

Acid-base disorders

The background of the slide is a light beige color. In the bottom right corner, there are several overlapping, wavy, light-colored lines that create a decorative, abstract pattern.

A bicarbonate of 2.3 mmol/l

- 78 yrs old male
 - PMHx: diabetes mellitus, hypertension
 - admission in very poor condition: desoriented, Kussmaul-breeding, BP 88/52 mmHg, oligo-anuric
- Serum Na 139 mmol/l, K 5,8 mmol/l, Cl 99 mmol/l
BUN 39 mmol/l, creatinine 504 umol/l,
bicarb 2,3 mmol/l
- **What should we do?**

A bicarbonate of 2.3 mmol/l

- Blood-gas analysis:

pH 6,97

HCO_3^- 2,3 mmol/l

pCO₂ 10,2 mmHg

- **What kind of acid-base disorder is this?**

A bicarbonate of 2.3 mmol/l

- Decrease in bicarbonate

$$25 - 2,3 = 22,7 \text{ mmol/l}$$

- Respiratory compensation: delta pCO₂

$$40 - p\text{CO}_2 = 40 - 10,2 = 29,8 \text{ mmHg}$$

- Anion gap

$$\text{Na}^+ - (\text{Cl}^- + \text{HCO}_3^-) = 139 - (99 + 2,3) = 37,7 \text{ mmol/l}$$

A bicarbonate of 2.3 mmol/l

- The patient has:
 - high anion gap metabolic acidosis
- His lactate level:
 - 12,2 mmol/l
- He was taking metformin, which caused lactic acidosis
- He was dialysed, survived, but remained dialysis dependent

Types of anion gap acidosis

Gain of acids

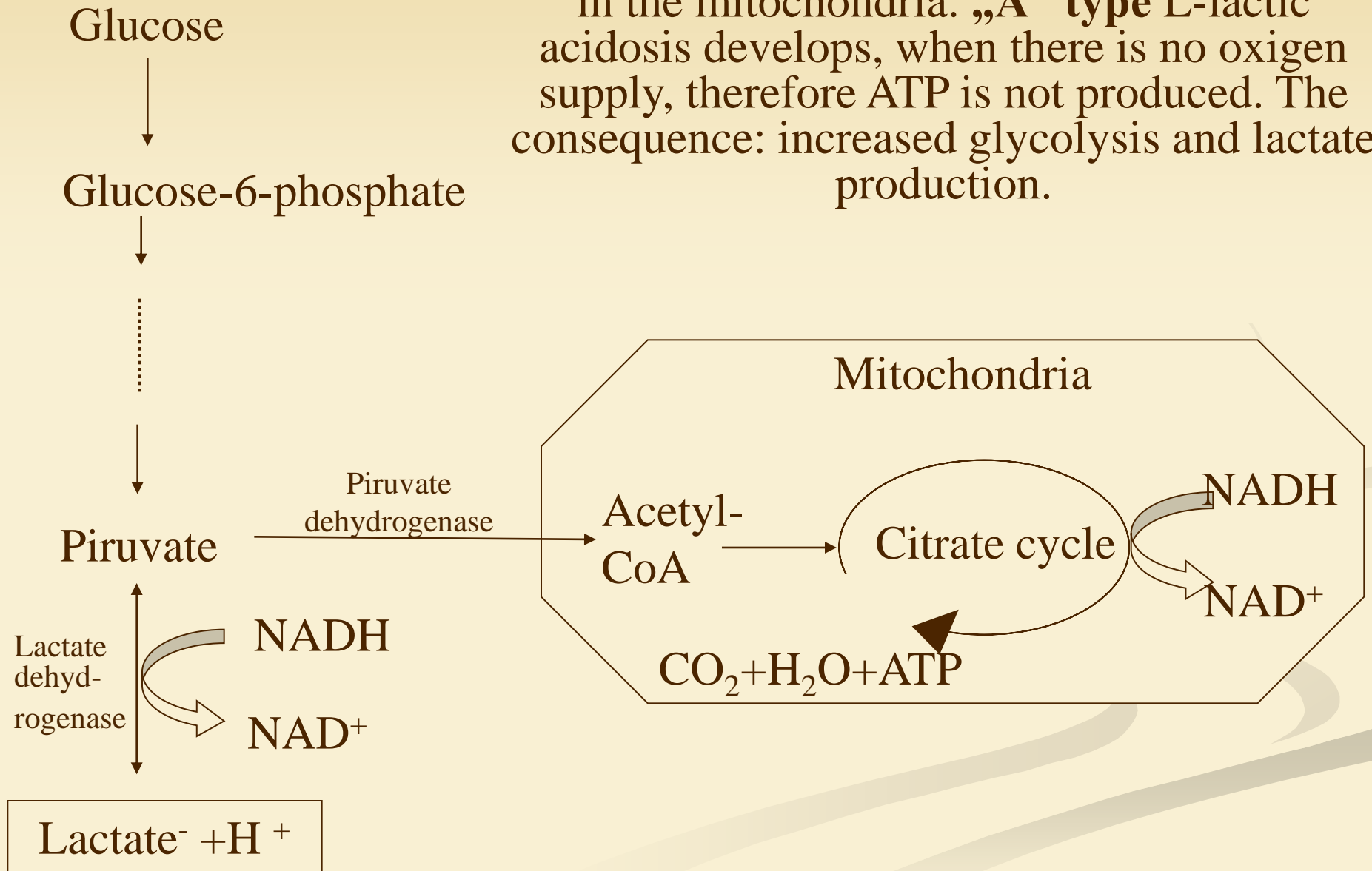
Endogenous acid production

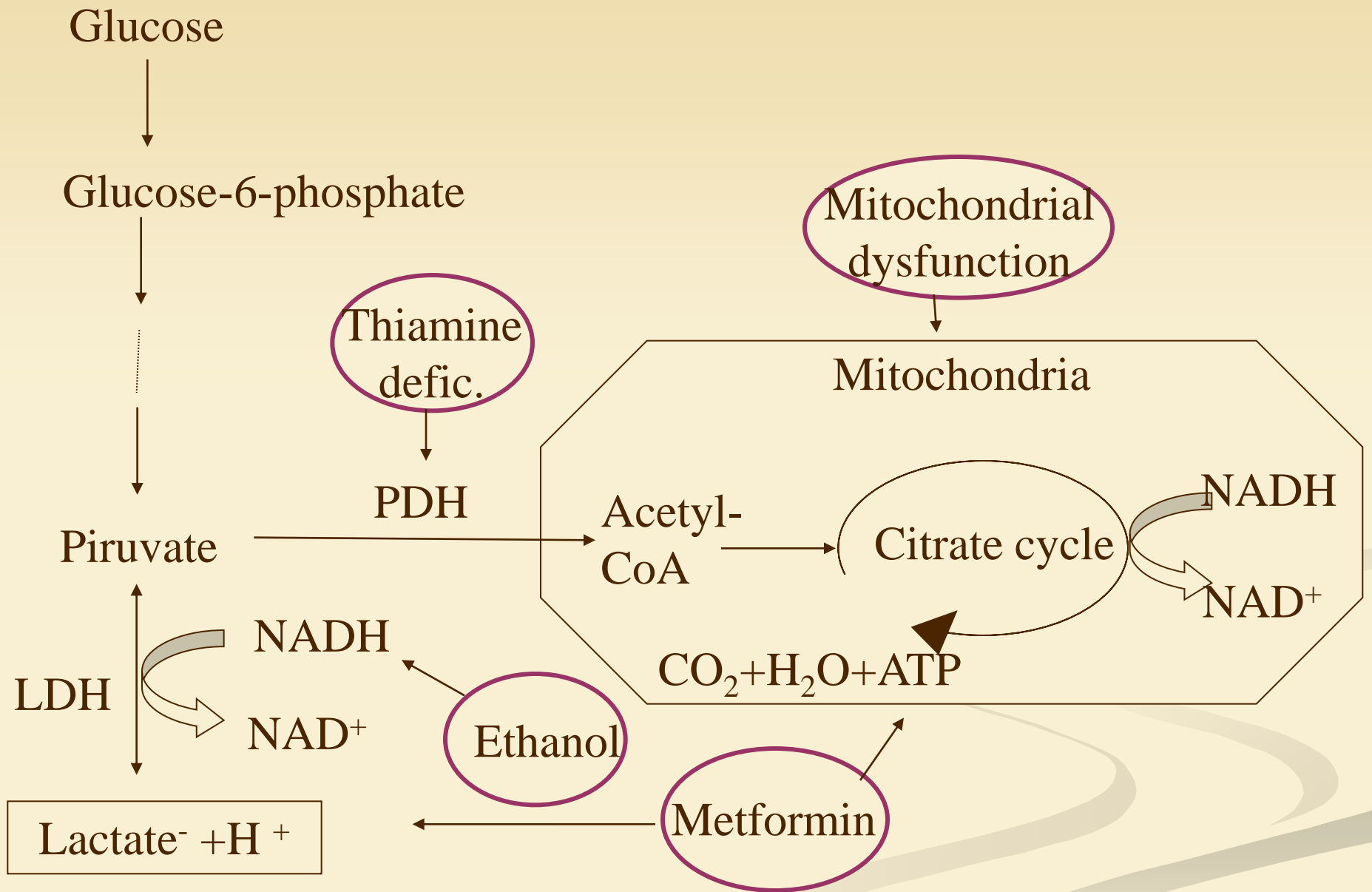
- ketoacidosis
- L-lactic acidosis (A and B-types)
- D-lactic acidosis
- advanced renal failure

Exogenous acids

- ethylene glycol
- methanol

Oxygen is necessary for the ATP production in the mitochondria. „A” type L-lactic acidosis develops, when there is no oxygen supply, therefore ATP is not produced. The consequence: increased glycolysis and lactate production.





The most frequent causes of „B” type L-lactic acidosis

Acidosis

pH ↓

$\text{HCO}_3^- \downarrow$

metabolic acidosis



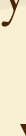
$\text{pCO}_2 \downarrow$

respiratory compensation

1:1

$\text{pCO}_2 \uparrow$

respiratory acidosis



$\text{HCO}_3^- \uparrow$

metabolic compensation

1:0,3

Alkalosis

pH ↑

HCO₃⁻ ↑

metabolic alkalosis



pCO₂ ↑

respiratory compensation

1:0,7

pCO₂ ↓

respiratory alkalosis



HCO₃⁻ ↓

metabolic compensation

1:0,5

Deteriorating pH

- A 42 yrs old male patient
- He was admitted to the gastroenterology department in rapidly deteriorating physical condition
- Medical Hx: joint problems, hip replacement, on NSAIDs, aethyl abusus
- Labs on admission:
 - Serum Na 138 mmol/l, K 2,9 mmol/l, Cl 121 mmol/l, BUN 24,8 mmol/l, creatinine 432 umol/l, Hgb 9,6 g%
 - Blood gas analysis:
pH 7,3, bicarbonate 11 mmol/l, pCO₂ 14 mmHg
- Gastroscopy revealed bleeding from gastric ulcers
- **What kind of acid-base disorder is this?**

Deteriorating pH

- Decrease in bicarbonate

$$25 - 11 = 14 \text{ mmol/l}$$

- delta pCO₂

$$40 - p\text{CO}_2 = 40 - 14 = 26 \text{ mmHg}$$

- Anion gap

$$\text{Na}^+ - (\text{Cl}^- + \text{HCO}_3^-) = 138 - (121 + 10,8) = 6,2 \text{ mmol/l}$$

Deteriorating pH

- Treatment: he got proton pump inhibitor, Na-bicarbonate infusion and furosemide, but did not improve
- Blood gas analysis few hours later:
pH 7,2, bicarb 9 mmol/l, pCO₂ 24 mmHg
(previous: pH 7,3, bicarb 11 mmol/l, pCO₂ 14 Hgmm)
- **What happend to the pH?**

Deteriorating pH

- Next day:
pH 6,84 bicarb 9 mmol/l, pCO₂ 58 mmHg
(previous: pH 7,3, bicarb 11 mmol/l, pCO₂ 14 Hgmm
pH 7,2, bicarb 9 mmol/l, pCO₂ 24 mmHg)
- **What kind of acid-base disorder is this now?**

A complicate case with dRTA

- 33 yrs old male patient
- PMHx: ileocecal reticulosarcoma, ileum and colon resection, irradiation colitis, moderate chronic renal failure, recently: suspected distal renal tubular acidosis
- Medical Hx: has usually 3-4 bowel movements/day
Had several GI tests performed recently, and severe, watery diarrhoea developed
Admitted in a very poor condition, severly volume depleted
- Labs: Serum Na 142 mmol/l, K 2,99 mmol/l, Cl 119 mmol/l, BUN 7,6 mmol/l, creatinine 224 umol/l,
- Blood gas analysis:
pH 7,2, bicarb 12 mmol/l, pCO₂ 32 mmHg

A complicate case with dRTA

- What kind of acid-base disorder is this?
- Why is he hypokalemic?

A complicate case with dRTA

- Combined metabolic and respiratory acidosis
 - delta bicarbonate: $25 - 12 = 13$ mmol/l
 - delta pCO₂: $40 - 32 = 8$ mmHg
- Anion gap: $142 - (119 + 12) = 9$
(lactate level 1,03 mmol/l)
- Urinary K excretion 9,9 mmol/day
 - non-renal K loss
- **Is this acidosis due to gastrointestinal bicarbonate loss or renal abnormality?**

Differential diagnosis of non-anion gap metabolic acidosis

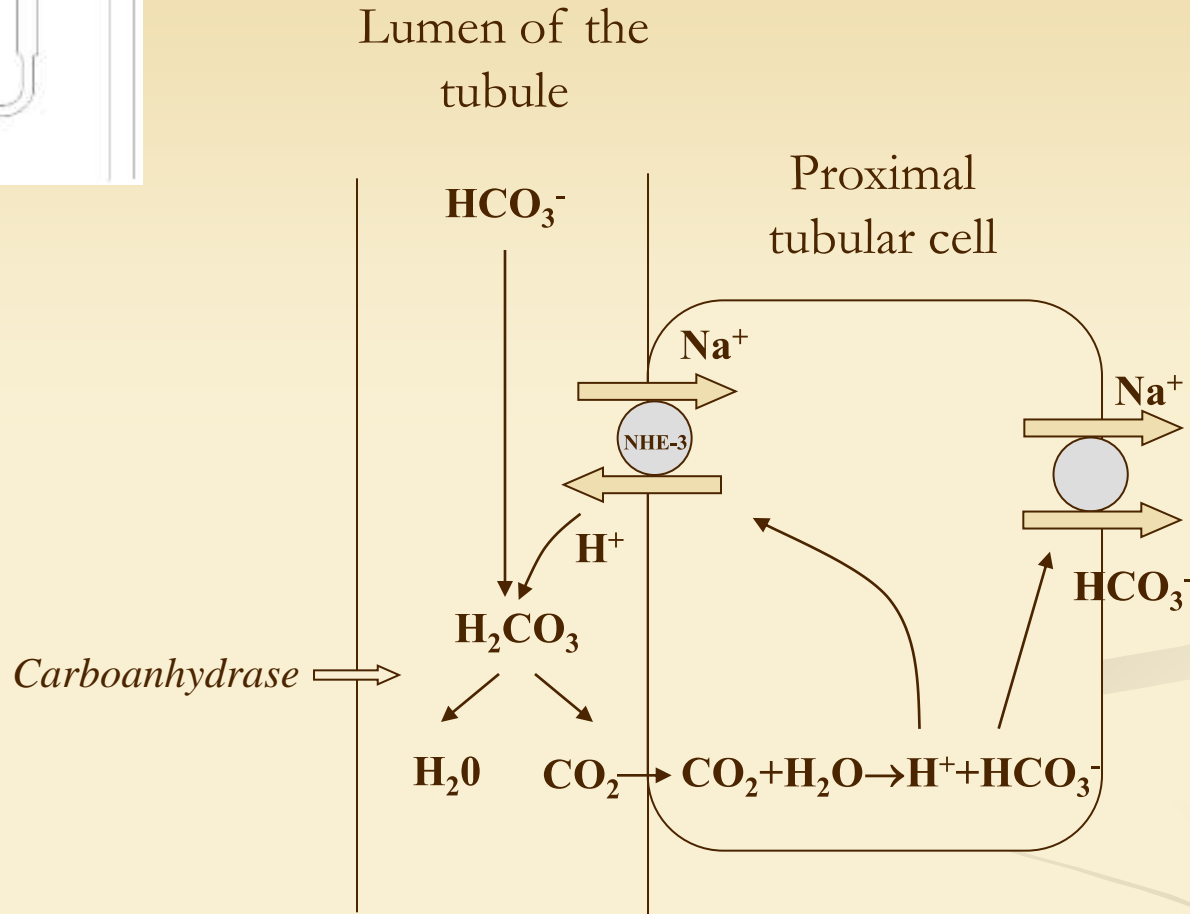
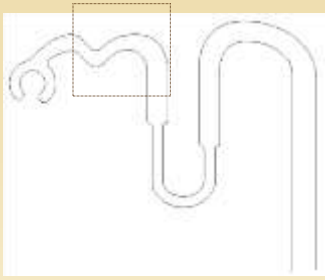
Low plasma bicarbonate level

Gastrointestinal bicarbonate loss

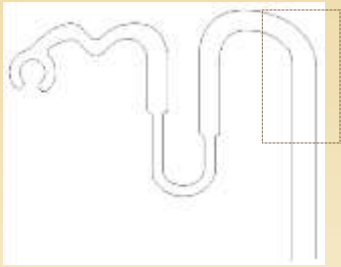
Urine $\text{Cl}^- \gg \text{Na}^+ + \text{K}^+$
(refers to NH_4^+ production)

Renal tubular abnormality

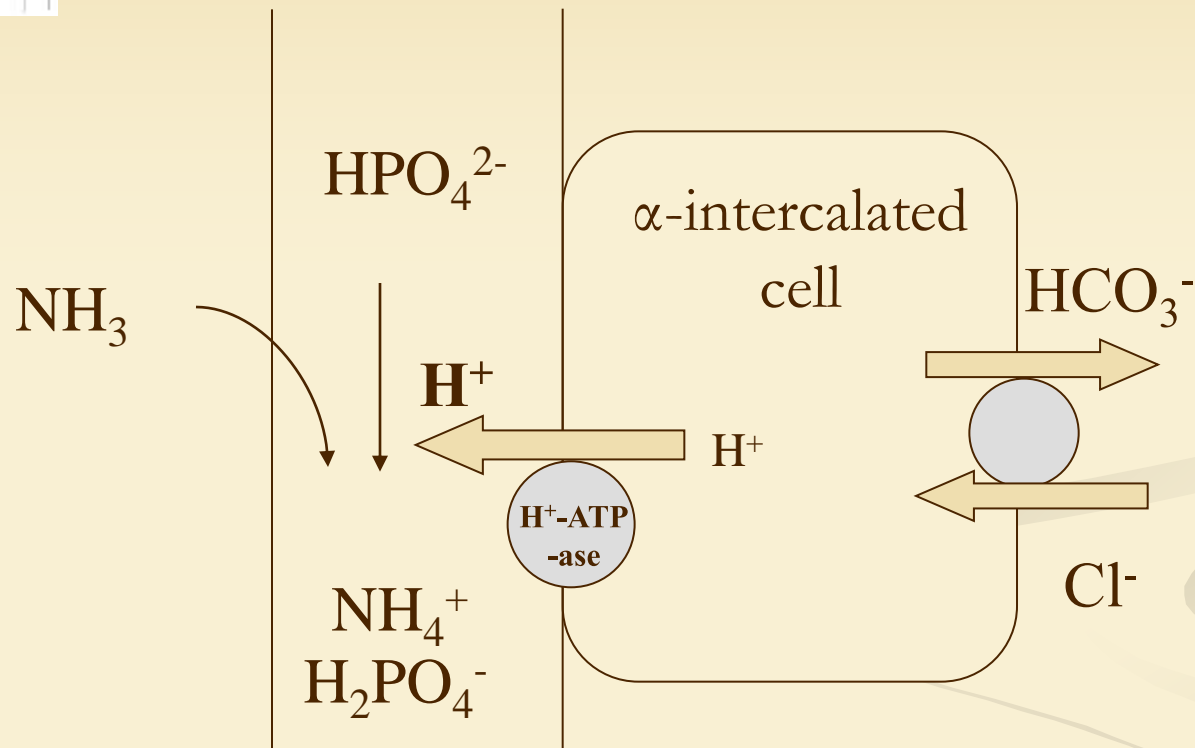
-pRTA
-dRTA
-insufficient NH_4^+ production



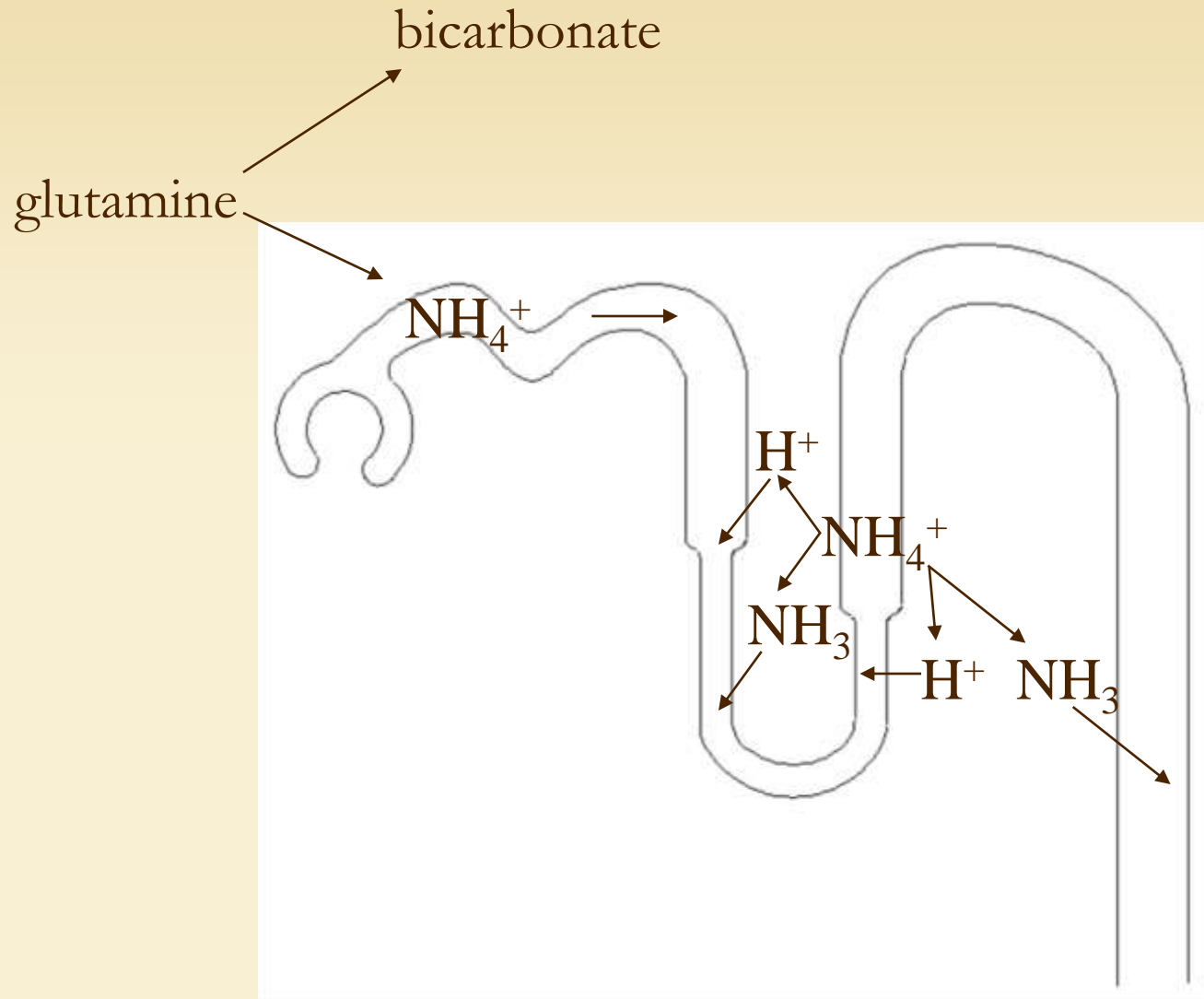
Physiologic bicarbonate reabsorption in the proximal tubule.
 pRTA develops due to reduced indirect bicarbonate reabsorption.



Collecting duct



dRTA develops if the H^+ secretion is disturbed in the collecting duct



Bicarbonate-ammonia production is necessary for H⁺ excretion

Severe hypokalemia due to dRTA

- 40 years old female, admitted to intensive care unit with fever, pneumonia, respiratory failure
- Serum Na 138 mmol/l, K 2,4 mmol/l, Cl 110 mmol/l
pH 7,32, HCO₃ 14,8 mmol/l, pCO₂ 28 Hgmm
AG 13,2 mmol/l
U_{Na} 244 mmol/day, U_K 52 mmol/day
- Bicarbonate loading test:
Baseline urine: pH 6,8, HCO₃ 6 mmol/l, pCO₂ 35 Hgmm
Urine after loading: 7,1 12 36
Blood after loading 7,3 22 37
- Urine pCO₂ – blood pCO₂ ≈ 0

Gastrointestinal disease diagnosed by blood pH

- A 41 year-old male patient, admitted to hospital secondary to nausea, vomiting and epigastric pain
- On admission he looked severely volume depleted, had a BP of 96/58 mmHg
- Labs: serum Na 123 mmol/l, K 3,5 mmol/l, BUN 24 mmol/l, creatinine 355 umol/l,
- Blood gas analysis: pH 7,65, bicarb 43 mmol/l, pCO₂ 55 mmHg.
- **What is your diagnosis?**

Gastrointestinal disease diagnosed by blood pH

- Metabolic alkalosis, hyponatremia and hypokalemia, acute renal failure
- According to the clinical picture – suspicion of pylorus stenosis
- Gastrosocopy confirmed this abnormality (ulcer causing pylorus stenosis)

Are both the patient and the doctor anxious?

- 70 years old male pt, HTN, stroke few months ago
- Presented with dizziness, fever and chills, shortness of breath
- Blood gas analysis:
pH 7,70, pCO₂ 11 mmHg, HCO₃ 14 mmol/l
- **What is the differential diagnosis?**

Differential diagnosis of respiratory alkalosis

- Hypoxia
 - pulmonary disease, congestive heart failure
- Pulmonary receptor stimulation
 - pneumonia, pulmonary embolism, asthma, fibrosis
- Drugs
 - salicylates, theophylline, catecholamines
- CNS disorders
- Miscellaneous
 - psychogenic hyperventilation, fever, Gram-negative sepsis, cirrhosis
- In our pt: CRP 182 mg/l, likely Gram-neg. sepsis+anxiety

The most difficult case for me

The image features a light beige background with a subtle gradient. In the lower right quadrant, there are several overlapping, wavy, light-colored lines that create a sense of movement and depth. The text 'The most difficult case for me' is centered horizontally and rendered in a bold, serif font with a slight shadow effect.

Sylvia

- 40 years old female, lives with severe hypotension (BP 78/53 mmHg)
- Serum potassium 1,4-1,8 mmol/l for 8 years
Cl 80 mmol/l, Mg 0,43 mmol/l, Ca 2,33 mmol/l, uric acid 830 umol/l
- Urinary electrolytes:
Na 94 mmol/day, K 31 mmol/day, uric acid 1159 umol/day, Ca 0,25 mmol/day
- Blood gas analysis:
pH 7,6-7,7, HCO₃ 36-51 mmol/l

Bartter and Gitelman syndromes

- Urinary potassium wasting → hypokalemia
- Salt diuresis → volume depletion, low BP
- Metabolic alkalosis
- Urinary magnesium wasting → hypomagnesemia

Bartter syndrome:

- like „taking continuously furosemide”
- high Ca excretion

Gitelman syndrome

- like „taking continuously thiazide”
- low Ca excretion