In nature, magnesium compounds are distributed in the form of such minerals as dolomite, magnesite, serpentinite, brucite, and in the form of such solutions as sea and ocean waters, salt lakes, bischofite. The main popular magnesium compounds are its chloride, hydroxide, and oxide. The most common technology for obtaining magnesium hydroxide is based on the action of alkaline reagents on solutions of magnesium. However, the technology has significant drawbacks. The main issue is that magnesium hydroxide cannot be obtained free of impurity metal ions, and the sediment itself has an extremely low filtration rate from the solution in which it is formed.

Magnesium hydroxide is an excellent sorbent collector for the ions of most metals, so it is possible to utilize this property of absorption of magnesium hydroxide to produce other highly pure compounds, for medicine and pharmacopeia. This paper investigates the processes of alkaline hydrolysis of magnesium chloride and the formation of magnesium hydroxide sediment, the kinetics of processes, as well as the properties of sediments. Empirical formulas for sediment production parameters with a high filtration coefficient that exceeds the known values of $1 \times 10^4$ have been derived. The study reported here has revealed the possibility of obtaining in a relatively simple way highly pure chloride, hydroxide, and magnesium oxide, sodium chlorides, potassium, calcium, in which the impurity 3d metals content does not exceed the value as a percentage of $(1-4) \times 10^{-7}$. Due to the properties of highly pure magnesium compounds to the sorption of metal ions, it is possible to create reference samples of chemical elements for the needs of metrology. For example, to create MR phantoms to verify measurements and check the operation of magnetic resonance imaging (MRI) in medical institutions. Especially pure magnesium hydroxide can be used for instrumental methods of analytical chemistry (X-ray fluorescent, neutron-activation methods) when fabricating chemical references for testing laboratories.

Keywords: magnesium compounds, alkaline hydrolysis of magnesium chloride, magnesium hydroxide, sorption of metal ions.
Taking into consideration the importance of the technology of obtaining magnesium hydroxide, an analysis of the state of scientific and experimental studies was carried out on the chemical aspects of obtaining it from solutions of its salts (mainly magnesium chloride). Technologies for obtaining high-purity magnesium hydroxide and magnesium oxide are rapidly developing in China regarding the use of magnesium hydroxide with a nanostructure to extract terbium lanthanoid from solutions of strategic importance. In Russia, where there are various raw materials sources of magnesium compounds, they pay great attention to their production [5]. Perfect technologies have been developed and large production facilities have been organized, which are described with a significant amount of information in publications. However, along with the existing descriptions of a number of technologies, there are no descriptions of chemical processes used to ensure the purity of reagents, types of such reagents and their features, and choice justification. Paper [6] reports existing technologies in Russia, where the production of magnesium products has a large volume and is in demand, but there are no data on the methods of filtration of magnesium hydroxide and research into the degree of purification of reagents, raw materials. It is not specified how the issues of obtaining high purity products and the peculiarities of filtration in technology are solved. The exception is data from [5], which indicates the use of the most accessible alkali, as a solution of lime (calcium hydroxide). However, it is known that the alkaline reagent calcium hydroxide also does not provide high-quality sediment of magnesium hydroxide, its purity. Fundamentally, it is impossible to obtain lime free from impurities, although most technologies are based on the use of lime as an alkaline reagent. There are needs for obtaining heat-resistant plastics and fire-resistant products from them when using magnesium hydroxide and its oxide in the form of structures with nanoscale particles [6]. At the same time, there are no requirements for the technology and its features for obtaining particles of hydroxide and magnesium oxide, the effect of their size for fillers in plastics with refractory properties.

It is indicated that the technology of nanostructured magnesium hydroxide is of great importance [7]. The properties of MgO and Mg(OH)\textsubscript{2}nanostructured powders were studied in [8]. A greater advantage of nanostructured hydroxide is shown compared to that of a multilayer structure and with MgO properties. Methods of synthesis of products from nanostructured hydroxide are highlighted in work [9]. The adsorption properties of the nanostructured Mg(OH)\textsubscript{2}and MgO were discussed in [10]. Work [11] reports methods for obtaining the nanostructure and the structure of [Mg\textsubscript{6}] clusters. Study [12] provides information about modern research in China on the use of magnesium hydroxide with a nanostructure to extract terbium lanthanoid from solutions of strategic importance. However, there is no information on research methods, only a general description of the method of obtaining magnesium hydroxide from diluted solutions of sodium alkali chloride for composite materials is given. The attention of scientists and technologists is attracted by the possibility of using natural solutions of magnesium salts to obtain highly pure magnesium compounds for the pharmaceutical industry [13]. For society, this need has grown significantly, there are new, non-traditional medicines that have a multifunctional purpose in the context of the development of viral epidemics, so the experiments and production of highly pure raw materials for pharmaceuticals are of great importance.

High-purity magnesium hydroxide can serve as a material for sorption and sediment-matrix formation for other metal ions, which are determined by the analysis of substances and solutions. That could increase the reliability of the results of X-ray fluorescent and atomic absorption tests [14]. It is shown that it is possible to create highly pure compounds for medicine (creation of MR phantoms for verification of measurements and verification of the work of magnetic resonance imaging scanners) [15].

In the production of magnesium hydroxide in alkaline precipitate from its saline natural solutions, there are two features. It is shown that the first of them is that magnesium hydroxide has a sufficiently high ability to sorption of ion forms of most 3-d metals [16]. That makes it possible to get solutions sufficiently free from foreign compounds, primarily 3-d metals [17] at a satisfactory sediment filtration rate. The use of adsorbents of an extraneous nature and composition requires the cost of their synthesis and restoration of properties [18]. The second feature of the sediment is associated with its structure, which is significantly watered and has the ability to deform water layers and environmental pressure and slows down filtration [19]. The low rate of filtration when the sediment of magnesium hydroxide is separated from the uterine fluid has always been the subject of separate studies and proposed methods of formation. To clean raw materials solutions, it is proposed to heat raw brine [20]. Partial deposition of magnesium hydroxide is known, which increases the filtration rate to 180 kg/m\textsuperscript{2} per hour [21]. Increasing the value of alkalinity to pH1012 during deposition leads to a decrease in the rate of filtration of magnesium hydroxide sediment, so it is important to regulate such deposition and not to carry it out under stoichiometric conditions. There are known methods to form the sediment of magnesium hydroxide under the conditions of using inactive alkaline reagents; the use of etching retur during deposition; the use of surfactants; heating the reaction volume; electrical deposition [22]. However such techniques to satisfy the filtration of magnesium hydroxide significantly complicate the entire technology, making it unprofitable. It is also known that the ability to sorption of foreign metals-ions with magnesium hydroxide is influenced by the state of the structure of hydroxide and the state of the forms of foreign metal-ions [23]. When an alkaline reagent is added to a magnesium solution, sediment is formed at a high rate, so foreign metal ions react mainly with the surface of magnesium hydroxide, without the participation of sorption-active centers of the sediment structure. It is important to ensure the conditions of uniform interaction of active centers in the structure of sediment with hydrolyzed forms of micro components. Work [15] earlier found that alkaline deposition of up to 5 % of magnesium ions from the solution makes it possible to completely isolate impurity elements on hydroxide. An effective way to regulate the rate of formation of sediment of a uniform structure with effective centers is the...
method of homogeneous deposition [23]. Consideration of all technological methods for obtaining magnesium hydroxide shows that the method of homogeneous deposition using a reagent arising throughout the reaction volume is lacking in the descriptions of magnesium compound technologies and is first proposed in the current paper.

The advantages of homogeneous deposition are that the sediment reagent agent is formed evenly throughout the solution and gradually achieves the pH value necessary for deposition. Compared to ordinary, direct alkaline deposition yields large-crystalline sediments. An even more effective technique of uniform introduction of small concentrations of the sedimentary is deposition from a homogeneous solution (homogeneous deposition) or a method of emerging reagents. In this method, the sediment agent is formed as a result of hydrolysis of a specially introduced reagent in a raw material solution. This method is called homogeneous deposition by the method of emerging reagents. Even those substances that, in an ordinary precipitate, form amorphous sediments, can be obtained in the form of crystals during homogeneous deposition. This type of deposition according to the method of emerging reagents makes it possible to obtain crystalline sediments of magnesium hydroxide with satisfactory adsorption and filtration properties. This technology for the deposition of magnesium hydroxide from solutions has been proposed for the first time and has not been investigated by anyone. Therefore, it is important to confirm the declared theoretical foundations by experimental studies or refute them after receiving the results.

3. The aim and objectives of the study

The purpose of this study is to determine the processes of magnesium hydrolysis in highly concentrated solutions of its chloride. This would make it possible to obtain solutions of magnesium hydroxocomplexes as reagents for homogeneous deposition of magnesium hydroxide. Such sediment has a high filtration coefficient and satisfactory sorption properties, which are suitable for the technology of obtaining highly pure magnesium compounds.

To accomplish the aim, the following tasks have been set:
- to investigate the processes of alkaline hydrolysis of magnesium ions in a wide range of concentrations of its chloride and the conditions for the formation of solutions of magnesium hydroxocomplexes;
- to explore the possibility of using solutions of magnesium hydroxocomplexes as a reagent in the method of homogeneous formation of its hydroxide;
- to investigate the peculiarities of sedimentation of magnesium hydroxide, its structure, the sorption and filtration properties of sediments;
- based on the results of the current research and their generalization to propose a simplified and cheaper scheme of technology for obtaining highly pure magnesium compounds and several alkali metals.

4. The study materials and methods

To tackle the set tasks, it is necessary to establish the composition and structure of hydrolyzed magnesium forms, to investigate the peculiarities of the process of sedimentation of its hydroxide by the method of the emerging reagent. Experiments were conducted with alkaline deposition of magnesium hydroxide from its chloride of varying degrees of concentration of 0.05–4.0 mol/dm³. In this case, a solution of sodium hydroxide of a constant concentration of 0.2 g/dm³ was used, which was added to the magnesium chloride solution. To study the processes of sedimentation, physical and chemical methods were used: solubility, conductometric and potentiometric titration, nephelometry, sedimentary volume, sedimentation analysis method. When determining the composition of sediments, the method of residual concentrations, the preparation and analytical method (a variant of the method of moisture sediment with an indifferent component) were also used. X-ray analysis of sediment samples was used to determine their composition and structure according to diffractograms acquired from the diffractometer DRON0.5, radiation SI-Kα with a Ni-filter, the identification was carried out by the radiographic database ASTM. The size of sediment crystals (primary particles and formed crystals) was estimated by the spread of X-ray diffraction peaks according to the Debye-Scherrer formula.

Infrared IR spectra were obtained using the Spekord-75 IR spectrometer. Samples were prepared by pressing tablets with dried at 150 °C compound KBr, grade “chemically pure 3–4”, the spectra were recorded in the spectral region of 400–4000 cm⁻¹.

A comprehensive thermal analysis was carried out on the derivatograph of the PauliqueErdey system, made by IOM (Hungary). We examined the structure of sediments and their size by optical microscopy methods at the electron microscope H-500 of Hitachi(Japan). The optical microscope MP-6 with an object micrometer was also used. A statistical analysis of the obtained photos was carried out, which was supplemented by data from the sedimentation analysis.

The physicochemical characteristics of the sediments were defined as the results of filtration of the suspension of magnesium hydroxide when diluted in a Bunsencup in the corresponding vacuum system. The filter factor was determined from the formula:

\[ Q = K_f \cdot S \cdot H \cdot t / l \cdot t \]

where \( Q \) is the volume of the filtrate, cm³;
\( K_f \) is the filtration factor, cm/s;
\( H \) is the liquefaction under the filter, cm H₂O;
\( l \) is the sediment thickness;
\( t \) is the length of filtration, time.

Specific electrical conductivity was measured using the OR-201 conductometer. Potentiometric, pH-metric studies with the use of ion-selective electrodes were carried out at the device “pH-meter”, brand pH-340 (Belarus). Photo-electro-colorimetric measurements were carried out at the devices SF-24 and KFC-3 (Russia).

The processing of graphical dependences of the properties of magnesium hydroxide sediment on supersaturation, in order to find empirical formulas, was carried out by an alignment method.

5. Results of studying conditions for the production of magnesium hydroxide as an effective sorbent with different filtration properties

5.1. Investigating the hydrolysis of magnesium ions in the solutions of its chloride

To solve the specified tasks, experiments were conducted with alkaline deposition of magnesium hydrox-
ide from its chloride of various degrees of concentration, 0.05–4.0 mol/dm³; a solution of sodium hydroxide of a constant concentration of 0.2 g/dm³ was used. For magnesium chloride concentrations (0.1–0.5 g/dm³), the existence of two main types of sediments at pH=8.22–9.86 was established. It was found that the first type of sediment, at the beginning of the deposition, with the product of solubility PS=(2.41–2.64)∙10⁻¹⁰, corresponds to the known labile (active) form of magnesium hydroxide with PS=3.47∙10⁻¹⁰. The second type of sediment (at the end of deposition) with PS=(1.5–1.54)∙10⁻¹¹ most corresponds to the value for the crystallized form of hydroxide with PS=1.48∙10⁻¹⁰[24]. The values of the products of solubility of sediments calculated according to the results of experiments were compared with those reported in chemical studies [25, 26]. Such results held for all concentrations of magnesium chloride in our experiments. It was found (there are no other data in the literature) that the active magnesium hydroxide has the ability to react with magnesium chloride and pass into the solution (without the formation of a solid phase).

This ability to dissolve the labile form of magnesium hydroxide is noticeable already at a concentration of magnesium chloride of 1 g-ion/dm³ and increases with an increase in the initial concentration of magnesium chloride (Fig. 1).

![Fig. 1. Dependence of the alkalinity of MgCl₂ solutions on their concentration when magnesium hydroxide is deposited](image)

In this case, the concentrations of magnesium chloride of 1.0–4.75 g-ion/dm³ make it possible, after dissolving the active hydroxide, to form it with alkali again until the dissolution process stops. This formation of metastable solutions indicates the interaction of concentrated solutions of magnesium chloride and the original labile form of magnesium hydroxide with the formation of magnesium hydroxo-complexes. The formation of such complexes corresponds to the theory of “core-link” [27]. The core is magnesium chloride (oxychloride), and the link is magnesium hydroxide of the labile (active) form, according to the following scheme:

\[ \text{MgCl}_2 + 2\text{OH}^- \rightarrow \text{(active) Mg(OH)}_2 + \text{Cl}^- \]

\[ \text{MgCl}_2 + 2\text{OH}^- \rightarrow \text{Mg(OH)}_2 \cdot \text{Cl}_2 \]

\[ \text{[Mg}_2\text{(OH)}_3\text{]}\text{Cl}^+ + \text{(active).Mg(OH)}_2 \rightarrow \text{[Mg}_2\text{(OH)}_3\text{]}\text{Cl}_2 \]

According to the provisions of the theory “core-link”, it was found that magnesium ions in solutions can exist in the form of hydroxo-complexes of the general formula \[ \text{[Mg}_{n+1}\text{(OH)}_{2n-2}\text{]}\text{Cl}_2 \], where \( n=1, 2, 3, ..., \) corresponding to the composition of magnesium compounds, which eventually fall out of such metastable solutions of magnesium complexes. The comparison of the composition of such sediments shows their genetic correspondence to the composition of hydroxocomplexes of the solution.

5.2. Studying the sedimentation of magnesium hydroxide by the method of “Emerging reagent”

Further studies of the homogeneous formation of magnesium hydroxide sediments according to the method of “Emerging reagent” showed that in the case of dilution of concentrated solutions of magnesium chloride containing hydroxo-complexes, magnesium hydroxide sediments occur according to the hydrolysis equation when diluting hydroxo-complexes solutions according to the following scheme:

\[ \text{[Mg}_n\text{(OH)}_{2n-2}\text{]}^{2+} + 2\text{H}_2\text{O} \rightarrow n\text{Mg(OH)}_2 + 2\text{H}^+ \]

These sediments have several specific properties and compositions depending on the time of their formation; the process of sediment formation takes place in three stages (Fig. 2).

![Fig. 2. Formation of magnesium hydroxide sediment during dilution of alkaline solutions MgCl₂ from 4 to 0.5 mol/dm³, \( \tau \) is the sediment formation time, min; 1 — the amount of Mg²⁺ in the sediment (A), % of the initial amount; 2 — electrical conductivity of the reaction medium (\( \chi \)), Ohm⁻¹·cm⁻¹·10⁻³; 3 — optical density (D); 4 — pH environment](image)

We studied sediments at three main stages of formation, their composition, and properties based on experimental data. Such data were obtained by pH-metric methods, electrical conductivity, optical properties (nephelometry) of the reaction medium. In addition, in the experiments, we used
Infrared spectroscopy, including deuteranalogues (Fig. 3), X-ray structure and X-ray analysis, thermography (Fig. 4), and electron microscopy (Fig. 5).

Infrared spectra (Fig. 3) allow us to draw conclusions about the composition and structure of solid phases of magnesium hydroxide when diluting alkaline solutions of its chloride. For the embryo phase, fluctuations in water molecules include a number of absorption bands at 3400–ν(OH) (H₂O), 1620–1650–δ(H₂O), 545–560 cm⁻¹-ρ(H₂O). Absorption bands in the region of 1020–1220 cm⁻¹ indicate deformation oscillation of coordinated Mg-OH⁺ hydroxogroups. Water molecules coordinated with OH-groups around magnesium atoms give a wide intense band at 3400 cm⁻¹. Our calculation, based on the method from [28], for wavenumbers at 3420 cm⁻¹ and 3250 cm⁻¹ give the value of hydrogen bond energy $E_{\text{act}(1)}=19.2$ kJ/mol and $E_{\text{act}(2)}=9.42$ kJ/mol.

These values differ from the hydrogen bond energy value for [Mg(OH)₂]⁺ 8.82 kJ/mol, indicating the proton-donor nature of the embryos compared to the crystallized form of magnesium hydroxide. The absence of a band of valence oscillations of free hydroxide groups at 3690 cm⁻¹ indicates the absence of a crystalline sediment structure consisting of [Mg(OH)₂]⁺.

The embryos are X-ray amorphous, unlike crystals, their thermographic analysis reveals (Fig. 4) that the release of structural water at 75–90 °C, $E_{\text{act}(1)}=9.42$ kJ/mol at 100–140 °C, $E_{\text{act}(2)}=19.6$ kJ/mol. The stage of formation of primary particles of the second stage proceeds up to 100 °C (activation energy, 28.5 kJ/mol, corresponding to $E_{\text{act}(1)}+E_{\text{act}(2)}$) [28]. Thermal decomposition of crystals from the third stage of sediment proceeds at 412–420 °C, when decomposing crystals from the third stage of sediment there is no dehydration. The decomposition of crystals during dilution is characterized by the activation energy of 113–120 kJ/mol, as opposed to the decomposition of sediment from fragments of surface or linear groups, $E=78.3–87$ kJ/mol, which indicates a perfect structure, grown during dilution [29].

The composition of the embryos (phase 1 particles) corresponds to Mg₅(OH)₆Cl₂·8H₂O with water molecules that give the embryos an active hydrogen bond and which are coordinated together with OH groups (Fig. 5). The structure of the embryos and their composition together with the obtained data can be imagined based on the structure of the lattice of magnesium hydroxide [29]. In the layers of the crystal lattice [30], a single isolated fragment is a cluster of composition [Mg₅(OH)₁₆]₆⁻, which is the coordination octahedron of hydroxide ions around each metal atom.

Such an octahedral structure persists in embryos, the composition of which, according to chemical, spectroscopic, and thermographic analyses, indicates the presence of OH groups and coordinated water molecules around magnesium atoms. The embryo formula (phase 1 particles) corresponds to Mg₅(OH)₆Cl₂·8H₂O with coordinated OH groups and water molecules that give the embryos an active hydrogen bond. The embryos have the structure and composition of Mg₅(OH)₆Cl₂·8H₂O with a critical embryo diameter of 2–3 nm, which corresponds to the average diameter of the formation of 4 magnesium atoms, 6 hydroxide groups, and 8 water molecules [31].

At the first stage of sediment formation, the solution is turbid – the formation of embryonal sediment particles and a decrease in the pH of the solution. Subsequently, at the second stage, the transition of embryo particles into primary particles proceedsthrough their growth and polymerization to (30–60) nm at a constant pH. At the third, final stage, the formation of sediment with the participation of particles
of the second stage and mononuclear hydroxocomplexes of magnesium MgOH\(^+\) bond continues. The formation of a crystallized stable structure by its oriented build-up from primary particles takes place after the transition of all embryo particles into primary particles. Sedimentation ends with the coalescence of all primary particles into a stable crystalline structure of magnesium hydroxide with a crystal size of 1000–4000 nm. Such features of the formation of magnesium hydroxide sediment when diluting concentrated solutions of magnesium chloride, which contain alkaline magnesium complexes, are also evidenced by the data on the negative charge of embryo particles of sediment by the value of the suspension effect, \(\Delta \text{pH}\) (Fig. 6).

![pH graph](image)

**Fig. 6.** Value of the medium pH in the formation of Mg(OH)\(_2\) when diluting alkaline solutions MgCl\(_2\): 1 – pH of the uterine solution; 2 – suspension pH; 3 – \(\Delta \text{pH}=\text{pH}_{\text{sol.}}-\text{pH}_{\text{susp.}}\)

It is established that the formation of magnesium hydroxide sediment when concentrated solutions of magnesium chloride are diluted (more than 1.0 g-mol/dm\(^3\)) takes place in a kinetic mode, with the order of reaction at the first stage \(n_1=4\); at the stage of crystallization, the reaction order is \(n_2=20\) [32]. The crystallization process is limited to \(S=\frac{a_{\text{Mg}}^0}{a_{\text{OH}}^0} / \text{DR Mg(OH)\(_2\)} \text{supersaturation, where } a_{\text{Mg}}^0\) is an active concentration of magnesium ions in the solution after dilution, which is determined using an ion-selective electrode, \(a_{\text{OH}}^0\) is the active concentration of hydroxideions in the solution determined by the potentiometric method.

It was established that sediment with crystals of size 1000–4000 nm and, accordingly, with the filtration rate \(K_f=8\times10^{-3}\) m/h during the separation of the solid phase from the solution, can be obtained by observing the conditions of dilution of alkaline solutions of magnesium chloride.

The results of our studies testify that the most important indicator is the amount of supersaturation at the values of \(S\leq 20\) (Fig. 7), which makes it possible to obtain large-crystalline precipitate. This method does not require the creation of any special conditions for it.

![Dependence of properties](image)

**Fig. 7.** Dependence of the properties of magnesium hydroxide sediment on the value of supersaturation, dm\(^3\): 1 – the size of sediment crystals (\(l, \mu\text{m}\)); 2 – specific filtration rate (\(V, \text{ml/min}\)); 3 – filtration factor \((K_f, \text{m/h}\times10^{-3})\), \(S\) when diluted from 2.0 to 0.1 mol/dm\(^3\)

Mathematical processing by equalization [32] of graphic experimental data (Fig. 8) has made it possible to express the dependence of the properties of the sediment on the value of...
the supersaturation, $S$, when diluted, in the form of empirical formulas:

$$l=60.3/S^{1.25},$$  \hspace{1cm} (2)$$

where $l$ is the sediment crystal size, mm;

$$V=620/(\sqrt{t}),$$  \hspace{1cm} (3)$$

where $V$ is the specific filtration rate, ml/min;

$$K_f=0.16/S^{1.4},$$  \hspace{1cm} (4)$$

Where $K_f$ is the filtration coefficient, m/g; $S$ is the supersaturation value.

5.3. Investigating conditions for obtaining highly pure magnesium compounds

Regarding the influence of the properties of sediment particles before the removal of micro quantities of admixture metal ions from the solution, it was established that the greatest ability to sorption is demonstrated by the particles of sediment of the first stage during the formation of the embryonic phase [12]. The embryos have a polymeric structure of $\text{Mg}(_2\text{OH})_2\text{Cl}_2\cdot\text{H}_2\text{O}$ clusters (Fig. 5), which contain active hydroxide groups and water molecules, have hydrogen bonds, and are capable of chemical interaction (sorption) with forms of impurity metals hydrolyzed at pH ~ 9.0.

Such interaction is basic and irreversible with further sedimentation and crystallization. Similar processes of interaction of micro quantities of hydrolyzed ionic forms of metals with negatively charged surfaces of hydroxides are known from studies reported in [16].

The degree of purification of magnesium chloride solutions depends both on the value of the original alkaline module $\alpha=\text{[OH]/[Mg}^{2+}]$, and the final, after dilution, concentration of $\text{MgCl}_2$ solution (Tables 1, 2).

Table 1

<table>
<thead>
<tr>
<th>Type</th>
<th>The content of impurities, wt. % (in terms of $\text{MgCl}_2$)</th>
<th>$\text{[OH]/[Mg}^{2+}]$ in the starting solution</th>
<th>$\text{[MgCl}_2]$ after dilution $^2$, mol/dm$^3$</th>
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</table>

Note: * $-\text{MgCl}_2$ concentration after dilution was 0.5 mol/dm$^3$; ** the $\text{[OH]/[Mg}^{2+}]$ ration in the original solution was 0.08

These values determine the composition and concentration of magnesium hydroxocomplexes and their interaction during hydrolysis in the process of diluting the MgCl$_2$ solution (Table 1), which, as a result, determines the speed of the crystal building process and the properties of magnesium hydroxide sediment.

Table 2

<table>
<thead>
<tr>
<th>Micro impurity type</th>
<th>Crude compound</th>
<th>Purified compound</th>
<th>Degree of purification</th>
<th>Crude compound</th>
<th>Purified compound</th>
<th>Degree of purification</th>
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<td>4·$10^{-7}$</td>
<td>5·$10^{-3}$</td>
<td>1·$10^{-3}$</td>
<td>5·$10^{-2}$</td>
</tr>
</tbody>
</table>

Note: * $-\text{MgCl}_2$ concentration after dilution was 0.5 mol/dm$^3$; ** the $\text{[OH]/[Mg}^{2+}]$ ration in the original solution was 0.08

The data in Table 2 demonstrate that when depositing 5–10 % of the original amount of magnesium when diluted to 0.5 mol/dm$^3$, a high degree of purification of MgCl$_2$ solution from impurities and MgO from such solution is achieved.

5.4. Schemes of technologies for obtaining highly pure compounds of magnesium and alkali metals

Alkaline polycondensation of magnesium ions in concentrated solutions of its chloride followed by dilution is a universal technique both for obtaining highly pure magnesium compounds in (Fig. 9) and for the possible purification of a number of salts – NaCl, KCl, CaCl$_2$ in pharmacy and medicine (Fig. 10).

Such technologies do not require complex equipment and special procedures and can simply be mastered on the existing raw material base of a number of galurgical enterprises in Ukraine. As for waste, it can also be used. The main one – magnesium hydroxide, as a collector of impurities of metal compounds in the given scheme, contains (wt%) (in terms of MgO): Fe – 4·$10^{-3}$, Mn – 4·$10^{-2}$, Cu – 1·$10^{-2}$, Ni – 6·$10^{-3}$, Co – 2·$10^{-2}$, Cr – 6·$10^{-2}$, V – 4·$10^{-3}$, B – 4·$10^{-3}$, Ca – 4·$10^{-2}$ S – 3·$10^{-3}$. In each cleaning cycle, this collector is removed from the process and can be used for appropriate needs – the production of heat-resistant ceramics, heat-insulating materials, transformer steel coating, etc. [1, 3]. The purified target solutions of salts can be repeatedly treated to an ultra-high degree of purity in accordance with the specific requirements and needs of the latest equipment in its production. Highly pure magnesium hydroxide can be used to concentrate metal ions and radionuclides on such a highly pure collector for chemical analyses regarding their contents. Sorption properties and satisfactory filtration rate are suitable for cleaning natural and technical solutions. The concentration of scattered elements, especially lanthanoids, expands their raw materials for the needs of the latest technology.
Variants of technological schemes for obtaining highly pure magnesium compounds, as well as salts of alkali metals (Figs. 9, 10) based on the studied hydrolytic processes involving magnesium chloride and alkalis have been proposed.

6. Discussion of results of studying the hydroxide sediment processes

Our experimental results of the formation of magnesium hydroxide sediment by homogeneous deposition have confirmed the possibility and economic attractiveness of such technology for obtaining highly pure compounds of magnesium and alkali metals. The achieved values of the filtration coefficient exceed known ones from other studies by $1 \times 10^6$, and the degree of purification of target products from impurities of 3-d metals reaches $(1-4) \times 10^4$ times. The experiments involving magnesium hydrolysis processes have shown the possibility of obtaining and applying a solution of hydrolyzed forms. Applying such a solution has shown that it is suitable for the method of emerging reagents in homogeneous deposition (Fig. 1). The use of the soluble reagent NaCl, KCl, CaCl$_2$ leads to the gradual formation of sediment and its crystallization not locally but throughout the reaction volume (Fig. 2). The initial particles of sediment carry out effective sorption of impurities of metal ions, which are removed by increased sediment of a crystalline shape with high filtration characteristics.

Existing and known methods of using magnesium hydroxocomplex solutions as a reagent were not used before and were not known from the available literary sources. We believe that the research in this area is pioneering. The proposed technology was issued a patent of Ukraine for the invention [23].

The caveat of this research is the need for a more thorough study of the properties of sediment at the stages of its formation, as well as its sorption capacity in solutions to several metal ions in exceptionally valuable technologies. We believe that such studies will be continued.

One should establish the composition and structure of hydrolysis forms of magnesium, features of the process of sedimentation. In practice, such studies open up the following opportunities:

– obtaining highly pure compounds of magnesium, potassium, sodium for the pharmaceutical industry and for medicine, with the content of admixture compounds in the form of 3-d metals no more than the value of $5 \times 10^{-7}$ as a percentage;
– devising an effective technology for treating the solutions and industrial wastewater from toxic and active metals, including radioactive substances, fire-resistant articles;
– creating the basis of technology of extraction and concentration of liquid and scattered metals for the needs of the latest technology.

The result of our studies is the proposal to use auxiliary reagents, not of extraneous nature and composition, but only those that are genetically related to raw materials with the sediment agent being a solution of hydrolyzed forms of magnesium in its chloride. In practice, such techniques are simple and not energy-consuming, and easy to execute.

Another very important result is to ensure the quality and reliability of results in establishing the content of metal ions in various objects by the methods of neutron-activation, X-ray-fluorescent, and atomic-absorption analyses. During
analyses, it is very important that the matrix collector does not have the content of impurity metal ions, which can lead to false test results. Based on a highly pure matrix of magnesium hydroxide, it is possible to manufacture reference samples for laboratory tests of various types of articles and to verify procedures for conducting chemical analyses.

Our experiments confirmed that it is possible to achieve high chemosorptional properties of magnesium hydroxide to most ions of impurity metals and the formation of magnesium sorbent hydroxide with high filtration properties during the deposition of the crystalline structure of magnesium sorbent. The results and their numerical values show the effectiveness of the proposed technology. The practical aspect of the proposed technology relates, for example to the following:

- for extraction from solutions of especially rare concentrating elements and extraction of rare earth compounds of lanthanoid group;
- or treating solutions and industrial wastewaters from toxic and active metals that are easily filtered on magnesium hydroxide.
- to obtain highly pure compounds for medical purposes;
- to create heat-resistant plastics and fire-protectivearticles, which is especially important in modern high-rise construction.

Our results related to obtaining hydrolyzed magnesium forms in the form of solutions of its hydroxocomplexes make it possible to apply them to obtain a highly effective collector of impurity 3-d metals with a high filtration coefficient, which has significant advantages compared to foreign sorbents that were used in [7, 8]. Unlike the use of foreign substances as a sorbent collector, the principle of formation of a sediment collector by the method of “Emerging reagent” makes it possible to use a chemically related sorption material. Such application is quite simple in terms of the chemical nature of reagents and technological procedures since it is effective in cleaning solutions (Tables 1, 2), as well as in terms of the filtration properties of the sediment (Fig. 7).

The proposed solution for the formation of solutions of magnesium hydroxocomplexes and the use of them as a reagent to obtain magnesium hydroxide sediment according to the “Emerging Reagent” method is new in the area of research into the technology of magnesium compounds. Therefore, we have accomplished our goal to obtain highly pure magnesium compounds and receive its hydroxide sediment as a collector for impurities of 3-d metals.

The limitations of the proposed technology include the need to use highly concentrated solutions of magnesium chloride at the stage of obtaining hydroxocomplexes.

During the experiments, we have not been able to deeply study the structure of the formation of magnesium hydroxide based on theembryo and primitive particles and the possibility of programming and controlling this process at the crystallization stage.

The current research can be advanced using the method of concentration and isolation from solutions of scattered and liquid metals, which are highly needed for new technologies.

7. Conclusions

1. Our study into the hydrolysis of magnesium ions in a wide range of concentrations has established that the primary product of hydrolysis is magnesium hydroxide in the labile (active) form. At a magnesium chloride concentration larger than 1 mol/dm$^3$, the labile form of hydroxide interacts with magnesium chloride and forms soluble hydroxocomplexes.

2. When a solution of hydroxocomplexes is diluted with water or a weak solution of magnesium chloride, according to the method of emerging reagents, the sediment of its hydroxide is formed. The formation of magnesium hydroxide sediment takes place gradually according to the scheme of homogeneous deposition: embryos→primaryparticles→crystals. The crystals themselves are formed from primary nanoparticles measuring 40–60 nm.

3. The embryos and primary particles are the most sorption active to 3d metal ions, irreversible sorption remains in the formation of sediment crystals, the degree of concentration is $10^4$ times.

4. The formation of crystals measuring 1000–4000 nm can significantly increase the filtration process from the uterine solution. The filtration factor increases from $K_f=8 \cdot 10^{-5} \text{m/h}$ to $K_f=8 \cdot 10^{-9} \text{m/h}$.

Our research results are suitable for technologies for obtaining highly pure magnesium compounds from various types of raw materials containing magnesium ions. Underlying the proposed technology is the application of the concentrated solutions of magnesium chloride hydrolyzed with alkali and saturated with magnesium hydroxocomplexes. The difference in technologies is combining different methods of their use (dilution) depending on the concentration of magnesium in the raw material or already in the purified solution of magnesium salt.

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References

2. Gidroksid magniya v tehnologii XXI veka. Available at: https://polit.ru/article/2015/07/07/nikohim/


8. Ruchec, A. N., Besarab, S. V., Macukevich, I. V. Adsorbционье svojstva nanostrukturirovannyh poroshkov Mg(OH)2 i MgO. Available at: https://rep.bntu.by/bitstream/handle/data/40450/Adsorbционье_svojstva_nanostrukturirovannyh_poroshkov_Mg_OH_2_1_MgO.pdf


