Water and minerals metabolism 304 Biochen

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Copper (Cu)

- Menke's syndrome:
 - An extreme form of copper deficiency an x-linked genetic defect in copper transport and storage. It is observed early in life (usually 3 months after birth) and death occurs within 5 years.

 The affected infants exhibit an <u>accumulation of copper in the</u> <u>duodenal mucosal tissues</u> as the result of defective copper absorption, with <u>low serum copper</u>, <u>low serum ceruloplasmin</u>, <u>low hepatic and cerebral copper</u> and <u>little or no cytochrome C</u> <u>oxidase activity in nervous tissues</u>.



• Wilson's disease (hepatolenticular degeneration):

 It is an autosomal recessive copper accumulation disease that usually presents between 6-40 years of age. It is suggested that decreased copper excretion via bile may be a factor in the pathogenesis of the disease.

- The disease is characterized by:

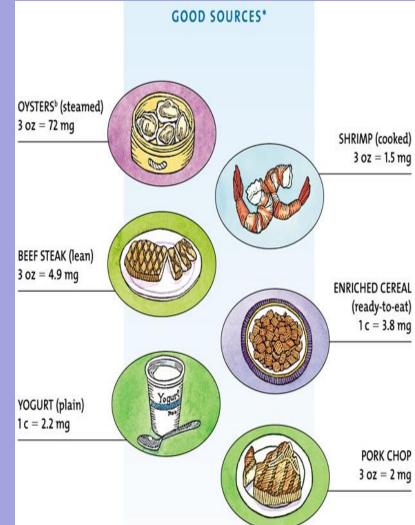
- increased concentrations of in the liver, brain, kidney, and cornea.
- Low total serum copper, and low serum ceruloplasmin.
- The disease symptoms include: neurologic disorders, liver cirrhosis, Kayser-Fleischer ring in the cornea (a green or golden pigment ring around the cornea caused by copper deposits)
- The disease is treated by penicillamine which increases the excretion of copper in urine.

Zinc (Zn)

- Adult male body weighing 60-70 kg contain about 2.0 - 2.3 g of zinc
- Distribution of zinc in the body:
 About 20% is present in the skin
 High concentrations are present in semen spermatozoa, prostate and epididymis (i.e. in male sex organs)
 A considerable proportion is present in the bones and teeth
- Food sources: animal protein containing foods and milk together with whole grain products are a good source of zinc

• RDA

- adults: 15 mg/day
- Pregnant and lactating women:
 25-30 mg/day



Zinc (Zn)

Functions:

- Essential component of a number of enzymes including:
 - Alcohol dehydrogenase
 - Alkaline phosphatase (ALP),
 - Carbonic anhydrase,
 - Carboxy peptidase,
 - Cytosolic SOD
 - Retinene reductase (required for formation of retinal (vit. A) during the rhodopsin cycle
- Essential for normal growth, reproduction, taste sensitivity, tissue repair, wound healing.
- Essential to maintain normal levels of vit. A in plasma and mobilization of vit A from the liver
- Insulin is sotred as zinc-insulin complex in beta cells of pancreas. Zinc adhering to insulin molecule increases the duration of insulin action when given by injection

Zinc (Zn)

• Excretion:

- it is mainly excreted in feces (unabsorbed fraction),
- smaller amounts are secreted in pancreatic juice, bile, and sweat.

• Deficiency :

Causes:

- 1. Marked reduced intake of zinc in diet
- 2. Malabsorption syndrome
- 3. Liver diseases as post-alcoholic cirrhosis

Symptoms:

- 1. Impaired taste sensitivity
- 2. Poor appetite
- 3. Growth retardation (dwarfism)
- 4. Delayed wound healing
- 5. Impaired sexual development (hypogonadism)

TOXICITY:

Loss of appetite, impaired immunity, reduced copper and iron absorption, low HDL cholesterol (a risk factor for heart disease)

Manganese (Mn)

 Adult male body weighing 60-70 kg contain about 0.012 – 0.02 g of manganese

Distribution of manganese in the body:
 Mitochondria are the main intracellular sites of manganese uptake.
 Kidney & liver are the main storage organs for manganese

- Manganese in blood: its blood level is 4-20 microgram/dl, it is bound to beta-globulin
- Food sources: found in nuts, whole grains and leafy vegetables and fruits
- RDA
 - adults: 2.5-7 mg/day
 - Pregnant and lactating women: 25-30 mg/day
- Absorption:
- It is absorbed through the small intestine by a mechanism similar to that for iron

Manganese (Mn)

- Excretion
- It is excreted mainly via bile &feces, and very little in urine
- Functions:
 - Essential for normal bone structure, reproduction, and the normal functions of central nervous system
 - Manganese is a cofactor of several different enzymes:
 - Phosphoglucomutase
 - Isocitric dehydrogenase
 - Cholinesterase
 - Intestinal peptidase
 - Carboxylases
 - ATPases
 - Mitochondrial SOD
- Deficiency :
 - rare

lodine (I₂)

Adult male body weighing 60-70 kg contain about 0.02 – 0.05 g of iodine

• Distribution of iodine in the body:

About 33% is in the thyroid gland. About 66% is distributed in all tissues particularly ovary, muscles and blood.

- Iodine in blood: most of iodine in blood is bound to protein at levels of 4 – 8 microgram/dl.
- Food sources: world's seas fish, seaweeds and iodized salt are best sources
- RDA
 - adults: 0.1-0.2 mg/day
 - Pregnant and lactating women: 25-30 mg/day