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500 Chipeta Way, Salt Lake City, Utah 84108-1221 phone: 801-583-2787, toll free: 800-522-2787 Jonathan R. Genzen, MD, PhD, Chief Medical Officer

Patient Age/Sex: 53 years Male

Specimen	Collected:	11-Mar-24	08:56
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Porphyrins and Porphobilinogen	Received: 11-Mar	-24 09:02 Repo	ort/Verified: 11-Mar-24 09:05
Procedure	Result	Units	Reference Interval
Hours Collected	24 ^{f1}	hr	
Total Volume	800	mL	
Creatinine,Urine -per volume	150	mg/dL	
Creatinine,Urine -per 24h	1200	mg/d	[800-2100]
Uroporphyrin -ratio to CRT	2	umol/mol CRT	[0-4]
Heptacarboxylate -ratio to CRT	2	umol/mol CRT	[0-2]
Coproporphyrin I -ratio to CRT	2	umol/mol CRT	[0-6]
Coproporphyrin III -ratio to C	RT 2	umol/mol CRT	[0-14]
Porphyrin Urine Interpretation	Negative ⁱ¹		
Porphobilinogen (PBG),Urine -p	er 2.0 ⁱ²	umol/L	
volume			
Porphobilinogen (PBG),Urine -p	er 1.6 ^H	umol/d	[0.4-1.5]
24h			
Porphobilinogen,Urine -ratio t	o 0.2	mmol/mol CRT	[0.0-0.2]
CRT			
Porphobilinogen Interpretation	See Note ⁱ³		

Result Footnote

fl: Hours Collected

Per 24h calculations are provided to aid interpretation for collections with a duration of 24 hours and an average daily urine volume. For specimens with notable deviations in collection time or volume, ratios of analytes to a corresponding urine creatinine concentration may assist in result interpretation.

Test Information

il: Porphyrin Urine Interpretation

INTERPRETIVE INFORMATION: Porphyrins, Fractionation and Quantitation, Urine

Results are normalized to creatinine concentration and reported as a ratio of amounts (micromoles of porphyrin/moles of creatinine).

This test was developed and its performance characteristics determined by ARUP Laboratories. It has not been cleared or approved by the US Food and Drug Administration. This test was performed in a CLIA certified laboratory and is intended for clinical purposes.

- i2: Porphobilinogen (PBG), Urine -per volume
- i3: Porphobilinogen Interpretation

INTERPRETIVE INFORMATION: Porphobilinogen Interpretation

Porphobilinogen (PBG), Urine Results for random urine specimens are normalized to creatinine (CRT) concentration and reported as a ratio of amounts (millimoles of PBG/mole of creatinine).Porphobilinogen (PBG) in a random urine specimen is used to evaluate an

*=Abnormal, #=Corrected, C=Critical, f=Result Footnote, H-High, i-Test Information, L-Low, t-Interpretive Text, @=Performing lab

Unless otherwise indicated, testing performed at: ARUP Laboratories 500 Chipeta Way, Salt Lake City, UT 84108 Laboratory Director: Jonathan R. Genzen, MD, PhD
 ARUP Accession:
 24-071-900020

 Report Request ID:
 19127599

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Jonathan R. Genzen, MD, PhD, Chief Medical Officer

Patient Age/Sex: 53 years Male

Test Information

i3: Porphobilinogen Interpretation

attack of acute porphyria. Slight increases in urinary PBG are associated with acute porphyrias other than acute intermittent porphyria (AIP) and may indicate a resolving or treated acute porphyria. Urinary PBG in excess of two times the upper reference limit is consistent with acute porphyria.

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