GENETIC TEST:
Alpha-1-antitrypsin deficiency (2 hot sopt mutations / p.Glu366Lys; p.Glu288Val)

FULL NAME:	Alpha-1-antitrypsin deficiency (2 hot sopt mutations / p.Glu366Lys; p.Glu288Val)
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Carrier diagnosis, Post-natal Diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA
METHOD CATEGORY:	Targeted variant analysis
METHOD TECHNIQUE:	Real-time PCR
RIZIV CODE:	565390-565401
ACCREDITATION (ISO 15189):	2021-07-08 / 2026-02-02
TURNAROUND TIME (MAXIMUM):	6 - 10 weeks
CREATED:	17 Jul 2019 - 11:18
CHANGED:	01 Mar 2023 - 14:36
URL:	https://laboboeken.nexuzhealth.com/pboek/internet/GHB%20CME/12349

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