

Comprehensive Neurotransmitter; urine



Order: SAMPLE REPORT

Client #: 12345

Doctor: Sample Doctor, MD

Doctors Data Inc. 3755 Illinois Ave. St. Charles, IL 60174 Patient: Sample Report

Age: 33 Sex: Male

Body Mass Index (BMI): 24.4

Sample Collection Date/Time Date Collected 01/07/2019

Wake Up Time 0800 Collection Time 0805

Collection Period2nd morning voidDate Received01/08/2019Date Reported01/09/2019

Analyte	Result	Unit per Creatinine	L	WRI	Н	Reference Interval
Phenethylamine (PEA)	27	nmol/g				26-70
Tyrosine	112	μmol/g				28 - 75
Tyramine	1.9	μmol/g				1.6 - 3.2
Dopamine	211	μg/g				110 - 200
3,4-Dihydroxyphenylacetic acid (DOPAC)	331	μg/g				330 - 1000
3-Methoxytyramine (3-MT)	175	nmol/g				82 - 174
Norepinephrine	21	μg/g		A		18 - 42
Normetanephrine	133	μg/g		<u> </u>		70 - 275
Epinephrine	4.3	μg/g				1.3 - 7.3
Metanephrine	55	μg/g		\		44 - 103
Norepinephrine / Epinephrine ratio	4.9			<u> </u>		< 12
Tryptamine	0.3	μmol/g				0.10 - 0.75
Serotonin	83	μg/g		<u> </u>		50 - 98
5-Hydroxyindolacetic acid (5-HIAA)	1450	μg/g				1600 - 6000
Glutamate	42	nmol/g				9.0 - 40.0
Gamma-aminobutyrate (GABA)	2.8	nmol/g				1.6 - 3.5
Glycine	2805	nmol/g				350 - 1500
Histamine	32	μg/g				12-30
Taurine	1111	μmol/g				240 - 900
Creatinine	125	mg/dL				35 - 240



Neurotransmitter Comments:

- Urinary neurotransmitter levels provide an overall assessment of the body's ability to make and break down neurotransmitters and are
 representative of whole body levels. They are required for neurotransmission throughout the body. Direct assessment of neurotransmitter levels
 and metabolism in the central nervous system is not clinically feasible and approximately twenty percent of the total urinary levels are derived
 from the brain. The enzymes, cofactors and precursors in neurotransmitter metabolism in general are the same in the periphery and in the
 central nervous system. Therefore, alterations in urinary neurotransmitter levels assessed in urine provide important clinical information, and may
 be associated with many symptoms including cognitive and mood concerns, diminished drive, fatigue and sleep difficulties, cravings, addictions
 and pain.
- Tyrosine is the non-essential amino acid precursor for dopamine, norepinephrine and epinephrine. Increased tyrosine may exacerbate migraine
 headaches and hyperthyroid conditions. Elevated tyrosine levels may occur due to supplementation (phenylalanine or tyrosine), heritable
 enzyme defects, or liver disease. Tyrosine hydroxylase converts tyrosine into the dopamine precursor L-DOPA; BH4, Vitamin D and iron are
 cofactors for that enzymatic activity.

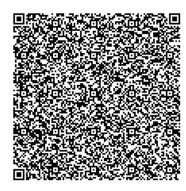
Notes:

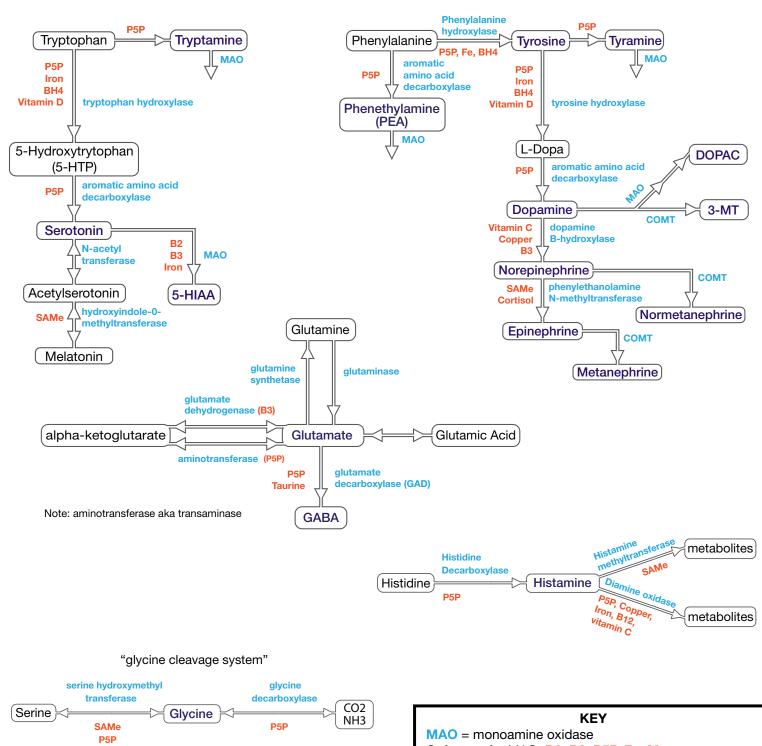
Results are creatinine corrected to account for urine dilution variations. Creatinine is not meant to be used as an indicator of renal function.

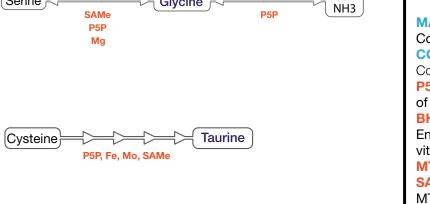
RI= Reference Interval, L (blue)= Low (below RI), WRI (green)= Within RI (optimal), WRI (yellow)= Within RI (not optimal), H (red)= High (above RI)

Methodology: LCMS QQQ, Creatinine by Jaffe Reaction

- Elevated dopamine may be associated with increased worry, distrust of others and decreased ability to interact socially and is often found in patients with attention deficits and hyperactivity. Medications that may increase dopamine levels include L-dopa, methyldopa, select antidepressants and some ADD/ADHD medications. L-theanine may modulate catecholamine effects. Metabolism requires vitamins B2, B3, SAMe, magnesium, and iron, while conversion to norepinephrine requires vitamin C, copper and vitamin B3.
- 3-MT may be increased if dopamine is high; rule out use of L-DOPA. 3-MT is formed by direct metabolism of dopamine by COMT. Very high levels of 3-MT may have stimulatory effects. 3-MT levels may increase during acute stress. Herbicides, such as paraquat, have been shown to increase 3-MT levels in animals. Consumption of foods rich in catecholamines (bananas, pineapple, walnuts) may acutely increase urinary levels of 3-MT. Deficiency or inhibition of MAO may increase 3-MT levels. MAO may be inhibited by cigarette smoke or medications such as monoamine oxidase inhibitors. Vitamins C, B2, B3, SAMe, magnesium, copper and iron are required for optimal dopamine metabolism.
- Low 5-HIAA may be associated with mood concerns including depression and anxiety, sleep changes, and poor concentration. Low 5-HIAA may
 be associated with low precursor serotonin, or compromised metabolism of serotonin by MAO-A. Some medications, including aspirin, MAOinhibitors, levodopa, and tricyclic antidepressants may decrease 5-HIAA levels. MAO may be inhibited by cigarette smoke. Vitamins B2, B3, B6,
 magnesium and iron are required for optimal serotonin metabolism. If MAO-A enzyme function is inhibited, serotonin may be elevated and 5HIAA will be low.
- Elevated glutamate may contribute to anxiety, poor concentration, attention deficits and hyperactive tendencies as well as poor sleep and nighttime awakening. Glutamate may be increased in association with hypoglycemia, Alzheimer's, ALS and chronic compromised blood flow to the brain. Possible sources of increased glutamate include MSG, yeast extract and other hidden sources of free glutamic acid. L-theanine may modulate elevated glutamate levels and attenuate glutamate signaling, and taurine may provide protection from excitotoxicity and neuroinflammation.
- Glycine is a non-essential amino acid that acts as an inhibitory neurotransmitter in the central nervous system. Elevated glycine levels may be
 associated with compromised cognitive processing. Elevated levels may be seen with glycine supplementation. Glycine may be given in
 conjunction with pharmaceutical agents when supporting schizophrenia or psychosis. Lipoic acid may enhance glycine break down. Break down
 of glycine requires vitamin B6 and tetrahydrofolate as cofactors. Note: High levels of glycine may interact with clozapine and decrease its clinical
 efficacy.
- Elevated histamine may be associated with allergy-like symptoms, gastro-intestinal concerns, skin itch/inflammation (pruritis), increased
 wakefulness and insomnia, and has been demonstrated in gastrointestinal blastocystis infections. Levels may be elevated due to use of
 histamine-releasing medications, consumption of allergenic and sulfite-rich foods and/or histamine-rich foods, dysbiotic bacterial production in
 the intestine and zinc deficiency. High urine (and blood) histamine levels have been associated with cluster and cyclic headaches. Break down of
 histamine requires SAMe and copper.
- Taurine is an essential amino acid that may have inhibitory effects on CNS neurons. High urinary levels of taurine may be associated with stress reactions, depression, autism and psychosis. Symptoms may include apathy, sleep changes, irritability, recklessness, poor concentration, aches and pains, or social withdrawal. Patients with Cushing's syndrome (high cortisol) may have elevated urinary taurine levels. Urinary taurine levels may be high with acute or chronic kidney damage, inherited kidney disorders, liver inflammation, or gastrointestinal dysbiotic bacterial or yeast over growth. Oral supplementation may raise taurine levels; taurine is an ingredient in many "energy drinks". High taurine levels may compete with glycine N-methyl-D-aspartate receptors (NMDR). Chronically high taurine excretion may deplete intracellular magnesium and calcium.
- Considerations to address the demonstrated imbalances beyond the identified co-factors and amino acid precursors may include dosage
 adjustments if indicated, as well as nervine and adaptogenic herbs, methylation support, vitamin D, and gastrointestinal health optimization.







Cofactors for MAO: B2, B3, P5P, Fe, Mg

COMT = catechol-o-methyl-transferase

Cofactors for COMT: SAMe, Mg

P5P = (pyridoxal-5-phosphate) activated form of vitamin B6

BH4 = (tetrahydrobiopterin)

Endogenous levels can be supported with SAMe, vitamin B3, C, Mo, Zn

MTHF = (methyltetrahydrofolate) active form of folate.

SAMe = endogenous levels can be supported with Mg, MTHF, and methylcobalamin supplementation.

Cofactors = Enzymes =