

For in vitro diagnostic use

IVD

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ABCB1 GENE VARIANT C3435T (MDR1)

ORDERING INFORMATIONS

REF: FGC-008-25 RDM Code: 2159865/R CND Code: W0106010499 Tests: 25 Reactions: 31 Manufacturer: BioMol Laboratories s,r,l,

CONTENTS OF THE KIT

The kit consists of: reagents for Real-Time PCR amplification *the reagents for the extraction of genomic DNA are not supplied in the kit.

PRODUCT CHARACTERISTICS

Device belonging to the family of in vitro medical devices REAL-TIME QUALITATIVE PCR-PHARMACOGENETICS TEST The FGC-008 kit allows the characterization of the C3435T genetic variant of the ABCB1 gene (rs1045642) by amplification with oligonucleotides and specific probes (allele-specific genotyping) and subsequent detection with qPCR-Real-time. Kit optimized for Real-Time PCR instrumentation Biorad CFX96 Dx, Biorad Opus Dx, Agilent AriaDx.

SCIENTIFIC BACKGROUND

screening Pharmacogenetic and/or drug-specific phenotyping of cancer patients eligible for treatment with chemotherapy drugs can identify patients likely to be reactive or resistant to proposed drugs. Similarly, identification of patients with an increased risk of developing toxicity allows for dose adaptation or application of other targeted therapies. Polymorphisms in genes encoding drug efflux transporters, such as Pglycoprotein, can affect the absorption and excretion of anticancer drugs. It is encoded by the multidrug resistance 1 (MDR1) gene (ABCB1, ATP-binding cassette transporter superfamily B member 1) located on chromosome 7g21. There are three main polymorphisms affecting P-gp activity: the c.2677G>T/A polymorphism in exon 21 which causes a substitution in the amino acid sequence Ala (G)/Ser (T) or Thr (A), with consequent possible increase in enzyme function. The second polymorphism is in exon 26 at position c.3435C>T, resulting in more than twofold expression of P-gp. The third C1236T polymorphism in exon 12 does not directly affect P-gp expression but has an indirect effect as it alters the stability of the mRNA encoding the protein.

CLINICAL SIGNIFICANCE

Docetaxel and paclitaxel are cytotoxic taxanes that inhibit mitosis causing the death of cancer cells. They are mainly used in the treatment of breast, ovarian and lung cancer. For taxanes, the ABCB1 gene is considered one of the best candidates to explain variations in clinical responses and toxicity.

Doxorubicin, an anthracycline widely used as mono- or combination therapy in the treatment of solid tumors including breast cancer, is also the substrate of P-gp. Significantly impaired clearance and decreased plasma concentration of doxorubicin have been observed in patients with one of the three above described polymorphisms of the ABCB1 gene. Irinotecan, a topoisomerase I inhibitor, plays an important role in the treatment of colorectal cancer in monotherapy or in combination with 5-FU. The antitumor activity is mainly due to the SN-38 metabolite which is metabolized to SN-38G, which has 1/100 of the antitumor activity and is practically inactive.

Aug 13,19,13240-33.
§Are pharmacogenomic biomarkers an effective tool to predict taxane toxicity and outcome in breast cancer patients? Literature review. Cancer Chemother Pharmacol. 2015. Oct,76(4):679-90. doi: 10.1007/s00280-015-2818-4. Epub 2015 Jul 22.







[§] Clinical utility of ABCB1 genotyping for preventing toxicity in treatment with irinotecan Pharmacol Res. 2018 Oct; 136:133-139.doi:10.1016/j.phrs.2018.08.026. Epub 2018 Sep 11.

[§] Genotypes Affecting the Pharmacokinetics of Anticancer Drugs. Clin Pharmacokinet. 2017,

[§] Genotypes Affecting the Pharmacokinetics of Anticancer Drugs. Clin Pharmacokinet. 2017, App;56(4):317-337. doi: 10.1007/s40262-016-0450-z. Review. § Influence of the ABCBI polymorphisms on the response to Taxane-containing chemotherapy: a systematic review and meta-analysis. Cancer Chemother Pharmacol. 2018, Feb;81(2):315-323.doi: 10.1007/s.00280-017-3496-1. Epub 2017 Dec 5. § Irinatecan Pathway Genotype Analysis to Predict Pharmacokinetics. Clin Cancer Res. 2003

⁻Aug 15;9(9):3246-53.



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DESCRIPTION	LABEL	VOLUME	STORAGE
		FGC-008-25	
Mix oligonucleotides and probes	Mix 10X C3435T ABCB1	1 x 85 µl	-20°C
Mix buffer and Taq-polymerase enzyme	Mix Real-Time PCR 2X	1 x 425 µl	-20°C
Deionized H ₂ O	Deionized H₂0	2 x 1 ml	-20°C
Genomic DNA or recombinant DNA	Control +1	1 x 22 µl	-20°C
Genomic DNA or recombinant DNA	Control +2	1 x 22 µl	-20°C
Genomic DNA or recombinant DNA	Control +3	1 x 22 µl	-20°C

TECHNICAL CHARACTERISTICS

COD. FGC-008-25

18 months	
Ready to use	
Genomic DNA extracted from whole blood, tissue, cells	
Recombinant DNA for at least 3 analytical sessions	
Real-time PCR; oligonucleotides and specific probes; 2 FAM/HEX fluorescence channels	
Biorad CFX96 Dx, Biorad Opus Dx e Agilent AriaDx	
85 min	
1 cycle (10 min) at 95 °C; 45 cycles at 95 °C (15 sec) + 60 °C (60 sec)	
Absence of non-specific pairings of oligonucleotides and probes; absence of cross-reactivity	
≥ 0,016 ng of DNA	
0% NCN	
99,9%	
100%/98%	



