



Need to talk? Call us **0845 2412173**

Monday to Friday
9am to 5pm



Supporting those affected by
Inherited Metabolic Disorders

Cutis Laxa, Autosomal Recessive Type 2B

This is a rare disorder which has varying severity. Cutis Laxa is a connective tissue disorder characterised by skin which has lost its elasticity causing it to hang loosely. Other symptoms may include abnormal growth and skeletal abnormalities, developmental delay and joint problems.

Synonyms

Alternative names for this condition are:

- ARCL2, Progeroid Type
- ARCL2B
- Cutis Laxa with Progeroid Features
- PYCR1-Related Cutis Laxa

Further information about this condition is available from Climb.

Disclaimer

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



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