

PHARMACOGENOMIC TEST REQUEST FORM

* mandatory fields



PATIENT DETAILS

Forename* _____ Surname* _____

Patient ID _____ DOB* (DD/MM/YYYY) _____

Biological Sex* _____ Hospital Sample ID _____

Ancestry¹

- European African American/Afro-Caribbean Sub-Saharan African Latino
 East Asian Central/South Asian Other _____

¹ As defined by the Clinical Pharmacogenetics Implementation Consortium (CPIC)

Sample ID Internal use only

HEALTH PRACTITIONER DETAILS

Account ID* _____ Address 1* _____

Full Name* _____ Address 2 _____

Phone* _____ City/town* _____

Email* _____ County/ State* _____

Institution* _____ Post Code* _____

Country* _____

TEST DETAILS

Gene to be analysed*²

- DPYD gene variants**
(*2A (c.1905+1A>G), *13 (c.1679T>G), HapB3 (c.1236G>A) and c.2846A>T)
- TPMT gene variants**
(*2 (c.238G>C), *3B (c.460G>A) and *3C (c.719A>G))
- CYP2C19 gene variants**
(*2, *3, *17)
- ² Please refer to our Laboratory User Guide for information on the sample requirements for this test.

CLINICAL INFORMATION

Referral Reason* _____

SAMPLE DETAILS

Sample Type Whole Blood (EDTA Tube) Genomic DNA, Source: _____

Date Collected (DD/MM/YYYY)* _____ Time Collected (HH:MM) _____

BILLING AND REPORT DETAILS

The invoice for this test will be sent to the default billing address associated with your Account ID. If your institution requires a PO number, please insert it here.

PO Number _____

The report for this test will be made available in your account on Genseq's online ordering portal. If you need other health practitioners to have access to the report, please ensure they are registered under your Account to ensure appropriate communication.

*Please, indicate how long you would like Genseq to store laboratory test raw data on your behalf:

- 6 months (default retention time where no option is chosen) 12 months

In addition, in order to fulfil your instructions to perform the genetic testing undertaken in the context of an accredited genetic testing service, you understand that other data types, such as patient data received on this Test Request Form, laboratory QC data and report data will be stored for a further period of years taking account of applicable law, regulation and industry guidance. In returning this Test Request Form for processing you are instructing us in writing to undertake such processing on your behalf

*I hereby confirm that I have obtained written informed consent from the patient for this test to be performed, including consent for health practitioners registered under my account to access the report.