
GENETIC TEST:
Peutz-Jeghers Syndrome (STK11 gene)

FULL NAME:	Peutz-Jeghers Syndrome (STK11 gene)
DESCRIPTION:	STK11 gene sequencing by NGS and CVN analysis by MLPA
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Post-natal Diagnosis, Predictive and Pre-symptomatic diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA, DNA
METHOD CATEGORY:	Sequence analysis: entire coding region Deletion/duplication analysis
METHOD TECHNIQUE:	Bi-directional Sanger Sequence analysis MLPA based techniques
RIZIV CODE:	565530-565541
ACCREDITATION (ISO 15189):	2022-02-24 / 2026-02-23

EQA:	• Hereditary Breast and Ovarian Cancer (panel testing), • Hereditary Breast and Ovarian Cancer (panel testing) , • Hereditary Breast and Ovarian Cancer (panel testing)
TURNAROUND TIME (MAXIMUM):	2 months
CREATED:	01 Aug 2019 - 11:01
CHANGED:	16 Jan 2024 - 14:49
URL:	https://www.chuliege.be/jcms/c_5225103/fr/syndrome-de-peutz-jeghers-gene-stk11

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