# Ministry of Health of the Republic of Belarus Educational institution ''Gomel State Medical University''

Department of General and Bioorganic Chemistry

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# METHODOLOGICAL MANUAL

for conducting the laboratory class with the first-year students of the Faculty of International Students studying in the specialty 7-07-0911-01 "Medical business" (FIS) English-speaking students in the discipline "Medical chemistry"

**Topic 8: General characteristics of solutions. Heterogeneous equilibria** 

Time: 2 hours

## THE TRAINING AND EDUCATIONAL GOALS, MOTIVATION TO STUDY THE TOPIC

## The purpose of the class:

To form an idea of the process of dissolution of different substances in water, as well as the influence of various factors on the solubility processes. To study heterogeneous equilibria in the system of a "hardly soluble electrolyte – its saturated solution".

# The tasks of the class:

As a result of the class, the student *must know:* 

1) solutions and their classification;

2) thermodynamics of the dissolution process;

3) features of dissolution of gases, liquids and solids in water;

4) solubility; factors affecting solubility;

5) heterogeneous equilibria between a hardly soluble electrolyte and its ions in a saturated solution;

6) conditions for precipitates formation;

7) formation of bone tissue.

## The student *must be able to:*

1) explain the reasons for the change in the solubility of various substances in water;

2) predict the heat effect of the dissolution of substances of various aggregate states in water;

3) determine the conditions of precipitation of hardly soluble electrolytes based on the values of their solubility-product constants;

4) to write the equations of chemical reactions of the formation of the mineral component of bone tissue;

5) explain the reasons for the violation of the formation of bone tissue from a chemical point of view.

# Motivation to study the topic:

The study of the properties of solutions helps the future doctor to form an idea of the important role of solutions and heterogeneous equilibria in the vital activity of the body.

The formation of bone tissue is a typical example of a heterogeneous process *in vivo*. Hydroxyapatite  $Ca_5(PO_4)_3OH$  is a mineral component of bones. Using the values of the solubility product constants ( $K_{sp}$ ), it is possible to estimate at what concentrations of ions, for example, the formation of gallstones, the deposition of cholesterol and calcium in the walls of blood vessels (atherosclerosis and calcification) and salts in the joints will occur.

Heterogeneous processes are also used to correct some pathological states. For example, precipitation reactions are used at poisoning with oxalic acid or its salts, introducing a solution of  $CaCl_2$  as an antidote; when poisoning with barium salts, the stomach is washed with a solution of MgSO<sub>4</sub>. Treatment of gout and urolithiasis is carried out with salts of citric acid, tartaric acid and its salts, lithium salts. Some hardly soluble compounds are used in pharmacotherapy. Their action is based on the combination of heterogeneous and protolytic equilibria. Aluminum hydroxide is an antacid. The dissolution of alumi-

num hydroxide stops at pH = 4, which prevents complete neutralization of gastric contents. At the same time, the activity of digestion decreases, but does not stop. The insoluble part of Al(OH)<sub>3</sub> has an enveloping and adsorbing effect.

Precipitation reactions are used in the quantitative analysis of drugs. The precipitation method is also applied in the clinical analysis of chlorides in urine, gastric juice, blood, in sanitary and hygienic practice – in the analysis of drinking water.

## NECESSARY EQUIPMENT

1. Methodological manual for students on the topic "General characteristics of solutions. Heterogeneous equilibria".

2. Reference materials of physico-chemical constants for the 1<sup>st</sup> year education international students.

3. Chemical reagents and equipment necessary for laboratory work.

# CONTROL QUESTIONS ON THE TOPIC OF THE CLASS

1. Solutions, types of solutions, thermodynamics of dissolution, solubility.

2. Solubility of gases in water. The effect of temperature, pressure and dissolved electrolytes on the solubility of gases.

3. Solubility of liquids in each other. The Nernst-Shilov's distribution Law as a theoretical basis for extraction.

4. Solubility of solids in water. Heterogeneous equilibrium "hardly soluble electrolyte – its ions in saturated solution".

# **COURSE OF THE CLASS**

## The theoretical part

# 1. SOLUTIONS, TYPES OF SOLUTIONS, THERMODYNAMICS OF DISSOLUTION, SOLUBILITY

**Solution** is a single-phase system of variable composition consisting at least of two components: a solute and a solvent.

**Solvent** is a component that determines the physical state of a solution. Normally it presents in the greatest amount.

A solution in which the solvent is water is an *aqueous solution*, whereas in a *nonaqueous* solution, the solvent is a substance other than water. Familiar examples of nonaqueous solvents are ethyl acetate, used in nail polish removers, and turpentine, used to clean paint brushes.

Water is the most important solvent in nature and the most abundant substance in the world. The loss of 2/3 volume of intra-cellular fluid may cause death. The excess of water is also dangerous; it may result in cell swelling and edema.

# **Classification of solutions:**

a) according to the aggregate state:

• gaseous: air;

 liquid: sugar in water, ethanol in water, all biological fluids (blood, plasma, lymph and etc.);

• solid: metal alloys.

## *b)* according to the degree of saturation:

• saturated: contain the maximum possible amount of solute that can dissolve under a given set of conditions; exist at equilibrium state with the excess of a solute;

When the maximum possible amount of solute has dissolved, the solution becomes saturated. If excess of solute is present, the rate of dissolution is equal to the rate of crystallization – an equilibrium state is maintained.



Figure 1 – Saturated solutions exist at equilibrium state with the excess of a solute

• unsaturated: contain less amount of a solute that saturated solution;

• supersaturated: contain the greater amount of a solute than saturated solution. They are unstable.

# Thermodynamics of dissolving process

Dissolution is a spontaneous physicochemical process consisting of three steps:

1) atomization – a destruction of solute's crystal lattice which runs with heat absorption ( $\Delta_{at}H > 0$ );

2) solvation (hydration) – a formation of solvates (hydrates) complexes which runs with heat releasing ( $\Delta_{sol}H < 0$ );

Solvation is the process in which solute particles are surrounded by solvent molecules. When the solvent is water, the process is called hydration.

3) a diffusion of solute particles through a solution – proceeds without any heat absorption or releasing ( $\Delta_{dif}H \approx 0$ ).

**Heat of solution** (the overall enthalpy change in the formation of the solution,  $\Delta_{soln}H$ ) is an integral magnitude which is calculated as:  $\Delta_{soln}H = \Delta_{at}H + \Delta_{sol}H + \Delta_{dif}H$ 

**Heat of solution** is amount of heat absorbed or released when one mole of a solute is dissolved in the endless amount of a solvent under the standard conditions.

Dissolution of most solids is endothermic  $(\Delta_{soln}H > 0)$  since the amount of heat absorbed in the first step exceeds the amount of heat released in the second step of a process.

*Dissolution of most gases* is exothermic ( $\Delta_{soln}H < 0$ ) since the first step of a process doesn't occur (gases do not form crystals under normal conditions).

Dissolution of one liquid in another is in fact a diffusion of one liquid into another. This process is accompanied by negligibly low heat effect ( $\Delta_{soln} H \approx 0$ ) as diffusion proceeds without any heat absorption or releasing.

**Solubility** (**S**, **g/l or mol/l**) is an ability of substance to be dissolved in a particular solvent. It's characterized by a solute's content in a saturated solution under certain temperature.

Solubility depends upon:

1) nature of solvents and solutes: the effect of nature is described by the rule «Like dissolves like";

Polar compounds are readily dissolved in polar solvents, while nonpolar compounds are readily dissolved in nonpolar solvents. For example, NaCl exhibits high solubility in water and low solubility in benzene; on the contrary, I<sub>2</sub> exhibits high solubility in benzene and low solubility in water.

2) temperature;

3) pressure (if gas is dissolved);

4) other solutes presence in a solution [2,3].

## 2. SOLUBILITY OF GASES IN WATER. THE EFFECT OF TEMPERATURE, PRESSURE AND DISSOLVED ELECTROLYTES ON THE SOLUBILITY OF GASES

Gas dissolution in water may be represented by a following scheme:

#### $A(g) + H_2O \rightleftharpoons A(aq), \Delta_{soln}H < 0$

According to Le Chatelier's principle, an increase in temperature shifts the equilibrium to the left, thereby reducing solubility, while a decrease in temperature favors solubility of gases.

According to Le Chatelier's principle, an increase in pressure shifts the equilibrium to the right, thereby increasing solubility, while a decrease in pressure reduces the solubility of gases.

According to Henry's Law the solubility of a gas in a liquid is proportional to the partial pressure of a gas above the solution:

$$S = k \times p$$
,

k – Henry's Law constant,

S – gas concentration in a solution,

p – partial pressure of a gas in the vapor phase above a solution.

Henry's Law explains gas exchange between a human body and its surroundings. Thus it gives reason for such a phenomenon as a caisson disease or bend. Deep-sea divers, astronauts and pilots of supersonic planes can suffer this disease. When a person appears in a high-pressure environment, blood and tissues are saturated with nitrogen. A caisson disease occurs when the pressure decreases rapidly (for example, when a diver rise from a depth to the surface too quickly). At the same time, the gas, previously dissolved in blood, forms gas bubbles in the blood vessels. Typical symptoms include pain and/or neurologic disorders. Severe cases can be fatal. Observance of safety rules by a diver is vital for the prevention of caisson disease. Divers avoid the bends by returning to the surface slowly to allow excess gas to escape from the blood without forming bubbles.

**Sechenov's equation** postulates that gas solubility in pure water is higher than in electrolytes' solutions:

$$S=S_o\times e^{-kc},$$

 $S(S_o)$  – gas solubility in a solution (in pure water),

e – a base of natural logarithm,

c – an electrolyte concentration,

k – Sechenov's equation constant.

Sechenov's equation makes clear why gas concentration in blood is lower than in pure water.

## 3. SOLUBILITY OF LIQUIDS IN EACH OTHER. THE NERNST-SHILOV'S DISTRIBUTION LAW AS A THEORETICAL BASIS FOR EXTRACTION

Liquids dissolution in water may be represented by the following scheme:

$$A(l) + H_2 O \rightleftharpoons A(aq), \Delta_{soln} H \approx 0$$

Increase in temperature initiates mutual diffusion of liquids thus increasing their solubility. Three types of liquids can be distinguished:

a) completely soluble in each other (miscible): for example, H\_2SO\_4/H\_2O, C\_2H\_5OH/H\_2O;

b) partially soluble in each other: for example,  $C_6H_6/H_2O$ ;

c) insoluble in each other (immiscible) for example, Hg/H<sub>2</sub>O.

**The Nernst-Shilov's distributional Law** deals with a heterogeneous equilibrium which is maintained when the third component is added into a system composed of two immiscible liquids. This Law is defined in the following way: *when the third component is added into a system of two immiscible liquids, the ratio of its concentrations in both liquids will stay constant at a certain temperature* (Fig. 2).



Figure 2 - A scheme of two-phase system composed of immiscible liquids, where [A]' and [A] are concentrations of the third component in both phases

The relationship of the third component concentrations in phases 1 and 2 is a constant value at a given temperature:

$$\frac{[A]'}{[A]} = K;$$
 K is distributional equilibrium constant

The Nernst-Shilov's distributional Law is a theoretical base for extraction -a method to separate mixtures into individual components [1,3].

# 4. SOLUBILITY OF SOLIDS IN WATER. HETEROGENEOUS EQUILIBRIUM "HARDLY SOLUBLE ELECTROLYTE – ITS IONS IN SATURATED SOLUTION"

The dissolution of solids in water can be represented by the following scheme:

$$A(s) + H_2 O \rightleftharpoons A(aq), \Delta_{sol} H > 0$$

According to the Le Chatelier's principle, an increase in temperature increases the solubility of solids in water.

If a hardly soluble electrolyte (salt, base, or acid) is dissolved, then the heterogeneous equilibrium (occurs when reagents and products are in different phases) is maintained between a solid and its ions in a saturated solution and it can be represented by the scheme:

$$A_n B_{m(s)} \rightleftharpoons n A^{m_+}{}_{(aq)} + m B^{n_-}{}_{(aq)}$$

Solubility equilibrium is characterized by the solubility-product constant (K<sub>sp</sub>):

$$\mathbf{K}_{\mathrm{sp}} = [\mathbf{A}^{\mathrm{m}+}]^{\mathrm{n}} \times [\mathbf{B}^{\mathrm{n}-}]^{\mathrm{m}}$$

K<sub>sp</sub> is useful for predicting solubility of hardly soluble electrolytes:

$$S = \sqrt[n+m]{\frac{K_{sp}}{n^n \times m^m}}$$

For a binary electrolyte n = m = 1:

$$AB_{(s)} \rightleftharpoons A^+_{(aq)} + B^-_{(aq)} K_{sp} = [A^+] \times [B^-]$$
, hence:

$$S = \sqrt{K_{sp}}$$

where solubility (S) is expressed in mol/l.

**Problem 1.** Calculate the solubility of CaCO<sub>3</sub> (in mol/l; g/l) in water at 298 K.  $K_{sp}$  (CaCO<sub>3</sub>) = 4.8×10<sup>-10</sup>

#### Solution:

1. Let's write a heterogeneous equilibrium in a saturated solution of CaCO<sub>3</sub>:  $CaCO_{3(s)} \rightleftharpoons Ca^{2+}(aq) + CO_{3}^{2-}(aq); \quad K_{sp} = [Ca^{2+}] \times [CO_{3}^{2-}] = 4.8 \times 10^{-10}$ 2. Let's assume that solubility of CaCO<sub>3</sub> = S mol/l; hence,  $[Ca^{2+}] = [CO_{3}^{2-}] = S$  mol/l  $K_{sp} = S^{2}; S = \sqrt{K_{sp}}; \quad S(CaCO_{3}) = \sqrt{4.8 \times 10^{-10}} = 2.2 \times 10^{-5}$  mol/l

3. To express the solubility in g/l we have to multiply solubility in mol/l by the molar mass of CaCO<sub>3</sub>:  $S(CaCO_3)$ ,  $g/l = S(CaCO_3)$ ,  $mol/l \times M(CaCO_3) = 2.2 \times 10^{-5} \times 100 = 2.2 \times 10^{-3} g/l$ 

Answer:  $S(CaCO_3) = 2.2 \times 10^{-5} \text{ mol/l}; S(CaCO_3) = 2.2 \times 10^{-3} \text{ g/l}$ 

The lower the  $K_{sp}$  value, the lower the solubility of the hardly soluble electrolyte, the better its precipitation (crystallization) occurs.

The conditions for the formation of a precipitate of a hardly soluble electrolyte can be formulated as follows: precipitation occurs from saturated and supersaturated solutions and never from unsaturated ones.

In saturated solutions:  $[A^+] \times [B^-] = K_{sp}$ In supersaturated solutions:  $[A^+] \times [B^-] > K_{sp}$ 

Total condition for precipitates formation:  $[A^+] \times [B^-] \ge K_{sp}$ 

**Problem 2.** 0.0005 M AgNO<sub>3</sub> solution is treated with equal volume of 0.0001 M KBr solution. Does AgBr precipitate from this solution?  $K_{sp}$  (AgBr) = 5.3×10<sup>-13</sup>.

Solution: 1.  $AgNO_3 + KBr \rightarrow \underline{AgBr} + KNO_3$  2. Conditions for formation of AgBr precipitate:

if  $[Ag^+] \times [Br^-] \ge K_{sp}$ , then AgBr precipitates from this solution.

3. Let's calculate the concentration of each ion in a given solution:

As a result of mixing two equal volumes of solutions, the volume of the final solution increased by 2 times, and the concentration of each substance decreased by 2 times:

 $[Ag^+] = 0.0005/2 = 0.00025 M; \ [Br^-] = 0.0001/2 = 0.00005 M$ 

4. The product of  $Ag^+$  and  $Br^-$  concentrations is equal to:

 $[Ag^+] \times [Br^-] = 0.00025 \times 0.00005 = 1.25 \times 10^{-8}$ 

5. Let's compare the calculated product of concentrations with  $K_{sp}(AgBr)$ :  $1.25 \times 10^{-8} > 5.3 \times 10^{-13}$ 

Answer: the solution is supersaturated; AgBr precipitates

**Bones tissue formation** is one of the most important heterogeneous processes *in vivo*. Hydroxy apatite is the mineral component of bones: Ca<sub>5</sub>(PO<sub>4</sub>)<sub>3</sub>OH.

In blood at pH 7.4 calcium cations  $Ca^{2+}$  and anions of phosphoric acid HPO<sub>4</sub><sup>2-</sup> and H<sub>2</sub>PO<sub>4</sub><sup>-</sup> are present. The amounts of both anions are approximately equal, but CaHPO<sub>4</sub> (K<sub>sp</sub> =  $2.7 \times 10^{-7}$ ) is less soluble than Ca(H<sub>2</sub>PO<sub>4</sub>)<sub>2</sub> (K<sub>sp</sub> =  $1 \times 10^{-3}$ ). Thus, it is proved that the formation of CaHPO<sub>4</sub> is the first step in the generation of bone tissue:

**1**)  $Ca^{2+} + HPO_4^{2-} \rightleftharpoons CaHPO_4$ 

In the next steps the following salts are produced:

2)  $3CaHPO_4 + Ca^{2+} + 2OH^- \rightleftharpoons Ca_4H(PO_4)_3 + 2H_2O$ 

3)  $Ca_4H(PO_4)_3 + Ca^{2+} + 2OH^- \rightleftharpoons Ca_5(PO_4)_3OH + H_2O$ 

Solubility of hydroxyapatite is very low ( $K_{sp} = 1.0 \times 10^{-58}$ ), so the bones are extremely stable. The excess of Ca<sup>2+</sup> shifts equilibrium to the right, which leads to calcification of the bones. Ca<sup>2+</sup> deficiency shifts equilibrium to the left, which leads to bone destruction. Children suffer from rickets, adults – osteoporosis.



Figure 3 – Bone tissue: a) osteoporosis: vertebrae are deformed, b) healthy bone tissue: vertebrae are not deformed; bones are dense

Calcium deficiency leads to incorporation of beryllium and strontium into the bone tissue. Their incorporation results in beryllium and strontium rickets (bones become fragile and brittle). The incorporation of Sr-90 radionuclide into the bone tissue results in its radioactive irradiation, which may cause leucosis and other oncological diseases. Thus, calcium prevents the accumulation of radioactive strontium in the human body [1-3].

#### The practical part

Safety instructions before laboratory work.

#### LABORATORY WORK No. 1

Preparation of hardly soluble sulfates of alkaline earth metals and comparison of their solubility

Sulfuric acid and all water-soluble sulfates form white crystalline precipitates of sulfates when interacting with solutions of calcium, strontium and barium salts:

 $\begin{array}{l} CaCl_{2} + H_{2}SO_{4} \rightarrow CaSO_{4} \downarrow + 2HCl \\ Ca^{2+} + SO_{4}^{2-} \rightarrow CaSO_{4} \downarrow \\ SrCl_{2} + H_{2}SO_{4} \rightarrow SrSO_{4} \downarrow + 2HCl \\ Sr^{2+} + SO_{4}^{2-} \rightarrow SrSO_{4} \downarrow \\ BaCl_{2} + H_{2}SO_{4} \rightarrow BaSO_{4} \downarrow + 2HCl \\ Ba^{2+} + SO_{4}^{2-} \rightarrow BaSO_{4} \downarrow \end{array}$ 

*Performing the experiment:* pour 5-6 drops of salt solutions containing  $Ca^{2+}$ ,  $Sr^{2+}$ ,  $Ba^{2+}$  ions into three test tubes and add 2-3 drops of sulfuric acid solution. Note the rate of precipitation. Using the values of solubility constants, explain why the  $BaSO_4$  precipitate falls out almost instantly, the  $SrSO_4$  precipitate – after some time, and the  $CaSO_4$  precipitate can only be obtained from concentrated solutions of calcium salts.

Add 1-2 drops of organic solvent (benzene, ethyl alcohol or acetone) to a test tube with a solution of calcium salt. What are you observing? Why does the addition of organic solvents contribute to the precipitation of calcium sulfate?

#### Report form:

1. Write the molecular and molecular-ionic equations of the interaction of calcium, strontium and barium salts with sulfuric acid. Specify the analytical effects of the reactions performed.

2. Make a conclusion about the solubility of calcium, strontium and barium sulfates in water based on the values of their solubility-product constants ( $K_{sp}$ ).

#### LABORATORY WORK No. 2

Conditions for precipitate formation of hardly soluble electrolyte magnesium carbonate

#### Performing the experiment:

a) pour 5 ml of 0.5 M magnesium sulfate solution to the test tube and add 5 ml of 0.5 M sodium carbonate solution. What are you observing? Pay attention to the color of the precipitate;

b) pour 5 ml of 0.005 M magnesium sulfate solution to the test tube and add 5 ml of 0.005 M sodium carbonate solution. What are you observing?

#### Report form:

1. Write the molecular and molecular-ionic equations of the interaction of magnesium sulfate and sodium carbonate. Note the color of the precipitate.

2. Calculate the product of concentrations of  $Mg^{2+}$  and  $CO_3^{2^-}$  ions in solutions obtained by mixing equal volumes of initial solutions of  $MgSO_4$  and  $Na_2CO_3$ . Make a conclusion about the nature of the prepared solutions and the possibility of the formation of a precipitate of hardly soluble  $MgCO_3$  from them. Do the theoretical calculations coincide with the experimental data?

## Control over the assimilation of the topic

It is conducted in the form of written independent work of students.

## **QUESTIONS FOR SELF-CONTROL OF KNOWLEDGE**

1. Solutions, types of solutions, thermodynamics of dissolution, solubility.

2. Solubility of gases in water. The effect of temperature, pressure and dissolved electrolytes on the solubility of gases.

3. Solubility of liquids in each other. The Nernst-Shilov's Distribution Law as a theoretical basis for extraction.

4. Solubility of solids in water. Heterogeneous equilibrium "hardly soluble electrolyte – its ions in saturated solution". Solubility-product constant. Conditions for the formation of a precipitate of a difficult-to-dissolve electrolyte. Heterogeneous equilibria in the formation of bone tissue and the formation of concretions in urolithiasis and cholelithiasis.

## Exercises for the self – control

**1.** What is the concentration of metal ions in the following saturated solutions (mol/l and g/l) at 298 K:

a) AgNCS, $K_{sp} = 1.1 \times 10^{-12}$ ;	c) BaCO <sub>3</sub> , $K_{sp} = 1.0 \times 10^{-10}$ ;
b) AgI, $K_{sp} = 8.3 \times 10^{-17}$ ;	d) PbS, $K_{sp} = 2.5 \times 10^{-27}$ ?

Answers: a)  $1.05 \times 10^{-6}$  M,  $1.13 \times 10^{-4}$  g/l; b)  $9.11 \times 10^{-9}$  M,  $9.84 \times 10^{-7}$  g/l; c)  $3.2 \times 10^{-5}$  M,  $4.38 \times 10^{-3}$  g/l; d)  $5.0 \times 10^{-14}$  M,  $1.04 \times 10^{-11}$  g/l

**2.** Oxalic acid (HOOC-COOH) is a toxic substance present in many plants and vegetables, including spinach. Calcium oxalate is hardly soluble salt ( $K_{sp} = 3.0 \times 10^{-9}$  at 25 °C), which is one of the components of kidney stones formed during urolithiasis.

Calculate:

a) the solubility of  $CaC_2O_4$  in water (mol/l and g/l);

b) the molar concentrations of calcium cations and oxalate ions in a saturated solution of this salt.

Answer: a) 
$$5.5 \times 10^{-5}$$
 M,  $7.04 \times 10^{-3}$  g/l; b)  $5.5 \times 10^{-5}$  M,  $5.5 \times 10^{-5}$  M

**3.** 0.005 M AgNO<sub>3</sub> solution is treated with equal volume of 0.001 M KNO<sub>2</sub>. Does AgNO<sub>2</sub> precipitate from this solution?  $K_{sp}$  (AgNO<sub>2</sub>) = 1.6×10<sup>-4</sup>.

Answer: AgNO<sub>2</sub> does not precipitate

**4.** 0.05 M Mg(NO<sub>3</sub>)<sub>2</sub> solution is treated with 0.05 M K<sub>2</sub>CO<sub>3</sub> solution. Does MgCO<sub>3</sub> precipitate from this solution?  $K_{sp}$  (MgCO<sub>3</sub>) = 4.0×10<sup>-5</sup>.

Answer: MgCO<sub>3</sub> precipitates

#### **QUESTIONS FOR INDEPENDENT WORK OF STUDENTS (IWS)**

1. Enthalpy and entropy factors of dissolution, their relationship with the mechanism of dissolution.

2. Heterogeneous equilibria in the vital activity of organisms (formation of calcium hydroxy apatite, formation of concretions in kidney stone and gallstone diseases).

#### LIST OF SOURCES USED

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