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# Familial anetoderma

INSERM

## Source

INSERM. (1999). *Orphanet: an online rare disease and orphan drug data base*. [Familial anetoderma](#). ORPHA:228277

Familial anetoderma is an extremely rare genetic skin disease characterized by loss of elastin tissue leading to localized areas of flaccid skin and a family history of the disorder.