



**M.SC**  
**Semester**  
**III Core**  
**Course XI**  
**Bio-Inorganic**  
**Chemistry**

**TOPIC:- Copper Metabolism**

**Department of Chemistry**  
**L.S COLLEGE MUZAFFARPUR**  
**B. R. A. BIHAR UNIVERSITY**  
**Dr. Priyanka**

# Characteristics

- **Copper is deposited in abnormal amounts in liver and lenticular nucleus of brain.**
- **This may lead to hepatic cirrhosis & brain necrosis.**
- **Low levels of copper and ceruloplasmin in plasma with increased excretion of copper in urine.**

- **Copper deposition in kidney causes renal damage.**
- **This leads to increased excretion of amino acids, glucose, peptides & hemoglobin in urine.**
- **Intestinal absorption of copper is very high, about 4-6 times higher than normal.**

# Probable causes of Wilson's disease

- **A failure to synthesize ceruloplasmin or an impairment in the binding capacity of copper to this protein or both.**
- **Copper is free in the plasma, it easily enters the tissues (liver, brain, kidney), binds with the proteins & gets deposited.**
- **Albumin bound copper is either normal or increased**

- **Copper accumulates particularly in liver, brain, kidney and eyes leading to copper toxicosis.**
- **Causes neurological symptoms, liver damage leading to cirrhosis, renal tubular damage and Kayser-Fleisher rings (brown pigment around the iris) at the edges of the cornea due to deposition of copper in the cornea.**

- **Treatment:**
- **Includes diet low in copper and administration of copper chelator, D-penicillamine.**