Climb™

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

Please read our disclaimer and information on data protection.

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