GENETIC TESTING LABORATORIES Ш

Biochemical Genetics Laboratory

ered By Patient Name: Test, urineAA				
Accession #: R5003	Specimen #: X5003			
	Specimen: Urine			
Birthdate: 08/05/2019	Age: 1			
Gender: Male				
MRN #: 151911	Collected: 08/04/2020			
Ethnicity: Caucasian	Received: 08/05/2020			
	Patient Name: Test, urineAA Accession #: R5003 Birthdate: 08/05/2019 Gender: Male MRN #: 151911 Ethnicity: Caucasian			

Urine Amino Acid Analysis - Quantitative

RESULTS

ANALYTE	REFERENCE RANGE*	RESULT*	FLAG	ANALYTE	REFERENCE RANGE*	RESULT*	FLAG
3-Methyl-histidine	0-682	18		Glycine	0-9207	71	
Alanine	0-2090	5		Guanidinoacetate	30-1200	75.9	
Alloisoleucine	0-25	10		Histidine	0-3879	81	
Alpha-aminoadipate	0-516	15		Homocitrulline	0-174	86	
Alpha-amino-n-butyrate	0-106	20		Homocystine	0-7	91.0	н
Anserine	0-820	3		Hydroxyproline	0-525	96	
Arginine	0-262	25		Isoleucine	0-100	101	н
Argininosuccinate	0-61	30		Leucine	0-269	106	
Asparagine	0-970	35		Lysine	0-666	111	
Aspartate	0-308	40		Methionine	0-69	116	н
Beta-alanine	0-496	0		Ornithine	0-119	121	н
Beta-Aminolsobutyrate	0-1742	491		Phenylalanine	0-326	126	
Citrulline	0-123	46		Proline	0-517	137	
Creatine/Creatinine Ratio	0-1.55	0.46		Sarcosine	0-103	142	н
Creatinine	5.8-85.8	19.8		Serine	0-2249	147	
Cystathionine	0-159	20		Sulfocysteine	0-87	2000	н
Cystine	0-212	15		Taurine	0-3852	152	
Delta-aminolevulinate	0-42	13		Threonine	0-953	157	
Gamma-amino-n-butyrate	0-43	30		Tryptophan	0-321	162	
Glutamate	0-376	61		Tyrosine	0-509	167	
Glutamine	0-3112	66		Valine	0-254	172	

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Test, urineAA

08/05/2020

*Values in micromols/g creatinine

*Creatinine value in mg/dl

*Creatine/Creatinine value in mol/mol ratio

INTERPRETATION

Mock Report

ASSAY INFORMATION

Method

Liquid chromatography tandem mass spectrometry (LC-MS/MS)

Limitations/Disclaimer

False negative results can occur in rare situations when diet and/or clinical condition masks or normalizes disease relevant analyte perturbations. In addition, false negatives may occur when disease presentation is intermittent or the result of a mild defect. Results should always be viewed in the context of clinical presentation and concurrent laboratory studies.

This test was developed and its performance characteristics determined by Indiana University Biochemical Genetics Laboratory. It has not been cleared or approved by the U.S. Food and Drug Administration. This test is used for clinical purposes. It should not be regarded as investigational or for research. The laboratory is certified under the Clinical Laboratory Improvement Amendments of 1988 (CLIA '88) as qualified to perform high complexity clinical laboratory testing. CLIA# 15D0647198 • CAP# 1678930

ELECTRONICALLY SIGNED BY

Marcus J. Miller, Director of the Biochemical Genetics Laboratory, 08/05/2020

