem inflammatory syndrome (MIS) (M35.81)
- Other specified systemic involvement of connective tissue (M35.89)
- Pneumonia due to coronavirus disease 2019 (J12.82)

These new codes will be effective January 1, 2021 to identify conditions resulting from COVID-19.

Please refer to the CDC recommendations below for further coding guidance. All applicable ICD-10 coding provided must be documented in the patient's medical chart.

Full CDC coding guidance can be found at: https://www.cdc.gov/nchs/icd/icd10cm.htm

Recently added symptoms:

- R68.83 Chills (without fever)
- M79.10 Myalgia, unspecified site
- R51 Headache
- J02.9 Acute pharyngitis, unspecified
- Loss of taste or smell R43.9, if loss of both R43.8
- a) For patients presenting with any signs/symptoms (such as fever, etc.) and where a definitive diagnosis has not been established, please provide ICD-10 codes that correlate with signs/symptoms that are documented in the patient's medical chart. U07.1 is only applicable if COVID-19 is confirmed positive at the time the testing is being ordered and documented in the patient's medical record (effective for dates of service on or after 04/01/20). Code only confirmed cases. Presumptive positive COVID-19 test should be coded as confirmed.
- b) When COVID-19 meets the definition of principal diagnosis, code U07.1, COVID-19, should be sequenced first, followed by the appropriate codes for associated manifestations.

c) Acute respiratory illness due to COVID-19

(i) Pneumonia

For a pneumonia case confirmed as due to the 2019 novel coronavirus (COVID-19), assign codes U07.1, COVID-19, and J12.89, Other viral pneumonia.

(ii) Acute Bronchitis

For a patient with acute bronchitis confirmed as due to COVID-19, assign codes U07.1 and J40, Bronchitis, not specified as acute or chronic.

(iii) Lower respiratory infection

If the COVID-19 is documented as being associated with the lower respiratory infection, not otherwise specified (NOS), or an acute respiratory infection, NOS, codes U07.1 and J22, Unspecified acute lower respiratory infection, should be assigned.

(iv) Acute respiratory distress syndrome

For acute respiratory distress syndrome (ARDS) due to COVID-19, assign codes U07.1 and J80, Acute respiratory distress syndrome.

d) Exposure to COVID-19

For cases where there is a concern about a possible exposure to COVID-19, but this is ruled out after evaluation, assign code Z03.818, Encounter for observation for suspected exposure to other biological agents ruled out.

For cases where there is an actual exposure to someone who is confirmed or suspected (not ruled out) to have COVID-19, and the exposed individual either tests negative or the test results are unknown, assign code Z20.828, Contact with and (suspected) exposure to other viral communicable diseases. If the exposed individual tests positive for the COVID-19 virus, see guideline a).

e) Screening for COVID-19

For asymptomatic individuals who are being screened for COVID-19 and have no known exposure to the virus, and the test results are either unknown or negative, assign code Z11.59, Encounter for screening for other viral diseases. For individuals who are being screened due to a possible or actual exposure to COVID-19, see guideline d).

f) Signs and symptoms without definitive diagnosis of COVID-19

For patients presenting with any signs/symptoms associated with COVID-19 (such as fever, etc.) but a definitive diagnosis has not been established, assign the appropriate code(s) for each of the presenting signs and symptoms such as:

- R05 Cough
- R06.02 Shortness of breath
- R50.9 Fever, unspecified

If a patient with signs/symptoms associated with COVID-19 also has an actual or suspected contact with or exposure to someone who has COVID-19, assign Z20.828, Contact with and (suspected) exposure to other viral communicable diseases, as an additional code. This is an exception to guideline I.C.21.c.1, Contact/Exposure.

g) Asymptomatic individuals who test positive for COVID-19

For asymptomatic individuals who test positive for COVID-19, assign code U07.1, COVID-19. Although the individual is asymptomatic, the individual has tested positive and is considered to have the COVID-19 infection.



UPDATE: SARS-CoV-2 Antibody (IgG), Spike, Semi-Quantitative

Detection of IgG antibodies may indicate prior exposure to a SARS-CoV-2 (COVID-19) infection or an immune response to a COVID-19 vaccination.

Testing is recommended at least 10 days after potential exposure to Coronavirus or the onset of symptoms, to allow for the development of IgG antibodies. The IgG antibody test provides insight into an individual's immune response after exposure to the SARS-CoV-2 virus (Coronavirus) but is not intended for diagnosis of active infection. Follow-up may be appropriate with a physician to determine if testing with a molecular diagnostic test should be considered to rule out an active infection.

For those vaccinated, the measurement of IgG levels can provide insight to an individual's immune response to the COVID-19 vaccine. The evaluation of an immune response should occur at least 14 days after the completion of the vaccine series. Although this test is designed to assess the level of an individual's immune response, studies are still needed to determine the level that indicates protective immunity as well as how long the immune response may last.

Interpretation of results:

- A **Nonreactive** result indicates that you have not developed a detectable level of SARS-CoV-2 IgG. It does not rule out a SARS-CoV-2 infection, particularly if you have been in contact with the virus. In addition, your immune response may be depressed if elderly, immunocompromised, or immunosuppressed.
- A Reactive result indicates that you have recovered from SARS-CoV-2 infection or have developed an immune
 response to a COVID-19 vaccine. Although uncommon, false-positive results may occur due to cross-reactivity from
 pre-existing antibodies to other human coronaviruses or other possible causes. A positive result does not mean you
 are immune to SARS-CoV-2 (Coronavirus) as additional studies are required to understand the immune status in
 relation to COVID-19.

Test 907097	SARS-CoV-2 Antik	oody (IgG), Spike, Semi-Quantitative	
Effective:	3/22/2021		
Specimen:	1 mL refrigerated se	1 mL refrigerated serum from a serum separator tube (SST) (0.5 mL min).	
Reference Ranges:	Nonreactive: <1.00		
	Reactive: ≥1.00		
Stability:	Room temperature:	Room temperature: 24 Hours	
	Refrigerated: 7 Day	vs .	
	Frozen: 30 Days		
Method:	Chemiluminescent	Immunoassay	
Setup:	Evenings: Monday	Evenings: Monday – Friday	
	Nights: Monday – S	Nights: Monday – Sunday	
Reports:	1-3 Days		
CPT*:	86769	86769	
Price:	Client: \$65.00 Pat	ient: \$99.00	
Interface Mapping:	Result Code	Result Name	
	10907097	SARS-CoV-2 lgG, Result	
	20907097	SARS-CoV-2 IgG, Interpretation	
Comment:	antibodies to the S ² measurement of Ig0 to a SARS-CoV-2 ir of an individual's im threshold that confe	gG assay is intended for qualitative and semi-quantitative detection of IgG I receptor binding domain (RBD) of the SARS-CoV-2 spike protein. The G levels can provide insight to an individual's adaptive immune response infection or vaccination. Although the assay is designed to assess the level immune response, studies are still needed to determine the index level ers protective immunity as well as how long the adaptive immune post-infection or via vaccination.	
	This test should not be used to diagnose or exclude an acute SARS-CoV-2 infection. If acute infection is suspected, direct testing by molecular methods for SARS-CoV-2 is necessary. False positive results for the test may occur due to cross-reactivity from pre-existing antibodies or other possible causes. This test has been authorized by the FDA under an Emergency Use Authorization (EUA) for use by authorized laboratories. The FDA authorized labeling is available at		
	www.SonoraQuest.		



ANNOUNCEMENT: Bordetella DFA Testing Delay

Due to a product backorder, testing for Bordetella DFA (test code 15053) will be unable to be performed after Saturday, March 13. Testing will resume once the backorder product is received which is expected to be Monday, March 29.

Please Note: Bordetella Culture testing will not be affected.

ANNOUNCEMENT: Scheduled Application Server Maintenance

On Sunday, March 28, 2021, Sonora Quest Laboratories will be performing application server maintenance. As a result, all computer systems will be unavailable beginning at 6:00 p.m. until approximately 11:00 p.m.

During this period, our Client Services Representatives will not have access to computerized information or patient results. All reporting will resume by normal processes once our systems are operational. Quanum users will have uninterrupted access to patient results that were completed prior to 6:00 p.m. on Sunday. Quanum will be updated with results completed during the downtime once the server maintenance is completed.

STAT courier services and STAT testing will remain available during this time and will be reported manually by phone via our established downtime processes.

During the downtime, please contact us as follows:

- Client Services: 602.685.5050
- Logistics (for pick-ups including Mobile Diagnostic Services STAT phlebotomy requests): 602.685.5052

ASSAY CHANGES:

Test 902280	Old Name: Sm Antibody New Name: Sm (Smith) Antibody
Effective:	3/15/2021

Test 9238	Hepatitis Be Antibody
Effective:	3/15/2021
Method:	Chemiluminescent Immunoassay
Reference Ranges:	Non-Reactive
Stability:	Ambient: 4 Days
	Refrigerated: 7 Days
	Frozen: Not Established



ANNOUNCEMENT: Cardio IQ Galectin-3 Testing Delay

Due to reagent unavailability at the referral laboratory, testing for Cardio IQ Galectin-3 (test code 906902) is currently delayed. Specimens will be frozen until reagent is received, which is expected to arrive mid to late April.

ANNOUNCEMENT: 12 mL Conical Bottom Skirted Polypropylene Urine Tube Backorder

Due to a manufacturer backorder, supply items 12 mL Conical Bottom Skirted Polypropylene No Cap (supply #42782) and Yellow Stopper/Cap (supply #42782) are currently unavailable. Supply item 10mL Round Bottom Urine Tube (supply #23247) is being sent as an alternative supply.

Please Note: Item #10845 – Urine Transfer Straw will be issued out with each tube of substitution item #23247 – 10mL Round Bottom Urine Tube.

We anticipate this to be a temporary substitution until the 12 mL Conical Bottom Skirted Polypropylene Urine Tubes (supply #42782) become available. Please continue to order item #42782 when placing supply orders.

Testing	Supplies
Urine Chemistry Testing	Supply #42782 – 12 mL Conical Bottom Skirted Polypropylene No Cap AND
(Protein, Urea Nitrogen,	Supply #42783 – Yellow Stopper/Cap
Magnesium, Chloride, Sodium, etc.);	
Toxicology/Drug Screen Testing	

For questions regarding supplies please contact our warehouse at the following numbers:

Phoenix: Phone (602) 685.5264; Fax (602) 685.5402 Toll-free (800) 766.6721, ext. 5264

Tucson: Phone (520) 784.8004; Fax (520) 296.5607 Toll-free (800) 266.8101

ANNOUNCEMENT: Temporary Patient Service Center Closures

Sonora Quest Laboratories is committed to ensuring the safety of our guests and communities along with our Sonora Quest team members. As we manage through this pandemic, we have made a difficult decision to **temporarily** close the following Patient Service Centers beginning Thursday, April 1 and anticipate reopening on Monday, May 3.

City	Address
Florence	3325 N. Hunt Hwy. (Inside Safeway), Florence, AZ 85132
Mesa	1225 W. Guadalupe Rd. (Inside Safeway), Mesa, AZ 85202

Additionally, our Patient Service Center located 1425 S. Greenfield Rd., #102, in Mesa will be temporarily closed beginning Friday, March 19 and anticipate reopening on Monday, April 5.

Patients with appointments at these locations will be contacted and rescheduled at alternative locations.

Patients can visit SonoraQuest.com for a complete listing of our Patient Service Centers and to schedule appointments.



URGENT ANNOUNCEMENT: Phone System Issues Resolved

Beginning this morning, and lasting until approximately 3:15 p.m. this afternoon, we experienced intermittent issues with our phone and fax systems. Some faxes and calls were being dropped mid-call or callers were unable to get through. The fax confirmations to senders may have shown that a fax sent to us was successful, even though it was not received.

Additionally, reporting completed by fax was affected by these issues and slight delays should be expected as the faxes are completed.

These issues have been resolved.

Due to the phone issues, the following is recommended to ensure a smooth experience for your patients visiting our Patient Service Centers:

Submission of faxed orders for patients visiting our Patient Service Centers: Orders faxed to our Patient
Service Centers today may not have been received; please re-fax patient orders to ensure they are received by our
Patient Service Centers.

For additional resources at any time, below are alternative methods of contacting us:

Client Services:

- You may fax requests or resolution to tests in question to our Client Services department at 602.685.5401.
- For add-on testing please use the <u>Add-On Form</u> located at SonoraQuest.com/Provider/Provider-Resources
 Laboratory Reference Materials.
- For requesting results, please use the <u>Patient Results Request Form</u> located at SonoraQuest.com/Provider/Provider-Resources > Laboratory Reference Materials.

Logistics:

- You may fax requests to our Logistics department at 602.685.5070.
- You may also email requests to Logistics at <u>SQLDispatchGroup@SonoraQuest.com</u>.

Billing:

You may use SonoraQuest.com/Contact-Us for Billing requests.

Anatomic Pathology & Cytology:

O You may fax requests to 602.685.5695.

Thank you for your patience and understanding as we worked through this issue.



ANNOUNCEMENT: 21st Century Cures Act "Information Blocking Rule" Effective April 5

As you should be aware, the Prohibition on Information Blocking, which is part of the 21st Century Cures Act, is effective April 5 (originally intended to be November 2020). This portion of the rule, available at https://www.healthit.gov/curesrule/final-rule-policy/information-blocking, requires that no healthcare entity delay any result access for patients.

For Sonora Quest, it has been determined that a patient creating a results portal account to access test results constitutes as a request for access. Sonora Quest will ensure that ALL results are available via our results portal immediately upon completion of testing.

Summarized from the Information Blocking Rule FAQ (link available below):

- Question: Are health care providers expected to release test results to patients through a patient portal or application programming interface (API) as soon as the results are available to the ordering clinician? (question asked/answered *1/15/2021*)
 - Answer: While the information blocking regulations do not require health care providers to proactively make electronic health information (EHI) available, once a request to access, exchange or use EHI is made, health care providers must timely respond to the request (for example, from a patient for their test results). Delays or other unnecessary impediments could implicate the information blocking provisions.

In practice, this could mean a patient would be able to access EHI such as test results in parallel to the availability of the test results to the ordering clinician.

For more information visit the https://www.healthit.gov/curesrule/ website and see the Information Blocking FAQs.



ANNOUNCEMENT: Blood Culture Bottles

In the interest of safety during specimen transport and testing, orders placed for glass blood culture bottles will been replaced with new plastic bottles. When you receive the plastic blood culture bottles, please store them in a cool, dry place (2-25° C/35.6-77° F), and out of direct light.

Please use the new supply numbers below when placing future orders. You may continue to use any of the glass bottles that you may have remaining.

Supply	Old Supply Number	New Supply Number
BLOOD CULTURE BOTTLE - AEROBIC	6121	33593
BLOOD CULTURE BOTTLE - ANAEROBIC/LYTIC	7507	33594
BLOOD CULTURE BOTTLE - PEDIATRIC	6088	33595

Supply orders can be made through our Provider Portal at SonoraQuest.com, our Quanum™ system, or by faxing a client supply requisition to our warehouse. For updated client supply requisitions, please call 602.685.5141.

For questions regarding supplies please contact our warehouse at the following numbers:

Phoenix: Phone (602) 685.5264; Fax (602) 685.5402 Toll-free (800) 766.6721, ext. 5264

Tucson: Phone (520) 784.8004; Fax (520) 296.5607 Toll-free (800) 266.8101

BILLING CHANGE:

Test 2044	Salicylate Level
Effective:	3/26/2021
CPT*:	80179

ASSAY CHANGES:

Test 902968	First Trimester Screen, hCG		
Effective:	4/12/2021		
Interfacing Mapping:	Ask at order entry question		
	Result Code Result Name Response Options		
	99906563	Pregnancy Result of IVF	Free Text

Test 901714	First Trimester Screen, H-hCG		
Effective:	4/12/2021		
Interfacing Mapping:	Ask at order entry question		
	Result Code Result Name Response Options		
	99906563	Pregnancy Result of IVF	Free Text

Test 906681	HIV-1 RNA, Qualitative Real-Time PCR
Effective:	Immediately
Method:	Real-Time Polymerase Chain Reaction
Specimen:	1.6 mL refrigerated serum collected from serum separator tube (SST) (0.8 mL min) or a red top tube or 1.6 mL refrigerated plasma collected from EDTA (lavender-top) or potassium EDTA (white-top) tube (0.8 mL min).
Stability:	Room temperature: 72 Hours Refrigerated: 5 Days Frozen: 42 Days

Test 906563	Integrated Screen, Part 1		
Effective:	4/12/2021		
Interfacing Mapping:	Ask at order entry question		
	Result Code	Result Name	Response Options
	99906563	Pregnancy Result of IVF	Free Text

Test 903323	Integrated Screen, Part 2		
Effective:	4/12/2021		
Interfacing Mapping:	Ask at order entry question		
	Result Code Result Name Response Options		
	98906563	Pregnancy Result of IVF	Free Text

Test 102259	Light Chain Screen w	Light Chain Screen w/Reflex Typing, Urine, 24 Hour	
Effective:	4/11/2021		
Interface Mapping:	Result Code	Result Name	
	11002482	Protein, Urine, Timed	
	10002533	Protein, Urine, 24 Hour	
	10009652	Albumin	
	19021734	% Albumin	
	19121733	M-spike (Beta Globulin)	
	19123733	M-Spike 2 (Beta Globulin)	
	19124733	M-Spike 3 (Beta Globulin)	
	19021733	M-spike (Gamma Globulin)	
	19422733	M-Spike 2 (Gamma Globulin)	
	19423733	M-Spike 3 (Gamma Globulin)	
	19221733	M-spike (Alpha-1-Globulin)	
	19241733	M-Spike 2 (Alpha-1-Globulin)	
	19251733	M-Spike 3 (Alpha-1-Globulin)	
	19321733	M-spike (Alpha-2-Globulin)	
	19331733	M-Spike 2 (Alpha-2-Globulin)	
	19341733	M-Spike 3 (Alpha-2-Globulin)	
	11021733	Globulin	
	19008469	Interpretation	

Test 209223	Light Chain Scree	Light Chain Screen w/Reflex Typing, Urine, Random	
Effective:	4/11/2021		
Interface Mapping:	Result Code	Result Name	
	2498	Creatinine, Urine, Random	
	10002482	Protein, Urine, Random	
	19002533	Protein, Urine, Normalized	
	11022372	Albumin	
	11021734	% Albumin	
	19122373	M-spike (Beta Globulin)	
	19133373	M-Spike 2 (Beta Globulin)	
	19134373	M-Spike 3 (Beta Globulin)	
	19022373	M-spike (Gamma Globulin)	
	19042373	M-Spike 2 (Gamma Globulin)	
	19052373	M-Spike 3 (Gamma Globulin)	
	19222373	M-spike (Alpha-2-Globulin)	
	19272373	M-Spike 2 (Alpha-2-Globulin)	
	19282373	M-Spike 3 (Alpha-2-Globulin)	
	19322373	M-spike (Alpha-1-Globulin)	
	19322373		
		M-Spike 2 (Alpha-1-Globulin)	
	19362373	M-Spike 3 (Alpha-1-Globulin)	
	11022373	Globulin	
	15008469	Interpretation	
Test 904992	Quad Screen		
Effective:	4/12/2021		
Interfacing Mapping:	Ask at order entry	question	
• •	Result Code	Result Name	Response Options
	99906563	Pregnancy Result of IVF	Free Text
			·
Test 903315	Serum Integrated Screen, Part 1		
Effective:	4/12/2021		
Interfacing Mapping:	Ask at order entry	question	
	Result Code	Result Name	Response Options
	99906563	Pregnancy Result of IVF	Free Text
	1		
Test 903316	Serum Integrated	Screen, Part 2	
Effective:	4/12/2021		
Interfacing Mapping:	Ask at order entry		
	Result Code	Result Name	Response Options
	98906563	Pregnancy Result of IVF	Free Text
Took 002205	Commontial Interme	stad Cayaan Dayt 1	
Test 903305	•	ted Screen, Part 1	
Effective:	4/12/2021		
Interfacing Mapping:			
	Result Code	Result Name	Response Options
	99906563	Pregnancy Result of IVF	Free Text
Test 903317	Sequential Integra	ited Screen, Part 2	
Effective:	4/12/2021	Sequential Integrated Screen, Part 2	
Interfacing Mapping:	_	Ask at order entry question	
michiacing mapping.	Result Code	Result Name	Response Options
	98906563	Pregnancy Result of IVF	Free Text
	30300000	Tregnancy Nesult Orivi	1100 101



ANNOUNCEMENT: Martin-Hopkins Calculation for LDL-C

Beginning Monday, April 26, Sonora Quest Laboratories will transition its current calculated LDL-C to the new Martin-Hopkins equation. This transition will result in greater accuracy in assessing concentrations of LDL-C and remove the necessity for patient fasting prior to the test. Please refer to the Martin-Hopkins Calculation information sheet located at Martin-Hopkins Calculation information sheet located at https://www.sonoraquest.com/provider/provider-resources/cardiovascular-disease/ for further information on this update.

The advantages of the Martin-Hopkins ensure identification of at-risk patients:

- ➤ Improved accuracy in assessing LDL-C levels in patients with very high or low LDL-C levels regardless of TG levels between 150 mg/dL and 400 mg/dL
- Comparable to a direct LDL-C measurement

Key benefits for patients include:

- Adjustable factor that allows for personalization of the LDL-C versus a one-size-fits all calculation
- Improved accuracy for non-fasting samples, which allows patients to skip the fast prior to their lipid panel blood draw

Please see below for testing that will have the new LDL calculation:

Test Code	Test Name
2902	Cardiac Risk Panel
906937	Cardio IQ Lipid Panel
906938	Cardio IQ Lipid Panel with reflex to Direct LDL
1877	Lipid Panel
902278	Lipid Panel with reflex to LDL Direct

ANNOUNCEMENT: Fulfillment Center Closed for Inventory

The Sonora Quest Laboratories Fulfillment Center will be closed on Thursday, April 22 and Friday, April 23, to conduct inventory. Any orders received after noon on Monday, April 19, will not be entered and processed until Monday, April 26, to allow sufficient time for order entry and processing. Orders will not be processed on the closure days, and Fulfilment Center staff will be unavailable during the closure. Slight delays may occur during the week following the inventory.



DISCONTINUED TEST:

Test 904602	Chromogranin A
Effective:	4/19/2021
Comment:	Test is being discontinued. Please see below for the recommended alternative testing, including initial
	rebaseline testing.

RECOMMENDED ALTERNATIVES:

Due to the methodology change of the alternative testing below, we recommend the following:

- Test Code 907233 Chromogranin A, Rebaseline for patients currently being monitored with Test Code 904602 Chromogranin A; to establish patients with new testing methodology (LC/MS/MS).
- Test Code 907232 Chromogranin A, LC/MS/MS for first time Chromogranin A patient orders and monitoring of patients with established Chromogranin A, Rebaseline (test code 907233).

Test 907233	Chromogranin A, Re	ebaseline	
Specimen:	1 mL frozen serum fro	m a plain (red top) tube (0.5 mL min.).	
Reference Ranges:	See Report		
Stability:	Room temperature: 7	Days	
	Refrigerated: Unaccep	otable	
	Frozen: 14 Days		
Method:	Immunoassay & Chroi	matography/Mass Spectrometry	
Setup:	Days: Monday - Satur	rday	
Reports:	4-6 Days	4-6 Days	
CPT*:	86316		
Price:	Client: \$152.00 Patient: \$152.00		
Interface Mapping:	Result Code	Result Name	
	10907232	Chromogranin A, LC/MS/MS, Serum	
	11904602	Chromogranin A	
Comment:	Test will be available until 4/15/2022		

Test 907232	Chromogranin A, LO	Chromogranin A, LC/MS/MS, Serum	
Specimen:	1 mL frozen serum fro	om a plain (red top) tube (0.5 mL min.).	
Reference Ranges:	See Report		
Stability:	Room temperature: 7	Days	
	Refrigerated: Unaccept	otable	
	Frozen: 31 Days		
Method:	Chromatography/Mas	Chromatography/Mass Spectrometry	
Setup:	Days: Monday - Satu	Days: Monday – Saturday	
Reports:	4-6 Days		
CPT*:	86316	86316	
Price:	Client: \$152.00 Patie	Client: \$152.00 Patient: \$152.00	
Interface Mapping:	Result Code	Result Name	
	10907232	Chromogranin A, LC/MS/MS, Serum	

ANNOUNCEMENT: Scheduled Application Server Maintenance

On Sunday, April 25, 2021, Sonora Quest Laboratories will be performing application server maintenance. As a result, all computer systems will be unavailable beginning at 6:00 p.m. until approximately 11:00 p.m.

During this period, our Client Services Representatives will not have access to computerized information or patient results. All reporting will resume by normal processes once our systems are operational. Quanum users will have uninterrupted access to patient results that were completed prior to 6:00 p.m. on Sunday. Quanum will be updated with results completed during the downtime once the server maintenance is completed.

STAT courier services and STAT testing will remain available during this time and will be reported manually by phone via our established downtime processes.

During the downtime, please contact us as follows:

- Client Services: 602.685.5050
- Logistics (for pick-ups including Mobile Diagnostic Services STAT phlebotomy requests): 602.685.5052

ASSAY CHANGE:

Test 907097	SARS-CoV-2 Antibody (IgG), Spike, Semi-Quantitative
Effective:	4/12/2021
Stability:	Room temperature: 72 Hours
	Refrigerated: 7 Days
	Frozen: 7 Days



ANNOUNCEMENT: Trace Element Vial Backorder

Due to a manufacturer backorder, supply #23994 - Vial Trace Element Serum w/Red Label is currently unavailable. At this time an alternative supply item has not been identified. Note that this vial is also included in our RestoreU METHOD Collection Kits (supply #40318). We expect a delivery of vials by approximately April 30 and will fill pending requests in the order received. Additional communications will be sent if the backorder extends beyond this week.

For questions regarding supplies please contact our warehouse at the following numbers:

Phoenix: Phone (602) 685.5264; Fax (602) 685.5402 Toll-free (800) 766.6721, ext. 5264

Tucson: Phone (520) 784.8004; Fax (520) 296.5607 Toll-free (800) 266.8101

NEW ASSAY: SARS-CoV-2 Ab IgG Nucleocapsid, QL

This new test offering is a qualitative assay that identifies COVID antibodies created in response to COVID-19 infection. It does not capture antibodies created in response to vaccine.

Please note, this testing would be useful to clients/patients that need to determine if their antibodies are from infection or from vaccine. If this is required both Test 907097 - SARS-CoV-2 Antibody (IgG), Spike, Semi-Quantitative and Test 907234 - SARS-CoV-2 Antibody (IgG), Spike, Semi-Quantitative should be ordered at the same time.

Test 907234	SARS-CoV-2 Ab IgG	Nucleocapsid, QL
Effective:	4/26/2021	
Specimen:	1 mL refrigerated seru	m from a serum separator tube (SST) (0.1 mL min).
Reference Ranges:	See Report	
Stability:	Room temperature: 4	Days
	Refrigerated: 7 Days	
	Frozen: 30 Days	
Method:	Immunoassay	
Setup:	Days: Monday – Friday	
Reports:	3-5 Days	
CPT*:	86769	
Price:	Client: \$169.00 Patie	nt: \$169.00
Interface Mapping:	Result Code	Result Name
	10907234	SARS-CoV-2 Ab IgG Nucleocapsid, QL
Comment:	Testing is performed a	at Quest Diagnostics.

ASSAY CHANGE:

Test 9360	Viscosity
Effective:	4/26/2021
Method:	Sonoclot Coagulation Analyzer



ASSAY CHANGES:

In order to enhance our Clinical Drug Monitoring test offerings, the below changes will go into effect beginning Monday, May 3. These changes will improve medMATCH® reporting and increase the number of prescribed drugs to submit to 10 drugs.

These changes include:

- A new comment field has been added to include 3 categories of results, making result interpretation easy:
 - Prescribed Consistent = Drugs prescribed and detected
 - Prescribed Inconsistent = Drugs prescribed but not detected
 - Not Prescribed Inconsistent = Drugs detected but not prescribed
- Update to the Prescribed Drug list (from 98 drugs to 145 drugs)
- Additional testing and an updated requisition will be available in the near future

The Prescribed Drug List has been updated to include the following drugs:

None Given	Diazepam	Lyrica™	Roxanol™
	Dilaudid™	Magnacet™	Roxicet™
Abstral™	Dolophine™	Marinol™	Roxicodone™
Aceta Codeine	Dronabinol™	Medical Marijuana	RoxyBond™
Actiq™	Duragesic™	Meperidine	Ryzolt™
Adderall™	Edluar™	Meprobamate	Sandoptal™
Alprazolam	Elavil™	Methadone	Secobarbital
Ambien™	Embeda™	Methamphetamine	Seconal™
Amitriptyline	Endep™	Methorphan	Serax™
Amobarbital	Endocet™	Methylphenidate	SOMA™
Amobarbital Secobarb	Equagesic™	Midazolam	Sublimaze™
Amphetamine	Esgic™	MorphaBond™	Sublocade™
Amytal™	Eszopiclone	Morphine	Suboxone™
Arymo™	Exalgo™	MS Contin™	Subutex™
Ativan	Fanatrex™	Mydayis™	Tapentadol
AventyI™	Fentanyl	Naltrexone	Temazepam
AVINŽA™	Fentora™	Nembutal™	Tramadol
Belbuca™	Fioricet™	Neurontin™	Tranxene™
Bunavail™	Fiorinal™	Norco™	Triazolam
Buprenex™	Flurazepam	Nortriptyline	Tylenol 3™
Buprenorphine	Gabapentin	Nucynta™	Tylenol 4™
Buprenorphine Nalox	Gralise™	Opana™	Tylox™
Butalbital	Halcion™	Oxazepam	Ultram™
Butrans™	Horizant™	Oxycodone	Valium™
Carisoprodol	Hycodan™	OxyContin™	Versed™
Chlordiazepoxide	Hydrocodone	OxyFAST™	Vicodin™
Clonazepam	Hydromorphone	Oxymorphone	Vicoprofen™
Clorazepate	Hysingla™	Pamelor™	Vivitrol™
Codeine	Intermezzo™	Pentobarbital	Vyvanse™
Concerta™	Ionsys™	Percocet™	Xanax™
Conzip™	Kadian™	Percodan™	Xtampza™
Dalmane™	Klonopin™	Perolox™	Zohydro™
Demerol™	Librium™	Phenergan™	Zolpidem
Desoxyn™	Lorazepam	Phenobarbital	Zolpimist™
Dexedrine™	Lorcet™	Pregabalin	Zubsolv™
Dextromethorphan	Lortab™	Restoril™	
Dextrostat™	Lunesta™	Ritalin™	

Please ADD the mapping below for the Drug Monitoring Template (This is a non-orderable code that will be automatically added in the reporting process):

Test 907230	Drug Monitoring Template	
Comment:	This is a non-orderable code that will be automatically added to every offering included in this	
	communication. Please build this test code and additional result codes in your system.	
Interface Mapping:	Result Code	Result Name
	10907230	Notes and Comments
	20907230	Patient Historical Report

Please ADD the mapping below by test/profile and note discontinued tests listed:

Test 906285	PDM Alcohol Metab	olites w/Confirmation
Interface Mapping:	Result Code	Result Name
monaco mapping.	13016217	Alcohol Metab Comments
	10010211	7 too for thouse comments
Test 906330	PDM Alcohol Metab	olite, w/Confirmation and medMATCH
Interface Mapping:	Result Code	Result Name
monaco mapping.	13016217	Alcohol Metab Comments
	10907229	Prescribed Drugs, medMATCH
	10001220	1 1000 ibou Brago, mount 1011
Test 906511	PDM Alcohol Metab	olites, Quantitative, Ur
Interface Mapping:	Result Code	Result Name
	13016217	Alcohol Metab Comments
-	<u>, </u>	<u>'</u>
Test 906512	PDM Alcohol Metab	olites, Quantitative w/medMATCH, Ur
Interface Mapping:	Result Code	Result Name
11 0	13016217	Alcohol Metab Comments
	10907229	Prescribed Drugs, medMATCH
		<u>-</u>
Test 906325	PDM Amphetamines	s w/Confirmation
Interface Mapping:	Result Code	Result Name
	14906264	Amphetamines Comments
Test 906324		s, w/Confirmation and medMATCH
Interface Mapping:	Result Code	Result Name
	14906264	Amphetamines Comments
	10907229	Prescribed Drugs, medMATCH
	1	
Test 906299	PDM Amphetamines	
Interface Mapping:	Result Code	Result Name
	14906264	Amphetamines Comments
Test 906264		s, Quantitative w/medMATCH
Interface Mapping:	Result Code	Result Name
	14906264	Amphetamines Comments
	10907229	Prescribed Drugs, medMATCH
T + 00004=	DD1/4	W/O (
Test 906847		s, W/ Confirmation, w/ D/L Isomers, Urine
Interface Mapping:	Result Code	Result Name
	14906264	Amphetamines Comments
Test 906848		s, w/ Conf w/ D/L Isomers, w/ MedMatch, Urine
Interface Mapping:	Result Code	Result Name
	14906264	Amphetamines Comments
	10907229	Prescribed Drugs, medMATCH

Test 906509	PDM Amphetamines, w D/l	L Isomers Quantitative, Ur	
Interface Mapping:	Result Code	Result Name	
	14906264	Amphetamines Comments	
Test 906510	PDM Amphetamines, w D/l	L Isomers Quantitative w/medMATCH, Ur	
Interface Mapping:	Result Code	Result Name	
	14906264	Amphetamines Comments	
	10907229	Prescribed Drugs, medMATCH	
Test 906323	PDM Barbiturates, w/Confi		
Interface Mapping:	Result Code	Result Name	
	15906300	Barbiturates Comments	
Test 906322	PDM Barbiturates, w/Confi	rmation and medMATCH	
Interface Mapping:	Result Code	Result Name	
interface mapping.	15906300	Barbiturates Comments	
	10907229	Prescribed Drugs, medMATCH	
	10001220	1100011000 21090, 111001111 11 011	
Test 906300	PDM Barbiturates, Quantita	ative	
Interface Mapping:	Result Code	Result Name	
	15906300	Barbiturates Comments	
Test 906265	PDM Barbiturates, Quantita	ative w/medMATCH	
Interface Mapping:	Result Code	Result Name	
	15906300	Barbiturates Comments	
	10907229	Prescribed Drugs, medMATCH	
T +000040	DDMD II I		
Test 906319	PDM Benzodiazepines, w/0		
Interface Mapping:	Result Code	Result Name	
	10906484	Benzodiazepines Comments	
Test 906318	PDM Renzodiazenines w//	Confirmation and medMATCH	
Interface Mapping:	Result Code	Result Name	
interface mapping.	10906484	Benzodiazepines Comments	
	10907229	Prescribed Drugs, medMATCH	
	1 1111111111111111111111111111111111111		
Test 906484	PDM Benzodiazepines, Qu	antitative, Ur	
Interface Mapping:	Result Code	Result Name	
	10906484	Benzodiazepines Comments	
Test 906266	PDM Benzodiazepines, Qu		
Interface Mapping:	Result Code	Result Name	
	10906484	Benzodiazepines Comments	
	10907229	Prescribed Drugs, medMATCH	
Took 000542	DDM Decrease and big a Co	n w/Confirmation III	
Test 906513		PDM Buprenorphine Screen, w/Confirmation, Ur	
Effective:		5/3/2021	
	I Lest is being discontinited	Please see recommended alternative testing below – Test 907218.	
Comment:	Test is being discontinued.		
		ofirmation and madMATCU	
Test 906316	PDM Buprenorphine, w/Co	nfirmation and medMATCH	
	PDM Buprenorphine, w/Co	nfirmation and medMATCH Please see recommended alternative testing without medMATCH below.	

	1==	
Test 907218	PDM, Buprenorphine an	
Interface Mapping:	Result Code	Result Name
	15906316	Buprenorphine Screen
	17906316	Buprenorphine
	19906316	Norbuprenorphine
	11907218	Naloxone
	12907191	Buprenorphine Comments
T / 00007/	DD14.D	
Test 906274	PDM Buprenorphine, Qu	uantitative
Effective:	5/3/2021	1 D)
Comment:	Test is being discontinue	ed. Please see recommended alternative testing below Test 907191
Test 906286	PDM Bunrenorphine Ou	uantitative w/medMATCH
Effective:	5/3/2021	Junitiative willicalvin troll
Comment:		ed. Please see recommended alternative testing without medMATCH below.
Comment.	rest is being discontinue	ed. I lease see recommended alternative testing without medivicing below.
Test 907191	PDM, Buprenorphine an	d Naloxone, Quant, Ur
Interface Mapping:	Result Code	Result Name
11 0	10906286	Buprenorphine
	10907191	Naloxone
	12906286	Norbuprenorphine
	12907191	Buprenorphine Comments
Test 906287	PDM Carisoprodol, Qua	
Interface Mapping:	Result Code	Result Name
	10906287	Carisoprodol Comments
Test 906275	PDM Carisoprodol Quantitative w/medMATCH	
Interface Mapping:	Result Code	Result Name
monaco mapping.	10906287	Carisoprodol Comments
	10907229	Prescribed Drugs, medMATCH
Test 906329	PDM Cocaine Metabolite	e, w/Confirmation
Interface Mapping:	Result Code	Result Name
11 0	10906328	Benzoylecgonine
	10906302	Cocaine Comments
	<u>.</u>	
Test 906328		e, w/Confirmation and medMATCH
Interface Mapping:	Result Code	Result Name
	10906302	Cocaine Comments
	10907229	Prescribed Drugs, medMATCH
Test 906302	PDM Cocaine Metabolite	e Quantitative Ur
Interface Mapping:	Result Code	Result Name
interface Mapping.	10906302	Cocaine Comments
	•	
Test 906267		e, Quantitative w/medMATCH
Interface Mapping:	Result Code	Result Name
	10906302	Cocaine Comments
	10907229	Prescribed Drugs, medMATCH
Test 907181	PDM Eszopiclone, Quar	ntitativa Urina
Interface Mapping:	Result Code	Result Name
mapping.	14907181	Eszopiclone Comments
		I LOZUDIOIUTE UUTITIETIIO

Test 906801	PDM Fentanyl Screen, Urin	ne w/ Rflx Confirmation
Interface Mapping:	Result Code	Result Name
	12906289	Fentanyl Comments
Test 906883		ne w/Confirm, w/medMATCH
Interface Mapping:	Result Code	Result Name
	12906289	Fentanyl Comments
	10907229	Prescribed Drugs, medMATCH
Test 706289	PDM Fentanyl, Quantitative	
Interface Mapping:	Result Code	Result Name
interiace mapping.	12906289	Fentanyl Comments
	12300203	1 charry comments
Test 906276	PDM Fentanyl, Quantitative	e w/medMATCH
Interface Mapping:	Result Code	Result Name
	12906289	Fentanyl Comments
	10907229	Prescribed Drugs, medMATCH
Test 906290	PDM Gabapentin, Quantita	
Interface Mapping:	Result Code	Result Name
	10906290	Gabapentin Comments
T	I pour our annual a	
Test 906277	PDM Gabapentin, Quantita	
Interface Mapping:	Result Code	Result Name
	10906290	Gabapentin Comments
	10907229	Prescribed Drugs, medMATCH
Test 906291	PDM Heroin Metabolite, w/	Confirmation
Interface Mapping:	Result Code	Result Name
interface mapping.	13906516	Heroin Metab Comments
	10000010	TIOTOTT MOULD COMMINING
Test 906278	PDM Heroin Metabolite, w/	Confirmation and medMATCH
Interface Mapping:	Result Code	Result Name
	13906516	Heroin Metab Comments
	10907229	Prescribed Drugs, medMATCH
	T==	
Test 906516	PDM Heroin Metabolite Qu	·
Interface Mapping:	Result Code	Result Name
	13906516	Heroin Metab Comments
Test 906514	PDM Heroin Metabolite Qu	ant w/ medMATCH_Ur
Interface Mapping:	Result Code	Result Name
intoriace mapping.	13906516	Heroin Metab Comments
	10907229	Prescribed Drugs, medMATCH
		· · · · · · · · · · · · · · · · · · ·
Test 906315	PDM Marijuana Metabolite,	w/Confirmation
Interface Mapping:	Result Code	Result Name
	11906303	Marijuana Comments
Test 906314	PDM Marijuana Metabolite,	
Interface Mapping:	Result Code	Result Name
	11906303	Marijuana Comments
	10907229	Prescribed Drugs, medMATCH
T 1000000	DDMM " 11 1 1 "	
Test 906303	PDM Marijuana Metabolite,	
Interface Mapping:	Result Code	Result Name
	11906303	Marijuana Comments

Test 906268	PDM Marijuana Metabolite, Quantitative w/medMATCH	
Interface Mapping:	Result Code	Result Name
	11906303	Marijuana Comments
	10907229	Prescribed Drugs, medMATCH
T +000000		e c
Test 906292	PDM MDMA/MDA, w/Con	
Interface Mapping:	Result Code 10906518	Result Name MDMA Comments
	10900310	MDMA Confinents
Test 906279	PDM MDMA/MDA, w/Con	firmation and medMATCH
Interface Mapping:	Result Code	Result Name
	10906518	MDMA Comments
	10907229	Prescribed Drugs, medMATCH
-		
Test 906518	PDM MDMA/MDA Quant,	
Interface Mapping:	Result Code	Result Name
	10906518	MDMA Comments
Test 906517	PDM MDMA/MDA Quant	w/ medMATCH_Ur
Interface Mapping:	Result Code	Result Name
interface Mapping.	10906518	MDMA Comments
	10907229	Prescribed Drugs, medMATCH
	1	
Test 906293	PDM Meperidine, Quantit	ative
Interface Mapping:	Result Code	Result Name
	12906293	Meperidine Comments
Test 906280	PDM Meperidine, Quantit	
Interface Mapping:	Result Code	Result Name
	12906293	Meperidine Comments
	10907229	Prescribed Drugs, medMATCH
Test 706313	PDM Methadone, w/Confi	rmation
Interface Mapping:	Result Code	Result Name
interface Mapping.	13906304	Methadone Comments
Test 906312	PDM Methadone, w/Confi	rmation and medMATCH
Interface Mapping:	Result Code	Result Name
	13906304	Methadone Comments
	10907229	Prescribed Drugs, medMATCH
T (00000)		
Test 906304	PDM Methadone, Quantit	
Interface Mapping:	Result Code	Result Name
	13906304	Methadone Comments
Test 906269	PDM Methadone, Quantit	ative w/medMATCH
Interface Mapping:	Result Code	Result Name
monaco mapping.	13906304	Methadone Comments
	10907229	Prescribed Drugs, medMATCH
	l	, ,
Test 906540	PDM Methamphetamine,	D/L Isomers, Ur
Interface Mapping:	Result Code	Result Name
	10906540	D/L Methamphet Comments
Test 906294	PDM Methylphenidate Me	
Interface Mapping:	Result Code	Result Name
	10906294	Ritalinic Acid Comments

Test 906332	PDM Methylphenidate, Quantitative w/medMATCH	
Interface Mapping:	Result Code	Result Name
	10906294	Ritalinic Acid Comments
	10907229	Prescribed Drugs, medMATCH
Test 907160	PDM Mitragynine Scree	n w/Rfly Confirmation
Interface Mapping:	Result Code	Result Name
interface mapping.	12907161	Mitragynine Comments
	12001101	militagy mil
Test 907161	PDM Mitragynine, Quan	
Interface Mapping:	Result Code	Result Name
	12907161	Mitragynine Comments
Test 907170	PDM Naltrexone, Quant	itative Ur
Interface Mapping:	Result Code	Result Name
mioriado mapping.	14907170	Naltrexone Comments
Test 906334	PDM Opiates, with Conf	
Interface Mapping:	Result Code	Result Name
	10906487	Opiates Comments
Test 906333	PDM Opiates, with Conf	and medMATCH Ur
Interface Mapping:	Result Code	Result Name
micriado mapping.	10906487	Opiates Comments
	10907229	Prescribed Drugs, medMATCH
	1	
Test 906281	PDM Opiates, Expanded	
Interface Mapping:	Result Code	Result Name
	10906487	Opiates Comments
	11906487	Oxycodone Comments
Test 906335	PDM Oniates Expanded	d Quant with medMATCH, Ur
Interface Mapping:	Result Code	Result Name
mioriaco mapping.	10906487	Opiates Comments
	10906487	Oxycodone Comments
	10907229	Prescribed Drugs, medMATCH
T+000000	DDM C 1 C	ec.e., H.
Test 906306 Effective:	PDM Oxycodone, Quan 5/3/2021	litative, Ur
Comment:		ed. Oxycodone is included in the Expanded Opiate testing above.
Johnnon.	Took is being disconding	ca. Oxygodono is included in the Expanded Opiate testing above.
Test 706271	PDM Oxycodone, Quan	titative w/medMATCH, Ur
Effective:	5/3/2021	
Comment:	Test is being discontinue	ed. Oxycodone is included in the Expanded Opiate testing above.
Test 906327	PDM Oxycodone, with O	
Interface Mapping:	Result Code	Result Name
	11906487	Oxycodone Comments
Test 906326	PDM Oxycodone with C	Conf and medMATCH, Ur
Interface Mapping:	Result Code	Result Name
onaco mapping.	11906487	Oxycodone Comments
	10907229	Prescribed Drugs, medMATCH
L	1	1

Test 706321	PDM Phencyclidine, w/Confirmation	
Interface Mapping:	Result Code	Result Name
	10906307	Phencyclidine Comments
	<u>.</u>	
Test 906320	PDM Phencyclidine, w/Confirmation and medMATCH	
Interface Mapping:	Result Code	Result Name
	10906307	Phencyclidine Comments
	10907229	Prescribed Drugs, medMATCH
Test 906307	PDM Phencyclidine, Qu	
Interface Mapping:	Result Code	Result Name
	10906307	Phencyclidine Comments
T (000070	DDM DI III O	CONTRACTOR
Test 906272		uantitative w/medMATCH
Interface Mapping:	Result Code	Result Name
	10906307 10907229	Phencyclidine Comments
	10907229	Prescribed Drugs, medMATCH
Test 906295	DDM Propobalia Over	titativo Ur
Interface Mapping:	PDM Pregabalin, Quan Result Code	Result Name
interrace mapping.	10906295	Pregabalin Comments
	10900293	Fregabalin Comments
Test 906282	PDM Pregabalin, Quan	titative w/medMATCH
Interface Mapping:	Result Code	Result Name
interface mapping.	10906295	Pregabalin Comments
	10907229	Prescribed Drugs, medMATCH
	10007220	1 Toodhbad Brago, maanii (1 ori
Test 906310	PDM Propoxyphene, w	/Confirmation
Interface Mapping:	Result Code	Result Name
	11906308	Propoxyphene Comments
	<u>.</u>	
Test 906308	PDM Propoxyphene Mo	etabolite, Quantitative, Ur
Interface Mapping:	Result Code	Result Name
	11906308	Propoxyphene Comments
Test 705653	PDM Synthetic Stimula	,
Interface Mapping:	Result Code	Result Name
	18705653	Syn Stimulants Comments
T+000000	DDMT () C	P4 - P
Test 906296	PDM Tapentadol, Quar	
Interface Mapping:	Result Code	Result Name
	10906296	Tapentadol Comments
Test 906331	DDM Tanantadal Over	atitativa w/modMATCH
Interface Mapping:	PDM Tapentadol, Quar Result Code	Result Name
interiace mapping.	10906296	Tapentadol Comments
	10907229	Prescribed Drugs, medMATCH
	10001220	1 100011000 Drugo, mount of t
Test 706297	PDM Tramadol, Quanti	tative Ur
Interface Mapping:	Result Code	Result Name
g.	10906495	Tramadol Comments
	1	
Test 906283	PDM Tramadol, Quanti	tative, w/medMATCH, Ur
Interface Mapping:	Result Code	Result Name
11.0	10906495	Tramadol Comments
	10907229	Prescribed Drugs, medMATCH
•	•	

Test 906298	PDM Tricyclic, Quantitativ	re
Interface Mapping:	Result Code	Result Name
	12906298	Tricyclic Antidep Comments
Test 906284	PDM Tricyclic, Quantitative w/medMATCH	
Interface Mapping:	Result Code	Result Name
	12906298	Tricyclic Antidep Comments
	10907229	Prescribed Drugs, medMATCH
Test 906115	PDM, Zolpidem Quantitati	ive. Ur
Interface Mapping:	Result Code	Result Name
	13906115	Zolpidem Comments
Profiles:		
Test 906258	PDM Base Profile with Co	onfirmation. Ur
Interface Mapping:	Result Code	Result Name
	10906484	Benzodiazepines Comments
	10906328	Benzoylecgonine
	10906302	Cocaine Comments
	10906487	Opiates Comments
	11906487	Oxycodone Comments
Test 906253	PDM Base Profile, with Co	onf and medMATCH Ur
Interface Mapping:	Result Code	Result Name
intoriace mapping.	10906484	Benzodiazepines Comments
	10906328	Benzoylecgonine
	20906328	medMATCH Benzoylecgonine
	10906302	Cocaine Comments
	10906487	Opiates Comments
	11906487	Oxycodone Comments
	10907229	Prescribed Drugs, medMATCH
	1	
Test 906254	PDM Profile 1, w/ Confirm Result Code	Result Name
Interface Mapping:	14906264	1 1000000
		Amphetamines Comments
	15906300	Barbiturates Comments
	10906484 10906328	Benzodiazepines Comments
		Benzoylecgonine
	10906302	Cocaine Comments
	11906303 13906304	Marijuana Comments
	10906487	Methadone Comments
	11906487	Opiates Comments
	10906307	Oxycodone Comments Phencyclidine Comments
	1	
Test 906249	•	mation and medMATCH, Ur
Interface Mapping:	Result Code	Result Name
	14906264	Amphetamines Comments
	15906300	Barbiturates Comments
	10906484	Benzodiazepines Comments
	10906328	Benzoylecgonine
	20906328	medMATCH Benzoylecgonine
	10906302	Cocaine Comments
	11906303	Marijuana Comments
	13906304	Methadone Comments
	10906487	Opiates Comments
	11906487	Oxycodone Comments
İ	10906307	Phencyclidine Comments
	10907229	Prescribed Drugs, medMATCH

Test 906508	PDM Profile 1, w/Confirma	ation and D/L Isomers, Ur	
Interface Mapping:	Result Code	Result Name	
•	14906264	Amphetamines Comments	
	15906300	Barbiturates Comments	
	10906484	Benzodiazepines Comments	
	10906328	Benzoylecgonine	
	10906302	Cocaine Comments	
	11906303	Marijuana Comments	
	13906304	Methadone Comments	
	10906487	Opiates Comments	
	11906487	Oxycodone Comments	
	10906307	Phencyclidine Comments	
Test 906507	PDM Profile 1, w/Conf and	D/L Isomers w/medMATCH, Ur	
Interface Mapping:	Result Code	Result Name	
	14906264	Amphetamines Comments	
	15906300	Barbiturates Comments	
	10906484	Benzodiazepines Comments	
	10906302	Cocaine Comments	
	11906303	Marijuana Comments	
	13906304	Methadone Comments	
	10906487	Opiates Comments	
	11906487	Oxycodone Comments	
	10906307	Phencyclidine Comments	
	10907229	Prescribed Drugs, medMATCH	
Test 906260	PDM Profile 2, withou	ut Confirmation, Ur	
Effective:	5/3/2021		
Comment:	Test is being discont	Test is being discontinued.	
Test 906255	PDM Profile 2, with 0	Confirmation, Ur	
Effective:	5/3/2021		
Comment:	Test is being discontinued.		
Test 906250	PDM Profile 2, with (PDM Profile 2, with Conf and medMATCH, Ur	
Effective:	5/3/2021		
Comment:	Test is being discont	Test is being discontinued.	
Test 906256	PDM Profile 3 with Confirmation, Ur		
Interface Mapping:	Result Code	Result Name	
	14906264	Amphetamines Comments	
	10906484	Benzodiazepines Comments	
	10906328	Benzoylecgonine	
	10906302	Cocaine Comments	
	11906303	Marijuana Comments	
	10906487	Opiates Comments	
	11906487	Oxycodone Comments	

Test 906251	PDM Profile 3, with Conf and medMATCH, Ur	
Interface Mapping:	Result Code	Result Name
	14906264	Amphetamines Comments
	10906484	Benzodiazepines Comments
	10906328	Benzoylecgonine
	20906328	medMATCH Benzoylecgonine
	10906302	Cocaine Comments
	11906303	Marijuana Comments
	10906487	Opiates Comments
	11906487	Oxycodone Comments
	10907229	Prescribed Drugs, medMATCH

Test 906257	PDM Profile 4, with Con	PDM Profile 4, with Confirmation, Ur	
Interface Mapping:	Result Code	Result Name	
	14906264	Amphetamines Comments	
	15906300	Barbiturates Comments	
	10906484	Benzodiazepines Comments	
	10906328	Benzoylecgonine	
	10906302	Cocaine Comments	
	13906304	Methadone Comments	
	10906487	Opiates Comments	
	11906487	Oxycodone Comments	
	10906307	Phencyclidine Comments	

Test 906252	PDM Profile 4, with Cor	PDM Profile 4, with Conf and medMATCH, Ur	
Interface Mapping:	Result Code	Result Name	
	14906264	Amphetamines Comments	
	15906300	Barbiturates Comments	
	10906484	Benzodiazepines Comments	
	10906328	Benzoylecgonine	
	20906328	medMATCH Benzoylecgonine	
	10906302	Cocaine Comments	
	13906304	Methadone Comments	
	10906487	Opiates Comments	
	11906487	Oxycodone Comments	
	10906307	Phencyclidine Comments	
	10907229	Prescribed Drugs, medMATCH	

For all testing that includes medMATCH, please ADD the Ask At Order Entry fields below:

Result Code	Result Name	Response Options	
90907229	PDM Prescription Drugs	Free Text	
91907229	PDM Prescription Drugs	Free Text	
92907229	PDM Prescription Drugs	Free Text	
93907229	PDM Prescription Drugs	Free Text	
94907229	PDM Prescription Drugs	Free Text	
95907229	PDM Prescription Drugs	Free Text	
96907229	PDM Prescription Drugs	Free Text	
97907229	PDM Prescription Drugs	Free Text	
98907229	PDM Prescription Drugs	Free Text	
99907229	PDM Prescription Drugs	Free Text	



ANNOUNCEMENT: Factor XIII Antigen Delay in Testing Due to Reagent Backorder

Due to a reagent backorder for Factor XIII Antigen testing performed at Sonora Quest Laboratories, there is turnaround testing (TAT) delay in the published 2-day TAT and will not be available for STAT ordering over the weekend due to a reagent shortage. Per the supplier, replacement reagent is expected on Tuesday, May 4.

The following tests are impacted:

Factor XIII Antigen testing (test code 907019)



ANNOUNCEMENT: Treponemal-based Testing Changes for Syphilis Due to Reagent Backorder

Due to a nationwide reagent backorder, treponemal-based testing for syphilis using the *T. pallidum* Antibody - Particle Agglutination (TPPA) will be cancelled and automatically replaced with the Fluorescent Treponemal Antibody Absorption (FTA-ABS), performed at Quest Diagnostics.

TPPA testing will resume at Sonora Quest once the backordered reagent is received, which is expected to arrive in early June.

Both the TPPA and the FTA-ABS are treponemal-based tests which can be used to confirm the presence of antibodies to *Treponema pallidum*. All CPT coding and billing will remain the same.

The following tests are impacted:

- Syphilis Screen w/reflex RPR and Titer, or TP-PA (905363)
- RPR Screen w/Reflex RPR Titer and TP-PA (1054)
- Treponema pallidum Antibody PA (900780)

Please add the mapping below by test:

Test 905363	Syphilis Screen w/reflex RPR and Titer, or TP-PA	
Interface Mapping:	Result Code 10091100	Result Name Treponema pallidum Total Antibodies, IFA

Test 1054 RI	RPR Screen w/Reflex RPR Titer and TP-PA	
•	Result Code 0091100	Result Name Treponema pallidum Total Antibodies, IFA

Test 900780	Treponema pallidum Antibody - PA	
Interface Mapping:	Result Code	Result Name
	10091100	Treponema pallidum Total Antibodies, IFA



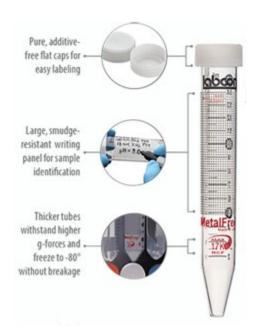
UPDATE: Trace Element Vial Backorder - Alternative Supply Identified

Due to a manufacturer backorder, supply #23994 - Vial Trace Element Serum w/Red Label is currently unavailable. We have identified an alternative supply item (pictured below) that will be automatically substituted for any supply orders received for the following:

- #23994 Viral Trace Element Serum w/Red Label, or
- #40318 RestoreU METHOD Collection Kits

Please continue to order supplies using the stock item numbers above.

Substitute Supply – 15mL Metal-Free w/Flat Cap



For questions regarding supplies please contact our warehouse at the following numbers:

Phoenix: Phone (602) 685.5264; Fax (602) 685.5402 Toll-free (800) 766.6721, ext. 5264

Tucson: Phone (520) 784.8004; Fax (520) 296.5607 Toll-free (800) 266.8101



ANNOUNCEMENT: Turnaround Time Delay - Prescription Drug Monitoring Testing

Due to the extensive changes to this test offering, our current turnaround time (TAT) is estimated to be approximately 5-7 days. We expect this to decrease over the next week until we return to the expected TAT of 3-5 days.

Discontinued Testing Replacement Update:

Effective 5/3/21, the discontinued tests below will automatically be cancelled and replaced with Test 907218:

Test 906513	PDM Buprenorphine Screen, w/Confirmation, Ur
Effective:	5/3/2021
Comment:	Test has been discontinued and will automatically be cancelled and replaced with Test 907218 below.

Test 906316	PDM Buprenorphine, w/Confirmation and medMATCH
Effective:	5/3/2021
Comment:	Test has been discontinued and will automatically be cancelled and replaced with Test 907218 below. Once Buprenorphine and Naloxone, w/Conf and medMATCH is available, the automatic replacement will include medMATCH.

Test 907218	PDM, Buprenorphine	PDM, Buprenorphine and Naloxone, w/Conf, UR	
Interface Mapping:	Result Code	Result Name	
	15906316	Buprenorphine Screen	
	17906316	Buprenorphine	
	19906316	Norbuprenorphine	
	11907218	Naloxone	
	12907191	Buprenorphine Comments	



UPDATE: Turnaround Time Delay – Prescription Drug Monitoring Testing

Due to the extensive changes to this test offering, our current turnaround time (TAT) is estimated to be approximately 10-14 days. We expect this to decrease over the next week until we return to the expected TAT of 3-5 days.

ANNOUNCEMENT: Scheduled Application Server Maintenance

On Sunday, May 23, 2021, Sonora Quest Laboratories will be performing application server maintenance. As a result, all computer systems will be unavailable beginning at 6:00 p.m. until approximately 11:00 p.m.

During this period, our Client Services Representatives will not have access to computerized information or patient results. All reporting will resume by normal processes once our systems are operational. Quanum users will have uninterrupted access to patient results that were completed prior to 6:00 p.m. on Sunday. Quanum will be updated with results completed during the downtime once the server maintenance is completed.

STAT courier services and STAT testing will remain available during this time and will be reported manually by phone via our established downtime processes.

During the downtime, please contact us as follows:

- Client Services: 602.685.5050
- Logistics (for pick-ups including Mobile Diagnostic Services STAT phlebotomy reguests): 602.685.5052



ANNOUNCEMENT: Temporary Patient Service Center Closures

Sonora Quest Laboratories has made the decision to **temporarily** close the following Patient Service Centers and will remain closed until further notice. Patients with appointments at these locations will be contacted and rescheduled at alternative locations.

City	Address
Apache Junction	2080 W. Southern Ave., Suite B6, Apache Junction, AZ 85248
Chandler	1060 E. Ray Rd., Chandler, AZ 85225
Mesa	1425 S. Greenfield Rd., #102, Mesa, AZ 85206
Scottsdale	7281 E. Earll Dr., Bldg. A, #1, Scottsdale, AZ 85251
Sun Lakes	10450 E Riggs Rd Suite 109, Chandler, AZ 85248

Additionally, we have made the decision to **temporarily** close the following Patient Service Centers beginning Monday, May 17, and which will remain closed until further notice. Patients with appointments at these locations will be contacted and rescheduled at alternative locations.

City	Address
Tucson	1940 E. Broadway Rd., Tucson, AZ 85719
Tucson	6565 E. Carondelet Dr., Suite 255, Tucson, AZ 85710

ASSAY CHANGES:

Test 709399	Calcium, Ionized	
Effective:	5/10/2021	
Interface Mapping:	Result Code	Result Name
	10003919	Calcium, Ionized
	10003920	pH
	11003919	Calcium, Ionized, pH adjusted

Test 903444	Drug Abuse Screen 7, Serum
Effective:	5/24/2021
Specimen:	10 mL frozen serum from a plain red-top tube (5 mL min).
Stability:	Room temperature: Unacceptable
-	Refrigerated: Unacceptable
	Frozen: 30 Days

Test 10352	Growth Hormone Antibody
Effective:	5/17/2021
Stability:	Room temperature: 7 Days Refrigerated: 14 Days Frozen: 28 Days



NEW PDM ASSAYS:

Test 907236	PDM, Buprenorphine and	I Naloxone, w/Conf, w/medMATCH, UR	
Effective:	5/17/2021		
Specimen:	20 mL refrigerated random	urine in a sterile screw-cap container (7 mL min).	
Reference Ranges:	See Report		
Stability:	Room temperature: 7 Days	3	
	Refrigerated: 14 Days		
	Frozen: 30 Days		
Method:	Immunoassay/Mass Spect	rometry	
Setup:	Days: Monday – Saturday		
Reports:	3-5 Days		
CPT*:	80307		
Price:	Client: \$95.00 Patient: \$1	15.00	
Interface Mapping:	Result Code	Result Name	
	15906316	Buprenorphine Screen	
	16906316	medMATCH Buprenorphine	
	17906316	Buprenorphine	
	18906316	medMATCH Buprenorphine	
	19906316	Norbuprenorphine	
	20906316	medMATCH Norbuprenorphine	
	11907218	Naloxone	
	11907191	medMATCH Naloxone	
	12907191	Buprenorphine Comments	
	10907229	Prescribed Drugs, medMATCH	

Test 907239	PDM, Buprenorphine and	PDM, Buprenorphine and Naloxone, Quant, w/medMATCH, Ur		
Effective:	5/17/2021			
Specimen:	20 mL refrigerated random	urine in a sterile screw-cap container (5 mL min).		
Reference Ranges:	See Report			
Stability:	Room temperature: 7 Days			
	Refrigerated: 14 Days			
	Frozen: 30 Days			
Method:	Immunoassay/Mass Spect	Immunoassay/Mass Spectrometry		
Setup:	Days: Monday – Saturday	Days: Monday – Saturday		
Reports:	3-5 Days	3-5 Days		
CPT*:	80348, 80362			
Price:	Client: \$125.00 Patient: \$151.50			
Interface Mapping:	Result Code	Result Name		
	10906286	Buprenorphine		
	11906286	medMATCH Buprenorphine		
	12906286	Norbuprenorphine		
	13906286	medMATCH Norbuprenorphine		
	10907191	Naloxone		
	11907191	medMATCH Naloxone		
	12907191	Buprenorphine Comments		
	10907229	Prescribed Drugs, medMATCH		

Test 907235	PDM Eszopiclone, Quant	PDM Eszopiclone, Quantitative, w/medMATCH, Urine	
Effective:	5/17/2021		
Specimen:	1 mL refrigerated random (urine in a sterile screw-cap container (0.5 mL min).	
Reference Ranges:	See Report		
Stability:	Room temperature: 48 Hou	urs	
	Refrigerated: 30 Days		
	Frozen: 30 Days		
Method:	Immunoassay/Mass Spect	Immunoassay/Mass Spectrometry	
Setup:	Days: Monday – Saturday	Days: Monday – Saturday	
Reports:	3-5 Days		
CPT*:	80368		
Price:	Client: \$125.00 Patient: \$	151.50	
Interface Mapping:	Result Code	Result Name	
	10907181	Eszopiclone	
	12907181	medMATCH Eszopiclone	
	11907181	Eszopiclone Metabolite	
	13907181	medMATCH Eszopiclone Met	
	14907181	Eszopiclone Comments	
	10907229	Prescribed Drugs, medMATCH	

Test 907237	PDM Mitragynine Screen	PDM Mitragynine Screen w/Rflx Confirmation, w/medMATCH	
Effective:	5/17/2021	5/17/2021	
Specimen:	2 mL refrigerated random	urine in a sterile screw-cap container (1 mL min).	
Reference Ranges:	See Report		
Stability:	Room temperature: 7 Days	3	
	Refrigerated: 14 Days		
	Frozen: 30 Days		
Method:	Immunoassay/Mass Spect	Immunoassay/Mass Spectrometry	
Setup:	Days: Tuesday Thursday,	Days: Tuesday Thursday, & Saturday	
Reports:	5-7 Days		
CPT*:	80307	80307	
Price:	Client: \$95.00 Patient: \$1	15.00	
Interface Mapping:	Result Code	Result Name	
	10907160	Mitragynine	
	12907160	medMATCH Mitragynine	
	11907160	Mitragynine Confirmation	
	13907160	medMATCH Mitragynine Conf.	
	12907161	Mitragynine Comments	
	10907229	Prescribed Drugs, medMATCH	

Test 907238	PDM Mitragynine, Quantitative, w/medMATCH		
Effective:	5/17/2021	5/17/2021	
Specimen:	2 mL refrigerated random	urine in a sterile screw-cap container (1 mL min).	
Reference Ranges:	See Report		
Stability:	Room temperature: 7 Days	S	
	Refrigerated: 21 Days		
	Frozen: 30 Days		
Method:	Immunoassay/Mass Spect	rometry	
Setup:	Days: Tuesday Thursday,	& Saturday	
Reports:	5-7 Days		
CPT*:	80323		
Price:	Client: \$125.00 Patient: \$151.50		
Interface Mapping:	Result Code	Result Name	
	10907161	Mitragynine	
	11907161	medMATCH Mitragynine	
	12907161	Mitragynine Comments	
	10907229	Prescribed Drugs, medMATCH	

Test 907240	PDM, Zolpidem Quantitat	PDM, Zolpidem Quantitative, w/medMATCH, Ur	
Effective:	5/17/2021		
Specimen:	1 mL refrigerated random ι	urine in a sterile screw-cap container.	
Reference Ranges:	See Report		
Stability:	Room temperature: 7 Days	3	
	Refrigerated: 30 Days		
	Frozen: 30 Days		
Method:	Immunoassay/Mass Spect	rometry	
Setup:	Days: Tuesday – Saturday	Days: Tuesday – Saturday	
Reports:	3-5 Days		
CPT*:	80368		
Price:	Client: \$125.00 Patient: \$	151.50	
Interface Mapping:	Result Code	Result Name	
	10906115	Zolpidem	
	11906115	medMATCH Zolpidem	
	20906115	Zolpidem Metabolite	
	12906115	medMATCH Zolpidem Metab	
	13906115	Zolpidem Comments	
	10907229	Prescribed Drugs, medMATCH will need all mapping	

PDM ASSAY CHANGES:

Test 802465	Drug Screen 6-Test, Urin	e w/Reflex Oxycodone Confirm	
Test 801996	Drug Screen 9-Test, Urine w/Reflex Oxycodone Confirm		
Test 801073	Drug Screen 10-Test, Urine w/Reflex Oxycodone Confirm		
Test 102011	Drug Screen 10-Test, Urine w/Reflex Confirm		
Test 906641	Drug Screen 11-Test, Uri		
Test 91149	Oxycodone Screen, Urine	e w/Reflex Confirm	
Effective:	5/17/2021	5/17/2021	
Interface Mapping:	Possible reflex:		
	Result Code	Result Code	
	14906335	Codeine	
	15906335	medMATCH Codeine	
	16906335	Hydrocodone	
	17906335	medMATCH Hydrocodone	
	12906335	Hydromorphone	
	13906335	medMATCH Hydromorphone	
	10906335	Morphine	
	11906335	medMATCH Morphine	
	31906480	Norhydrocodone	
	33906480	medMATCH Norhydrocodone	
	10906487	Opiates Comments	
	32906480	Noroxycodone	
	34906480	medMATCH Noroxycodone	
	10906271	Oxycodone	
	11906271	medMATCH Oxycodone	
	12906271	Oxymorphone	
	13906271	medMATCH Oxymorphone	
	11906487	Oxycodone Comments	



MAY HOLIDAY SCHEDULE

Saturday, May 29, 2021

All Patient Service Centers normally operating on Saturdays will be open.

Monday, May 31, 2021

- Limited Courier Services will be provided for Phoenix and Tucson metro area clients on a WILL-CALL BASIS ONLY. For pick-up, please call 602.685.5052 or 520.886.8101.
- STAT testing will be provided for Phoenix and Tucson metro area clients ONLY. For STAT pick-up, please call 602.685.5052 or 520.886.8101.
- Mobile Diagnostics Services STAT ONLY phlebotomy services will be available where offered.
- Billing Department will be closed.
- Warehouse services will not be available.
- All Patient Service Centers will be closed.

Sonora Quest Laboratories wishes you a safe and happy holiday!



ANNOUNCEMENT: AHCCCS: Referring, Ordering, Prescribing, Attending (ROPA)

Sonora Quest Laboratories has received notification from the Patient Protection and Affordable Care Act (ACA) and the 21st Century Cures Act (Cures) that effective June 1, 2021, all health care providers who refer Medicaid (Arizona Health Care Cost Containment System/AHCCCS) members for services must be registered as AHCCCS providers. AHCCCS calls this initiative, and these providers, "ROPA" (Referring, Ordering, Prescribing, Attending).

This affects the following services for any Medicaid/AHCCCS member:

- An item or service
- Order for non-physician services
- Prescribing medications
- Attend/certify medical necessity for services and/or
- > Taking primary responsibility for the members' medical care

Please ensure to properly enroll as an AHCCCS provider to ensure continued laboratory services for your patients.

For your convenience, AHCCCS offers online enrollment at https://www.azahcccs.gov/PlansProviders/APEP/ProviderEnrollment.html. This website also has AHCCCS contact information for any questions you may have regarding ROPA and their requirements.

ASSAY CHANGES:

Test 903444	Drug Abuse Screen 7, Serum	
Effective:	Immediately	
Comment:	Please remove the below Interface Mapping code.	
Interface Mapping:	Result Code	Result Name
	38903444	Ecgonine Methyl Ester

Test 901713	Lactoferrin, Stool (Qualitative)	
Effective:	6/7/2021	
Comment:	Please remove the below Interface Mapping code.	
Interface Mapping:	Result Code	Result Name
	10004005	Consistency



ANNOUNCEMENT: Sonora Quest Laboratories Cardiovascular Risks and COVID-19 Webinar

Beginning Wednesday, May 19, we invite you to join Robert Sherman, product manager at Sonora Quest Laboratories and Dr. Penn, founder of Cleveland HeartLab, as they discuss cardiovascular disease in times of COVID-19, and how to keep patients healthy by lowering their risks.

View the 14-minute recorded webinar now or at a time that is convenient for you by visiting SonoraQuest.com/CARDIOCOVID.

ANNOUNCEMENT: T-SPOT.TB Testing Holiday Schedule

In observance of the upcoming holidays, we ask that you do not submit samples for Test 906927 - T-SPOT.TB on the days listed below as the performing laboratory, Quest Diagnostics T-SPOT®.TB Lab, will be closed.

- Friday, May 28
- Saturday, May 29
- Sunday, May 30
- Monday, May 31

These samples have critical stability and should only be collected Monday – Friday after 10:00 a.m., and not collected on holidays or weekends.



ANNOUNCEMENT: Exagen Avise Testing Holiday Schedule

In observance of the upcoming holidays, we ask that you do not submit samples for Exagen Avise testing on the days listed below as the performing laboratory will be closed.

- Friday, May 28
- Saturday, May 29
- Sunday, May 30
- Monday, May 31

These samples have critical stability and should only be collected Monday – Friday after 10:00 a.m., and not collected on holidays or weekends.

DISCONTINUED TESTS:

 Test 906840
 Glvantage Hereditary Colorectal Cancer Panel

 Effective:
 Immediately

 Comment:
 Test is being discontinued. Alternative testing is currently being evaluated.

Test 906745	Myvantage™ Hereditary Comprehensive Cancer Panel	
Effective:	Immediately	
Comment:	Test is being discontinued. Alternative testing is currently being evaluated.	

REMINDER: AHCCCS: Referring, Ordering, Prescribing, Attending (ROPA)

Sonora Quest Laboratories has received notification from the Patient Protection and Affordable Care Act (ACA) and the 21st Century Cures Act (Cures) that effective June 1, 2021, all health care providers who refer Medicaid (Arizona Health Care Cost Containment System/AHCCCS) members for services must be registered as AHCCCS providers. AHCCCS calls this initiative, and these providers, "ROPA" (Referring, Ordering, Prescribing, Attending). For more information visit https://providers.sonoraquest.com/provider-resources/ahcccs-ropa-enrollment/.

This affects the following services for any Medicaid/AHCCCS member:

- An item or service
- Order for non-physician services
- Prescribing medications
- Attend/certify medical necessity for services and/or
- Taking primary responsibility for the members' medical care

Please ensure to properly enroll as an AHCCCS provider to ensure continued laboratory services for your patients.

For your convenience, AHCCCS offers online enrollment at https://www.azahcccs.gov/PlansProviders/APEP/ProviderEnrollment.html. This website also has AHCCCS contact information for any questions you may have regarding ROPA and their requirements.



ANNOUNCEMENT: Scheduled Laboratory Information System Downtime

On Sunday, June 13, 2021, Sonora Quest Laboratories will implement a scheduled downtime to upgrade our laboratory information system (LIS). As a result, all internal computer systems will be unavailable beginning at 10 a.m. on Sunday, June 13, until approximately 10 p.m. Sonora Quest is starting the downtime at 10 a.m. to ensure early morning routine reporting is completed prior to resulting being interrupted.

Benefits of this upgrade include:

- Positions Sonora Quest to add new Anatomic Pathology and Genetics functionality including developing discrete reporting of results
- Enables future enhancements to our Patient Service Center and In-Office Phlebotomy systems to streamline processes and improve the patient experience
- Enhances our Application Programming Interface (API) abilities to support the CURES Act requirements and will provide patients with additional access points to their data in the future

During the downtime, our Client Services Representatives will not have access to computerized information or patient results. All reporting will resume by normal processes once our systems are operational. Quanum™ users will have uninterrupted access to patient results that were completed prior to 10 a.m. on Sunday. Quanum will be updated with results completed during the downtime once the version update is completed.

STAT courier services and STAT testing will remain available during this time in areas where these services are available and will be reported manually by phone via our established downtime processes.

Results for specimens submitted in a preservative for microbiology culture (urine, stool, swabs, etc.) may be delayed by as many as 1-2 days beyond the normal result turnaround time. Irretrievable and/or unpreserved specimens (CSF, sterile body fluids, tissues, and sputum) should not be impacted as these cultures will follow normal processing protocols throughout the downtime event.

Results for SARS-CoV-2 RNA (COVID-19), Qualitative, NAAT testing will not be available during this downtime. Results for samples received Saturday and Sunday may be slightly delayed beyond the current turnaround time but are expected to be reported within the published 2-4 day timeframe.

During the downtime, please contact us as follows:

- Client Services: 602.685.5050
- Logistics (for pick-ups including Mobile Diagnostic Services STAT phlebotomy requests): 602.685.5052



ANNOUNCEMENT: Referral Testing Delays

The following tests are sent to Quest Diagnostics and are currently delayed. Please see the individual comments below regarding impact to result turnaround time. Updates will be sent as we receive additional information on this testing.

Test 907232	Chromogranin A, LC/MS/MS, Serum
Comment:	Testing is delayed due to instrumentation issues. Samples are being held frozen until testing resumes. The target resolution date is 6/4/21, however, results may be delayed once testing resumes on the testing backlog.
Test 906531	Liver Fibrosis, FibroTest – ActiTest Panel
Comment:	Result turnaround time may extend past the published timeframe of 2-4 days due to intermittent testing delays. The target resolution date is 6/4/21, however, results may be delayed dependent upon the testing backlog.
Test 18786	Kappa/Lambda Light Chains, Total, Serum
Comment:	Result turnaround time may extend past the published timeframe of 3-6 days due to intermittent testing delays. The target resolution date is 6/4/21, however, results may be delayed dependent upon the testing backlog.
Test 904253	Vitamin D, 1,25-Dihydroxy, LC/MS/MS
Comment:	Result turnaround time is expected to take an additional 5 days beyond the published timeframe of 4 days due to intermittent testing delays. The target resolution date is 6/30/21.



NEW ASSAYS:

Test 907220	FISH: ABL1 Gene Rearrangement	
Effective:	6/7/2021	
Specimen:	5 mL room temperature bone marrow in a green-top (Na hep) tube (3 mL min).	
Reference Ranges:	See Report	
Stability:	Room temperature: 7 Days	
	Refrigerated: 7 Days	
	Frozen: Unacceptable	
Method:	Fluorescence In Situ Hybridization (FISH)	
Setup:	Days: Sunday – Friday	
Reports:	1-3 Days	
CPT*:	88271 (x2), 88275, 88291	
Price:	Client: \$429.96 Patient: \$429.96	
Interface Mapping:	Result Code Result Name	
	90722001 Results:	
	90722002 Interpretation:	
	90722003 Comments:	
	90722004 Cytogenetics Director:	
	50905837 Pathologist:	

Test 907224	FISH: ABL2 Gene Rearrangement		
Effective:	6/7/2021		
Specimen:	5 mL room temperatur	re bone marrow in a green-top (Na hep) tube (3 mL min).	
Reference Ranges:	See Report		
Stability:	Room temperature: 7	Days	
	Refrigerated: 7 Days	Refrigerated: 7 Days	
	Frozen: Unacceptable		
Method:	Fluorescence In Situ Hybridization (FISH)		
Setup:	Days: Sunday – Friday		
Reports:	1-3 Days		
CPT*:	88271 (x2), 88275, 88291		
Price:	Client: \$429.96		
Interface Mapping:	Result Code	Result Name	
	90722401	Results:	
	90722402	Interpretation:	
	90722403	Comments:	
	90722404	Cytogenetics Director:	
	50905837	Pathologist:	

Test 907225	FISH: MYH11/CBFB Rearrangement	
Effective:	6/7/2021	
Specimen:	5 mL room temperatur	re bone marrow in a green-top (Na hep) tube (3 mL min).
Reference Ranges:	See Report	
Stability:	Room temperature: 7	Days
	Refrigerated: 7 Days	
	Frozen: Unacceptable	
Method:	Fluorescence In Situ Hybridization (FISH)	
Setup:	Days: Sunday – Friday	
Reports:	1-3 Days	
CPT*:	88271 (x2), 88275, 88291	
Price:	Client: \$429.96 Patient: \$429.96	
Interface Mapping:	Result Code	Result Name
	90722501	Results:
	90722502	Interpretation:
	90722503	Comments:
	90722504	Cytogenetics Director:
	50905837	Pathologist:

Test 907221	FISH: NUP98 Gene R	FISH: NUP98 Gene Rearrangement	
Effective:	6/7/2021	6/7/2021	
Specimen:	5 mL room temperatur	re bone marrow in a green-top (Na hep) tube (3 mL min).	
Reference Ranges:	See Report		
Stability:	Room temperature: 7	Days	
	Refrigerated: 7 Days		
	Frozen: Unacceptable		
Method:	Fluorescence In Situ I	Fluorescence In Situ Hybridization (FISH)	
Setup:	Days: Sunday – Frida	Days: Sunday – Friday	
Reports:	1-3 Days	1-3 Days	
CPT*:	88271 (x2), 88275, 88	88271 (x2), 88275, 88291	
Price:	Client: \$429.96 Pation	Client: \$429.96 Patient: \$429.96	
Interface Mapping:	Result Code	Result Name	
	90722101	Results:	
	90722102	Interpretation:	
	90722103	Comments:	
	90722104	Cytogenetics Director:	
	50905837	Pathologist:	



DISCONTINUED TESTS:

Test 900933	Honey (Rf247), IgE
Effective:	6/21/2021
Comment:	Test is being discontinued. There is no alternative testing at this time.

Test 709332	Cholinesterase, RBC
Effective:	6/28/2021
Comment:	Test is being discontinued. There is no alternative testing at this time.

ASSAY CHANGES:

Test 907134	Cardio IQ Troponin T, High Sensitivity
Effective:	6/28/2021
Reference Range:	Males: ≤19 ng/L
_	Females: ≤11 ng/L
	Relative Risk:
	Female:
	Moderate: 6-11
	High: >11
	Male:
	Moderate: 6-19
	High: >19

Test 907083	Troponin T, High Sensitive
Effective:	6/28/2021
Reference Range:	Males: ≤19 ng/L
_	Females: ≤11 ng/L

Test 803137	Maternal Serum Screen 1			
Effective:	6/21/2021	6/21/2021		
Interfacing Mapping:	Ask at order entry question			
	Result Code	Result Name	Response Options	
	10003715	Pregnancy Result of IVF?	Yes/No	
	10003716	Retrieval Date	mm/dd/yyyy	
	10003717	Transfer Date	mm/dd/yyyy	

Test 803138	Maternal Serum Screen 3		
Effective:	6/21/2021		
Interfacing Mapping:	Ask at order entry question		
	Result Code Result Name Response Options		
	10003715	Pregnancy Result of IVF?	Yes/No
	10003716	Retrieval Date	mm/dd/yyyy
	10003717	Transfer Date	mm/dd/yyyy

Test 803139	Maternal Serum Screen 4			
Effective:	6/21/2021	6/21/2021		
Interfacing Mapping:	Ask at order entry question			
	Result Code Result Name Response Options			
	10003715	Pregnancy Result of IVF?	Yes/No	
	10003716	Retrieval Date	mm/dd/yyyy	
	10003717	Transfer Date	mm/dd/yyyy	



DISCONTINUED TEST:

Test 2096	17-Hydroxycorticosteroids w/Creatinine, 24 Hr Urine	
Effective:	Immediately	
Comment:	Test is being discontinued. Please see recommended alternative testing below.	

RECOMMENDED ALTERNATIVE:

Test 8004	Cortisol, Free, LC/MS/MS, 24-Hour Urine	
Interface Mapping:	Result Code	Result Name
	10009308	Cortisol, Free, Urine
	10717343	Creatinine, 24-Hour Urine

DISCONTINUED TEST:

Test 9330	Cholinesterase, Plasma
Effective:	6/28/2021
Comment:	Test is being discontinued. There is no alternative testing at this time.



ANNOUNCEMENT: Scheduled Application Server Maintenance

On Sunday, June 27, 2021, Sonora Quest Laboratories will be performing application server maintenance. As a result, all computer systems will be unavailable beginning at 6:00 p.m. until approximately 11:00 p.m.

During this period, our Client Services Representatives will not have access to computerized information or patient results. All reporting will resume by normal processes once our systems are operational. Quanum users will have uninterrupted access to patient results that were completed prior to 6:00 p.m. on Sunday. Quanum will be updated with results completed during the downtime once the server maintenance is completed.

STAT courier services and STAT testing will remain available during this time and will be reported manually by phone via our established downtime processes.

During the downtime, please contact us as follows:

- Client Services: 602.685.5050
- Logistics (for pick-ups including Mobile Diagnostic Services STAT phlebotomy requests): 602.685.5052



JULY HOLIDAY SCHEDULE

Saturday, July 3, 2021

• All Patient Service Centers normally operating on Saturdays will be open.

Sunday, July 4, 2021

Mobile Diagnostics Services STAT ONLY phlebotomy services will be available where offered.

Monday, July 5, 2021

- Limited Courier Services will be provided for Phoenix and Tucson metro area clients on a WILL-CALL BASIS ONLY. For pick-up, please call 602.685.5052 or 520.886.8101.
- STAT testing will be provided for Phoenix and Tucson metro area clients ONLY. For STAT pick-up, please call 602.685.5052 or 520.886.8101.
- Mobile Diagnostics Services business as usual
- Billing Department will be closed.
- Fulfillment Center/Warehouse services will not be available. Orders received after 2 p.m. on Thursday, July 1, will not be entered and
 processed until Tuesday, July 6, to allow sufficient time for order entry and processing. Slight delays may occur during the week following
 the closure.
- All Patient Service Centers will be closed.

T-SPOT.TB Testing Holiday Schedule

In observance of the upcoming holidays, we ask that you do not submit samples for Test 906927 - T-SPOT.TB on the days listed below as the performing laboratory, Quest Diagnostics T-SPOT®.TB Lab, will be closed.

- Friday, July 2
- Saturday, July 3
- Sunday, July 4
- Monday, July 5

These samples have critical stability and should only be collected Monday – Friday after 10:00 a.m., and not collected on holidays or weekends.

Exagen Avise Testing Holiday Schedule

In observance of the upcoming holidays, we ask that you do not submit samples for Exagen Avise testing on the days listed below as the performing laboratory will be closed.

- Friday, July 2
- Saturday, July 3
- Sunday, July 4
- Monday, July 5

These samples have critical stability and should only be collected Monday – Friday after 10:00 a.m., and not collected on holidays or weekends.

Sonora Quest Laboratories wishes you a safe and happy holiday!



ANNOUNCEMENT: Cardio IQ F2 Isoprostane/Creatinine Testing Delay

Due to reagent unavailability at the referral laboratory, testing for Cardio IQ F2 Isoprostane/Creatinine (test code 906899) is currently delayed. Specimens will be frozen until reagent is received, which is expected to arrive mid to late July.

REMINDER: Lock Boxes and High Temperatures

With the summer heat in full swing, placing two frozen ice packs in your lock box will ensure **refrigerated** specimens stay cool and viable for testing. Using paper to separate the refrigerated specimens from the ice packs will keep the specimens from freezing.

Additionally, please remember to never leave **frozen** samples in your lock box unless they are stored in Sonora Quest Laboratories' ConstanTemp Frozen Specimen Totes. These totes can be requested by contacting our Logistics Department at 602.685.5052 or 520.886.8101 and can also be ordered through our Provider Portal at SonoraQuest.com.



UPDATE: Cardio IQ F2 Isoprostane/Creatinine Testing Delay

As announced in Client Gram volume 42, due to reagent unavailability at the referral laboratory, testing for Cardio IQ F2 Isoprostane/Creatinine (test code 906899) is delayed. During the backorder period all outstanding testing will be cancelled. Reports will be released with amended results once testing resumes, which is expected to be mid to late July.

ANNOUNCEMENT: Urine Preservative Tubes for Chemistry and Toxicology/Drug Screen Testing

To improve efficiency and quality through our automation capabilities, we have made a change to the following supplies for urine chemistry and toxicology/drug screen testing:

Discontinued Supplies	New Supplies	
Supply #42782 - 12 mL Conical Bottom Skirted Polypropylene Urine Tube (No Cap)	Supply #23247 – 10 mL Round Bottom Urine Tube	BD Vacutainer An to Addition And The Addition
Supply #42783 - Yellow Stopper/Cap	Supply #10845 – Urine Transfer Straw	

Please order the new supplies for requests moving forward. You may continue to use any of the discontinued supplies that you may have remaining for urine chemistry and toxicology/drug screen testing.

Supply orders can be made through our SonoraQuest.com Provider Portal, our Quanum™ system, or by faxing a client supply form to our warehouse. For updated client supply forms, please call 602.685.5141.

For questions regarding supplies please contact our warehouse at the following numbers:

Phoenix: Phone (602) 685.5264; Fax (602) 685.5402 Toll-free (800) 766.6721, ext. 5264

Tucson: Phone (520) 784.8004; Fax (520) 296.5607 Toll-free (800) 266.8101



UPDATE: Scheduled Laboratory Information System Update

On Sunday, June 13, 2021, Sonora Quest Laboratories implemented a scheduled downtime to upgrade our laboratory information system (LIS).

We are continuing to work an identified issue impacting reporting. The software update has altered the font size and formatting on some test results. This issue is being addressed by our software vendor, but there is no ETA for a resolution at this time. Results can still be received by calling Client Services at 602.685.5050 or by filling out the fax request form located at https://providers.sonoraquest.com/provider-resources/laboratory-reference-materials/.

Thank you for your patience as we continue to focus on resolving this issue. We will continue to update you on our progression. If you are experiencing any issues not covered above, please contact your Account Manager, or our Sales Support Department at 602.685.5285 or by email at SQLMarketing@SonoraQuest.com.

ASSAY CHANGES:

Test 902968	First Trimester Screen, hCG		
Effective:	Immediately	Immediately	
Comment:	Please remove th	Please remove the below Interface Mapping code.	
Interface Mapping:	Result Code	Result Code Result Name	
	55127600	Insulin Dependent Diabetic	

Test 5687	HTLV I/II Antibody, w/Reflex Confirmatory Assay
Effective:	7/19/2021
Specimen:	1 mL refrigerated serum from a serum separator tube (SST) (0.5 mL min).

REMINDER: Lock Boxes and High Temperatures

With the summer heat in full swing, placing two frozen ice packs in your lock box will ensure **refrigerated** specimens stay cool and viable for testing. Using paper to separate the refrigerated specimens from the ice packs will keep the specimens from freezing.

When submitting samples for Test 906927 - T-SPOT.TB lock box usage is not recommended, but if necessary, configure the ice packs in an A-Frame or Lean-to formation. Place in a separate bag.

Additionally, please remember to never leave **frozen** samples in your lock box unless they are stored in Sonora Quest Laboratories' ConstanTemp Frozen Specimen Totes. These totes can be requested by contacting our Logistics Department at 602.685.5052 or 520.886.8101 and can also be ordered through our Provider Portal at SonoraQuest.com.



ANNOUNCEMENT: UnitedHealthcare Non-Contracted Network Plans

As of June 2021, we expanded our prior authorization support services to include genetic and molecular testing for UnitedHealthcare Community Plan and UnitedHealthcare Community Plan Medicare. Currently, Sonora Quest Laboratories is not contracted with these payers.

Providers requesting laboratory testing for UnitedHealthcare Community Plan and UnitedHealthcare Community Plan Medicare should be directing their patients to the contracted laboratory for the patient's plan. Please note that Sonora Quest Laboratories will not perform any testing without receipt of a prior authorization. If testing is received without receipt of a prior authorization, the patient will be contacted, and Discounted Pay at Time of Service pricing will be offered. Please visit https://www.sonoraquest.com/patient/knowledge-center/discounted-pay-at-time-of-service-rates/ for a list of tests and pricing.

UPDATE: UnitedHealthcare Genetic and Molecular Prior Authorization/Notification Program

As of June 2021, UnitedHealthcare will manage prior authorizations for genetic and molecular tests. Beacon Laboratory Benefit Solutions, Inc. (Beacon LBS) will no longer manage the prior authorizations for genetic and molecular testing. Ordering providers are still required to obtain prior authorizations with UnitedHealthcare by phone at 877.303.7736 or online at www.UHCprovider.com.

In addition, UnitedHealthcare has added these additional tests to their Genetic and Molecular Prior Authorization/Notification Program:

- Test 906966 JAK2, Exon 12 Mutation Analysis CPT Code 81279
- Test 906965 MPL Mutation Analysis CPT Code 81339

If you have any questions regarding the information listed above, please contact our Prior Authorization Department at 866.202.9181. For a list of UnitedHealthcare impacted tests, please visit https://www.sonoraquest.com/prior-authorization/.



ASSAY CHANGE: T-Spot® TB Now Being Performed at Our Main Laboratory in Phoenix

Benefits of this test and testing being performed at our main laboratory include:

- Timely: Performing this test locally decreases turnaround time by 1 to 2 days
- Reliable: Performing this test locally reduces risk of cancellation due to short sample stability and transportation
- Convenient: Room temperature single tube collection vs. 4 tubes for the QuantiFERON-TB Gold Plus test
- Accuracy: Higher sensitivity in specific populations, such as children under the age of 2, as well as those with compromised immune systems

Test 906927	T-Spot® TB	
Effective:	7/19/2021	
Specimen:	Adults and children 10 years and older: 9 mL lithium heparin tube (6 mL minimum). The use of smaller volume lithium heparin tubes is acceptable. Children 2-9 years: 4 mL minimum in 1 or more lithium heparin tubes. Children <2 years: 2 mL minimum in 1 or more lithium heparin tubes. Critical stability. Due to short specimen stability, specimens must be collected Monday through Friday. Large weekend collections must be coordinated with your Sales Representative prior to occurrence. Apply the neon green T-Spot label provided with the tubes to the outside of the specimen transport bag. Do not refrigerate or freeze. Do not spin or centrifuge samples. Lockbox usage is not recommended, but if necessary, configure the ice packs in an A-Frame or Lean-to formation.	
Reference Ranges:	Negative	
Stability:	Room temperature: Refrigerated: Unacc Frozen: Unacceptab	eptable
Method:	Enzyme Linked Immunospot	
Setup:	Days: Tuesday – Saturday Nights: Monday – Friday	
CPT*:	86481	
Reports:	2-4 Days	
Interface Mapping:	Result Code 50907231	Result Name Interpretation
Comment:	Testing is now performed at our main laboratory in Phoenix.	



NEW ASSAY: Malaria Rapid EIA w/ Blood Parasite smears

Included in this panel is rapid antigen testing for malaria and the microscopic examination of whole blood for parasites.

Malaria rapid antigen test: This test detects a *Plasmodium falciparum* specific antigen and a pan-malarial antigen, common to *P. vivax, P. ovale,* and *P. malariae*. The assay is intended to aid in the rapid diagnosis of human malaria infections and to assist in the differential diagnosis of *P. falciparum* infections from other less virulent malarial infections.

Microscopic examination: Thick and thin blood smears will be prepared from whole blood and examined using a 100x oil immersion objective to determine the presence or absence of blood parasites (i.e., *Plasmodium, Babesia,* microfilaria species, trypanosomes, etc.).

Test 907074	Malaria Rapid EIA	w/ Blood Parasite smears		
Effective:	7/21/2021			
Specimen:	4.0 mL refrigerated whole blood in lavender top (EDTA) tube. Optimal specimen is drawn during febrile episodes. Anti-coagulated specimen tube should be filled to their entirety, to ensure no anti-coagulant dilution of parasites. Ordering Physician to provide direct phone number in case the pathologist needs to speak with them.			
Reference Ranges:	Negative			
Stability:	Room temperature: 3 Days Refrigerated: 3 Days Frozen: Information not available			
Method:		Immunoassay, Microscopy		
Setup:	Days, Evenings, and Nights: Monday – Sunday			
Reports:	1-3 Days			
CPT*:	87899, 87207, and 87015			
Price:	Client: \$175.00 Patient: \$200.00			
Interface Mapping:	Result Code Result Name 10907250 Malaria Rapid EIA Test 10008167 Blood Parasites 11003162 Parasitemia 21008167 Path Review 31008167 CDC Report			
	Ask at order entry question			
	Result Code 99907250	Result Name Ordering physician direct phone number	Response Options Free Text	
Comment:	Test will be available STAT in the Phoenix Metro area but may have a greater than 4-hour STAT turnaround time.			



DISCONTINUED TEST:

Test 907200	SARS-CoV Serology (Covid 19) Ab (IgG,IgM) Immunoassay	
Effective:	7/26/2021	
Comment:	Testing is being discontinued. Please see below for the recommended alternative testing.	

What's the Right Test for You?

Patient history	Expected Result*		
-	SARS-CoV-2 IgG Antibody, Spike	SARS-CoV-2 IgG Antibody, Nucleocapsid	
Previous COVID-19 Infection	Positive	Positive	
COVID-19 Vaccination	Positive	Negative	

^{*} It is recommended to wait at least 10-14 days after potential exposure or onset of symptoms, or vaccination to allow for the development of IgG antibodies. The IgG antibody test options provide insight into an individual's IgG response to the SARS-CoV-2 virus or COVID-19 vaccination, but these tests are not intended to diagnose an active infection or determine whether someone has immunity to COVID-19.

RECOMMENDED ALTERNATIVES:

Test 907234	SARS-CoV-2 Ab IgG Nucleocapsid, QL		
Specimen:	1 mL refrigerated serum from a serum separator tube (SST) (0.1 mL min).		
Comment:	This assay for Nuc	say for Nucleocapsid proteins is helpful in determining if the patient has developed an IgG	
	response to a SAF	to a SARS-CoV-2 infection (COVID-19).	
Interface Mapping:	Result Code	Result Name	
	10907234	SARS-CoV-2 Ab IgG Nucleocar	osid, QL
	Ask at order entry	y question	
	Result Code	Result Name	Response Options
	80907080	Patient Symptomatic	Yes/No/Unknown/No (Pre procedure test)
	99981230	County of Residence	Free Text
	99981231	Patient Race	White/Black or African American/American
			Indian or Alaska Native/Asian Native
			Hawaiian or Other Pacific
			Islander/Unknown
	99981232	Ethnicity	Hispanic/Non-Hispanic /Unknown
	99981233	Pregnant?	Yes/No/Unknown/Not Applicable
	99981235	First Test?	Yes/No/Unknown
	99981236	Employed in Healthcare?	Yes/No/Unknown
	99981238	If yes, date of symptom onset	Date in format (dd/mm/yy)
	99981239	Hospitalized?	Yes/No/Unknown
	99981240	ICU?	Yes/No/Unknown
	99981241	Congregate care setting?	Yes/No/Unknown
	99981242	Is Patient Uninsured?	Yes/No/Unknown
	99981243	ID Type	SSN/DL/State ID/Refused

Test 907097	SARS-CoV-2 Ant	ibody (IgG), Spike, Semi-Quanti	tative	
Specimen:	1 mL refrigerated	1 mL refrigerated serum from a serum separator tube (SST) (0.5 mL min).		
Comment:	antibodies to the S measurement of Iç to a SARS-CoV-2 of an individual's i threshold that con	The SARS-CoV-2 IgG assay is intended for qualitative and semi-quantitative detection of IgG antibodies to the S1 receptor binding domain (RBD) of the SARS-CoV-2 spike protein. The measurement of IgG levels can provide insight to an individual's adaptive immune response to a SARS-CoV-2 infection or vaccination. Although the assay is designed to assess the level of an individual's immune response, studies are still needed to determine the index level threshold that confers protective immunity as well as how long the adaptive immune response may last post-infection or via vaccination.		
	This test should not be used to diagnose or exclude an acute SARS-CoV-2 infection. If acute infection is suspected, direct testing by molecular methods for SARS-CoV-2 is necessary. False positive results for the test may occur due to cross-reactivity from pre-existing antibodies or other possible causes.			
Interface Mapping:	Result Code	Result Name		
	10907097	SARS-CoV-2 lgG, Result		
	20907097	SARS-CoV-2 IgG, Interpretation	SARS-CoV-2 IgG, Interpretation	
	Ask at order entr	Ask at order entry question		
	Result Code	Result Name	Response Options	
	80907080	Patient Symptomatic	Yes/No/Unknown/No (Pre procedure test)	
	99981238	If yes, date of symptom onset	Date in format (dd/mm/yy)	
	99981235	First Test?	Yes/No/Unknown	
	99981236	Employed in Healthcare?	Yes/No/Unknown	
	99981239	Hospitalized?	Yes/No/Unknown	
	99981240	ICU?	Yes/No/Unknown	
	99981241	Congregate care setting?	Yes/No/Unknown	
	99981233	Pregnant?	Yes/No/Unknown/Not Applicable	
	99981231	Patient Race	White/Black or African American/American	
			Indian or Alaska Native/Asian Native	
			Hawaiian or Other Pacific	
			Islander/Unknown	
	99981232	Ethnicity	Hispanic/Non-Hispanic /Unknown	
	99981230	County of Residence	Free Text	
	99981242	Is Patient Uninsured?	Yes/No/Unknown	
	99981243	ID Type	SSN/DL/State ID/Refused	



ANNOUNCEMENT: Scheduled Application Server Maintenance

On Sunday, July 25, 2021, Sonora Quest Laboratories will be performing application server maintenance. As a result, all computer systems will be unavailable beginning at 6:00 p.m. until approximately 11:00 p.m.

During this period, our Client Services Representatives will not have access to computerized information or patient results. All reporting will resume by normal processes once our systems are operational. Quanum users will have uninterrupted access to patient results that were completed prior to 6:00 p.m. on Sunday. Quanum will be updated with results completed during the downtime once the server maintenance is completed.

STAT courier services and STAT testing will remain available during this time and will be reported manually by phone via our established downtime processes.

During the downtime, please contact us as follows:

- Client Services: 602.685.5050
- Logistics (for pick-ups including Mobile Diagnostic Services STAT phlebotomy requests): 602.685.5052

UPDATE: Provider Portal Available at Providers. Sonora Quest.com

Please note that our Provider Portal is now accessed at Providers. Sonora Quest.com.

Visit <u>Providers.SonoraQuest.com</u> today and register for an account which will allow you to:

- Order supplies we make it easy with images for each supply item and the ability to save your commonly ordered supplies for ease-of-use when reordering
- Securely view and download detailed client invoices and make payments
- Order lab requisitions and educational patient brochures for your office
- View laboratory testing updates and sign up to receive them via email to ensure you are always current with specimen requirements, new testing, and billing changes
- Submit demographic changes to your account, such as change of address or phone/fax number

Please visit <u>Providers.SonoraQuest.com</u> today to take advantage of these complimentary services available through our Provider Portal.



ANNOUNCEMENT: Urinalysis Testing Delays

Due to instrumentation issues at our main laboratory in Phoenix, the testing below is currently experiencing delays. Parts have been ordered and we expect to be fully operational by Monday, July 26. Specimens will be kept refrigerated for stability.

Please Note: This only affects testing performed at our main laboratory in Phoenix.

Test Code	Test Name
3320	pH, Urine
104391	Protein, Qualitative, Urine
3315	Specific Gravity, Urine
203302	Urinalysis
203405	Urinalysis Complete
3360	Urinalysis Microscopic
233405	Urinalysis with Reflex to Culture
243405	Urinalysis with Reflex to Culture (OB)
800411	Urinalysis, Diabetic, w/Reflex to Urine Albumin
7093	Urobilinogen, Urine Qualitative
3305	Urogram
23305	Urogram with Reflex to Culture
33305	Urogram with Reflex to Culture (OB)
3300	Urogram with Reflex to Microscopic
43305	Urogram with Reflex to Microscopic and Culture
53305	Urogram with Reflex to Microscopic and Culture (OB)



UPDATE: Urinalysis Testing Delays Resolved

As stated in Client Gram vol. 50, due to instrumentation issues at our main laboratory in Phoenix, the testing below was experiencing delays. Instrumentation has been fixed and we are fully operational.

Test Code	Test Name
3320	pH, Urine
104391	Protein, Qualitative, Urine
3315	Specific Gravity, Urine
203302	Urinalysis
203405	Urinalysis Complete
3360	Urinalysis Microscopic
233405	Urinalysis with Reflex to Culture
243405	Urinalysis with Reflex to Culture (OB)
800411	Urinalysis, Diabetic, w/Reflex to Urine Albumin
7093	Urobilinogen, Urine Qualitative
3305	Urogram
23305	Urogram with Reflex to Culture
33305	Urogram with Reflex to Culture (OB)
3300	Urogram with Reflex to Microscopic
43305	Urogram with Reflex to Microscopic and Culture
53305	Urogram with Reflex to Microscopic and Culture (OB)