Adrenal Hyperplasia (21Hydroxylase deficiency)

General:

21-hydroxylase enzyme mutations are located on the CYP21 gene on chromosome 6p21.3 and can be identified in AGS.

Indication:	Congenital adrenal hyperplasia, AGS, pseudopubertas praecox, virilization, hirsutism.
Material:	5 ml EDTA blood, amniotic fluid, chorionic villus sampling
TAT:	2-3 weeks*
Method:	MLPA (deletions/duplications) of CYP21A2, sequence analysis CYP21A2
Ref.- range:	see report

For complete list of laboratory test offered at Freiburg Medical Laboratory, please visit http://www.fml-dubai.com/parameter-listings/

Updated 24/03/2025