

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

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

NAME	: Mrs. MEENA	PATIENT ID	: 1697504
AGE/ GENDER	: 57 YRS/FEMALE	REG. NO./LAB NO.	: 012412120032
COLLECTED BY	:	REGISTRATION DATE	: 12/Dec/2024 03:32 PM
REFERRED BY	:	COLLECTION DATE	: 12/Dec/2024 03:36PM
BARCODE NO.	: 01522360	REPORTING DATE	: 12/Dec/2024 06:16PM
CLIENT CODE.	: KOS DIAGNOSTIC LAB		
CLIENT ADDRESS	: 6349/1, NICHOLSON ROAD, AMBALA CANTT		

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	273.24^H	U/mL	0.00 - 41.0
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




DR.VINAY CHOPRA
 CONSULTANT PATHOLOGIST
 MBBS, MD (PATHOLOGY & MICROBIOLOGY)


DR.YUGAM CHOPRA
 CONSULTANT PATHOLOGIST
 MBBS, MD (PATHOLOGY)



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

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

NAME	: Mrs. MEENA	PATIENT ID	: 1697504
AGE/ GENDER	: 57 YRS/FEMALE	REG. NO./LAB NO.	: 012412120032
COLLECTED BY	:	REGISTRATION DATE	: 12/Dec/2024 03:32 PM
REFERRED BY	:	COLLECTION DATE	: 12/Dec/2024 03:36PM
BARCODE NO.	: 01522360	REPORTING DATE	: 12/Dec/2024 05:32PM
CLIENT CODE.	: KOS DIAGNOSTIC LAB		
CLIENT ADDRESS	: 6349/1, NICHOLSON ROAD, AMBALA CANTT		

Test Name	Value	Unit	Biological Reference interval
-----------	-------	------	-------------------------------

CARCINO EMBRYONIC ANTIGEN (CEA)

CARCINO EMBRYONIC ANTIGEN (CEA): SERUM 3.64 ng/mL < 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
2. 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 colorectal 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 ***




DR. VINAY CHOPRA
 CONSULTANT PATHOLOGIST
 MBBS, MD (PATHOLOGY & MICROBIOLOGY)


DR. YUGAM CHOPRA
 CONSULTANT PATHOLOGIST
 MBBS, MD (PATHOLOGY)

