

Client: Example Client ABC123  
123 Test Drive  
Salt Lake City, UT 84108  
UNITED STATES

Physician: Doctor, Example

**Patient: Patient, Example**

**DOB:** Unknown  
**Gender:** Male  
**Patient Identifiers:** 01234567890ABCD, 012345  
**Visit Number (FIN):** 01234567890ABCD  
**Collection Date:** 00/00/0000 00:00

**Pharmacogenetics Panel: Psychotropics, with GeneDose Access**

ARUP test code 3006366

PGX PSYCH Specimen	whole blood
CYP2C19 Genotype	*1/*2
CYP2C19 Phenotype	<b>Intermediate</b> *
CYP2C9 Genotype	*1/*5
CYP2C9 Phenotype	<b>Intermediate</b> *
CYP2D6 Genotype	*1/*4
CYP2D6 Phenotype	<b>Intermediate</b> *
CYP3A4 Genotype	*1/*22
CYP3A4 Phenotype	<b>Intermediate</b> *
CYP3A5 Genotype	*1/*3
CYP3A5 Phenotype	<b>Intermediate</b> *
CYP2B6 Genotype	*1/*6

H=High, L=Low, \*=Abnormal, C=Critical

Unless otherwise indicated, testing performed at:

CYP2B6 Phenotype	<b>I</b> ntermediate	*
UGT2B15_1902023	<b>T/G</b> Hetero	*
ANKK1 rs1800497	<b>G/A</b> Hetero	*
COMT rs4680	<b>G/A</b> Hetero	*
DRD2 rs1799978	<b>A/G</b> Hetero	*
GRIK4 rs1954787	<b>T/C</b> Hetero	*
HTR2A rs6311	<b>G/A</b> Hetero	*
HTR2A rs7997012	<b>T/C</b> Hetero	*
HTR2C rs3813929	<b>T</b> Hemizygous	*
MTHFR rs1801133	<b>C/T</b> Hetero	*
MTHFR rs1801131	<b>A/C</b> Hetero	*
OPRM1 Genotype, Interpretation	<b>AG</b>	*
OPRM1 Phenotype, Interpretation	See Note	

**PGX PSYCH Interpretation**

See Note

The following CYP2C19 allele(s) were detected: \*2/Neg. This result predicts the intermediate metabolizer phenotype.

Recommendation: Guidelines for genotype-based dosing are published by the Clinical Pharmacogenetics Implementation Consortium (CPIC) and other organizations. See : <https://www.pharmgkb.org/>

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The following CYP2C9 allele(s) were detected: \*5/Neg. This result predicts the intermediate metabolizer phenotype, with an activity score of 1.5 of 2.

Recommendation: Guidelines for genotype-based dosing are published by the Clinical Pharmacogenetics Implementation Consortium (CPIC) and other organizations. See: <https://www.pharmgkb.org/>

The following CYP2D6 allele(s) were detected: \*4/Neg. This result predicts the intermediate metabolizer phenotype with an activity score estimated at 1 of 2.

Recommendation: Guidelines for genotype-based dosing are published by the Clinical Pharmacogenetics Implementation Consortium (CPIC) and other organizations. See: <https://www.pharmgkb.org/>

The following CYP3A4 allele(s) were detected: \*22/Neg. This result predicts the intermediate metabolizer phenotype.

The following CYP3A5 allele(s) were detected: \*3/Neg. This result predicts the intermediate metabolizer phenotype.

Recommendation: Guidelines for genotype-based dosing are published by the Clinical Pharmacogenetics Implementation Consortium (CPIC) and can be found at: <https://www.pharmgkb.org/>.

The following CYP2B6 alleles were detected: \*6/Neg. This result predicts the intermediate metabolizer phenotype.

Recommendation: Guidelines for gene-based dosing are published by the Clinical Pharmacogenetics Implementation Consortium (CPIC) and other organizations. See [https://www.pharmgkb.org](https://www.pharmgkb.org/)

This test interrogates the UGT2B15 c.253T>G (rs1902023) variant, and the results are T/G. As such, the interpretation is heterozygous.

The UGT2B15 gene codes for the UDP glucuronosyltransferase family 2 member B15 (UGT2B15) that is involved in conjugative metabolism of many medications, such as the anxiolytics oxazepam and lorazepam. See PharmGKB.org for more information.

The following ANKK1 c.2137G>A (rs1800497) alleles were detected: G/A. As such, the interpretation is heterozygous.

The ANKK1 gene codes for the TAQ1A polymorphism that affects the expression of binding sites for dopamine on the dopamine D2 receptor. Variants may influence the likelihood for toxicity and response to drugs that target the dopaminergic system. Variants are also associated with risk of substance use disorders. See PharmGKB.org for more information.

The following COMT c.472G>A (rs4680) alleles were detected: G/A. As such, the interpretation is heterozygous.

The COMT gene codes for the catechol-O-methyltransferase (COMT) enzyme, which is involved in metabolism of catecholamines such as dopamine and norepinephrine. Variants are associated with variance in response to many drugs as well as tolerance to pain.

Recommendation: Guidelines for genotype-based dosing are published by the Clinical Pharmacogenetics Implementation Consortium (CPIC) and can be found at: <https://cpicpgx.org/> and <https://www.pharmgkb.org/>.

The following DRD2 c.-585A>G (rs1799978) alleles were detected A/G. As such, the interpretation is heterozygous.

The DRD2 gene codes for the dopamine D2 receptor. Variants may influence likelihood for toxicity and response to drugs that

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target the dopaminergic system. See PharmGKB.org for more information.

The following GRIK4 c.83-10039T>C (rs1954787) alleles were detected: T/C. As such, the interpretation is heterozygous.

The GRIK4 gene codes for the subunit 4 of the kainite (glutamate) receptor. Variants are associated with variance in response to some antidepressants. See PharmGKB.org for more information.

This test interrogates two variants. For HTR2A c.-998G>A (rs6311) the results are G/A. As such, the interpretation is heterozygous. For HTR2A c.614-2211T>C (rs7997012) the results are T/C. As such, the interpretation is heterozygous.

The HTR2A gene codes for the serotonin receptor 2A. Variants may influence response to some antipsychotics and antidepressants. See PharmGKB.org for more information.

This test interrogates the HTR2C c.-850C>T (rs3813929) variant, and the result is T. As such, the interpretation is hemizygous.

The HTR2C gene codes for the serotonin 2C receptor that is involved in response to psychotropic medications, particularly antipsychotics. See PharmGKB.org for more information.

This test interrogates two variants. For MTHFR c.665C>T (rs1801133, previously designated as C677T) the results are C/T. As such, the interpretation is heterozygous. For MTHFR c.1286A>C (rs1801131, previously designated A1298C) the results are A/G. As such, the interpretation is heterozygous.

The MTHFR gene codes for methylenetetrahydrofolate reductase (MTHFR), an enzyme that metabolizes folate. Variants are associated with variance in response to many drugs as well as symptoms of depression and hyperhomocysteinemia.

Indication for testing: predict opioid sensitivity.

Interpretation: One copy of the OPRM1 A allele and one copy of the G allele (rs1799971) were detected in this sample. Further studies are needed to determine the clinical significance of this genotype; however, it is possible this patient may require higher or more frequent doses of opioid receptor agonists (e.g., morphine) to achieve adequate pain control. He/she may also be more likely to respond to opioid antagonists (e.g., naltrexone) in the treatment of alcohol and/or opioid dependency. This association of OPRM1 and drug sensitivity is not definitive and may be different for individual opioids.

Recommendation: Annotations for clinical application of this OPRM1 allele are available through the Pharmacogenomics Knowledge Base at: <https://www.pharmgkb.org/gene/PA31945>

For ANNK1, DRD2, GRIK4, HTR2A, HTR2C, and UGT2B15, clinical evidence is limited for the drug associations described thus far, and gene-based dosing guidelines are not currently published.

This result has been reviewed and approved by [REDACTED]

Background Information for Pharmacogenetics Panel:Psychotropics:

CHARACTERISTICS: Variation in genes affecting pharmacokinetics and/or pharmacodynamics (pharmacogenetics) may influence medication selection and dose planning. For example, variants in genes that code for metabolizing enzymes (CYP2B6, CYP2C19, CYP2C9, CYP2D6, CYP3A4, CYP3A5, and UGT2B15) may be associated with altered (slower or faster) metabolism which would affect the kinetics of medication activation, inactivation, and/or

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elimination. Other genes in this panel may predict risk of side effects and/or likelihood of response (ANKK1, COMT, DRD2, GRIK4, HTR2A, HTR2C, MTHFR, and OPRM1). This information may guide medication and dose selection for many prescription medications, including medications relevant to psychiatry such as psychostimulants (e.g., ADHD medication), antidepressants, antipsychotics, and anxiolytics.

Inheritance: Autosomal codominant.

Cause: Gene variants affect enzyme function.

Genes included: ANKK1, COMT, CYP2B6, CYP2C19, CYP2C9, CYP2D6, CYP3A4, CYP3A5, DRD2, GRIK4, HTR2A, HTR2C, MTHFR, OPRM1, and UGT2B15.

Variants Tested:

(Variants are numbered according to the following transcripts: ANKK1 NM\_178510, COMT NM\_000754, CYP2B6 NM\_000767, CYP2C19 NM\_000769, CYP2C9 NM\_000771, CYP2D6 M33388 sequence, CYP3A4 NM\_017460 and CYP3A5 NM\_000777, DRD2 NM\_000795, GRIK4 NM\_014619, HTR2A NM\_000621, HTR2C NM\_001256760, MTHFR NM\_005957, OPRM1 NM\_000914, UGT2B15 NM\_001076).

\*1: Indicative of no detected targeted variants and an assumption of functional allele.

ANKK1: rs1800497, c.2137G>A

COMT: rs4680, c.472G>A

CYP2B6\*4: rs2279343, c.785A>G  
CYP2B6\*6: rs3745274, c.516G>T; rs2279343, c.785A>G  
CYP2B6\*7: rs3745274, c.516G>T; rs2279343, c.785A>G; rs3211371, c.1459C>T  
CYP2B6\*9: rs3745274, c.516G>T  
CYP2B6\*18: rs28399499, c.983T>C  
CYP2B6\*22: rs34223104, c.-82T>C  
CYP2B6\*36: rs34223104, c.-82T>C; rs3745274, c.516G>T; rs2279343, c.785A>G

CYP2C19\*2: rs4244285, c.681G>A; rs12769205, c.332-23A>G  
CYP2C19\*3: rs4986893, c.636G>A  
CYP2C19\*4A: rs28399504, c.1A>G  
CYP2C19\*4B: rs28399504, c.1A>G; rs12248560, c.-806C>T  
CYP2C19\*5: rs56337013, c.1297C>T  
CYP2C19\*6: rs72552267, c.395G>A  
CYP2C19\*7: rs72558186, c.819+2T>A  
CYP2C19\*8: rs41291556, c.358T>C  
CYP2C19\*9: rs17884712, c.431G>A  
CYP2C19\*17: rs12248560, c.-806C>T  
CYP2C19\*35: rs12769205, c.332-23A>G

CYP2C9\*2: rs1799853, c.430C>T  
CYP2C9\*3: rs1057910, c.1075A>C  
CYP2C9\*4: rs56165452, c.1076T>C  
CYP2C9\*5: rs28371686, c.1080C>G  
CYP2C9\*6: rs9332131, c.818del  
CYP2C9\*8: rs7900194, c.449G>A  
CYP2C9\*11: rs28371685, c.1003C>T  
CYP2C9\*12: rs9332239, c.1465C>T

CYP2D6\*2: rs16947, g.2850C>T; rs1135840, g.4180G>C  
CYP2D6\*2A: rs1080985, g.-1584C>G; rs16947, g.2850C>T; rs1135840, g.4180G>C  
CYP2D6\*3: rs35743686, g.2549del  
CYP2D6\*4: rs1065852, g.100C>T; rs3892097, g.1846G>A; rs1135840, g.4180G>C  
CYP2D6\*5: gene deletion  
CYP2D6\*6: rs5030655, g.1707del; rs1135840, g.4180G>C  
CYP2D6\*7: rs5030867, g.2935A>C  
CYP2D6\*8: rs5030865, g.1758G>T; rs16947, g.2850C>T; rs1135840, g.4180G>C  
CYP2D6\*9: rs5030656, g.2615\_2617del  
CYP2D6\*10: rs1065852, g.100C>T; rs1135840, g.4180G>C  
CYP2D6\*11: rs1080985, g.-1584C>G; rs201377835, g.883G>C;

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rs16947, g.2850C>T;  
rs1135840, g.4180G>C  
CYP2D6\*13: a CYP2D7-derived exon 1 conversion  
CYP2D6\*14: rs5030865, g.1758G>A; rs16947, g.2850C>T; rs1135840, g.4180G>C  
CYP2D6\*15: rs774671100, g.137\_138insT  
CYP2D6\*17: rs28371706, g.1023C>T; rs16947, g.2850C>T; rs1135840, g.4180G>C  
CYP2D6\*29: rs16947, g.2850C>T; rs59421388, g.3183G>A; rs1135840, g.4180G>C  
CYP2D6\*35: rs769258, g.31G>A; rs16947, g.2850C>T; rs1135840, g.4180G>C; rs1080985, g.-1584C>G  
CYP2D6\*36: a CYP2D6\*10 carrying a CYP2D7-derived exon 9 conversion  
CYP2D6\*36-\*10: a CYP2D6\*36 and a CYP2D6\*10 in tandem  
CYP2D6\*40: rs28371706, g.1023C>T, rs72549356, c.1863\_1864insTTTCGCCCTTCGCCCC, rs16947, g.2850C>T; rs1135840, g.4180G>C;  
CYP2D6\*41: rs16947, g.2850C>T; rs28371725, g.2988G>A; rs1135840, g.4180G>C  
CYP2D6\*42: rs16947, g.2850C>T; rs72549346, g.3260\_3261insGT;  
rs1135840, g.4180G>C  
CYP2D6\*49: rs1065852, g.100C>T; rs1135822, g.1611T>A; rs1135840, g.4180G>C  
CYP2D6\*69: rs1065852, g.100C>T; rs16947, g.2850C>T; rs28371725, g.2988G>A; rs1135840, g.4180G>C  
CYP2D6\*114: rs1065852, g.100C>T; rs5030865, g.1758G>A; rs16947, g.2850C>T; rs1135840, g.4180G>C  
DUP: complete gene duplications

CYP3A4\*1B: rs2740574, c.-392G>A  
CYP3A4\*22: rs35599367, c.522-191C>T

CYP3A5\*3: rs776746, c.219-237A>G  
CYP3A5\*6: rs10264272, c.624G>A  
CYP3A5\*7: rs41303343, c.1035dup

DRD2: rs1799978, c.-585A>G  
DRD2: rs1079598, c.-31-870T>C  
DRD2: rs1799732, c.-486dup  
DRD2: rs2734841, c.1139-134T>G  
GRIK4: rs1954787, c.83-10039T>C

HTR2A: rs6311, c.-998G>A  
HTR2A: rs7997012, c.614-2211T>C

HTR2C: rs3813929, c.-850C>T

MTHFR: rs1801131, c.1286A>C  
MTHFR: rs1801133, c.665C>T

OPRM1: rs1799971, c.118A>G

UGT2B15: rs1902023, c.253T>G

Clinical sensitivity: Drug dependent.  
Methodology: Polymerase chain reaction (PCR) and fluorescence monitoring. Long-range PCR and Sanger sequencing is only performed if needed to characterize a duplicated CYP2D6 gene.  
Analytic Sensitivity and Specificity: Greater than 99 percent.  
Limitations: Only the targeted variants will be detected by this panel, and assumptions about phase and content are made to assign alleles. Publicly available sources such as the [www.pharmvar.org](http://www.pharmvar.org) or [www.pharmgkb.org](http://www.pharmgkb.org) provide guidance on phenotype predictions and allele frequencies. A combination of the CYP2D6\*5 (gene deletion) and a CYP2D6 gene duplication cannot be specifically identified; however, this combination is not expected to adversely affect the phenotype prediction. Diagnostic errors can occur due to rare sequence variations.

Risk of therapeutic failure or adverse reactions with gene substrates may be affected by genetic and nongenetic factors that are not detected by this test.

**H=High, L=Low, \*=Abnormal, C=Critical**

For ANKK1, DRD2, GRIK4, HTR2A, HTR2C, and UGT2B15, clinical evidence is limited for the drug associations described thus far, and gene-based dosing guidelines are not currently published. This result does not replace the need for therapeutic drug or clinical monitoring.

Please note the information contained in this report does not contain medication recommendations and should not be interpreted as recommending any specific medications. Any dosage adjustments or other changes to medications should be evaluated in consultation with a medical provider.

This test was developed and its performance characteristics determined by ARUP Laboratories. It has not been cleared or approved by the U.S. Food and Drug Administration. This test was performed in a CLIA-certified laboratory and is intended for clinical purposes.

Counseling and informed consent are recommended for genetic testing. Consent forms are available online.

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PGX PSYCH, GeneDose Link

See Note

Interpretive Information: PGX Panel: Psychotropics with GeneDose  
GeneDose LIVE content is provided by Coriell Life Sciences and not by ARUP Laboratories.

Any dosage adjustments or other changes to medication should be evaluated in consultation with a medical provider.

This test was developed and its performance characteristics determined by ARUP Laboratories. It has not been cleared or approved by the U.S. Food and Drug Administration. This test was performed in a CLIA-certified laboratory and is intended for clinical purposes.

Counseling and informed consent are recommended for genetic testing. Consent forms are available online.

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EER PGX Panel: Psych, GeneDose

See Note

Authorized individuals can access the ARUP Enhanced Report using the following link:

[REDACTED]

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VERIFIED/REPORTED DATES

Procedure	Accession	Collected	Received	Verified/Reported
PGX PSYCH Specimen	23-324-115679	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
CYP2C19 Genotype	23-324-115679	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
CYP2C19 Phenotype	23-324-115679	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
CYP2C9 Genotype	23-324-115679	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
CYP2C9 Phenotype	23-324-115679	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
CYP2D6 Genotype	23-324-115679	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
CYP2D6 Phenotype	23-324-115679	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
CYP3A4 Genotype	23-324-115679	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
CYP3A4 Phenotype	23-324-115679	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
CYP3A5 Genotype	23-324-115679	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
CYP3A5 Phenotype	23-324-115679	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
CYP2B6 Genotype	23-324-115679	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
CYP2B6 Phenotype	23-324-115679	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
UGT2B15_1902023	23-324-115679	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
ANKK1 rs1800497	23-324-115679	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
COMT rs4680	23-324-115679	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
DRD2 rs1799978	23-324-115679	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
GRIK4 rs1954787	23-324-115679	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
HTR2A rs6311	23-324-115679	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
HTR2A rs7997012	23-324-115679	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
HTR2C rs3813929	23-324-115679	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
MTHFR rs1801133	23-324-115679	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
MTHFR rs1801131	23-324-115679	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
OPRM1 Genotype, Interpretation	23-324-115679	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
OPRM1 Phenotype, Interpretation	23-324-115679	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
PGX PSYCH Interpretation	23-324-115679	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
PGX PSYCH, GeneDose Link	23-324-115679	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00
EER PGX Panel: Psych, GeneDose	23-324-115679	00/00/0000 00:00	00/00/0000 00:00	00/00/0000 00:00

END OF CHART

H=High, L=Low, \*=Abnormal, C=Critical

Unless otherwise indicated, testing performed at:

ARUP LABORATORIES | 800-522-2787 | aruplab.com  
500 Chipeta Way, Salt Lake City, UT 84108-1221  
Jonathan R. Genzen, MD, PhD, Laboratory Director

Patient: Patient, Example  
ARUP Accession: 23-324-115679  
Patient Identifiers: 01234567890ABCD, 012345  
Visit Number (FIN): 01234567890ABCD  
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