

TEST CHANGE

Urea Nitrogen, Fluid 0020183, FL UN	
Specimen Requirements:	
Patient Preparation:	
Collect:	Peritoneal/ <u>Ascites</u> fluid.
Specimen Preparation:	Centrifuge and separate to remove cellular material. Trans <u>feport</u> 1 mL body fluid <u>to an ARUP Standard Transport</u> <u>Tube</u> (Min: 0.2 mL)
Transport Temperature:	Refrigerated.
Unacceptable Conditions:	<u>SpecimenDialysate and any specimen</u> types other than those listed. <u>Specimens too viscous to be aspirated by instrument.</u>
Remarks:	Indicate source on test request form.
Stability:	Ambient: <u>1 week</u> 24 hours; Refrigerated: <u>1 week5 days</u> ; Frozen: 1 year
Methodology:	Quantitative Spectrophotometry
Performed:	Sun-Sat
Reported:	Within 24 hours
Note:	
CPT Codes:	84520
New York DOH Approval Status:	This test is New York DOH approved.
Interpretive Data:	

interpretive Data.

For information on body fluid reference ranges and/or interpretive guidance visit https://www.aruplab.com/bodyfluids_Reference ranges for this assay have not been established for body fluid. Results should be interpreted in comparison to the concentration in blood and in conjunction with clinical context.

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.

Reference Interval:

Reference Intervals have not beenNot established.



HOTLINE NOTE: There is a component change associated with this test. One or more components have been added or removed. Refer to the Hotline Test Mix for interface build information.