

PATIENT INFORMATION
NAME: Sample Patient
DOB: 16/Feb/2000
SEX AT BIRTH: Female

SPECIMEN DETAILS
BARCODE: GNL-DL-00000
SAMPLE ID: 0000
TYPE: Copan FLOQSwab
COLLECTED: 02/Dec/2023

ORDERED BY
Nordic Laboratories
REPORT
GENERATED: 07/May/2024

Summary of Genetic Lab Data & Phenotypes

Gene	Allele Result	Phenotype Result
CYP3A4	*1/*1	Normal Metabolizer
CYP2D6	*2/*4	Intermediate Metabolizer
CYP2C9	*2/*3	Poor Metabolizer
CYP2C19	*1/*1	Normal Metabolizer
SLCO1B1	*1/*1	Normal Function
CYP2B6	*1/*1	Normal Metabolizer
CYP3A5	*3/*3	Poor Metabolizer

This is a short summary of the full medication report. The patient's results are now accessible within the clinical decision support software, TreatGX and ReviewGx, and can be used with other clinical information to enable precision prescribing and medication management. The final genotype/phenotype call is at the discretion of the laboratory director. Medication changes should only be initiated at the discretion of the patient's healthcare provider after a full assessment.

Methods

DNA was extracted from dried blood spot (DBS) card by Chemagic 360 system (Revvity) and processed in a Biomark X platform (Standard Biotools) with Advanta™ Pharmacogenomics Assay.

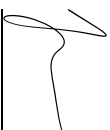
Limitations

The annotations and interpretations provided in this report are based on scientific literature and do not take into account drug-drug interactions, medical conditions or other clinical factors that may affect medication response. Gene-drug interactions are ranked according to guidelines, level of evidence and clinical utility. GenXys reports and TreatGX Clinical Decision Support are regularly updated. Current predicted phenotype and allele functionality may change in the future depending on new evidence. Phenotype annotations for CYP2C9 are based on total activity scores as defined by CPTC⁹. Genetic test results and interpretation may be inaccurate for individuals who have undergone or are receiving non-autologous blood transfusion, tissue, or organ transplant therapies.

The report includes alleles of proteins involved in the metabolism of many medications. In rare cases, a variant that is not covered may be typed as *1 or other variants. In the case of pseudogenes and mutations in the untranslated regions of genes, incorrect allele typing may occur despite proper SNP detection. Preferential amplification of one allele over another present in the sample may also lead to incorrect genotyping.

Liability Disclaimer

This test was developed and its performance characteristics determined by GenXys Health Care Systems. It has not been cleared or approved by the US Food and Drug Administration. The report is not a diagnostic test, and TreatGX is not a prescribing system. You should discuss your pharmacogenetic information with a physician or other health care provider before you act upon the pharmacogenetic information resulting from this report. The medication brand names are not an exhaustive list and do not include combination therapies. Not all medications in this report are included in the TreatGX or ReviewGx software or other GenXys derivative works.



Dr. Juha Matilainen, Laboratory
Director, PhD

Date of Signature

07/May/2024



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	1 Mild or no known interaction	2 Moderate gene-drug interaction Consider alternative medications				3 Serious drug-gene interaction: evaluate and consider alternative medications
Analgesia	Alfentanil	Meloxicam Piroxicam Tenoxicam		Amitriptyline Celecoxib Desipramine Flurbiprofen Ibuprofen Imipramine Nortriptyline	Celecoxib Flurbiprofen Ibuprofen Meloxicam Piroxicam Tenoxicam	
	Carsisoprodol					
	Codeine					
	Fentanyl					
	Hydrocodone					
Autimmune	Morphine					
	Tramadol					
	Venlafaxine					
Cancer	Cyclosporine			Siponimod		
	Tacrolimus					
Cardiovascular	Erdafitinib	Tamoxifen	Tamoxifen	Tamoxifen		
	Atorvastatin Carvedilol Clopidogrel Lovastatin Nebivolol Pitavastatin Pravastatin Propranolol	Warfarin	Flecainide Fluvastatin Warfarin	Flecainide Metoprolol Propafenone Warfarin	Flecainide Fluvastatin Metoprolol Propafenone Warfarin	



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	Rosuvastatin Simvastatin						
Gastroenterology	Metoclopramide Ondansetron	May require an increased dose	Dexlansoprazole Lansoprazole Omeprazole Pantoprazole	Dronabinol Medicine	Dronabinol Medicine	Dronabinol Medicine	
Infection	Efavirenz Voriconazole						
Mental Health	Amoxapine Amphetamine Aripiprazole lauroxil Atomoxetine Citalopram Clonazepam Escitalopram Lofexidine Protriptyline Risperidone Sertraline Venlafaxine			Amiripityline Clomipramine Desipramine Doxepin Imipramine Nortriptyline Paroxetine Trimipramine		Alprazolam Amiripityline Aripiprazole Asenapine Brexpiprazole Bromazepam Cariprazine Chlordiazepoxide Chlorpromazine Clomipramine Clonazepam Clorazepate Clonazepam Desipramine Diazepam	Zuclopenthixol

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	May require an increased dose	May require a reduced dose	May reduce efficacy	May increase adverse events	
				Doxepin Flupentixol Fluphenazine Flurazepam Fluvoxamine Haloperidol lloperidone Imipramine Lorazepam Loxapine Lurasidone Methotrimeprazine Molindone Nitrazepam Nortriptyline Olanzapine Oxazepam Paliperidone Paroxetine Perphenazine Pimozide Prochlorperazine Promethazine Quetiapine Temazepam Thioridazine Triazolam	

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Neurology	Brivaracetam						
	Clobazam						
	Deutetrabenazine						
	Donepezil						
	Galantamine						
Other	Avatrombopag						Eliglustat
	Cevimeline						
	Elagolix						
	Eltrombopag						
	Filbanserin						
Rheumatology	Oral contraceptives						



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	1	2				3	
	Mild or no known interaction	Moderate gene-drug interaction	May require an increased dose	May require a reduced dose	May reduce efficacy	May increase adverse events	Serious drug-gene interaction: evaluate and consider alternative medications
		Consider alternative medications			Piroxicam Tenoxicam	Piroxicam Tenoxicam	
Urology	Darifenacin Fesoterodine Mirabegron Tamsulosin Tolterodine						