DISEASE:
Autosomal recessive ataxia, Beauce type

NAME:	Autosomal recessive ataxia, Beauce type
DESCRIPTION:	A rare disorder characterised by a slowly progressive pure cerebellar ataxia associated with dysarthria. It has been described in 53 individuals from 26 families of Canadian origin. The mode of transmission is autosomal recessive. Positional cloning has led to the identification of several gene mutations.
ORPHACODE:	88644
SYNONYMS:	ARCA1 Autosomal recessive cerebellar ataxia type 1 SCAR8
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	SYNE1
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