

Update Notes

Update Summary

Update Existing Test	12/14/2020	UPBGQ - "Porphobilinogen, 24 Hr Urine"
Update Existing Test	12/14/2020	UPBGR - "Porphobilinogen, Rand Ur"
Inactivate Test With Replacement	1/26/2021	ASGPP - "Aspergillus IgG Precipitins Panel" replaced by ASIGP - "Aspergillus IgG Precipitins"
Inactivate Test Without Replacement	12/18/2020	GLUTA - "Glutathione"

Update Existing Test

Effective Date	12/14/2020
Name	Porphobilinogen, 24 Hr Urine
Code	UPBGQ
Interface Order Code	3426500
Legacy Code	UPBG24Q
Notes	Updated specimen requirements, stability, methodology, reference range, days performed and TAT

Required Testing Changes

Specimen Required	Collect 24 hour urine in a screw-capped plastic container, PROTECTED FROM LIGHT . Add 5 g sodium carbonate (Na ₂ CO ₃) at the start of the collection to maintain a pH of 6 - 7. Refrigerate during collection. Send 2.0 mL preserved urine (1.0 minimum). Provide 24 hour urine volume.
Alternate Specimen	24 hour urine (no preservative) in screw-capped container. Send 2.0 mL urine (1.0 mL minimum) refrigerated. PROTECT FROM LIGHT.
Rejection Criteria	Received room temperature, not protected from light
Stability	Room temperature: Unacceptable; Refrigerated: 7 days; Frozen: 30 days
Methodology	Chromatography/Mass Spectrometry
Reference Range	24 hour Porphobilinogen: Adult: <0.34 mg/24 hr Pediatric: Not Established
Performed Days	Tuesday, Thursday, Saturday
Turnaround Time	5 - 8 days

Update Existing Test	
Effective Date	12/14/2020
Name	Porphobilinogen, Rand Ur
Code	UPBGR
Interface Order Code	3426600
Legacy Code	UPBGRQ
Notes	Updated specimen requirements, stability, methodology and reference range.
Required Testing Changes	
Specimen Required	Collect random urine (no preservative) in a screw-capped plastic container, PROTECTED FROM LIGHT . Send 2.0 mL urine (1.0 mL minimum) refrigerated.
Alternate Specimen	Random clean catch urine collected with 0.5 g sodium carbonate (Na ₂ CO ₃) per 100 mL of sample to maintain a pH of 6 - 7. Send 2.0 mL urine (1.0 mL minimum) refrigerated. PROTECT FROM LIGHT.
Stability	Room temperature: Unacceptable; Refrigerated: 7 days; Frozen: 30 days
Methodology	Chromatography/Mass Spectrometry
Reference Range	Adult: <0.22 mg/g creat Pediatric: <0.36 mg/g creat

Inactivate Test With Replacement			
Effective Date	1/26/2021		
Inactivated Test			
Name	Aspergillus IgG Precipitins Panel		
Code	ASGPP		
Legacy Code¹	ASGPP		
Interface Order Code	3350670		
Notes			
Replacement Test			
Name	Aspergillus IgG Precipitins		
Code	ASIGP		
CPT Code(s)	86331 x6		
Notes			
Specimen Requirements			
Specimen Required	Draw blood in a SST. Centrifuge, remove serum from cells and send 1.0 mL serum (0.5 mL minimum) refrigerated in a screw-capped plastic vial.		
Stability	Room temperature: 4 weeks; Refrigerated: 4 weeks; Frozen: 4 weeks		
Performing Information			
Methodology	Ouchterlony Gel Immunodiffusion		
Reference Range	See report		
Performed Days	Tuesday, Friday		
Turnaround Time	6 - 8 days		
Performing Laboratory	Viracor Eurofins		
Interface Information			
Legacy Code¹	ASIGP		
Interface Order Code	3300182		
Result Code	Name	LOINC Code	AOE/Prompt ²
3300183	Aspergillus amstelodami/glaucus Gel Diffusion*	43823-4	No
3300184	Aspergillus flavus Gel Diffusion*	23820-4	No
3300185	Aspergillus fumigatus Mix Gel Diffusion*	23821-2	No
3300186	Aspergillus nidulans Gel Diffusion*	24508-4	No
3300187	Aspergillus niger Gel Diffusion*	10894-4	No
3300188	Aspergillus versicolor Gel Diffusion*	43911-7	No



LABORATORY REPORT

Example Client, XYZ123
1234 Warde Road
ANN ARBOR MI 48108

EXAMPLE, REPORT
WX0000085261 F 12/23/1970 49 Y

Referral Testing

Collected: 12/11/2020 16:17 Received: 12/11/2020 16:17

Table with 6 columns: Test Name, Result, Flag, Ref-Ranges, Units, Site. Contains Aspergillus IgG Precipitins Panel results.

The gel diffusion method was used to test this patient's serum for the presence of precipitating antibodies(IgG) to the antigens indicated. These antibodies are serological markers for exposure and immunological sensitization. The clinical significance varies, depending on the history and symptoms. *This test was developed and its performance characteristics determined by Viracor Eurofins. It has not been cleared or approved by the U.S. Food and Drug Administration.

Testing Performed At:
Viracor Eurofins
1001 NW Technology Drive
Lee's Summit, MO 64086
(800) 305-5198
Lab Director: Michelle Altrich PhD HCLD(ABB)
CLIA# 26D-0983643

LAB: L - LOW, H - HIGH, AB - ABNORMAL, C - CRITICAL, . - NOT TESTED

Inactivate Test Without Replacement	
Effective Date	12/18/2020
Name	Glutathione
Code	GLUTA
Legacy Code	GLUTA
Interface Code	3724260
Notes	