



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



Supporting those affected by
Inherited Metabolic Disorders

Carnosinaemia

Carnosinaemia is a very rare disorder that is characterised by developmental delays and intellectual difficulties of varying severity. Other symptoms may include seizures, extreme drowsiness, growth delays, and involuntary muscle spasms in the arms, legs or head (myoclonic seizures). Some individuals with Carnosinaemia may have no symptoms.

Synonyms

Alternative names for this condition are:

- Carnosinase Deficiency
- Homocarnosinosis

Further information about this condition is available from Climb.

Disclaimer

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