# Pharmacogenomics Test List

Test Name	Test No.
Cytochrome P450 Testing	
Cytochrome P450 2D6/2C19 Genotyping	512255
Cytochrome P450 2D6 Genotyping	512150
Cytochrome P450 2C19 Genotyping	512212
Cytochrome P450 3A4/3A5 Genotyping	512260
Cytochrome P450 2C9 Genotyping	512143
Cytochrome P450 2C9 Genotyping Siponimod	512215
HLA Testing	
HLA B*58:01, Allopurinol Hypersensitivity	167351
HLA B*57:01, Abacavir Hypersensitivity HLA Association Test	006926
Carbamazepine sensitivity HLA Associations (HLA B*15:02, HLA A*31:01)	167443
Other	
TPMT and NUDT15 Genotyping	512300
UGT1A1 Irinotecan Toxicity	511200
DPYD Genotyping	512275

## **Result Interpretation**

Pharmacogenomic result interpretations vary depending on the test/genes involved.

### Cytochrome P450 Enzymes

Genetic variation in cytochrome P450 (CYP450) genes can affect metabolic activity. CYP450 drug metabolizing enzyme activity can range from the total absence of metabolism to ultrarapid metabolism of certain drugs. Results include genotype and predicted metabolic activity.

#### **Metabolic Activity**

Depending on the CYP450 gene, metabolic activity categories include all or several of the following:

- "Ultrarapid": Ultrarapid metabolizer (UM) Increased activity
- "Rapid": Rapid metabolizer (RM) Slightly increased activity
- "Normal": Normal metabolizer (NM) Normal activity
- "Intermediate": Intermediate metabolizer (IM) Reduced activity
- "Poor": Poor metabolizer (PM) Significantly reduced or absent activity

**Note:** "Rapid" is a metabolic activity category for CYP2C19 only. For CYP2C19 there are also "Likely Intermediate" (LIM) and "Likely Poor" (LPM) categories. For CYP2D6, predicated metabolic activities may also be reported as a range or may be categorized as indeterminate.

#### HLA

Positive or negative for allele(s) associated with adverse events from specific drug therapies

#### Gene variants tested

- CYP2D6: \*2, \*3, \*4, \*5 (deletion), \*6, \*7, \*8, \*9, \*10, \*11, \*12, \*13 (hybrid) \*14, \*15, \*17, \*29, \*31, \*35, \*36 (hybrid), \*36 + \*10 (hybrid), \*40, \*41, \*42, \*49, \*53, \*59, \*68 (hybrid), copy number determination
- CYP2C19: \*2, \*3, \*4, \*5, \*6, \*7, \*8, \*9, \*10, \*17, \*35
- CYP2C9: \*2, \*3, \*5, \*6, \*8, \*11, \*13
- CYP3A4: \*22
- CYP3A5: \*3, \*6, \*7
- TPMT: \*2, \*3A, \*3B, \*3C
- NUDT15: \*2 or \*3, \*4
- UGT1A1: \*28 , \*36, \*37
- DPYD: c.1905+1G>A (previously \*2A, rs3918290), c.1679T>G (previously \*13, rs55886062), c.2846A>T (rs67376798), c.1236G>A (in HapB3 w c.1129-5923C>G, rs56038477), c.557A>G (rs115232898)

**Note:** \*1 in genotype results denotes detection of the normal (reference) sequence at all the variant sites assessed.

#### TPMT and NUDT15

*TPMT:* Genotype (including \*2, \*3A, \*3B and \*3C) and predicted metabolic activity: Normal, Intermediate, Poor. *NUDT15:* Genotype (including \*2 or \*3, and \*4) and predicted metabolic activity: Normal, Intermediate, Possible Intermediate, Poor, Indeterminate.

For both *TPMT* and *NUDT15*, decreased metabolic activity is associated with increased risk of adverse events (myelosuppression) from thiopurine drugs.

#### UGT1A1

One copy (heterozygous), two copies (homozygous), or negative for the \*28 allele associated with reduced *UGT1A1* enzyme activity and increased risk for irinotecan toxicity. \*36 and \*37 variant alleles are also detected.

#### DYPD

Genotype (including variants noted above) and predicted metabolic activity (Normal, Intermediate and Poor). Intermediate metabolizers have decreased DPD enzyme activity and Poor metabolizers have complete DPD deficiency. Both have increased risk for severe or even fatal drug toxicity when treated with fluoropyrimidine drugs.



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