

Partial or complete lack of dihydropyrimidine dehydrogenase (DPD) enzyme functionality can increase the risk of severe drug toxicity during treatment with fluoropyrimidines (5-FU, capecitabine, tegafur)¹. Understanding *DPYD* gene status and its impact on drug metabolism is a rapidly growing focus area in clinical research.

The VeriDose® DPYD Panel targets a set of 9 *DPYD* variants associated with an increased risk of severe toxicity, including the 5 variants recommended by the European Medical Agency² and the 4 variants of primary relevance as noted by the Clinical Pharmacogenetics Implementation Consortium³.

The Agena Bioscience® Solution

- Single well multiplex panel for the detection of 9 SNP genotypes associated with increased risk of toxicity
- > Use as a stand-alone or in combination with other PGx panels on the same run, creating workflow efficiencies
- > Cost effective testing with minimal waste



ASSAY COMPONENTS

Contains all high toxicity risk markers mandated by guidelines plus additional markers of interest.

rsID	Nucleotide change	Allelic Variant	
rs3918290	c.1905+1G>A	DPYD*2A	
rs55886062	c.1679T>G	DPYD*13, p.I560S	
rs67376798	c.2846A>T	p.D949V	
rs75017182	c.1129-5923C>G	HapB3	
rs115232898	c.557A>G	p.Y186C	
rs56038477	c.1236G>A	E412E	
rs115349832	c.959-51T>C		
rs6668296	c.680+139G>A		
rs56276561	c.483+18G>A		

ORDERING INFORMATION

The assay panel sets are available in both 96 and 384 reaction formats to support varying sample throughput.

Catalog No.	Item	# Samples	Format
13330F	VeriDose DPYD Panel Set – CPM (10x96)	960	96 CPM
13331D	VeriDose DPYD Panel Set - CPM (10x384)	3840	384 CPM

References

- 1. https://doi.org/10.1159/000510258
- $2. \ \ https://www.ema.europa.eu/en/news/ema-recommendations-dpd-testing-prior-treatment-fluorouracil-capecitabine-tegafur-flucytosine$
- 3. https://doi.org/10.1002/cpt.911

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Agena Bioscience, Inc.

4755 Eastgate Mall
San Diego, CA 92121
Phone: +1.858.882.2800

Order Desk: +1.858.202.9301
Order Desk Fax: +1.858.202.9220
Orderdesk@agenabio.com
Web: www.agenabio.com

