



LABORATORY REPORT

Example Client, XYZ123
1234 Warde Road
Ann Arbor MI 48108

EXAMPLE, REPORT W
WX0000003827 M 07/08/1978 45 Y

Referral Testing

Collected: 08/30/2023 11:35 Received: 08/30/2023 11:35

Table with 6 columns: Test Name, Result, Flag, Ref-Ranges, Units, Site. Row 1: Carcinoembryonic Ag, Fluid; Carcinoembryonic Antigen, Fluid; 1.4; ng/mL; ARRL

INTERPRETIVE INFORMATION: Carcinoembryonic Antigen, Fluid

The Roche CEA electrochemiluminescent immunoassay was used. Results obtained with different assay methods or kits cannot be used interchangeably. The CEA assay value, regardless of level, should not be interpreted as evidence for the presence or absence of malignant disease.

For information on body fluid reference ranges and/or interpretive guidance visit http://aruplab.com/bodyfluids/

This test was developed and its performance characteristics determined by ARUP Laboratories. It has not been cleared or approved by the US Food and Drug Administration. This test was performed in a CLIA certified laboratory and is intended for clinical purposes.

Performed By: ARUP Laboratories
500 Chipeta Way
Salt Lake City, UT 84108
Laboratory Director: Jonathan R. Genzen, MD, PhD
CLIA Number: 46D0523979

Source, Fluid Peritoneal ARRL

Performing Site:
ARRL: ARUP REFERENCE LAB 500 Chipeta Way Salt Lake City UT 841081221

Reported Date: 2023.08.30 11:35 CEAFI

LAB: L - LOW, H - HIGH, AB - ABNORMAL, C - CRITICAL, . - NOT TESTED