

1-800-533-1710

<b>PATIENT NAME</b> TESTING, 91545		<b>PATIENT NUMBER</b> L3MRNW4123564		<b>AGE</b> 51	<b>SEX</b> F	<b>ACCESSION #</b> W4123564
<b>ORDERING PHYSICIAN</b>		<b>CLIENT ORDER #</b>			<b>ACCOUNT #</b> LIAISONS	
<b>COLLECTION</b> 08/02/11 01:52 P	<b>RECEIVED</b> 08/02/11 01:52 P	<b>REPORT PRINTED</b> 08/03/11 08:11 A		<b>SPECIMEN INFORMATION</b> DATE OF BIRTH: 6/19/1960		
<b>DATE</b>	<b>TIME</b>	<b>DATE</b>	<b>TIME</b>			
Test Client Attn: Mayo Liaisons 200 First Street SW Rochester, MN 55905 507-284-8202						

TEST REQUESTED	HI	LO	REF RANGE	PERFORM SITE *
<b>Myositis Ab 2 Panel</b>			<b>REPORTED: 08/02/11 01:55 P</b>	
JO-1	Negative		Negative	REF
	Negative	<20 units		
	Weak Positive	20-39 units		
	Moderate Positive	40-80 units		
	Strong Positive	>80 units		
MI-2	Negative		Negative	REF
PL-7	Negative		Negative	REF
PL-12	Negative		Negative	REF
EJ	Weak Positive		Negative	REF
OJ	Negative		Negative	REF
SRP	Negative		Negative	REF
KU	Negative		Negative	REF
PM/SCL	Negative		Negative	REF
U2 SN RNP	Negative		Negative	REF

This test was developed and its performance characteristics validated by RDL Inc. The FDA has determined that approval for this test is not necessary. This is an analyte specific reagent (ASR) test.

Test Performed by: RDL Reference Laboratory, Inc.  
 10755 Venice Blvd  
 Los Angeles, CA 90034

\* PERFORMING SITE

<b>PATIENT NAME</b> TESTING, 91545	<b>ORDER STATUS</b> Final	<b>COLLECTION DATE AND TIME</b> 08/02/11 01:52 P
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