GENETIC TEST:
Microphthalmia, syndromic 5; Retinal dystrophy, early-onset, and pituitary dysfunction

FULL NAME:	Microphthalmia, syndromic 5; Retinal dystrophy, early-onset, and pituitary dysfunction
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Post-natal Diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA
METHOD CATEGORY:	Sequence analysis: entire coding region
METHOD TECHNIQUE:	Next Generation Sequencing (NGS)
RIZIV CODE:	565456-565460
TURNAROUND TIME (MAXIMUM):	20 - 60 days
CREATED:	07 Aug 2019 - 14:27
CHANGED:	25 Apr 2022 - 09:30

RELATED CONTENT

Related Diseases

- Butterfly-shaped pigment dystrophy
- Combined pituitary hormone deficiencies, genetic forms
- Syndromic microphthalmia type 5

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)

Related Analytes

- orthodenticle homeobox 2