## MECHANISMS FOR CONCENTRATING & DILUTING URINE

Regulation of ECF osmolarity

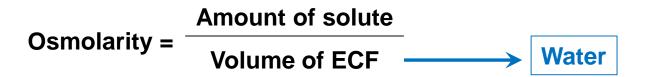
## **Objectives**

At the end of this session, students should be able to;

- Identify and describe that the loop of Henle is referred to as countercurrent multiplier and the vasa recta as countercurrent exchanger systems in concentrating and diluting urine.
- Explain what happens to osmolarity of tubular fluid in the various segments of the loop of Henle when concentrated urine is being produced.
- Explain the factors that determine the ability of loop of Henle to make a concentrated medullary gradient.
- Differentiate between water diuresis and osmotic diuresis.
- Appreciate clinical correlates of diabetes mellitus and diabetes insipidus.

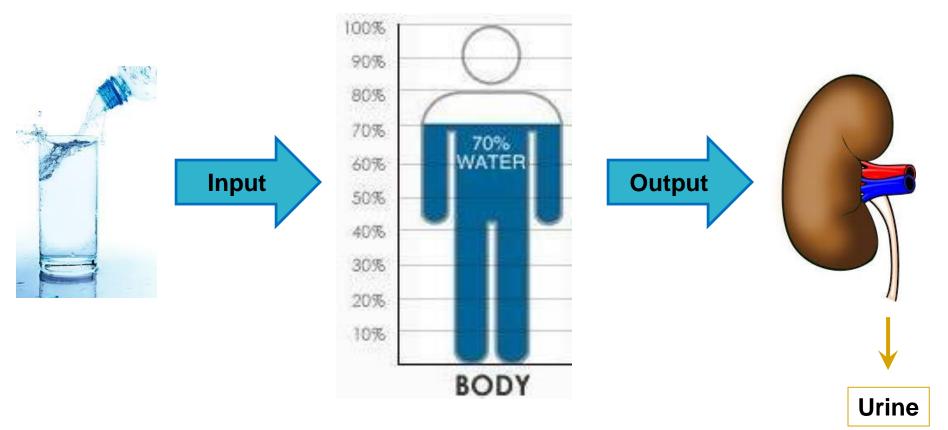


- Maintaining a constant concentration of solutes & electrolytes in the ECF is important for normal cellular function.
- The concentration of solutes in the ECF = osmolarity.
- Normal ECF osmolarity ≈ 300 mOsm/L
- What determines the osmolarity of ECF?

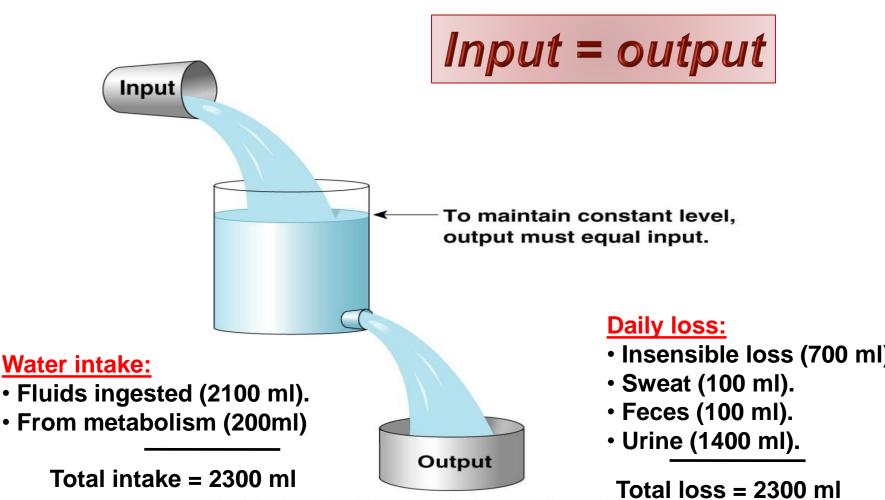


### **Water Balance**

Simplified version!

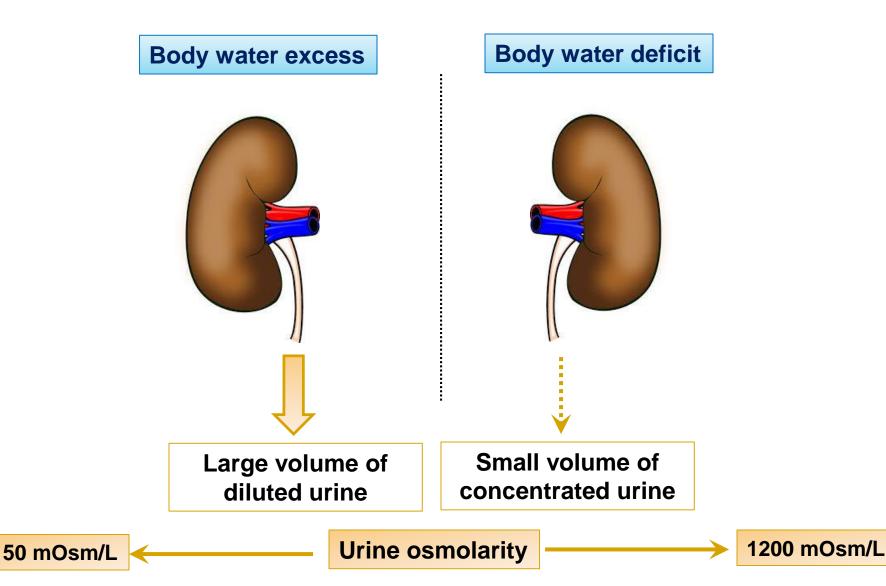


### **Water Balance**

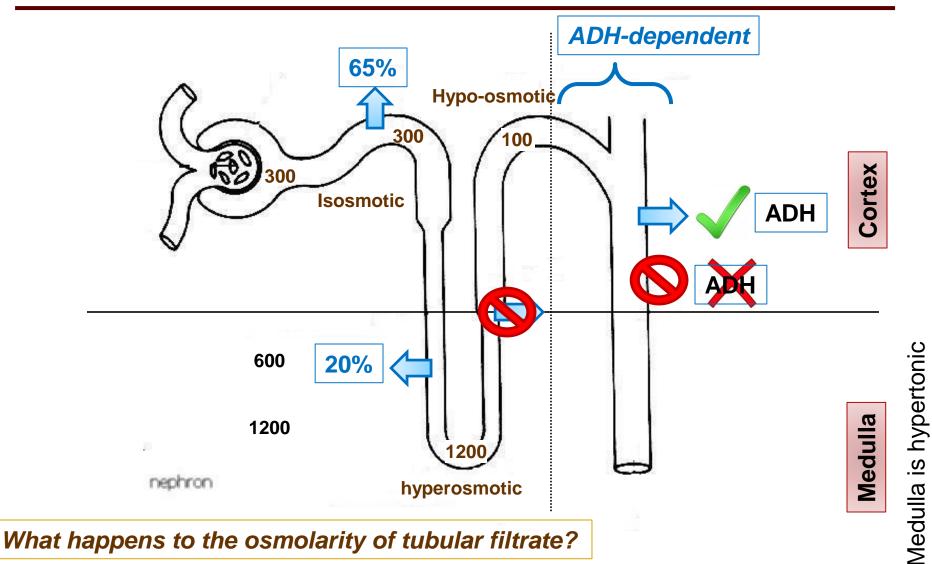


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### **Regulation of H2O by the Kidney**



## H<sub>2</sub>O Handling by the Kidney



## **Obligatory urine volume**

- The minimal volume of urine that must be excreted to rid the body of waste products of metabolism.
- A 70-Kg human needs to excrete 600 mOsm of solutes per day.
- What is the obligatory urine volume?

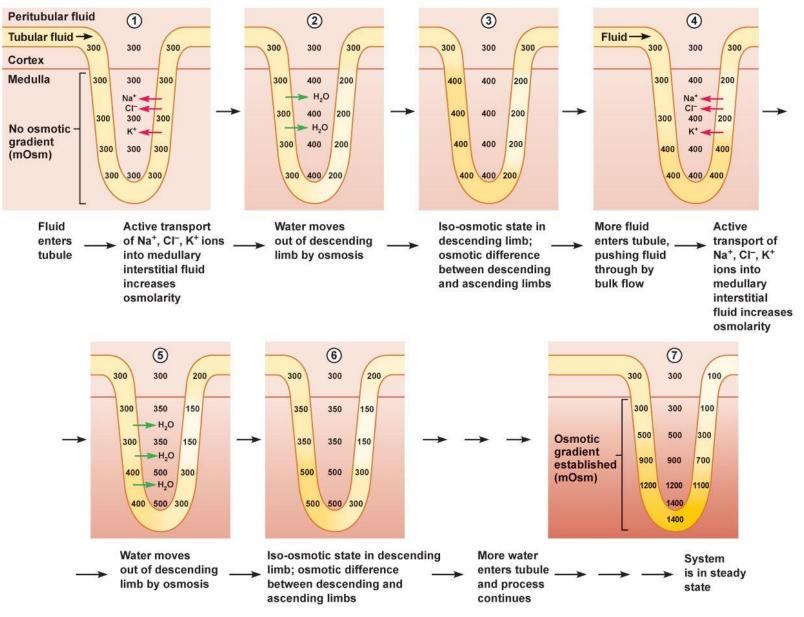
600 mOsm/d 1200 mOsm/L = 0.5 L/day

## **Forming a Concentrated Urine**

### • Requires:

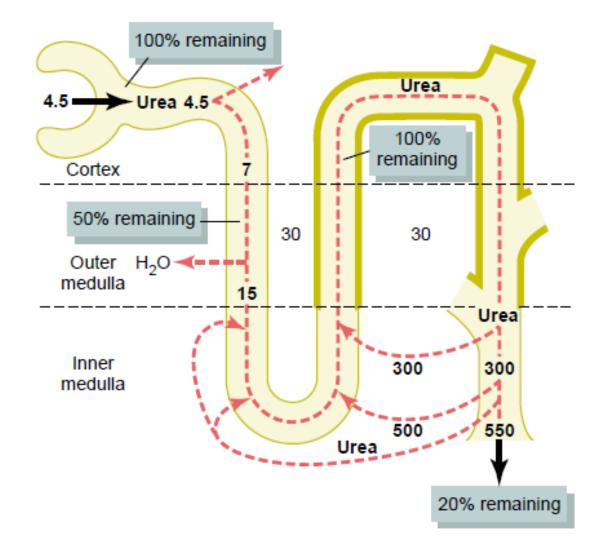
- 1. High levels of ADH.
- 2. Hyperosmotic renal medulla.
  - a. Countercurrent mechanism.
  - b. Urea recirculation.
- IF around the body has an osmolarity of ≈ 300 mOsm/L..
   How did the renal medullary interstitium become hyperosmotic?

## **Countercurrent Multiplier Mechanism**



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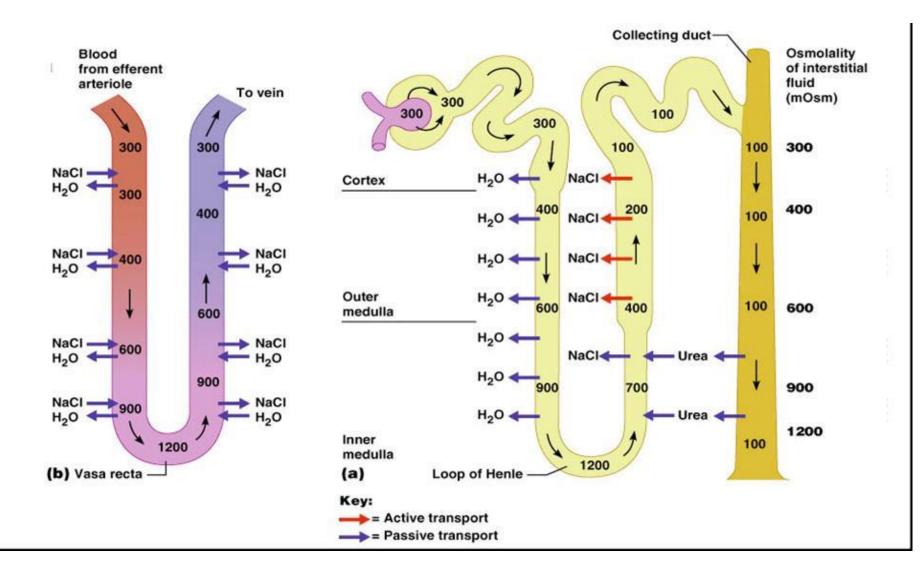
### **Recirculation of Urea**



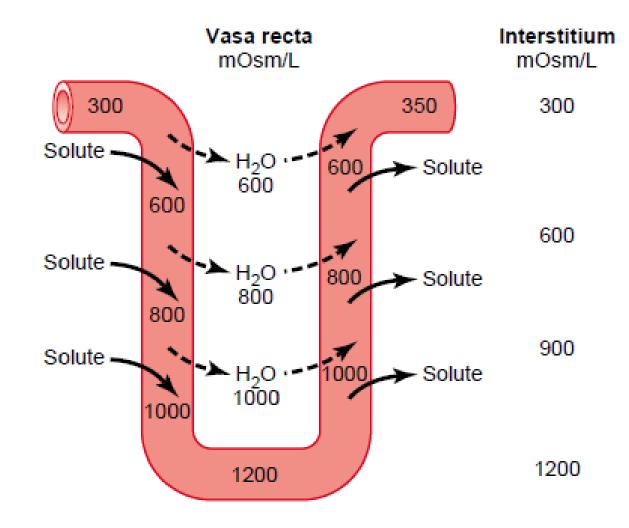
### **The Vasa Recta**

- Why doesn't the blood flowing through the vasa recta into the renal medulla wash out the medullary hyperosmotic gradient?
  - 1. Medullary blood flow is low (<5%) of renal blood flow.
  - 2. The vasa recta serve as countercurrent exchangers.

### The Vasa Recta Countercurrent exchanger

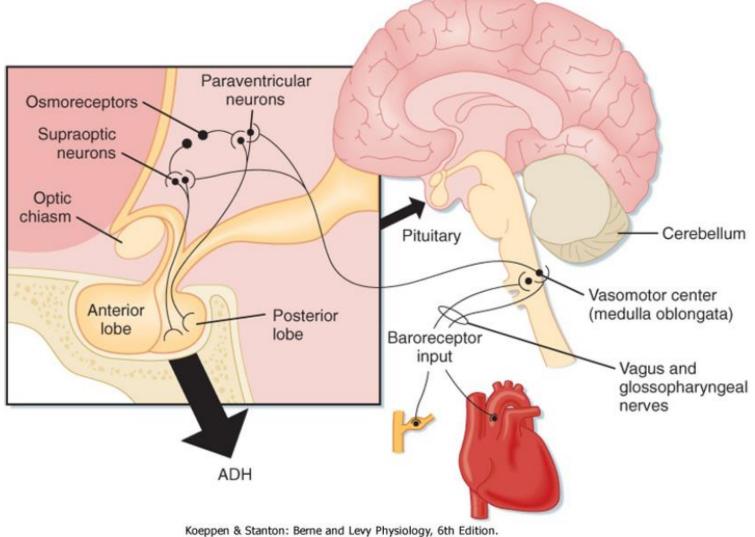


### **The Vasa Recta Countercurrent Exchanger**



# REGULATION OF ECF OSMOLARITY

## **Antidiuretic Hormone (ADH)**



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## **Stimulants for ADH Secretion**

### Osmotic (most important)

- Osmolarity of ECF.
- 1% change in osmolarity can alter ADH secretion significantly.

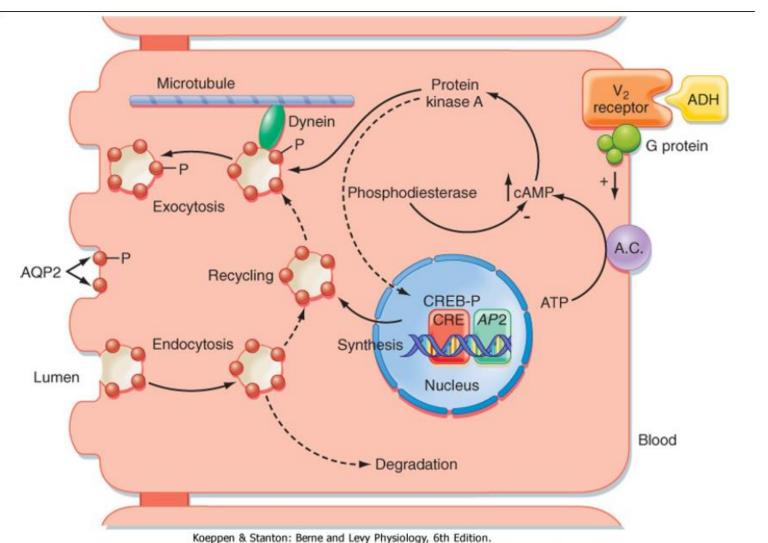
### Hemodynamic

- Volume & pressure in the vascular system.
- 5-10% decrease in BP or BV is required before ADH secretion is stimulated.

### **Factors That Can Alter ADH Secretion**

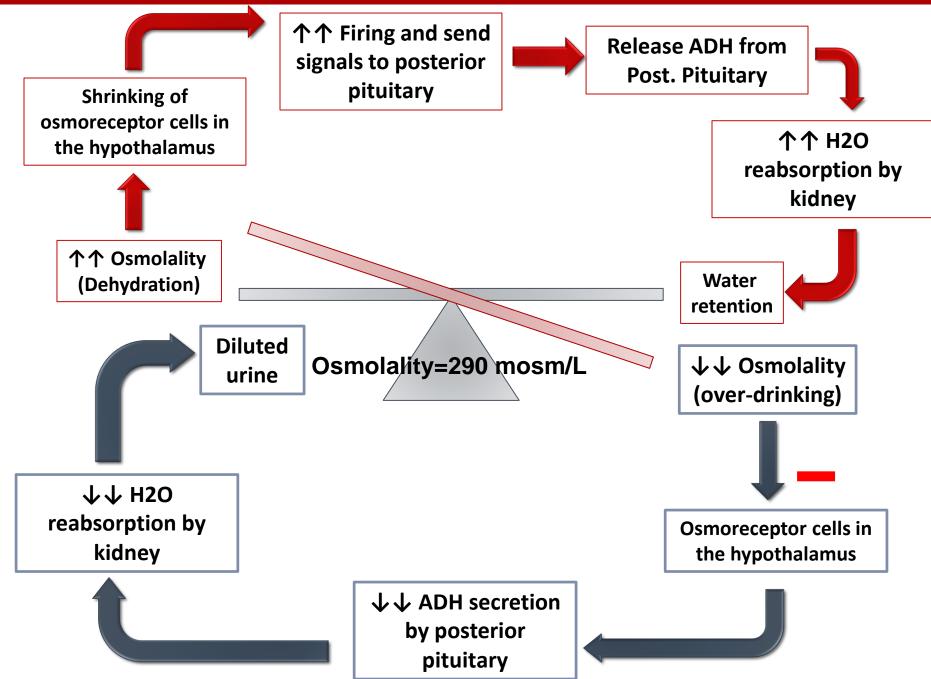
Increase ADH	Decrease ADH
Nausea	ANP
Hypoxia	
Angiotensin-II	
Drugs; Morphine Nicotine	Drugs; Alcohol

### **ADH Mechanism of Action**

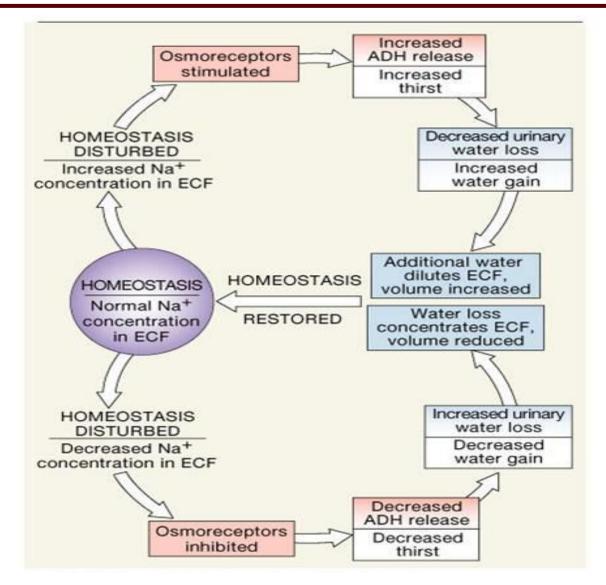


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#### Water Balance



### Feedback Mechanisms Involved in Regulation of Water Balance



### **Control of ADH Secretion & Thirst**

#### Regulation of ADH Secretion

Increase ADH	Decrease ADH
↑ Plasma osmolarity	↓ Plasma osmolarity
↓ Blood volume	↑ Blood volume
↓ Blood pressure	↑ Blood pressure
Nausea Hypoxia	
Drugs:	Drugs:
Morphine	Alcohol
Nicotine	Clonidine (antihypertensive drug)
Cyclophosphamide	Haloperidol (dopamine blocker)

#### **Control of Thirst**

#### Increase Thirst

↑ Osmolarity ↓ Blood volume ↓ Blood pressure ↑ Angiotensin Dryness of mouth

#### Decrease Thirst

drug)

↓ Osmolarity 1 Blood volume ↑ Blood pressure ↓ Angiotensin II Gastric distention

#### **Abnormalities in ADH Secretion Abnormalities in ADH secretion Excessive ADH** No ADH effect "SIADH" "Diabetes insipidus" High ADH levels. Water retention. ECF hypo-osmotic Nephrogenic Central Urine hyperosmotic Mutations in V2 $\downarrow \downarrow$ ADH from receptors or AQP2. posterior pituitary. Cannot respond to Polyuria ADH. Polydipsia Polyuria Polydipsia

### Water Diuresis vs Osmotic Diuresis

Water diuresis	Osmotic diuresis
Increased urine flow rate (No change in urine excretion of solutes	Increase urine flow rate as well as the excretion of solutes
<ul> <li>Causes:</li> <li>Excess ingestion of water</li> <li>Lack of ADH</li> <li>Defect in ADH receptors in Distal segment of nephron (nephrogenic Diabetes Insipidus)</li> </ul>	<ul> <li>Causes:</li> <li>Increase plasma glucose level (DM)</li> <li>Increase level of poorly reabsorbed solutes/ anions</li> <li>Diuretic drugs (Lasix)</li> </ul>
Diuresis is mainly due to decrease in water reabsorption in distal segment of nephron. No change to the water reabsorbed proximally	Diuresis is mainly due to decrease reabsorption of solute in PCT or LOH. Decrease solute reabsorption results in decrease in water reabsorption proximally as well as distally

### Water Diuresis vs Osmotic Diuresis

Water diuresis	Osmotic diuresis
Increase urine volume results from increased excretion of pure water	Increase urine volume results from increased excretion of osmotically active solutes which pulls water with it.
Urine osmolality falls far below plasma osmolality.	Urine osmolality falls but remains above plasma osmolality.
Only about 15% filtered load of water reaching distal segments may remain unabsorbed and excreted in urine (maximum urine volume 20 ml/min)	Due to decreased water reabsorption in all segments of nephron, a much greater fraction of filtered water may be excreted volume more than 20 ml/min
ADH administration will stop diuresis if it is due to lack of ADH. ADH administration will not be effective in Nephrogenic Diabetes Insipidus.	ADH administration will not stop diuresis.

# **THANK YOU**