



RN LABS 1190885 Requisition #: Practitioner: 05/02/2023 Patient Name: Veyan Saini Date of Collection: 7 Date of Birth: 08/25/2015 Patient Age: Time of Collection: 07:15 AM Patient Sex: Report Date: 05/19/2023



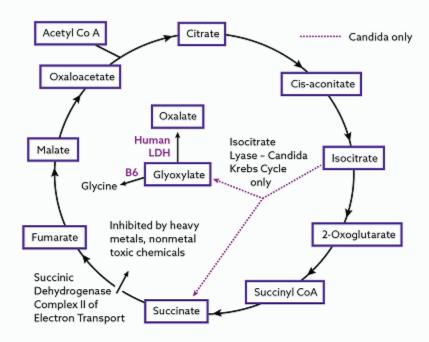
Organic Acids Test - Nutritional and Metabolic Profile

| Metabolic Markers in Urine | Reference Range mmol/mol creatinine) | Patient Value | Reference Population - Males Under Age 13 |
|---|--------------------------------------|------------------|---|
| Intestinal Microbial Overgrow | rth . | | |
| Yeast and Fungal Markers | | | |
| 1 Citramalic | ≤ 5.0 | 0.80 | 0.80 |
| 2 5-Hydroxymethyl-2-furoic (Aspergillus) | ≤ 28 | 1.5 | 1.5 |
| 3 3-Oxoglutaric | ≤ 0.46 | 0.08 | (0.08) |
| 4 Furan-2,5-dicarboxylic (Aspergillus) | ≤ 18 | 2.0 | 2.0 |
| 5 Furancarbonylglycine (Aspergillus) | ≤ 3.1 | 1.6 | (1.6) |
| 6 Tartaric (Aspergillus) | ≤ 6.5 | 1.1 | 1.1 |
| 7 Arabinose | ≤ 50 | 39 | 39 |
| 8 Carboxycitric | ≤ 25 | 0.29 | 0.23 |
| 9 Tricarballylic (Fusarium) | ≤ 1.3 | 0.07 | 0.07 |
| Bacterial Markers | | | |
| 10 Hippuric | ≤ 680 | 54 | 54 |
| 11 2-Hydroxyphenylacetic | ≤ 0.86 | 0.39 | (33) |
| 12 4-Hydroxybenzoic | ≤ 3.0 | 0.88 | (0.88) |
| 13 4-Hydroxyhippuric | ≤ 30 | H 40 | 40> |
| 14 DHPPA (Beneficial Bacteria) | ≤ 0.59 | 0.05 | 0.05 |
| Clostridia Bacterial Markers | | | |
| 15 4-Hydroxyphenylacetic (C. difficile, C. stricklandii, C. lituseburense | 2.0 - 32 e & others) | 22 | 22 |
| 16 HPHPA (C. sporogenes, C. caloritolerans, C. botuli | ≤ 220 inum & others) | 7.2 | 7.2 |
| 17 4-Cresol (C. difficile) | ≤ 84 | 0.67 | (0.67) |
| 18 3-Indoleacetic (C. stricklandii, C. lituseburense, C. subter | 0.60 - 14 minale & others) | L 0.51 | 0.51 |

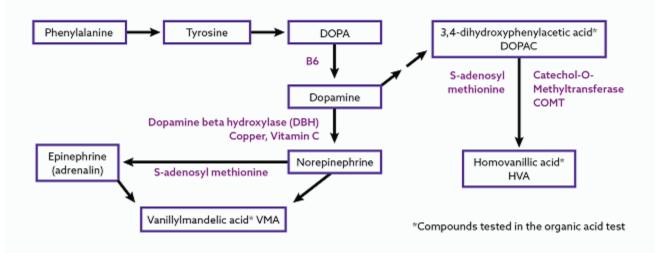
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Human Krebs Cycle showing Candida Krebs Cycle variant that causes excess Oxalate via Glyoxylate



Major pathways in the synthesis and breakdown of **catecholamine neurotransmitters** in the absence of microbial inhibitors



39

Quinolinic

40 Kynurenic

1190885 **RN LABS** Requisition #: Practitioner: Patient Name: Veyan Saini Date of Collection: 05/02/2023 **Metabolic Markers in Urine** Reference Range **Patient** Reference Population - Males Under Age 13 (mmol/mol creatinine) **Value Oxalate Metabolites** 19 Glyceric 0.74 - 13 3.5 3.5 20 Glycolic 27 - 221 97 97 21 Oxalic 35 - 185 88 88 Glycolytic Cycle Metabolites 22 Lactic 2.6 48 19 **(19)** 23 Pyruvic 0.32 - 8.8 2.8 Mitochondrial Markers - Krebs Cycle Metabolites 24 Succinic ≤ 23 5.5 (5.5) 25 Fumaric ≤ 1.8 0.36 0.36 26 Malic ≤ 2.3 0.95 (0.95) 27 2-Oxoglutaric ≤ 96 27 **<27**> 28 Aconitic 9.8 - 39 10.0 29 Citric ≤ 597 189 (189) Mitochondrial Markers - Amino Acid Metabolites 30 3-Methylglutaric 0.01 - 0.97 0.28 0.28 31 3-Hydroxyglutaric ≤ 16 4.3 4.3 32 3-Methylglutaconic ≤ 6.9 1.8 1.8> **Neurotransmitter Metabolites Phenylalanine and Tyrosine Metabolites** 33 Homovanillic (HVA) 0.49 - 13 2.9 (dopamine) 34 Vanillylmandelic (VMA) 0.72 - 6.4 2.7 (2.7) (norepinephrine, epinephrine) 35 HVA / VMA Ratio 0.23 - 2.8 1.1 (1.1) 36 Dihydroxyphenylacetic (DOPAC) 0.13 - 4.9 2.0 $\langle 2.0 \rangle$ 37 HVA/ DOPAC Ratio 0.37 - 3.3 1.4 (1.4) **Tryptophan Metabolites** 38 5-Hydroxyindoleacetic (5-HIAA) ≤ 11 1.8 1.8

(2.2)

3.7

3.7

2.2

0.48

- 8.8

≤ 4.2

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| etabolic Markers in Urine Pyrimidine Metabolites - F | Reference Rang (mmol/mol creation) olate Metabolism | ine) | Patient Value | Reference Population - Males Under Age 13 |
|--|---|--------|------------------|---|
| 1 Uracil | | ≤ 16 | 1.8 | 1.8 |
| 2 Thymine | | ≤ 0.91 | 0.14 | 0.19 |
| Ketone and Fatty Acid Oxi | dation | | | |
| 3 3-Hydroxybutyric | | ≤ 4.8 | 2.2 | 2.2 |
| 4 Acetoacetic | | ≤ 10 | 0.39 | 0.39 |
| 5 Ethylmalonic | 0.06 | - 4.8 | 4.2 | 4.2 |
| 6 Methylsuccinic | | ≤ 4.0 | 2.2 | 2.2 |
| 7 Adipic | 0.19 | - 6.5 | 5.4 | 5.4 |
| 8 Suberic | | ≤ 7.0 | H 7.9 | 7.9 |
| 9 Sebacic | | ≤ 0.61 | 0.16 | (j.16) |
| Nutritional Markers | | | | |
| itamin B12 0 Methylmalonic * | | ≤ 5.2 | 2.9 | 2.9 |
| itamin B6 1 Pyridoxic (B6) | | ≤ 53 | 1.6 | 1.6 |
| itamin B5 2 Pantothenic (B5) | | ≤ 14 | 3.4 | 3.4 |
| itamin B2 (Riboflavin) 3 Glutaric * | | ≤ 1.4 | 0.60 | (0.6) |
| itamin C 4 Ascorbic | 10 | - 200 | L 5.2 | 5.2 |
| tamin Q10 (CoQ10) 5 3-Hydroxy-3-methylglutaric # | | ≤ 88 | 21 | 21 |
| utathione Precursor and Chelati N-Acetylcysteine (NAC) | ng Agent | ≤ 0.34 | 0 | 0.00 |
| otin (Vitamin H) 7 Methylcitric * | | ≤ 5.7 | 1.4 | 1.4 |

^{*} A high value for this marker may indicate a deficiency of this vitamin.

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| Metabolic Markers in Urine | Reference Range (mmol/mol creatinine) | Patient Value | Reference Population - Males Under Age 13 |
|-------------------------------------|---------------------------------------|------------------|---|
| Indicators of Detoxification | n | | |
| Glutathione | | | |
| 58 Pyroglutamic * | 13 - 62 | 34 | 34 |
| Methylation, Toxic exposure | | | |
| 59 2-Hydroxybutyric ** | 0.19 - 2.0 | 1.1 | 1.1 |
| Ammonia Excess | | | |
| 60 Orotic | 0.04 - 0.80 | 0.19 | 0.19 |
| Aspartame, salicylates, or GI bacte | eria | | |
| 61 2-Hydroxyhippuric | ≤ 1.2 | H 1.3 | 1.3 |

^{*} A high value for this marker may indicate a Glutathione deficiency.

Amino Acid Metabolites 62 2-Hydroxyisovaleric ≤ 2.0 0.19 63 2-Oxoisovaleric ≤ 2.5 0.08 64 3-Methyl-2-oxovaleric ≤ 2.0 0.74 0.74 65 2-Hydroxyisocaproic ≤ 2.0 0 66 2-Oxoisocaproic ≤ 2.0 0.15 67 2-Oxo-4-methiolbutyric ≤ 2.0 0.27 Mandelic ≤ 2.0 68 0.17 **Phenyllactic** ≤ 2.0 69 0.12 ≤ 4.0 Phenylpyruvic 0 70 0.00 Homogentisic ≤ 2.0 0.11 4-Hydroxyphenyllactic ≤ 2.0 0.96 (0.96) N-Acetylaspartic ≤ 38 2.5 74 Malonic ≤ 18 7.3 75 4-Hydroxybutyric ≤ 4.7 3.1

Mineral Metabolism

76 Phosphoric 1,000 - 7,300 2,097

^{**} High values may indicate methylation defects and/or toxic exposures.

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Indicator of Fluid Intake

77 *Creatinine 82 mg/dL

*The creatinine test is performed to adjust metabolic marker results for differences in fluid intake. Urinary creatinine has limited diagnostic value due to variability as a result of recent fluid intake. Samples are rejected if creatinine is below 20 mg/dL unless the client requests results knowing of our rejection criteria.

Explanation of Report Format

The reference ranges for organic acids were established using samples collected from typical individuals of all ages with no known physiological or psychological disorders. The ranges were determined by calculating the mean and standard deviation (SD) and are defined as ± 2SD of the mean. Reference ranges are age and gender specific, consisting of Male Adult (≥13 years), Female Adult (≥13 years), Male Child (<13 years), and Female Child (<13 years).

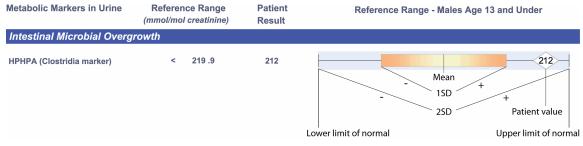
There are two types of graphical representations of patient values found in the new report format of both the standard Organic Acids Test and the Microbial Organic Acids Test.

The first graph will occur when the value of the patient is within the reference (normal) range, defined as the mean plus or minus two standard deviations.

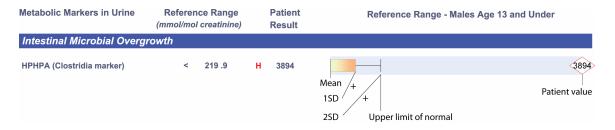
The second graph will occur when the value of the patient exceeds the upper limit of normal. In such cases, the graphical reference range is "shrunk" so that the degree of abnormality can be appreciated at a glance. In this case, the lower limits of normal are not shown, only the upper limit of normal is shown.

In both cases, the value of the patient is given to the left of the graph and is repeated on the graph inside a diamond. If the value is within the normal range, the diamond will be outlined in black. If the value is high or low, the diamond will be outlined in red.

Example of Value Within Reference Range



Example of Elevated Value



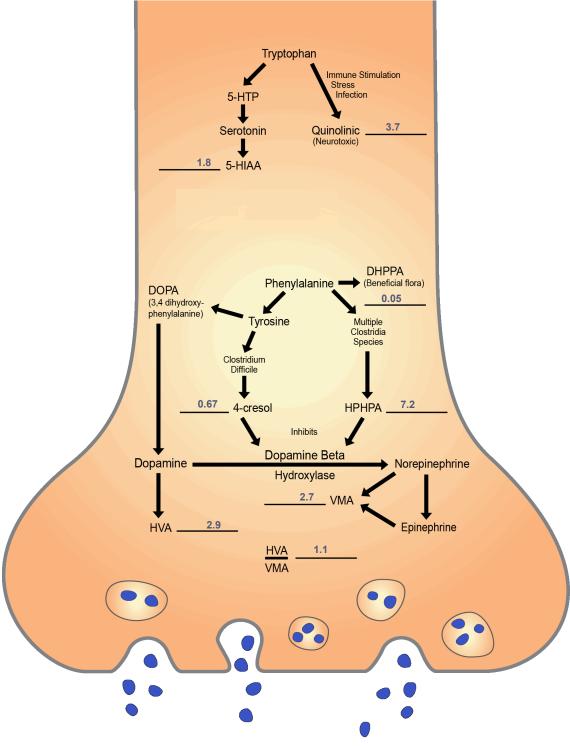
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Neurotransmitter Metabolism Markers



The diagram contains the patient's test results for neurotransmitter metabolites and shows their relationship with key biochemical pathways within the axon terminal of nerve cells. The effect of microbial byproducts on the blockage of the conversion of dopamine to norepinephrine is also indicated.

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Interpretation

High 4-hydroxybenzoic acid and/or 4-hydroxyhippuric acid (12,13) may be due to bacterial overgrowth of the GI tract, intake of fruits such as blueberries rich in polyphenols (anthocyanins, flavonols, and hydroxycinnamates), or may be from paraben additive exposure. Parabens are 4-hydroxybenzoic acid alkyl esters with antimicrobial properties.

4-Hydroxybenzoic acid may be excreted as its glycine conjugate 4-hydroxyhippuric acid. High levels of these paraben metabolites in urine (>10 mmol /mol creatinine) may result from excessive exposure to parabens. Parabens are common preservatives allowed in foods, drugs, cosmetics and toiletries, but they also have a long history of use in a variety of pharmaceutical products for injection, inhalation, oral, topical, rectal or vaginal administration. Some individuals experience skin reactions as most parabens are readily and completely absorbed through the skin and the GI tract. Parabens have been considered safe because of their low toxicity profile and their long history of safe use; however, recent studies challenge this view. In 1998, Routledge et.al., (Toxicol.Appl.Pharmacol. 153,12-19), reported parabens having estrogenic activity in vitro. A number of in vivo studies have further elucidated potential endocrine disruption by parabens affecting reproduction or promote tumor growth. Parabens have been found at high levels in breast cancer biopsies, although a definitive relationship with breast cancer has not been demonstrated. Parabens may contribute to mitochondrial failure by uncoupling oxidative phosphorylation and depleting cellular ATP. 4-Hydroxyhippuric acid has been found to be an inhibitor of Ca2+-ATPase in end-stage renal failure. Eliminate all sources of parabens. To accelerate paraben excretion, use sauna therapy, the Hubbard detoxification protocol employing niacin supplementation, or glutathione supplementation (oral, intravenous, transdermal, or precursors such as N-acetyl cysteine [NAC]).

Low or low normal citric acid (29) may be due to impaired function of the Krebs cycle, low dietary intake of citrate-containing foods such as citrus fruits and juices, potassium deficiency, acidosis (especially renal tubular acidosis), chronic kidney failure, diabetes, hypoparathyroidism, or excessive muscle activity. Low values may indicate increased risk of oxalate kidney stone formation, especially if oxalic acid is elevated also. Supplement with calcium or magnesium citrate if oxalic acid is elevated.

Homovanillic acid (HVA) levels (33) below the mean indicate low production and/or decreased metabolism of the neurotransmitter dopamine. Homovanillic acid is a metabolite of the neurotransmitter dopamine. Low production of HVA can be due to decreased intake or absorption of dopamine's precursor amino acids such as phenylalanine and/or tyrosine, decreased quantities of cofactors needed for biosynthesis of dopamine such as tetrahydrobiopterin and vitamin B6 coenzyme or decreased amounts of cofactors such as S-adenosylmethionine (Sam-e) needed to convert dopamine to HVA. In addition, a number of genetic variations such as single nucleotide polymorphisms (SNPs) or mutations can cause reduced production of HVA due to enzymes with decreased function. HVA values below the mean but which are much higher than VMA values are usually due to impairment of dopamine beta hydroxylase due to excessive Clostridia metabolites, the mold metabolite fusaric acid, pharmaceuticals such as disulfiram, or food additives like aspartame or deficiencies of cofactors such as vitamin C or copper. Values may also be decreased in patients on monoamine oxidase (MAO) inhibitors. In addition, a number of genetic variations such as single nucleotide polymorphisms (SNPs) or mutations in MAO or COMT genes can cause reduced production of HVA. Such SNPs are available on The Great Plains DNA methylation pathway test which can be performed on a cheek swab.

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VanillyImandelic acid (VMA) levels (34) below the mean indicate low production and/or decreased metabolism of the neurotransmitters norepinephrine and epinephrine. VanillyImandelic acid is a metabolite of the neurotransmitters norepinephrine and epinephrine. Low production of VMA can be due to decreased intake or absorption of norepinephrine's and epinephrine's precursor amino acids such as phenylalanine and/or tyrosine, decreased quantities of cofactors needed for biosynthesis of norepinephrine and epinephrine such as tetrahydrobiopterin and vitamin B6 coenzyme or decreased amounts of cofactors such as S-adenosylmethionine (Sam-e) needed to convert norepinephrine and epinephrine to VMA. In addition, a number of genetic variations such as single nucleotide polymorphisms (SNPs) or mutations in MAO or COMT genes can cause reduced production of VMA. Such SNPs are available on The Great Plains DNA methylation pathway test which can be performed on a cheek swab. VMA values below the mean but which are much lower than HVA values are usually due to impairment of dopamine beta hydroxylase due to Clostridia metabolites, the mold metabolite fusaric acid, pharmaceuticals such as disulfiram, or food additives like aspartame or deficiencies of cofactors such as vitamin C or copper. Values may be decreased in patients on monoamine oxidase (MAO) inhibitors. Another cause for a low VMA value is a genetic variation (single nucleotide polymorphism or SNP) of the DBH enzyme. Patients with low VMA due to Clostridia metabolites or genetic DBH deficiency should not be supplemented with phenylalanine, tyrosine, or L-DOPA.

5-hydroxyindoleacetic acid (5HIAA) (38) levels below the mean may indicate lower production and/or decreased metabolism of the neurotransmitter serotonin. 5-hydroxy-indoleacetic acid is a metabolite of serotonin. Low values have been correlated with symptoms of depression. Low production of 5 HIAA can be due to decreased intake or absorption of serotonin's precursor amino acid tryptophan, decreased quantities of cofactors needed for biosynthesis of serotonin such as tetrahydrobiopterin and vitamin B6 coenzyme. In addition, a number of genetic variations such as single nucleotide polymorphisms (SNPs) or mutations can cause reduced production of 5HIAA. Such SNPs are available on The Great Plains DNA methylation pathway test which can be performed on a cheek swab. Values may be decreased in patients on monoamine oxidase (MAO) inhibitors that are drugs or foods that contain tyramine such such as Chianti wine and vermouth, fermented foods such as cheeses, fish, bean curd, sausage, bologna, pepperoni, sauerkraut, and salami.

Slight elevation in suberic acid (48) is consistent with overnight fasting or increased fat in the diet. Regardless of cause, supplementation with L-carnitine or acetyl-L-carnitine may be beneficial.

Pyridoxic acid (B6) levels below the mean (51) may be associated with less than optimum health conditions (low intake, malabsorption, or dysbiosis). Supplementation with B6 or a multivitamin may be beneficial.

Pantothenic acid (B5) levels below the mean (52) may be associated with less than optimum health conditions. Supplementation with B5 or a multivitamin may be beneficial.

Ascorbic acid (vitamin C) levels below the mean (54) may indicate a less than optimum level of the antioxidant vitamin C. Individuals who consume large amounts of vitamin C can still have low values if the sample is taken 12 or more hours after intake. Supplementation with buffered vitamin C taken 2 or 3 times a day is suggested.

High 2-hydroxyhippuric acid (61) may result from ingestion of aspartame (Nutrasweet®), salicylates (aspirin), dietary salicylates, or from GI bacteria converting tyrosine or phenylalanine to salicylic acid. For more information about salicylates in foods go to http://www.feingold.org/salicylate.php. 2-Hydroxyhippuric acid is a conjugate of hydroxybenzoic acid (salicylic acid) and glycine. Very high 2-hydroxyhippuric also inhibits dopamine beta-hydroxylase resulting in elevated HVA, decreased VMA, and elevated HVA/VMA ratio.

Low citramalic, 2-hydroxyphenylacetic, 4-hydroxyphenylacetic, 4-hydroxybenzoic, 4-hydroxyhippuric, 3-indoleacetic, glyceric, glycolic, oxalic, lactic, pyruvic, 3-Methylglutaric, 3-methylglutaconic, 2-hydroxybutyric, fumaric, malic, aconitic, quinolinic, kynurenic, thymine, ethylmalonic, methylsuccinic, adipic, suberic, glutaric, 3-hydroxy-3-methylglutaric, methylcitric, or orotic values have no known clinical significance.

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