

## Amino Acids (Plasma)

<b>Description</b>	A profile of amino acids is provided: alanine, $\alpha$ -aminobutyric acid, arginine, asparagine, aspartic acid, carnosine, citrulline, cystine, glutamic acid, glutamine, glycine, histidine, homocystine, hydroxylysine, isoleucine, leucine, lysine, methionine, 1-methylhistidine, 3-methylhistidine, ornithine, phenylalanine, phosphoethanolamine, proline, sarcosine, serine, taurine, threonine, tyrosine, tryptophan, valine.
<b>Indication</b>	In general, unless looking for a disorder of renal transport, plasma amino acid analysis is more useful than urine. This test is mainly used in paediatrics to screen for inborn errors of metabolism. In adults, it may be used to screen for late presenting metabolic disorders, such as suspected aminoacidopathies. Plasma amino acids may also aid in investigating hyperammonaemia, suspected disorders of energy metabolism, renal disorders (nephrolithiasis, Fanconi syndrome), a positive urine nitroprusside test, epileptic encephalopathy and for monitoring of a protein restricted diet.
<b>Additional Info</b>	Functions of amino acids include the basic structural units of proteins, metabolic intermediates and neurotransmission.
<b>Concurrent Tests</b>	Urine amino acids
<b>Dietary Requirements</b>	Fasting sample preferred
<b>Interpretation</b>	<p>Values depend on metabolic state.</p> <p>Non-specific changes may arise from haemolysis, delayed separation or shipping at room temperature.</p> <p><u>Homocystinuria</u>: Increased plasma homocysteine and methionine.</p>
<b>Collection Conditions</b>	No restrictions.
<b>Frequency of testing</b>	Repeat measurement inappropriate <i>except</i> in acute presentation of undiagnosed suspected metabolic disorder.