MEDIUM-CHAIN ACYL-COA DEHYDROGENASE DEFICIENCY (MCADD)
Supporting information

This guideline has been prepared with reference to the following:

NHS Newborn Blood Spot Screening Programme. The MCADD programme. 2021.

http://newbornbloodspot.screening.nhs.uk/mcadd

British Inherited Metabolic Disease Group. Medium Chain Acyl CoA Dehydrogenase Deficiency (MCADD). 2016. BIMDG

http://www.bimdg.org.uk/store/guidelines/ER-MCFAO-v4_340405_09092016.pdf

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