

<h2>Total Calcium (Serum)</h2>	
<b>Description</b>	Part of bone profile Reported as both total calcium and calcium relative to normalised albumin (adjusted for changes in albumin concentration)
<b>Indication</b>	Diagnosis and monitoring of disorders of calcium homeostasis and bone metabolism
<b>Additional Info</b>	Serum calcium is 40% protein bound (albumin 32%, globulin 8%), 10% in the form of freely diffusible inorganic complexes and 50% free (ionised). Total serum calcium is adjusted using a population derived formula
<b>Concurrent Tests</b>	PTH, Vitamin D, magnesium, phosphate
<b>Dietary Requirements</b>	N/A
<b>Interpretation</b>	<p>The gut, bone and kidney are three major organ systems involved in homeostasis of calcium, controlled via PTH and Vitamin D.</p> <p><b>Hypercalcaemia</b> most frequently caused by hyperparathyroidism, malignancy and over-treatment with vitamin D, particularly vitamin D analogues Alfacalcidol and Calcitriol.. Other causes include thyrotoxicosis, sarcoidosis, tertiary hyperparathyroidism, lithium, thiazide diuretics, Paget's disease with immobility, milk alkali syndrome, and more rarely Addison's disease and acromegaly.</p> <p><b>Hypocalcaemia</b> results from hypoalbuminaemia, chronic kidney disease (ionised levels normal), vitamin D deficiency, acute renal failure, hypoparathyroidism, magnesium deficiency, acute pancreatitis, blood transfusion, altered vitamin D metabolism, rhabdomyolysis and conditions resulting in shock.</p> <p>Familial hypocalciuric hypercalcaemia (FHH) is a rare condition where patients have defective calcium sensing receptors leading to a raised serum calcium level by which PTH release is triggered. They exhibit a mild PTH-dependent hypercalcaemia. As the receptor is expressed in renal tubules, inappropriate calcium reabsorption occurs leading to hypocalciuria.</p>
<b>Collection Conditions</b>	N/A
<b>Frequency of testing</b>	N/A