

RETINITIS PIGMENTOSA AND CONE ROD DYSTROPHY IN 2 SIBLINGS

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ABSTRACT

Cone rod dystrophies (CRDs) (prevalence 1/40,000) are inherited retinal dystrophies that belong to the group of pigmentary retinopathies. CRDs are characterized by retinal pigment deposits visible on fundus examination, predominantly localized to the macular region. Two siblings from the same family presented with rod-cone dystrophy in our regular Outpatient department.

KEYWORDS: *Rods, Cones, Retinopathy, Sibling, Dystrophy*

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