GENETIC TEST:
Gilbert disease / Irinotecan sensitivity / Raltegravir toxicity - Pharmacogenetics

FULL NAME:	Gilbert disease / Irinotecan sensitivity / Raltegravir toxicity - Pharmacogenetics
DESCRIPTION:	UGT1A1 (*rs8175347)* genotyping (identification of the UGT1A1*1, *6,*28,*36,*37 alleles)
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Carrier diagnosis, Drug Response, Post-natal Diagnosis, Therapeutic Management
SPECIMEN:	Peripheral (whole) blood on EDTA
METHOD CATEGORY:	Targeted variant analysis
METHOD TECHNIQUE:	PCR based technique
RIZIV CODE:	565390-565401

TURNAROUND TIME (MAXIMUM):	3 weeks
CREATED:	19 Jul 2019 - 11:38
CHANGED:	23 Jan 2023 - 12:06
URL:	https://www.chuliege.be/jcms/c_6989208/fr/genotypage-du-variant-ugt1a1-28-rs817…

RELATED CONTENT

Related Diseases

- Gilbert syndrome (NON RARE IN EUROPE)
- Irinotecan toxicity

Related Laboratories

- Centre de Génétique Humaine - CHU Sart-Tilman

Related Analytes

- UDP glucuronosyltransferase family 1 member A1