sciensano **Belgian Genetic Tests database**

GENETIC TEST:
Oculopharyngeal Muscular Dystrophy - GCN repeats expansion

FULL NAME:	Oculopharyngeal Muscular Dystrophy - GCN repeats expansion
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Post-natal Diagnosis, Predictive and Pre-symptomatic diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA
METHOD CATEGORY:	Targeted variant analysis
METHOD TECHNIQUE:	PCR based technique
RIZIV CODE:	565456-565460
TURNAROUND TIME (MAXIMUM):	8 weeks
CREATED:	22 Jul 2019 - 11:03
CHANGED:	19 Oct 2021 - 09:51
URL:	https://laboboeken.nexuzhealth.com/pboek/internet/GHB%20CME/13281

Source URL: http://gentest.healthdata.be/genetic_test/127

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- Oculopharyngeal muscular dystrophy

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- poly(A) binding protein nuclear 1