



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



Supporting those affected by
Inherited Metabolic Disorders

Beta-Ureidopropionase Deficiency

Beta-Ureidopropionase Deficiency is a very rare disorder of pyrimidine metabolism which causes a wide range of symptoms which are of extremely varied severity. Some individuals may have no symptoms at all and only be diagnosed through laboratory testing, others may have severe neurological and developmental disorders. Fewer than 10 patients have been reported.

Synonyms

Alternative names for this condition are:

- Beta-alanine Synthase Deficiency
- Deficiency of Beta-Ureidopropionase

Further information about this condition is available from Climb.

Disclaimer

[Please read our disclaimer and information on data protection.](#)



Updated 02/06/17

Telephone: **0845 241 2173**
Email: contact@climb.org.uk

Freephone: **0800 652 3181**
Website: www.climb.org.uk

Children Living with Inherited Metabolic Diseases is a charity registered in England and Wales (1089588) in Scotland (SC044634) and a Company Limited by Guarantee 4267454