

292/05, Tulsidas Marg, Basement Chowk, Lucknow-226 003

Phone: 0522-4062223, 9305548277, 8400888844 9415577933, 9336154100, Tollfree No.: 8688360360

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CMO Reg. No. RMEE 2445133 NABL Reg. No. MC-2491 Certificate No. MIS-2023-0218

Patient Name : Mr. DAYAL SHARAN SINGH

Age/Gender : 75 Y/M

Lab No : 10144222

Referred By : Dr.WE CARE HOSPITAL

Refer Lab/Hosp : CHARAK NA

P.R.

Doctor Advice : 2D ECHO,USG KUB,PSA-TOTAL,URIC ACID,UACR

Visit No : CHA250046927

Registration ON : 17/Mar/2025 11:33AM

Sample Collected ON : 17/Mar/2025 11:50AM

Sample Received ON : 17/Mar/2025 12:07PM

Report Generated ON : 17/Mar/2025 04: 28PM



Test Name	Result	Unit	Bio. Ref. Range	Method		
URIC ACID						
Sample Type : SERUM						
SERUM URIC ACID	8.4	mg/dL	2.40 - 5.70	Uricase,Colorimetric		
URINE ALBUMIN CREATININE RATIO						
URINE FOR MICRO ALBUMIN	24	MG/L	< 20 MG/L			
URINARY CREATININE	86.80	mg/dL	20-320 mg/dL			
URINE ALBUMIN CREATININE RATIO	27.6	mg/g		calculated		









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Report Generated ON : 17/Mar/2025 02:31PM



Test Name	Result	Unit	Bio. Ref. Range	Method
PSA-TOTAL				
PROSTATE SPECIFIC ANTIGEN	0.4	ng/mL	0.2-4.0	CLIA

COMMENT: 1. Prostate specific antigen (PSA) is useful for diagnosis of disseminated CA prostate & its equential measurement is the most sensitive measure of monitoring treatment of disseminated CA prostate with its shorter half life (half life of 2.2 days only) it is superior to prostatic acis phosphatase(PAP). PSA is elevated in nearly all patients with stage D carcinoma whereas PAP is elevated in only 45 % of patient. Mild PSA elevation are also reported in some patients of BHP. 2. Blood samples should be obtained before prostate biopsy or prostatecomy or prostatic massage or digital pre rectal examination as it may result intrasient levation of PSA value for few days.

NOTE: - PSA values obtained in different types of PSA assay methods cannot be used interchangeably as the PSA value in a given sample varies with assays from different manufactures due to difference in assay methodology and reagent specificity. If in the course of monitoring a patient the assay method used for determination is changed, additional sequential testing should be carried out to confirm baseline value.

DONE BY:

Enhanced Chemiluminescence "VITROS ECI"

\*\*\* End Of Report \*\*\*

CHARAK



