GENETIC TEST:
Hypophosphatasia

FULL NAME:	Hypophosphatasia
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Carrier diagnosis, Mutation confirmation, Post-natal Diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA, DNA
METHOD CATEGORY:	Sequence analysis: entire coding region
METHOD TECHNIQUE:	Bi-directional Sanger Sequence analysis
RIZIV CODE:	565471-565482
ACCREDITATION (ISO 15189):	2022-10-07 / 2027-10-06
TURNAROUND TIME (MAXIMUM):	20-60 days
CREATED:	12 Dec 2019 - 16:34
CHANGED:	04 Nov 2022 - 11:01

RELATED CONTENT

Related Diseases

- Adult hypophosphatasia
- Childhood-onset hypophosphatasia
- Infantile hypophosphatasia
- Odontohypophosphatasia
- Perinatal lethal hypophosphatasia
- Prenatal benign hypophosphatasia

Related Laboratories

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

Related Analytes

- alkaline phosphatase, biomineralization associated