

- Depending on what our body needs at the time solutes can be filtrated, reabsorbed, secreted or excreted
- Creatinine is a true waste product

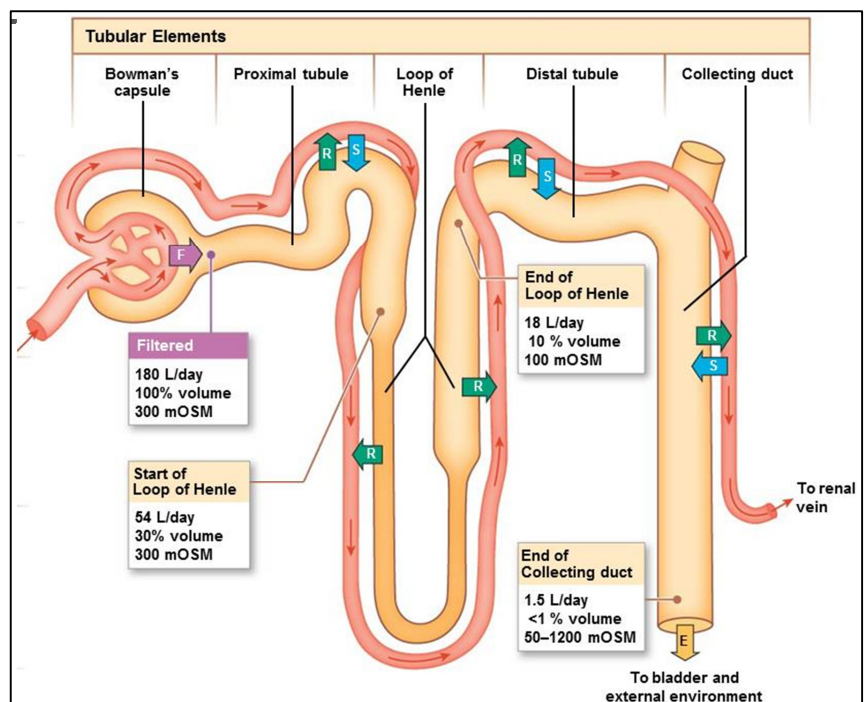
PRS: The plasma is MOST similar in chemical composition to the fluid in the:

- A. Proximal tubule
- B. Collecting duct
- C. Distal tubule
- D. Bowman's capsule**
- E. Ascending limb of the loop of Henle

	Filtration	Reabsorption	Secretion	Excretion
Water	+	+		1%
Na <sup>+</sup>	+	+		0.5%
K <sup>+</sup>	+	+	+	10%
Ca <sup>2+</sup>	+	+		2%
Phosphate	+	+		20%
Glucose	+	+		0%
Creatinine	+			100%
Urea	+	+		50%

### Changes in Filtrate along nephron

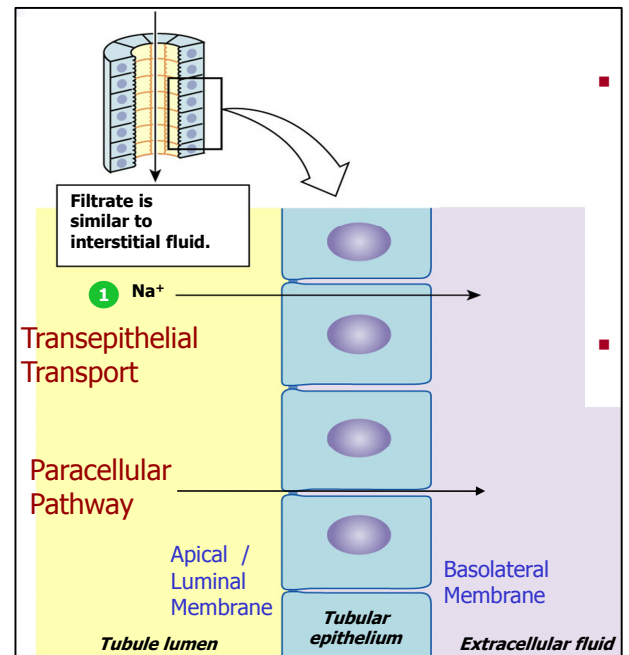
- Only consider the 20% of plasma that is filtered → assume the other 80% doesn't exist so 100% = all the plasma that gets filtered
- 300 mOsm = isosmotic
- Proximal tubule – bulk absorption and isosmotic absorption (reabsorb 70% of our fluid but it stays the same osmolarity) → made possible by the microvilli in the proximal tubule
- Once past the loop of Henle we further reabsorb 20% of the total filtrate but it is only 100 mOsm (hyposmotic → we have reabsorbed a lot more solute than we have fluid) – Controlled by counter current multiplier (discussed later)
- At the end of the collecting duct we reabsorb another 9% (approx.) – hormone controlled
- Wide range of osmolarity for the fluid coming out of the collecting duct and essentially ends up being urine – osmolarity can be from 50-1200 mOsm → can greatly dilute or concentrate our urine
  - This is important depending on our diet (i.e. eaten a lot of salt etc and get rid of without affecting the



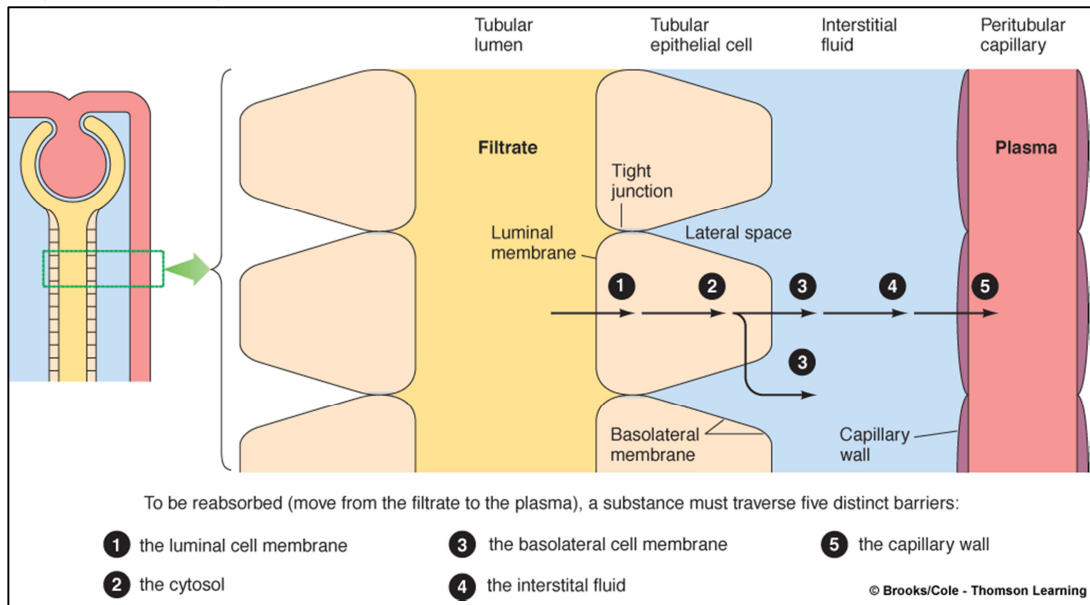
rest of the body much)

## Pathways for Tubular Reabsorption

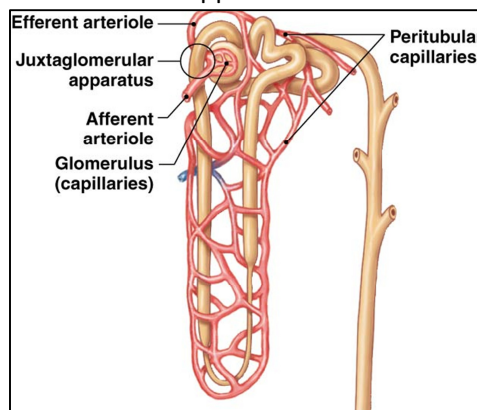
- Transepithelial transport
  - Substances cross both apical and basolateral membrane
  - Has to cross a cell
  - Usually an active process because we have to create concentration gradients
- Paracellular pathway
  - Substances pass through the junction between two adjacent cells
  - Create tight junctions



## Steps of Transepithelial Transport

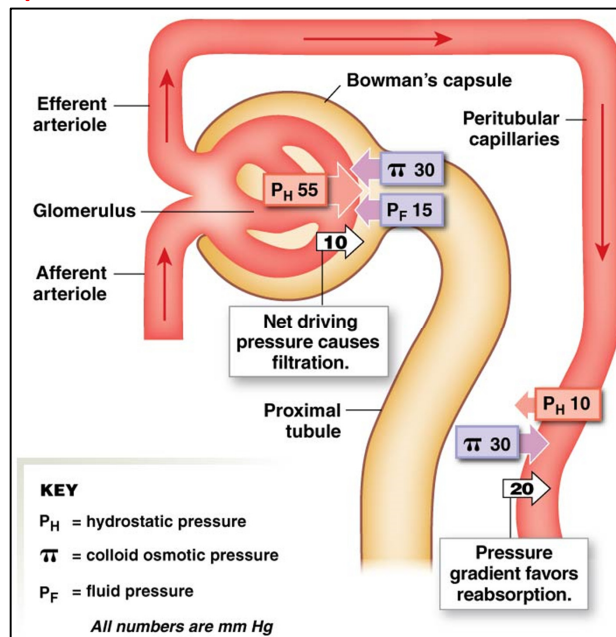


- First need to go through luminal membrane
- Either 4 or 5
  - 4 – stay in interstitial fluid
  - 5 – but if full reabsorption needs to happen cross the interstitial fluid into the plasma



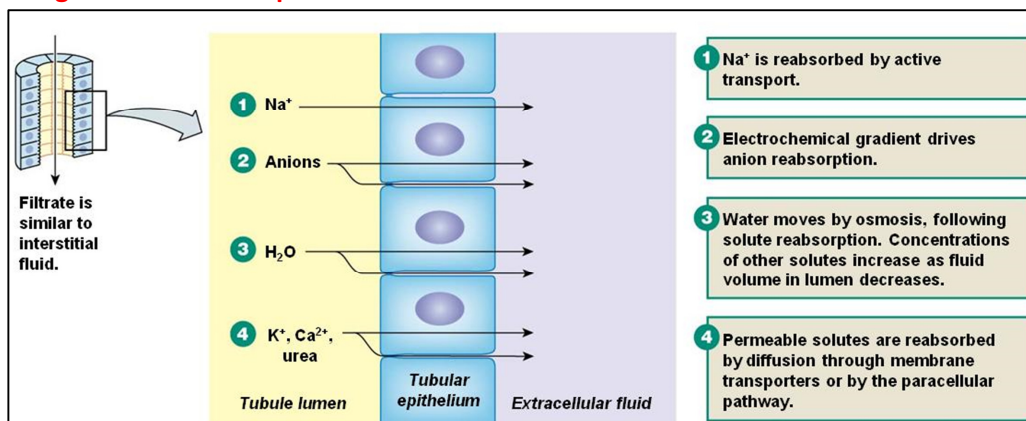
- Position of blood vessels around the nephron
- Efferent arteriole forms a large part of our peritubular capillary – it sits intertwined very close to the tubule which decreases diffusion distance and helps to drive reabsorption

## Reabsorption into Peritubular Capillaries



- Hydrostatic pressure is much lower when we go down the efferent tubule than in our bowman's capsule

## Principles governing Tubular Reabsorption



- $\text{Na}$  is the first thing that moves in and it helps to move all the other stuff
- $\text{Na}$  reabsorption = ACTIVE PROCESS
- When  $\text{Na}$  is reabsorbed, there is a net positive charge in the ECF so anions naturally want to follow → electrochemical gradient
- Now we have an osmolarity gradient as the osmolarity in the ECF has increased due to the reabsorption of the solutes
  - For 3 – i.e. in proximal tubule (however not in all parts of nephron → i.e. in collecting duct which are usually impermeable to water can become permeable to water depending on what hormones are circulating and contacting them)

PRS: Which one/s of these solutes is/are dependent on active transport to be reabsorbed? (active reabsorption)

- Amino acids
- Glucose
- Sodium
- Anions

Explanation: Active reabsorption

There are 2 types of active transport: primary and secondary

- $\text{Na}$  undergoes primary active transport as it is the molecule that is transported by the ATPase but it is the energy created by the  $\text{Na}$  (i.e. electrochemical and osmotic gradients) that forces the movement of the other solutes