Climb™

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.

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Telephone: **0845 241 2173** Freephone: **0800 652 3181**
Email: contact@climb.org.uk Website: www.climb.org.uk