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**Lab Dept:** Chemistry

**Test Name:** THIOPURINE METHYLTRANSFERASE, RBC

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***General Information***

**Lab Order Codes:** TMT

**Synonyms:** TPMT (Thiopurine Methyltransferase); Myelotoxicity

**CPT Codes:** 83789 – Mass Spectrometry, Tandem Mass Spectrometry ,NOS

**Test Includes:** Thiopurine Methyltransferase level reported in U/mL RBC.

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***Logistics***

**Test Indications:** Useful for detection of individuals with low TPMT activity who will have excessive myelosuppression or severe hematopoietic toxicity when taking azathioprine (Imuran) or 6-MP (Purinethol).

**Lab Testing Sections:** Chemistry - Sendouts

**Referred to:** Mayo Medical Laboratories (MML Test#: TPMT/80291), temporary forward to ARUP (FATPM)

**Phone Numbers:** MIN Lab: 612-813-6280

STP Lab: 651-220-6550

**Test Availability:** Daily, 24 hours

**Turnaround Time:** 2 – 4 days

**Special Instructions:** N/A

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***Specimen***

**Specimen Type:** Whole blood

**Container:** Green top (Sodium Heparin) tube  
Alternate: Lavender top (EDTA) or Green top (Lithium Heparin) tube

**Draw Volume:** 5 mL (Minimum: 3 mL) blood

**Processed Volume:** Same as Draw Volume

**Collection:** Routine venipuncture

**Special Processing:** Lab Staff: **Do Not** Centrifuge. Store specimen at refrigerated temps before shipping. Stable for 6 days at 2-8 degrees centigrade. Forward specimen to reference lab on a cold pack. **Do Not** Freeze. Forward promptly.

**Patient Preparation:** None

**Sample Rejection:** Gross hemolysis; mislabeled or unlabeled specimens

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**Interpretive**

**Reference Range:**

Reference ranges apply to all ages	
Normal	24.0 – 44.0 U/mL Individuals are predicted to be at low risk of bone marrow toxicity (myelosuppression) as a consequence of standard thiopurine therapy; no dose adjustment is recommended.
Intermediate:	17.0 – 23.9 U/mL Individuals are predicted to be at intermediate risk of bone marrow toxicity (myelosuppression) as a consequence of standard thiopurine therapy; a dose reduction and therapeutic drug management is recommended.
High:	>44.0 U/mL Individuals are not predicted to be at risk for bone marrow toxicity (myelosuppression) as a consequence of standard thiopurine dosing, but may be at risk for therapeutic failure due to excessive inactivation
Low:	<17.0 U/mL Individuals are not predicted to be at high risk for bone marrow toxicity (myelosuppression) as a consequence of standard thiopurine dosing. It is recommended to avoid use of thiopurine drugs.

**Critical Values:** N/A

**Limitations:** There is a partial overlap for the distribution of thiopurine methyltransferase activities observed between patients with normal and heterozygous genotypes; thus, results that fall within the low normal range can occur because of assay variability or biological variation.

**Methodology:** Enzymatic End Point/Liquid Chromatography-Tandem Mass Spectrometry (LC-MS/MS)

**References:** [Mayo Medical Laboratories](#) December 2015

**Updates:**  
3/8/2012: Tube type changed from Green to Lavender. Temporary change due to test issue at Mayo. Mayo is forwarding testing to Prometheus until further notification.  
12/26/2012: Test reinstated at MML. Updated reference ranges.  
12/7/2015: Updated for temporary test referral to ARUP. Internal test as MML is unavailable.