# Protein & Enzyme Lab (BBT 314)

# **Experiment 1**

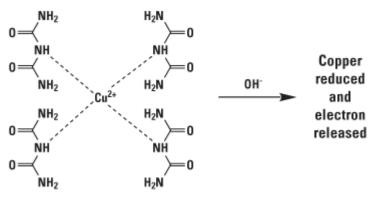
Estimation of total protein in unknown sample by biuret method using standard curve

#### **Background:**

Plasma proteins are synthesized predominantly in the liver plasma cells, lymph nodes, spleen and in bone marrow. In the course of disease the total protein concentration and also the percentage represented by individual fractions can significantly deviate from the normal values. Hypoproteinemia can be caused by diseases and disorders such as loss of blood, nephritic syndrome, severe burns, salt retention syndrome and kwashiorkor (acute protein deficiency). Hyperproteinemia can be observed in case of severe dehydration and illness such as multiple myeloma changes in the relative percentages of one plasma protein fraction. Often in such cases the amount of total protein does not change. The A/G is commonly used as in index of the distribution of albumin and globulin fractions. Marked changes in this ratio can be observed in cirrhosis of the liver, glomerulonephritis, nephrotic syndrome, acute hepatitis and lupus erythrometosus as well as in certain acute and chronic inflammations. The total protein measurements are used in the diagnosis and treatment of a variety of diseases involving the liver, kidney or bone marrow as well as other metabolic or nutritional disorders.

## Principle:

Biuret methods are based on the formation of coloured complexes between peptide bonds and cupric ions, where divalent copper reacts with the peptide bonds of protein under alkaline conditions to form an intensive violet-blue complex. Iodine is included as an antioxidant.



**Biuret Reaction** 

## Reagents:

Protein	Bovine serum albumin (BSA)	5 g/dl
standard		
R Biuret	Sodium iodide	100 mmol/1
	Potassium iodide	5 mmol/1
	Copper (II) sulphate	19 mmol/1

#### **Procedure:**

**Specimen:** Bovine Serum Albumin (BSA)

**Standard preparation**: Using the given BSA solution, prepare 5 standards of different concentrations (200, 400, 600, 800 and 1000 mg/dl) using dilution from a stock solution of 5 g/dl.

Order	Desired Concentration (mg/dl)	Volume of BSA solution (µl)	dH <sub>2</sub> O (μl)	Total volume (ml)
5.	1000			1.0
4.	800			1.0
3.	600			1.0
2.	400			1.0
1.	200			1.0
Blank	0			1.0

**Standards:** BSA solution in different concentration 200mg/dl, 400mg/dl, 600mg/dl, 800mg/dl & 1000mg/dl.

## Assay preparation:

Take 7 test tubes and level them as B (Blank), Std1, Std2, Std3, Std4, Std5, and Unknown Solution.

Prepare the test tubes as shown in the table below:

Tube	В	1	2	3	4	5	SP
Reagents(R)	1ml	1ml	1ml	1ml	1ml	1ml	1ml
Standards	50μ1	50 μl of	50μ1 of	50μ1 of	50μ1 of	50μ1 of	50μ1 of
	of	200mg/dl	400mg/dl	600mg/dl	800mg/dl	1000mg/dl	unknown
	dH <sub>2</sub> O						Protein
							solution
Absorbance							

Mix well and incubate for 5 minutes at 37°C or 10 minutes at room temperature.

Measure the absorbance of the blank and standards at 545 nm.

Colour is stable for at least 30 minutes.

**Graph preparation:** Plot an abs vs. concentration graph using the values,

Y-axis concentration of standards

X-axis absorbance of the standards

# **Experiment 2**

Estimation of total protein in serum by enzymatic method using semi-automated Biochemistry Analyzer

#### Reagents:

Protein standard	Bovine albumin	5 g/d1
R Biuret	Sodium iodide	100 mmol/1
	Potassium iodide	5 mmol/1
	Copper (II)	19 mmol/1
	sulphate	

#### Procedure:

Sample: Serum

**Sample collection**: Sample must be collected from any healthy individual. Specimen may be stored at 2-8°C prior to analysis.

**Sample preparation:** Blood cells are removed by centrifugation. After that the top clear layer of serum is collected by pipetting carefully.

Prepare the tube as shown below:

	Blank	Standard	Sample
R	1.0 ml	1.0 ml	1.0 ml
Standard	-	10 μ1	-
Sample	-	-	10 μ1
dH₂O	10 μ1	-	-
Absorbance			

Mix well and incubate for 5 minutes at 37°C or 10 minutes at room temperature.

Measure the absorbance of sample at 545 nm. Colour is stable for at least 30 minutes.

#### **Calculation:**

Calculate the total Protein concentration by using the following formula:

**Total Protein concentration** = (Absorbance of sample/Absorbance of standard) x [Standard]

(Unit conversion:  $mg/dl \times 1.45 = \underline{\hspace{1cm}} mmol/L$ )

Expected value:

Adults 6.3-8.3 g/dL

Children

> 1 year 6.0-8.0 g/dL

< 1 year 4.6-7.6 g/dL