

ACUMEN DIAGNOSTICS PTE LTD.

Tel: 6980 0080 HCI: 2010257 41 Science Park Road #01-02 Singapore 117610

Patient Name MOCK
NRIC/FIN No. XXXXX999A

Passport No.

Nationality SINGAPOREAN D.O.B. 01-JAN-1999

Sex (M/F) M

Accession No.MOCK SAMPLEClinic / HospitalMOCK SAMPLELab-Use IDMOCK SAMPLEOrdering DoctorMOCK SAMPLE

Date of SamplingDD-MMM-YYYYDate of Receipt at LabDD-MMM-YYYY hh:mm:ssSample TypeWhole blood in EDTA tubeDate of ResultsDD-MMM-YYYY hh:mm:ss

Report Section 1: Overall Genotyping Results Genotyping by PCR - Pharmacogenomics Test for Statins

Gene Tested	Diplotype Detected*	Overall Functional Effect
SLCO1B1	*5/*37	Decreased enzyme function
ABCG2 (c.421G>T)	G/G	Normal enzyme function
CYP2C9	*3/*3	Poor enzyme function
		See the following pages for CPIC® Medication Insights and Therapeutic Recommendations

*Allelic variants tested:

SLCO1B1 *5 (c.521T>C, rs4149056), SLCO1B1 *15/*37 (c.388A>G, rs2306283); ABCG2 allele c.421G>T (rs2231142)

Disclaimers:

This genotying test is a clinical test intended to provide genetic information to the healthcare provider to aid in the dose selection of drugs. These genetic variations detected under this test do not account for all of the variability in drug pharmacokinetics. This test is not intended as diagnostic, nor is it capable of being an advice on any specific problem or a recommendation for the prescription of any specific drug or a replacement thereof. The healthcare provider shall exercise professional judgement and careful interpretation of the test result in determining their advise to the patient and in the dose selection of drugs.



Source of Therapeutic Recommendations and Medical Insights: CPIC® cpicpgx.org/guidelines/cpic-guideline-for-statins/

Results Verified By: MR LOUIS ONG [Title]

Results Approved By: DR CHIEW YOKE FONG Medical Director



ACUMEN DIAGNOSTICS PTE LTD.

Tel: 6980 0080 HCI: 2010257 41 Science Park Road #01-02 Singapore 117610

Report Section 2:

Therapeutic Recommendations based on SLCO1B1 genotyping results

Therapeutic Recommendations based on SLCO1B1 genotyping results			
Medications Affected	Medication Insights^	Therapeutic Recommendations^ (Classification of Recommendations)	
Atorvastatin	Increased atorvastatin exposure as compared to normal function which may translate to increased myopathy risk.	Prescribe ≤40mg as a starting dose and adjust doses of atorvastatin based on disease-specific guidelines. Prescriber should be aware of possible increased risk for myopathy especially for 40mg dose. If dose >40mg needed for desired efficacy, consider combination therapy (i.e., atorvastatin plus non-statin guideline directed medical therapy). (Moderate)	
Lovastatin	Increased lovastatin acid exposure as compared to normal function which may translate to increased myopathy risk.	Prescribe an alternative statin depending on the desired potency (see Page 4 for recommendations for alternative statins). If lovastatin therapy is warranted, limit dose to ≤20mg/day. (Moderate)	
Pitavastatin	Increased pitavastatin exposure as compared to normal function which may translate to increased myopathy risk.	Prescribe ≤ 2mg as a starting dose and adjust doses of pitavastatin based on disease-specific guidelines. Prescriber should be aware of possible increased risk for myopathy especially for doses >1mg. If dose >2mg needed for desired efficacy, consider an alternative statin (see Page 4 for recommendations for alternative statins) or combination therapy (i.e. pitavastatin plus non- statin guideline directed medical therapy). (Moderate)	
Pravastatin	Increased pravastatin exposure as compared to normal function; Typical myopathy risk with doses ≤40 mg.	Prescribe desired starting dose and adjust doses of pravastatin based on disease-specific guidelines. Prescriber should be aware of possible increased risk for myopathy with pravastatin especially with doses >40mg per day. (Moderate)	
Simvastatin	Increased simvastatin acid exposure as compared to normal function; increased risk of myopathy.	Prescribe an alternative statin depending on the desired potency (see Page 4 for recommendations for alternative statins). If simvastatin therapy is warranted, limit dose to <20mg/day. (Strong)	

Anterpretive drug information is only available for certain genetic results. Medical Insights and Therapeutic Recommendations information are extracted from CPIC® guidelines.



ACUMEN DIAGNOSTICS PTE LTD.

Tel: 6980 0080 HCI: 2010257 41 Science Park Road #01-02 Singapore 117610

Report Section 3:

Therapeutic Recommendations based on SLCO1B1/ABCG2 and SLCO1B1/CYP2C9 genotyping result combinations

Medications Affected	Therapeutic Recommendations^ (Classification of Recommendations)
Rosuvastatin	Prescribe desired starting dose and adjust doses of rosuvastatin based on disease-specific
(Based on SLCO1B1-ABCG2 genotyping result combinations)	and specific population guidelines. Prescriber should be aware of possible increased risk for myopathy especially for doses>20mg. (Strong)
Fluvastatin	
(Based on SLCO1B1-CYP2C9	Prescribe an alternative statin depending on the desired potency (see Page 4 for recommendations for alternative statins). (Optional)

genotyping result
combinations)

Aleterpretive drug information is only excitable for contain genetic results. Medical Insights and Therepretic Recom

^Interpretive drug information is only available for certain genetic results. Medical Insights and Therapeutic Recommendations information are extracted from CPIC® guidelines.



ACUMEN DIAGNOSTICS PTE LTD.

Tel: 6980 0080 HCI: 2010257 41 Science Park Road #01-02 Singapore 117610

Reference:

SLCO1B1 recommendations with intensity and statin dose stratified by SLCO1B1 phenotype; all doses assume adult dosing according to the CPIC®



https://doi.org/10.1002/cpt.2557

SLCOIBI decreased function

High intensity statina

Low SAMS risk with: Rosuvastatin 20 mg

Moderate SAMS risk with: Atorvastatin 40 mg Rosuvastatin 40 mg

High SAMS risk with: Atorvastatin 80 mg

Moderate intensity statin^a

Low SAMS risk with: Atorvastatin 10-20 mg Pitavastatin I mg Pravastatin 40 mg Rosuvastatin 5-10 mg

Moderate SAMS risk with: Fluvastatin 80 mg Pitavastatin 2 mg Pravastatin 80 mg

High SAMS risk with: Lovastatin 40-80 mg Pitavastatin 4 mg Simvastatin 20-40 mg

Low intensity statin^a

Low SAMS risk with: Fluvastatin 20-40 mg Pravastatin 10-20 mg

Moderate SAMS risk with: Lovastatin 20 mg Simvastatin 10 mg

Legend: Light gray boxes: Prescribe stated starting dose. Dark gray boxes: Prescriber should be aware of possible increased risk of increased exposure and myopathy. Black boxes: Consider a reduced dose or alternative statin. All boxes: Doses indicated are total daily dose. Dose recommendations are based on clinical toxicity data when available. aStatin intensity as recommended by current American College of Cardiology/American Heart Association guidelines.

SLCO IBI poor function

High intensity statina

Low SAMS risk with: Rosuvastatin 20 mg

High SAMS risk with: Atorvastatin 40-80 mg Rosuvastatin 40 mg

Moderate intensity statin^a

Low SAMS risk with: Atorvastatin 10-20 mg Pitavastatin I mg Pravastatin 40 mg Rosuvastatin 5-10 mg

Moderate SAMS risk with: Fluvastatin 80 mg Pravastatin 80 mg

High SAMS risk with: Lovastatin 40-80 mg Pitavastatin 2-4 mg Simvastatin 20-40 mg

Low intensity statin^a

Low SAMS risk with: Fluvastatin 20-40 mg Pravastatin 10-20 mg

High SAMS risk with: Lovastatin 20 mg Simvastatin 10 mg