DISEASE:
Aplasia cutis congenita

NAME:	Aplasia cutis congenita
DESCRIPTION:	A rare skin disorder characterized by localized absence of skin that is usually located on the scalp but can occur anywhere on the body including the face, trunk and extremities. Aplasia cutis congenita (ACC) may occasionally be associated with other anomalies.
ORPHACODE:	1114
XREF(S):	Orphanet OMIM MeSH ICD-10 OMIM
ANALYTE(S):	PLEC ITGB4 BMS1 DLL4 UBA2
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Source URL: http://gentest.healthdata.be/disease/1338

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