

GENETIC TEST:
Cystic Fibrosis, newborn screening (12 hot spot mutations; CFTR)

FULL NAME:	Cystic Fibrosis, newborn screening (12 hot spot mutations; CFTR)
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Newborn screening
SPECIMEN:	Dried blood spot card
METHOD CATEGORY:	Targeted variant analysis
METHOD TECHNIQUE:	Next Generation Sequencing (NGS)
RIZIV CODE:	565353-565364
EQA:	• CF blood spot, • CF blood spot, • CF blood spot
TURNAROUND TIME (MAXIMUM):	5 days
CREATED:	29 Nov 2019 - 13:05

CHANGED:	22 Jan 2024 - 13:41
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