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Epidermolytic Hyperkeratosis

National Cancer Institute

Source

National Cancer Institute. *Epidermolytic Hyperkeratosis*. NCI Thesaurus. Code C62569.

An autosomal dominant inherited skin disorder caused by mutations in the KRT 1 and KRT 10 genes. It is manifested at birth and is characterized by generalized erythema, skin blisters and skin fragility.