DISEASE:
Progressive cone dystrophy

NAME:	Progressive cone dystrophy
DESCRIPTION:	A rare retinal dystrophy characterized by photophobia, progressive loss of visual acuity, nystagmus, visual field abnormalities, abnormal color vision, and psychophysical and electrophysiological evidence of abnormal cone function. Progressive cone dystrophy usually presents in childhood or early adult life, and patients tend to develop rod photoreceptor dysfunction in later life.
ORPHACODE:	1871
SYNONYMS:	Cone dystrophy
XREF(S):	Orphanet OMIM OMIM OMIM OMIM OMIM ICD-10
ANALYTE(S):	PDE6C GNAT2 CNGB3 GUCA1A

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