



# Bilirrubina Total

AA

DPD method for total bilirubin determination in serum or plasma

## SUMMARY

Bilirubin, a compound produced by the degradation of hemoglobin, is captured by the liver for its conjugation and bile excretion. Hepatocellular disorders or biliary obstructions may lead to hyperbilirubinemia. Erythroblastosis fetalis or hemolytic anemia of the newborn, is a pathology produced by mother-fetus blood incompatibility, in which an excessive destruction of red blood cells occurs. This results in a severe increase of serum bilirubin, with the consequent risk of pigment diffusion to the central nervous system. Therefore, bilirubin determination in these newborns is extremely important.

## PRINCIPLE

Indirect bilirubin, bound to albumin, is released by a surfactant. Total bilirubin (free and conjugated) reacts with dichlorophenyldiazonium salt (DPD) forming a red azocompound in an acid solution.

## PROVIDED REAGENTS

- A. Reagent A:** vials containing dichlorophenyldiazonium salt.  
**B. Reagent B:** aqueous solution containing 90 mmol/l hydrochloric acid and surfactant.  
**C. Reagent C** (Reagent for Sample Blank): aqueous solution containing 90 mmol/l hydrochloric acid and surfactant.

### Final concentrations

hydrochloric acid..... 90 mmol/l  
surfactant..... 4 %  
dichlorophenyldiazonium ..... 1 mmol/l

## NON-PROVIDED REAGENTS

Wiener lab's **Calibrador A plus** or **Bilirrubina Standard**.

## INSTRUCTIONS FOR USE

**Reagent B and C:** ready to use.

**Working Reagent:** reconstitute each vial of Reagent A with 10 ml of Reagent B. Cap and shake until complete dissolution. Homogenize and date.

## WARNINGS

Reagents are for "in vitro" diagnostic use.

Reagent B and Reagent C are corrosive. H315+H320: Causes skin and eye irritation. H314 Causes severe skin burns and eye damage. P262 Do not get in eyes, on skin, or on clothing. P305 + P351 + P338 IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing. P302 + P352 IF ON SKIN: Wash with plenty of soap and water. P280 Wear protective gloves/protective clothing/eye protection/face protection.

Use the reagents according to the working procedures for clinical laboratories.

The reagents and samples should be discarded according to the local regulations in force.

## INSTABILITY OR DETERIORATION OF REAGENTS

Absorbance readings of the Blank Reagent, higher than 0.100 O.D. (at 546 nm) indicate its deterioration. Discard in such case.

Turbidity indicates reagents deterioration. Discard in such case.

## STABILITY AND STORAGE INSTRUCTIONS

**Provided Reagents:** stable in refrigerator (2-10°C) until the expiration date shown on the box.

**Working Reagent:** is stable for 21 days in refrigerator (2-10°C).

## SAMPLE

Serum or plasma

**a) Collection:** obtain as usual. Protect from natural or artificial light covering the tube with black paper.

**b) Additives:** if plasma is used, use heparin for collection.

**c) Known interfering substances:** no interference have been observed by hemoglobin up to 350 mg/dl, triglycerides up to 1000 mg/dl (10 g/l) or heparin up to 50 U/ml.

See Young, D.S. in References for effect of drugs on the present method.

**d) Stability and storage instructions:** sample should be preferably fresh. If assay is not performed immediately, serum can be stored up to 48 hours in refrigerator (2-10°C) and whole blood no more than 24 hours in refrigerator (2-10°C) or 12 hours at room temperature (< 25°C).

The action of light is capable of destroying up to a 50% of the bilirubin present in the sample. Consequently, it should be carefully protected from light.

## REQUIRED MATERIAL (non-provided)

- Spectrophotometer or photocolormeter
- Micropipettes and pipettes for measuring the stated volumes
- Tubes or spectrophotometric cuvettes
- Stopwatch

## ASSAY CONDITIONS

- Wavelength: 546 nm in spectrophotometer or 520-550 nm in photocolormeter with green filter
- Reaction temperature: room temperature (< 25°C)
- Reaction time: 10 minutes
- Sample volume: 40 ul
- Final reaction volume: 0.54 ml

## PROCEDURE

In 4 photocolormeter tubes labeled B<sub>C</sub> (Calibrator or Standard Blank), C (Calibrator or Standard), B<sub>U</sub> (Unknown Blank) and U (Unknown), place:

	B <sub>C</sub>	C	B <sub>U</sub>	U
<b>Reagent C</b>	0.5 ml	-	0.5 ml	-
<b>Reagent B</b>	-	0.4 ml	-	0.4 ml
<b>Calibrator</b>	40 ul	40 ul	-	-
<b>Sample</b>	-	-	40 ul	40 ul
<b>Working Reagent</b>	-	0.1 ml	-	0.1 ml

Mix and incubate for 10 minutes at room temperature (< 25°C). Read in spectrophotometer at 546 nm or in photocolormeter with green filter (520-550 nm), setting the instrument to zero with the Reagent Blank (B<sub>R</sub>). Volumes may be proportionally increased.

Note: process a Reagent Blank (B<sub>R</sub>) for each series of determinations, mixing 0.4 ml Reagent B + 40 ul distilled water + 0.1 ml Working Reagent.

## STABILITY OF FINAL REACTION

Reaction color is stable for 30 minutes. Therefore, absorbance should be read within that period of time.

## CALCULATIONS

Total Bilirubin (mg/l) = (U - B<sub>U</sub>) x f

$$f = \frac{X^* \text{ mg/l}}{C - B_C}$$

\*total bilirubin concentration in Wiener lab.'s **Calibrator A plus**. If Wiener lab.'s Bilirubina Standard is used, X = 100 mg/l.

Example:

B<sub>C</sub> = 0.002

C = 0.258

B<sub>D</sub> = 0.002

D = 0.092

X = 48 mg/l

$$f = \frac{48 \text{ mg/l}}{0,258 - 0,002} = 188 \text{ mg/l}$$

Total bilirubin (mg/l) = (0,092 - 0,002) x 188 = 17 mg/l

## QUALITY CONTROL METHOD

Each time the test is performed, analyze two levels of a quality control material (**Standatrol S-E 2 niveles**) with known direct bilirubin concentration.

## REFERENCE VALUES

**Total bilirubin in serum or plasma**

Adults: up to 10 mg/l

Newborns:

	Term infant	Preterm infant
umbilical cord blood	< 20 mg/l	< 20 mg/l
up to 24 hrs	14 - 87 mg/l	< 80 mg/l
up to 48 hrs	34 - 115 mg/l	< 120 mg/l
from 3 <sup>rd</sup> to 5 <sup>th</sup> day	15 - 120 mg/l	< 160 mg/l

Values start to decrease to reach the adult's average level within a month after birth.

In premature babies, levels take longer to reach normality, depending on the hepatic immaturity degree.

It is recommended that each laboratory establishes its own reference values.

## UNITS CONVERSION

Bilirubin (mg/l) = Bilirubin (mg/dl) x 10

Bilirubin (mg/dl) x 17.1 = Bilirubin (umol/l)

## PROCEDURE LIMITATIONS

See Known interfering substances under SAMPLE.

The action of light, both on samples and on standard solutions, is capable of destroying up to 50% of bilirubin in one hour.

## PERFORMANCE

**a) Reproducibility:** processing replicates of the same samples on the same day, the following values were obtained:

Level	S.D.	C.V.
6.0 mg/l	± 0.09 mg/l	1.50 %
56.3 mg/l	± 0.55 mg/l	0.98 %

Processing the same sample on different days, were obtained:

Level	S.D.	C.V.
5.5 mg/l	± 0.15 mg/l	2.68 %
56.1 mg/l	± 0.60 mg/l	1.08 %

**b) Linearity:** the reaction is linear up to 200 mg/l bilirubin. For higher values, dilute the sample 1:2 or 1:4 with saline solution, repeat the determination and multiply the obtained result by 2 or 4 accordingly.

**c) Recovery:** adding known amounts of bilirubin to different sera, a recovery between 96 and 107% was obtained.

**d) Analytical sensitivity:** 0.8 mg/l.

## PARAMETERS FOR AUTOANALYZERS

For programming instructions check the user's manual of the autoanalyzer in use.

For calibration, it must be used Wiener lab.'s **Calibrator A plus**. The kit provides the Reagent for the Sample Blank (Reagent C) which is only required in some autoanalyzers. In other analyzers this Reagent is not necessary. For further information consult your analyzer's applications manual for Wiener lab. reagents.

## WIENER LAB. PROVIDES

- 4 vials Reagent A
- 4 x 10 ml Reagent B
- 4 x 40 ml Reagent C
- Cat. 1009305


- 4 vials Reagent A
- 4 x 50 ml Reagent B
- 2 x 100 ml Reagent C
- Cat. 1120005

## REFERENCES

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## SYMBOLS

The following symbols are used in the packaging for Wiener lab. diagnostic reagents kits.

 This product fulfills the requirements of the European Directive 98/79 EC for "in vitro" diagnostic medical devices

 Authorized representative in the European Community

 "In vitro" diagnostic medical device


 Contains sufficient for <n> tests

 Use by

 Temperature limitation (store at)

 Do not freeze

 Biological risks

 Volume after reconstitution

 Contents

 Batch code

 Manufactured by:

 Harmful

 Corrosive / Caustic

 Irritant

 Consult instructions for use


 Calibrator

 Control

 Positive Control

 Negative Control

 Catalog number

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