DISEASE:
Autosomal recessive bestrophinopathy

NAME:	Autosomal recessive bestrophinopathy
DESCRIPTION:	A rare retinal dystrophy, characterized by central visual loss in the first 2 decades of life, associated with an absent electrooculogram (EOG) light rise and a reduced electroretinogram (ERG).
ORPHACODE:	139455
SYNONYMS:	Retinopathy, Burgess-Black type
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	BEST1
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