Need to talk? Call us 0845 2412173 Monday to Friday 9am to 5pm

Climb™

Supporting those affected by Inherited Metabolic Disorders

Aminoacylase 1 Deficiency

Aminoacylase 1 Deficiency is a rare metabolic disorder which is characterised by increased levels of N-acetylated amino acids in the urine. This disorder often presents in childhood and causes neurological and movement problems. Severity varies greatly with some having no symptoms at all (asymptomatic). Less than 20 cases have been reported so far.

Synonyms

Alternative names for this condition are:

- ACY1D

Further information about this condition is available from Climb.

Disclaimer

Please read our disclaimer and information on data protection.

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