DISEASE:
Bietti crystalline dystrophy

NAME:	Bietti crystalline dystrophy
DESCRIPTION:	A rare progressive autosomal recessive tapetoretinal degeneration disease, occurring in the third decade of life, characterized by small sparkling crystalline deposits in the posterior retina and corneal limbus in addition to sclerosis of the choroidal vessels and manifesting as nightblindness, decreased vision, paracentral scotoma, and, in the end stages of the disease, legal blindness.
ORPHACODE:	41751
SYNONYMS:	BCD Bietti crystalline corneoretinal dystrophy Bietti crystalline retinopathy
XREF(S):	Orphanet OMIM MeSH ICD-10
ANALYTE(S):	CYP4V2
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

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