



Prostate-Specific Antigen (PSA) Diagnostic, Serum

Patient ID	Patient Name		Birth Date	Gender	Age
SA00591936	TESTINGRNV, REPORTS NORM		1952-08-16	M	67
Order Number	Client Order Number	Ordering Physician	Report Notes		
SA00591936	SA00591936	CLIENT, CLIENT			
Account Information		Collected			
C7028846 DLMP Rochester		24 Sep 2019 09:00			

Prostate-Specific Ag Diagnostic, S

SDL

3.1 ng/mL

 $\begin{array}{c} \text{Reference Value} \\ \leq 4.5 \end{array}$

ADDITIONAL INFORMATION

The testing method is an electrochemiluminescence assay manufactured by Roche Diagnostics Inc. and performed on the Modular or Cobas system.

Values obtained with different assay methods or kits may be different and cannot be used interchangeably.

Test results cannot be interpreted as absolute evidence for the presence or absence of malignant disease.

Received: 25 Sep 2019 13:28 **Reported:** 25 Sep 2019 13:28

Performing Site Legend

Code	Laboratory	Address	Lab Director	CLIA Certificate
SDL	Mayo Clinic Laboratories - Rochester Superior Drive	3050 Superior Drive NW, Rochester MN 55901	William G. Morice M.D. Ph.D	24D1040592