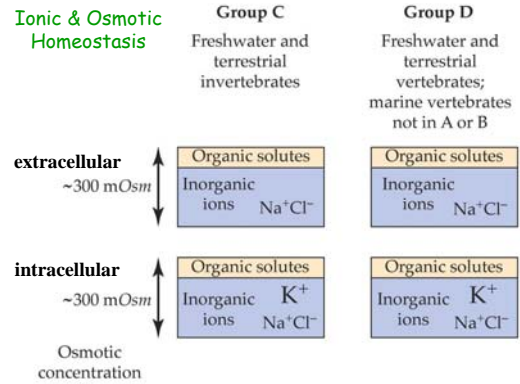


# osmoregulation mechanisms in gills, salt glands, and kidneys

## Ionic & Osmotic Homeostasis



ANIMAL PHYSIOLOGY, Figure 35.10 (Part 2) © 2014 Sinauer Associates, Inc.

## Salt Secretion:

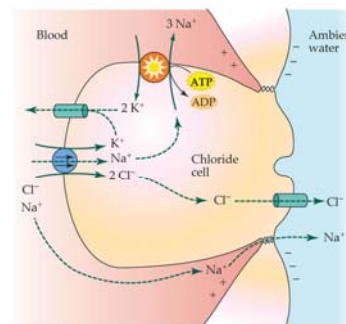
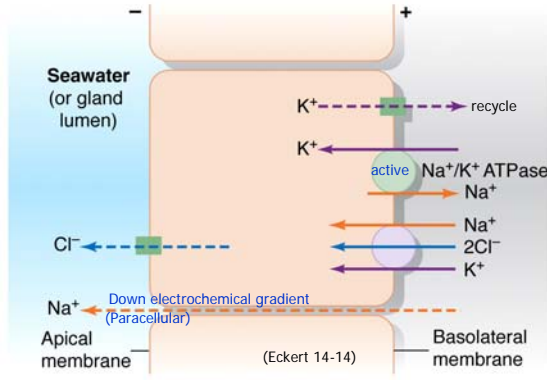


Figure in Box 26.2 Hill et al. 2004

## Chloride Cell

ANIMAL PHYSIOLOGY, Box 26.2 © 2014 Sinauer Associates, Inc.

## Salt Glands

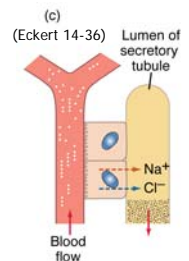
- Shark **rectal glands** to dispose of excess NaCl
- blood **hyperosmotic** to seawater, but **less salt**
- more **urea** and **TMAO** (trimethylamine oxide)
- NaCl actively secreted



## Shark Rectal Salt Glands

### Salt-secreting cells:

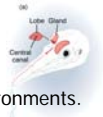
- Na<sup>+</sup>/K<sup>+</sup>-ATPase pump in basolateral membrane
- generates gradient for Na<sup>+</sup> by which Na<sup>+</sup>/2Cl<sup>-</sup>/K<sup>+</sup> cotransporter drives up [Cl<sup>-</sup>] in cell
- Cl<sup>-</sup> across apical membrane
- Na<sup>+</sup> follows paracellularly down electrochemical gradient (and H<sub>2</sub>O)
- apical membrane **impermeable** to urea and TMAO
- therefore **iso-osmotic** secretion with lots of NaCl



... slightly different in birds and lizards →

Salt Glands

(Eckert 14-36)



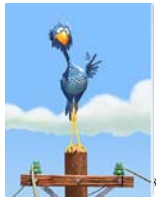
Nasal/orbital salt glands of birds and reptiles -especially species in desert or marine environments.

Hypertonic NaCl secretions (2-3x plasma osmolarity)

Allows some birds to drink salt water and end up with osmotically free water



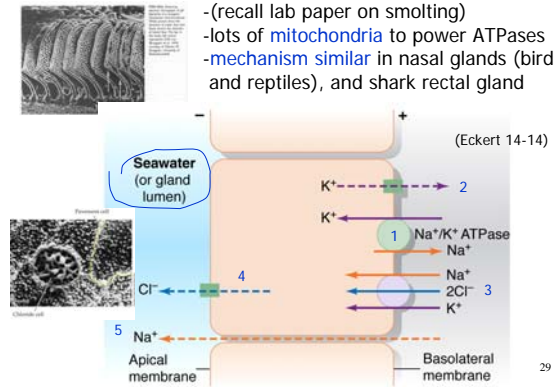
Amblyrhynchus cristatus



Fish Gills

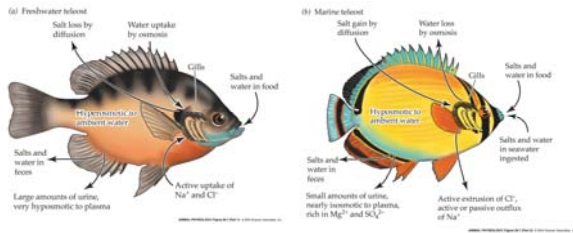
Chloride cells involved in osmoregulation

- (recall lab paper on smolting)
- lots of mitochondria to power ATPases
- mechanism similar in nasal glands (birds and reptiles), and shark rectal gland



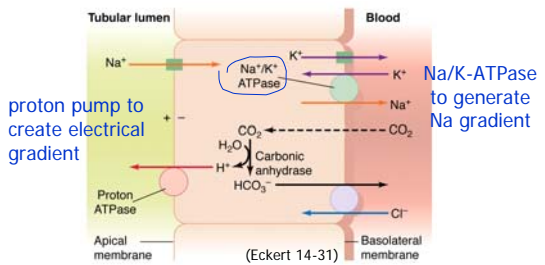
Freshwater fish:

The mechanism basically reversed to allow uptake of salt from water against concentration gradient



Hill et al. 2004, Fig 26.7

30



31

Sea ↔ Freshwater

(recall lab paper on smolting)

Switch between getting rid of excess salt in seawater and taking up salt in freshwater



Growth hormone and cortisol for → sea (more active chloride cells with more Na/K-ATPase activity)

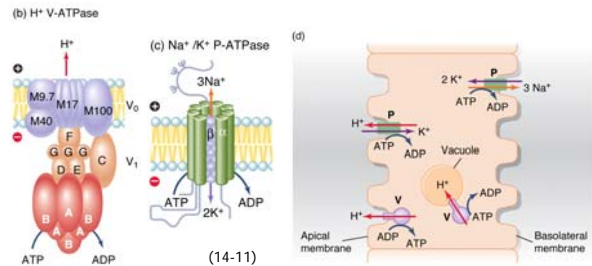
Prolactin for → freshwater



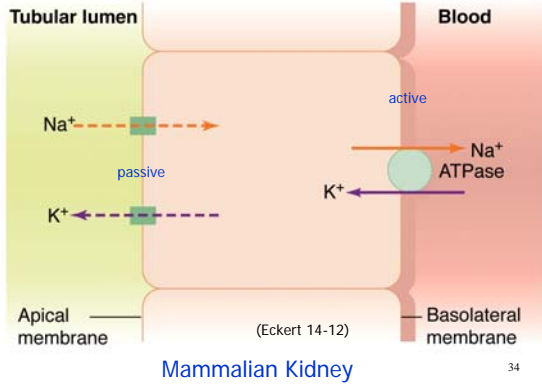
Osmoregulatory Mechanisms

Apical surface (faces lumen and outside world)  
Basal surface (faces body and extracellular fluid)

- Active movement of ions/salts requires ATP
- Movement of water follows movement of ions/salts

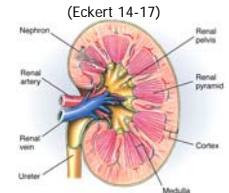


Gradients established and used...to move ions, water



Kidney Functions:

- Osmoregulation
- Blood volume regulation
- Maintain proper ion concentrations
- Dispose of metabolic waste products
- pH regulation (at ~ 7.4)
- Dispose of toxins and foreign substances



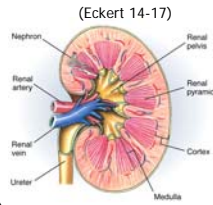
How does the kidney accomplish this?

Mammalian Kidney

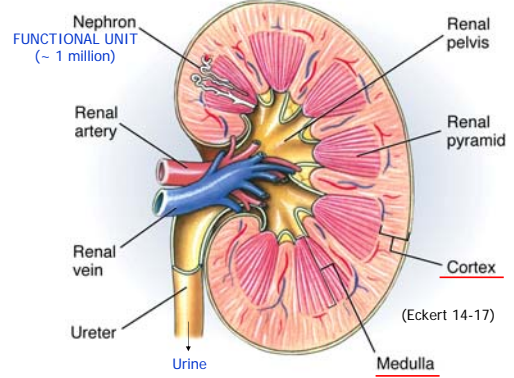
- Paired
- 1% body mass
- 20% blood flow

-urine contains:  
water  
metabolic byproducts (e.g., urea)  
excess salts etc.

-from ureter to urinary bladder  
(smooth muscle, sphincter, inhibition)  
-out via urethra during micturition

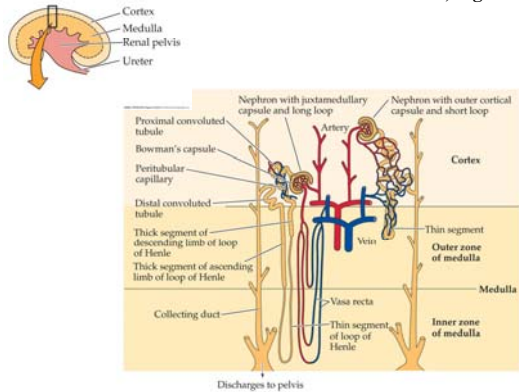


Mammalian Kidney Anatomy



(a) Kidney in cross section

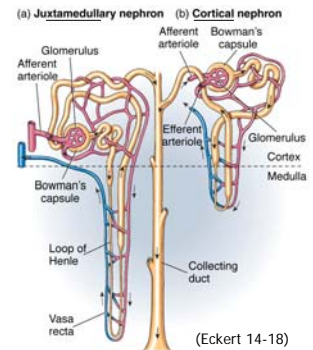
Hill et al. 2004, Fig. 27.6



Nephron Anatomy

- 1 -Proximal tubule
- 2 -Loop of Henle  
-descending  
-ascending
- 3 -Distal tubule

-numerous nephrons empty into collecting duct  
-collecting ducts empty into renal pelvis



## Nephron Anatomy

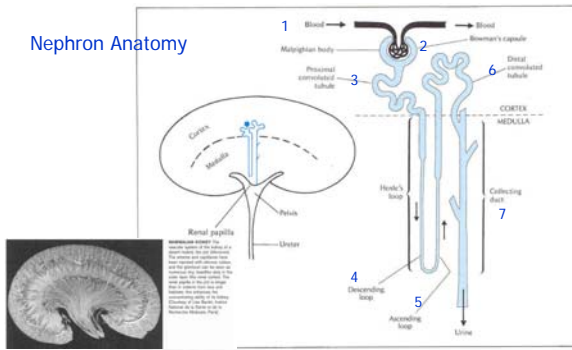


Figure 9.8 Diagram of a mammalian kidney. The kidney contains a large number, up to several million, of single nephrons. Only one nephron is indicated in this diagram and is shown enlarged to the right. The outer layer of the kidney, the cortex, contains the Malpighian bodies and the proximal and the distal convoluted tubules. The capillary network within the Malpighian body is known as the glomerulus. The inner portion, the medulla, contains Henle's loops and collecting ducts. The urine is initially formed by ultrafiltration in the Malpighian bodies. The filtrate fluid is modified and greatly reduced in volume as it passes down the renal tubule and into the collecting ducts. These empty the urine into the renal pelvis, from where it is conveyed to the bladder. Knut Schmidt\_Nielsen 1997

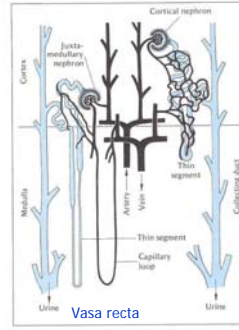
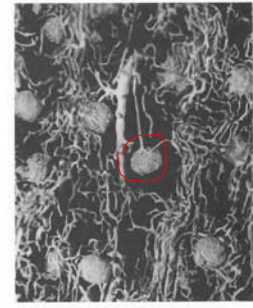


Figure 9.11 Diagram of a nephron showing the vasa recta and countercurrent exchange. The vasa recta are a network of blood vessels that facilitate countercurrent exchange between the descending and ascending limbs of the loop of Henle. The diagram illustrates the cortical nephron and the juxtamedullary nephron. Labels include: Cortex, Juxtamedullary nephron, Cortical nephron, Thin segment, Thick segment, Capillary loop, Vasa recta, Artery, Vein, and Collecting duct. Knut Schmidt\_Nielsen 1997

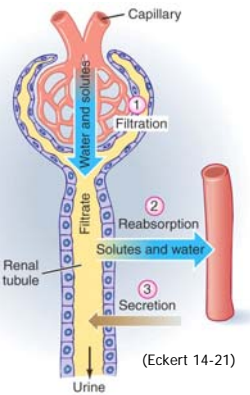


DOG KIDNEY Close-up view of glomeruli from a dog kidney after arterial injection of silicone rubber. The glomerulus in the center of the photo shows the slightly thicker vessel leading into the glomerulus and the somewhat thinner vessel leaving it. The diameter of these vessels is about 15 to 20  $\mu\text{m}$ ; the diameter of the glomerulus is about 150  $\mu\text{m}$ . [Courtesy of A. Clifford Berger, Harvard University]. Knut Schmidt\_Nielsen 1997

## Kidney Processes- overview

1. **FILTRATION**  
blood --> filtrate
2. **REABSORPTION**  
filtrate --> blood
3. **SECRETION**  
blood --> filtrate

All 3 involved in final Urine Composition



(Eckert 14-21)

Mosm = x1000 U/P

Animal	Urine maximum osmotic concentration (Osm liter <sup>-1</sup> )	Urine/plasma concentration ratio
Beaver <sup>a</sup>	0.52	2
Pig <sup>a</sup>	1.1	3
Human <sup>a</sup>	1.4	4
White rat <sup>a</sup>	2.9	9
Cat <sup>a</sup>	3.1	10
Kangaroo rat <sup>a</sup>	5.5	14
Sand rat <sup>a</sup>	6.3	17
Hopping mouse <sup>a</sup>	9.4	25

<sup>a</sup> B. Schmidt-Nielsen and O'Dell (1961).  
<sup>b</sup> K. Schmidt-Nielsen (1964).  
<sup>c</sup> MacMillen and Lee (1997).

Table 9.2 The maximum concentrating of various mammals is correlated with the animal, desert animals having the concentrations and fresh-water animals.

## Filtration plus secretion

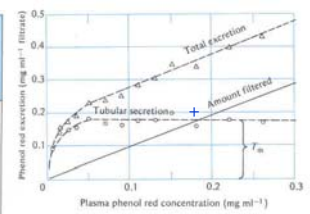


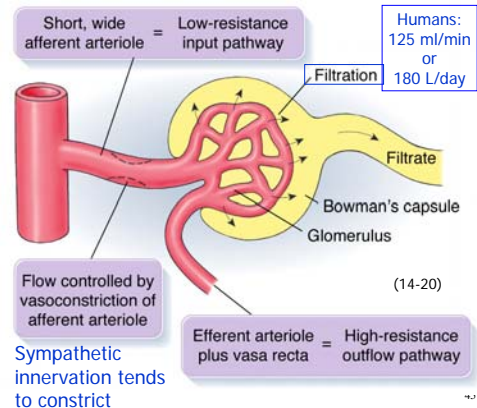
Figure 8.10 Excretion of phenol red by the bullfrog. The amount filtered increases in proportion to the plasma concentration. In addition, phenol red is added to the urine by active tubular transport, increasing the amount in the urine. The amount added by tubular secretion remains constant at plasma concentrations above 0.05 mg per milliliter; this indicates that the tubular maximum for phenol red has been reached. [Forster 1940]

Knut Schmidt\_Nielsen 1997

U/P ratio	Implications for excretion		
	Effects on water excretion	Effects on solute excretion	Effects on composition of blood plasma
U/P = 1 (isotonic urine)	Water is excreted in the same relation to solutes as prevails in the blood plasma.	Solutes are excreted in the same relation to water as prevails in the blood plasma.	The formation of urine leaves the ratio of solutes to water in the blood plasma unchanged, thus does not alter the plasma osmotic pressure.
U/P < 1 (hypoosmotic urine)	Water is preferentially excreted. Urine contains more water relative to solutes than plasma.	Solutes are preferentially held back from excretion. Urine contains less solutes relative to water than plasma.	The ratio of solutes to water in the plasma is shifted upward. The osmotic pressure of the plasma is raised.
U/P > 1 (hyperosmotic urine)	Water is preferentially held back from excretion. Urine contains less water relative to solutes than plasma.	Solutes are preferentially excreted. Urine contains more solutes relative to water than plasma.	The ratio of solutes to water in the plasma is shifted downward. The osmotic pressure of the plasma is lowered.

Hill et al. 2004, Fig 25.7

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Flow controlled by vasoconstriction of afferent arteriole  
Sympathetic innervation tends to constrict

Humans: 125 ml/min or 180 L/day

Filtration:

Bowman's capsule

- 3 layers
  1. **Glomerular endothelial cells**
    - 100x leakier than other capillary walls
  2. **Basement membrane**
    - negatively charged glycoproteins
    - repel plasma proteins by charge
  3. **Epithelial cells**
    - podocytes create slits

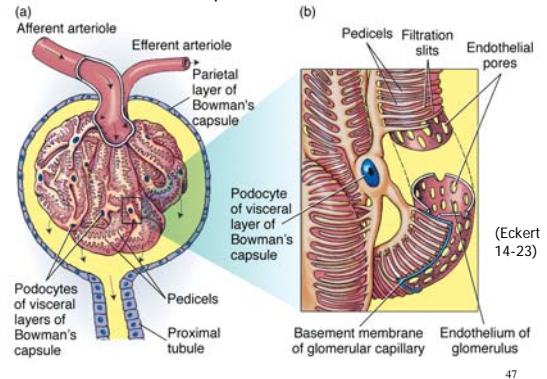
Filtrate = protein-free and cell-free plasma

Glomerular Filtration Rate (GFR)

Humans: 125 ml/min or 180 L/day (60x plasma vol.)

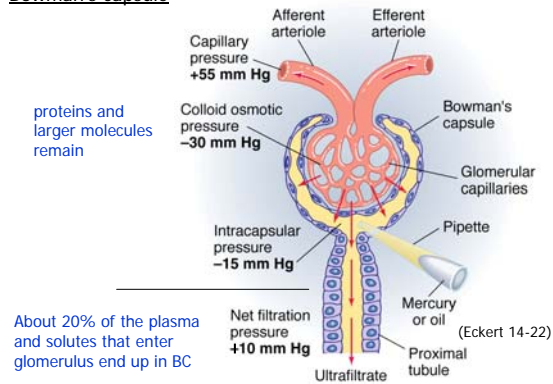
46

Bowman's capsule

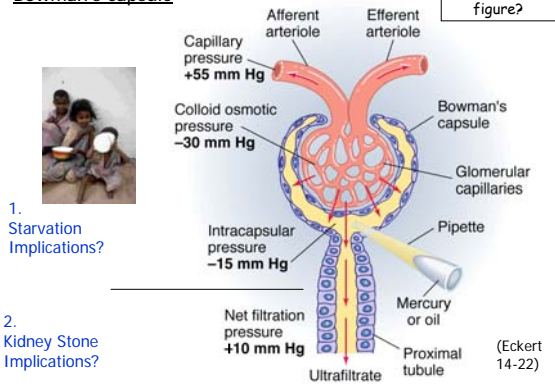


47

Bowman's capsule

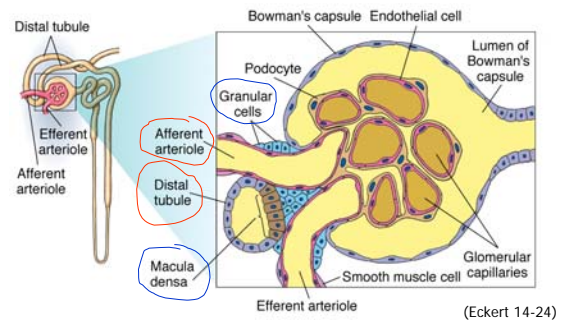


Bowman's capsule



Filtration Regulation:

1. Myogenic props. of afferent arteriole resist stretch
2. Secretions from cells of **juxtaglomerular apparatus** (where distal tubule passes near bowman's capsule)
  - Macula densa** cells (distal tubule)
    - monitor osmolarity and flow in distal tubule
    - paracrine hormonal activity on afferent arteriole
  - Granular or juxtaglomerular** cells (afferent arteriole)
    - release **renin** which alters blood pressure...



50

51

**Filtration Regulation:**

**Renin** (from granular cells) released in response to  
 -low renal BP,  
 -low solute [ ] in distal tubule,  
 -or sympathetic activation

Renin leads to activation of **Angiotensin II** which causes systemic **vasoconstriction** to inc. BP  
 stimulates **aldosterone** from adrenal cortex  
**vasopressin (ADH)** from post. pit.  
 (these promote salt, water reabsorption)

3. Sympathetic innervation (reduce GFR)  
 -afferent vasoconstriction  
 -decreased space between podocytes

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**Renal Clearance:**

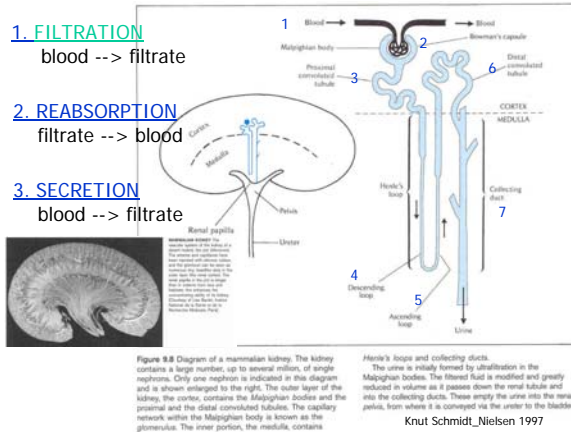
Volume of plasma cleared of a substance by the kidney.

(Filtration, Reabsorption, Secretion)

Inulin (=GFR) b/c neither reabsorbed nor secreted

If clearance > GFR = secretion  
 If clearance < GFR = reabsorption

53



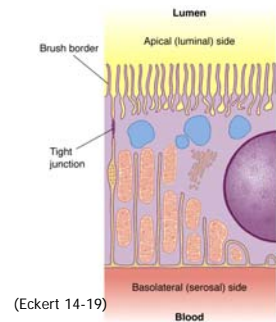
**Reabsorption:**

of 180 L/day filtered, ~178.5 L reabsorbed in humans

Lots of active transport of salts and other substances

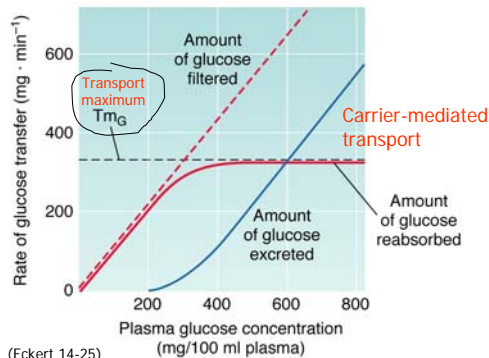
Tight junctions not so tight in proximal tubule, so water can move from filtrate to plasma

Because of reabsorption (and secretion), Renal clearance does NOT often equal GFR

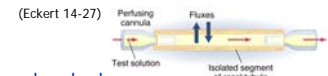


**Reabsorption limit – Glucose example**

T<sub>m</sub> at 300 mg/min/100ml plasma



**Reabsorption:**



**Proximal Tubule**

70% filtered Na<sup>+</sup> actively reabsorbed (by Na<sup>+</sup>/K<sup>+</sup>ATPase pump in basolateral membrane) Cl<sup>-</sup> and water follow

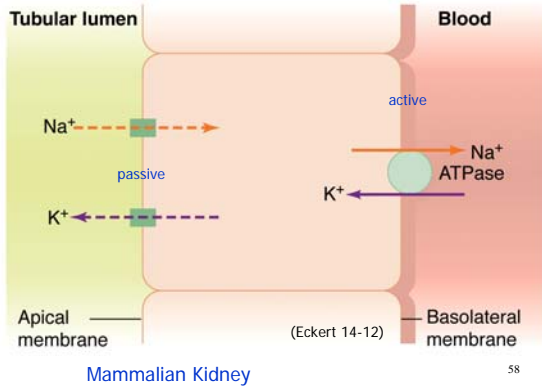
75% of filtrate is reabsorbed including glucose and amino acids (Na<sup>+</sup> dependent) also, phosphates, Ca<sup>+</sup>, electrolytes as needed

Parathyroid hormone controls phosphate and Ca<sup>+</sup> reabsorp. triggers calcitriol production (Vit. D) for Ca<sup>+</sup>

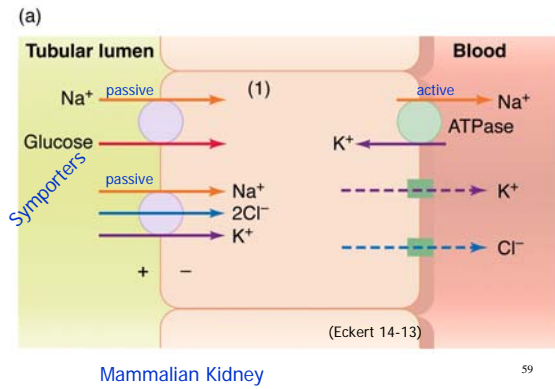
At end of proximal tubule filtrate is **isosmotic** with plasma (~300mOsm) however, remaining substances are 4x concentrated

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Gradients established and used:



Gradients established and used:



Reabsorption:

Loop of Henle

Descending limb

- no active NaCl transport
- low urea and NaCl permeability
- permeable to water

Ascending thin limb

- no active NaCl transport
- but permeable to NaCl
- low urea permeability
- low water permeability

Ascending thick limb

- NaCl transported out of tubule
- low water permeability

Countercurrent multiplier

One driver of concentrating mechanism of nephron

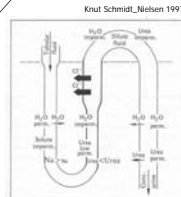
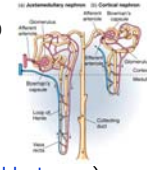


Figure 9-11 Diagram of the concentrating mechanism in the loop of Henle in the mammalian kidney using the principle of countercurrent multiplier. Active transport of chloride ion is indicated by heavy arrows; passive flow of water and ions by light arrows. (Kisler and Taylor 1976)

Reabsorption (and Secretion):

Distal Tubule

- K<sup>+</sup>, H<sup>+</sup>, NH<sub>3</sub> into tubule
- under endocrine control
- Na<sup>+</sup>, Cl<sup>-</sup>, HCO<sub>3</sub><sup>-</sup> back into body
- water follows
- (Na<sup>+</sup> reabsorption facilitated by aldosterone)



Angiotensinogen  $\xrightarrow{\text{Renin}}$  Ang. I  $\xrightarrow{\text{ACE in lung}}$  Ang. II  $\rightarrow$  aldosterone from adrenal cortex  
 ADH from post. pit.

Collecting Duct

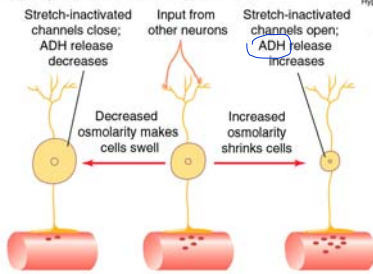
- permeable to water
- hormone control (ADH/vasopressin)
- water (via aquaporins) follows osmotic gradient

- permeable to Urea in inner medulla

Another driver of concentrating mechanism of nephron

-ADH role in water reabsorption/urine concentration

(a) Magnocellular neurons of hypothalamus



-Renin  $\rightarrow$  Ang. II  $\rightarrow$  ADH

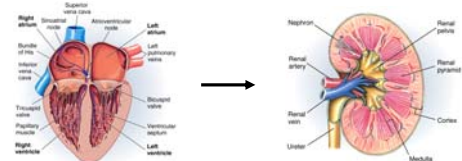
-Baroreceptor input (atrial and arterial)

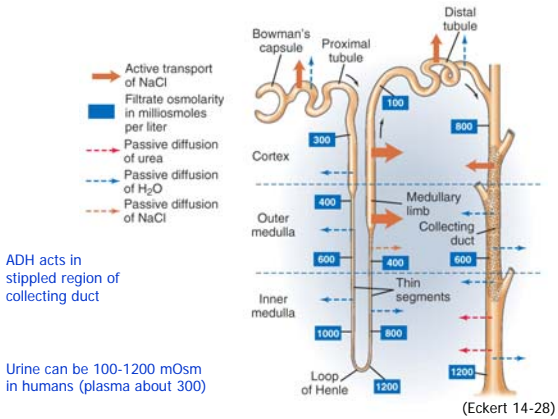
-EtOH inhibits ADH release

Atrial Natriuretic Peptide (ANP)

- released by atrium cells in response to stretch (elevated BP)

- opposite effect of renin-angiotensin system
- decreases sodium reabsorption
- therefore increased urine production
- ANP inhibits release of ADH, renin, aldosterone





Secretion:

From plasma into tubule of nephron

K<sup>+</sup>, H<sup>+</sup>, NH<sub>3</sub>, organic acids, organic bases

Organic anions (OA<sup>-</sup>):

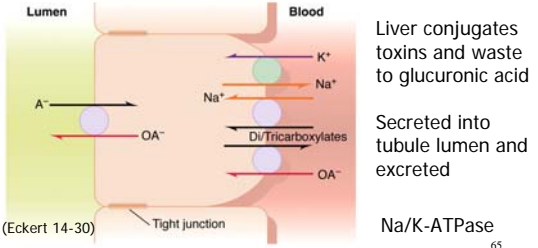


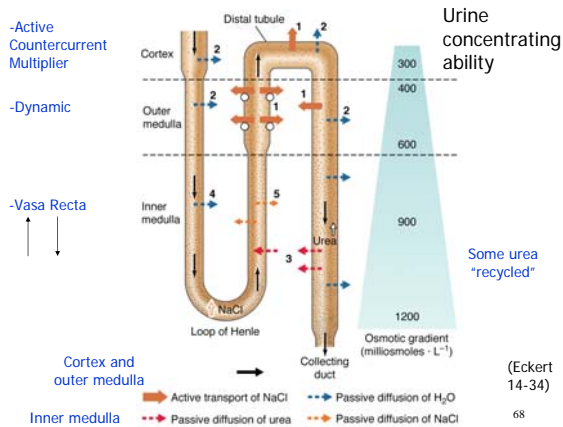
Table 14-9 Some organic ions secreted by the proximal tubule  
Eckert

Anions	Cations
<i>Endogenous</i>	<i>Endogenous</i>
Urates	Dopamine
Hippurates	Epinephrine
Oxalate	Norepinephrine
Prostaglandins	Creatinine
cAMP	
<i>Exogenous</i>	<i>Exogenous</i>
Furosemide	Morphine
Bumetanide	Amiloride
Penicillin	Quinine
Aspirin	Atropine
Chlorothiazides	Isoproterenol

Countercurrent Exchangers (passive)

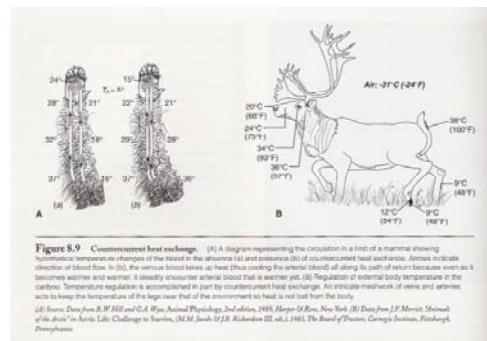
Countercurrent Multipliers (active)

See p.736 in your text



Endotherms in the COLD...

Countercurrent Heat Exchange



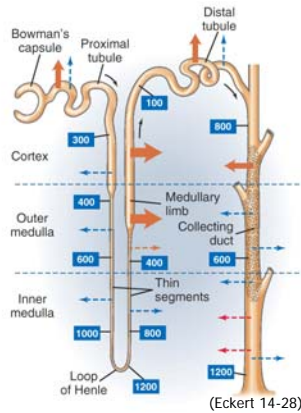


Urine concentrating ability

- Active transport of NaCl
- Filtrate osmolarity in milliosmoles per liter
- - - Passive diffusion of urea
- - - Passive diffusion of H<sub>2</sub>O
- - - Passive diffusion of NaCl

ADH acts in stippled region of collecting duct

Urine can be 100-1200 mOsm in humans (plasma about 300)

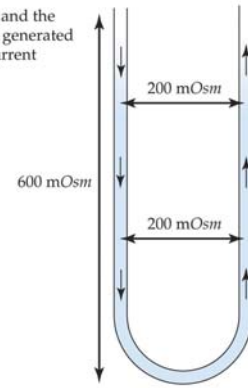


(Eckert 14-28)

(a) The single effect and the end-to-end gradient generated from it by countercurrent multiplication

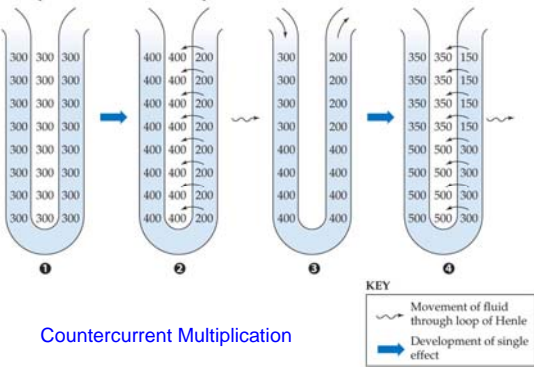
1. Single Effect
2. End-to-end Gradient

Figure 27.12 Hill et al. 2004



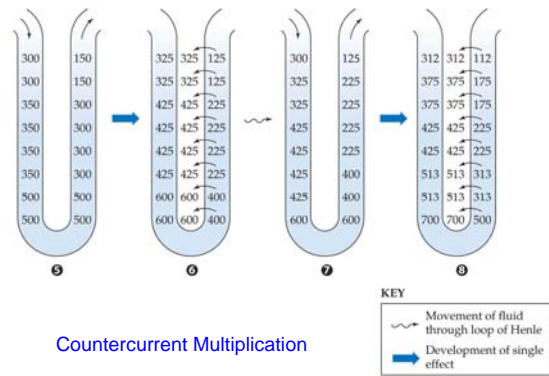
ANIMAL PHYSIOLOGY: Figure 27.12 (Part 1) © 2004 Sinauer Associates, Inc.

(b) The process of countercurrent multiplication



Countercurrent Multiplication

ANIMAL PHYSIOLOGY: Figure 27.12 (Part 2) © 2004 Sinauer Associates, Inc.



Countercurrent Multiplication

ANIMAL PHYSIOLOGY: Figure 27.12 (Part 3) © 2004 Sinauer Associates, Inc.

Same story, different picture

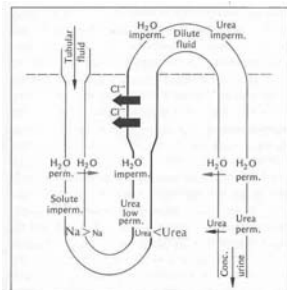


Figure 9.12 Diagram of the concentrating mechanism in the loop of Henle in the mammalian kidney during the formation of concentrated urine. Active transport of chloride ion is indicated by heavy arrows; passive flux of water and urea by light arrows. [Kokko and Tisher 1976]

Knut Schmidt-Nielsen 1997

Non-mammalian kidneys:

-Only birds also have loops of henle

-Freshwater fish with more and larger glomeruli to make lots of dilute urine



-Some marine fish without glomeruli or bowman's capsule – urine formed by secretion, ammonia secreted by gills

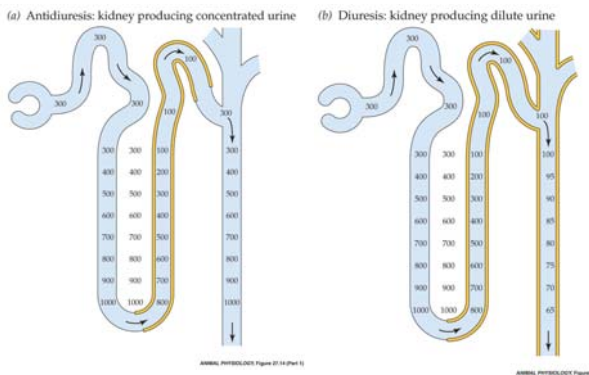
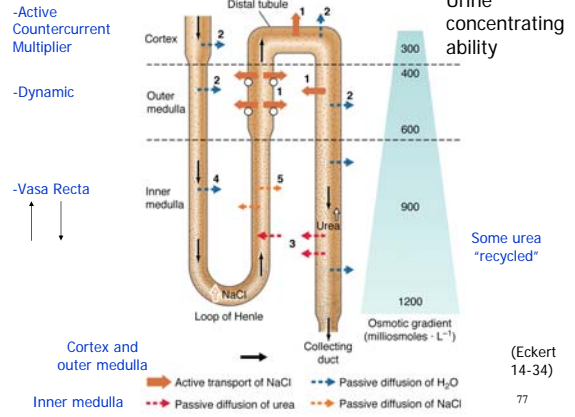
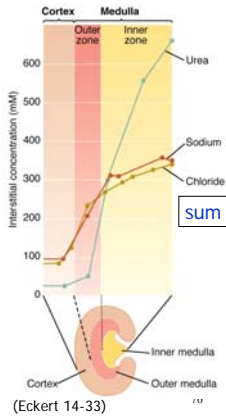
-Osmoregulation also via extrarenal organs

Urine concentrating ability

1200 mOsm in humans  
 9000 mOsm on kangaroo rats  
 9600 mOsm in *Perognathus* (mouse)



-Length of loops of henle  
 -Corticomedullary concentration gradient



Hill et al. 2004, Fig. 27.14

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Urine concentrating ability

-Vasa Recta

-Loops of Henle only in Mammals and Birds -> Hyperosmotic Urine

