

Patient Name TESTINGRNV,GTPMT NEG	Patient ID SA00065671	Age 39	Gender M	Order # SA00065671
Ordering Phys CLIENT,CLIENT				DOB 01/01/1975
Client Order # SA00065671	Account Information			Report Notes
Collected 02/18/2014 06:00	C7028846-DLMP Rochester SDSC 2 - Client Support			
Printed 03/10/2014 13:51	Rochester, MN 55901			

Test	Flag	Results	Unit	Reference Value	Perform Site*
TPMT Genotype, B					
RECEIVED: 02/19/2014 08:28 REPORTED: 02/19/2014 08:58					
TPMT Genotype Result		1/1			MCR
TPMT *2, *3A, *3B and *3C were not detected. This individual most likely has a TPMT *1/*1 genotype.					
TPMT Interpretation		This patient most likely has extensive (normal) TPMT activity. Dosing guidance for thiopurines can be found at: CPIC guidelines http://www.pharmgkb.org/gene/PA356 .			MCR
Since some adverse reactions to thiopurine drugs, including myelosuppression, are not explained by TPMT (Genbank # NM_000367.2), regular monitoring of complete blood count (CBC) and liver function tests is still essential.					
This test detects TPMT*2, *3A, *3B, and *3C. If these alleles are not detected, the patient most likely has the *1/*1 genotype. There is a small residual risk that other rare alleles may be present which are not detected by this assay and which might affect the patients response to thiopurine drugs. This genotyping method will not distinguish between a heterozygous *3A and the very rare *3B/*3C, which is associated with poor (deficient) enzyme activity. Evaluation of enzyme activity is necessary to definitively identify this rare genotype (Order Mayo Test ID TPMT, secondary ID 80291, Published name: Thiopurine Methyltransferase (TPMT), Erythrocyte using the specimen requirements for this test).					
Inhibitors: Co-prescription of allopurinol might inhibit TPMT activity. Drugs that have been shown to inhibit TPMT activity include: naproxen, ibuprofen, ketoprofen, furosemide, sulfasalazine, mesalamine, olsalazine, mefenamic acid, thiazide diuretics, and benzoic acid inhibitors.					
If the patient has had an allogeneic blood or marrow transplant or a recent (i.e. less than 6 weeks from time of sample collection) heterologous blood transfusion these results may be inaccurate due to the presence of donor DNA. Laboratory developed test.					
Reviewed By		See Below			MCR
Result:Linnea M. Baudhuin, Ph.D.					

Performing Site Legend on Last Page of Report

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* Report times for Mayo performed tests are CST/CDT

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>> Accession SA00065671 - Continued From Previous Page <<
 >> Do Not Discard <<

* Performing Site:

MCR	Mayo Clinic Laboratories - Rochester Main Campus 200 First St SW Rochester, MN 55905	Lab Director:
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Patient Name TESTINGRNV,GTPMT NEG	Collection Date and Time 02/18/2014 06:00	Report Status Final
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