Kidneys in regulation of homeostasis

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This presentation includes only the most important terms and facts. Its content by itself is not a sufficient source of information required to pass the Physiology exam.

A42. Kidney in regulation of homeostasis

A3. Compartmentalization of body fluids
A4. Differences between intra- and extracellular fluids

B70. Regulation of body fluid volume
B71. Regulation of constant osmotic pressure

B53. Formation and secretion of posterior pituitary hormones
B58. Adrenal cortex. Functions, malfunctions.
B62. Natriuretic peptides

B61. Bone formation and resorption. Regulation of calcaemia.

A30. Homeostasis (acid-base balance)

Homeostasis

= maintainance of stable conditions in the internal body environment

Maintainance of Constant Volume and Composition of Body Fluids

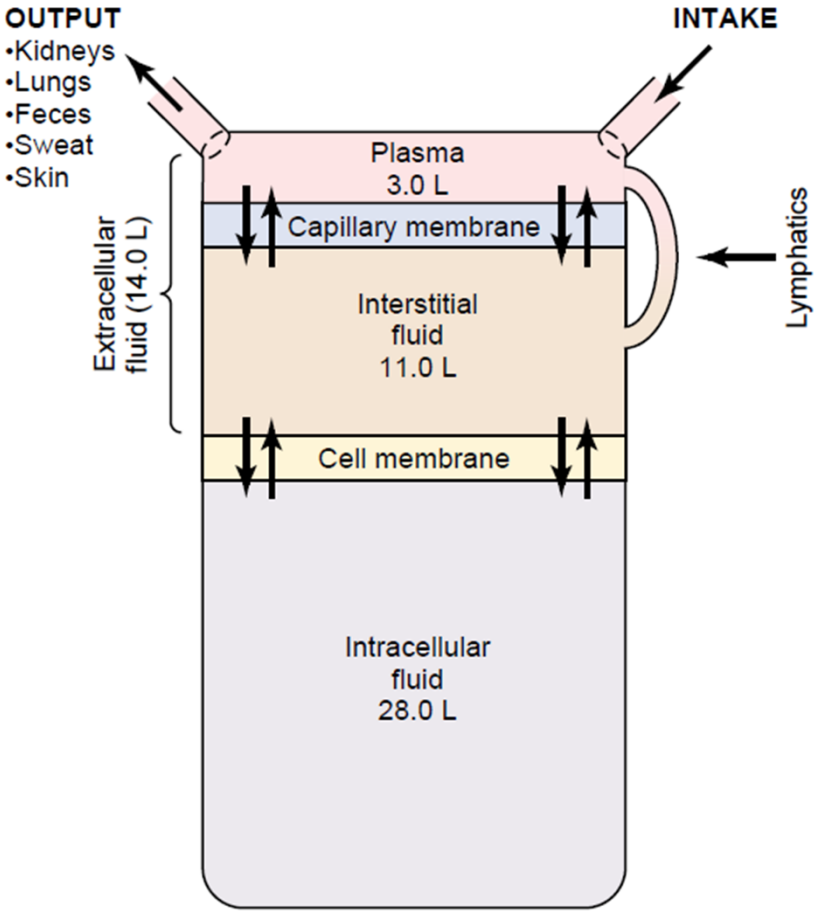
Maintainance of Acid-Base Balance

Constant Volume and Composition of Body Fluids
- Regulation by Kidneys -

Body Fluids – Types and Volumes

Compartments

60 % of the body weight in total



5 % of the body weight

15 % of the body weight

Transcellular fluid (1-2 l) – special type of ECF

40 % of the body weight

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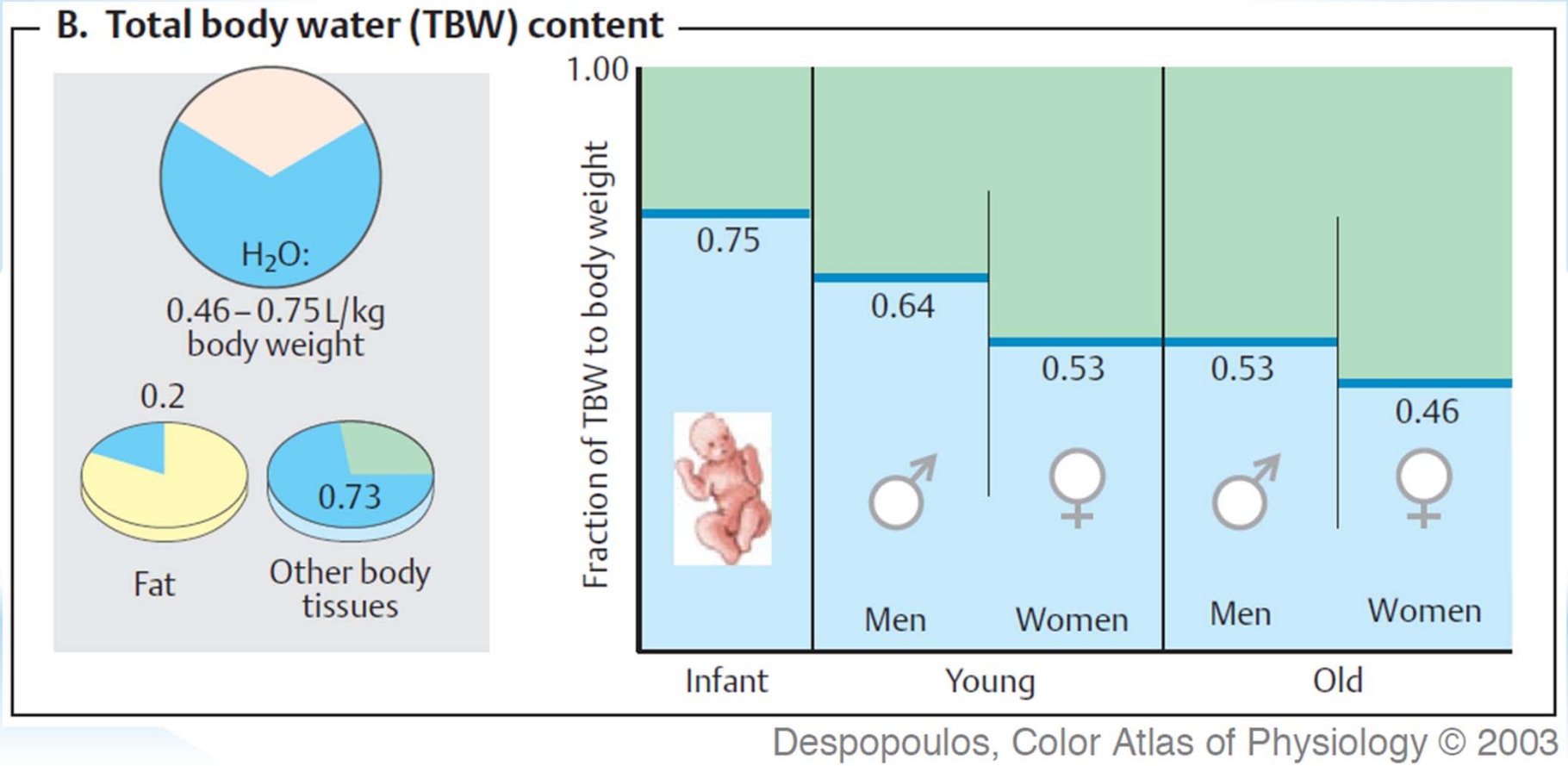
Body Fluids – Types and Volumes

Changes with aging



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Body Fluids – Types and Volumes

Balance between Input and Output of Fluid

Daily Intake and Output of Water (ml/day)

	Normal	Prolonged, Heavy Exercise
Intake		
Fluids ingested	2100	?
From metabolism	200	200
Total intake	2300	?
Output		
Insensible—skin	350	350
Insensible—lungs	350	650
Sweat	100	5000
Feces	100	100
Urine	1400	500
Total output	2300	6600

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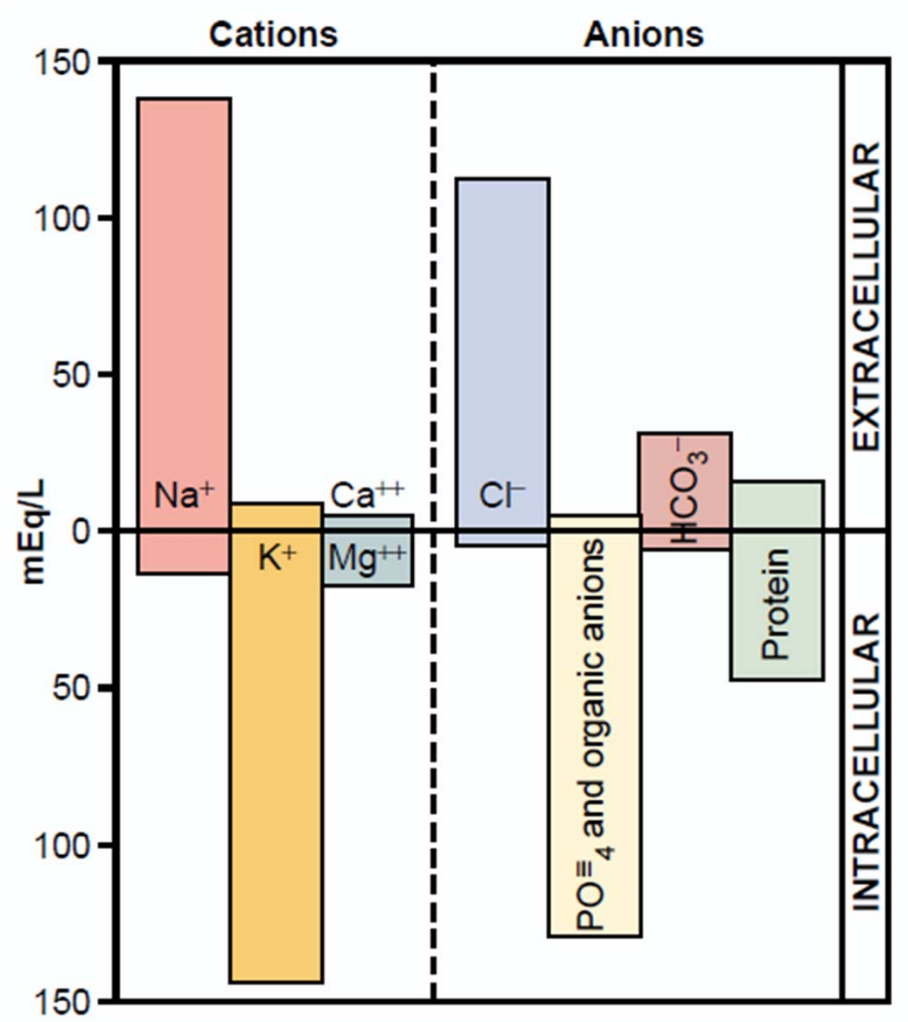
Body Fluids – Composition

ECF vs. ICF



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Body Fluids – Composition

plasma vs. ISF

	Plasma (mOsm/L H_2O)	Interstitial (mOsm/L H_2O)
Na^+	142	139
K^+	4.2	4.0
Ca^{++}	1.3	1.2
Mg^+	0.8	0.7
Cl^-	108	108
HCO_3^-	24	28.3
HPO_4^-, $H_2PO_4^-$	2	2
SO_4^-	0.5	0.5
Phosphocreatine		
Carnosine		
Amino acids	2	2
Creatine	0.2	0.2
Lactate	1.2	1.2
Adenosine triphosphate		
Hexose monophosphate		
Glucose	5.6	5.6
Protein	1.2	0.2
Urea	4	4
Others	4.8	3.9

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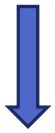
Body Fluids – Composition

osmolality 285 mosm/kg H_2O

↑ ↑ NaCl intake, loss of water → shrinking of cells

↓ ↓ NaCl intake, ↑ water input → cell edema



Precise regulation of osmolality of ESF is necessary!

- **osmoreceptors**
- **kidneys** (target organ for the action of hormones below)
- **antidiuretic hormone**
- **aldosteron**
- **natriuretic peptides**

Humoral Regulation of Body Fluids

Antidiuretic Hormone

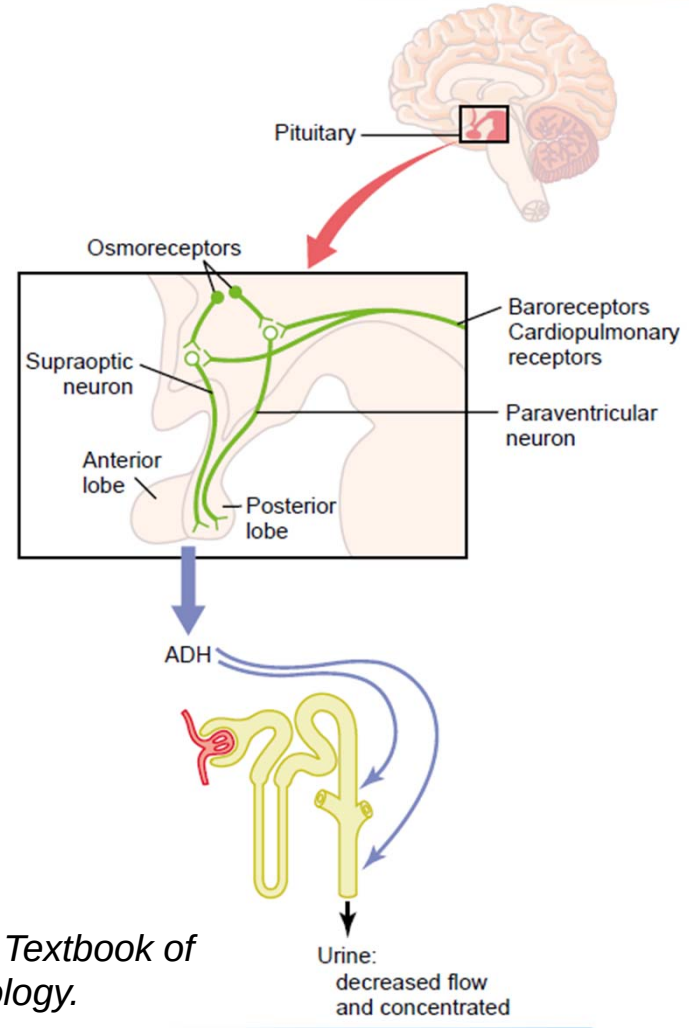
(vasopressin)

- effects:

→ water reabsorption in kidneys (aquaporin 2)

→ control of blood pressure

→ ↑ glycogenolysis, mediator in the brain, ↑ secretion of ACTH in adenohypophysis



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Humoral Regulation of Body Fluids

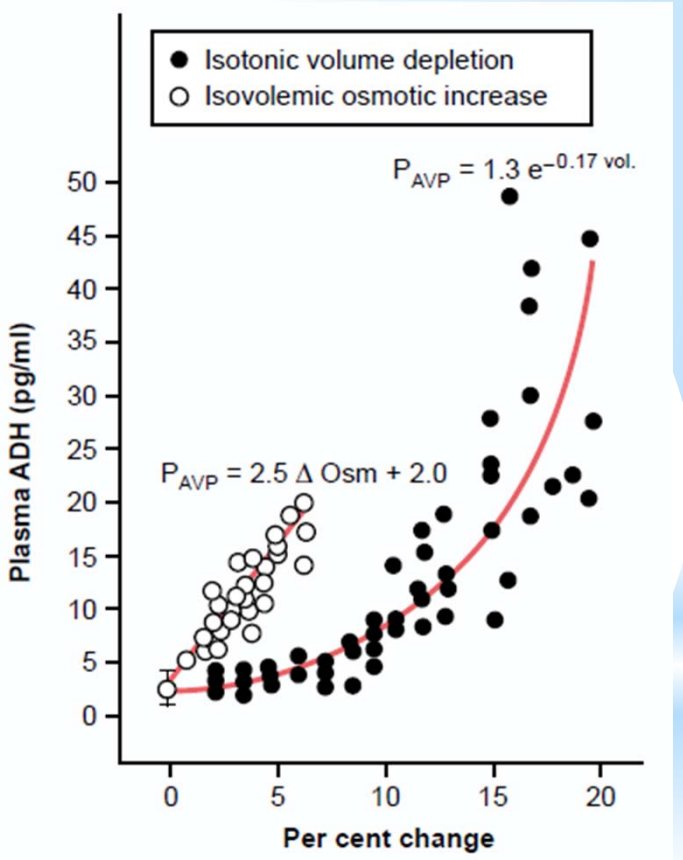
Antidiuretic Hormone

(vasopressin)

- regulation of secretion:

↑
- ↑ osmolality
- ↓ volume of ECF
- pain, emotions, stress (surgical), physical exertion; standing
- nausea, vomitting
- angiotensine II
- morphin, nicotine, barbiturates, …

↓
- ↓ osmolality, ↑ volume of ECF
- alcohol; antagonists of opioids



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Humoral Regulation of Body Fluids

Antidiuretic Hormone

(vasopressin)

- pathology:

↑ SIADH

↓ *diabetes insipidus*

Humoral Regulation of Body Fluids

Aldosteron

- the most important steroid with the mineralocorticoid effect

- mechanism of action:

  binding to the mineralocorticoid receptor

  → synthesis of proteins:

  - namely Na^+/K^+-ATPase
  - ↑ number of amiloride-inhibited Na^+-channels
  - ↑ activity of H^+-pump
  - ↑ activity of Na^+/H^+-antiport

Humoral Regulation of Body Fluids

Aldosteron

- the most important steroid with the mineralocorticoid effect

- effects:

→↑ Na^+ reabsorption (urine, sweat, saliva, gastric juice)
→↑ K^+ urine excretion, ↑ acidity of urine (exchange for Na^+)
→↑ K^+ content and ↓ Na^+ content in muscle and brain cells

Humoral Regulation of Body Fluids

Aldosteron

- the most important steroid with the mineralocorticoid effect

- regulation of its secretion:
  - ACTH (transient effect)
  - direct stimulatory effect of ↑ plasmatic concentration of K^+ and ↓ Na^+ (lower sensitivity)
  - renin-angiotensine-aldosteron system
  - atrial natriuretic peptide
  - other hormones od adenohypophysis (maintenance of reactivity of *zona glomerulosa*)

Humoral Regulation of Body Fluids

Aldosteron - Pathology

Primary hyperaldosteronism (Conn´s syndrome)

- tumors of adrenal cortex which secretes aldosteron

Secondary hyperaldosteronism

- patients with the congestive heart failure, nephrosis, liver cirhosis, renal artery constriction, hypertension, with the salt-losing form of adrenogenital syndrome

Hyporeninemic hypoaldosteronism

Pseudohypoaldosteronism

Humoral Regulation of Body Fluids

Atrial Natriuretic Peptide

- one of natriuretic peptides (BNP – cardiac ventricles, CNP – brain)
- secreted by atrial cardiomyocytes, found also in the brain
- receptors
- short half-life

Humoral Regulation of Body Fluids

Atrial Natriuretic Peptide

- one of natriuretic peptides (BNP – cardiac ventricles, CNP – brain)

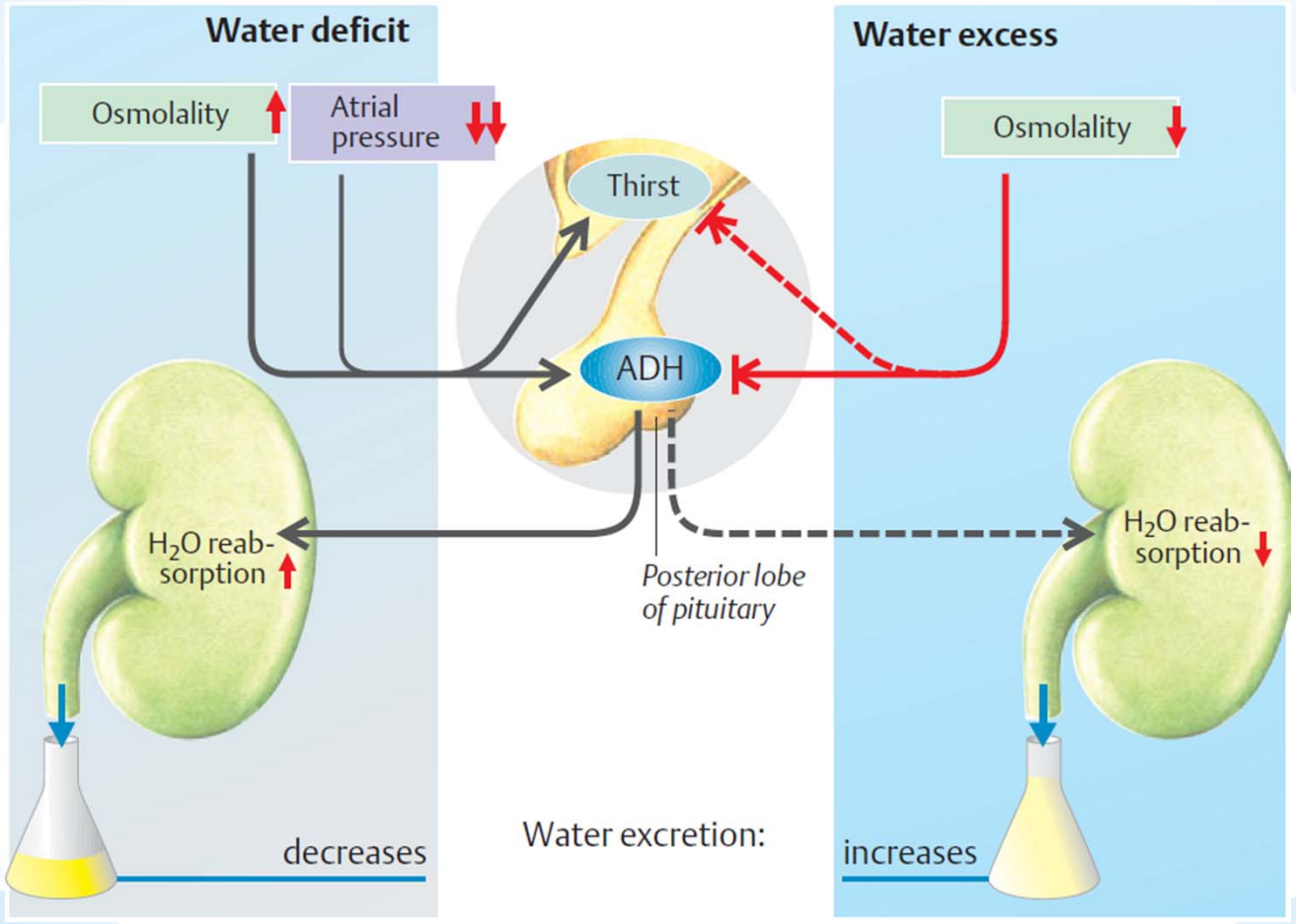
- effects (through ↑ cGMP): → ↓ **BP** (also through the brain stem)
  - → natriuresis
  - → ↓ reactivity of vascular smooth muscles for vasocontrictive substances
  - → inhibition of renin secretion, ↓ reactivity of *zona glomerulosa* for stimuli ↑ aldosteron secretion
  - → inhibition of ADH secretion → ↑ water excretion

- regulation of its secretion:
  - ↑ - ↑ ECF volume
  - ↓ - ↓ CVP at orthostasis

Humoral Regulation of Body Fluids

Water Homeostasis

water intoxication



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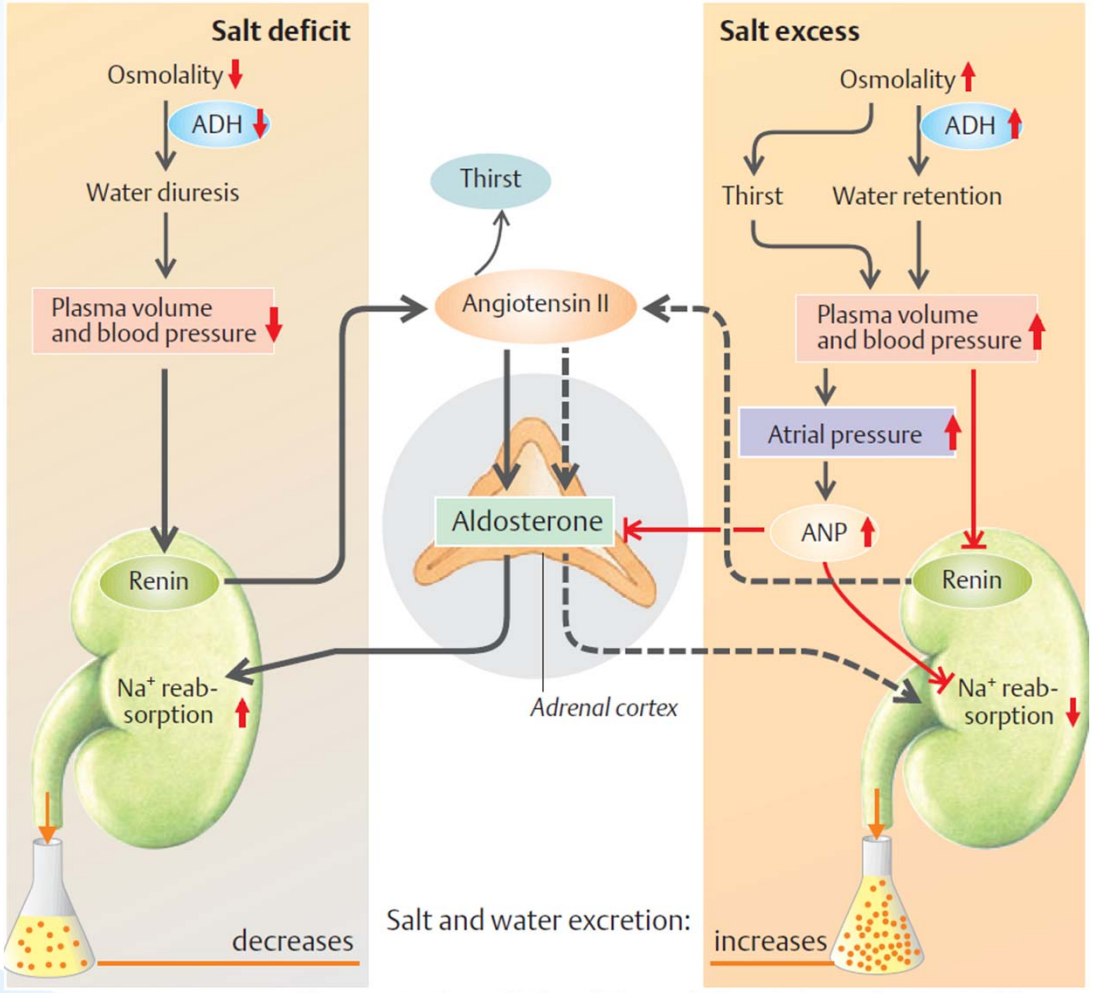
Humoral Regulation of Body Fluids

Salt Homeostasis



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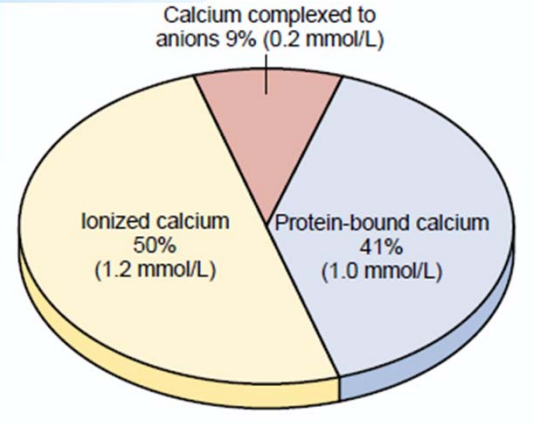
Humoral Regulation of Body Fluids

Calcium in the Body



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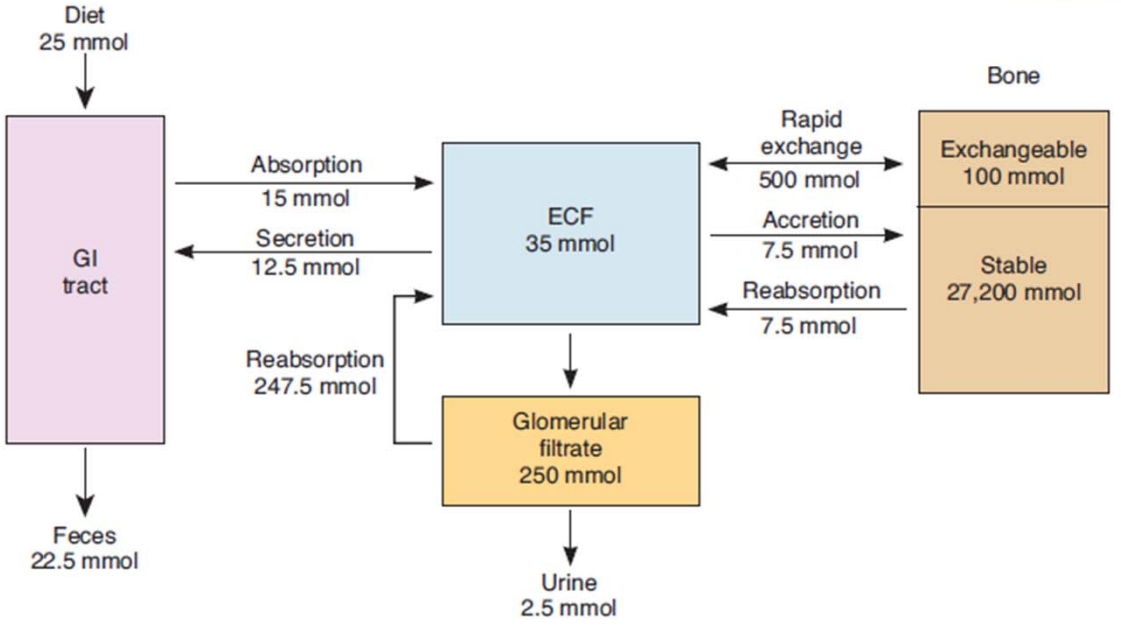


Ganong´s Review of Medical Physiology

hypocalcemia

hypercalcemia

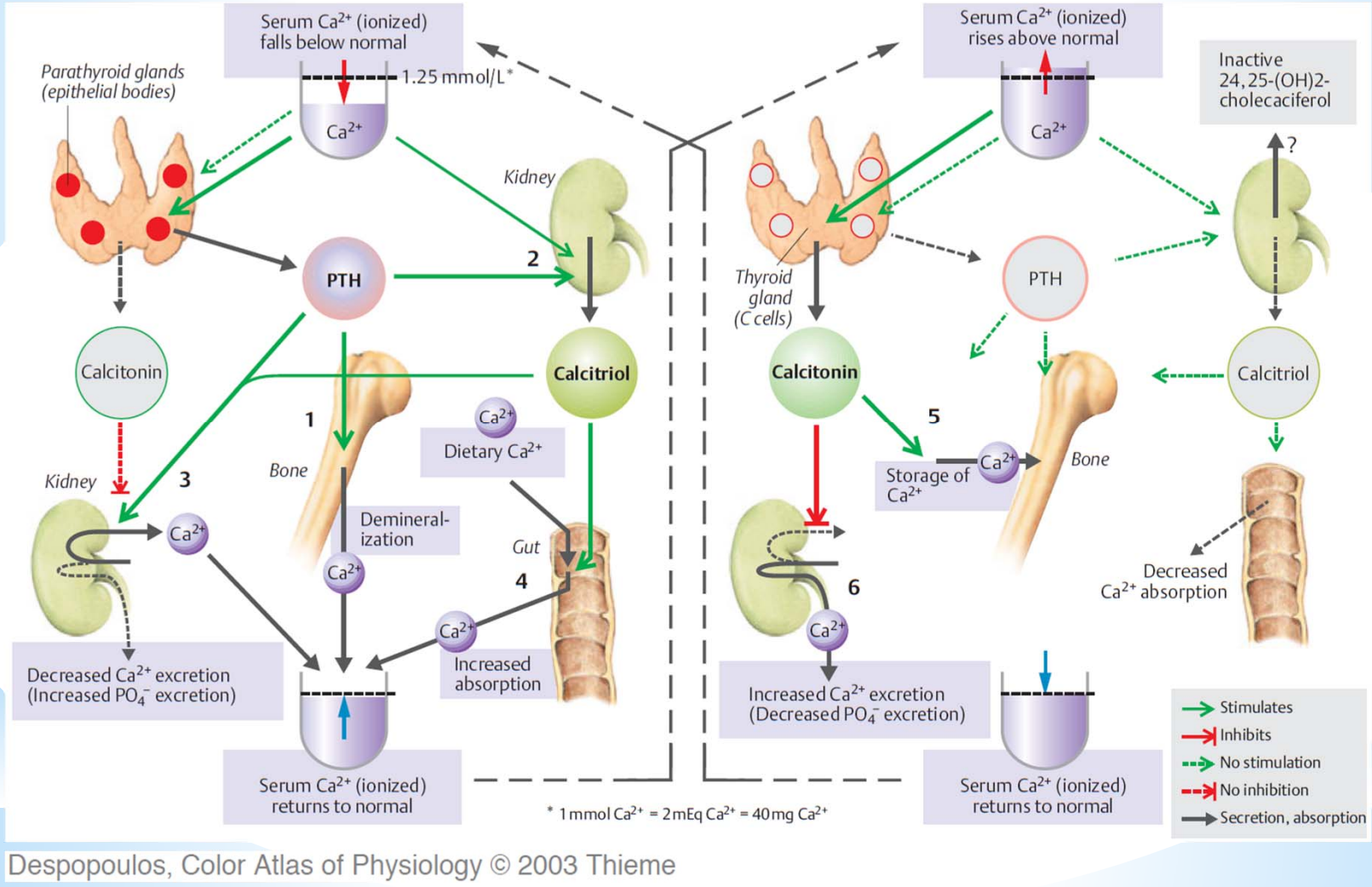
Humoral Regulation of Body Fluids

Hormonal Regulation of Calcemia

Parathormone
Vitamin D
Calcitonin

Humoral Regulation of Body Fluids

Hormonal Regulation of Calcemia



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Acid-Base Balance
- Regulation by Kidneys -

Acid-Base Balance and its Regulation

Acid-base balance is regulated by:

1) Buffers
   - fast regulation (seconds)
   - pH changes attenuated by binding and release of H^+:

     buffer + H^+ $\rightleftharpoons$ H - buffer

     $\uparrow[H^+]$ direction to the right favoured till free buffer is available

     $\downarrow[H^+]$ direction to the left favoured, H^+ released

2) Lungs
   - fast regulation (minutes even hours)
   - elimination of CO_2 from the body ($H_2CO_3 \rightarrow H_2O + CO_2$)

3) Kidneys
   - slower regulation (hours even days) but the most powerful
   - elimination of acids and bases from the body

Acid-Base Balance and its Regulation

Regulation of Acid-Base Balance by Kidneys

- by excretion of acid or alkalic urine
- a high amount of HCO_3^- still filtered in the glomerulus

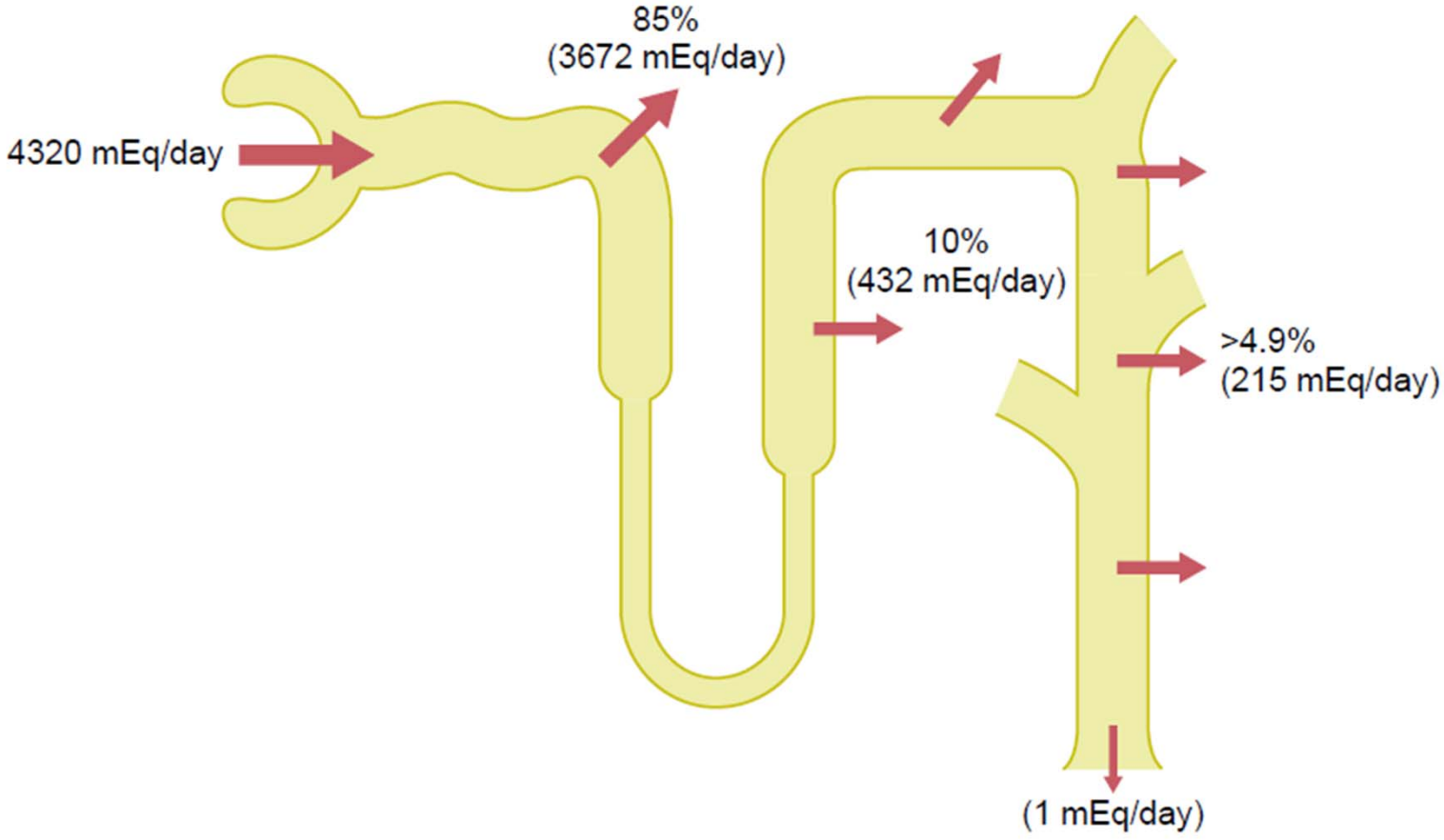
  GFR 180 l/day, $[HCO_3^-]_{plasma}$ 24 mEq/l $\rightarrow$ 4320 mEq HCO_3^- filtered per day – almost all ordinarily reabsorbed
- a high amount of H^+ still secreted in renal tubules

  about 80 mEq of non-volatile acids are formed in the course of metabolic processes per day – have to be excreted by kidneys
- filtered HCO_3^- / secreted H^+

Acid-Base Balance and its Regulation

Regulation of Acid-Base Balance by Kidneys

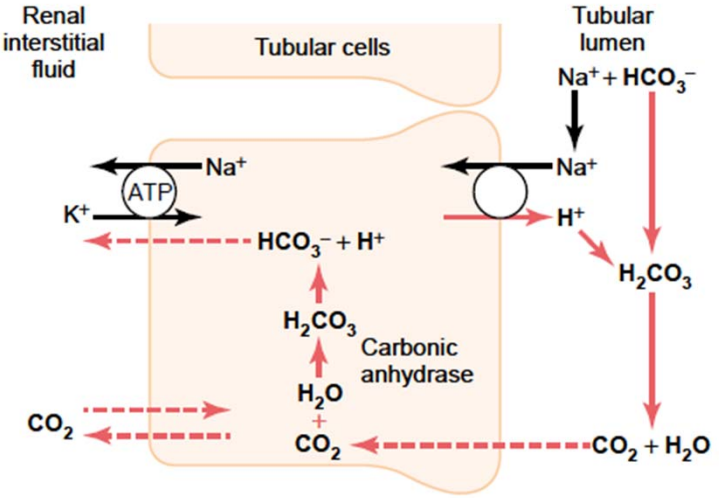
1) Secretion of H^+
2) Reabsorption of HCO_3^-



Acid-Base Balance and its Regulation

Regulation of Acid-Base Balance by Kidneys

1) Secretion of H^+
2) Reabsorption of HCO_3^-
   - ❖ in the proximal tubule, thick loop of Henle and at the beginning of the distal tubule



Na^+/H^+-antiport

>90% HCO_3^- reabsorbed – only a slight acidification of the urine!

Reabsorption of HCO_3^- across the basolateral membrane facilitated by:

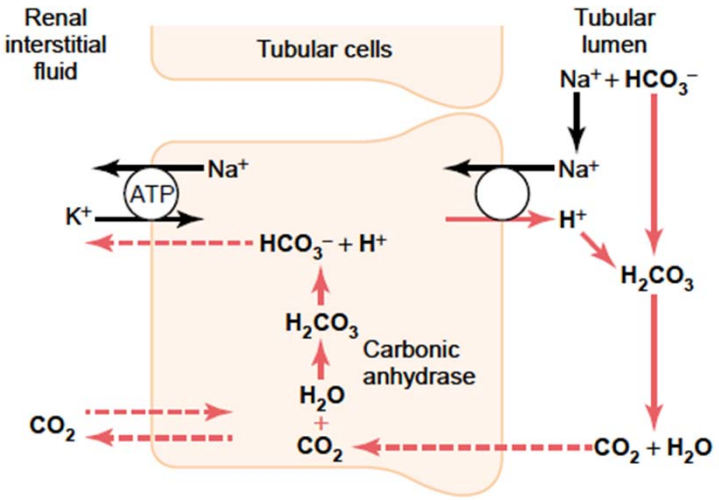
- Na^+-HCO_3^- co-transport
  (the proximal tubule)
- Cl^--HCO_3^- exchanger
  (the end of proximal tubule and the following parts of tubulus except for the thin loop of Henle)

Acid-Base Balance and its Regulation

Regulation of Acid-Base Balance by Kidneys

1) Secretion of H^+
2) Reabsorption of HCO_3^-

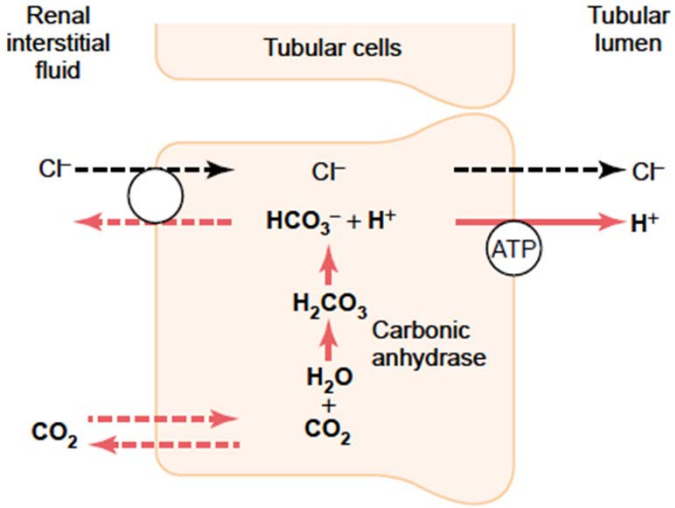
- in the proximal tubule, thick loop of Henle and at the beginning of the distal tubule



Na^+/H^+-antiport

>90% HCO_3^- reabsorbed – only a slight acidification of the urine!

- in the final part of distal tubule and in the collecting duct



primary active transport of H^+ (intercalated cells)

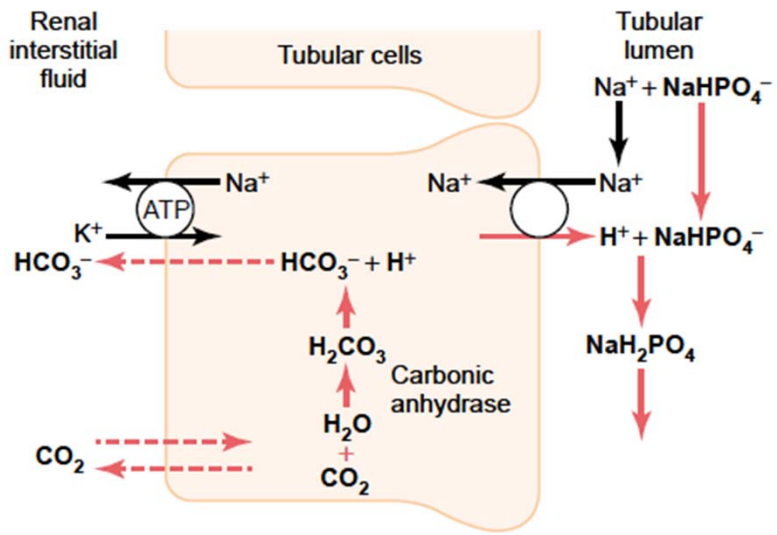
acidification of urine

Acid-Base Balance and its Regulation

Regulation of Acid-Base Balance by Kidneys

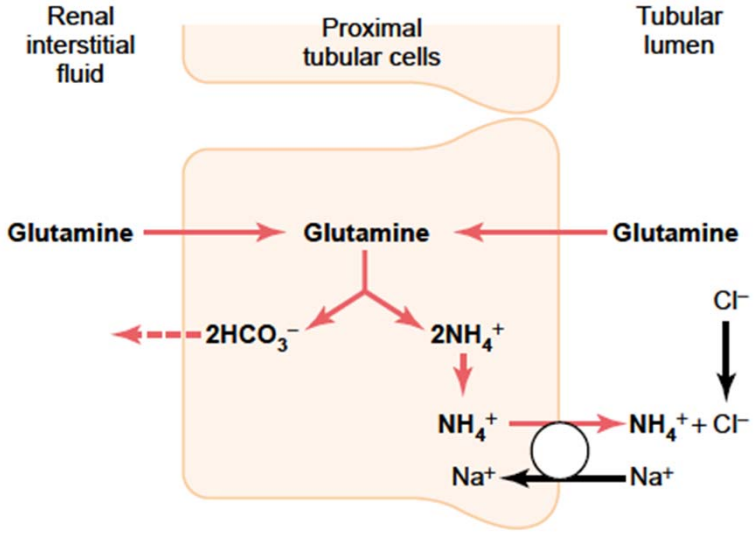
1) Secretion of H^+
2) Reabsorption of HCO_3^-
3) Production of HCO_3^- *de novo*

❖ Phosphate buffer (HPO_4^{2-}, $H_2PO_4^-$)



HPO_4^{2-} and $H_2PO_4^-$ are reabsorbed less than water ⇒ their concentration in the tubular fluid gradually rises

❖ Ammonium buffer (NH_3, NH_4^+)



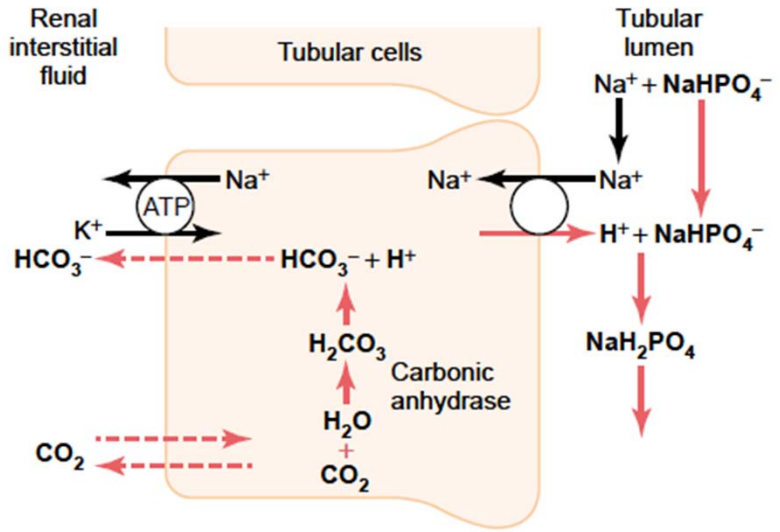
NH_4^+ originates from glutamine – the proximal tubule, thick ascending loop of Henle and distal tubule

Acid-Base Balance and its Regulation

Regulation of Acid-Base Balance by Kidneys

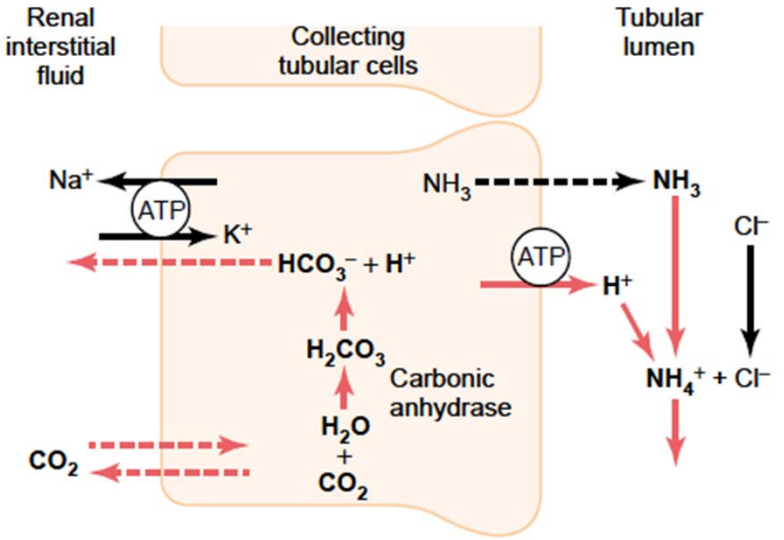
1) Secretion of H^+
2) Reabsorption of HCO_3^-
3) Produkce nového HCO_3^-

❖ Phosphate buffer (HPO_4^{2-}, $H_2PO_4^-$)



HPO_4^{2-} and $H_2PO_4^-$ are reabsorbed less than water ⇒ their concentration in the tubular fluid gradually rises

❖ Ammonium buffer (NH_3, NH_4^+)



the collecting duct (permeable for NH_3 but far less for NH_4^+ - excreted by urine)

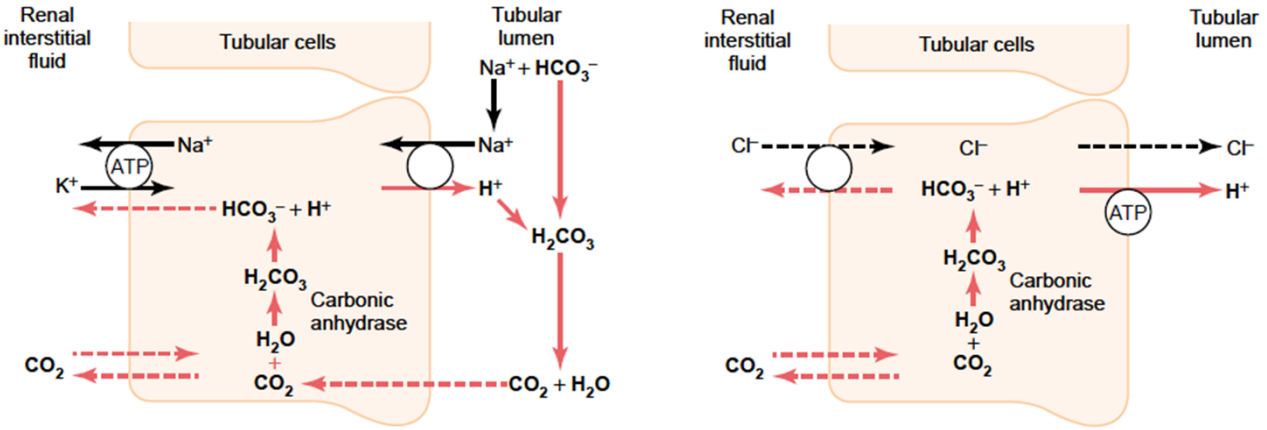
50% of H^+ secretion and HCO_3^- formed *de novo*!

Acid-Base Balance and its Regulation

Regulation of Acid-Base Balance by Kidneys

Regulation of H^+ secretion

↑ - ↑ **pCO_2 in ECF** (respiratory acidosis; direct stimulation due to ↑ formation of H^+ in tubular cells)



- ↓ **pH in ECF** (respiratory or metabolic acidosis)
- ↑ **secretion of aldosteron** (stimulates H^+ secretion in intercalated cells of collecting ducts; Conn´s syndrome - alkalosis)

Acid-Base Balance and its Regulation

Regulation of Acid-Base Balance by Kidneys

Regulation of H^+ secretion

Factors That Increase or Decrease H^+ Secretion and HCO_3^- Reabsorption by the Renal Tubules

Increase H^+ Secretion and HCO_3^- Reabsorption	Decrease H^+ Secretion and HCO_3^- Reabsorption
↑ P_{CO_2}	↓ P_{CO_2}
↑ H^+, ↓ HCO_3^-	↓ H^+, ↑ HCO_3^-
↓ Extracellular fluid volume	↑ Extracellular fluid volume
↑ Angiotensin II	↓ Angiotensin II
↑ Aldosterone	↓ Aldosterone
Hypokalemia	Hyperkalemia

RAS

↑ activity of Na^+/H^+ antiport

↑ activity of H^+ ATPase

↓

tendency to alkalosis

Acid-Base Balance and its Regulation

Regulation of Acid-Base Balance by Kidneys

Acidosis – correction by kidneys

$$\downarrow \; pH = 6.1 + \log \frac{HCO_3^-}{0.03 \times P_{CO2}} \; \downarrow$$

- metabolic acidosis: due to $\downarrow HCO_3^-$
  renal correction : $\downarrow HCO_3^-$ in ECF $\rightarrow \downarrow$ filtered $HCO_3^- \rightarrow$ complete reabsorption of HCO_3^- + its formation *de novo* (HCO_3^- not excreted) + $\uparrow H^+$ excretion $\rightarrow$ pH normalization
- respiratory acidosis: due to $\uparrow P_{CO2}$ (hypoventilation)
  renal correction: $\uparrow P_{CO2}$ in ECF $\rightarrow \uparrow P_{CO2}$ in tubular cells $\rightarrow \uparrow$ formation of H^+ and HCO_3^- in tubular cells $\rightarrow \uparrow H^+$ secretion + $\uparrow HCO_3^-$ reabsorption $\rightarrow$ pH normalization

Acid-Base Balance and its Regulation

Regulation of Acid-Base Balance by Kidneys

Alkalosis – correction by kidneys

$$\uparrow \; pH = 6.1 + \log \frac{HCO_3^-}{0.03 \times P_{CO2}} \; \uparrow$$

- metabolic alkalosis: due to $\uparrow HCO_3^-$

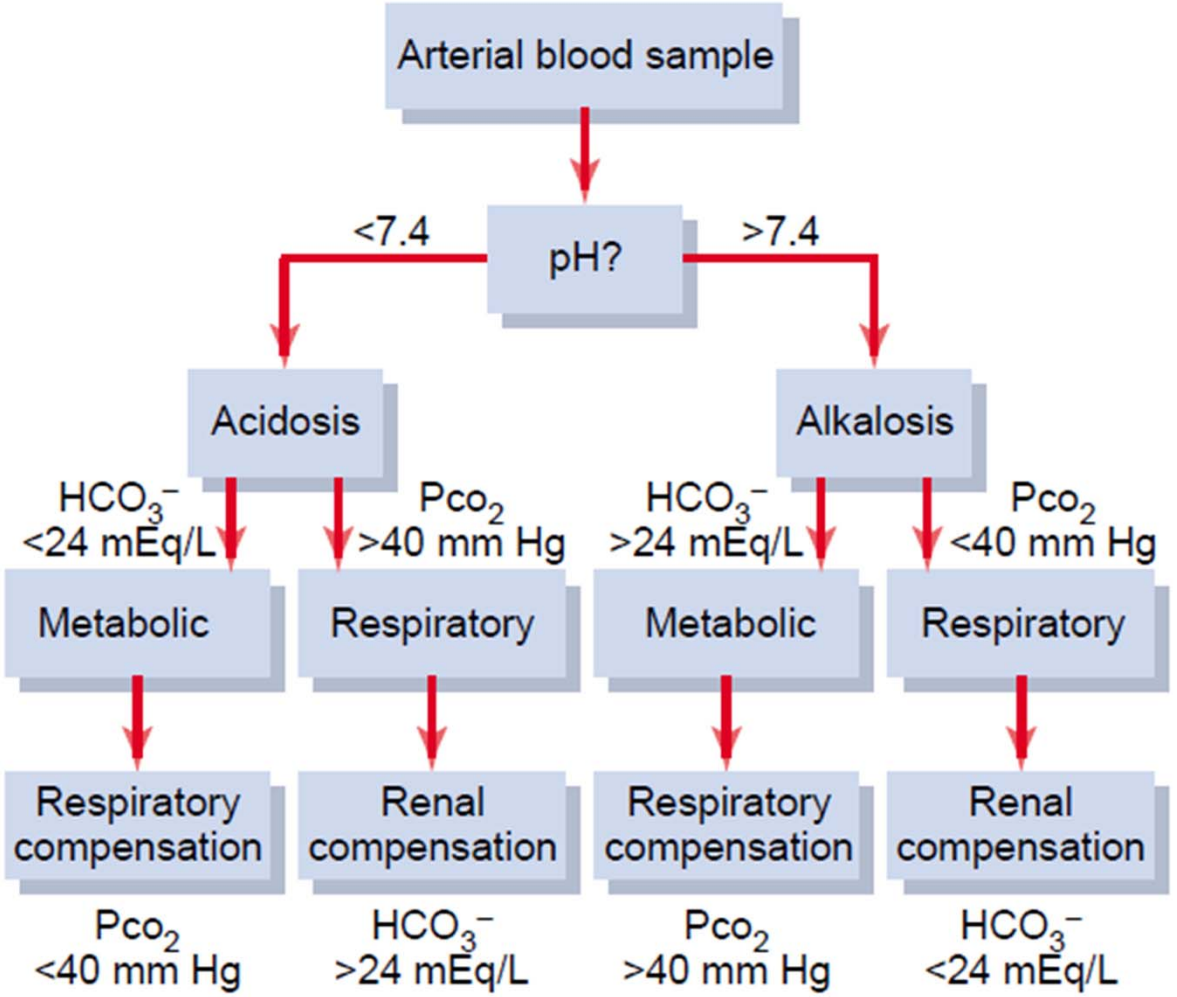
  renal correction: $\uparrow HCO_3^-$ in ECF $\rightarrow \uparrow$ filtered $HCO_3^- \rightarrow$ incomplete HCO_3^- reabsorption (lack of H+) $\rightarrow \uparrow HCO_3^-$ excretion by urine $\rightarrow$ pH normalization

- respiratory alkalosis : due to $\downarrow P_{CO2}$ (hyperventilation)

  renal correction: $\downarrow P_{CO2}$ in ECF $\rightarrow \downarrow P_{CO2}$ in tubular cells $\rightarrow \downarrow$ formation of H^+ and HCO_3^- in tubular cells $\rightarrow \downarrow H^+$ secretion + $\downarrow HCO_3^-$ reabsorption $\rightarrow$ pH normalization

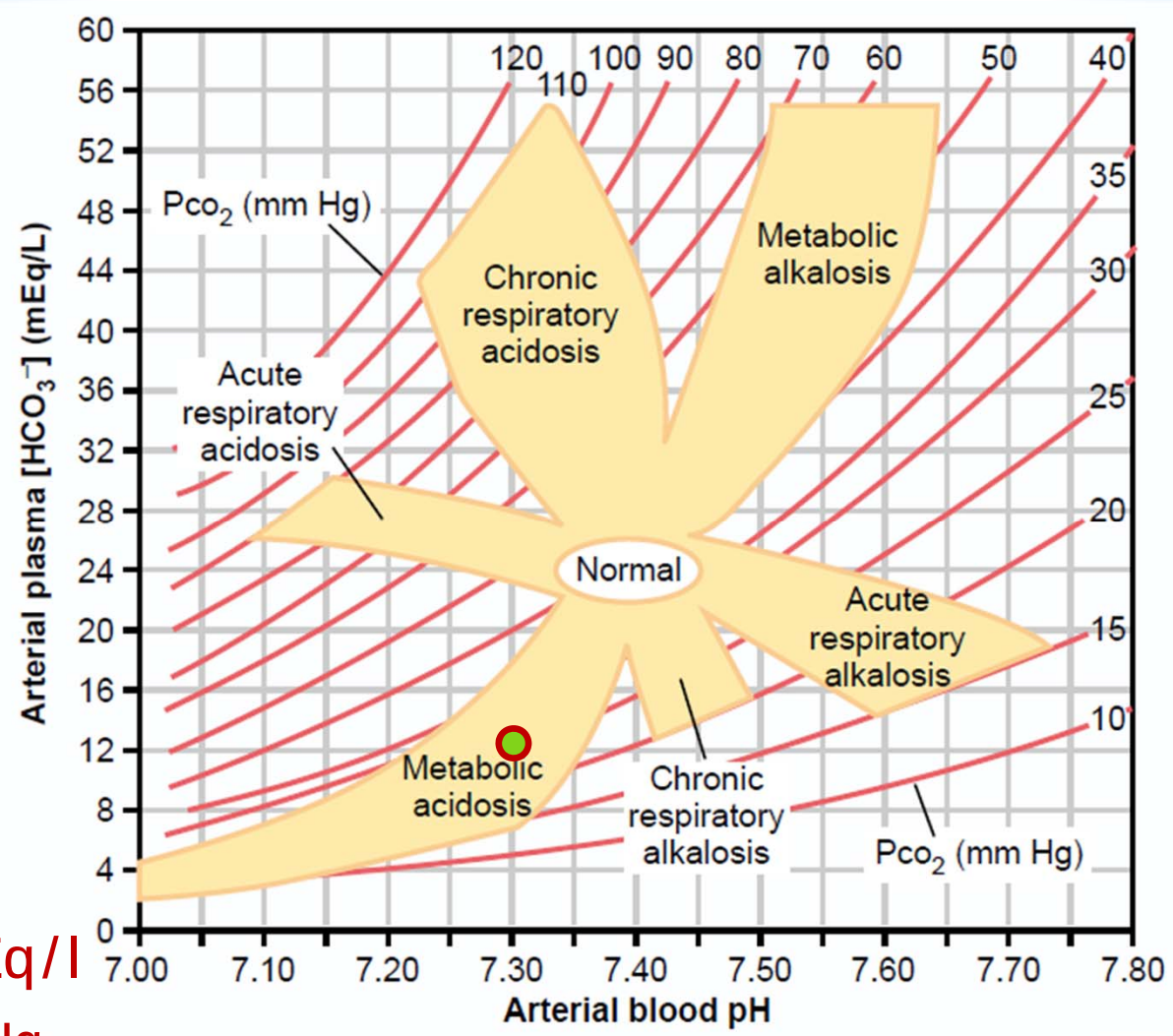
Acid-Base Balance and its Regulation

Diagnostics



Acid-Base Balance and its Regulation

Diagnostics



pH: 7,3

HCO_3^-: 12 mEq/l

P_{CO2}: 25 mmHg

Acid-Base Balance and its Regulation

Diagnostics - Siggaard–Andersen nomogram