Investigation of water structure changes in carbamide and nitrates [NaNO₃, Ca(NO₃)₂, NH₄NO₃] solutions by ultraacoustic method

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Carbamide or urea is the final product of an albumen metabolism of a human being and majority of mammalian. It is produced in liver. It is found in blood, muscles and other tissues, in saliva, lymph and milk. Carbamide regulates water metabolism in an organism an hydratation of tissues. Microorganisms of alimentary canal of ruminant animals use carbamide in biosynthesis of albumen. Carbamide is used in chemical industry for production of formaldehyde resins, medicine, hydrazine, hydrazoformamide, melanine, some sorts of paints and for synthesizing of other combinations, for oil deparaffination and stabilization of smokeless powder. Toothpastes and cosmetical creams contain carbamide too. Carbamide is one of the best concentrated nitrogen fertilizers. In industry carbamide is produced from ammonium hydrate and carbon dioxide. The most usual form of this material granules. Carbamide is used in the main and supplementary fertilization of crops. Nitrates of natrium, calcium and ammonium are also used as nitrogen fertilizers. Complex fertilizers, e. i., mixtures of various fertilizers, are widely used nowadays. The aim of our work is to investigate the influence of carbamide and natrium, calcium and ammonium nitrates on water structure, while dissolving them separately and together at the temperatures 25, 30, 40° C, at the concentration of nitrates 0,5 m (0,5 substance of mole per 1kg of H2O), and changing concentration of carbamide at intervals from 0,5 to 15 m.

Water is compressive, but uncompressive particles - ions - appear after dissolving of salts.

Ions influence water molecules and hydrogen connections between molecules, they can weaken or strengthen the connections, and that is the reason why compression of solution changes.

If volume of the solution is V, and uncompressive volume in it is v_1 , the uncompressive part in the solution is equal to:

$$\alpha = \frac{v_1}{V}.\tag{1}$$

The uncompressive part of the solution volume can be calculated from the adiabatic compression coefficients of water and solution β_0 and β [1 - 10]:

$$\alpha = \frac{\Delta \beta}{\beta_o} = \frac{v_1}{V} \,, \tag{2}$$

where $\Delta \beta = \beta_0 - \beta$.

From the formula (2) Follows that the uncompressive volume in the solution is equal to:

$$v_1 = \frac{\Delta \beta}{\beta_o} V \tag{3}$$

If salts of mass m_o are dissolved in water, it has N molecules ($N=m_oN_A/\mu$, μ is a mole mass, N_A is Avogadro's constant). The uncompressive volume for one molecule is

$$V_{v} = \frac{\Delta \beta V}{\beta_{o} N}.$$
 (4)

Introducing rotation n=N/V (that is a concentration of molecules) we have:

$$V_{v} = \frac{\Delta \beta}{\beta_{O} n}.$$
 (5)

If two salts instead of one are dissolved in the solution, the average uncompressive volume in the solution for one molecule is equal to:

$$V_{\nu} = \frac{\Delta \beta}{\beta_{o}(n_1 + n_2)}.\tag{6}$$

Here $\Delta\beta = \beta_0 - \beta$, β_0 is the coefficient of adiabatic compression of water, β is the coefficient of adiabatic compression of carbamide and nitrate solution, n_1 and n_2 -concentrations of carbamide and nitrate molecules.

An average uncompressive volume, per one molecule, when carbamide and nitrate are dissolved separately, is equal to:

$$V_{\nu} = \frac{1}{2} \left(V_{\nu_1} + V_{\nu_2} \right), \tag{7}$$

where V_{ν_1} is the uncompressive volume of one molecule of carbamide, when dissolved separately, V_{ν_2} is the uncompressive volume of one molecule of nitrate, when dissolved in water separately.

The evaluation of change of water structure is based on the difference of an average uncompressive volume of one molecule, when the substances are dissolved together and separately:

$$\Delta V = \frac{\Delta \beta}{\beta_O (n_1 + n_2)} - \frac{1}{2} \left(\frac{\Delta \beta_1}{\beta_O n_1} + \frac{\Delta \beta_2}{\beta_O n_2} \right) \tag{8}$$

here $\Delta\beta_1 = \beta_0 - \beta_1$; $\Delta\beta_2 = \beta_0 - \beta_2$. β_1 and β_2 are the adiabatic compression coefficients of carbamide and nitrate, dissolved separately.

The coefficients of adiabatic compression (β_0 , β , β_1 , β_2) have been measured by the ultraacoustic method according to the formula $\beta=1/(\rho v^2)$. Here ρ is density of the solution, v is speed of ultrasound in the solution. Speed of ultrasound has been measured by an ultrasound interferometer to within ± 0.01 % (v=2 MHz). The density has been measured by a pyknometer to within 0.001 % (volume of the pyknometer is 25 cm³). Volume of the

solution has been calculated by the formula $V=m/\rho$ (here m is the mass of the solution, ρ is its density).

An uncompressive part of the solution v_1 consists of ions (cations and anions). The volume of these ions can be calculated from their crystallographic radius. Cations of nitrates are surrounded with two layers of water molecules of different structures [11]. Anions have only one water molecules' layer of different structure (exept NO₃, which does not destroy water structure at all [7, 12, 13, 14]). Water molecules which are in the first layer around the cation are only polarized and deformed [11,13]. The dispersive Van der Waals forces appear as a result of deformation. These forces destroy hydrogen connections of water molecules. This phenomenon is called hydratation. Ions of some salts strengthen hydrogen connections between water molecules positive hydratation, other ions weaken while hvdrogen connections between water molecules negative Water compressibility decreases hydratation. hydratation is positive, and increases when hydratation is negative [11]. The uncompressive volume of the ions themselves must not be dependent neither concentration of salts nor from temperature. The polarization layer of cations has to increase when temperature increases, as the Debay screening radius is directly proportional to square root from absolute temperature [15].

Carbamide destroys water structure [16–19] and forms more compact combinations with water molecules, therefore, adiabatic compressibility of the solution decreases, and speed of ultrasound increases. Carbamide, as an organic substance, can have different influence on hydrogen connections from that of nitrate ions can. Thus, the process of hydratation is a complicated one. In our opinion, different ΔV , when salts are dissolved together or separately, occurs only because of changes in hydratation process.

The dependence of v and β on the concentration of carbamide and the dependence of ΔV on temperature and on the concentration of carbamide are depicted in Fig. 1, 2,

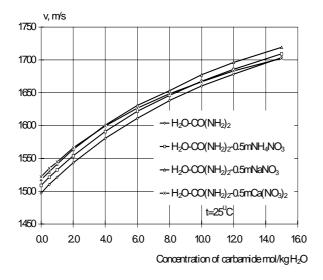


Fig. 1. Dependence of ultrasound ν on the concentration of carbamide at the temperature of 25° C.

3 and 4.

In Fig. 1 the dependence of ultrasound speed v on the concentration of carbamide is shown. We see, that at the increasing concentration of carbamide, the ultrasound velocity increases also. The ultrasound velocity is increased by nitrates in the solution, and sodium nitrate is

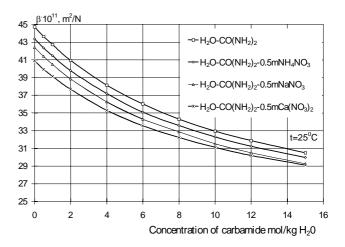
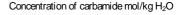


Fig. 2. Dependence of the coefficient of adiabatical compressibility of the solution on the concentration of carbamide at the temperature of 25° C.



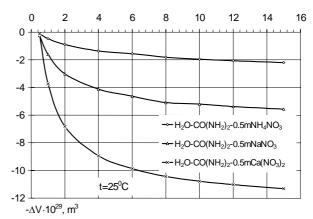


Fig. 3. Dependence of ΔV on the concentration of carbamide at the temperature of 25° C.

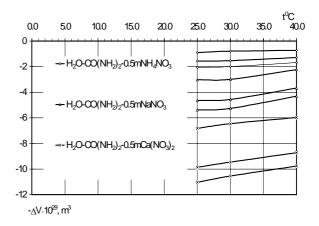


Fig. 4. Dependence of ΔV on the temperature, when concentration of the solution is 2, 6 and 12m.

of the strongest influence here.

Fig. 2 shows the dependence of the solutions' adiabatic compressibility coefficient β on the concentration of carbamide. We see that at the increasing concentration of carbamide, the adiabatic compressibility of solutions decreases. The adiabatic compressibility of solutions is decreased by nitrates, and calcium nitrate is of the strongest influence.

The dependence of ΔV on the concentration of carbamide is shown in the Fig 3. The graph shows that if concentration of carbamide increases, ΔV decreases. When the concentration of carbamide exceeds 10 m, ΔV almost does not change. It means the change of hydratation to be very slight as the number of free water molecules is small.

The Fig. 4 presents the dependency of ΔV on temperature at different carbamide concentrations. It is clear that when temperature increases, ΔV also increases. This dependency of ΔV is not related to the concentration of carbamide. It is possible that the change of hydratation process increases at the increasing temperature.

References

- 1. **Пасинский А. Г.** Сжимаемость и сольватация растворов электролитов. // ЖФХё 1938, т.11,в.5, 606-627.
- Дубинина Е. Ф., Кудрявцев Б. Б. Скорость распространения ультразвука и гидратация растворов. // ЖФХ, 1957, т.31, в.10, 2191-2199.
- 3. **Михайлов И. Г., Соловьев В. А.** Основы молекулярной акустики. М.:Наука, 1964.
- Стюэр Дж., Егер Е. Свойства газов, жидкостей и растворов. Под ред. У.Мезона. М.: Мир, 1969, 375-379.
- 5. Робинсон Р., Стокс Р. Растворы электролитов. М.:ил, 1963.
- Некоторые проблемы современной электрохимии. Пер. с англ., под ред. Дж. Бокриса.
- Михайлов И. Г., Савина Л. И. и др. Скорость звука и сжимаемость концентрированных растворов сильных электролитов. // Вестн.ЛГУ, 1957, в. 22, 25-42.
- Яненас В. К. Исследование гидратации электролитов ультраакустическим методом. // ЖФХ, 1979, т.53, N 4, с. 963-965.

- 9. **Яненас В. К.** Применение ультраакустического и кинематического методов для изучеия гидратации электролитов. // ЖФХ, т. 52, N 6, 1462-1465.
- Яненас В. К. Определение чисел гидратации ионов по коэфициентам сжимаемости и вязкости. // Химия и химическая технология. Т. 21, в. 6, 826-830.
- Janėnas V. K. Nitratų hidratacijos tyrimas ultraakustiniu metodu. // Ultragarsas. K.: Technologija. 1966, Nr. 1.
- 12. **Липидинова И. И.** Уранил и его соединения. М.: АН СССРё 1959ё 250-254.
- 13. Хорн Р. Морская химия. М.: Мирё 1972.
- 14. **Janz J., James D. W.** // Elektrochim. Acta, 1962, vol. 7, p. 427.
- 15. Химическая энциклопедия. М.: 1990ё том 2ё с. 7.
- Frank S., Franks F. Structural approach to the solvent power of water for hydracarbons; urea as a structure breaker // J. Chem. Phys., 1968, 48, 10, 4746-4757.
- 17. **Hammes G. G., Schimmel P. R.** An investigation of water-urea and water-polyethylen glycol interaction // Journal of the American Chemical Society /89:2/, January 18, 1967, 442-446.
- 18. **Beauregard D. V., Barett R. E.** Ultrasonics and water structure in urea solutions //J. Chem. Phys., 1968, 49, 12, 5241-5244.
- Herskovits T. T., Kelly T. M. Viscosity studies of aqueous urea solution // J. Amer. Chem. Soc., 1972, 94, 13, p. 4424.
- V. Janėnas, V. Abaravičiūtė, V. Kasperiūnas,
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Ultraakustinis vandens struktūros pakitimo karbamido ir nitratų $[NaNO_3, Ca(NO_3)_2, NH_4NO_3]$ tirpaluose tyrimas

Reziumė

Ištirta, kaip veikia vandens struktūrą karbamido ir natrio, kalcio bei amonio nitratai, ištirpinti kartu ir atskirai, esant 25, 30 ir 40°C temperatūrai. Nitratų koncentracija buvo keičiama 0,5 m, o karbamido koncentracija - 0,5-15 m intervale. Vandens struktūros pokytis įvertintas iš vienos molekulės vidutinio nespūdaus tūrio skirtumo ΔV , ištirpinus medžiagas kartu ir atskirai. Nustatyta, kad kai karbamido koncentracija didesnė nei 10 m, ΔV beveik nekinta. Tai rodo, kad, esant didesnei karbamido koncentracijai, vandens struktūra jau nebekinta.

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