Zmyome[™] Proactive Health MEDICATION RESPONSE[™]

TEST CODE: PR32005

Overview

MyOme Proactive Health Medication Response report uses a Blended Genome-Exome (BGE) backbone to identify variants associated with medication response. This allows MyOme to re-query a patient's data as healthcare needs change and new information about the genome is discovered.

Clinical Use

Test is intended to facilitate the use of pharmacogenomic guidance in a general care setting. MyOme reports variants and star alleles in 15 pharmacogenes that are known to impact an individual's response to medication. The results of this test should be interpreted by a trained healthcare provider based on the full context of a patient's medical situation.

Method

Genomic DNA obtained from submitted samples is used to construct a PCR-free whole genome library and an exome library and sequenced using Illumina technology. Reads are aligned to the GRCh38 reference assembly. MyOme's Medication Response variant calling pipeline is used to call variants, star alleles, copy number variants, and assign pharmacogenomic diplotypes.

Sample Types

- Blood (2 EDTA tubes)
- Saliva (2 tubes)
- Buccal (2 swabs)

Turn Around Time

• From initial sample received, approximately 6-8 weeks

Included

- Analysis of key genes with known pharmacogenomic implications for more than 70 drugs, including antidepressants, statins and opioids
- 96% Tier 1 variant alleles recommended by the Association for Molecular Pathology (AMP) PGx Working Group for CYP2C19¹, CYP2C9², CYP2D6³ (excluding deletions/duplications), TPMT and NUDT15⁴
- All Tier 2 variant alleles recommended by the AMP PGx Working Group for CYP2C19¹, CYP2C9², CYP2D6 (excluding hybridizations)³
- Independent healthcare provider review of analysis to make any potential drug recommendations based on gene-drug interactions outlined in the Clinical Pharmacogenetics Implementation Consortium (CPIC)⁵ guidelines and FDA Table of Pharmacogenomic Associations⁶
- 15 genes analyzed: CYP2B6: *4; *6; *9; *18; *22 CYP2C9: *2; *3; *4; *5; *6; *8; *11; *12; *13; *15; *16; *26; *28; *29; *30; *31; *42; *55 CYP2C19: *2; *3; *4; *5; *6; *7; *8; *9; *10; *17; *35 CYP2D6: *2; *3; *4; *6; *7; *8; *9; *10; *11; *12; *14; *15; *17; *21; *31; *40; *41; *42; *49; *56; *59; *100; *114; CYP3A4: *22 CYP3A5: *3; *6 CYP4F2: *3 DPYD: rs3918290; rs55886062; rs59086055; rs67376798; rs75017182+rs56038477; rs112766203; rs115232898; rs146356975; rs183385770 F5: rs6025 IFNL3: rs12979860 NUDT15: *3; *4; *9, SLC01B1: *5; *9; *14; *20 TPMT: *2; *3A; *3B; *3C; *4; *11; *29 UGT1A1: *6; *27 VKORC1: rs9923231

Test Performance⁷

- >98.5% accuracy for diplotypes
- 98% of core pharmacogenomic variants at ≥20x depth
- 60x average exome-wide coverage

1. Pratt VM et al. Recommendations for Clinical CYP2C19 Genotyping Allele Selection: A Report of the Association for Molecular Pathology. J Mol Diagn. 2018 May;20(3):269-276. doi: 10.1016/j. jmoldx.2018.01.011. 2. Pratt VM et al. Recommendations for Clinical CYP2C9 Genotyping Allele Selection. J Mol Diagn. 2019 Sep;21(5):746-755. doi: 10.1016/j.jmoldx.2019.04.003. 3. Pratt VM et al. Recommendations for Clinical CYP2D6 Genotyping Allele Selection. J Mol Diagn. 2021 Sep;23(9):1047-1064. doi: 10.1016/j.jmoldx.2021.05.013. 4. Pratt VM et al. TPMT and NUDT15 Genotyping Recommendations. J Mol Diagn. 2022 Oct;24(10):1051-1063. doi: 10.1016/j.jmoldx.2022.06.007. 5. CPIC. Genes-Drugs. Web. cpicpgx.org/genes-drugs. Accessed 2024 Dec. 6. US Food & Drug Administration. Table of Pharmacogenetic Associations. 2022 Oct;24(10):1051-1016/j.jmoldx.2024 Dec. 7.MyOme Inc, Data on File

The test described above was developed, and its performance characteristics determined, by MyOme, Inc., a clinical laboratory certified under the Clinical Laboratory Improvement Amendments of 1988 (CLIA) to perform high complexity clinical laboratory testing. This test has not been cleared or approved by the U.S. Food and Drug Administration (FDA).

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DRUG-GENE IMPACT

The list of drugs below can lead to a major or moderate drug-gene interaction based on guidelines described above.^{5,6}

-	pelow can lead to a m			
Behavioral Health				
Drug Name	Gene(s)			
amitriptyline	CYP2D6, CYP2C19			
amoxapine	CYP2D6			
amphetamine	CYP2D6			
aripiprazole	CYP2D6			
aripiprazole lauroxil	CYP2D6			
atomoxetine	CYP2D6			
brexpiprazole	CYP2D6			
citalopram	CYP2C19, CYP2D6			
clobazam	CYP2C19			
clomipramine	CYP2D6			
clozapine	CYP2D6			
desipramine	CYP2D6			
diazepam	CYP2C19			
doxepin	CYP2D6, CYP2C19			
duloxetine	CYP2D6			
escitalopram	CYP2C19			
fluoxetine	CYP2D6			
fluvoxamine	CYP2D6			
iloperidone	CYP2D6			
imipramine	CYP2D6			
lofexidine	CYP2D6			
nortriptyline	CYP2D6			
paroxetine	CYP2D6			
perphenazine	CYP2D6			
protriptyline	CYP2D6			
sertraline	CYP2C19			
thioridazine	CYP2D6			
trimipramine	CYP2D6, CYP2C19			
valbenazine	CYP2D6			
venlafaxine	CYP2D6			
vortioxetine	CYP2D6			
U	rology			
Drug Name	Gene(s)			
darifenacin	CYP2D6			
fesoterodine	CYP2D6			
mirabegron	CYP2D6			
tamsulosin	CYP2D6			
tolterodine	CYP2D6			
Reproductive and Sexual Health				
Drug Name	Gene(s)			
flibancarin				

or moderate drug-gene interaction bas				
Gastro	ben	terology		
Drug Name		Gene(s)		
dexlansoprazole		CYP2C19		
dronabinol		CYP2C9		
esomeprazole		CYP2C19		
lansoprazole		CYP2C19		
metoclopramide		CYP2D6		
nateglinide		CYP2C9		
omeprazole		CYP2C19		
ondansetron		CYP2D6		
pantoprazole		CYP2C19		
rabeprazole		CYP2C19		
Infecti	ious	s Disease		
Drug Name		Gene(s)		
atazanavir		UGT1A1		
dolutegravir		UGT1A1		
efavirenz		CYP2B6		
peginterferon alfa-2a		IFNL3		
peginterferon alfa-2b		IFNL3		
quinine		CYP2D6		
voriconazole		CYP2C19		
Cardiology				
Drug Name	Ge	ene(s)		
atorvastatin	SLCO1B1			
carvedilol	CY	CYP2D6		
clopidogrel	CYP2C19			
fluvastatin	CY	CYP2C9, SLCO1B1		
Iovastatin	SL	SLCO1B1		
mavacamten	CY	P2C19		
pitavastatin	SL	SLC01B1		
pravastatin	SL	SLC01B1		
rosuvastatin	SL	SLC01B1		
simvastatin	SL	CO1B1		
warfarin	CYP2C9, CYP4F2, VKORC1			
Mise	cell	aneous		
	Ge	ene(s)		
Drug Name	CYP2C19			
Drug Name abrocitinib	СҮ	'P2C19		
	-	'P2C19 'P2D6		
abrocitinib	СҮ	-		
abrocitinib cevimeline	CY CY	P2D6		
abrocitinib cevimeline eliglustat	CY CY CY	P2D6 P2D6		

on guidelines described above.5,6			
Neurology			
Drug Name	Gene(s)		
brivaracetam	CYP2C19		
deutetrabenazine	CYP2D6		
donepezil	CYP2D6		
fosphenytoin	CYP2C9		
modafinil	CYP2D6		
phenytoin	CYP2C9		
pimozide	CYP2D6		
pitolisant	CYP2D6		
siponimod	CYP2C9		
tetrabenazine	CYP2D6		
valbenazine	CYP2D6		
Pain Management			
Drug Name	Gene(s)		
carisoprodol	CYP2C19		
celecoxib	CYP2C9		
codeine	CYP2D6		
elagolix	SLCO1B1		
flurbiprofen	CYP2C9		
ibuprofen	CYP2C9		
lornoxicam	CYP2C9		
meloxicam	CYP2C9		
oliceridine	CYP2D6		
piroxicam	CYP2C9		
tenoxicam	CYP2C9		
tramadol	CYP2D6		
Hematology/Oncology			
Drug Name	Gene(s)		
capecitabine	DPYD		
erdafitinib	CYP2C9		
eltrombopag	F5		
fluorouracil	DPYD		
gefitinib	CYP2D6		
mercaptopurine	TPMT, NUDT15		
tamoxifen	CYP2D6		
thioguanine	TPMT, NUDT15		
Transplant			
Drug Name	Gene(s)		
azathioprine	TPMT, NUDT15		
tacrolimus	CYP3A5		

5. CPIC. Genes-Drugs. Web. cpicpgx.org/genes-drugs. Accessed 2024 Dec. 6. US Food & Drug Administration. Table of Pharmacogenetic Associations. 2022 Oct. Web. Accessed 2024 Dec. The test described above was developed, and its performance characteristics determined, by MyOme, Inc., a clinical laboratory certified under the Clinical Laboratory Improvement Amendments of 1988 (CLIA) to perform high complexity clinical laboratory testing. This test has not been cleared or approved by the U.S. Food and Drug Administration (FDA). MyOme is not responsible for the content or accuracy of third-party websites.

CYP2D6

tropisetron

CYP2C19, CYP2C9, CYP2D6

flibanserin