Climb™

Supporting those affected by Inherited Metabolic Disorders

Alpha-Ketoglutarate Dehydrogenase Deficiency

Alpha-Ketoglutarate Dehydrogenase Deficiency is a rare metabolic disease that is linked with a defect in the OGDH gene. This is related to a few different processes in the body and affects various tissues including the salivary gland and skeletal muscle. This condition may also be a feature of another disorder known as Dihydrolipoamide Dehydrogenase (DLD) Deficiency.

Synonyms

- Alternative names for this condition are:
- Alpha KGD Deficiency
- 2 Alpha Ketoglutarate Dehydrogenase Deficiency
- 2 Oxoglutarate Dehydrogenase Deficiency
- Oxoglutaric Aciduria

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