

STUDYING THE PHYSICAL AND CHEMICAL REGULARITIES OF THE INTERACTION OF CALCIUM OXIDE WITH WATER (p. 4-15)

Valery Shaporev, Inna Pitak, Mykhailo Vasilyev, Oleg Pitak

The paper presents an analysis of the experiment findings in the study of the processes of water hydration of CaO free at different ratios of CaO/H₂O and different activity of CaO.

The analysis of the regularities in the processes of obtaining lump CaO and hydration of the samples proved that the reactivity of CaO is predetermined by the concentration of point defects in the lattice, and the ionicity proportion in the Ca-O compound. The highest reactivity was achieved at 1473 K, the lowest – at 1773 K.

The experimental data on the hydration of CaO of different activity at various ratios of CaO/H₂O allowed confirming the fact that the mechanism of interaction between CaO and water has a series of sequential steps, such as: (1) the capillary water absorption by the sample, (2) chemisorption of water on the surface of CaO crystals, and (3) formation of end products of Ca(OH)₂ solid or Ca(OH)₂ slurry.

The proposed equation is aimed at determining the velocity of the capillary water absorption depending on the reactivity of the CaO samples and properties of the liquid phase. It is found that the rate of the specific surface of the hydrated sample is commensurate with the rate of Ca(OH)₂ formation.

Keywords: process, hydratin, calcium oxide, reaction, water, temperature, kinetics, crystals.

References

- Zaytsev, I. D., Tkach, G. A., Stoev, N. D. (1986). *Proizvodstvo sody*, 311.
- Shaporev, V. P., Tkach, H. A., Khytrova, Y. V., Pohorelov, Yu. Y. (1987). Certificate of authorship 280206 SSSR. (Publykatsiya v otkrytoi pechati zapreshchena). 3181221; declared. 25.09.1987; registered 1.08.1988.
- Cormos, A.-M., Cormos, C.-C., Agachi, P. Ş. (2007). Making soda ash manufacture more sustainable. A modeling study using ASPEN Plus. *Computer Aided Chemical Engineering*, 24, 551–556. doi: 10.1016/s1570-7946(07)80115-5
- Yakymenko, Ya. B. (2014). Deiki zakonirnosti hidratatsiinoi aktyvnosti kaltsiiu oksydu. *Eastern-European Journal of Enterprise Technologies*, 4 (6 (70)), 30–35. doi: 10.15587/1729-4061.2014.26239
- Cizer, O., Campforts, J., van Balen, K., Elsen, J., van Gemert, D. (2006). Hardening of Calcium Hydroxide and Calcium Silicate Binders Due to Carbonation and Hydration. *Brittle Matrix Composites*, 8, 589–599. doi: 10.1533/9780857093080.589
- Rant, Z. (1968). *Die Erzeugung von Soda nach dem Solvay Verfahren*. Sarajevo: Forschungsinstitut Fur Bergbau und Chemische Technologie Tuzla, 543.
- Tkach, H. A., Shaporev, V. P., Tytov, V. M. (1998). *Proizvodstvo sodi po malootkhodnoi tekhnolohyy*. Kharkov: HGPU, 430.
- Shaporev, V. P., Haidash, N. Y., Tkach, H. A. et al. (1987). *Yssledovanye protsessa hasheniya yzvesty vodoi v proizvodstve sodi*. Kharkov: NIOHIM, 65, 127–134.
- Monteagudo, S. M., Moragues, A., Gálvez, J. C., Casati, M. J., Reyes, E. (2014). The degree of hydration assessment of blended cement pastes by differential thermal and thermogravimetric analysis. Morphological evolution of the solid phases. *Thermochimica Acta*, 592, 37–51. doi: 10.1016/j.tca.2014.08.008
- Kohut, M. D. (1997). Rozrobka promyslovoho protsessu vyrobnytstva vapniakovoi suspensiiu metodom hasinnia vapna rozchynamy elektrolitiv. Kharkiv, 25.
- Oldovskyi, P. P., Malkova, L. A. (1981). O deformatsiyakh tverdoy fazy dispersnykh system pod vliyaniem adsorbyrovannoi

- vodi y orhanycheskykh soedyneni. *Inzhenerno-fyzycheskyi zhurnal*, 2, 304–315.
- Shaporev, V. P., Khytrova, Y. V., Darmakian, A. V. (1990). Yssledovanye reaktsyonnoi sposobnosti oksyda kaltsiya, poluchennoho termicheskyim razlozhenyem yzvestniaka. *Vestnyk KhPY. Seriya: Khymycheskaia tekhnika*, 2, 22–26.
 - Speziale, S., Reichmann, H. S., Schiling, H. R., Wenk, H. R., Monterio, P. J. (2008). Determination of the elastic constants of portlandite by Brillouin spectroscopy. *Cement and Concrete Research*, 38, 1148–1153. doi: 10.1016/j.cemconres.2008.05.006
 - Shaporev, V. P. (1995). *Tekhnolohiia neorhanichnykh armuichykh napovniuvachiv*. Kharkiv, 365.
 - Manzano, N., Peleng, R. S. M., Ulm, E. J., Buchiler, M. J., van Duin, A. C. T. (2007). Hydration of calcium oxide surface predicted by reactive force field molecular dynamics. *Langmuir*, 28, 4187–4197. doi: 10.1021/la204338m
 - Pleva, M. (1981). Einflub des Schnellbrandes auf die Eigenschaften des Branntkalkes. *Baustoffindustrie*, 24, 10–12.
 - Luhynyna, Y. H., Barbaniahe, V. D. (1963). Yssledovanye elektroprovodnosti oksyda kaltsiya. *Izvestiya vuzov. Seriya: Khymiya y khymycheskaia tekhnolohiya*, 6, 1011–1015.
 - Horelyk, S. S., Rastorhiev, L. N., Skakov, Yu. A. (1970). *Renthenografycheskyi y elektronografycheskyi analiz*. Moscow: Nauka, 140.
 - Yzuchenye s pomoshchiu metoda EPR vliyaniya temperaturi obzhyha na nekotorye svoystva produktov obzhyha yzvestniaka (1972). *Trudy Vsem. in-ta po proektirovaniyu i nauch.-issled. rabotam «Juzhgirocement»*. Kharkiv: Juzhgirocement, 13, 126–133.
 - Maruchevskaia, N. N., Tabunshchykov, N. P., Aizen, A. M. (1976). Termichesкое razlozhenye polydispersnoho kuskovoho materyala. *Inzhenerno-fyzycheskyi zhurnal*, 30 (3), 441–446.
 - Solodovnykov, A. O., Kakhkharov, V. K., Hulyn, Y. N. (1981). Radyoaktyvnyi yzmyritel stepeni dekarbonizatsyy materyala vo vrashchayushchykhia pechakh. *Sbornik trudov Tal. NYIstromproekt. Seriya: Sovershenstvovanye tekhnolohyy stroytelnykh materyalov*, 7, 54–62.
 - Ostrovskiy, N. M., Chumakova, N. A., Bukhavtsova, N. M., Vernikovskaya, N. V., Aristov, Yu. I. (2007). Vlyaniye kapyl-liarnoi kondensatsyy na protsess sorbtsyy vodi kompozitnyimi sorbentamy. «Khloryd kaltsiya v porystoi matrytse». *Teoreticheskiye osnovy khymycheskoi tekhnolohyy*, 41 (2), 213–216. doi: 10.1134/S0040579507020145
 - Delmon, B. (1972). *Kynetyka heterohennikh reaktsiy*. Moscow: Mir, 554.
 - Budnykov, P. P., Hynstlynh, A. M. (1971). *Reaktsyy v smesiakh tverdikh veshchestv*. Moscow: Strojizdat, 487.
 - Pashchenko, A. A., Tuliaev, S. Kh. (1985). Kontaktnie vzaymode-istviya v tvrdeishchykh smesiakh dlia zakladky v podzemnykh virabotok. *Zhurnal prykladnoi khymyy*, 6, 1273–1278.

DEVELOPMENT OF TECHNOLOGY OF PORTLAND CEMENT USING GAS CONDENSATE SLUDGE RECYCLING WASTE (p. 16-21)

Tatiana Rishenko, Konstantin Vyatkin

The paper deals with the topical issue of increasing the energy efficiency of manufacturing construction materials, such as Portland cement, applying organic-mineral additives. Using gas industry waste – gas sludge as the organic-mineral additive is proposed. Gas sludge due to its chemical composition accelerates the rate of chemical reactions during firing the raw mix of Portland cement clinker, thereby saving energy. Energy saving is associated with a decrease in temperature of the relevant physicochemical reactions during the Portland cement clinker firing.

The paper contains the analysis of theoretical and practical aspects of the problem, proposes solutions to improve the

energy efficiency of Portland cement clinker production by applying the organic-mineral additive, in particular gas sludge, which lowers the temperature of chemical reactions, and also considers the features of chemical reactions in the process of Portland cement clinker firing using the gas sludge waste in the raw mix.

Keywords: gas sludge, organic-mineral additive, Portland cement clinker.

References

1. Leont'ev, S. A., Galikeev, R. M., Fominykh, O. V. (2010). Calculation of process units of the system of collection and preparation of borehole production. Tyumen, 256.
2. Korzhubayev, A. G. (2012). Napravleniya i mehanizmy obespecheniya tehnologicheskoy bezopasnosti neftegazovogo kompleksa Rossii. *Burenie i nef't*, 4, 8–10.
3. Klassen, V. K. (2012). Tehnologiya i optimizacija vyrobnyctva cementu. Belgorod: BDTU, 308.
4. Lugynyna, V. R. (2004). Himiya i himichna tehnologiya neorganichnykh v'jazhuchykh materialiv. Belgorod: BDTU, 1, 240.
5. Bajkov, N. M. (2007). Utilizacija nefljanogo i uglekislogo gaza dlja povysheniya nefteodachi na mestorozhdenijah SShA i Kanady. *Neftjanoe hozjajstvo*, 6, 105–108.
6. Ratinov, V. B., Rozenberg, T. I., Rubinina, I. M. (1962). *Cementy. DAN SSSR*, 145 (5).
7. Kalashnikov, V. I., Guljaeva, E. V., Valiev, D. M., Volodin, V. M., Hvastunov, A. V. (2011). Vysokoeffektivnyye poroshkovo-aktivirovannyye betony razlichnogo funkcional'nogo naznachenija s ispol'zovaniem super-plastifikatorov. *Stroitel'nye materialy*, 11, 44–47.
8. Hodzhomurodov, S. (2003). Osnovnyye polozhenija i primenenie OTDV mikrodur v stroitel'stve Puti razvitiya promyshlennosti stroitel'nykh materialov respublik. Dushanbe, 3–7.
9. Burlov, A. Ju., Samchenko, S. V. (2011). Vlijanie al'ternativnogo topliva na klinkeroobrazovanie i kachestvo klinkera. *Materialy seminarov-konkursov molodykh uchenykh i aspirantov. St. Petersburg: Alitinform*, 30–35.
10. Brown, W. E., Chow, L. C. (2003). A new calcium phosphate fast-setting cement. *Journal of Dental Research*, 62.
11. Association of portlandcement. Available at: <http://www2.cement.org>
12. Romeo, L. M., Catalina, D., Lisbona, P., Lara, Y., Martínez, A. (2011). Reduction of greenhouse gas emissions by integration of cement plants, power plants, and CO₂ capture systems. *Greenhouse Gases: Science and Technology*, 1 (1), 72–82. doi: 10.1002/ghg3.5
13. Oberhauser, S., Kather, S. (2011). CO₂-Capture from cement plants applying oxyfuel concepts. 2nd International Conference on Energy Process Engineering. Efficient Carbon Capture for Coal Power Plants. Frankfurt/Main.

SIMULATION OF THE VAPOR PHASE COMPOSITION IN HNO₃ – H₂SO₄ – H₂O SYSTEM WITH LOW CONTENT OF NITRIC ACID (p. 22-26)

Sergiy Kondratov, Tet'yana Khlyakina

Based on the analysis and processing of literature data, the mathematical model describing the vapor phase composition of the ternary system nitric acid – sulfuric acid – water from the liquid phase composition and temperature at a low mole fraction of nitric acid was developed. It was shown that even with low content of nitric acid in the system, deviations of the ideal solution and the execution of the Raoult's law to within a factor (activity coefficient), the value of which depends on the sulfuric acid concentration are observed. It was found that with an increase in the mass concentration of sulfuric acid from 60 to 80 %, the mean values of the activity coefficients increase linearly with the coefficient of determination 0,998. The general equation of the model – dependence of the partial pressure of the nitric acid on the temperature (t, °C), the mole fraction of nitric acid (x), and the mass concentration of sulfuric acid

(C_{H₂SO₄}, % mass) has the form of a non-linear dependence, obtained by combining the Antoine, Raoult's equations and the dependence of the nitric acid activity coefficient on sulfuric acid concentration.

It was found that using the equations of the model for the ternary system HNO₃ – H₂SO₄ – H₂O at mole fraction of nitric acid in the solution of up to 0.02, it is possible to predict normal boiling points with the maximum error not exceeding 5 °C, and mole fractions of nitric acid in the vapor phase with a relative error no more than 10 %.

The model can be used as a subsystem in developing the mathematical model of the reactor for adiabatic nitration of aromatic compounds, as well as in calculating and designing the regeneration systems of spent sulfuric acid.

Keywords: mathematical modeling, HNO₃ – H₂SO₄ – H₂O system, vapor phase, composition, temperature.

References

1. Zhilin, V., Zbarskiy V. (2006). Sintez i tehnologiya nitroprodukovnykh benzola i toluola. *Rossiiskii Khimichskii Jurnal*, 50 (3), 104–115.
2. Olah, G., Malhorta, R., Narang, S. (1989). Nitration. *Methods and Mechanisms*. New York: VCH, 330.
3. Alexanderson, V., Trecek, J. B., Vanderwaart, C. M. (1977). Adiabatic process for nitration of nitratable aromatic compounds. United States Patent CA1069137A, CA1069137A1, DE2655197A1, DE2655197C2. № 4021498; declared 09.12.1975; published 03.05.1977.
4. Alexanderson, V., Trecek, J. B., Vanderwaart, C. M. (1978). Continuous adiabatic process for the mononitration of benzene. United States Patent CA1083608A, CA1083608A1, DE2821571A1. № 4091042; declared 19.08.1977; published 23.05.1978
5. Munnig, J., Pennemann, B., Rausch, A. K. (2013). Process for the continuous preparation of nitrobenzene. United States Patent CN102153476A, CN102153476B, DE102010006984A1, EP2354117A1, EP2354117B1, US20110196177. № 8357827; declared 02.02.2011; published 22.01.2013.
6. Rausch, A., Knauf, T., Bolton, J., Racoos, A. (2010). Process for the preparation of nitrobenzene by adiabatic nitration. United States Patent CN101456817A, DE102007059513A1, EP2070907A1, EP2070907B1, US20090187051. № 7781624; declared 08.12.2008; published 24.08.2010.
7. König, B.-M., Judat, H., Blank, H. U. (1997). Process for the adiabatic preparation of mononitrotoluenes. United States Patent CA2142257A1, CA2142257C, CN1070472C, CN1111614A, DE4410417A1, EP0668263A1, EP0668263B1. № 5648565; declared 12.07.1996; published 15.07.1997.
8. Gattrell, M. (2014). Process for adiabatic production of mononitrotoluene. United States Patent CN104220416A, DE112013001622T5, US20130253233, WO2013140369A1, WO2013140369A9. № 8907144; declared 22.03.2012; published 09.12.2014.
9. Blank, H. U., Judat, H., König, B.-M. (1998). Process for the adiabatic preparation of mononitrohalogenobenzenes. United States Patent DE4411064A1, EP0675104A1, EP0675104B1. № 5714647; declared 21.01.1997; published 3.02.1998.
10. Washburn, E. W. (Ed.) (1928). *International critical tables of numerical data, physics, chemistry and technology*. New York: McGraw Hill, 3, 444.
11. Ellis, S. R. M., Thwaites, J. M. (1957). Vapour-liquid equilibria of nitric acid – water-sulphuric acid mixtures. *Journal of Applied Chemistry*, 7 (4), 152–160. doi: 10.1002/jctb.5010070402
12. Zhang, R., Wooldridge, P. J., Molina, M. J. (1993). Vapor pressure measurements for the H₂SO₄/HNO₃/H₂O and H₂SO₄/HCl/H₂O systems: incorporation of stratospheric acids into background sulfate aerosols. *Journal of Physical Chemistry*, 97 (32), 8541–8548. doi: 10.1021/j100134a026
13. Kondratov, S. A., Krasil'nikova, A. A. (2013). Model of adiabatic nitration of aromatic compounds. *Eastern-European Journal of Enterprise Technologies*, 3 (6 (66)), 16–20. Available at: <http://journals.urau.ru/eejet/article/view/14783/12585>

14. Perry, R. H., Green, D. W. eds. (1999). Perry's chemical engineers' handbook. New York: McGraw-Hill, 2–79.
15. Hoggett, J. G., Moodie, R. B., Penton, J. R., Schofield, K. (1971). Nitration and aromatic reactivity. Cambridge: University Press, 246.

PHENOMENOLOGICAL STUDY OF VISCOSITY AND DENSITY OF ALKALI METAL HYDROXIDE SOLUTIONS (p. 27-33)

Vladimir Nefedov, Alexander Atapin, Dmitri Golovko

Alkaline solutions are widely used in many industries. For dilute solutions, physicochemical properties, such as solubility of gases, hydration numbers and mobility of ions are well known. A relatively new trend is using concentrated alkaline solutions to obtain ferrates. In these processes, the solution concentration of more than 8 M is used. Studying the processes of transport of matter and charge in concentrated alkaline solutions requires updating the data on the hydration numbers and radii of ions, the effect of concentration, viscosity and nature of electrolytes on them.

A method for determining the hydration numbers of alkali metal cations in hydroxide solutions, based on calculating the number of water moles per mole of cations at concentrations close to the limit was proposed. In addition, the influence of ions, strengthening and breaking the solution structure on the density and viscosity of alkali metal solutions was examined. It was shown that primary hydration of ions has the predominant impact on the solution viscosity in dilute solutions, and secondary – in concentrated.

It was found that a sharp increase in the solution viscosity with increasing concentration occurs with decreasing distance between ions up to $3,7 \times 10^{-7}$ m.

At high concentrations, the solution density is higher in hydrated cations with smaller dimensions. High density of lithium hydroxide solutions in concentrations of from 2 to 4 M can be explained by the fact that the positively hydrated cation is embedded in the structure of water without a significant increase in its volume.

Keywords: concentrated alkaline solution, crystallographic radius, hydration of ions, density, viscosity, structure strengthening and structure breaking ions.

References

1. Kleperis, J. (2012). Electrolysis. Rijeka: InTech, 290. doi: 10.5772/2820
2. Ropp, R. C. (2013). Encyclopedia of the alkaline earth compounds. Elsevier, 1187. doi: 10.1016/B978-0-444-59550-8.01001-2
3. Sharma, V. K (2008). Ferrates: synthesis, properties, and applications in water and wastewater treatment. Oxford: University Press, 509. doi: 10.1021/bk-2008-0985.fw001
4. Alsheyab, M. Jia-Qia, J., Stanford, C. (2009). On-line production of ferrate with an electrochemical method and its potential application for wastewater treatment – A review. Journal of Environmental Management, 90 (3), 1350–1356. doi: 10.1016/j.jenvman.2008.10.001
5. Lapique, F., Valentin, G. (2002). Direct electrochemical preparation of solid potassium ferrate. Electrochemistry Communications, 4 (10), 764–766. doi: 10.1016/s1388-2481(02)00438-1
6. He, W., Liu, G., Cui, W., Tang, Y. (2011). Effect of KIO3 additive on the direct electrosynthesis of K₂FeO₄. Russian Journal of electrochemistry, 47 (11), 1287–1292. doi: 10.1134/s1023193511110097
7. Golovko, D. A., Nefedov, V. G., Girenko, D. V., Cherenkova, O. A. (2013). Elektroliz kontsentririvannykh shelochnykh rastvorov. Soobshchenie 1. Vliyaniye faktorov na razmeryi vyidelyayushchih-sya kislorodnykh puzyrei. Voprosy himii i himicheskoy tehnologii, 4, 144–148.
8. Nefedov, V. G., Atapin, A. G., Golovko, D. A. (2015). Elektroliz koncentririvannykh shelochnykh rastvorov. 2. Faktori, vliyaushie na razmeryi videlyayushchih-sya vodorodnykh pyzirei. Voprosy himii i himicheskoy tehnologii, 2, 51–52.

9. Butyirskaya, E. V., Shaposhnik, V. A., Butyirskiy, A. M. (2004). Sravnitelnyiy analiz struktur gidratnykh obolochek kationov litiya i kaliya. Vestnik Samarskogo Gosudarstvennogo Universiteta. Seriya "Himiya, Biologiya, Farmatsiya", 2, 25–27.
10. Butyirskaya, E. V., Shaposhnik, V. A., Butyirskiy, A. M., Rozhkova, A. G. (2006). Kvantovohimicheskiy raschet gidratatsii soley shelochnykh metallov. Zhurnal strukturnoi himii, 47, 89–93.
11. Klugman, I. Yu. (1999). Ekvivalentnaya elektroprovodnost vodnykh rastvorov. Elektrohimiya, 1 (35), 85–92.
12. Hall, S., McMahon, B. (Eds.) (2005). International Tables for Crystallography. Vol. G: Definition and exchange of crystallographic data. Springer, 598. doi: 10.1107/97809553602060000107
13. Dansurun, D. H. (1994). Opredeleniye chisel gidratatsii nekotorykh otricatelnih ionov mass–spektrograficheskim metodom polekogo ispareniiya ionov iz rastvora. Moscow, 29.
14. Samoylov, O. Ya. (1957). Struktura vodnykh rastvorov elektrolitov i gidratatsiya ionov. Moscow: Izd-vo AN SSSR, 185.
15. Klygman, I. U. (1997). Vyazkost rastvorov silnykh elektrolitov tipa 1:1. Elektrohimiya, 33 (3), 337–345.
16. Volkov, A. I., Zharskiy, I. M. (2005). Bolshoy himicheskii spravochnik. Minsk: Sovremennyya shkola, 608.
17. Shishelova, T. I., Korzyn, N. L., Tolstoi, M. U. (2014). Perspektivy i napravleniya v issledovanii vodi. Mezhdunarodnyi jurnal prikladnykh i fundamentalnykh issledovaniy, 3, 231–231.
18. Sipo, P. M., Hefter, G., May, P. M. (2000). Viscosities and Densities of Highly Concentrated Aqueous MOH Solutions (M⁺ Na⁺, K⁺, Li⁺, Cs⁺, (CH₃)₄N⁺) at 25.0 °C. Journal of Chemical and Engineering Data, 45 (4), 613–617. doi: 10.1021/je000019h
19. Zacepina, G. N. (1998). Fizicheskie svoystva i stryktyra vodi. Moscow: Izd-vo Mosk. In-ta, 184.

INCREASE OF THE REACTIVITY OF A LOW-ACTIVE PART OF LIME SLURRY IN SODA PRODUCTION (p. 33-39)

Musii Tseytlin, Oleksii Shestopalov, Valentina Rayko

The study focuses on the reactivity of low-active parts of lime in the solid phase of lime slurry in soda production. It is determined that low-active lime, that is compounds of calcium with oxides of aluminum, iron, silicon and magnesium, is concentrated in fractions with particles over 60 microns in size. It is found that the maximum rate of decomposition of the low-active components of lime is achieved with the increase of its residence time in the reactor to 40–50 minutes at low pH – 8–8.5.

It is shown that the reactivity of lime milk increases at its dispersion in a vapor-liquid injector due to the cavitation effect. Favorable conditions for the use of low-active parts of lime are possible in recycling the solid unreacted phase in a mixer reactor through a vapor-liquid injector. It is determined that accumulation of the solid recirculating phase in the reactor promotes the removal of supersaturation of the sulfate ion that crystallizes on the surface of the slurry solid particles.

The suggested solutions allow both maximizing the use of low-active lime and reducing the reactor operation time to 10 minutes.

Keywords: soda production, ammonia regeneration, low-active lime, vapor-liquid injector, mixer reactor, recycling.

References

1. Kasikowski, T. (2004). Cleaner production in the ammonia-soda industry: an ecological and economic study. Journal of Environmental Management, 73 (4), 339–356. doi: 10.1016/j.jenvman.2004.08.001
2. Tkach, G. A., Shaporev, V. A., Titov, V. M. (1998). Proizvodstvo sody po maloethodnoj tehnologii. Kharkov: HGPU, 429.
3. Tabunshhikov, N. P. (1974). Proizvodstvo izvesti. Moscow: Himiya, 240.
4. Zubahina, Z. K., Gerasimenko, E. I., Revenko, S. S. (1978). Vzaimodeystvie maloaktivnoy izvesti s hloridom ammoniya. Himicheskaya promyshlennost, 6, 47.

5. Mihajlova, E. N. (2003). Povyshenie stepeni ispol'zovaniya karbonatnogo syr'ja pri regeneracii ammiaka v sodovom proizvodstve. Trudy NIOHIM. Kharkov: NIOHIM, 73, 64–65.
6. Kuzenko, Ju. N., Olesjuk, V. I., Rusina, E. L., Kuhtenkov, K. M. (2003). K voprosu ob usovershenstvovanii tehnologij podachi izvestkovogo moloka na distilljaciju v Sterlitamaskom OAO «Soda». Trudy NIOHIM. Kharkov: NIOHIM, 73, 77–86.
7. Zaikin, A. P., Gridasov, V. N., Annopol'skij, V. F. (2010). Dispergiruvanie izvestkovej suspenzii, ispol'zuej dlja razlozhenija hlorda ammonija na stadii distilljacii v sodovom proizvodstve. Trudy NIOHIM. Kharkov: NIOHIM, 76, 25–31.
8. Ponomarenko, V. V. (2013). Hidrodinamichnyj aktyvator vapnjakovogo moloka. Tsukor Ukrainy, 9, 8–11.
9. Nemirovich, P. M., Zheplinska, M. M., Matiyaschuk, A. M., Homichak, L. M. (2008). Zastosuvannja parokondensacijnoi kavitacii v burjakocukrovomu vyrobnyctvi. Vibracii v tehnicji ta tehnologijah, 2, 102–104.
10. Filonenko, D. V., Shestopalov, O. V. (2013). Analiz napravljenij usovershenstvovanija reaktora-smesitelja otdelenija distilljacii sodovogo proizvodstva. Visnyk Nacional'nogo tehničnogo universytetu «HPI». Serija : Novi rishennja v suchasnyh tehnologijah, 26, 194–198.
11. Filonenko, D. V., Shestopalov, O. V. (2014). Shhodo pytannja rozkladannja hlorydu amoniju v reaktori-zmishuvachi viddilennja dystyljacii sodovogo vyrobnyctva. Visnyk Nacional'nogo tehničnogo universytetu «KhPI», 7, 120–126.
12. Filonenko, D. V., Shestopalov, O. V. (2014). Matematyčeskoe opisanie reaktora-smesitelja otdelenija distilljacii proizvodstva sody i ego komp'juternaja realizacija. Komp'juterne modeljuvannja v himii, tehnologijah i systemah stalogo rozvytku – KMHT-2014. Kyiv, 126–131.
13. Onac, B. P., Forti, P. (2011). Minerogenetic mechanisms occurring in the cave environment: an overview. International Journal of Speleology, 40 (2), 79–98. doi: 10.5038/1827-806x.40.2.1
14. Gainey, R. J., Thorp, C. A., Cadwallader, E. A. (1963). CaSO₄ seeding prevents CaSO₄ scale. Industrial and Engineering Chemistry, 55 (3), 39–43. doi: 10.1021/ie50639a009
15. Lager, G. A. (1984). A crystallographic study of the low-temperature dehydration products of gypsum CaSO₄·0.5H₂O and γ-CaSO₄. Am. Mineral, 69, 910–918.
16. Kasikowski, T., Buczkowski, R., Dejewska, B., Peszynska-Bialczyk, K., Lemanowska, E., Iglinski, B. (2004). Utilization of distiller waste from ammonia-soda processing. Journal of Cleaner Production, 12 (7), 759–769. doi: 10.1016/s0959-6526(03)00120-3
17. Südmalis, M., Sheikholeslami, R. (2000). Coprecipitation of CaCO₃ and CaSO₄. Canadian Journal of Chemical Engineering, 78 (1), 21–31. doi: 10.1002/cjce.5450780106
18. Höfling, V., Augustin, W., Bohnet, M. (2004). Crystallization fouling of the aqueous two-component system CaSO₄/CaCO₃. International Journal of Transport Phenomena, 6 (2), 99–109.
19. Filonenko, D. V., Rayko, V. F., Shestopalov, O. V. (2014). Snizhenie inkrustacii distillera putem snjatija peresyshhenija po sul'fatu v reaktore-smesitele otdelenija distilljacii. Visnyk NTU «HPI». Serija: Mehaniko-tehnologični systemy ta kompleksy, 60 (1102), 102–110.

INVESTIGATION OF DIFFUSION MECHANISMS DURING PERVAPORATION OF MOLECULES OF ORGANIC SOLVENTS IN POLYMER MEMBRANES (p. 40-47)

Inessa Burtina, Otar Gachechiladze

With the help of the original experimental setup using a pervaporation process, parameters which allow to indirectly determine the mechanisms and models of diffusion of solvent molecules in the polymer membrane body were investigated.

The time dependencies of these parameters during pervaporation were obtained for both unsaturated and saturated membranes. It was shown that the mechanism of diffusion of the solvent molecules in the polymer membrane for the unsaturated membranes consists of two components: diffusion through thermal activation of the system and diffusion occurring through the free volume. Moreover, the rate of thermally activated diffusion is much higher and is observed in the beginning of the pervaporation process, whereas the diffusion using the free-volume model begins to emerge in the middle of the process and runs more slowly, but it is a source of transfer of the majority of solvent molecules. For saturated membranes, the main transfer mechanism is diffusion through the structural voids of the polymer. These results allow to interpret diffusion models in polymer membranes.

Keywords: polymer, membrane, pervaporation, thermal activation, free volume, diffusion, voids, jump technology.

References

1. Burtina, I. A., Hachechiladze, O. O. (2014). Transfer processes in polymer membranes. Part 4. Eastern-European Journal of Enterprise Technologies, 2/6(68), 4–11. doi: 10.15587/1729-4061.2014.23383
2. Cranc, J., Park, G. S. (1968). Methods of measurement in Diffusion in Polymers. New York: Academic Press, 1–39.
3. Marion, K. Buckley-Smith (2006). The Use of Solubility Parameters to select membrane materials for Pervaporation of organic mixtures. The University of WAIKATO. Hamilton, New Zealand, 18–56.
4. Burtina, I. A., Hachechiladze, O. O. (2013). Transfer processes in polymer membranes. Part 3. Eastern-European Journal of Enterprise Technologies, 6/6(66), 4–7. Available at: <http://journals.urau.ua/eejet/article/view/18713/17124>
5. Burtina, I. A., Hachechiladze, O. O. (2014). Experimental determination of possible models of diffusion during pervaporation process in polymer membranes. Eastern-European Journal of Enterprise Technologies, 5/6(71), 26–32. doi: 10.15587/1729-4061.2014.27661
6. Binning, R. C., Lee, R. J., Jennings, J. F., Martin, E. C. (1961). Separation of Liquid Mixtures by Permeation. Industrial & Engineering Chemistry, 53 (1), 45–50. doi: 10.1021/ie50613a030
7. Shao, D., Huang, R. Y. M. (2006). Polymeric membrane Pervaporation. Department of Chemical Engineering University of Waterloo, Ont, Canada, 162–175.
8. Koros, W. J. (1990). Barrier Polymers and Structures. ACS Publications. Washington: D. C., 1, 1–21. doi: 10.1021/bk-1990-0423.ch001
9. Meares, P. (1965). Polymers: Structure and Bulk Properties. London. D. Van Nostrand Co. LTD, 12, 381.
10. Brandt, W. W. (1959). Self – Diffusion and Inter diffusion in Polymers. Journal of Physical Chemistry, 63, 1080–1088.
11. DiBenedetto, A. T., Paul, D. R. (1964). Diffusion in amorphous polymers. Journal of Polymer Science, A (2), 1001–1015.
12. Turnbull, D., Cohen, M. H. (1961). Free-volume model of the amorphous phase. Journal of Chemical Physics, 34, 120–131. doi: 10.1063/1.1731549
13. Cohen, M. H., Turnbull, D. (1959). Molecular transport in liquids and glasses. Journal of Chemical Physics, 31 (5), 1164–1169. doi: 10.1063/1.1730566
14. Duda, J. L., Vrentas, J. S., Ju, S. T., Liu, H. T. (1982). Prediction of diffusion Coefficients for polymers solvent systems. AIChE Journal, 28 (2), 279–285. doi: 10.1002/aic.690280217
15. Reis, R. A., Oliveira, J., Nobrega, R. (2001). Diffusion Coefficients in Polymer – Solvent Systems for Highly Concentrated Polymer Solutions. Brazilian Journal of Chemical Engineering, 18 (4), 221–232. doi: 10.1590/s0104-66322001000400003
16. Burtina, I. A. (2013). Transfer processes in polymer membranes. Part 2. Eastern-European Journal of Enterprise Technologies, 2/11(61), 41–44. Available at: <http://journals.urau.ua/eejet/article/view/11731/9873>

DEVELOPMENT AND RESEARCH OF THE METHOD FOR PROPANE-BUTANE GAS CONVERSION INTO METHANOL (p. 48-52)

Alexei Tselischev, Marina Loriya,
Peter Eliseev, Vladimir Nosach,
Ayodezhi Adebayo Ijagbujii, Roman Fedotov

The schematic diagram and the physicochemical study of the process of propane-butane gas conversion into methanol in the presence of hydrogen peroxide solution were proposed. In general, the process of methanol formation is described in the reaction: $(C_3H_8 + C_4H_{10}) + H_2O_2 \rightarrow 2CH_3OH + C_2H_4 + C_3H_6 + H_2$. It should be noted that other alcohols and other oxygen-containing products except methanol were not found in all presented experiments. The analysis of the liquid reaction products was carried out by chromatography in a specialized laboratory by the instruments LHM and "Tsvet 500" using the 3 m long nozzle "Polisorb". The analysis of experimental data led to the conclusion that direct conversion of propane (butane) into methanol allows to achieve the conversion level of ~ 10 % in one pass. The reactor design and the flow diagram of the laboratory setup for methanol synthesis by the cavitation method in the total absence of reaction byproducts were developed. The applied cavitation method is extremely promising for using propane-butane gas as a raw material in a fundamentally new production of methanol.

Keywords: hydroxyl radical, propane-butane gas, methyl radical, cavitation, methanol.

References

- Rozovskij, A. Ja., Lin, G. I. (1990). *Teoreticheskie osnovy processa sinteza metanola*. Moscow: Himija, 272.
- Lebedev, N. N. (1988). *Himija i tehnologija osnovnogo organicheskogo i neftehimicheskogo sinteza*. Moscow: Himija, 592.
- Shilov, A. E., Shul'pin, G. B. (2000). *Activation and catalytic reactions of saturated hydrocarbons in the presence of metal complexes*. Dordrecht: Kluwer Academic Publishers, 232.
- Fokins, A. A., Schreiner, P. R. (2002). Selective alkane transformations via radicals and radical cations: Insight into the activation step from experimental theory. *Chemical Reviews*, 102 (5), 1551–1594. doi: 10.1021/cr000453m
- Arutjunov, N. S., Krylov, L. V. (2005). Okislitel'naja konversija metana uspehi himii. *Uspehi himii*, 74 (12), 1216–1274.
- Zamilov, M. F., Godin, S. (2012). Investigation of hydrodynamic cavitation as a means of natural crude oil and synthetic biofuel upgrading. *Quantum Potential Corporation*, 1–21.
- Celishhev, A. B., Zaharova, O. I., Loriya, M. G., Zaharov, I. I. (2009). Fiziko-himicheskie osnovy foto-avtokataliticheskogo processa okislenija metana v metanol. *Voprosy himii i him-tehnologii*, 4, 43–55.
- Celishhev, A. B., Zaharov, I. I., Loriya, M. G., Idzhagbudzhi, A. A. (2012). Analiz sposobov aktivacii metana v «mjagkih» uslovijah. *Voprosy himii i himicheskoj tehnologii*, 2, 39–44.
- Zakharov, I. I., Ijagbujii, A. A., Tselishev, A. B., Loriya, M. G., Fedotov, R. N. (2015). The new pathway for methanol synthesis: Generation of methyl radicals from alkanes. *Journal of Environmental Chemical Engineering*, 3 (1), 405–412. doi: 10.1016/j.jece.2014.08.008
- Celishhev, A. B., Loriya, M. G., Zaharov, I. I. (2011). Analiz fiziko-himicheskij metodov poluchenija gidroksil'nogo radikala. *Visnik Nacional'nogo tehnicnogo universitetu «KhPI»*, 65, 111–124.
- Aseev, D. G., Kenzhin, R. M., Stojanovskij, V. O., Batoeva, A. A., Volodin, A. M. (2011). Detektirovanie ON radikalov v processe gidrodinamicheskij kavitacii i v sonoreaktore ljuminiscentnymi metodami. *Sovremennaja himicheskaja fizika. Tuapse (Rossija): Iz-vo MGU*, 81–83.
- Celishhev, A. B., Zaharov, I. I., Loriya, M. G., Idzhagbudzhi, A. A., Eliseev, P. I., Nosach, V. A. (2014). Motornye topliva: kavitacionnyj sposob povyshenija ih kachestva. *Himichna promyslovist' Ukrainy*, 121 (2), 39–42.

- Loriya, M. G., Ijagbujii, A. A., Tselischev, A. B., Zakharov, I. I. (2013). Autocatalytic photo-oxidation process of C₃–C₄ fraction to methanol. *Advanced Materials Research*, 660, 51–56. doi: 10.4028/www.scientific.net/amr.660.51
- NIST Chemistry: WebBook. Available at: <http://webbook.nist.gov/chemistry>.
- Landau, L. D., Lifshic, E. M. (1986). *Teoreticheskaja fizika. Gidrodinamika*. Moscow: Nauka, 6, 503–506.
- Abramovich, G. N. (1991). *Prikladnaja gazovaja dinamika*. Moscow: Nauka, 600.
- Han, L. B., Tsubota, S., Harula, M. (1995). Effect of the addition of nitrogen dioxide on the gas-phase partial oxidation of methane with oxygen under normal pressure. *Chemistry Letters*, 24 (10), 931–932. doi: 10.1246/cl.1995.931
- Taylor, C. E. (2003). Methane conversion via photo-catalytic reaction. *Catalysis Today*, 84 (1–2), 9–15. doi: 10.1016/s0920-5861(03)00295-5

DEVELOPMENT OF A MATHEMATICAL MODEL OF THE PROCESS OF BIOLOGICAL TREATMENT OF GASEOUS EMISSIONS (p. 53-61)

Anna Bakhareva, Oleksii Shestopalov,
Olesya Filenko, Tetyana Tykhomirova

In experimental studies, the kinetic characteristics of methane oxidation by the immobilized microbial association in gaseous emissions were determined. The obtained quantitative values of specific oxidation rate of CH₄ indicate a technological possibility of using the fluidized-bed bioreactor as the stage of the installation, designed for treatment of gaseous emissions from methane in drainage networks. It was found that the oxidation rate of CH₄ varied from 60 ml/g·h in the region of minimum concentrations of CH₄ in the medium to a maximum value of 260 ml/g·h. The presence of dependence of the specific oxidation rate of methane on its concentration in air was revealed.

Based on experimental studies, a mathematical description of the processes occurring in the reactor due to changes in the concentration of incoming pollutants was developed. It was found that persistent cyclic changes in the concentration at the bioreactor inlet will lead to the persistent cycle of changes in the pollution concentration at the outlet. The results of checking calculations show the transformation of fairly smooth concentration variations of the methane at the bioreactor inlet into dramatic changes in its concentration at the end of the biotreatment process, consideration of which is necessary in designing gas-treatment equipment.

Keywords: mathematical model, biological treatment of emissions, specific oxidation rate, concentration, harmful substance, bioreactor.

References

- Myakenkiy, V. I., Kurdish, I. K. (1991). *Mikrobiologicheskoe okislenie metana ugolnyih shaft. Kyiv, Nauk. dumka*, 148.
- Peinado, P. A., Moreno, J. J., Villaba, J. M., Gonzales-Reyes, J. A., Ortega, J. M., Mauricio, J. C. (2006). A new immobilization method and their application. *Enzyme Microb Tech*, 40, 79–84.
- Abbasi, T., Abbasi, S. A. (2011). Sources of Pollution in Rooftop Rainwater Harvesting Systems and Their Control. *Critical Reviews in Environmental Science and Technology*, 41 (23), 2097–2167. doi: 10.1080/10643389.2010.497438
- Yang, J., Spanjers, H., Jeison, D., Van Lier, J. B. (2013). Impact of Na⁺ on Biological Wastewater Treatment and the Potential of Anaerobic Membrane Bioreactors: A Review. *Critical Reviews in Environmental Science and Technology*, 43 (24), 2722–2746. doi: 10.1080/10643389.2012.694335
- Papirio, S., Villa-Gomez, D. K., Esposito, G., Pirozzi, F., Lens, P. N. L. (2013). Acid Mine Drainage Treatment in Fluidized-Bed Bioreactors by Sulfate-Reducing Bacteria: A Critical Review. *Critical Reviews in Environmental Science and Technology*, 43 (23), 2545–2580. doi: 10.1080/10643389.2012.694328

6. Oturan, M. A., Aaron, J.-J. (2014). Advanced Oxidation Processes in Water/Wastewater Treatment: Principles and Applications. A Review. *Critical Reviews in Environmental Science and Technology*, 44 (23), 2577–2641. doi: 10.1080/10643389.2013.829765
7. Kennes, C., Rene, E. R., Veiga, M. C. (2009). Bioprocesses for air pollution control. *Journal of Chemical Technology & Biotechnology*, 84 (10), 1419–1436. doi: 10.1002/jctb.2216
8. Shestopalov, O. V., Pitak, I. V. (2014). Analysis of existent processes and devices of bioscrubbing gas emissions. *Technology audit and production reserves*, 3, 5, 49–52.
9. Seedorf, J. (2013). Biological exhaust air treatment systems as a potential microbial risk for farm animals assessed with a computer simulation. *Journal of the Science of Food and Agriculture*, 93 (12), 3129–3132. doi: 10.1002/jsfa.6106
10. Iranpour, R., Cox, H. H. J., Deshusses, M. A., Schroeder, E. D. (2005). Literature review of air pollution control biofilters and biotrickling filters for odor and volatile organic compound removal. *Environmental Progress*, 24 (3), 254–267. doi: 10.1002/ep.10077
11. Mohammad, B. T., Veiga, M. C., Kennes, C. (2007). Mesophilic and thermophilic biotreatment of BTEX-polluted air in reactors. *Biotechnology and Bioengineering*, 97 (6), 1423–1438. doi: 10.1002/bit.21350
12. Rojo, N., Muñoz, R., Gallastegui, G., Barona, A., Gurtubay, L., Prenafeta-Boldú, F. X., Elias, A. (2012). Carbon disulfide biofiltration: Influence of the accumulation of biodegradation products on biomass development. *Journal of Chemical Technology & Biotechnology*, 87 (6), 764–771. doi: 10.1002/jctb.3743
13. Malhautier, L., Cariou, S., Legrand, P., Touraud, E., Geiger, P., Fanlo, J. L. (2014). Treatment of complex gaseous emissions emitted by a rendering facility using a semi-industrial biofilter. *Journal of Chemical Technology and Biotechnology*. doi: 10.1002/jctb.4593
14. Engesser, K.-H., Plaggemeier, T. (2008). Microbiological Aspects of Biological Waste Gas Purification. *Biotechnology: Environmental Processes III*, 11c, 275–302.
15. Banerle, V., Fisher, H., Baroltki, D. (1986). Biologische Abluftreinigung mit Hilfe eines membranartigen Permeationsreactors. *Stand-Reinhaltung der Luft*, 46 (5), 233–235.
16. Don, T. A. (1983). Biofiltratie – ein milieu lijhe effectieve en relatief goedje manier van luchtreinigung. *Innovative*, 13 (53), 4–5.
17. Gabrieland, D., Deshusses, M. A. (2004). Technical and economical analysis of the conversion of a full-scale scrubber to a biotrickling filter for odour control. *Water Science and Technology*. Portland: IWAPublishing, 4, 309–318.
18. Londong, J. (1992). Strategies for optimized nitrate reduction with primary denitrification. *Water Science and Technology*. Portland: IWAPublishing, 5-6, 1087–1096.
19. Sotomayor, O. A. Z., Park, S. W., Garcia, C. (2001). A simulation benchmark to evaluate the performance of advanced control techniques in biological wastewater treatment plants. *Brazilian Journal of Chemical Engineering*, 18 (1). doi: 10.1590/s0104-66322001000100008
20. Wentzel, M. C., Ekama, G. A., Marais, G. V. R. (1992). Processes and modeling of nitrification-denitrification biological excess phosphorus removal systems – a review. *Water Science and Technology*. Portland: IWAPublishing, 6, 59–82.
21. Jan, R., Ng, V. L., Chen, X. G., Geng, A. L., Gouhd, W. D., Duan, H. Q., Ling, D. T., Koe, L. C. (2004). Bath experiment on H₂S degradation by bacteria immobilised on activated carbons. *Water Science and Technology*. Portland: IWAPublishing, 4, 299–308.
22. Barbosa, V. L., Dufol, D., Callan, J. L., Sneath, R., Stuetz, R. M. (2004). Hydrogen sulphide removal by activated sludge diffusion. *Water Science and Technology*. Portland: IWAPublishing, 4, 199–205.
23. Krichkovska, L. V., Shestopalov, O. V., Bakhareva, G. Y., Slis, K. V. (2013). Prozesi ta aparati biologichnoy ochistki ta dezodorazii gazopovitryanih vikidiv. Kharkiv: NTU «KhPI», 200.
24. Krichkovska, L. V., Vaskovez, L. A., Gurenko, I. V. et al. (2014). Proektni rishennya u rozrobzi aparativ biologichnoy ochistki gazopovitryanih vikidiv. Kharkiv: NTU «KhPI», 208.
25. Bahareva A. Yu., Shestopalov O. V., Semenov E. O., Bukatenko N. O. (2015). Macrokinetic mathematical model development of biological treatment process of gasiform emissions. *Science-Rise*, 2/2(7), 12–15. doi: 10.15587/2313-8416.2015.37057
26. Kuznetsov, S. I., Dubinina, G. A. (1989). *Metodyi izucheniya vodnyih mikroorganizmov*. Moscow: Nauka, 286.

STUDY OF WATER QUALITY IN THE DISTRIBUTION NETWORK OF THE CENTRALIZED WATER SUPPLY SYSTEM IN THE CITY OF LVIV (p. 62-70)

Oksana Matsiyevska

The problem of water quality change in the distribution network of the centralized water supply system in the city of Lviv is considered, and some results of our research in this area are presented. The main purpose of the research is identifying the main factors of water quality change in the distribution network of the city. During the research, collection, analysis and synthesis of data on technological parameters of the distribution network were carried out, and the data of the analysis of eight water samples taken from residents in different districts of the city of Lviv were used. The results suggest that the increase in total hardness and alkalinity of water is caused by mixing of water from different water intakes in the distribution network; water oxidability growth - by probable accidents with entry of organic pollutants into the pipeline; increase in the concentration of total iron - by corrosion of long-life and great-length sites of the distribution network; reduction of free residual chlorine - by its spending on oxidation of chemicals contained in water and elimination of pathogenic microorganisms of biofilms on the inner walls of pipes. In some water samples, an excess of requirements of physiological full value of the mineral composition of drinking water by total hardness and probable value of solids, due to their increased values in the source water was observed. However, the values of water quality indices in the studied sites do not exceed the norms applicable to drinking water. To prevent the deterioration of the drinking water quality, and, therefore, the health of residents, the introduction of modern systems of monitoring and modeling of water quality change in the distribution network is recommended.

Keywords: water quality, physiological full value, distribution network, centralized water supply, human health.

References

1. Matsiyevska, O. O. (2013). Yakist pytnoi vody, shcho nadkhodyt u merezhu tsentralizovanoho vodopostachannia m. Lviv. *Kharchova nauka i tekhnolohiia*, 1 (22), 87–89.
2. Summerscales, I. M., McBean, E. A. (2011). Application of risk assessment tools to small drinking water systems in British Columbia. *Water Quality Research Journal of Canada*, 46 (4), 332–344. doi: 10.2166/wqrjc.2011.102
3. Malm, A., Axelsson, G., Lars, B., Ljungqvist, J., Forsberg, B., Bergstedt, O., Pettersson, Th. J.R. (2013). The association of drinking water treatment and distribution network disturbances with Health Call Centre contacts for gastrointestinal illness symptoms. *Water Research*, 47 (13), 4474–4484. doi: 10.1016/j.watres.2013.05.003
4. Helbling, D. E., VanBriesen, J. M. (2009). Modeling Residual Chlorine Response to a Microbial Contamination Event in Drinking Water Distribution Systems. *Journal of Environmental Engineering*, 135 (10), 918–927. doi: 10.1061/(asce)je.1943-7870.0000080
5. Khadse, G. K., Kalita, M., Pimpalkar, S. N., Labhsetwar, P. K. (2011). Drinking water quality monitoring and surveillance for safe water supply in Gangtok, India. *Environmental Monitoring and Assessment*, 178 (1-4), 401–414. doi: 10.1007/s10661-010-1699-6
6. Fallahzadeh, R. A., Gholami, M., Madreseh, E., Ghaneian, M. T., Farahzadi, M. H., Askarnejad, A. A., Sadeghi, Sh. (2015). Comparison of Using an Electronic System and Conventional Moni-

- toring Method for Monitoring the Quality of Drinking Water and Defects Discovery in Rural Area Water Distribution Network of Abarkouh, Iran. *Health*, 07 (01), 35–40. doi: 10.4236/health.2015.71005
7. Goyal, R. V., Patel, H. M. (2014). Analysis of residual chlorine in simple drinking water distribution system with intermittent water supply. *Applied Water Science*, 5 (3), 311–319. doi:10.1007/s13201-014-0193-7
 8. Khovanskyi, S. O., Nenia, V. H. (2010). Systemnyi analiz kompleksu podachi i rozpodilu vody v zhytlovo-komunalnomu hospodarstvi. *Eastern-European Journal of Enterprise Technologies*, 4 (4(46)), 56–59. Available at: <http://journals.urau.ua/eejet/article/view/2967/2770>
 9. Mandel, P., Maurel, M., Chenu, D. (2015). Better understanding of water quality evolution in water distribution networks using data clustering. *Water Research*, 87, 69–78. doi: 10.1016/j.watres.2015.08.061
 10. Wasowski, J., Kowalski, D., Kowalska, B., Kwietniewski, M., Zawilska, M. (2012). Water Quality Variations in Cement-lined Water-pipe Networks. *Ochrona Srodowiska*, 1 (38), 53–58.
 11. Girol, M. M., Kovalski, D., Khomko, V. Ye., Girol, A. M. (2008). Problemy yakosti vody v vodoprovodnyh mrezha. *Vodopostachannya ta vodovidvedennya*, 2, 15–21.
 12. Niu, Z. B., Wang, Y., Zhang, X. J., Chen, C., Wang, S. H. (2007). Effect on iron release in drinking water distribution systems. *Huan Jing Ke Xue*, 28 (10), 2270–2274.
 13. Lashena, M. R., Sharabyb, C. M., El-Kholyb, N. G., Elsherifa, I. Y., El-Wakeela, S. T. (2008). Factors influencing lead and iron release from some Egyptian drinking water pipes. *Journal of Hazardous Materials*, 160 (2-3), 675–680. doi: 10.1016/j.jhazmat.2008.03.040
 14. Yonov, V. S. (2005). O roly trub v reforme ZhKKh. *Santekhnika*, 1, 50–53.
 15. Dong, J., Wang, G., Yan, H., Xu, J., Zhang, X. (2015). A survey of smart water quality monitoring system. *Environmental Science and Pollution Research*, 22 (7), 4893–4906. doi: 10.1007/s11356-014-4026-x
 16. Lee, A., Francisque, A., Najjaran, H., Rodriguez, M. J., Hoorfar, M., Imran, S. A., Sadiq, R. (2012). Online monitoring of drinking water quality in a distribution network: a selection procedure for suitable water quality parameters and sensor devices. *International Journal of System Assurance Engineering and Management*, 3 (4), 323–337. doi: 10.1007/s13198-012-0133-6
 2. Dolmatova, L. A., Guseva, M. A. (2004). Organicheskie veshchestva v snegovom pokrove pribrezhnoj chasti r. Barnaulki. *Polzunovskij Vestnik*, 2, 150–154.
 3. Jurchenko, V. A., Mel'nikova, O. G., Jachnik, M. V. (2014). Issledovaniya poverhnostnyh stochnyh vod s avtomobil'nyh dorog i dorozhno-infrastrukturnykh kompleksov. *Avtoshljahovyk Ukrainy*, 5 (241), 43–47.
 4. Polkowska, Z., Skarzyńska, K., Dubiella-Jackowska, A., Staszek, W., Namieśnik, J. (2007). Evaluation of pollutant loading in the runoff waters from a major urban highway (Gdansk Beltway, Poland). *Global NEST Journal*, 9 (3), 269–275.
 5. Golwer, A., Schmoll, O., Howard, G., Chilton, J., Chorus, I. (2006). Traffic and transport: Control and protection. World Health Organization. *Protecting Groundwater for Health: Managing the Quality of Drinking-water Sources*. London: IWA Publishing, 653–662.
 6. Pravyta pryymannya stichnykh vod pidpryemstv u komunal'ni ta vidomchi systemy kanalizatsiyi naselenykh punktiv Ukrainy № 37 vid 19.02.02. Available at: <http://zakon5.rada.gov.ua/laws/show/z0403-02>
 7. Mostepan, E. V. (2009). Issledovanie vliyanija livnevnyh vod s vodosbornykh territorij goroda na sostojanie vodnyh obektov. *Jeftektivnye materialy, tehnologii, mashiny i oborudovanie dlja stroitel'stva i jekspluatacii sovremennyh transportnyh sooruzhenij*. Belgorod, 261–263.
 8. Bruen, M., Johnston, P., Quinn, M., Desta, M., Higgins, N., Bradley, C., Burns, S. (2006). Impact assessment of highway drainage on surface water quality. Wexford: Environmental Protection Agency, 272.
 9. Perevoznikov, B. F., Il'ina, A. A. (2002). Sooruzhenija systemy vodootvoda s proezzhej chasti avtomobil'nyh dorog. Moscow, 60.
 10. Il'ina, A. A. (2004). Meroprijatija po snizheniju urovnja zagriznenija livnevnyh i talyh poverhnostnyh stokov s avtomobil'nyh dorog. *Novosti v dorozhnom dele*. Moscow, 3.
 11. *Avtomobil'nye dorogi* (1987). SNiP 2.05.02-85. Moscow: GUP CPP.
 12. Brannvall, E. (2007). Improvement of storm water runoff treatment system with natural mineral sorbent. *Geologija*. Vilnius, 59, 72–76.
 13. Piguet, P. (2007). Road runoff over the shoulder diffuse infiltration. Real-scale experimentation and optimization. Suisse, 277.
 14. *Kanalizatsiya* (2013). Zovnishni mrezhi ta sporudy. Osnovni polozhennya proektuvannya. DBN V.2.5.-75:2013. Kyiv, 128.
 15. Lur'e, Ju. Ju., Rybnikova, A. I. (1974). *Himicheskij analiz proizvodstvennyh stochnyh vod*. Moscow: Himija, 336.
 16. Turlough, F. G. (1999). The extraction of aged polycyclic aromatic hydrocarbon (PAH) residues from a clay soil using signification and a soxhlet procedure: a comparative study. *Journal of Environmental Monitoring*, 1 (1), 63–67. doi: 10.1039/a807307d
 17. Nazarov, A. V. (2007). Vlijanie neftjanogo zagriznenija pochvy na rastenija. *Vestnik permskogo universiteta*, 5 (10), 14–16.
 18. Muhachjov, V. A. (2007). Planirovanie i obrabotka rezul'tatov jeksperimenta. Tomsk: Tomskij gosudarstvennyj universitet sistem upravlenija i radioelektroniki, 118.

RESEARCH OF MECHANICAL TREATMENT OF STORMWATER RUNOFFS FORMED ON OBJECTS OF AUTOMOBILE AND ROAD COMPLEX (p. 71-77)

Valentina Iurchenko, Oksana Melnikova,
Anna Bakhareva, Mariia Yachnyk

Experimental research revealed that the concentration of oil products in surface wastewater, formed in winter and summer on the objects of automobile and road complex – gas stations, car parks and roads, greatly exceeds the MPC for discharge into water bodies. In stormwater runoffs at gas stations and car parks, the concentration of oil products has higher values than in surface wastewater, formed on the roads, despite the low traffic level. In surface wastewater, formed at gas stations and car parks, conditionally heavy fractions of oil products, more efficiently removable during settling prevail. The effect of surface wastewater treatment from oil products (maximum 80 %) depends on several parameters, the most significant of which is the concentration of suspended solids.

Keywords: automobile and road complex, objects of road infrastructure, stormwater runoffs, mechanical treatment, oil products.

References

1. Pshenin, V. N., Kovalenko, V. I. (2007). Zagriznenie livnevnyh stokov s avtomobil'nyh dorog. *Vestnik INZhJeKONA*, 6 (19), 140–145.

INFLUENCE OF REDOX POTENTIAL OF THE MEDIUM ON PHOSPHORUS MIGRATION IN SLUDGE LIQUOR (p. 78-84)

Valentina Yurchenko, Oleksandr Smyrnov, Anna Bakhareva

The characteristics of the influence of the redox potential of the medium on the process of biological wastewater treatment from phosphates were investigated. The research was aimed at identifying new control actions to intensify wastewater treatment from phosphorus compounds and protect natural water bodies from pollution. The quantitative relationships were determined in the laboratory simulation of aerobic and anaerobic cultivation conditions of sludge liquor, as well as when examining existing sewage treatment facilities. It was found that the lower the redox potential values in the anaerobic wastewater treat-

ment zones, the more intensive the phosphate mobilization from activated sludge into the aqueous medium and, consequently, the more phosphates will be immobilized from the aqueous medium into sludge under aerobic conditions. The obtained quantitative characteristics allow to use the redox potential of wastewater (controlled variable) to manage and intensify the deep removal of phosphorus compounds during biological wastewater treatment.

Keywords: wastewater, activated sludge, phosphates, redox potential, volatile fatty acids.

References

1. Chemical-physical methods for the removal of phosphorus from wastewater (2004). Standard ATV-DVWK-A 202E, 25.
2. Zaljotova, N. A. (2011). Osobennosti himicheskogo udaleniya fosfora pri biologicheskoy ochildki stochnyh vod. *Vodosnabzhenie i sanitarnaya tehnika*, 11, 40–46.
3. Henze, M., Harremoës, P., Janssen, P. M. J., Arvin, E. (1995). *Wastewater treatment – biological and chemical processes*. Springer Verlag.
4. Ruston, G., Fort, C. (2012). Engineering considerations for phosphorus removal. IWEA O&M: seminar, 14.
5. Barnard, J. L. (2006). Biological nutrient removal: where we have been, where we are going? *Proceedings of the Water Environment Federation (WEFTEC)*, 13, 1–25. doi: 10.2175/193864706783710578
6. Imhoff, K. R. (1996). Surface water quality objectives and standards in relation to improvement projects. *Metropolitan Areas and Rivers : general reports*. Roma: Maggio, 1, 33–57.
7. Randall, C. W., Barnard, J. L., Stensel, H. D. (1992). Design and retrofit of wastewater treatment plants for biological nutrient removal. *Water quality management library*. Lancaster: TECHNOMIC Publication, 5, 420.
8. Keller, J., Subramaniam, K., Gosswein, J., Greenfield, P. F. (1997). Nutrient removal from industrial wastewater using single tank sequencing batch reactors. *Water Science and Technology*, 35 (6), 137–144. doi: 10.1016/s0273-1223(97)00104-2
9. Mahdy, A. M., Elkhatib E. A., Fathi, N. O. (2007). Drinkig water treatment residuals as an amendment to alkaline soils: Effects on the growth of corn and phosphorus extractabilit. *International Journal of Environmental Science and Technology*, 4 (4), 489–496. doi: 10.1007/bf03325985
10. Griffiths, P. C., Stratton, H. M., Seviour, R. J. (2002). Environmental factors contributing to the “G bacteria” population in full-scale EBPR plants. *Water Science and Technology*, 46 (4-5), 185–192.
11. de-Bashan, L. E., Bashan, Y. (2004). Recent advances in removing phosphorus from wastewater and its future use as fertilizer (1997–2003). *Water Research*, 38 (19), 4222–4246. doi: 10.1016/j.watres.2004.07.014
12. Banu, R. J., Do, K. U., Yeom, I. T. (2008). Phosphorus removal in low alkalinity secondary effluent using alum. *International Journal of Environmental Science and Technology*, 5 (1), 93–98. doi: 10.1007/bf03326001
13. Chemical-Physical Methods for the Removal of Phosphorus from Wastewater (2004). Standard ATV-DVWK-A 202E, 25.
14. Christensson, M. (1997). Enhanced biological phosphorous. Department of Biotechnology, Lund University. Lund, Sweden, 79.
15. Janssen, P. M. J., Meinema, K., van der Roest, H. F. (2002). *Biological phosphorus removal: manual for design and operation*. London: IWA Publishing, 26.
16. Deinema, M. H., Loosdrecht, M. V., Scholten, A. (1985). Some physiological characteristics of *Acinetobacter* spp. accumulating large amounts of phosphate. *Water Science and Technology*, 17, 119–125.
17. He, S., Gu A. Z., McMahon, K. D. (2006). Fine-scale differences between *Accumulibacter*-like bacteria in enhanced biological phosphorous removal activated sludge. *Water Science and Technology*, 54 (1), 111–117. doi: 10.2166/wst.2006.378
18. Blackall, L. L., Crocetti, G. R., Saunders, A. M., Bond, P. L. (2002). A review and update of the microbiology of enhanced biological phosphorous removal in wastewater treatment plants. *Antonie Van Leeuwenhoek*, 81, 681–691.
19. Cloete, T. E., Oosthuizen, D. J. (2001). The role of extracellular exopolymers in the removal of phosphorous from activated sludge. *Water Resources*, 35 (15), 3595–3598. doi: 10.1016/s0043-1354(01)00093-8
20. Lur'e, Y. Y., Rybnikova, A. I. (1974). *Chimicheskiy analiz proizvodstvenih stochnih vod*. Moscow: Chimiya, 336.