

RENAL PHYSIOLOGY INTRODUCTION



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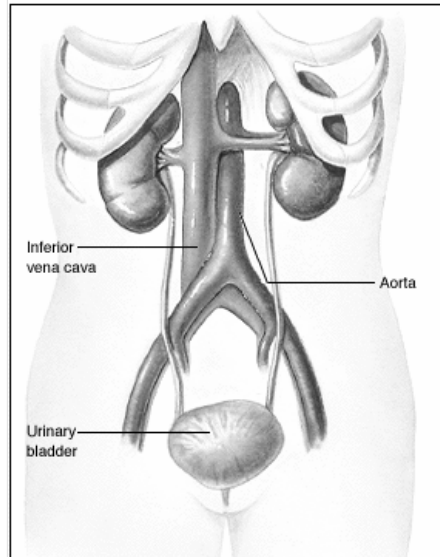
OBJECTIVES

At the end of this lecture you should be able to describe:

- **Physiologic anatomy of Urinary system**
- **Roles of the kidney**
- **Structure, Parts and Types of Nephrons**
- **Juxtaglomerular Apparatus**
- **Filtration Membrane**



URINARY SYSTEM

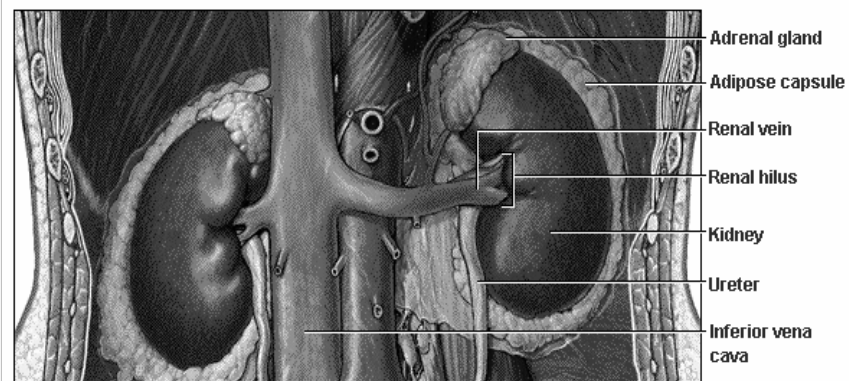


Kidneys: filter blood and remove wastes, producing urine

Ureters: carry urine to bladder from kidneys

Urethra: carries urine from bladder to the exterior

PHYSIOLOGIC ANATOMY OF KIDNEYS



ROLES OF THE KIDNEY

- 1. Regulation of water (extracellular fluid volume)**
- 2. Maintenance of electrolyte balance (Na^+ , K^+ , HCO_3^- , Ca^{++})**
- 3. Regulation of arterial pressure**
- 4. Regulation of blood pH**

ROLES OF THE KIDNEY (Cont.)

- 5. Excretion of wastes (creatinine, urea, benzoate, penicillin, saccharin)**
- 6. Secretion, metabolism, and excretion of hormones**
 - ⊕ Hormone production (Erythropoietin, Renin)**
 - ⊕ Activation of Vitamin D**
- 7. Gluconeogenesis**

Naturally occurring vitamin D (cholecalciferol)



25-hydroxycholecalciferol (25-OHD3)

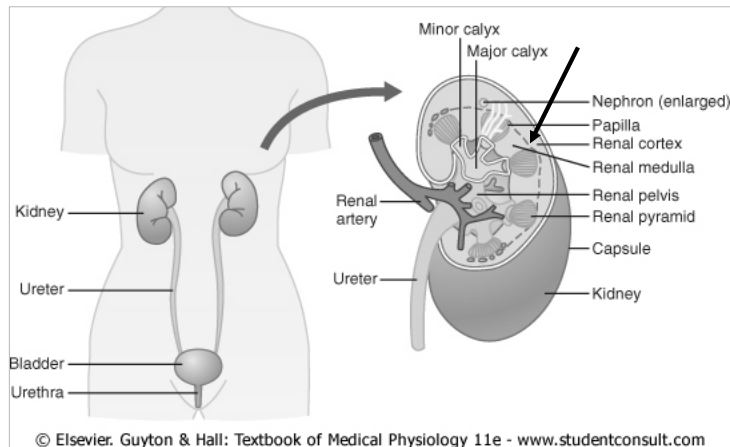
PTH →



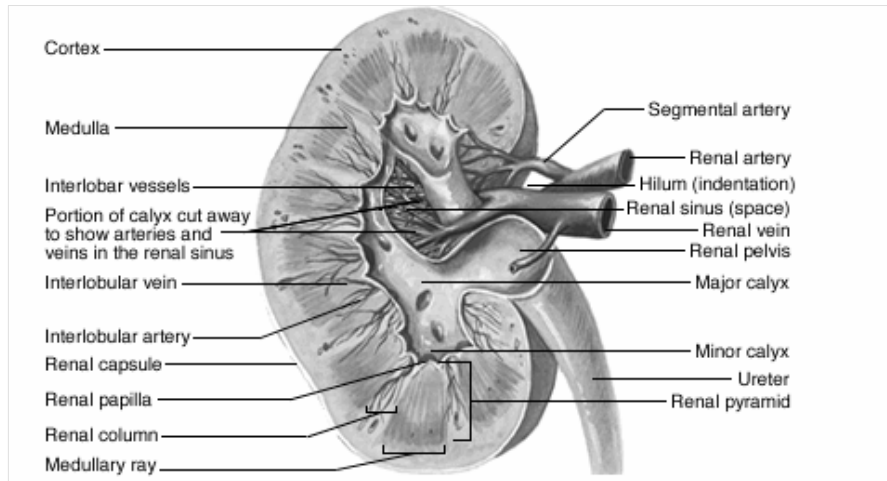
1,25-dihydroxycholecalciferol (1,25-(OH)2D3)

PHYSIOLOGIC ANATOMY OF KIDNEYS

- Size Clenched Fist
- Weight 150 grams

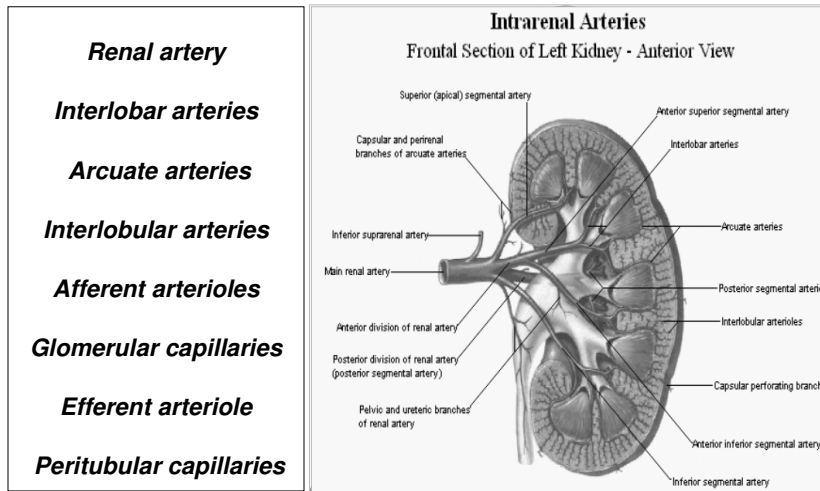


PHYSIOLOGIC ANATOMY OF KIDNEYS



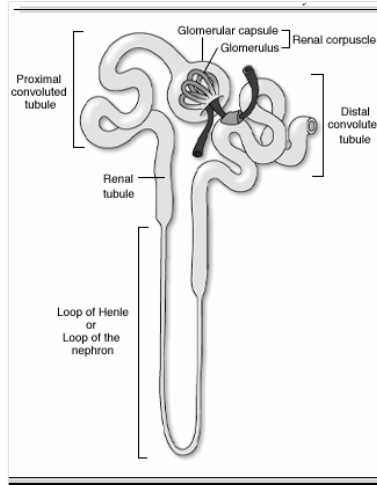
RENAL BLOOD SUPPLY

22 per cent of the cardiac output, or 1100 ml/min



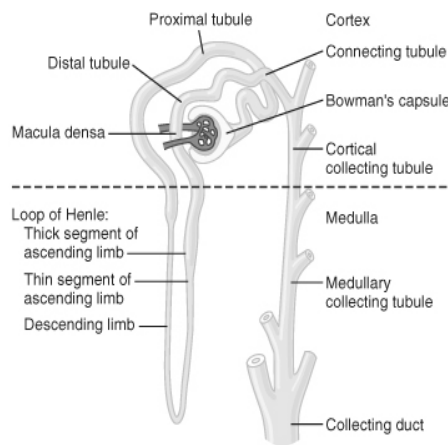
NEPHRON

- ❖ Each kidney in the human contains about 1.3 million nephrons, each capable of forming urine.
- ❖ The kidney cannot regenerate new nephrons.
- ❖ After age 40, the number of functioning nephrons usually decreases about 10 per cent every 10 years; thus, at age 80, many people have 40 per cent fewer functioning nephrons than they did at age 40.

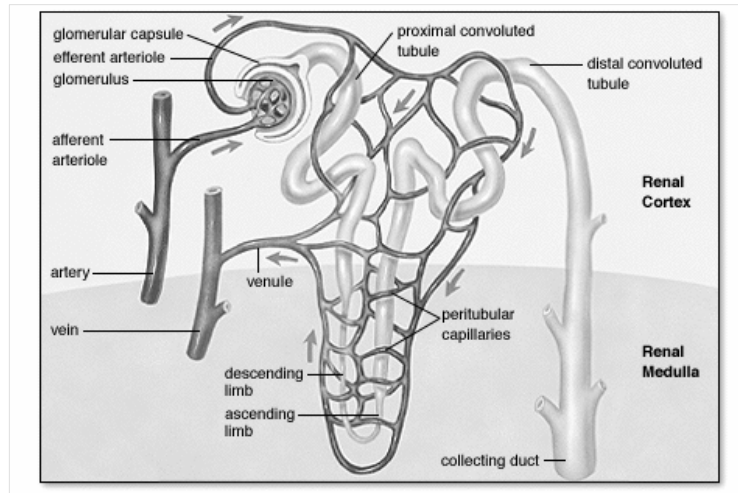


PARTS OF NEPHRON

- The initial parts of 8 to 10 cortical collecting ducts join to form a single larger collecting duct that runs downward into the medulla and becomes the *medullary collecting duct*.
- The *collecting ducts merge to form progressively larger ducts that eventually empty into the renal pelvis through the tips of the renal papillae.*



RENAL PORTAL SYSTEM

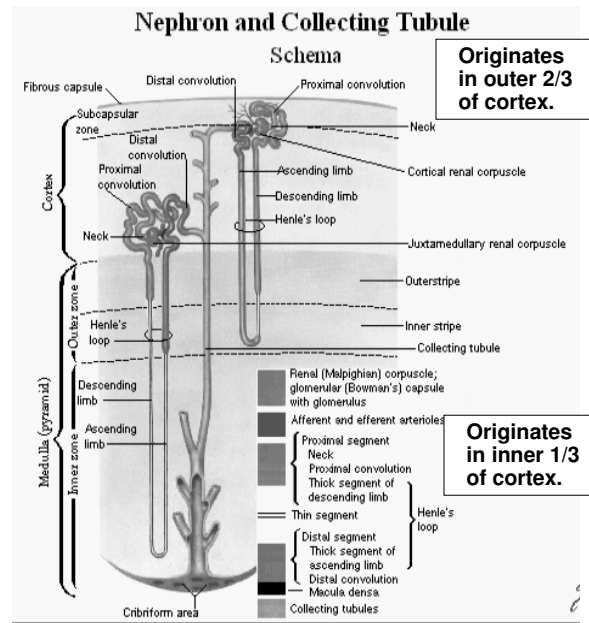


NEPHRON TYPES

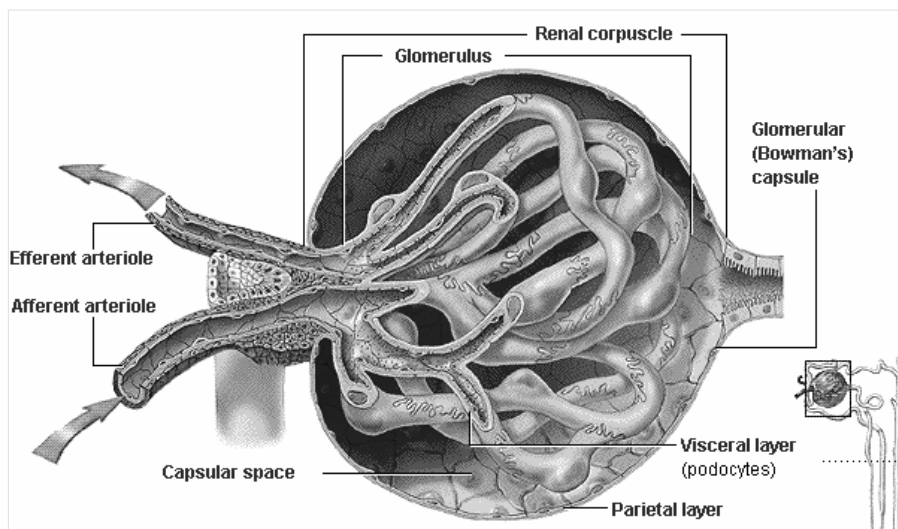
- ± Superficial (cortical) [85 %]
 - Capable of forming dilute urine
- ± Juxtamedullary [15 %]
 - Capable of forming concentrated (> 300 mOsm/kg) urine

NEPHRON TYPES
Cortical and
Juxtamedullary
Nephrons

**1-2 % Blood
Flows
Through
Juxta Medullary
Nephrons**



BOWMAN'S CAPSULE



THE RENAL CORPUSCLE COMPRISES FOUR MAIN CELL TYPES

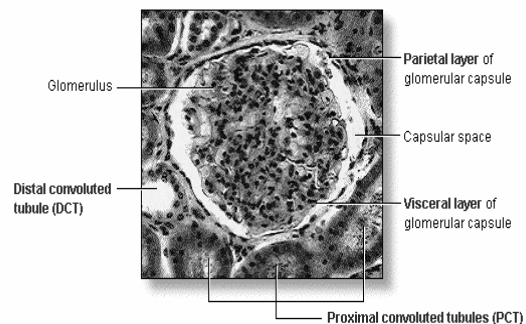
- 1) Endothelial cells which are fenestrated with
- 2) Visceral epithelial cells (podocytes) which support the delicate glomerular basement membrane by means of foot processes
- 3) Parietal epithelial cells which cover the Bowman's capsule;
- 4) Mesangial cells are contractile cells

Type I Medullary Interstitial Cells secrete PGE2

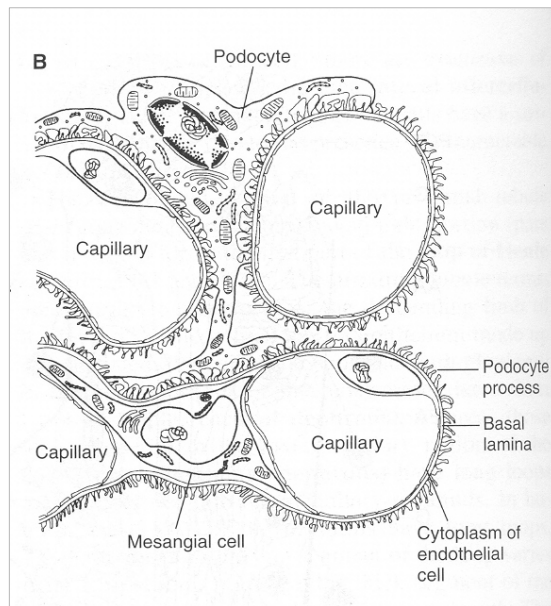
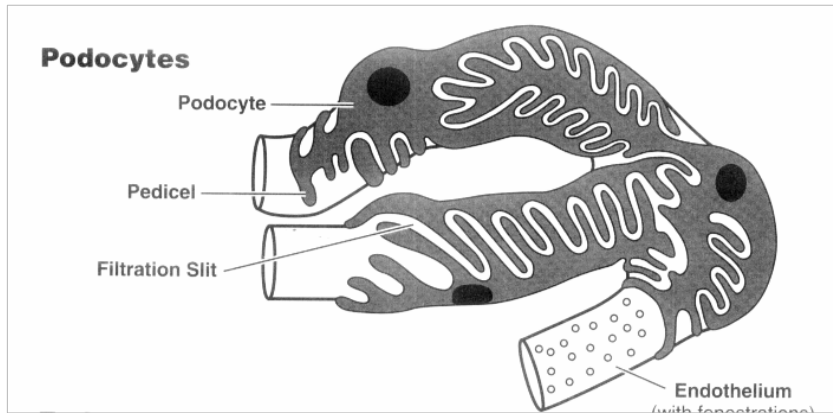
ULTRASTRUCTURE OF GLOMERULUS

PHOTOMICROGRAPH OF GLOMERULUS AND ADJACENT TUBULES

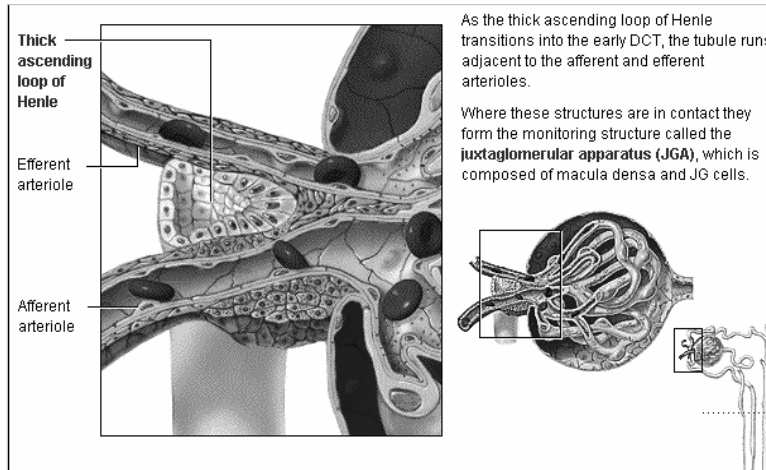
This photomicrograph shows a cross section of a glomerulus surrounded by a glomerular capsule.



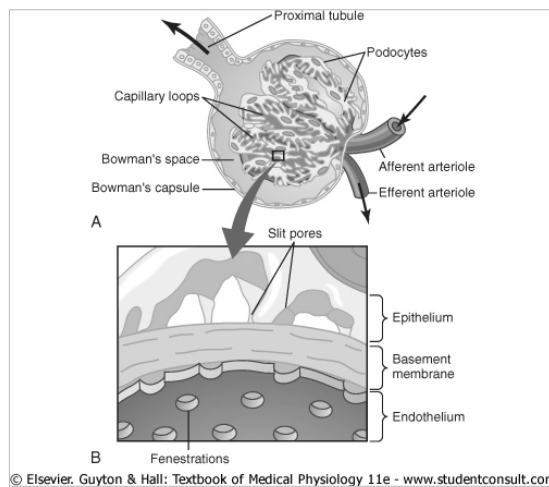
PODOCYTES

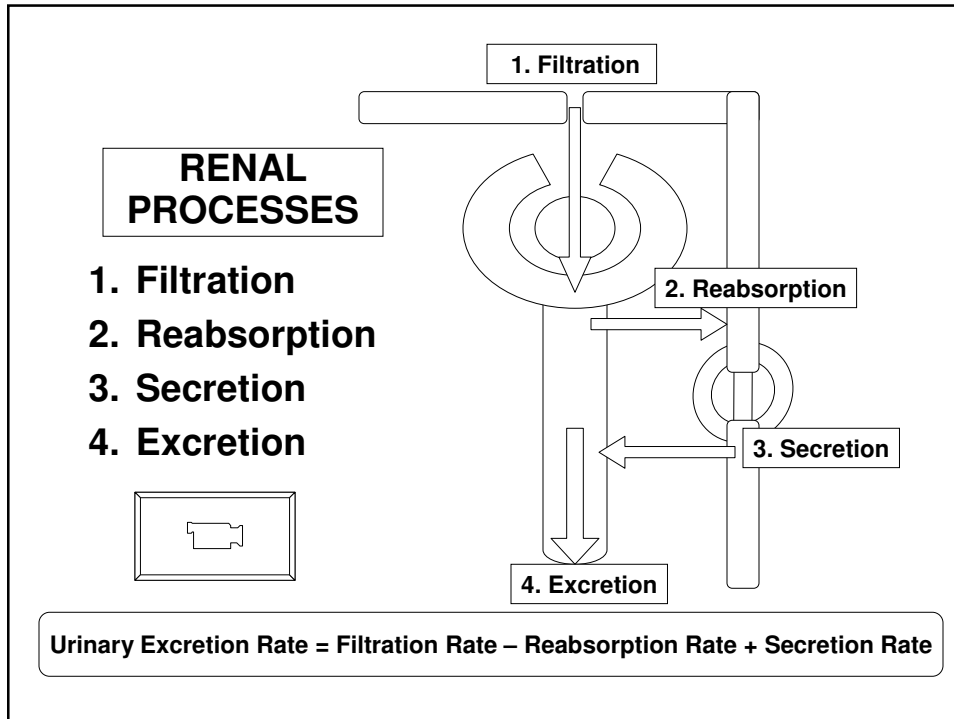


JUXTA GLOMERULAR APPARATUS



RENAL CORPUSCLE





RENAL PHYSIOLOGY
GLOMERULAR FILTRATION

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OBJECTIVES

At the end of this lecture you should be able to describe:

- **Filtration Membrane**
- **GFR, F_f, K_f, Plasma Clearance**
- **Control of GFR**
- **Regulation of Renal Blood Flow and GFR**

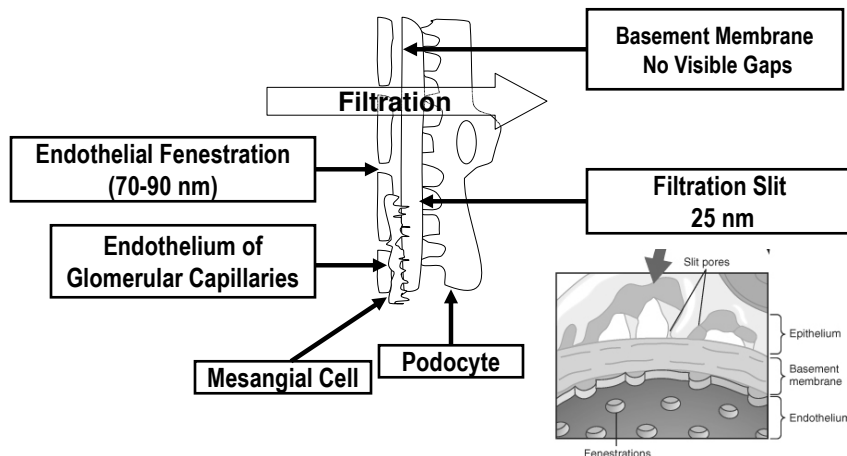


FILTRATION AND OSMOSIS

At a molecular level, filtration is the bulk flow of fluid through a membrane or other barrier that selectively impedes the movement of some molecules, the largest being impeded most. This process is sometimes called ULTRAFILTRATION.

FILTRATION MEMBRANE

4-8 nm size particles can be filtered easily

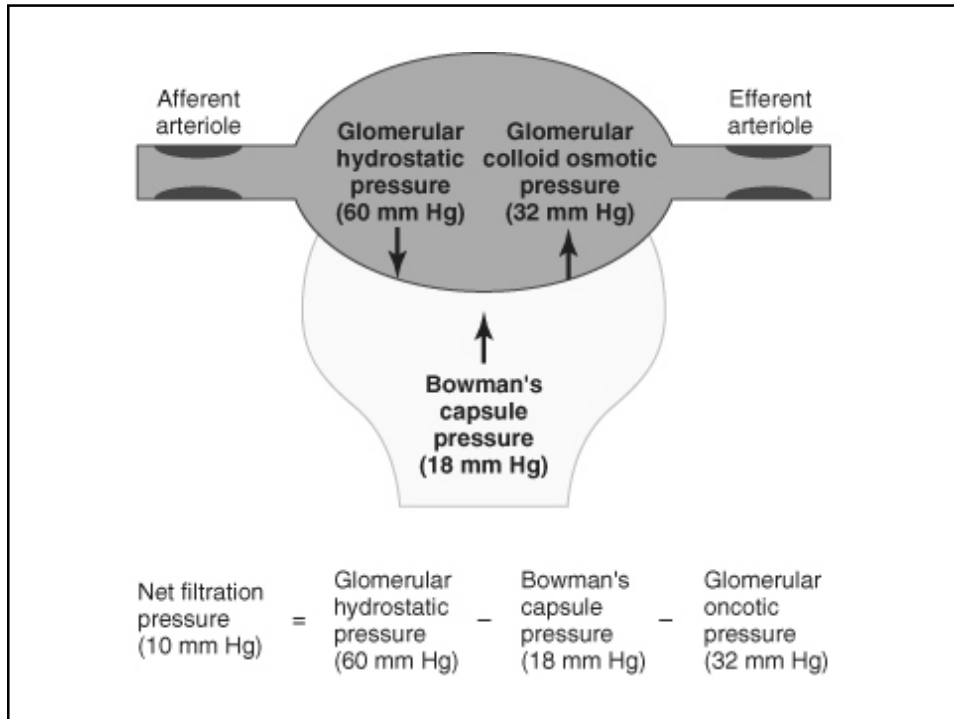


- **GFR=125ML/MIN**
- **OR $125 \times 60 \times 24 = 180000 = 180 \text{L/DAY}$**
- **Normal Urinary Output=1.5 L/day**
- **Daily Reabsorption $180 - 1.5 = 178.5 \text{ L/day}$**
- **Percent Reabsorbed = $178.5 \times 100 / 180 = 99.2\%$**
- **Percent Excreted = $100 - 99.2 = 0.8\%$
(Less than 1 % becomes urine)**

- **Obligatory Urinary Output=0.5-0.6L**
- **Oliguria= <300 ml/day**
- **Anuria= Less than 50 ml/day**
- **RBF=1100 ML/MIN**
- **RPF=1100x0.55=620 ML/MIN**
- **FILTRATION FRACTION =125/620X100=20%**
- **137x60x24=198 L/day if tubular reabsorption remains constant urine volume will increase from 1.5 to 19.5 L/day**

FILTRATION PRESSURES

- **PROMOTED BY:**
 - **HYDROSTATIC PRESSURE (from blood pressure)**
- **OPPOSED BY:**
 - **PLASMA COLOID OSMOTIC PRESSURE (oncotic pressure: proteins remain in capillaries)**
 - **HYDROSTATIC PRESSURE IN TUBULE**



DETERMINANTS OF GFR

Forces Favoring Filtration (mm Hg)	
Glomerular hydrostatic pressure	60
Bowman's capsule colloid osmotic pressure	0
Forces Opposing Filtration (mm Hg)	
Bowman's capsule hydrostatic pressure	18
Glomerular capillary colloid osmotic pressure	32
Net Filtration Pressure = 60 - (18 + 32) = +10 mm Hg	

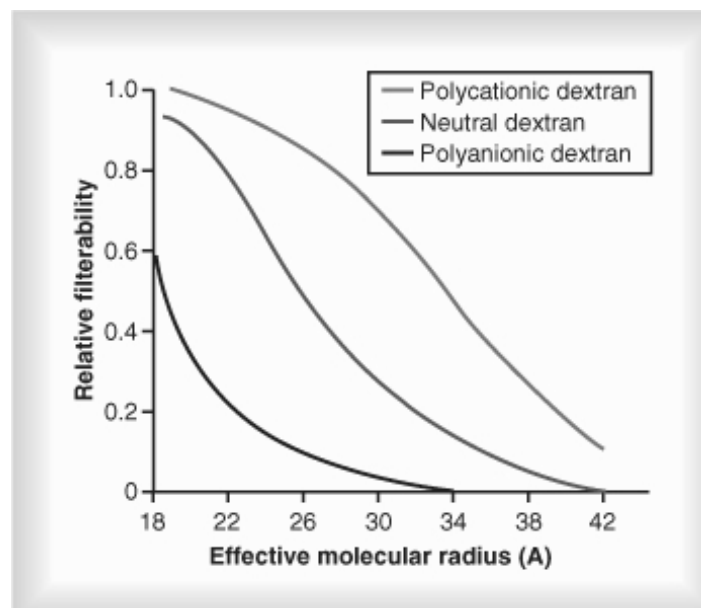
IgA Nephropathy

SUBSTANCE	MOLECULAR WEIGHT	MOLECULAR SIZE nm	FILTERABILITY
Water	18	0,15	1.0
Sodium	23	0,1	1.0
Glucose	180	0,33	1.0
Inulin	5,500	1.48	1.0
Myoglobin	17,000	1.88	0.75
Albumin (6 nm)	69,000	3.55	0.005

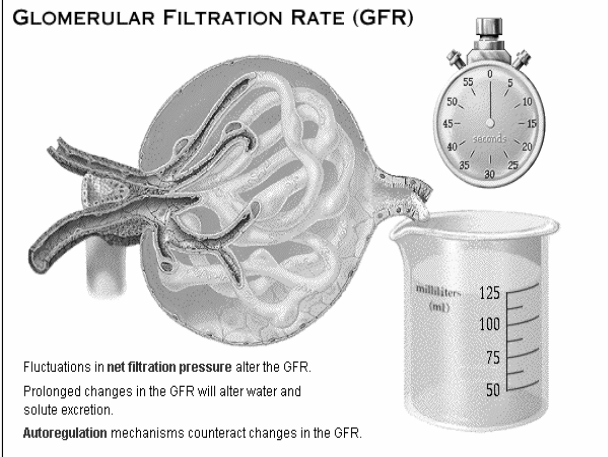
Filterability of Solutes Is Inversely Related to Their Size

Negatively Charged Large Molecules Are Filtered Less Easily Than Positively Charged Molecules of Equal Molecular Size.

Dextrans are polysaccharides that can be manufactured as neutral molecules or with negative or positive charges.



GLOMERULAR FILTRATION RATE



FILTRATION FRACTION

- Fraction of renal plasma that becomes Glomerular Filtrate

$$\begin{aligned} F_f &= \text{GFR} / \text{Renal Plasma Flow} \\ &= 125 \text{ ml per min} / 650 \text{ ml per min} \\ &= 19.2 \text{ or approximately } 20 \% \end{aligned}$$

About 20 per cent of the plasma flowing through the kidney is filtered through the glomerular capillaries

PLASMA CLEARANCE

The Volume of Plasma that is completely cleared of any substance by the Kidneys per minute is called the clearance of that particular substance

$$\text{Clearance} = \text{Urine Conc.} \times \text{Vol of Urine} / \text{Plasma Conc}$$

CONTROL OF GFR

$$\text{GFR} = K_f \times [(P_G - P_B) - (\pi_G - \pi_B)]$$

$$\text{GFR} = K_f \times [(60-18) - (32- 0)]$$

- (1) Hydrostatic pressure inside the glomerular capillaries (glomerular hydrostatic pressure, P_G), which promotes filtration
- (2) The hydrostatic pressure in bowman's capsule (P_B) outside the capillaries, which opposes filtration
- (3) The colloid osmotic pressure of the glomerular capillary plasma proteins (π_G), which opposes filtration
- (4) The colloid osmotic pressure of the proteins in bowman's capsule (π_B), which promotes filtration

FILTRATION COEFFICIENT

- **Glomerular Filtration Rate in both kidneys per mm Hg Filtration Pressure**

$$\begin{aligned} K_f &= \text{GFR} / \text{Filtration Pressure} \\ &= 125 \text{ ml per min} / 10 \text{ mm Hg} \\ &= 12.5 \text{ ml/min/mm Hg of filtration Pr} \end{aligned}$$

Increased Glomerular Capillary Filtration Coefficient Increases GFR

The Kf is a measure of the product of the Permeability and surface area of the glomerular capillaries. The Kf cannot be measured directly

$$\mathbf{K_f = GFR / \text{net filtration pressure}}$$

increased Kf raises GFR and decreased Kf reduces GFR

FACTORS AFFECTING GFR

- | |
|---|
| ❑ Changes in renal blood flow |
| ❑ Changes in Glomerular Capillary hydrostatic pressure |
| ❑ Changes in hydrostatic pressure in bowman's Capsule |
| ❑ Changes in concentration of plasma proteins (Dehydration, Hypoproteinemia etc.) |
| ❑ Changes in kf (Permeability and Surface Area) |

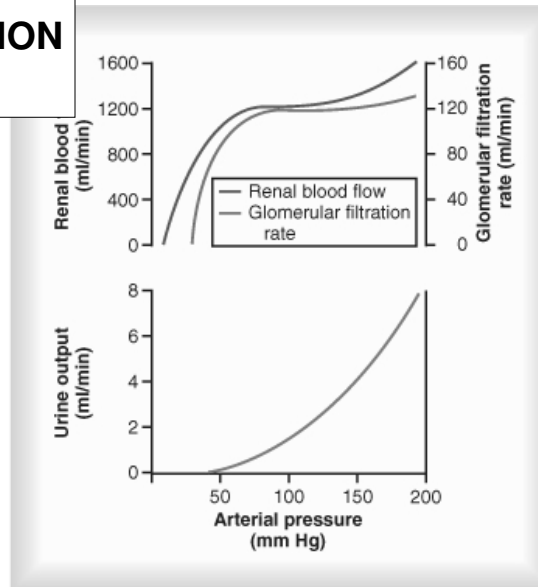
Physical Determinants*	Physiologic/Pathophysiologic Causes
$\downarrow K_f \rightarrow \downarrow \text{GFR}$	Renal disease, diabetes mellitus, hypertension
$\uparrow P_B \rightarrow \downarrow \text{GFR}$	Urinary tract obstruction (e.g., kidney stones)
$\uparrow \pi_G \rightarrow \downarrow \text{GFR}$	\downarrow Renal blood flow, increased plasma proteins
$\downarrow P_G \rightarrow \downarrow \text{GFR}$	
$\downarrow A_p \rightarrow \downarrow P_G$	\downarrow Arterial pressure (has only small effect due to autoregulation)
$\downarrow R_E \rightarrow \downarrow P_G$	\downarrow Angiotensin II (drugs that block angiotensin II formation)
$\uparrow R_A \rightarrow \downarrow P_G$	\uparrow Sympathetic activity, vasoconstrictor hormones (e.g., norepinephrine, endothelin)

K_f, glomerular filtration coefficient; **P_B**, Bowman's capsule hydrostatic pressure; **π_G** , glomerular capillary colloid osmotic pressure; **P_G**, glomerular capillary hydrostatic pressure; **A_p**, systemic arterial pressure; **R_E**, efferent arteriolar resistance; **R_A**, afferent arteriolar resistance.

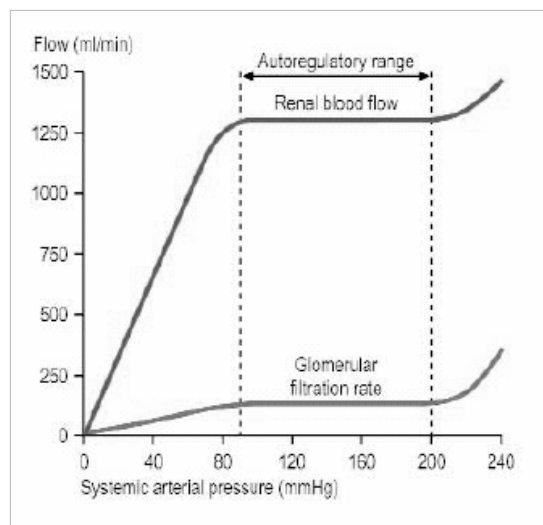
* Opposite changes in the determinants usually increase GFR.

AUTOREGULATION OF GFR

- GFR remains constant over a large range of values 75-160
- Autoregulation largely occurs by the regulation of renal vascular resistance



AUTOREGULATION OF RBF AND GFR



GFR REGULATION: *INTRINSIC*

1) MYOGENIC:

- Intrinsic property of mesangial (smooth muscle) cells and in afferent arteriole
- Reflex contraction induced by increased blood pressure reduces filtration

2) TUBULOGLOMERULAR FEEDBACK:

- Flow rate sensed by macula densa (part of juxtaglomerular apparatus), sends chemical signal to alter afferent arteriole resistance

GFR REGULATION: *EXTRINSIC*

± SYMPATHETIC INNERVATION

- Sympathetic stimulation/ epinephrine released from adrenal medulla cause arteriole vasoconstriction and reduced GFR

± ANGIOTENSIN II

- Produced as a result of renin release from kidney
- Constricts efferent ↑ arteriole; prevent decrease in GFR

GFR REGULATION: *EXTRINSIC*

(Cont.)

- ⚡ **NITRIC OXIDE (NO)**
 - Causes arteriolar vasodilation
 - Elevated NO may result in hyperfiltration of early Diabetes Mellitus
 - Reduced NO after salt intake may lead to hypertension

GFR REGULATION BY HORMONES OR AUTACOIDS

Hormone or Autacoid	Effect on GFR
Norepinephrine	↓
Epinephrine	↓
Endothelin	↓
Angiotensin II	↔ (prevents ↓)
Endothelial-derived nitric oxide	↑
Prostaglandins	↑

AGENTS AFFECTING MESANGIAL CELLS

CONTRACTION	RELAXATION
▪ Endothelins	▪ ANP
▪ Angiotensin II	▪ Dopamine
▪ Vasopressin	▪ PGE2
▪ Norepinephrine	▪ cAMP
▪ Platelet-activating factor	
▪ Thromboxane A2	
▪ PGF2	
▪ Histamine	
▪ Leukotriene C4&D4	

TUBULOGLOMERULAR FEEDBACK

