

KOS Diagnostic Lab

(A Unit of KOS Healthcare)



Dr. Vinay Chopra
MD (Pathology & Microbiology)
Chairman & Consultant Pathologist

Dr. Yugam Chopra MD (Pathology) CEO & Consultant Pathologist

NAME : Mr. BALRAJ KHURANA

AGE/ GENDER : 65 YRS/MALE PATIENT ID : 1605983

COLLECTED BY : REG. NO./LAB NO. : 042409080001

 REFERRED BY
 : 08/Sep/2024 08:48 AM

 BARCODE NO.
 : A0465446
 COLLECTION DATE
 : 08/Sep/2024 11:20AM

 CLIENT CODE.
 : KOS DIAGNOSTIC SHAHBAD
 REPORTING DATE
 : 08/Sep/2024 01:08PM

CLIENT ADDRESS: 6349/1, NICHOLSON ROAD, AMBALA CANTT

Test Name Value Unit Biological Reference interval

CLINICAL CHEMISTRY/BIOCHEMISTRY LIVER FUNCTION TEST (COMPLETE)

BILIRUBIN TOTAL: SERUM by DIAZOTIZATION, SPECTROPHOTOMETRY	0.52	mg/dL	INFANT: 0.20 - 8.00 ADULT: 0.00 - 1.20
BILIRUBIN DIRECT (CONJUGATED): SERUM by DIAZO MODIFIED, SPECTROPHOTOMETRY	0.15	mg/dL	0.00 - 0.40
BILIRUBIN INDIRECT (UNCONJUGATED): SERUM by CALCULATED, SPECTROPHOTOMETRY	0.37	mg/dL	0.10 - 1.00
SGOT/AST: SERUM by IFCC, WITHOUT PYRIDOXAL PHOSPHATE	23.2	U/L	7.00 - 45.00
SGPT/ALT: SERUM by IFCC, WITHOUT PYRIDOXAL PHOSPHATE	18.4	U/L	0.00 - 49.00
AST/ALT RATIO: SERUM by CALCULATED, SPECTROPHOTOMETRY	1.26	RATIO	0.00 - 46.00
ALKALINE PHOSPHATASE: SERUM by Para nitrophenyl phosphatase by amino methyl propanol	84.87	U/L	40.0 - 130.0
GAMMA GLUTAMYL TRANSFERASE (GGT): SERUM by SZASZ, SPECTROPHTOMETRY	12.7	U/L	0.00 - 55.0
TOTAL PROTEINS: SERUM by BIURET, SPECTROPHOTOMETRY	5.92 ^L	gm/dL	6.20 - 8.00
ALBUMIN: SERUM by BROMOCRESOL GREEN	3.65	gm/dL	3.50 - 5.50
GLOBULIN: SERUM by CALCULATED, SPECTROPHOTOMETRY	2.27 ^L	gm/dL	2.30 - 3.50
A : G RATIO: SERUM by CALCULATED, SPECTROPHOTOMETRY	1.61	RATIO	1.00 - 2.00

INTERPRETATION

NOTE:- To be correlated in individuals having SGOT and SGPT values higher than Normal Referance Range.

USE:- Differential diagnosis of diseases of hepatobiliary system and pancreas.

INCREASED:

DRUG HEPATOTOXICITY	> 2
ALCOHOLIC HEPATITIS	> 2 (Highly Suggestive)
CIRRHOSIS	1.4 - 2.0



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Test Name	Value	Unit	Biological Reference interval
INTRAHEPATIC CHOLESTATIS		> 1.5	
HEPATOCELLULAR CARCINOMA & CHRONIC HEPATITIS		> 1.3 (Slightly Increased)	
DEODEACED	•		<u> </u>

DECREASED:

- 1. Acute Hepatitis due to virus, drugs, toxins (with AST increased 3 to 10 times upper limit of normal)
- 2. Extra Hepatic cholestatis: 0.8 (normal or slightly decreased).

PROGNOSTIC SIGNIFICANCE:

NORMAL	< 0.65
GOOD PROGNOSTIC SIGN	0.3 - 0.6
POOR PROGNOSTIC SIGN	1.2 - 1.6



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Test Name Value Unit Biological Reference interval

TUMOUR MARKER

CANCER ANTIGEN 19.9 (CA 19.9): PANCREATIC CANCER MARKER

CANCER ANTIGEN (CA) -19.9: SERUM

1.788 U/mL 0.0 - 42.0

by CMIA (CHEMILUMINESCENCE MICROPARTICLE IMMUNOASSAY)

INTERPRETATION:

- 1.CA 19.9 isolated originally from colon cancer cell line has greatest utility in detecting pancreatic cancers and hence is the most useful circulating tumour marker for evaluating chronic pancreatic disorders.
- 2. The specificity and positive predictive value for cancers increase with higher CA 19.9 values.
- 3.Tumour size and histological grade affect the values, being higher in tumors > 3cms in diameter and in differentiated tumors.
- 4. High levels suggest tumour is unresectable. Used in conjunction with CT scan and other imaging modalities to decide about tumor resection.
- 5.Useful in predicting survival and recurrence after surgery. A persistent elevation following surgery may be indicative of occult metastasis or recurrence of disease.

INCREASED LEVELS ARE SEEN IN:

- 1.Pancreatic Cancer
- 2.. Cancers of bile duct, stomach, colon and oesophagus
- 3. Some non-gastrointestinal cancers
- 4.Hepatomas
- 5. Non-malignant conditions like hepatitis, cirrhosis, acute cholangitis pancreatitis and cystic fibrosis.

NOTE

- 1.CA 19.9 assay should be used as an adjunct with other diagnostic information in the management of pancreatic cancer.
- 2. The results obtained with different analytical techniques and different equipments cannot be used interchangeably due to difference in assay methods and reagent specificity.
- 3.In course of monitoring, the assay method preferably should not be changed



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Test Name Value Unit **Biological Reference interval**

CARCINO EMBRYONIC ANTIGEN (CEA)

CARCINO EMBRYONIC ANTIGEN (CEA): SERUM

0.52 < 5.0

by CLIA (CHEMILUMINESCENCE IMMUNOASSAY)

INTERPRETATION:

1. Carcinoembryonic antigen (CEA) is a glycoprotein normally found in embryonic entodermal epithelium.

2. Increased levels may be found in patients with primary colorectal cancer or other malignancies including medullary thyroid carcinoma and

breast, gastrointestinal tract, liver, lung, ovarian, pancreatic, and prostatic cancers.

3. Serial monitoring of CEA should begin prior to initiation of cancer therapy to verify post therapy decrease in concentration and to establish a baseline for evaluating possible recurrence. Levels generally return to normal within 1 to 4 months after removal of cancerous tissue.

CLINICAL SIGNIFICANCE:

1. Monitoring colorectal cancer and selected other cancers such as medullary thyroid carcinoma

May be useful in assessing the effectiveness of chemotherapy or radiation treatment.

NOTE:

1. Carcinoembryonic antigen levels should not be used for screening of the general population for undetected cancers.

2. Grossly elevated carcino-embryonic antigen (CEA) concentrations (>20 ng/mL) in a patient with compatible symptoms are strongly suggestive of the presence of cancer and also suggest metastasis.

3. Most healthy subjects (97%) have values < or =3.0 ng/mL.

4. After removal of a colorated tumor, the serum CEA concentration should return to normal by 6 weeks, unless there is residual tumor.

5. Increases in test values over time in a patient with a history of cancer suggest tumor recurrence.

End Of Report ***



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