

Patient ID SA00166153	Patient Name TESTINGRNV, SMPU A	Birth Date 1980-09-09	Sex M	Age 43
Order Number SA00166153	Client Order Number SA00166153	Ordering Physician CLIENT,CLIENT	Report Notes	
Account Information C7028846 DLMP Rochester		Collected 31 Jan 2024 06:00		

Protein Electrophoresis, 24 HR, U

Albumin	SDL	A/G Ratio	SDL
60.0 mg/24 h		0.06	
Alpha-1 globulin	SDL	M spike	SDL
10.0 mg/24 h		 181 mg/24 h	
Alpha-2 globulin	SDL	Impression	① SDL
60.0 mg/24 h		M-spike in gamma fraction.	
Beta globulin	SDL	See Isotype.	
90.0 mg/24 h		Received: 31 Jan 2024 10:19	Reported: 31 Jan 2024 10:28
Gamma globulin	SDL		
770.0 mg/24 h			

Performing Site Legend

Code	Laboratory	Address	Lab Director	CLIA Certificate
SDL	Mayo Clinic Laboratories - Rochester Superior Drive	3050 Superior Drive NW, Rochester MN 55905	William G. Morice M.D. Ph.D.	24D1040592
MCR	Mayo Clinic Laboratories - Rochester Main Campus	200 First Street SW, Rochester, MN 55905	William G. Morice M.D. Ph.D.	24D0404292



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M-protein Screen, 24 HR, U

M-protein Mass-Fix, 24 HR, U

M-protein Isotype MS, 24 HR, U

Kappa, monoclonal

ADDITIONAL INFORMATION

The submitted sample was assayed by five separate immunopurifications for IgG, IgA, IgM, kappa and lambda. The result reflects the findings of either no monoclonal protein detected or those monoclonal immunoglobulins that were detected.

② SDL

Flag M-protein Isotype MS, 24 HR, U



Positive

SDL

Reference Value
Negative

Protein, Total, 24 HR, U

Total Protein, 24 HR, U



1000 mg/24 h

MCR

Reference Value
<229

Urine Volume

1000 mL

MCR

Received: 31 Jan 2024 10:19

Reported: 31 Jan 2024 10:23

Collection Duration

24 h

MCR

Laboratory Notes

- ① This test has been modified from the manufacturer's instructions. Its performance characteristics were determined by Mayo Clinic in a manner consistent with CLIA requirements. This test has not been cleared or approved by the U.S. Food and Drug Administration.
- ② This test was developed and its performance characteristics determined by Mayo Clinic in a manner consistent with CLIA requirements. This test has not been cleared or approved by the U.S. Food and Drug Administration.

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