



TEST PATIENT

GUa d'Y`HYghBUa Y
 Sex : :
 DUHY Collected : 00-00-0000
 111 H9GH ROAD TEST SUBURB
@AB =8: 00000000 UR#:0000000

TEST PHYSICIAN

DR JOHN DOE
 111 CLINIC STF 99H
 7@-B=7`GI 6I F 6`J =7` \$\$\$

P: 1300 688 522
 E: info@nutripath.com.au
 A: PO Box 442 Ashburton VIC 3142

INTEGRATIVE MEDICINE

URINE, 24 HOUR	Result	Range	Units	
Total Branched Chain AAs	311 *L	424 - 557	umol/L	
GABA	<1.0	0.0 - 1.0	mmol/molCr	
Hydroxylysine	2.2	2.0 - 5.0	umol/L	
AMINO ACIDS, Urine.				
24hr Urine Volume	3250	693 - 3741	mL	
Creatinine Concentration	1385.0	600.0 - 2000.0	mg/24hr	
Specimen Validity				
24hr Urinary Ammonia	34200	11000 - 60000	umol/24h	
Glutamine/Glutamate	0.3 *L	5.0 - 160.0	RATIO	
Taurine, Urine	88.3 *H	16.0 - 80.0	mmol/molCr	
Threonine, Urine	<1.0 *L	7.0 - 29.0	mmol/molCr	
Valine, Urine	1.8 *L	3.0 - 13.0	mmol/molCr	
Cysteine, Urine	<1.0 *L	3.0 - 17.0	mmol/molCr	
Methionine, Urine	2.1	2.0 - 16.0	mmol/molCr	
Isoleucine, Urine	1.9	0.0 - 4.0	mmol/molCr	
Leucine, Urine	<1.0 *L	2.0 - 11.0	mmol/molCr	
Phenylalanine, Urine	<1.0 *L	2.0 - 19.0	mmol/molCr	
Histidine, Urine	23.8 *L	26.0 - 153	mmol/molCr	
Tryptophane, Urine	<1.0	0.0 - 7.0	mmol/molCr	
Arginine, Urine	<1.0	0.0 - 5.0	mmol/molCr	
Lysine, Urine	10.1	7.0 - 58.0	mmol/molCr	
Aspartate, Urine	1.3 *L	2.0 - 7.0	mmol/molCr	
Hydroxyproline, Urine	<1.0	< 13.0	mmol/molCr	
Serine, Urine	<1.0 *L	21.0 - 50.0	mmol/molCr	
Asparagine, Urine	<1.0	0.0 - 23.0	mmol/molCr	
Glutamate, Urine	10.6	0.0 - 12.0	mmol/molCr	
Glutamine, Urine	2.9 *L	20.0 - 76.0	mmol/molCr	
Proline, Urine	<1.00	0.00 - 9.00	mmol/molCr	
Glycine, Urine	41.0 *L	43.0 - 173	mmol/molCr	
Alanine, Urine	10.2 *L	16.0 - 68.0	mmol/molCr	
Tyrosine, Urine	<1.0 *L	2.0 - 23.0	mmol/molCr	
Citrulline, Urine	<1.00	< 4.00	mmol/molCr	
Ornithine, Urine	<1.0	< 5.0	mmol/molCr	
1 Methyl Histidine, Urine	334 *H	< 40.0	mmol/molCr	
3 Methyl Histidine, Urine	23.2	18.0 - 47.0	mmol/molCr	

(*) Result outside normal reference range

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Amino Acids Comment

Glutamine/Glutamate LOW

The low glutamine : glutamate ratio in this specimen may indicate specimen decay. When aged, warmed or improperly preserved, urine glutamine readily breaks down to glutamate and ammonia, hence the test results may not be truly representative for this patient. However, high levels of glutamate can be associated with gout or metabolic acidosis, and thereby are not an indication of specimen decay. Likewise, chronic alcoholism and negative nitrogen balance are associated with physiologically low levels of glutamine.

Histidine Low - check dietary protein, or malabsorption if other essential AAs are low. Low histidine is associated with rheumatoid arthritis and folate deficiency.
Treatment: Folate 800mcg; Histidine 500mg TID.

Leucine Low - potential catabolism of skeletal muscle. Check 3-methylhistidine to confirm this.
Treatment: Use a balanced or custom mixture of essential amino acids,

Phenylalanine Low - can result in altered thyroid function and catecholamine deficits including symptoms of depression, cognitive disorders, memory loss, fatigue, and autonomic dysfunction. Reduce lifestyle stressors and supplement phenylalanine.
Treatment: Use a balanced or custom mixture of essential amino acids,

Threonine Low - can result in hypoglycemic symptoms, particularly if glycine or serine is also low.

Threonine is the precursor of serine and glycine, and is required in the formation of glycoproteins that are essential in immune function. Threonine is slowly absorbed and is often low as a result of rapid transit time, maldigestion or insufficient quality or quantity of dietary protein.

Meats, poultry, fish, some nuts and peanuts and, cheeses are good sources of Threonine.

Treatment: Use a balanced or custom mixture of essential amino acids,

Tryptophan Low - commonly correlated with depression, insomnia, and schizophrenia. Supplementation with 5-hydroxy-tryptophan (5-HTP) may help. 5-HTP is one enzymatic step away from serotonin.
Treatment: 5HTP 50mg TID.

Valine Low - deficiency in this or other BCAAs indicates potential muscle loss. If several essential Amino Acids (AAs) are low, check for adequate stomach acid.
Treatment: Supplement the BCAAs.

Glycine Low - possible generalized tissue loss, glycine being part of the nitrogen pool and important in gluconeogenesis. Supplement glycine.
Treatment: Glycine 1000mg TID.

Serine Low - can lead to disordered methionine metabolism and deficits in acetylcholine synthesis. If simultaneous high threonine or phosphoserine, then need for vitamin B6, folate, and manganese is indicated.
Serine is plentiful in dietary protein and is also formed endogenously from dietary phosphoserine (magnesium dependent), glycine and threonine. In addition, serine is derived from glycolysis provided that the status of B-6 and magnesium are good. Serine is also required for proper metabolism of methionine; a blatant serine deficiency would be expected to be associated with low cysteine and cystathionine and, homocystinurea (elevated plasma homocysteine). Elevated phosphoserine: serine is a good indicator of functional magnesium insufficiency. Low urinary serine is usually

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associated with insufficient protein intake or malabsorption or magnesium deficiency.
Treatment: B6 100mg; Mn 15mg; Folate 800mcg.

Taurine High - may be due to excessive inflammation in the body or to supplementation of other amino acids.

Elevated urinary taurine is usually associated with impaired renal conservation (wasting) due to competition by elevated levels of B-alanine (check B-alanine). Excessive levels of B-alanine are commonly associated with dysbiosis (bacterial and/or fungal). However, first rule out oral supplementation of taurine. B-alanine could also accumulate and compete for retention of taurine with a frank B-6 deficiency; in such a case one would also expect to see elevations in other amino acids that require transamination (eg. leucine, isoleucine, valine). Urinary wasting of taurine can be associated with low intracellular taurine that can negatively impact on intracellular electrolytes (magnesium, potassium, calcium, sodium). Taurine accounts for about 50% of the free amino acids in cardiac tissue, therefore taurine deficiency can result in arrhythmias.

Taurine is also an important antioxidant, neurotransmitter (CNS), and a component of bile acids (fat and fat soluble vitamin absorption). Taurine is a key scavenger of hypochlorite ions, thus a shortage of taurine after viral or bacterial infections, or exposure to xenobiotics (eg. chlorine, chlorite, alcohol, aldehydes) can result in excessive inflammation or chemical sensitivity.

It can be futile to simply supplement Taurine (or magnesium) without correcting the cause of renal wasting of taurine, therefore a Comprehensive Stool Analysis test may be warranted.

Treatment: Vit E 800IU; Vit C 1g TID; b-carotene 25,000 IU; CoQ10 30mg; Lipoate.

Tyrosine Low - implicated in depression, hypothyroidism, and blood pressure disorders. If phenylalanine is normal or high (barring PKU), iron, vitamin C, and niacin supplementation might be indicated to help convert phenylalanine to tyrosine.
Treatment: Iron 30mg; Tyrosine 500mg TID, Vit C 1g BID; Niacin 50mg.

Asparagine Low - can reflect functional need for magnesium in the conversion from aspartic acid. Supplement magnesium.
Treatment: Mg 200mg BID.

Aspartate Low - inhibits ammonia detoxification in the urea cycle. Can be converted to oxaloacetate using B6 and a-KetoGluterate and thus enter the Krebs cycle. Low levels can reflect decreased cellular energy generation, seen as fatigue. Citric and aspartic acids can drive the Krebs (citric acid) cycle, when combined with B6 and a-KetoGluterate.
Treatment:a-KetoGluterate 600mg BID; B6 100mg.

Glutamine Low - deficient intake or absorption of essential amino acids (glutamine is derived from histidine). Check overall amino acid level of diet.
Glutamine is derived directly from dietary protein, and also formed endogenously by addition of ammonia to glutamate. In the CNS the formation of glutamine from glutamate provides a disposal mechanism that protects against excess accumulation of cytotoxic ammonia.
Low glutamine can be a result of protein malnutrition or negative nitrogen balance, incomplete digestive proteolysis or other malabsorption syndromes, or chronic alcoholism. Glutamine can also be low as a result of renal acidosis (low pH, high H+ ion concentration) that is associated with increased renal glutaminase activity and

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increased ammonia excretion.

Glutamine can also be artifactually low as a result of sample decay in which glutamine is broken down to glutamate and ammonia as a result of improper preservation of the urine specimen.

Cystine Low - possible dietary deficiency of methionine and/or cystine. Low cystine can impair Taurine synthesis.

Treatment: N-Acetyl Cystine (NAC) 500mg BID.

Alanine Low - may point to hypoglycemic conditions because of its role in gluconeogenesis. Supplement with alanine and the branched chain amino acids leucine, isoleucine and valine.

Treatment: Use a balanced or custom mixture of essential amino acids,