

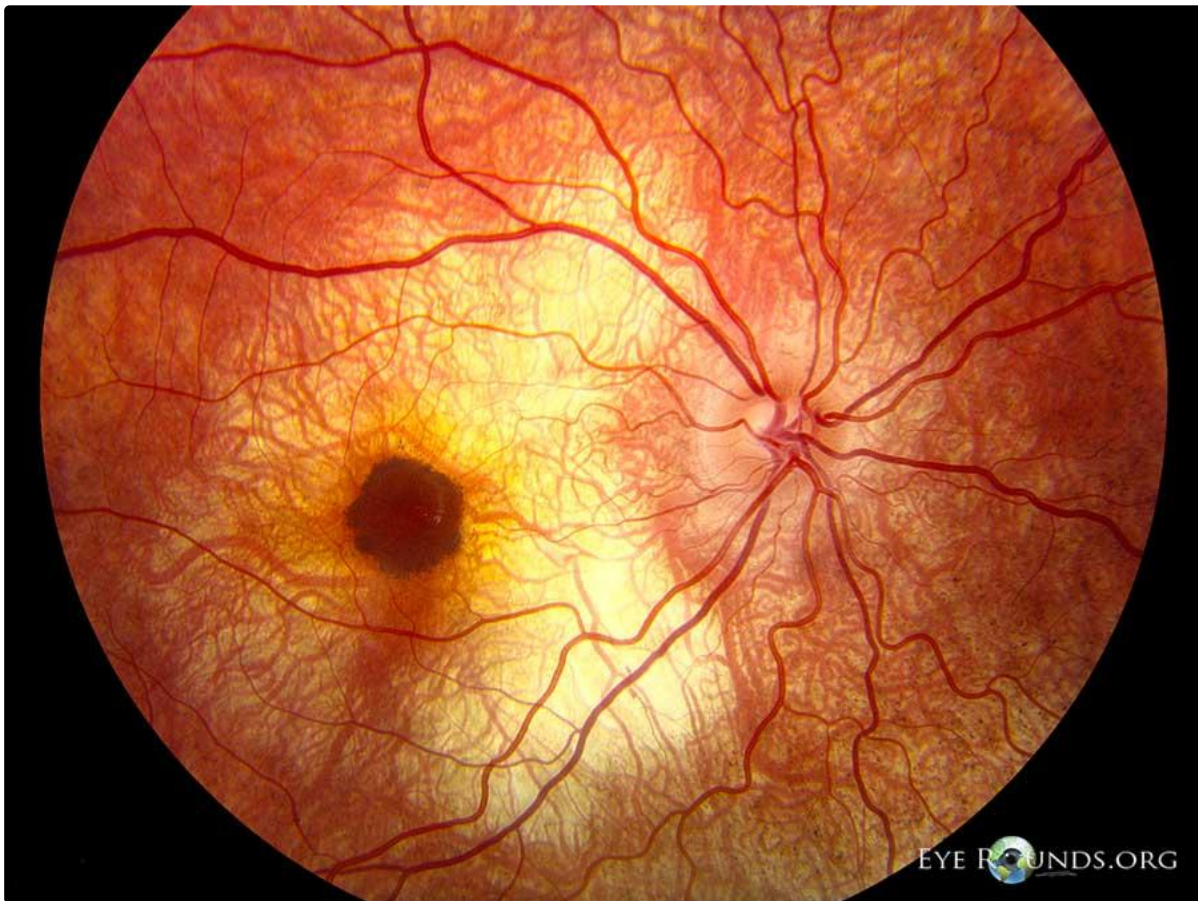
LCHADD: long chain 3-hydroxyacyl CoA dehydrogenase deficiency

Category(ies): Genetics, Inherited Eye Disease, Retina

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[OMIM #609016](#)



7-year-old female with long chain 3-hydroxyacyl CoA dehydrogenase deficiency (LCHADD), Vision 20/40

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