GENETIC TEST:
Temple syndrome / Kagami-Ogata Syndrome

FULL NAME:	Temple syndrome / Kagami-Ogata Syndrome
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Post-natal Diagnosis, Prenatal diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA, Amniotic fluid
METHOD CATEGORY:	Methylation analysis Deletion/duplication analysis
METHOD TECHNIQUE:	MLPA based techniques
RIZIV CODE:	565456-565460
TURNAROUND TIME (MAXIMUM):	3 months
CREATED:	26 Jul 2019 - 10:10
CHANGED:	20 Jan 2023 - 13:31

Source URL: http://gentest.healthdata.be/genetic_test/225

RELATED CONTENT

Related Diseases

- Kagami-Ogata syndrome due to maternal 14q32.2 hypermethylation
- Kagami-Ogata syndrome due to maternal 14q32.2 microdeletion
- Kagami-Ogata syndrome due to paternal uniparental disomy of chromosome 14
- Temple syndrome due to maternal uniparental disomy of chromosome 14
- Temple syndrome due to paternal 14q32.2 hypomethylation
- Temple syndrome due to paternal 14q32.2 microdeletion

Related Laboratories

- Centre de Génétique Médicale UCL

Related Analytes

- delta like non-canonical Notch ligand 1
- maternally expressed 3
- retrotransposon Gag like 1