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Autosomal Recessive Congenital Ichthyosis 2

National Cancer Institute

Source

National Cancer Institute. <u>Autosomal Recessive Congenital Ichthyosis 2</u>. NCI Thesaurus. Code C132827.

An autosomal recessive condition caused by mutation(s) in the ALOX12B gene, encoding arachidonate 12-lipoxygenase, 12R-type. It is characterized by dry, thickened, scaly skin.

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