

## Overview

### Useful For

Identifying individuals with genetic variants in *DPYD* who are at increased risk of toxicity when prescribed 5-fluorouracil (5-FU) or capecitabine chemotherapy treatment

### Genetics Test Information

[This is a pharmacogenomics test associated with 5-fluorouracil and capecitabine drug sensitivity. Biallelic variation in the \*DPYD\* gene is also associated with dihydropyrimidine dehydrogenase deficiency.\(1\) Individuals who have variations identified in the \*DPYD\* gene may benefit from genetic consultation.](#)

### Special Instructions

- [Informed Consent for Genetic Testing](#)
- [Pharmacogenomic Association Tables](#)
- [Multiple Genotype Test List](#)
- [Informed Consent for Genetic Testing \(Spanish\)](#)

### Method Name

Real-Time Polymerase Chain Reaction (PCR) with Allelic Discrimination Analysis

### NY State Available

Yes

## Specimen

### Specimen Type

Varies

### Ordering Guidance

This test does not detect or report variants other than the \*2A, \*7, \*8, \*10, \*13, rs67376798, rs75017182, and rs115232898 alleles. Sequencing of the full gene is available for detection of additional variants as well as the alleles listed: order DPYDZ / Dihydropyrimidine Dehydrogenase, *DPYD* Full Gene Sequencing, Varies.

### Specimen Required

Multiple genotype tests can be performed on a single specimen after a single extraction. See [Multiple Genotype Test List](#) for a list of tests that can be ordered together.

**Submit only 1 of the following specimens:**

**Specimen Type:** Whole blood

**Container/Tube:** Lavender top (EDTA)

**Specimen Volume:** 3 mL

**Collection Instructions:**

1. Invert several times to mix blood.
2. Send whole blood specimen in original tube. **Do not aliquot.**

**Specimen Stability Information:** Ambient (preferred) 9 days/Refrigerated 30 days

**Specimen Type:** Saliva

**Patient Preparation:** Patient should not eat, drink, smoke, or chew gum 30 minutes prior to collection.

**Supplies:** Saliva Swab Collection Kit (T786)

**Specimen Volume:** 1 Swab

**Collection Instructions:** Collect and send specimen per kit instructions.

**Specimen Stability Information:** Ambient 30 days

**Specimen Type:** Extracted DNA

**Container/Tube:** 2-mL Screw top tube

**Specimen Volume:** 100 mcL (microliters)

**Collection Instructions:**

1. The preferred volume is 100 mcL at a concentration of 50 ng/mcL.
2. Provide concentration of DNA and volume on tube.

**Specimen Stability Information:** Frozen (preferred)/Ambient/Refrigerated

## Forms

1. **New York Clients-Informed consent is required.** Document on the request form or electronic order that a copy is on file. The following documents are available:

-[Informed Consent for Genetic Testing](#) (T576)

-[Informed Consent for Genetic Testing \(Spanish\)](#) (T826)

2. If not ordering electronically, complete, print, and send a [Therapeutics Test Request](#) (T831) with the specimen.

## Specimen Minimum Volume

Blood: 0.4 mL; Saliva, extracted DNA: see Specimen Required

## Reject Due To

All specimens will be evaluated at Mayo Clinic Laboratories for test suitability.

## Specimen Stability Information

Specimen Type	Temperature	Time	Special Container
Varies	Varies		

## Clinical & Interpretive

## Clinical Information

5-Fluorouracil (5-FU) and its orally administered prodrug, capecitabine, are fluoropyrimidine-based chemotherapeutic agents that are widely used for the treatment of colorectal cancer and other solid tumors.

The dihydropyrimidine dehydrogenase (*DPYD*) gene encodes the rate-limiting enzyme for fluoropyrimidine catabolism and eliminates over 80% of administered 5-FU. Dihydropyrimidine dehydrogenase (*DPYD*) activity is subject to wide variability, mainly due to genetic variation. This results in a broad range of enzymatic deficiency from partial (3%-5% of population) to complete loss (0.2% of population) of enzyme activity.(2-5) Patients who are deficient in *DPYD* are at an increased risk for side effects and toxicity when undergoing 5-FU treatment.(6) In addition, pathogenic homozygous or compound heterozygous variants within *DPYD* are associated with dihydropyrimidine dehydrogenase (*DPD*) deficiency. *DPD* deficiency shows large phenotypic variability, ranging from no symptoms to a convulsive disorder with motor and intellectual disabilities.

The following table displays the *DPYD* variants detected by this assay, the corresponding star allele, and the effect on *DPYD* enzyme activity. Other or novel variations, besides those listed here, may also impact fluoropyrimidine-related side effects and tumor response.

Table. Enzyme Activity of Individual Star Alleles

<b><i>DPYD</i> allele</b>	<b>cDNA nucleotide change</b>	<b>Effect on enzyme activity</b>
*1	None (wild type)	Normal activity
*2A	1905+1G>A	No activity
*7	299_302delTCAT	No activity
*8	703C>T	No activity
*10	2983G>T	No activity
*13	1679T>G	No activity
rs67376798	2846A>T	Decreased activity
rs75017182	1129-5923C>G	Decreased activity
rs115232898	557A>G	Decreased activity

### Reference Values

*DPYD* Phenotype: Normal metabolizer

*DPYD* Activity Score: 2.00

*DPYD* Genotype: No variants were detected in the *DPYD* gene.

An interpretive report will be provided.

### Interpretation

An interpretive report will be provided.

For additional information regarding pharmacogenomic genes and their associated drugs, see [Pharmacogenomic Associations Tables](#). This resource also includes information regarding enzyme inhibitors and inducers, as well as potential alternate drug choices.

### Cautions

Rare genetic variants may be present that could lead to false-negative or false-positive results. Other variants in the

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primer binding regions can affect the testing, and ultimately, the genotype assessment made.

Specimens may contain donor DNA if obtained from patients who received heterologous blood transfusions or allogeneic blood or marrow transplantation. Results from specimens obtained under these circumstances may not accurately reflect the recipient's genotype. For individuals who have received blood transfusions, the genotype usually reverts to that of the recipient within 6 weeks. For individuals who have received allogeneic blood or marrow transplantation, a pretransplant DNA specimen is recommended for testing.

Dihydropyrimidine dehydrogenase (*DPYD*) genetic test results in patients who have undergone liver transplantation may not accurately reflect the patient's *DPYD* status.

Test results should be interpreted in the context of clinical findings, family history, and other laboratory data. Large deletions or rearrangements are not detected by this assay, and these may affect *DPYD* protein expression and their impact on fluoropyrimidine-related side effects and tumor response.

This test is not designed to provide specific dosing or drug selection recommendations and is to be used as an aid to clinical decision making only. Drug-label guidance should be used when dosing patients with medications regardless of the predicted phenotype.

### Clinical Reference

1. OMIM: Dihydropyrimidine dehydrogenase; *DPYD*. 2009. Updated December 13, 2023. Accessed February 22, 2024. Available at [www.omim.org/entry/612779](http://www.omim.org/entry/612779)
2. Amstutz U, Henricks LM, Offer SM, et al. Clinical Pharmacogenetics Implementation Consortium (CPIC) Guideline for Dihydropyrimidine Dehydrogenase Genotype and Fluoropyrimidine Dosing: 2017 Update. *Clin Pharmacol Ther*. 2018;103(2):210-216. doi:10.1002/cpt.911
3. Lunenburg CATC, van der Wouden CH, Nijenhuis M, et al. Dutch Pharmacogenetics Working Group (DPWG) guideline for the gene-drug interaction of *DPYD* and fluoropyrimidines. *Eur J Hum Genet*. 2020;28(4):508-517. doi:10.1038/s41431-019-0540-0
4. Morel A, Boisdron-Celle M, Fey L, et al. Clinical relevance of different dihydropyrimidine dehydrogenase gene single nucleotide polymorphisms on 5-fluorouracil tolerance. *Mol Cancer Ther*. 2006;5(11):2895-2904. doi:10.1158/1535-7163.MCT-06-0327
5. Offer SM, Fossum CC, Wegner NJ, Stuflessen AJ, Butterfield GL, Diasio RB. Comparative functional analysis of *DPYD* variants of potential clinical relevance to dihydropyrimidine dehydrogenase activity. *Cancer Res*. 2014;74(9):2545-2554. doi:10.1158/0008-5472.CAN-13-24826
6. U.S. Food and Drug Administration: Table of Pharmacogenomic Biomarkers in Drug Labeling. FDA; Updated February 2, 2024. Accessed February 22, 2024. Available at [www.fda.gov/drugs/scienceresearch/researchareas/pharmacogenetics/ucm083378.htm](http://www.fda.gov/drugs/scienceresearch/researchareas/pharmacogenetics/ucm083378.htm)

### Performance

### Method Description

Genomic DNA is extracted from whole blood or saliva. Genotyping for *DPYD* alleles is performed using a polymerase

chain reaction (PCR)-based 5'-nuclease assay. Fluorescently labeled detection probes anneal to the target DNA. PCR is used to amplify the section of DNA that contains the variant. If the detection probe is an exact match to the target DNA, the 5'-nuclease polymerase degrades the probe, the reporter dye is released from the effects of the quencher dye, and a fluorescent signal is detected. Genotypes are assigned based on the allele-specific fluorescent signals that are detected. (Unpublished Mayo method)

**PDF Report**

No

**Day(s) Performed**

Monday through Friday

**Report Available**

3 to 10 days

**Specimen Retention Time**

Whole blood/saliva: 2 weeks; Extracted DNA: 2 months

**Performing Laboratory Location**

Rochester

**Fees & Codes****Fees**

- Authorized users can sign in to [Test Prices](#) for detailed fee information.
- Clients without access to Test Prices can contact [Customer Service](#) 24 hours a day, seven days a week.
- Prospective clients should contact their account representative. For assistance, contact [Customer Service](#).

**Test Classification**

This test was developed and its performance characteristics determined by Mayo Clinic in a manner consistent with CLIA requirements. It has not been cleared or approved by the US Food and Drug Administration.

**CPT Code Information**

81232

**LOINC® Information**

Test ID	Test Order Name	Order LOINC® Value
DPYDQ	DPYD Genotype, V	93199-8

Result ID	Test Result Name	Result LOINC® Value
610138	DPYD Phenotype	79719-1
610139	DPYD Activity Score	104665-5

## Test Definition: DPYDQ

Dihydropyrimidine Dehydrogenase Genotype,  
Varies

613999	DPYD Genotype	45284-7
610140	Interpretation	69047-9
610141	Additional Information	48767-8
610142	Method	85069-3
610143	Disclaimer	62364-5
610144	Reviewed by	18771-6