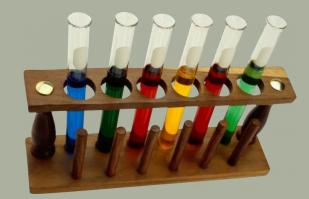


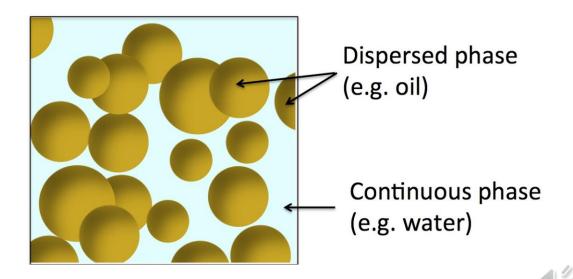
COLLOIDS Lecture 6



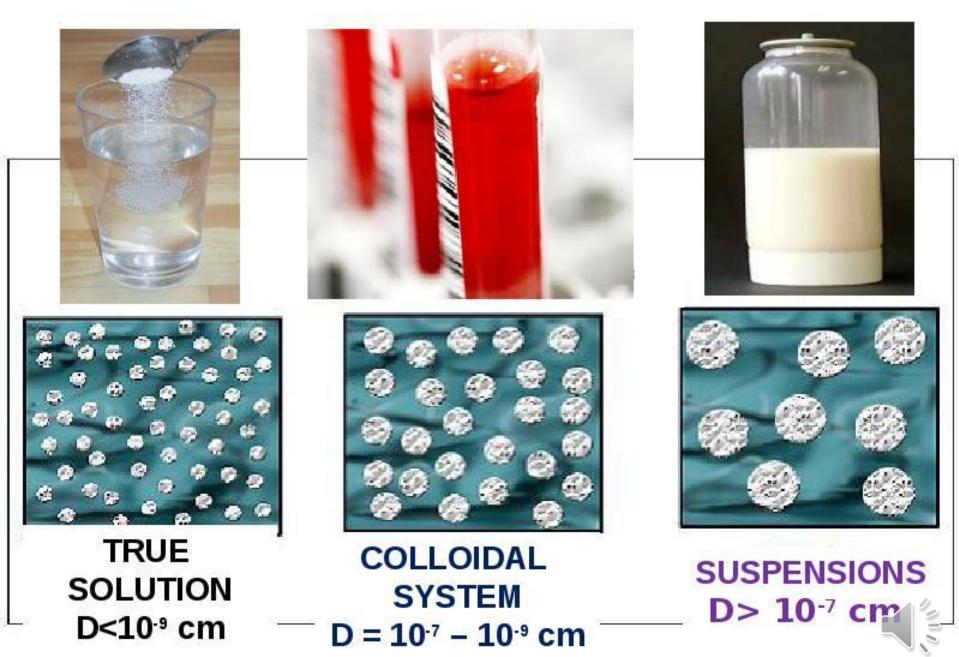


THE CONCEPT OF DISPERSED SYSTEMS

Dispersed systems are heterogeneous (inhomogeneous) systems in which one substance, being in a finely divided state (dispersed phase), is evenly distributed in continuous phase (dispersed medium).



TYPES OF DISPERES SYSTEMS BY PARTICLE SIZE



COLLOIDAL AND TRUE SOLUTIONS

micelles.

Colloidal solutions are True Solution is solution solutions related to in which particles of a dispersed systems, solute are in water or where the particles of other solvent in the form the dispersed phase of molecules, atoms or are in the dispersion ions. For example medium in the form of solutions of lowmolecular weight compounds (salts, acids, alkalis).

Heterogeneous mixture

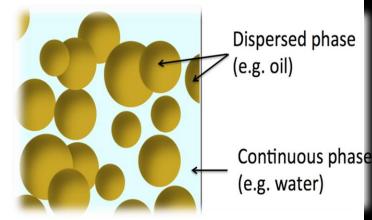
Homogeneous mixture

BASIC DEFINITIONS

- So Colloidal solutions are dispersed systems with a particle size of the dispersed phase exceeding the molecular size (10⁻⁹ 10⁻⁷ m).
 - A dispersed system is a heterogeneous system in which one of the phases is represented by small particles evenly distributed in the volume of another homogeneous phase.
 - A phase is a homogeneous part of the system.
 - Dispersed phase is finely divided particles, evenly distributed in the dispersion medium.

SoDispersion medium (continuous phase)

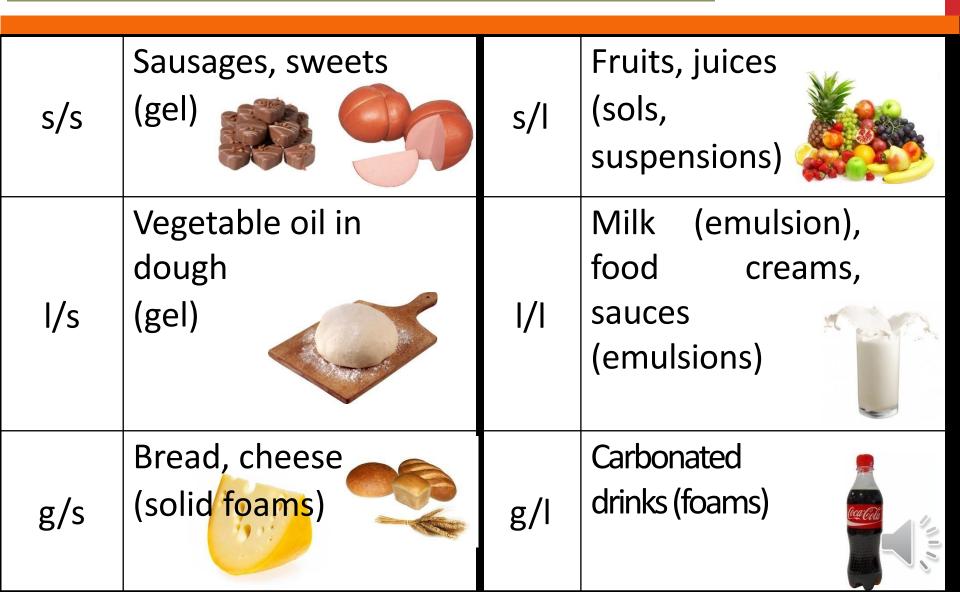
is a homogeneous continuous phase, in which the particles of the dispersed phase are distributed.



CLASSIFICATION

Dispersed systems		Type of dispersed	Examples of dispersed systems
Dispersion phase	Dispersion medium	system, its	
		designation	
Solid	Gas (g)	Aerosol (s/g)	Dust, smoke, snow flakes
	Liquid (l)	Suspensions (s/l)	Clay, toothpaste, lipstick.
		Colloidal solutions	Egg white solution, blood plasma,
		(s/l)	chlorophyll alcohol extract, silicic
		True solutions	acid.
		(s/l)	Solutions of salts, alkalis, sugar.
	Solid (s)	Solid solutions (s/s)	Alloys, minerals, colored glasses.
Liquid	Gas (g)	Spray can (l/g)	Fog, clouds, drizzling rain, spray
			from an aerosol can.
	Liquid (1)	Emulsion (1/1)	Milk, butter, mayonnaise, cream,
		True solutions (l/l)	ointments, emulsion paints.
			Lower alcohols+water, acetone +
			water, acid solutions.
	Solid (s)	Solid emulsion (l/s)	Pearls, opal.
Gas	Gas (g)	No dispersed system	-
		is formed	
	Liquid (l)	Foam (g/l)	Soda water foam, lather, whipped
			cream, whipped cream, candy.
	Solid (s)	Solid foam (g/s)	Styrofoam, foam concrete, foam
			glass, pumice, lava.

EXAMPLES OF DISPERSED SYSTEMS AMONG MEAL

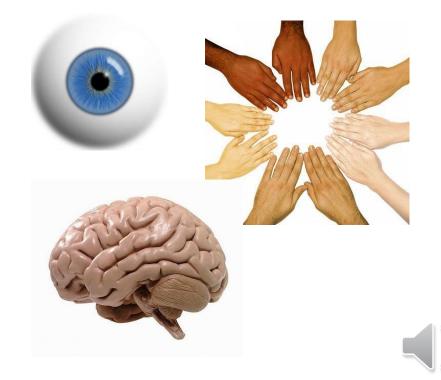


EXAMPLES OF DISPERSED SYSTEMS IN A BODY

Sols are blood, lymph, spinal liquid, saliva, cytosol



Solve and so



COLLOID FORMATION CONDITIONS ARE ...

- Mutual insolubility of the substances forming the system;
- 2) The presence of a stabilizer;
- 3) Very small particle sizes 10⁻⁹-10⁻⁶m.

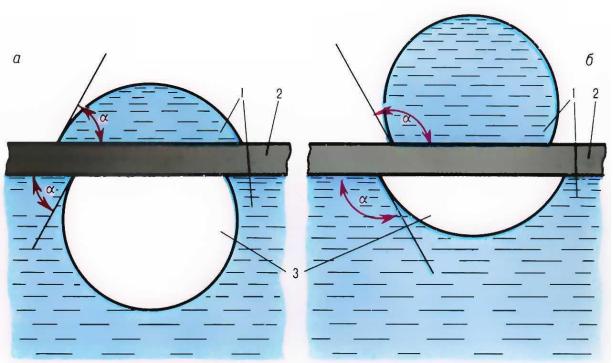
But even if these conditions are met, colloidal solutions, in contrast to true ones, are **thermodynamically unstable**.

The reasons for low stability are **small particle size**, **large specific surface area**.



Classification by degree of interaction dispersion phase with the dispersion medium

 Lyophilic - a systems, where interaction of the particles of the dispersed phase with the solvent is highly expressed.
Lyophobic - dispersion phase interacts weakly with the dispersion medium.



Hydrophilic (a) and hydrophobic (b) surface in a three phase system - water solid - air; 1 - Water 2; - Solid; 3 - air; a - wetting angle.



STRUCTURE OF COLLOID MICELLES

According to the standard theory of micellar sol consists of 2 parts:

•<u>Micella</u> is a colloidal structural unit, surrounded by an electric double layer.

•Intermicellar fluid is the dispersion medium, separating the micelles, where the electrolytes, non-electrolytes and surfactants are soluted.



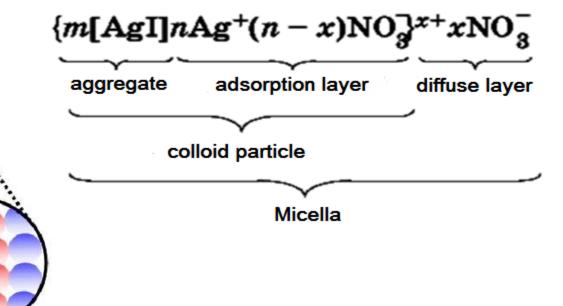
STRUCTURE COLLOID MICELLES

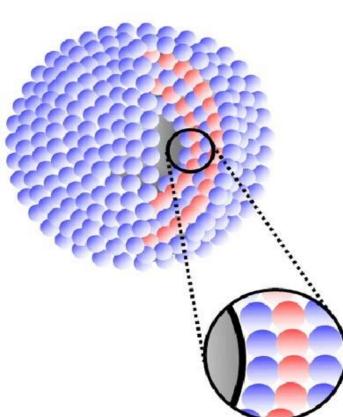
Micelle structure can be considered only as a first approximation, because it has no specific composition

 $KI + AgNO_3(e_X) = AgI\downarrow + KNO_3$

 $m\text{KI} + (m + n)\text{AgNO}_3 \rightarrow m\text{AgI} \cdot n\text{Ag}^+ +$ + $n\text{NO}_3^- + m\text{KNO}_3$

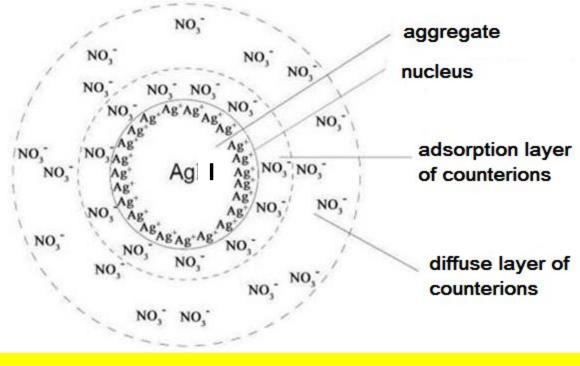
 ${m[AgI]nAg^+(n-x)NO_3^-}^{x+xNO_3^-}$





With an excess of one reactant microchip adsorbs its ions, which do not form a precipitate.

As a result of this microchip acquires a charge, ions, informing him that the charge potential-called, and he charged crystal - core micelles.



Charged core attracts ions from the solution with the opposite charge - counterions; interfacial electrical double layer is formed.

Some part of counterions adsorbed on the surface of the nucleus, forming a so-called adsorption layer counterions; nucleus together with adsorbed thereon are called counterions colloidal particles or granules. The remaining counter, the number of which is determined on the basis of the rules of electrical micelles constitute a diffuse layer of counterions; counterions adsorption and diffusion layers are in a state of dynamic equilibrium adsorption - desorption.

RULE PANETH-FAJANS

"On the surface of the solid assembly is primarily adsorbed ions which:

{[AgI] m n I [n-x)K+]

- included in the assembly;
- able to complete construction of the crystal lattice of the unit;
- form compounds with ions of the unit;
- are isomorphic with the ions of the unit. "

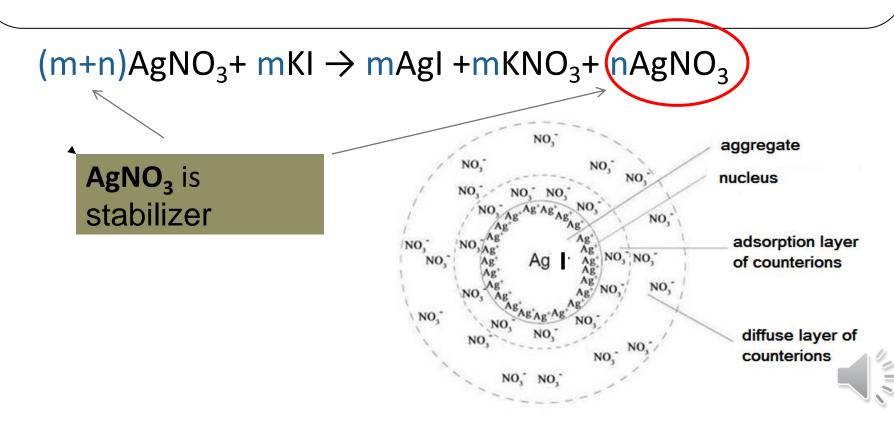
 $mAgNO_3 + (m+n)KI \rightarrow mAgI + mKNO_3 + (nKI)$

KI is stabilizer

RULE PANETH-FAJANS

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- included in the assembly;
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PROPERTIES OF COLLOIDS

Solution Molecular kinetic properties:

Brownian motion (the collision of particles is an exchange of energy and as a result the average kinetic energy is set, same for all particles) Diffusion Osmosis

Solution Optical properties:

ability to scatter light (this phenomenon was observed Faraday (1857) in the study of gold sol. The phenomenon Tyndall in 1868.

> **Solution** Solution



STABILITY OF DISPERSED SYSTEMS

So Sedimentation stability characterizes the ability of particles dispersed phase be in suspension and do not settle under the influence of gravity Aggregate stability characterizes the ability of particles of the dispersed phase oppose them adhesion to each other and thereby maintain unchanged their sizes

The stability of dispersed systems is characterized by the ability of the dispersed phase to maintain the state of uniform distribution of particles of the dispersed phase in the entire volume of the dispersion medium

STABILITY OF DISPERSED SYSTEMS

- 1) Coarsely dispersed systems are heterogeneous and unstable, spontaneously stratified.
- 2) True solutions are homogeneous and indefinitely stable.
- 3) Colloidal solutions occupy an intermediate position on stability.



So COAGULATION IS THE PROCESS OF ADHESION OF COLLOIDAL PARTICLES TO FORM LARGER AGGREGATES DUE TO THE LOSS OF AGGREGATE STABILITY.

THE CONSEQUENCE OF COAGULATION IS A DECREASE SEDIMENTATION STABILITY Coagulation in the body:
Sclerotic changes
vessels as a result of coagulation of colloidal solution

cholesterol



2) Blood clotting process





50 THE COAGULATION THRESHOLD IS THE MINIMUM AMOUNT OF ELECTROLYTE THAT MUST BE ADDED TO THE COLLOIDAL SOLUTION TO INDUCE COAGULATION (CLOUDINESS OR DISCOLORATION OF THE SOLUTION)

$$C_{n\kappa} = \frac{c_{\scriptscriptstyle \mathfrak{I}}V_{\scriptscriptstyle \mathfrak{I}}}{V_{\scriptscriptstyle \kappa p} + V_{\scriptscriptstyle \mathfrak{I}}}$$

- $\sim C_{\pi\kappa}$ coagulation threshold
- $\sim C_{3\pi}$ initial concentration of electrolyte solution

 $\sim V_{3\pi}$ – the volume of electrolyte solution added to the colloidal solution

$$\sim V_{\kappa p}$$
 – colloidal solution volume



COLLOIDAL PROTECTION

- When added to the sols of some high-molecular compounds (as well as surfactants), their resistance to the action of electrolytes increases significantly.
- This phenomenon is called colloidal protection.
- The protected sol can be concentrated and even evaporated to dryness (and becomes thermodynamically stable, as if acquiring the properties of high-molecular compounds).





