
Overview**Useful For**

Evaluation of peritoneal fluid or abdominal drain fluid as a screening test for bile leakage

May aid in the distinction between a transudative and an exudative pleural effusion

Method Name

Photometric, Diazonium Salt

NY State Available

Yes

Specimen**Specimen Type**

Body Fluid

Ordering Guidance

For bilirubin testing on amniotic fluid specimens, order AFBIL / Bilirubin, Amniotic Fluid. Testing will be changed to AFBIL if this test is ordered on amniotic fluid specimens.

For bilirubin testing on urine specimens, order UBILU / Bilirubin, Random, Urine. Testing will be changed to UBILU if this test is ordered on urine specimens.

Shipping Instructions

Ship specimen in amber vial to protect from light.

Necessary Information

1. Date and time of collection are required.

2. Specimen source is required.

Specimen Required

Supplies: Amber Frosted Tube, 5 mL (T192)

Preferred Source:

-Peritoneal fluid (peritoneal, abdominal, ascites, paracentesis)

-Pleural fluid (pleural, chest, thoracentesis)

-Drain fluid (drainage, JP drain)

-Pericardial fluid

Acceptable Source: Write in source name with source location (if appropriate)

Collection Container/Tube: Sterile container

Submission Container/Tube: Opaque, amber vial

Specimen Volume: 1 mL

Collection Instructions:

1. Centrifuge to remove any cellular material and transfer into an amber vial to protect from light.
2. Indicate the specimen source and source location on label.

Specimen Minimum Volume

0.5 mL

Reject Due To

Gross hemolysis	Reject
Gross lipemia	Reject
Gross icterus	OK
Anticoagulant or additive, amniotic fluid, breast milk, saliva, sputum, cerebrospinal fluid, bronchoalveolar lavage (BAL) or bronchial washings, colostomy, ostomy, gastric secretions, nasal secretions, urine, feces, vitreous fluid, or synovial fluid	Reject

Specimen Stability Information

Specimen Type	Temperature	Time	Special Container
Body Fluid	Frozen (preferred)	70 days	LIGHT PROTECTED
	Refrigerated	14 days	LIGHT PROTECTED

Clinical and Interpretive

Clinical Information

Peritoneal fluid:

Bilirubin is typically measured in peritoneal fluid of patients with suspected bile duct leak or gallbladder perforation as a screening test prior to imaging or cholescintigraphy. If the value is higher than that of serum and is greater than 6 mg/dL, and the ascitic fluid amylase is not elevated (indicating upper intestinal perforation), it can be assumed that the gallbladder has perforated into the peritoneum (choleperitoneum) and/or bowel or biliary perforation has occurred.(1) Furthermore, biliary leakage after laparoscopic cholecystectomy is the most common post-operative complication.(2) While endoscopy is a beneficial first-line treatment for the management of bile leaks there often are logistical issues which hinder the procedure from being performed rapidly. Post-cholecystectomy patients generally have a drain in place (particularly a Jackson Pratt or JP drain) and may undergo bilirubin testing on the drain fluid as an objective assessment of a bile leak. A body fluid/serum bilirubin ratio of greater than 5 in a JP drain fluid is highly sensitive and specific for bile leak.(3)

Pleural fluid:

Measurement of bilirubin in pleural fluid has been investigated to aid in the differentiation of transudative and exudative effusions in pursuit of more specific biomarkers than traditional light criteria measuring total protein and lactate dehydrogenase. Bilirubin values tend to be higher in exudates than in transudates, although there is some overlap between groups which limits the usefulness of its measure.(4)

Other fluids:

Determination of body fluid bilirubin concentration can aid in the distinction between a transudative and an exudative fluid or identify the presence of bile in other fluid compartments.

Reference Values

An interpretive report will be provided.

Interpretation

Bilirubin may be measured in other fluids although the decision limits are not well defined in fluids other than pleural fluid. Fluid to serum bilirubin ratios are expected to be less than or equal to 1.0 and should be interpreted in conjunction with other clinical findings.

Cautions

Bilirubin is photosensitive. Failure to protect from light may cause decreased results.

In very rare cases, gammopathy, in particular type IgM (Waldenstrom macroglobulinemia), may cause unreliable results.

Cyanokit (Hydroxocobalamin) may cause false low results.

Clinical Reference

1. Runyon BA: Ascitic fluid bilirubin concentration as a key to choleperitoneum. *J Clin Gastroenterol.* 1987 Oct;9(5):543-545
2. Koch M, Garden OJ, Padbury R, et al: Bile leakage after hepatobiliary and pancreatic surgery: a definition and grading of severity by the International Study Group of Liver Surgery. *Surgery* 2011. May;149(5):680-688. doi: 10.1016/j.surg.2010.12.002
3. Darwin P, Goldberg E, Uradomo L: Jackson Pratt drain fluid-to-serum bilirubin concentration ratio for the diagnosis of bile leaks. *Gastrointest Endosc.* 2010 Jan;71(1):99-104. doi: 10.1016/j.gie.2009.08.015
4. Metintas M, Alatas O, Alatas F, Colak O, Ozdemir N, Erginel S: Comparative analysis of biochemical parameters for differentiation of pleural exudates from transudates Light's criteria, cholesterol, bilirubin, albumin gradient, alkaline phosphatase, creatine kinase, and uric acid. *Clin Chim Acta.* 1997 Aug 29;264(2):149-162. doi: 10.1016/s0009-8981(97)00091-0

Performance

Method Description

Total bilirubin, in the presence of a suitable solubilizing agent, is coupled with 3,5-dichlorophenyl diazonium in a strongly acidic medium to produce azobilirubin. The intensity of the color of the azobilirubin produced is proportional to the total bilirubin concentration and is measured at 546/600 nm.(Package insert: Bilirubin Total Gen. 3. Roche Diagnostics; V9.0. 01/2020)

PDF Report

No

Day(s) Performed

Monday through Sunday

Report Available

Same day/1 to 2 days

Specimen Retention Time

1 week

Performing Laboratory Location

Rochester

Fees and Codes**Fees**

- Authorized users can sign in to [Test Prices](#) for detailed fee information.
- Clients without access to Test Prices can contact [Customer Service](#) 24 hours a day, seven days a week.
- Prospective clients should contact their Regional Manager. For assistance, contact [Customer Service](#).

Test Classification

This test has been modified from the manufacturer's instructions. Its performance characteristics were determined by Mayo Clinic in a manner consistent with CLIA requirements. This test has not been cleared or approved by the US Food and Drug Administration.

CPT Code Information

82247

LOINC® Information

Test ID	Test Order Name	Order LOINC Value
BFBL	Bilirubin, BF	1974-5

Result ID	Test Result Name	Result LOINC Value
BRNBF	Bilirubin (BF)	1974-5
FLD14	Fluid Type:	14725-6