Chemical Examination of Urine

Renal Block 1 Lecture

Objectives

Upon completion of this lecture, students should be able to:

Differentiate between normal and abnormal constituents of urine including: Proteins, sugars, ketone bodies, nitrite, bile pigments, blood etc.
Know the clinical conditions in different types of proteinuria, blood urea and glycosuria etc.

Overview

- Introduction
- Normal composition of urine
- Abnormal composition of urine
- Proteinuria:
 - Pre-renal (multiple myeloma)
 - Renal
 - Post-renal
- Glycosuria: fructosuria, galactosuria
- Ketonuria
- Hematuria: hemoglobinuria

Urine

 Urine is a fluid excreted by most animals including humans

It is formed in the kidneys (renal glomeruli)

 The fluid undergoes chemical changes before it is excreted as urine

 Normal urine excretion by a healthy person is about 1.5 L per day

Normal composition of urine

Urine contains organic and inorganic constituents:

- Major inorganic constituents:
 - Sodium
 - Potassium
 - Chloride
 - Small amounts of Ca, Mg, sulfur and phosphates
 - Traces of Fe, Cu, Zn, I
 - Major organic constituents:
 - Non-protein nitrogen (NPN) compounds
 - Organic acids
 - Sugars

Traces of proteins, vitamins, hormones, pigments

Abnormal composition of urine

Proteins:

- Normal urine contains small amount of protein:
 < 200 mg/day
- Excretion of more than this level causes proteinuria

Proteinuria:

- Excretion of abnormal amounts of protein in urine
- Proteinuria has three types:
 - Pre-renal
 - Renal
 - Post-renal

Pre-renal proteinuria

- Some diseases or conditions increase plasma protein levels not involving the kidneys
- Causes increased filtration of these proteins in the kidneys
- This exceeds the normal reabsorptive capacity of renal tubules
- Results in overflow of proteins in the urine

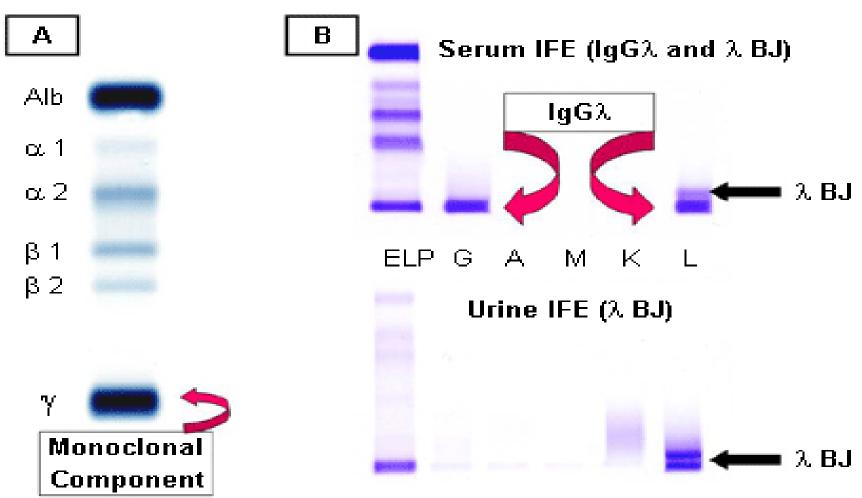
Multiple myeloma:

- Cancer of the antibody-producing plasma cells
- Causes pre-renal proteinuria

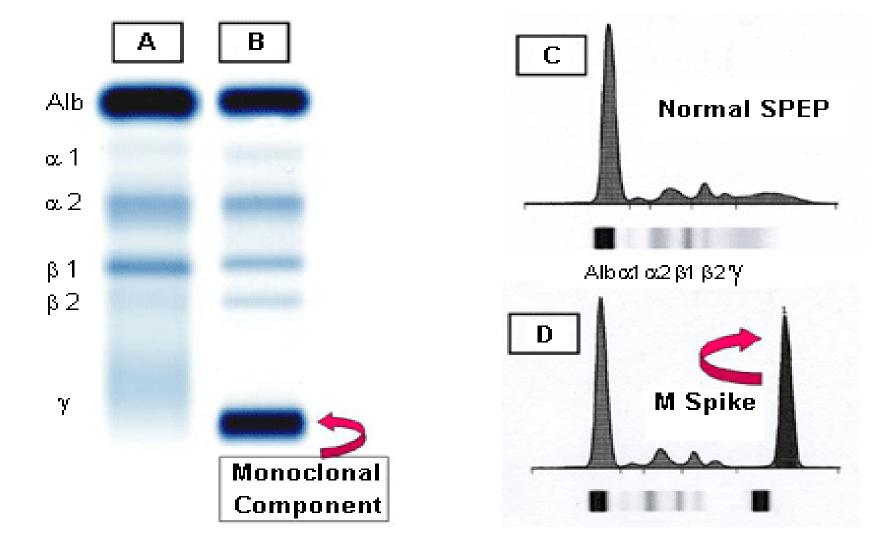
Pre-renal proteinuria

- The serum contains elevated levels of light-chain monoclonal antibodies called Bence-Jones protein
 This protein is filtered in the kidneys in high amounts
 Exceeding the tubular reabsorption capacity
 Hence excreted in the urine
 Bence-Jones protein coagulates at 40–60 °C and dissolves at 100 °C
- Multiple myeloma cases are diagnosed by:
 - Serum electrophoresis
 - Immunoelectrophoresis

Multiple myeloma



A: Serum protein electrophoresis showing the M component B: Serum and urine immunofixation electrophoresis



A: Normal serum B: Multiple myeloma (M component in γ region) C: Densitometry of "A" D: Densitometry of "B" (M component is called M Spike)

Renal proteinuria

Associated with renal disease
 Glomerular proteinuria:

- High glomerular permeability causes filtration of high molecular weight proteins
 - Example: glomerulonephritis

Tubular proteinuria:

- Low tubular reabsorption with normal glomerular permeability
- Causes excretion of low mol. wt. proteins
 Example: chronic nephritis

Renal proteinuria

Orthostatic (postural) proteinuria:A form of benign or physiological proteinuria

 Occurs frequently in young adults due to periods spent in a vertical posture (body position) or during muscular exercise

• Increased pressure on the renal vein in the vertical position causes orthostatic proteinuria

• Disappears in horizontal posture

Renal proteinuria

Microalbuminuria:

Presence of small amounts of albumin in the urine
20-200 mg/L

• Cannot be detected by ordinary urine testing

Needs special tests for detection

Early indicator of glomerular dysfunction due to:
 Uncontrolled diabetes mellitus

Hypertension

Post-renal proteinuria

Proteins are added to the urine after kidney filtration

• While passing through the lower urinary tract (ureters, bladder, urethra, prostate, vagina)

• Due to:

• Lower urinary tract infection

- Trauma
- Tumors
- Stones

Glycosuria

Presence of sugar in urine Glucosuria: Presence of detectable amount of glucose in urine

Due to diabetes mellitus
Plasma glucose level exceeds the renal threshold

Due to renal disease (renal glucosuria)

- Normal plasma glucose level with proximal tubular malfunction
- Decreased renal threshold as observed in gestational diabetes and Fanconi's syndrome

Glycosuria

Fructosuria:

• Presence of fructose in urine

• Nutritional cause:

High fructose intake

• Metabolic cause:

• Low fructokinase or aldolase B in the liver

Galactosuría:

Presence of galactose in urine

- Nutritional cause: high galactose intake
- Metabolic cause:

 Low galactokinase or galactose -1-PO4 uridyl transferase in the liver

Ketonuria

 Presence of ketones, acetone, acetoacetic acid and βhydroxybutyric acid in urine

Due to:

- Diabetic ketoacidosis
- Starvation
- Dietary imbalance: high fat and low CHO dietPhenylketonuria (PKU)

Choluria

• Presence of bile, bilirubin and bile salts in urine Bilirubin: normally no bilirubin is detected in urine • It is detected in: Hepatocellular damage Obstruction of bile duct due to stones (extrahepatic) and hepatic tumors (intrahepatic) Urobilinogen: normally present in trace amounts • High urobilinogen is found in: Hemolytic anemia Hepatocellular damage Nitrites:

• Positive nitrite test indicates bacteria in urine

Hematuria

• Presence of detectable amount of blood in urine

Due to:

- Acute / chronic glomerulonephritis
- Local disorders of kidney and genito-urinary tract
 Trauma, cystitis, renal calculi, tumors
 Bleeding disorders
 - Hemophilia

HematuriaHemoglobinuria:Presence of hemolysed blood in urine

Due to:

- Hemoglobinopathies
 Sickle cell anemia
 Thalassemia
 Malaria
- Transfusion reactionBlood group incompatibility