

A Description of Reference Ranges for Organic Acids in Urine Samples from A Pediatric Population in Iran

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Abstract

Background: Organic acids refer to a family of compounds that are intermediates in a variety of metabolic pathways. Many organic acids are present in urine from clinically normal individuals. Elevated levels of urine organic acids cause to the organic acidurias, disorders in which some metabolic pathways in organic acid metabolism are blocked. The present work identified major and minor urinary acidic metabolites in normal subjects, and their quantitative ranges in a pediatric population of Iran.

Methods: Two hundred and fifty-one healthy subjects, including 132 males and 119 females, from 2 days to 15 years of age were enrolled. Urinary organic acids were extracted from urine with organic solvents and identified and quantified by gas chromatography-mass spectrometry.

Results: The results provide a foundation on which to check results for patients with potentially abnormal organic acidurias. By this method 98 organic acids were identified in a pediatric population of Iran.

Conclusions: The present work identifies and quantifies major and minor urinary metabolites excreted by normal subjects. We also analyzed urine from 30 patients with organic acid metabolism abnormalities and compared the concentrations of specific organic acids with those from urines of normal individuals.

Keywords: Gas chromatography/mass spectrometry, Iran population, Normal individuals, Urine organic acid analysis, Urine organic acids range

Introduction

Approximately 50 diseases have been described in which an inherited single enzyme defect causes a high concentration of acidic metabolites in the blood or urine. Dysfunction in any protein complex that involves the absorption, transportation, activation, and/or application of a vitamin can result in an elevated urinary organic acid (1). With this in mind, organic acid concentrations in urine can serve as markers to diagnose the numerous genetic metabolic disorders known as organic acidurias (2). Organic acids refer to a family of compounds that are intermediates in a variety of metabolic pathways

including glycolysis and citric acid cycle metabolites, fatty acid oxidation, ketone metabolites and cofactors, and markers of detoxification (3, 4). Many organic acids are present in urine from clinically normal individuals (5). Elevated levels of urine organic acids cause to the organic acidurias, disorders in which some metabolic pathways in organic acid metabolism are blocked. An analysis of urinary organic acids is usually a key test in the assessment of patients with doubtful genetic disorders of organic acid metabolism and is frequently used in the diagnosis of persons with possible genetic disorders of

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Received: Jul30, 2016; Accepted: Aug28, 2016

mitochondrial fatty acid metabolism, amino acid disorders, and disorders of mitochondrial oxidative phosphorylation. Disorders of organic acid metabolism have various clinical presentations and ages of onset. Some of the common presentations include lethargy, fasting intolerance, myalgia, myopathy, exercise intolerance, cardiomyopathy, and various types of developmental or neurological disabilities such as seizures or vision deficits (6, 7).

Gas Chromatography/Mass Spectrometry (GC/MS) is the most reliable method for urine organic acid analysis (8). Its primary asset is that it allows precise and simultaneous quantification of different compounds in samples. It has contributed greatly to the understanding of many disease states. GC/MS was initially developed for qualitative purposes, and quantitative analytical criteria have seldom been extensively studied, if at all (9-11). Clearly, an encyclopedic analysis such as this compounds the potential information on the physiological and pathophysiological situation of different metabolic pathways and their interrelationships in the body, and may provide clinical relevant information for patient care (12, 13).

Knowledge of biological variation allows evaluation of population-based reference ranges. For example, it would be useful to know that a result found in a patient would likely differ from that of a healthy individual. The present work identified major and minor urinary acidic metabolites in normal subjects, and their quantitative ranges in a pediatric population of Iran. This could provide age-dependent reference intervals for laboratory staff and physicians using current methods and equipment, as currently available data is rather dated.

Materials and Methods

Subjects

Two hundred and fifty-one healthy subjects, including 132 males and 119 females, from 2 days to 15 years of age were enrolled. All of them were referred to our laboratory for organic acid analysis, and their results were normal. All procedures followed were in accordance with the ethical standards of the committee on human experimentation MUMS (Mashhad University of Medical Science). Informed consent was obtained from all patients for being included in the study.

Chemical, Reagents and instrument

In this study chemicals and reagents with highest purity available were purchased. Pentadecanoic acid (PDA) and O-(2,3,4,5,6-Pentafluorobenzyl) hydroxylamine hydrochloride (PFBH) were obtained from Sigma-Aldrich. Tetramethylsilane (TMS) Tri-Sil (BSA: pyridine) from Thermo Scientific was used as the derivatizing reagent. Ethyl acetate and n-Hexane was from Merck Millipore. We use a Clarus 500 GC-MS from PerkinElmer with an Agilent fused silica column 30 m x 0.25 mm x 0.25 μ m.

Extraction, Derivatization, and Gas chromatography

The organic acids were extracted from urine by organic solvent extraction and then identified and quantified by GC/MS. Initially, the urinary pH was adjusted to 2 to 5, 50 μ l of 50 mg/ml pentafluoro benzyl hydroxylamine-hydrochloride solution were added, and the sample was incubated at room temperature for one hour. Subsequently, the pH was adjusted to 1 by dropwise addition of HCl. The acidic sample was extracted two times with 4 ml of ethyl acetate by vortexing the sample and centrifuging at 700 x g for 5 min. The organic supernatant layers from each extraction were transferred to a second vial, 25 μ l of pentadecanoic acid (PDA) used as an internal standard were added, and the solution was evaporated to dryness under nitrogen at 40 $^{\circ}$ C. Finally, the sample was silylated by adding 100 μ l of tetramethylsilane (TMS), vortexed, and incubated at 60 $^{\circ}$ C for 30 min. After the sample cooled to room temperature, 500 μ l of hexane were added and the sample was analyzed by GC/MS. The temperatures for the injectors and detectors were 250 and 300 $^{\circ}$ C, respectively. The GC temperature program was as follows: initial temperature was 70 $^{\circ}$ C, held for 4 min, increased to 180 $^{\circ}$ C at a rate of 20 $^{\circ}$ C/min, then to 200 $^{\circ}$ C at a rate of 4 $^{\circ}$ C/min, held for one min and finally to 275 $^{\circ}$ C at a rate of 3 $^{\circ}$ C/min and held for 10 min (8, 14).

Statistical Analysis

Statistical analysis was performed using SPSS version 16. The compounds were identified and their concentration ranges were determined for all individuals.

Results

Analysis of normal urines

We identified 98 compounds in urine samples from 251 healthy subjects. Compounds detected in 60-80% of the samples were Aconitic Acid, Adipic Acid, Succinic Acid, Citric Acid, 2,3-Dihydroxy Butyric Acid, 3,4-Dihydroxy Butyric Acid, Ethylmalonic Acid, Glycolic Acid, Hippuric Acid, Homovanillic Acid, 4-Hydroxy Benzoic Acid, 3-Hydroxy Isobutyric Acid, 2-Hydroxy Glutaric Acid, 4-Hydroxy Phenyl Acetic Acid, 3-Methyl Adipinic Acid, Methyl Succinic Acid, Methylmalonic Acid, 2-Methyl-3-Hydroxy Butyric Acid, Octadecarboxylic Acid, Oxalic Acid, 2-Oxoglutaric Acid, 5-Oxoproline, Pimelic Acid, Suberic Acid, VanillylMandelic Acid, Lactic Acid and Pyruvate. Compounds detected in 20-60% of the samples included Malic Acid, Benzoic Acid, Decadienic Dicarboxylic Acid, Azelaic Acid, 2,4-Dihydroxy Butyric Acid, 2-Ethyl-3-hydroxy Propionic Acid, Fumaric Acid, Glutaric Acid, Hydroxy Malic Acid, 3-Hydroxy Adipic Acid, 4-Hydroxy Hippuric Acid, HydroxyDecadecarboxylic Acid, 5-Hydroxy Hexanoic Acid, 3-Hydroxy-3-methyl Glutaric Acid, 3-Hydroxy Phenyl Acetic Acid, 4-Hydroxy Phenyl Lactic Acid, 3-Hydroxy Propionic Acid, (3-hydroxy phenyl)-3-Hydroxy Propionic Acid, 3-Hydroxy Isovaleric Acid, 2-Hydroxy Isovaleric Acid, Furan-2,5-Dicarboxylic Acid, 5-Hydroxymethyl-2-Furanoic Acid, 3-Methyl-4-Hydroxy Benzoic Acid, Phenyl Acetic Acid, PhenylLactic Acid and Phosphoric Acid.

Compounds detected in 0.5-20% of the samples included Carbamazepin, Cyclohexandiol, Decendicarboxylic Acid, Chloralhydrate Glucuronide, 4-Cresol, Erythronic Acid, 2-Furoylglycin, Hexenoic Dicarboxylic Acid, Homogentisic Acid, 2-Hydroxyadipinic Acid, 3-Hydroxy Benzoic Acid, 3-Hydroxy Butyric Acid, 4-Hydroxy Butyric Acid, 2-Hydroxy Butyric Acid, 3-Hydroxy Hippuric Acid, 5-Hydroxy Indole Acetic Acid, 3-Hydroxy Adipic Acid Lacton, 2-Hydroxy-3 methyl-Valeric Acid, 4-Hydroxy Phenyl Pyruvate, 4-Hydroxy Phenyl Propionic Acid, 2-Hydroxy Valeric Acid, Levulinic Acid, Mevalonic, N-acetylaspartate, 3-Methyl Glutaric Acid, 3-(3-methyl-4-hydroxyphenyl) 3-Hydroxy Propionic Acid, 3-Methyl-4-Hydroxy-Phenyl Lactic Acid, 2-Oxo adipinic Acid, 2-Oxo-3-Methylvaleric Acid, 2-Oxoisocaproic Acid, Palmitic Acid, Stearic Acid, Uracil, 4-Hydroxy Cyclohexyl Carbonic Acid, 3-Methyl Glutaconic Acid, Urate, 2-HydroxyIsocaproic Acid, HydroxyDecadecarboxylic Acid, Tricarballic Acid, Tartaric Acid, Malonic Acid, Phenobarbital, Lauric Acid, 4-Hydroxy Cyclo Acetic, Mandelic Acid, and 2-Hydroxy Isobutyric Acid. Frequency, percentage (number of urines in which the compounds were detected), range of concentration, and pathological area of these compounds are listed in Table 1. A typical chromatogram of urinary organic acids from a normal individual is shown in Figure 1.

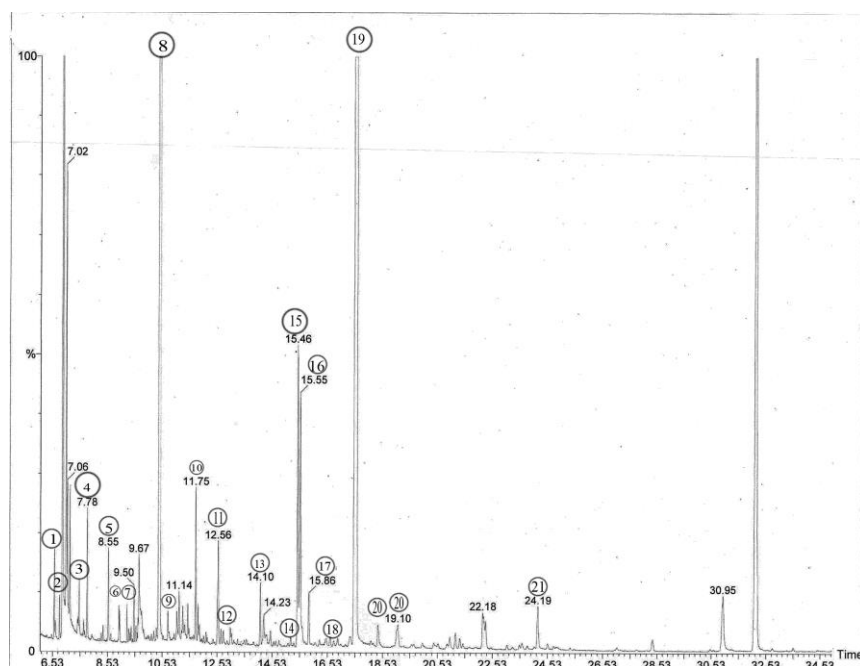


Fig 1. A typical GC-MS chromatogram of urinary organic acids from a normal individual. The compounds noted in the figure are TMS-derivatives of the following compounds: 1: Lactic acid, 2: Glycolic acid, 3: Oxalic acid, 4: 3-hydroxy butyric acid, 5: Urea, 6: Phosphoric acid, 7: Succinic acid, 8: Pentafluorobenzyl hydroxyl amine, 9: Pyruvic acid, 10: 2-hydroxy glutaric acid, 11: 4-hydroxy phenyl acetic acid, 12: Tartaric acid, 13: Aconitic acid, 14: Homovanillic acid, 15: Citric acid, 16: Hippuric acid, 17: 3-(3-hydroxyphenyl)-3-hydroxyl propionic acid, 18: 3-methoxy-4-hydroxyl mandelic acid, 19: Internal standard, 20: 2-oxoglutaric acid, 21: 4-hydroxyl hippuric acid.

Table 1. Frequency, percentage and range of concentration of compounds in normal urine

	OA Name		A R	F	%	RR ($\mu\text{mol}/\text{mmolCreat}$)	PA ($\mu\text{mol}/\text{mmolCreat}$)
1	Aconitic Acid	Yes	2 D-1 Y	110	44	<25	>60
		No	>1 Y	86	34	<17	
2	Adipic Acid	Yes	2 D-1 Y	120	48	<19	> 60
		No	>1 Y	63	25	<9	
3	Malic Acid	Yes	2 D-1 Y	34	13.5	<12	> 30
		No	>1 Y	11	4.5	<9	
4	Benzoic Acid	Yes	2 D-1 Y	31	12.35	<34	> 250
		No	>1 Y	15	5.98	<90	
5	Succinic Acid	Yes	2 D-1 Y	120	47.8	<200	> 500
		No	>1 Y	86	34.26	<83	
6	Carbamazepin	Yes	2 D-1 Y	-	-	-	> 72
		No	>1 Y	4	1.59	<24	
7	Cyclohexandiol	Yes	2 D-1 Y	17	6.77	<13	> 30
		No	>1 Y	2	0.8	<10	
8	DecadienicDicarboxylac Acid	Yes	2 D-1 Y	55	21.9	<10	> 25
		No	>1 Y	20	7.97	<8	
9	Decendicarboxylic Acid	Yes	2 D-1 Y	8	3.19	<12	> 25
		No	>1 Y	2	0.8	<8	
10	Chloralhydrate Glucuronide	Yes	2 D-1 Y	2	0.8	<80	> 400
		No	>1 Y	7	2.79	<236	
11	Citric Acid	Yes	2 D-1 Y	115	45.82	<87	> 250
		No	>1 Y	91	36.25	<72	
12	4- Cresol	Yes	2 D-1 Y	5	1.99	<4	> 15
		No	>1 Y	31	12.35	<10	
13	Azelaic Acid	Yes	2 D-1 Y	41	16.33	<11	> 25
		No	>1 Y	38	15.14	<9	
14	2,3- Dihydroxy Butyric Acid	Yes	2 D-1 Y	92	36.65	<10	> 25
		No	>1 Y	76	30.28	<8	
15	2,4-Dihydroxy Butyric Acid	Yes	2 D-1 Y	58	23.11	<5	> 15
		No	>1 Y	36	14.34	<4	
16	3,4-Dihdroxy Butyric Acid	Yes	2 D-1 Y	104	41.43	<7	> 15
		No	>1 Y	52	20.72	<4	
17	Ethylmalonic Acid	Yes	2 D-1 Y	111	44.22	<17	> 45
		No	>1 Y	81	32.27	<13	
18	Erythronic Acid	Yes	2 D-1 Y	26	10.36	<9	> 15
		No	>1 Y	4	1.59	<5	
19	2-Ethyl-3-hydroxy Propionic Acid	Yes	2 D-1 Y	35	13.94	<12	> 30
		No	>1 Y	16	6.37	<10	
		No		200	79.68		

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20	Fumaric Acid	Yes	2 D-1 Y	56	22.31	< 14	> 30
			>1 Y	30	11.95	< 7	
21	2-Furoylglycin	No		165	65.74		
		Yes	2 D-1 Y	2	0.8	< 4	> 12
			>1 Y	3	1.19	< 4	
22	Glutaric Acid	No		246	98		
		Yes	2 D-1 Y	76	30.28	< 8	> 20
			>1 Y	39	15.54	< 5	
23	Glycolic Acid	No		136	54.18		
		Yes	2 D-1 Y	116	46.25	< 14	> 45
			>1 Y	92	36.65	< 16	
24	Hippuric Acid	No		43	17.13		
		Yes	2 D-1 Y	53	21.11	< 158	> 500
			>1 Y	65	25.9	< 253	
25	Hexenoic Dicarboxylic Acid	No		133	52.99		
		Yes	2 D-1 Y	47	18.72	< 5	> 15
			>1 Y	1	0.399	-	
26	Homogentisic Acid	No		203	80.88		
		Yes	2 D-1 Y	1	0.399	-	> 25
			>1 Y	-	-		
27	Homovanillic Acid	No		250	99.6		
		Yes	2 D-1 Y	124	49.4	< 14	> 25
			>1 Y	91	36.25	< 8	
28	2-Hydroxyadipinic Acid	No		34	13.55		
		Yes	2 D-1 Y	6	2.39	< 5	> 5
			>1 Y	-	-		
29	Hydroxy Malic Acid	No		245	97.6		
		Yes	2 D-1 Y	34	13.55	< 10	> 20
			>1 Y	30	11.95	< 7	
30	3-Hydroxy Adipic Acid	No		187	74.5		
		Yes	2 D-1 Y	91	36.25	< 10	> 15
			>1 Y	14	5.58	< 5	
31	3-Hydroxy Benzoic Acid	No		146	58.17		
		Yes	2 D-1 Y	7	2.79	< 6	> 30
			>1 Y	21	8.37	< 8	
32	4-Hydroxy Benzoic Acid	No		223	88.84		
		Yes	2 D-1 Y	76	30.28	< 12	> 50
			>1 Y	58	23.11	< 16	
33	3-Hydroxy Butyric Acid	No		117	46.61		
		Yes	2 D-1 Y	28	11.15	< 63	> 150
			>1 Y	8	3.19	< 55	
34	4-Hydroxy Butyric Acid	No		215	85.66		
		Yes	2 D-1 Y	1	0.399	-	> 25
			>1 Y	-	-		
35	2-Hydroxy Butyric Acid	No		250	99.6		
		Yes	2 D-1 Y	14	5.58	< 8	> 25
			>1 Y	3	1.19	< 8	
36	3-Hydroxy Isobutyric Acid	No		234	93.23		
		Yes	2 D-1 Y	89	35.46	< 28	> 60
			>1 Y	72	28.68	< 15	
37	3-Hydroxy Hippuric Acid	No		90	35.86		
		Yes	2 D-1 Y	10	3.98	< 11	> 45
			>1 Y	31	12.35	< 18	
38	4-Hydroxy Hippuric Acid	No		210	83.66		
		Yes	2 D-1 Y	50	19.92	< 15	> 50
			>1 Y	45	17.93	< 20	
39	HydroxyDecadicarboxylic Acid	No		156	62.15		
		Yes	2 D-1 Y	63	25.1	< 25	> 50
			>1 Y	10	3.98	< 10	
		No		178	70.92		

40	2-Hydroxy Glutaric Acid	Yes	2 D-1 Y	114	45.42	< 19	> 50
		No	>1 Y	76	30.28	< 8	
41	5-Hydroxy Hexanoic Acid	Yes	2 D-1 Y	44	17.53	< 6	> 15
		No	>1 Y	16	6.37	< 3	
42	5-Hydroxy Indole Acetic Acid	Yes	2 D-1 Y	11	4.38	< 8	> 8
		No	>1 Y	4	1.59	< 6	
43	3-Hydroxy Adipic Acid Lacton	Yes	2 D-1 Y	39	15.54	< 15	> 40
		No	>1 Y	3	1.19	< 7	
44	3-Hydroxy-3-methyl Glutaric Acid	Yes	2 D-1 Y	91	36.25	< 16	> 40
		No	>1 Y	33	13.15	< 6	
45	3-Hydroxy Phenyl Acetic Acid	Yes	2 D-1 Y	6	2.39	< 5	> 30
		No	>1 Y	58	23.11	< 11	
46	4-Hydroxy Phenyl Acetic Acid	Yes	2 D-1 Y	133	52.99	< 104	> 300
		No	>1 Y	100	39.84	< 87	
47	4-Hydroxy Phenyl Lactic Acid	Yes	2 D-1 Y	68	27.09	< 28	> 100
		No	>1 Y	21	8.37	< 8	
48	4-Hydroxy Phenyl Pyruvate	Yes	2 D-1 Y	5	1.99	< 15	> 60
		No	>1 Y	-	-	-	
49	3-Hydroxy Propionic Acid	Yes	2 D-1 Y	27	10.76	< 5	> 15
		No	>1 Y	21	8.37	< 3	
50	3-(3-hydroxy phenyl)-3-Hydroxy Propionic Acid	Yes	2 D-1 Y	27	10.76	< 16	> 100
		No	>1 Y	70	27.89	< 31	
51	4-Hydroxy Phenyl Propionic Acid	Yes	2 D-1 Y	1	0.399	-	> 80
		No	>1 Y	-	-	-	
52	3-Hydroxy Isovaleric Acid	Yes	2 D-1 Y	44	17.53	< 24	> 80
		No	>1 Y	28	11.15	< 12	
53	2-Hydroxy Isovaleric Acid	Yes	2 D-1 Y	44	17.53	< 6	> 20
		No	>1 Y	18	7.17	< 5	
54	2-Hydroxy Valeric Acid	Yes	2 D-1 Y	1	0.399	-	> 20
		No	>1 Y	-	-	-	
55	Furan-2,5-Dicarboxylic Acid	Yes	2 D-1 Y	31	12.35	< 11	> 50
		No	>1 Y	46	18.33	< 11	
56	5-Hydroxymethyl-2-Furanoic Acid	Yes	2 D-1 Y	40	15.94	< 24	> 80
		No	>1 Y	56	22.31	< 21	
57	Levulinic Acid	Yes	2 D-1 Y	2	0.8	< 7	> 7
		No	>1 Y	-	-	-	
58	3-Methyl Adipinic Acid	Yes	2 D-1 Y	110	43.82	< 8	> 25
		No	>1 Y	82	32.67	< 5	
59	Methyl Succinic Acid	Yes	2 D-1 Y	76	30.28	< 6	> 18
		No	>1 Y	69	27.49	< 6	
				106	42.23		

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60	Methylmalonic Acid	Yes	2 D-1 Y	79	31.47	<11	> 40
			>1 Y	50	19.92	<6	
61	Mevalonic	No		122	48.6		
		Yes	2 D-1 Y	1	0.399	-	
62	N-acetylaspartate		>1 Y	-		-	
		No		250	99.61	-	
63	3-Methyl Glutaric Acid	Yes	2 D-1 Y	25	9.96	<5	> 50
			>1 Y	2	0.797	<16	
64	3-(3-methyl-4-hydroxyphenyl) 3-Hydroxy Propionic Acid	No		224	89.24		
		Yes	2 D-1 Y	11	4.38	<4	> 12
65	2-Methyl-3-Hydroxy Butyric Acid		>1 Y	9	3.58	<3	
		No		231	92.04		
66	3-Methyl-4-Hydroxy Benzoic Acid	Yes	2 D-1 Y	4	1.59	<13	> 50
			>1 Y	24	9.56	<9	
67	3-Methyl-4-Hydroxy-Phenyl Lactic Acid	No		223	88.85		
		Yes	2 D-1 Y	81	32.27	<7	> 20
68	3-Methyl-4-Hydroxy Benzoic Acid		>1 Y	59	23.5	<5	
		No		111	44.23		
69	Oxalic Acid	Yes	2 D-1 Y	14	5.58	<29	> 100
			>1 Y	29	11.55	<10	
70	2-Oxoadipinic Acid	No		208	82.87		
		Yes	2 D-1 Y	6	2.39	<5	> 8
71	2-Oxo-3-Methylvaleric Acid		>1 Y	-			
		No		245	97.61		
72	2-Oxoglutaric Acid	Yes	2 D-1 Y	116	46.21	<14	> 40
			>1 Y	52	20.72	<7	
73	2-Oxoisocaproic Acid	No		83	33.07		
		Yes	2 D-1 Y	114	45.42	<30	> 80
74	5-Oxoproline		>1 Y	81	32.27	<18	
		No		56	22.31		
75	Palmitic Acid	Yes	2 D-1 Y	3	1.19	<5	> 25
			>1 Y	3	1.19	<5	
76	Phenyl Acetic Acid	No		245	97.6		
		Yes	2 D-1 Y	3	1.19	<2	> 5
77	Phenyl Lactic Acid		>1 Y	2	0.8	<5	
		No		246	98.01		
78	Pimelic Acid	Yes	2 D-1 Y	96	38.25	<136	> 500
			>1 Y	63	25.1	<46	
79	Stearic Acid	No		92	36.65		
		Yes	2 D-1 Y	2	0.8	<2	> 2
80	Stearic Acid		>1 Y	-			
		No		249	99.2		
81	Stearic Acid	Yes	2 D-1 Y	80	31.87	<9	> 20
			>1 Y	57	22.71	<5	
82	Stearic Acid	No		114	45.42		
		Yes	2 D-1 Y	6	2.39	<5	> 5
83	Stearic Acid		>1 Y	1	0.4		
		No		244	97.21		
84	Stearic Acid	Yes	2 D-1 Y	27	10.76	<11	> 50
			>1 Y	21	8.37	<10	
85	Stearic Acid	No		203	80.88		
		Yes	2 D-1 Y	61	24.31	<6	> 20
86	Stearic Acid		>1 Y	17	6.77	<4	
		No		173	68.92		
87	Stearic Acid	Yes	2 D-1 Y	76	30.28	<5	> 15
			>1 Y	49	19.52	<5	
88	Stearic Acid	No		126	50.2		
		Yes	2 D-1 Y	12	4.78	<7	> 30
89	Stearic Acid		>1 Y	12	4.78	<8	
		No		227	90.44		

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80	Suberic Acid	Yes	2 D-1 Y	103	41.03	<12	> 30
		No	>1 Y	30	11.95	<10	
81	VanillylMandelic Acid	Yes	2 D-1 Y	122	48.6	<12	> 30
		No	>1 Y	69	27.49	<5	
82	Lactic Acid	Yes	2 D-1 Y	90	35.86	<60	> 200
		No	>1 Y	82	32.67	<50	
83	Phosphoric Acid	Yes	2 D-1 Y	38	15.14	<31	> 60
		No	>1 Y	44	17.53	<14	
84	Pyruvate	Yes	2 D-1 Y	169	67.33	<32	> 100
		No	>1 Y	82	32.67	<15	
85	Uracil	Yes	2 D-1 Y	19	7.57	<20	> 50
		No	>1 Y	49	19.52	<10	
86	4-Hydroxy Cyclohexyl Carbonic Acid	Yes	2 D-1 Y	6	2.39	<20	> 30
		No	>1 Y	15	5.98	<18	
87	3-Methyl Glutaconic Acid	Yes	2 D-1 Y	29	11.55	<9	> 30
		No	>1 Y	15	5.98	<14	
88	Urate	Yes	2 D-1 Y	7	2.79	<4	> 8
		No	>1 Y	1	0.4	-	
89	2- Hydroxy Isocaproic Acid	Yes	2 D-1 Y	243	96.81	<3	> 8
		No	>1 Y	2	0.8	<3	
90	HydroxyDecadicarboxylic Acid	Yes	2 D-1 Y	249	99.2	<10	> 10
		No	>1 Y	3	1.19	<10	
91	Tricarballic Acid	Yes	2 D-1 Y	11	4.38	<12	> 30
		No	>1 Y	7	2.79	<4	
92	Tartaric Acid	Yes	2 D-1 Y	233	92.83	<4	> 20
		No	>1 Y	13	5.18	<4	
93	Malonic Acid	Yes	2 D-1 Y	231	92.03	<14	> 14
		No	>1 Y	2	0.8	<14	
94	Phenobarbital	Yes	2 D-1 Y	249	99.2	-	> 20
		No	>1 Y	1	0.4	-	
95	Lauric Acid	Yes	2 D-1 Y	250	99.6	<3	> 3
		No	>1 Y	1	0.4	<3	
96	4-Hydroxy Cyclo Acetic acid	Yes	2 D-1 Y	249	99.6	<3	> 3
		No	>1 Y	2	-	<3	
97	Mandelic Acid	Yes	2 D-1 Y	6	-	<74	> 80
		No	>1 Y	2	-	<64	
98	2-Hydroxy Isobutyric Acid	Yes	2 D-1 Y	243	-	<9	> 20
		No	>1 Y	7	-	<9	

OA Name=Organic Acids Name, AR= Age Range, F=Frequency, %=Percent, RR= Reference Range, PA=Pathgenical Area, Creat=Creatinine, D=Day, Y=Year,

Analysis of abnormal urines

In addition to the urines from healthy individuals, we also analyzed 30 abnormal urines from patients with methylmalonic aciduria, propionic aciduria, ethylmalonic aciduria, glutaric aciduria, maple syrup urine disease (MSUD), and isovaleric aciduria as described above.

Urine from 30 patients with methylmalonic aciduria (MMA, OMIM 251000) contained 35-3717 $\mu\text{mol}/\text{mmol}$ Creatinine of methylmalonic acid, which were 3-330 folds greater than the normal levels found in our study. Also methylcitrate, which was not present in the urine of normal individuals, was found in the MMA patients.

Urine from five patients with propionic aciduria (PA, OMIM 606054) contained 28-115 $\mu\text{mol}/\text{mmol}$ Creatinine of 3-hydroxy propionic acid, which was 5-23 fold greater than that in normal levels.

Urine from three cases with ethylmalonic aciduria contained 65-197 $\mu\text{mol}/\text{mmol}$ Creatinine ethylmalonic acid, which were 3-12 folds greater than normal levels.

Urine from three patients with glutaric aciduria (OMIM 231670) contained 415-2800 $\mu\text{mol}/\text{mmol}$ Creatinine glutaric acid and 7-116 $\mu\text{mol}/\text{mmol}$ Creatinine of 3-Hydroxy glutaric acid. The glutaric acid concentrations 51-350 fold greater than normal levels. 3-hydroxy glutaric acid is not seen in normal individuals.

Urine from the patient with isovaleric aciduria (IVA, OMIM 243500) contained 2556 $\mu\text{mol}/\text{mmol}$ Creatinine of isovaleryl glycine and 162 $\mu\text{mol}/\text{mmol}$ Creatinine of 3-hydroxy isovaleric acid. The concentration of 3-Hydroxy isovaleric acid was 6-12 folds greater than normal. Isovaleryl glycine is not seen in normal individuals.

Urine from three patients with MSUD (OMIM 248600) contained 20-2600 $\mu\text{mol}/\text{mmol}$ Creatinine of 2-hydroxy isovaleric acid, 10-80 $\mu\text{mol}/\text{mmol}$ Creatinine of 2-hydroxy isocaproic acid, and 190-1905 $\mu\text{mol}/\text{mmol}$ Creatinine of 3-hydroxy butyric acid. These values were 3-30 folds greater than normal.

Discussion

Most often, laboratories do not quantify organic acids, or they express semi-quantitative results equivalents of an internal standard. Sweetmann considered that errors in quantitative results as great

as 50% would be acceptable for the diagnosis of inherited disorders but the error for organic acids should be <20%. In our opinion, lower analytical errors are desirable for some differential diagnoses, and for the diagnoses of patients having moderate hyper excretions. Moreover, essential biological variables such as the normal excretion concentrations and their variations according to age, genetic factors, and nutritional status have yet to be unequivocally established (12).

Dagleish et al. proposed the use of gas chromatography for the study of the trimethylsilyl and methyl esters of organic acids extracted from physiological fluids, and reported data on a wide range of synthetic reference compounds including butyric acid; di- and tri-carboxylic acids; hydroxy acids and keto acids; polyhydroxy and alicyclic compounds such as glycerol, inositol, quinic acid, shikimic acid, ascorbic acid, and sugar alcohols; aromatic hydroxy and acidic compounds, both benzenoid and indolic; sesquiterpenes; steroids; glycine conjugates; mercapturic acids; and glucuronides (15).

Tanaka et al. analyzed samples from 50 normal subjects and showed that compounds detectable in essentially all urines are succinic, adipic, p-hydroxyphenylacetic, hippuric, and citric acids (8). They used PDA as a qualitative and quantitative internal standard, as described in this study. The benefit of using pentadecanoic acid as an internal standard is that it elutes later than the organic acids; therefore, it does not overlap with them in the GC/MS elution.

In a similar study, Lawson AM et al. was studied the qualitative pattern of urinary organic acids in normal persons. Oxalic acid, sulfate, benzoic acid, phosphate, succinic acid, 4-deoxytetronic acid, 3-oxo-tetronic acid, 2-deoxytetronic acid, 5-hydroxymethyl-2-furoic acid, tetronic acids, 2-oxoglutaric acid, 4-hydroxyphenylacetic acid, tartaric acid, 2-deoxypentonic acid, aconitic acid, hippuric acid, citric acid, glucono-1,5-lactone, glucuronic acid and gluconic plus glucaric acids were identified in urine from normal persons. In addition, small amounts of lactic, pyruvic, methylmalonic, oxalic, 3-hydroxybutyric, and furan-2,5-dicarboxylic acids were also found (16). This finding is consistent with our study, but we

quantify major and minor urinary metabolites excreted by normal subjects.

Chalmers et al. assessed variations of urinary organic acids excretion in normal persons and the effects of alterations in dietary combination on these metabolites. They found that dietary alterations produced small changes in the excretion patterns of these metabolites. Alteration in ranges of compounds excreted in urine for the normal population depend more on individual metabolic variations. These findings provide a basis for determination of the normal ranges (17).

Christou et al. proposed the use of gas chromatography for the determination of ten free organic acids by GC-MS with the aim to create a method for organic acid profiling in human urine. This method was used to the quantitative analysis of urinary organic acids in hospitalized children (18).

Conclusion

Routine GC-MS is acceptable for quantifying urinary organic acids. With the extraction method described here, urinary organic acids were analyzed both qualitatively and quantitatively. This method also made it possible to establish a more accurate concentration range of organic acids in urine than was

References

1. Kuhara T. Diagnosis and monitoring of inborn errors of metabolism using urease- pretreatment of urine, isotope dilution, and gas chromatography-mass spectrometry. *Chromat.* 2002 Dec; 781(1-2): 497- 517.
2. Greter J, Jacobson CE. Urinary Organic Acids: Isolation and Quantification for Routine Metabolic Screening. *Clin Chem.* 1987 Apr; 33 (4): 473- 80.
3. Song YZ, Li BX, Hao H, Xin RL, Zhang T, Zhang CH, et al. Selective screening for inborn errors of metabolism and secondary methylmalonic aciduria in pregnancy at high risk district of neural tube defects: A human metabolome study by GC-MS in China. *Clin Biochem.* 2008 May; 41 (7-8): 616- 20.
4. Wajner M, Coelho DM, Ingrassia R, Oliveira AB, Busanello ENB, Raymond K, et al. Selective screening for organic acidemias by urine organic acid GC-MS analysis in Brazil: Fifteen-year experience. *Clinica Chimica Acta.* 2009 Feb; 400 (1-2): 77- 81.

previously available. From the above results, we suggest that this efficient method will be suitable for the quantitative analysis of organic acids for diagnoses and follow-up studies of patients with new or ill-defined disorders in most clinical laboratories. For successful quantification, one must ensure a constant fragmentation by use of adequate internal standards and perform external calibration. Generally, isotope standards would be preferable for an exact quantitation but this is not feasible for all acids (19, 20). Therefore, a single standard for diagnostic purposes and follow-up is acceptable. Standardization of this method would allow comparisons of values from different labs.

Our studies showed a diverse range of compounds; in addition to the previously recognized urinary organic acids, most of the compounds identified in our study had not been previously reported in a pediatric population of Iran.

Acknowledgment

This research was supported and funded by the Metabolic Department, Pardis Clinical and Genetic Laboratory, Mashhad, Iran.

All authors declare they have no conflicts of interest.

5. Lord RS, Nelson-Dooley C, Morris C, Bralley JA. Weekly biological variability of urinary organic acids. *N A J Med Sci.* 2012 Jul; 5 (3): 148-56.
6. Verhaghe BJ, Lleverer MF, DeLeenheer AP. Solid-Phase Extraction with Strong Anion-Exchange Columns for Selective Isolation and Concentration of Urinary Organic Acids. *Clin Chem.* 1988 Jun; 34 (6): 1077- 83.
7. Wasant P, Liammongkolkul S, Kuptanon C, Vatanavicharn N, Sathienkijakanchai A, Shinka T. Organic acid disorders detected by urine organic acid analysis: Twelve cases in Thailand over three-year experience. *Clinica Chimica Acta.* 2008 Jun; 392 (1-2): 63- 8.
8. Tanaka K, West-Dull A, Hine DJ, Lynn TB, Lowe T. Gas-Chromatographic Method of Analysis for Urinary Organic Acids. II. Description of the Procedure, and Its Application to Diagnosis of Patients with Organic Acidurias. *Clin Chem.* 1980 Dec; 26 (13): 1847- 53.
9. Duez P, Kumps A, Mardens Y. GC-MS profiling of urinary organic acids evaluated as a

- quantitative method. *Clin Chem.* 1996 Oct; 42 (10): 1609-15.
10. Kumps A, Duez P, Genin J, Mardens Y. Gas Chromatography–Mass Spectrometry Analysis of Organic Acids: Altered Quantitative Response for Aqueous Calibrators and Dilute Urine Specimens. *Clin Chem.* 1999 Aug; 8 (45): 1297- 1300.
11. Suh JW, Lee SH, Chung BC. GC-MS determination of organic acids with solvent extraction after cation-exchange chromatography. *Mol Pathol Genet.* 1997 Dec; 43 (12): 2256- 61.
12. Sweetman L. Organic acid analysis. In: Hommes FA., editors. *Techniques in diagnostic human biochemical genetics. A laboratory manual.* New York, Wiley-Liss, 1991; pp 143-76.
13. Hoffmann G, Aramaki S, Blum-Hoffmann E, Nyhan WL, Sweetman L. Quantitative analysis for organic acids in biological samples: batch isolation followed by gas chromatographic-mass spectrometric analysis. *Clin Chem.* 1989 Apr; 35 (4): 587-95.
14. Rinaldo P. Organic acids. In: Blau N, Duran M, Gibson KM. *Laboratory guide to the methods in biochemical genetics.* Berlin, Heidelberg, Springer, 2008; pp 137- 70.
15. Dagleish CE, Horning EC, Horning MG, Knox KL, Yarger K. A gas-liquid- chromatographic procedure for separating a wide range of metabolites occurring in urine or tissue extracts. *Biochem.* 1966 Dec; 101 (3): 792-810.
16. Lawson AM, Chalmers RA, Watts RWE. Urinary organic acids in man. I. Normal patterns. *Clin Chem.* 1976 Aug; 22 (8): 1283-87.
17. Chalmers RA, Healy MJR, Lawson AM, Watts RWE. Urinary organic acids in man.II. Effects of individual variation and diet on the urinary excretion of acidic metabolites. *Clin Chem.* 1976 Aug; 22 (8): 1288-91.
18. Christou C, Gika HG, Raikos N, Theodoridis G. GC-MS analysis of organic acids in human urine in clinical settings: a study of derivatization and other analytical parameters. *J Chromatogr B Analyt Technol Biomed Life Sci.* 2014 Aug;964 (1): 195-201.
19. Hyman DB, Saunders AM, Tanaka K. A rapid spot test for urinary methylmalonic acid collected on-ion-exchange filter paper. *Clinica ChimicaActa.* 1983 Aug; 132 (3): 219- 27.
20. Inoue Y, Kuhara T. Rapid and sensitive method for prenatal diagnosis of propionic acidemia using stable isotope dilution gas chromatography–mass spectrometry and urease pretreatment. *J Chrom.* 2002 Aug; 776 (1): 71- 7.