# Familial keratoacanthoma

INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Familial keratoacanthoma. ORPHA:493*

Multiple familial keratoacanthoma (KA) of Witten and Zak is a rare a rare inherited skin cancer syndrome and is characterized by the coexistence of features characteristic of both multiple KA, Ferguson Smith type and generalized eruptive keratoacanthoma (see these terms), such as multiple small miliary-type lesions, larger self-healing lesions, and nodulo-ulcerative lesions .Lesions do not have a predilection for the mucosal surfaces. Transmission is autosomal dominant.