

Patient Name : Mr.SANTOSH PURI	Visit No : CHA250046651
Age/Gender : 56 Y/M	Registration ON : 17/Mar/2025 09:11AM
<b>Lab No : 10143946</b>	Sample Collected ON : 17/Mar/2025 09:13AM
Referred By : Dr.D DALELA**	Sample Received ON : 17/Mar/2025 09:13AM
Refer Lab/Hosp : CHARAK NA	Report Generated ON : 17/Mar/2025 10:25AM
Doctor Advice : URINE COM. EXMAMINATION,APTT,PSA-TOTAL,PT/PC/INR	



Test Name	Result	Unit	Bio. Ref. Range	Method
<b>PT/PC/INR</b>				
PROTHROMBIN TIME	13 Second		13 Second	Clotting Assay
Prothromin concentration	100 %		100 %	
INR (International Normalized Ratio)	1.00		1.0	

<b>APTT</b>
<b>Sample Type : SODIUM CITRATE</b>

<b>APTT</b>				
APTT Patient Value	29 Seconds	Seconds	26 - 38	Clotting Assay

**INTERPRETATION**

Determination of APTT helps in estimating abnormality in most of the clotting factors of the intrinsic pathway including congenital deficiency of factor VIII, IX, XI, and XII and is also a sensitive procedure for generating heparin response curve for monitoring heparin therapy.

**Causes of a prolonged APTT:**

- Disseminated intravascular coagulation.
- Liver disease.
- Massive transfusion with stored blood.
- Administration of heparin or contamination with heparin.
- A circulating anticoagulant.
- Deficiency of a coagulation factor other than factor VII.
- APTT is also moderately prolonged in patients on oral anticoagulant drugs and in the presence of Vitamin K deficiency.

**Limitations of assay:**

- Abnormalities of coagulation factor VII, factor XIII and platelets are not detected by this test procedure.
- Platelet factor IV, a heparin neutralizing factor can be released due to platelet aggregation or damage and may influence the test.
- Decrease in APTT time is observed in males under estrogen therapy and oral contraceptive administration in females.
- APTT based heparin therapeutic range is not established for this assay.

[Checked By]

Print.Date/Time: 17-03-2025 11:10:58

\*Patient Identity Has Not Been Verified. Not For Medicolegal



*Sharma*

DR. NISHANT SHARMA DR. SHADAB Dr. SYED SAIF AHMAD  
PATHOLOGIST PATHOLOGIST MD (MICROBIOLOGY)

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Test Name	Result	Unit	Bio. Ref. Range	Method
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**URINE EXAMINATION REPORT**

Colour-U	STRAW		Light Yellow	
Appearance (Urine)	CLEAR		Clear	
Specific Gravity	<b>1.005</b>		1.005 - 1.025	
pH-Urine	Acidic (6.0)		4.5 - 8.0	
PROTEIN	Absent	mg/dl	ABSENT	Dipstick
Glucose	Absent			
Ketones	Absent		Absent	
Bilirubin-U	Absent		Absent	
Blood-U	Absent		Absent	
Urobilinogen-U	0.20	EU/dL	0.2 - 1.0	
Leukocytes-U	Absent		Absent	
NITRITE	Absent		Absent	
<b>MICROSCOPIC EXAMINATION</b>				
Pus cells / hpf	Nil	/hpf	< 5/hpf	
Epithelial Cells	2-3	/hpf	0 - 5	
RBC / hpf	Nil		< 3/hpf	

CHARAK

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<b>Lab No : 10143946</b>	Sample Collected ON : 17/Mar/2025 09:13AM
Referred By : Dr.D DALELA**	Sample Received ON : 17/Mar/2025 09:28AM
Refer Lab/Hosp : CHARAK NA	Report Generated ON : 17/Mar/2025 10:37AM
Doctor Advice : URINE COM. EXMAMINATION,APTT,PSA-TOTAL,PT/PC/INR	



Test Name	Result	Unit	Bio. Ref. Range	Method
<b>PSA-TOTAL</b>				
PROSTATE SPECIFIC ANTIGEN	1.00	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



MC-2491

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