

<b>Patient Name</b> TESTINGRNV,QTBG_ABNORM	<b>Patient ID</b> SA00060672	<b>Age</b> 29	<b>Gender</b> M	<b>Order #</b> SA00060672
<b>Ordering Phys</b> CLIENT,CLIENT				<b>DOB</b> 05/05/1984
<b>Client Order #</b> SA00060672	<b>Account Information</b>			<b>Report Notes</b>
<b>Collected</b> 08/13/2013 00:00	C7028846-DLMP Rochester 3050 Superior Drive Rochester, MN 55901			
<b>Printed</b> 09/17/2013 12:27				

Test	Flag	Results	Unit	Reference Value	Perform Site*
<b>M. tuberculosis by QuantiFERON, B</b>	AB	Indeterminate		Negative	SDL
<b>RECEIVED:</b> 08/14/2013 07:23 <b>REPORTED:</b> 08/14/2013 08:02 Indeterminate result due to High Nil (negative control) value. This may occur due to heterophile antibody effects or non-specific gamma interferon in the patient's blood sample. Suggest retesting of new specimen if clinically indicated.					
Tuberculosis Antigen Value		0.50	IU/mL		SDL
This is a qualitative test. The TB antigen IU/mL value is required for documentation on certain government reporting forms (e.g., Form I-693), but this value should not be used to monitor disease progression or response to therapy.					
Diagnosing or excluding tuberculosis disease, and assessing the probability of LTBI, require a combination of epidemiological, historical, medical, and diagnostic findings that should be taken into account when interpreting QuantiFERON-TB results.					

\* Performing Site:

SDL	Mayo Clinic Laboratories - Rochester Superior Drive 3050 Superior Dr. NW Rochester, MN 55901	Lab Director:
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<b>Patient Name</b> TESTINGRNV,QTBG_ABNORM	<b>Collection Date and Time</b> 08/13/2013 00:00	<b>Report Status</b> Final
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\* Report times for Mayo performed tests are CST/CDT