



P K R JAIN HEALTHCARE INSTITUTE

NASIRPUR, Hissar Road, AMBALA CITY- (Haryana)

A PIONEER DIAGNOSTIC CENTRE

☎ 0171-2532620, 8222896961 ✉ pkrijainhealthcare@gmail.com

TEST PERFORMED AT KOS DIAGNOSTIC LAB, AMBALA CANTT.

NAME	: Mrs. KAVITA	PATIENT ID	: 1644724
AGE/ GENDER	: 42 YRS/FEMALE	REG. NO./LAB NO.	: 122410160005
COLLECTED BY	:	REGISTRATION DATE	: 16/Oct/2024 10:36 AM
REFERRED BY	:	COLLECTION DATE	: 16/Oct/2024 10:37AM
BARCODE NO.	: 12505195	REPORTING DATE	: 16/Oct/2024 04:52PM
CLIENT CODE.	: P.K.R JAIN HEALTHCARE INSTITUTE		
CLIENT ADDRESS	: NASIRPUR, HISSAR ROAD, AMBALA CITY - HARYANA		

Test Name	Value	Unit	Biological Reference interval
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CLINICAL CHEMISTRY/BIOCHEMISTRY

CREATININE PHOSPHOKINASE-MB (CPK-MB)

CPK-MB - SERUM <i>by EFIA (ENZYM FLUORESCENT IMMUNOASSAY)</i>	3.01	ng/mL	0.0 - 5.0
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Interpretation:-

- Alternative name of Creatine Kinase (CK) is Creatine phospho-kinase(CPK).
- Creatine Kinase (CK) is a dimeric enzyme composed of two types of monomer sub-units (i.e. M-Muscular & B-Brain), which combine to form three distinct CK isoenzymes.
 - CK-BB(CK-I), is produced primarily by brain, lungs and smooth muscles, and enters the blood only on injury to these organs like cerebrovascular accidents or pulmonary infarctions.
 - CK-MB (CK-II), is produced primarily by heart muscle;
 - CK-MM (CK-III), is produced primarily by skeletal muscle.
- Normally very little CK is found circulating in the blood. Elevated levels indicate damage to either muscle or brain possibly from a myocardial infarction, muscle disease, or stroke.
- CK levels are reduced in first half of pregnancy, and increased in second half of pregnancy.

Increased:-

Physiological:-

- Strenuous physical activity.
- New Born.

Pathological :-

- Myocardial & pulmonary infarction
- Accident and recent surgery.
- Drugs:- Statins.
- Convulsions & brain tumour.
- Myopathies
- Malignant hyperthermia
- Hypothyroidism & Hyperthyroidism

5).CK-MB (CK-II) levels increase significantly 4-6 hours following a myocardial infarction and peak at around 12-24 hours after the infarct. The levels return to normal, in case of no further myocardial damage, after 24 to 48 hours. Hence the increased levels of CK-MB along with elevated levels of total CK is a good indicator of myocardial infarction.

6).For diagnosis of MI with high sensitivity and specificity, serial sampling over a period of 8 to 12 hours is required. For accurate diagnosis of myocardial infarction, CK-MB activity along with total CK should be measured. If the total CK activity is raised and CK-MB contributes more than 6% of the total activity, then myocardial infarction is considered highly probable.



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IMMUNOPATHOLOGY/SEROLOGY

TROPONIN I ULTRASENSITIVE (QUANTITATIVE)

TROPONIN I ULTRASENSITIVE (QUANTITATIVE) 0.07 ng/mL < 0.50

by ELFA (ENZYME LINKED FLUORESCENT IMMUNOASSAY), NEXT GENERATION, ULTRASENSITIVE

INTERPRETATION:

NOTE:

1.False negative / positive results are observed in patients receiving mouse monoclonal antibodies for diagnosis or therapy.

COMMENTS

- 1.Troponin is a regulatory complex of 3 proteins that resides at regular intervals in the thin filament of striated muscle.
- 2.Cardiac Troponin is a cardiospecific, highly sensitive marker of myocardial damage and has never shown to be expressed in normal, regenerating or diseased skeletal muscle.
- 3.In cases of acute myocardial damage, Troponin I levels rise in serum about 3-4 hours after appearance of cardiac symptoms and remain elevated upto 10 days.
- 4.It is an independent prognostic marker which can predict near, mid and long term outcome in patients with Acute Coronary Syndrome (ACS).

INCREASED LEVELS

- 1.Congestive Heart Failure
- 2.Cardiomyopathy
- 3.Myocarditis
- 4.Heart contusion
- 5.Interventional therapy like cardiac surgery and drug induced cardiotoxicity

USES

- 1.To differentiate patients with Non ST elevation Myocardial Infarction (NSTMI) from Unstable angina-patients with ACS with elevated Troponin I and / or CK-MB are considered to have NSTMI whereas the diagnosis of Unstable angina is established if Troponin I and CK-MB are within the normal range.
- 2.Ideally Troponin I should be measured at presentation (0 hour) and repeated after 6-9 hours & 12-24 hours if earlier specimens are normal and the clinical suspicion is high.
3. Risk stratification of patients presenting with ACS and for cardiac risk in patients with Chronic Renal Failure. As it offers powerful risk assessment, in ACS, Troponin I monitoring should be included in practice guidelines.
4. For selection of more intensive therapy and intervention in patients with elevated Troponin I.



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SPECIAL INVESTIGATIONS

N-TERMINAL PRO B TYPE NATRIURETIC PEPTIDE (NT-PRO BNP)

N-TERMINA PRO B TYPE NATRIURETIC PEPTIDE (NT-PRO BNP) 203 pg/mL < 300

by ELFA (ENZYME LINKED FLOURESCENT ASSAY)

INTERPRETATION:

AGE AND CONDITION RELATED CUT OFF VALUES FOR NT-PRO BNP

IN ACUTE HEART FAILURE		
AGE (Years)	UNITS (pg/mL)	OPTIMAL CUT OFF VALUE
< 50	pg/mL	450
50 - 75	pg/mL	900
>75	pg/mL	1800
IN CHRONIC HEART FAILURE		
< 75	pg/mL	125
>75	pg/mL	450
NEGATIVE PREDICTIVE VALUE CUT OFF FOR NT-PRO BNP: < 300 pg/ml (HEART FAILUE UNLIKELY)		

The N-terminal of the prohormone brain natriuretic peptide (NT-proBNP), is a 76 amino acid terminal inactive protein that is cleaved from proBNP to release brain natriuretic peptide.

The main physiological function of NP is homeostasis and protection of among others the cardiovascular (CV) system from the effects of volume overload. They play an important role in regulating blood pressure (BP) and body fluid volume by their natriuretic and diuretic actions, arterial dilatation, and inhibition of the renin angiotensin system.

Concentrations of NP increase in patients with congestive heart failure (CHF) and other CV diseases owing to pressure and volume overload, whereas levels below cutoff are a strong negative predictor for CHF.

Both BNP and NT-proBNP levels in the blood are used for screening, diagnosis of acute congestive heart failure (CHF) and may be useful to establish prognosis in heart failure, as both markers are typically higher in patients with worse outcome. The plasma concentrations of both BNP and NT-proBNP are also typically increased in patients with asymptomatic or symptomatic left ventricular dysfunction and is associated with coronary artery disease and myocardial ischemia

It can be used, along with other cardiac biomarkers test, to detect heart stress and damage and/or along with lung function tests to distinguish



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between causes of shortness of breath. Heart failure can be confused with other conditions, and it may co-exist with them. BNP and NT-proBNP levels can help doctors differentiate between heart failure and other problems, such as lung disease. An accurate diagnosis is important because the treatments are often different and must be started as soon as possible.

A BNP or NT-proBNP test may be ordered when a person has signs and symptoms that could be due to heart failure. These may include:

1. Difficulty breathing, shortness of breath
2. Fatigue
3. Swelling in the feet, ankles, legs, abdomen

NOTE:


1. Lack of NT-ProBNP elevation has been reported if Congestive Heart Failure (CHF) is very acute (first hour) or if there is Ventricular inflow obstruction
2. As per a number of studies, threshold for NT-ProBNP is 125 pg/mL
3. BNP and NT-proBNP levels decrease in most people who are taking drug therapies for heart failure, such as angiotensin-converting enzyme (ACE) inhibitors, beta blockers and diuretics.
4. Levels of both BNP and NT-proBNP tend to increase with age.
5. Levels of NT-proBNP and BNP may be increased in persons with kidney disease due to reduced clearance.
6. While both BNP and NT-proBNP will rise with left ventricle dysfunction and either can be measured for diagnosis or monitoring therapy, they are not interchangeable and the results cannot be directly compared.
7. Results to be clinically correlated.


CLINICAL USE:

1. As an aid in the diagnosis of suspected cases of CHF
2. Detection of mild forms of cardiac dysfunction
3. To assess severity of heart failure in already diagnosed cases of CHF
4. For risk stratification of patients with Acute Coronary Syndrome & CHF For monitoring therapy in patients with Left Ventricular dysfunction

*** End Of Report ***




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