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MAGNETICALLY SENSITIVE NANOCOMPOSITES FOR TARGETED ANTITUMOR THERAPY WITH APPLICATION OF GEMCITABINE

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The aim of the work is synthesis and study on the properties of polyfunctional magnetosensitive nanocomposites (NC) and target-directed magnetic fluids (MF) based on physiological solution (PS), magnetite, gemcitabine (GEM) and HER2 antibodies (AB), promising for use in targeted antitumor therapy against MDA-MB-231 aggressive tumor cells of triple-negative human breast cancer (BC) with high proliferative and metastatic activity.

The specific surface area (S_{sp}) of samples was determined by the method of thermal desorption of nitrogen using a device KELVIN 1042 of "COSTECH Instruments". The size of nanoparticles (NP) has been estimated by the formula $D_{BET} = 6/(\rho S_{BET})$, where ρ is the density of NC particle, S_{BET} is the value of the specific surface area calculated by the polymolecular adsorption theory of Brunauer, Emmett and Teller (BET). The surface condition of nanodispersed samples was studied by IR spectroscopy ("Perkin Elmer" Fourier spectrometer, a model 1720X). To calculate the concentration of hydroxyl groups on the surface of nanostructures, the method of differential thermal analysis was used in combination with differential thermogravimetric analysis. The thermograms were recorded using a derivatograph Q-1500D of MOM firm (Hungary) in the temperature range of 20–1000 °C at a heating rate of 10 deg/min. X-ray phase analysis of nanostructures was performed using a diffractometer DRON-4-07 (CuK_{α} radiation with a nickel filter in a reflected beam, the Bragg-Brentano focusing). The size and shape of NP were determined by electron microscopy (a transmission electron microscope (TEM) JEM-2100F (Japan)). The hysteresis loops of the magnetic moment of the samples were measured using a laboratory vibration magnetometer of Foner type at room temperature. Measurement of optical density, absorption spectra and GEM concentration in solutions was performed by spectrophotometric analysis (Spectrometer Lambda 35 UV/Vis Perkin Elmer Instruments). The amount of adsorbed substance on the surface of magnetite was determined using a spectrophotometer at $\lambda = 268$ nm from a calibration graph. The thickness of the adsorbed layer of GEM in the composition of $Fe_3O_4@GEM$ NC was determined by magnetic granulometry. To study the direct cytotoxic/cytostatic effect of a series of experimental samples of MF based on PS, Fe_3O_4 NP, GEM, HER2 AB, as well as MF components in mono- or complex use, onto MDA-MB-231 cells in vitro, IC_{50} index was determined.

MF were synthesized on the basis of single-domain Fe_3O_4 and PS, stabilized with sodium oleate (Ol.Na) and polyethylene glycol (PEG), containing GEM and HER2 ($Fe_3O_4@GEM/Ol.Na/PEG/HER2+PS$). The cytotoxic/cytostatic activity of MF against MDA-MB-231 cells was studied. It was found that as a result of application of synthesized MF composed of $Fe_3O_4@GEM/Ol.Na/PEG/HER2+PS$ at the concentration of magnetite of 0.05 mg/mL, GEM - 0.004 mg/mL and HER2 AB - 0.013 μ g/mL, a synergistic effect arose, with reduction of the amount of viable BC cells to 51 %. It has been proved that when using MF based on targeted $Fe_3O_4/GEM/HER2$ complex, the increased antitumor efficacy is observed compared to traditional use of the drug GEM, with a significant reduction (by four times) of its dose. The high cytotoxic/cytostatic activity of $Fe_3O_4/GEM/HER2$ complexes is explained by the fact that endogenous iron metabolism disorders play a significant role in the mechanisms of realization of the apoptotic program under the influence of nanocomposite. Thus, when the nanocomposite system contains $Fe_3O_4/GEM/HER2$ complexes in MDA-MB-231 cells, a significant increase is observed in the level of "free iron", which favours formation of reactive oxygen species and causes oxidative stress (Fenton reaction). The consequences of oxidative stress are induction of apoptosis, enhancement of lipid peroxidation processes, as well as structural and functional rearrangement of biological membranes. The prospects

have been shown of further studies of $\text{Fe}_3\text{O}_4@\text{GEM}/\text{OI.Na}/\text{PEG}/\text{HER2}+\text{PS MF}$ in order to create on their basis a magnetically carried remedy for use in targeted antitumor therapy.

Keywords: gemcitabine, nanosized single-domain magnetite, core-shell nanocomposites, magnetic fluids, HER2 antibody, targeted antitumor therapy

INTRODUCTION

One of the main problems of modern antitumor therapy with gemcitabine (GEM) is its toxic effects on the human body and the occurrence of severe adverse reactions in patients [1–4]. Therefore, a generally accepted alternative to traditional GEM chemotherapy has been the use of targeted delivery methods [5–11], which allows us to create a therapeutic dose of the drug in the disease locus, to carry out therapy at a much lower total dose and minimize side effects. Among the variety of modern methods of targeted drug delivery and local therapy of diseases at the level of organs, cells and genes, the most developed at this time are methods using magnetically sensitive nanostructures conjugated with antitumor drugs of different mechanism of action.

To date, a concept has been substantiated of chemical construction of magnetosensitive nanocomposites (NC) with multilevel hierarchical nanoarchitecture, characterized by the functions of “nanoclinics” and medico-biological nanorobots [12–17]: recognition of microbiological objects in biological media; targeted delivery of drugs to specific cells and organs, and deposition; complex local chemo-, immuno-, neutron-capture, hyperthermic, photodynamic therapy and magnetic resonance imaging in real time, detoxification of the body by adsorption of cell decomposition residues, viral particles, heavy metal ions, *etc.* and their removal with the aid of magnetic field [18–24]. The urgency of the topic is due to its focus on the creation of the latest medical theranostic remedies for targeted delivery of drugs with different mechanisms of action, and local complex therapy, primarily for the needs of oncology.

For the manufacture of magnetically sensitive multifunctional NC, a significant interest of researchers is drawn to nanostructures of core-shell type based on single-domain magnetite (Fe_3O_4), which are characterized by a unique set of physical, chemical and biological properties, the capability to create magnetic fluids (MF) on their basis, containing

oncological remedies with various functional purpose and mechanisms of action [25, 26].

In recent years, there has been a significant increase in the use of natural mechanisms of endocytosis and nanostructures of various types to deliver drugs to tumors, involving ligands such as hormones, vitamins and growth factors against tumor-associated receptors that are overexpressed on tumor cell surfaces, with limited distribution in normal tissues [9, 11, 27]. At present, magnetically sensitive and nonmagnetic conjugates are actively being developed by the methods of modern nanotechnology to deliver GEM in a targeted manner to malignant cells, which will allow to reduce its systemic toxicity. The advantage of the use of drug conjugation on the surface of nanoparticles to deliver drugs *in vivo* is a slow release of the drug, which potentially provides a longer presence of the drug in the bloodstream at the required level. A positive result of the use of hyperthermia in combination with prolonged drug release may be an improvement in the patient’s condition and a decrease in the frequency of drug reception.

Despite a large number of successful studies on the use of magnetic nanoparticles as a theranostic material, as well as repeated successful results in small animal models, they still do not meet clinical needs. However, upon reaching high drug capacity, increased specificity and affinity for tumor cells, a combination of imaging and multimodal local therapy, magnetic nanoparticles may become convenient for clinical use in the near future and significantly affect the effectiveness of cancer treatment.

We note that a review of scientific publications is contained in [28] on the synthesis, study of properties of magnetosensitive NC based on GEM, promising for use in the method of targeted delivery, and their applications in medicine, in particular antitumor therapy.

The aim of this work is to synthesize and study the properties of magnetically sensitive polyfunctional NC and target-directed magnetic fluids based on magnetite and GEM, promising for use in targeted antitumor therapy against

MDA-MB-231 cells of aggressive tumor with high proliferative and metastatic activity, three times negative human breast cancer (BC).

EXPERIMENTAL PART AND DISCUSSION

Research methods and results. Gemcitabine ((2-deoxy-2',2')difluorocytidine monochloride) is a cytotoxic drug, an antimetabolite from the group of pyrimidine antagonists, that has been chosen as the drug of chemotherapeutic mechanism of action. GEM belongs to the List of basic medicines of the World Health Organization and the most effective and safe drugs needed in the health care system. It is characterized by considerable antitumor activity in the treatment of solid tumors (non-small cell lung cancer, pancreatic, bladder, breast, ovarian cancer), satisfactory tolerability and the possibility of successful combination with other antitumor drugs. It is used for treatment of cholangiocarcinoma and other types of bile duct cancer. Currently, the possibility is being actively studied [5–11, 28] to use GEM in the composition of magnetosensitive NC to create polyfunctional antitumor drugs for targeted delivery and local therapy, e.g. breast cancer, hepatocellular carcinoma, osteosarcoma, etc.

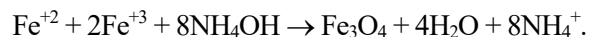
The antibody (AB) HER2 (Neu, ErbB-2, CD340) is a membrane protein, tyrosine protein kinase of EGFR/ErbB epidermal growth factor receptor family, which is encoded by ERBB2 human gene, it has been chosen as a drug of immunotherapeutic mechanism of action. HER2 gene amplification plays an important role in the pathogenesis and progression of certain aggressive types of cancer [29–31]. HER2 is an important biomarker and therapeutic target of the disease, associated with tumor aggressiveness and unfavorable prognosis. HER2 AB is considered to be an optimal for the treatment of diseases such as human BC, in particular, in the presence of metastases. Therefore, for *in vitro* studies, we chose HER2 AB in combination with GC in composition of MF based on magnetite.

To test the antitumor activity of the synthesized MF *in vitro*, in this work we chose MDA-MB-231 cells of the aggressive tumor with high proliferative and metastatic activity, three times negative human BC.

Samples of magnetically sensitive nanostructures (ensembles of Fe₃O₄ nanoparticles (NP) and Fe₃O₄@GEM NC) and MF stabilized with sodium oleate (Ol.Na) and

polyethylene glycol (PEG) were synthesized for research.

The synthesis of nanodispersed magnetite was carried out according to the technique [10] by coprecipitation of iron salts by the reaction:



The synthesized ensembles of Fe₃O₄ NP were characterized by sizes of 3–23 nm. The average size (d_0) of NP depended on the synthesis conditions and was 6–13 nm, and the size distribution could be controlled technologically.

The specific surface area (S_{sp}) of the samples was determined by the method of thermal desorption of nitrogen on a device KELVIN 1042 of “COSTECH Instruments” firm. The size of NP has been estimated by the formula $D_{\text{BET}} = 6/(\rho S_{\text{BET}})$, where ρ is the density of NC particle, S_{BET} is the value of the specific surface area calculated by the polymolecular adsorption theory of Brunauer, Emmett and Teller (BET). The specific surface area (S_{sp}) of synthesized magnetite was $S_{sp} = 90\text{--}180 \text{ m}^2/\text{g}$, dependent on the average particle size, in this work we used samples with $S_{sp} = 110 \pm 1 \text{ m}^2/\text{g}$.

The surface condition of nanodispersed samples was studied by IR spectroscopy (“Perkin Elmer” Fourier spectrometer, a 1720X model). OH functional groups were revealed by the study on IR spectra of magnetite surface. To calculate the concentration of hydroxyl groups on the surface of nanostructures, the method of differential thermal analysis was used in combination with differential thermogravimetric analysis. Thermograms were recorded using a Q-1500D derivatograph of MOM firm (Hungary) in the temperature range of 20–1000 °C at a heating rate of 10 deg/min. It has been found that the concentration of OH functional groups on the surface of magnetite is 2.4 mmol/g.

X-ray phase analysis of nanostructures was performed using a diffractometer DRON-4-07 (CuK α radiation with a nickel filter in a reflected beam, the Bragg-Brentano focusing). The size of the crystallites was determined from the width of the corresponding most intense line according to the Scherrer equation. The value of the average diameter of Fe₃O₄ NP, calculated from the results of investigation of X-ray diffraction patterns according to the Scherrer formula, D_{XRD} was 10.5 nm.

To study the morphology and size distribution of NP, their dispersions in water were used. The size and shape of NP were determined by electron microscopy (a transmission electron microscope (TEM) JEM-2100F (Japan)). Fe₃O₄ NP in ensemble were characterized by spheroidal shape, their average size was ~10.8 nm.

The hysteresis loops of the magnetic moment of the samples were measured using a laboratory vibration magnetometer of Foner type at room temperature. Description of the installation and the measurement technique are

set out in [32]. Demagnetized nanoparticles were distributed in a paraffin matrix with a volume concentration of ~0.05 to prevent interaction. For comparison, we used materials with a known value of the specific saturation magnetization (σ_s): a tested sample of nickel and Fe₃O₄ NP (98 %) manufactured by “Nanostructured & Amorphous Materials Inc.”, USA. The error in measuring σ_s did not exceed 2.5 % in relation to the reference sample. Table 1 shows the magnetic characteristics of the synthesized ensemble of magnetite nanoparticles.

Table 1. Magnetic characteristics of the ensemble of magnetite nanoparticles

Sample	H_c , Oe	σ_s , emu/g	σ_r , emu/g	σ_r/σ_s	$\alpha_{Fe_3O_4}^{calc}$, %
Fe ₃ O ₄	41	57.7	10.4	0.18	100

H_c , Oe is the coercive force; σ_s , emu/g is the specific saturation magnetization of NC; σ_r , emu/g is the residual specific magnetization of NC; σ_r/σ_s is the relative residual magnetization; $\alpha_{Fe_3O_4}^{calc}$ is the calculated mass concentration of Fe₃O₄ in NC, %

Measuring of optical density, absorption spectra and GEM concentration in solutions was performed by spectrophotometric analysis (a spectrometer Lambda 35 UV/Vis Perkin Elmer Instruments).

The adsorption capacity of samples A (mg/g) has been calculated by the formula: $A = (C_0 - C_{eq}) \cdot V/m$, where C_0 and C_{eq} are the concentration of the initial solution and the solution after adsorption (mg/L), V is the volume of the solution (L), m is the mass of the sorbent (g). Adsorption isotherms were constructed on the basis of experimental results. The coefficients of distribution E (mL/g) of GEM between the surface of nanostructures and the solution, the extraction extent R (%) were determined by the formulas: $E = A/C_{eq}$, $R = (1 - C_{eq}/C_0) \cdot 100$ %, respectively.

Gemcitabine TEVA (Pharmachemie BV, the Netherlands) was used for research in this work. Adsorption of GEM on the surface of magnetite Fe₃O₄ was performed in physiological solution in the concentration range $C_0 = 0.02$ – 0.67 mg/mL ($m = 0.03$ g, $V = 5$ mL, pH = 3.0) for 2 h in a dynamic mode at room temperature. The level of pH of the medium of GEM solution in 0.9 % NaCl was set by 0.1N HCl. The amount of adsorbed substance on the surface of magnetite was determined using a spectrophotometer at $\lambda = 268$ nm from a calibration graph.

The experimental values of adsorption capacity A of magnetite surface were ~37.2 mg/g, extraction extent R – 33.13 %, distribution coefficient $E = 82.58$ mL/g.

The magnetic characteristics of magnetite with adsorbed GEM (Fe₃O₄@GEM NC) are given in Table 2.

Table 2. Magnetic characteristics of Fe₃O₄@GEM NC

Sample	H_c , Oe	σ_s , emu/g	σ_r , emu/g	σ_r/σ_s	$\alpha_{Fe_3O_4}^{calc}$, %
Fe ₃ O ₄ @GEM	44	39.0	6.07	0.15	68

Using the method of magnetic granulometry and assuming that the thickness of GEM layers depends a little on the diameter of Fe₃O₄ NP, we have estimated the average value of the thickness

of the adsorbed GEM layer in the composition of Fe₃O₄@GEM NC, which is 2.4 ± 0.1 nm.

Synthesis and properties of magnetic fluids. MF were synthesized based on magnetite and

physiological solution (PS), stabilized with Ol.Na and PEG, containing GEM ($\text{Fe}_3\text{O}_4@\text{GEM}/\text{Ol.Na}/\text{PEG}+\text{PS}$).

As the dispersed phase of MF, nanosized magnetite in the single-domain state was used at the concentration of 14 mg/mL (for comparison), or $\text{Fe}_3\text{O}_4@\text{GEM}$ NC based on it.

To prevent aggregation, Fe_3O_4 and NC nanoparticles were stabilized with sodium oleate ($\text{C}_{18}\text{H}_{33}\text{O}_2$), dynamic mode, 1 h), and polyethylene glycol (PEG-2000).

To stabilize the surface of NP and NC in the composition of MF, the mass of sodium oleate $m_{\text{Ol.Na}}$ was calculated taking into account the concentration of reactive hydroxyl groups on the surface of magnetite. The calculation was performed according to the formula: $m_{\text{Ol.Na}} = B \cdot M \cdot m$, where B is the concentration of hydroxyl groups (2.2 mmol/g) on the surface of starting nanosized magnetite, M is the molecular mass of sodium oleate (304 Da), m is the mass of Fe_3O_4 or NC.

It is known that PEG prevents the adsorption interactions of fluid components with protein molecules [33], which is important in medical applications of MF. Additional modification with PEG-2000 was carried out in a dynamic mode using a shaker, the amount of polymer was 10–15 % by weight of the sample of Fe_3O_4 NP, or NC [34, 35]. The double stabilizing layer of NC ($\text{Ol.Na}@\text{PEG}$) was considered to be continuous with thickness ≈ 3 nm in water and PS, and ≈ 1 nm in dry residues [35].

Putting of HER2 AB into the composition of $\text{Fe}_3\text{O}_4@\text{GEM}/\text{Ol.Na}/\text{PEG}+\text{PS}$ MF and obtaining of $\text{Fe}_3\text{O}_4@\text{GEM}/\text{Ol.Na}/\text{PEG}/\text{HER2 AB}+\text{PS}$ MF was performed in a dynamic mode using a shaker.

To obtain optimal experimental MF based on magnetite, we used compositions with the following parameters: concentration of Fe_3O_4 – 14 mg/mL, particle size of Fe_3O_4 – 4–22 nm, average particle size of Fe_3O_4 – 10.8 nm; average particle size of Fe_3O_4 stabilized with sodium oleate – 16.8 nm; saturation magnetization $M_\infty = 14.1 \pm 2.5$ % Gs, hypsometric height – 25 ± 10 % cm, viscosity $\eta = 1.14 \pm 3$ % mPa·s, density $\rho_{\text{MF}} = 1.14 \pm 1.0$ % g/cm³, the concentration of GEM was 1.25 mg/mL, HER2 AB – 3.75 µg/mL.

The concentration of GEM and HER2 AB in such MF is determined by the therapeutic need. In the starting MF, the concentration of GEM

and HER2 AB has been 1.25 mg/mL and 3.75 µg/mL, respectively, which allows us to provide the required doses of the drug in the test samples by diluting the starting MR. In addition, this fluid is characterized by satisfactory rheological properties and sedimentation stability.

We note that the conditions used in these studies for the synthesis of magnetite, adsorption immobilization of GEM on the surface of nanosized Fe_3O_4 , production of MF, as well as the magnetic properties of the research samples are described in more detail in [9, 11, 25, 26].

Investigation of the influence of experimental samples on the viability of MDA-MB-231 cells of human BC in vitro. To study the direct cytotoxic/cytostatic effect of a series of experimental samples of MF based on PS, Fe_3O_4 NP, GEM, $\text{Fe}_3\text{O}_4@\text{GEM}$ NC, HER2 AB, in mono- or complex application onto MDA-MB-231 cells *in vitro*, IC_{50} index was determined.

The description of the used samples, materials, devices and methods is given below.

Samples

1. MF: $\text{Fe}_3\text{O}_4@\text{Ol.Na}/\text{PEG} + \text{PS}$ (control 1),
2. Gemcitabine + PS (control 2),
3. HER2 AB + PS (control 3),
4. MF + GEM: $\text{Fe}_3\text{O}_4@\text{GEM}/\text{Ol.Na}/\text{PEG} + \text{PS}$,
5. MF + AB: $\text{Fe}_3\text{O}_4@\text{Ol.Na}/\text{PEG} + \text{PS} + \text{HER2}$,
6. MF + GEM + AB: $\text{Fe}_3\text{O}_4@\text{GEM}/\text{Ol.Na}/\text{PEG} + \text{PS} + \text{HER2}$,
7. GEM + HER2 + PS (control 4),
8. PS (control 5).

For all the corresponding systems, the concentration of Fe_3O_4 was $C_{\text{Fe}_3\text{O}_4} = 3$ mg/mL, the concentration of GEM $C_{\text{GEM}} = 0.25$ mg/mL, antibodies - $C_{\text{HER2 AB}} = 0.75$ µg/mL.

Materials

Medium: DMEM High glucose (Biowest, France, catalog No. L0102-500).

Serum: fetal calf serum (FCS) (Biowest, France, catalog No. S181B-500).

Solutions: Versene solution (BioTestLaboratory, Ukraine), phosphate-saline buffer (Sigma, USA, cat. No. D1408), physiological solution (Lekhim, Ukraine, 71033007 series), ethyl alcohol (Ukrspirt, Ukraine).

Dyes: crystalline violet (Sigma, USA, cat. No. C6158).

Other materials: plastic ware for cell culture (TPP, Italy), 96-well plates for cell culture (SPL, Korea).

Devices

CO₂ incubator (Heal Force, China), an inverted microscope Axiovert 25 (Carl Zeiss, Germany), a Goryaev camera (Pharmmedtech, Ukraine), a mini-shaker PSU-2T (BioSan, Latvia), a multiwell spectrophotometer (Labsystems Multiskan PLUS, Finland), automatic pipettes up to 20 µL (Eppendorf AG, Germany), 200 µL (Thermo Fisher Scientific Oy, Finland) and 1000 µL (Eppendorf AG, Germany).

Methods

The tested cells were cultured in complete DMEM medium with 10 % FCS and 40 µg/mL of gentamicin in a plastic ware in humidified atmosphere at 5 % CO₂ and 37 °C. Changing of the medium and resowing of cells were performed according to standard technique [36]. The cells in the exponential phase of growth were used for experiments. In 24–48 h after the last resowing, the cells were planted out for cultivation at a concentration of 1–1.5×10⁴ cells/well of a 96-well plate. In 24 h, different amounts of experimental substances were added to the respective wells according to the study scheme. Syringe filters were used when making the solutions. The cells were incubated for

another 48 h, and the cytotoxic effect of each experimental substance was determined.

The viability of cells in the experiment was evaluated by colorimetric method, staining the cells with crystalline violet. Adhesive cells are separated from the substrate after death. This property is used to determine the number of living cells after treatment with test agents. One way to detect attached (living) cells is to stain them with a crystalline violet dye that binds to proteins and DNA. The results were evaluated using a multi-well spectrophotometer (wavelength 540 nm). The percentage of viable cells (I_R) was calculated by the formula:

$I_R = (A_{540}(\text{experiment})/A_{540}(\text{control})) \times 100 \%$, where A_{540} is absorption (the value proportional to cell concentration) at wavelength of 540 nm.

Cytotoxic and antiproliferative activity was determined using the index “inhibitory concentration 50” (IC₅₀ – the concentration of the test substance at which 50 % of cells die), which was calculated using Excel program. Table 3 shows the results of the study of the effect of concentrations of the studied preparations when used in mono-regime and their combinations on the viability of MDA-MB-231 cells.

Table 3. Influence of concentrations of the studied preparations in mono-regime and their combinations on the viability of MDA-MB-231 cells

Sample			MF	GEM	HER2	MF+ GEM	MF+ HER2	MF+ GEM+ HER2	GEM+ HER2	PS
Concentration										
MF*, mg/mL	GEM, mg/mL	HER2, µg/mL	Amount of living cells, % **							
1.5	0.125	0.38	—***	38.0±	89.7 ±	—***	—***	—***	36.5±	96.0 ±
				1.7	1.9				1.8	3.1
0.75	0.063	0.19	—***	35.0±	88.7± 2.9	—***	—***	—***	35.0±	89.7 ±
				2.2					2.4	6.6
0.38	0.031	0.1	—***	38.6±	98.8± 4.6	—***	—***	—***	36.9±	97.3 ±
				1.6					4.1	1.3
0.19	0.016	0.05	—***	48.5±	98.4± 5.9	—***	—***	—***	46.4±	94.5 ±
				2.2					3.2	1.9
0.1	0.008	0.025	91.3±	72.4±	100.0±	70.7±	85.9 ±	55.0±	63.9±	90.9 ±
			6.6	1.9	2.3	6.7	4.3	5.7	2.5	6.2
0.05	0.004	0.013	90.1±	74.3±	100.0±	68.7±	88.0 ±	51.2±	72.6±	96.7 ±
			7.0	2.4	4.5	5.6	7.7	2.9	3.9	1.8
0.025	0.002	0.007	97.3±	88.3±	100.0±	73.9±	95.1 ±	66.5±	85.0±	96.4±
			9.7	2.2	2.9	4.8	10.0	4.8	1.6	2.1

Note: * – the concentration of MF was determined by the content of magnetite, ** – in comparison with the cells of the control group, which were cultured without adding the substances indicated in Table 3 (100 %), *** – it is not determined, or it is not determined reliably

It has been shown by analysis of the data that HER2 antibody at concentrations less than 0.1 $\mu\text{g/mL}$ and PS in monoapplication do not affect significantly the viability or proliferation of MDA-MB-231 BC cells. Therefore, the IC_{50} is not determined for these ingredients, at the studied concentrations.

The IC_{50} was determined experimentally for gemcitabine to be $C_{\text{GEM}} = 0.016 \text{ mg/mL}$, and for GEM/HER2 composition ($C_{\text{GEM}} = 0.016 \text{ mg/mL}$, $C_{\text{HER2}} = 0.05 \text{ }\mu\text{g/mL}$) with respect to MDA-MB-231 BC cells.

In the experiment, the IC_{50} value was not determined for MF at concentrations less than 0.1 mg/mL in the absence of toxicity. Cultivating MDA-MB-231 cells in the presence of MF and HER2 AB, simultaneously, we observed a slight synergistic effect of these components up to 4 %, compared with control 1.

Close to optimal concentrations of all components (MF, GEM and HER2 antibodies) were determined to achieve the highest cytotoxic effect of MF. Experimental studies have shown that the combination of concentrations of components $C_{\text{MF}} = 0.05 \text{ mg/mL}$, $C_{\text{GEM}} = 0.004 \text{ mg/mL}$, $C_{\text{HER2}} = 0.013 \text{ }\mu\text{g/mL}$ creates the best result of cytotoxic activity, at which about 51.2 % of MDA-MB-231 cells survive.

The results show that the proliferation of human breast carcinoma MDA-MB-231 cells is inhibited by GEM in individual use better than by other studied components. Under the conditions of complex action of GEM with other experimental substances, it has been found that:

- MF allows to increase the antitumor activity of GEM *in vitro* by 5.6–14.4 %;
- MF allows to increase the antitumor activity of HER2 antibody *in vitro* by 4.9–12.0 %,
- target-directed MF/GEM/HER2 complex causes a synergistic antitumor effect and increases the cytotoxic activity of GEM by more than 23 %.

This can be explained by the fact that in the mechanisms of realization of the apoptotic

program under the influence of nanocomposite, a significant role is played by disorders of endogenous iron metabolism. Thus, when the nanocomposite system contains $\text{Fe}_3\text{O}_4/\text{GEM}/\text{HER2}$ complexes, in human breast carcinoma MDA-MB-231 cells a significant increase is observed in the level of “free iron”, which favours formation of reactive oxygen species and causes oxidative stress (Fenton reaction). The consequence of oxidative stress is the induction of apoptosis and enhancement of lipid peroxidation processes. It is proved that the nanocomposite is capable to initiate the apoptotic program in cells by mitochondrial pathway and cause structural and functional rearrangements of biological membranes [37].

CONCLUSIONS

Magnetic fluids were synthesized based on Fe_3O_4 and physiological solution, stabilized with sodium oleate and polyethylene glycol, containing GEM and HER2 antibody ($\text{Fe}_3\text{O}_4@\text{GEM}/\text{Ol.Na}/\text{PEG}/\text{HER2}+\text{PS}$). The cytotoxic/cytostatic activity of MF was studied against MDA-MB-231 aggressive tumor cells of triple-negative human breast cancer with high proliferative and metastatic activity. It was found that as a result of application of synthesized MF composed of $\text{Fe}_3\text{O}_4@\text{GEM}/\text{Ol.Na}/\text{PEG}/\text{HER2}+\text{PS}$ at the concentration of magnetite of 0.05 mg/mL , GEM – 0.004 mg/mL and HER2 AB – 0.013 $\mu\text{g/mL}$, a synergistic effect arose, with reduction of the amount of viable BC cells to 51 %.

It has been proved that when using MF based on targeted $\text{Fe}_3\text{O}_4/\text{GEM}/\text{HER2}$ complex, the increased antitumor efficacy is observed compared to traditional use of the drug GEM, with a significant reduction (by four times) of its dose. The prospects have been shown of further studies of $\text{Fe}_3\text{O}_4@\text{GEM}/\text{Ol.Na}/\text{PEG}/\text{HER2}+\text{PS}$ MF in order to create on their basis a magnetically carried remedy for use in targeted antitumor therapy.

Магніточутливі наноккомпозити для адресної протипухлинної терапії з використанням гемцитабіну

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Метою роботи є синтез і дослідження властивостей поліфункціональних магніточутливих наноккомпозитів (НК) та мішень-спрямованих магнітних рідин (МР) на основі фізіологічного розчину (ФР), магнетиту, гемцитабіну (ГЦ) та антитіла (АТ) HER2, перспективних для використання в адресній протипухлинній терапії проти клітин лінії MDA-MB-231 агресивної пухлини з високою проліферативною та метастатичною активністю тричі негативного раку молочної залози (РМЗ) людини.

Питому поверхню ($S_{\text{пит}}$) зразків визначали методом термодесорбції азоту за допомогою приладу KELVIN 1042 фірми "COSTECH Instruments". Розмір наночастинок (НЧ) оцінювали за формулою $D_{\text{ВЕТ}} = 6/(\rho S_{\text{ВЕТ}})$, де ρ – густина частинки НК, $S_{\text{ВЕТ}}$ – значення питомої площі поверхні, розрахованої за теорією полімолекулярної адсорбції Брунауера, Еммета і Теллера (БЕТ). Дослідження стану поверхні нанодисперсних зразків здійснювали методами ІЧ-спектроскопії (Фур'є-спектрометр "Perkin Elmer", модель 1720X). Для розрахунку концентрації гідроксильних груп на поверхні наноструктур використовували метод диференціального термічного аналізу в поєднанні з диференціальним термогравіметричним аналізом. Реєстрацію термограм здійснювали за допомогою дериватографа Q-1500D фірми МОМ (Угорщина) в інтервалі температур 20–1000 °C за швидкості нагрівання 10 град/хв. Рентгенофазовий аналіз наноструктур виконували за допомогою дифрактометра ДРОН-4-07 (випромінювання CuK_α з нікелевим фільтром у відбитому пучку, фокусування за Бреггом-Брентано). Розмір та форму НЧ визначали методом електронної мікроскопії (просвічуючий електронний мікроскоп (ПЕМ) JEM-2100F (Японія)). Петлі гістерезису магнітного моменту зразків вимірювали за допомогою лабораторного вібраційного магнітометра фонерівського типу при кімнатній температурі. Вимірювання оптичної густини, спектрів поглинання та концентрації ГЦ в розчинах здійснено методами спектрофотометричного аналізу (Spectrometer Lambda 35 UV/Vis Perkin Elmer Instruments). Кількість адсорбованої речовини на поверхні магнетиту визначали за допомогою спектрофотометра при $\lambda = 268$ нм за калібрувальним графіком. Товщину адсорбованого шару ГЦ у складі НК Fe_3O_4 @ГЦ визначали методом магнітної гранулометрії. Для вивчення прямої цитотоксичної/цитостатичної дії серії експериментальних зразків МР на основі ФР, НЧ Fe_3O_4 , ГЦ, АТ HER2, а також компонентів МР в моно- або комплексному застосуванні, на клітини лінії MDA-MB-231 *in vitro*, визначали показник IC_{50} .

Синтезовано МР на основі одномоментного Fe_3O_4 і ФР, стабілізовані олеатом натрію (Ол.На) і поліетиленгліколем (ПЕГ), що містять ГЦ та АТ HER2 (Fe_3O_4 @ГЦ/Ол.На/ПЕГ/HER2+ФР). Вивчено цитотоксичну/цитостатичну активність МР щодо клітин лінії MDA-MB-231. Встановлено, що застосування синтезованих МР складу Fe_3O_4 @ГЦ/Ол.На/ПЕГ/HER2+ФР за концентрації магнетиту 0.05 мг/мл, ГЦ – 0.004 мг/мл та АТ HER2 – 0.013 мкг/мл виявляло синергічний ефект та зменшувало кількість життєздатних клітин РМЗ до 51 %. Доведено, що використання МР на основі мішень-спрямованого комплексу Fe_3O_4 @ГЦ/HER2 характеризується підвищеною ефективністю протипухлинної дії, порівняно з традиційним застосуванням препарату ГЦ, при істотному зменшенні (у чотири рази) його дози. Високу цитотоксичну/цитостатичну активність комплексів Fe_3O_4 @ГЦ/HER2 пояснено тим, що в механізмах реалізації апоптичної програми за впливу наноккомпозиту суттєву роль відіграють порушення обміну ендогенного заліза. Так, за наявності в наноккомпозитній системі комплексів Fe_3O_4 @ГЦ/HER2 у клітинах MDA-MB-231 спостерігається значне підвищення рівня «вільного заліза», що сприяє утворенню активних форм кисню та спричиняє оксидативний стрес (реакція Фентона). Наслідками оксидативного стресу є індукція апоптозу, посилення процесів перекисного окиснення ліпідів та структурно-функціональна перебудова біологічних мембран. Показано перспективність подальших досліджень МР

$\text{Fe}_3\text{O}_4@ \text{ГЦ/Ол.На/ПЕГ/HER2} + \text{ФР}$ з метою створення на їхній основі магнітокерованого лікарського засобу для використання в адресній протипухлинній терапії.

Ключові слова: гемцитабін, нанорозмірний ододоменний магнетит, нанокомпозити ядро-оболонка, магнітні рідини, антитіло HER2, адресна протипухлинна терапія

Магниточувствительные нанокомпозиты для адресной противоопухолевой терапии с использованием гемцитабина

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Цель работы – синтез и исследование свойств полифункциональных магниточувствительных нанокомпозитов (НК) и мишень-направленных магнитных жидкостей (МЖ) на основе физиологического раствора (ФР), магнетита, гемцитабина (ГЦ) и антитела (АТ) HER2, перспективных для использования в адресной терапии трижды негативного рака молочной железы (РМЖ) человека – агрессивной опухоли с высокой пролиферативной и метастатической активностью (клетки линии MDA-MB-231).

Удельную поверхность ($S_{уд}$) образцов определяли методом термодесорбции азота с помощью прибора KELVIN 1042 фирмы «COSTECH Instruments». Размер наночастиц (НЧ) оценивали по формуле $D_{ВЕТ} = 6/(\rho S_{ВЕТ})$, где ρ – плотность частицы НК, $S_{ВЕТ}$ – значение удельной площади поверхности, рассчитанной по теории полимолекулярной адсорбции Брунауэра, Эммета и Теллера (БЭТ). Исследование состояния поверхности нанодисперсных образцов осуществляли методами ИК-спектроскопии (Фурье-спектрометр «Perkin Elmer», модель 1720X). Для расчета концентрации гидроксильных групп на поверхности наноструктур использовали метод дифференциального термического анализа в сочетании с дифференциальным термогравиметрическим анализом. Регистрацию термограмм осуществляли с помощью дериватографа Q-1500D фирмы MOM (Венгрия) в интервале температур 20-1000 °C при скорости нагрева 10 град/мин. Рентгенофазовый анализ наноструктур выполняли с помощью дифрактометра ДРОН-4-07 (излучение $\text{Si}_{K\alpha}$ с никелевым фильтром в отраженном пучке, фокусировка по Брэггу-Брентано). Размер и форму НЧ определяли методом электронной микроскопии (просвечивающий электронный микроскоп (ПЭМ) JEM-2100F (Япония)). Петли гистерезиса магнитного момента образцов измеряли с помощью лабораторного вибрационного магнитометра фонеровского типа при комнатной температуре. Измерение оптической плотности, спектров поглощения и концентрации ГЦ в растворах осуществлено методами спектрофотометрического анализа (Spectrometer Lambda 35 UV/Vis Perkin Elmer Instruments). Количество адсорбированного вещества на поверхности магнетита определяли с помощью спектрофотометра при $\lambda = 268$ нм по калибровочному графику. Толщину адсорбированного слоя ГЦ в составе НК $\text{Fe}_3\text{O}_4@ \text{ГЦ}$ определяли методом магнитной гранулометрии. Для изучения прямого цитотоксического/цитостатического действия серии экспериментальных образцов МЖ на основе ФР, НЧ Fe_3O_4 , ГЦ, НК $\text{Fe}_3\text{O}_4@ \text{ГЦ}$, АТ HER2, а также компонентов МЖ в моно- или комплексном применении, на клетки линии MDA-MB-231 in vitro, определяли показатель IC_{50} .

Синтезированы МЖ на основе Fe_3O_4 и ФР, стабилизированные олеатом натрия (Ол.На) и полиэтиленгликолем (ПЭГ), содержащие ГЦ и АТ HER2 ($\text{Fe}_3\text{O}_4@ \text{ГЦ/Ол.На/ПЭГ/HER2} + \text{ФР}$). Изучена цитотоксическая/цитостатическая активность МЖ по отношению к клеткам линии MDA-MB-231. Установлено, что применение синтезированных МЖ состава $\text{Fe}_3\text{O}_4@ \text{ГЦ/Ол.На/ПЭГ/HER2} + \text{ФР}$ при концентрации магнетита 0.05 мг/мл, ГЦ – 0.004 мг/мл и АТ HER2 – 0.013 мкг/мл приводило к синергическому эффекту и уменьшению количества жизнеспособных клеток РМЖ до 51 %. Доказано, что

использование МЖ на основе мишень-направленного комплекса $\text{Fe}_3\text{O}_4/\text{ГЦ}/\text{HER2}$ характеризуется повышением эффективности противоопухолевого действия, по сравнению с традиционным применением препарата ГЦ, при существенном уменьшении (в четыре раза) его дозы. Высокая цитотоксическая/цитостатическая активность комплексов $\text{Fe}_3\text{O}_4/\text{ГЦ}/\text{HER2}$ объяснена тем, что в механизмах реализации программы апоптоза при воздействии наноконкомпозита существенную роль играют нарушения обмена эндогенного железа. Так, при наличии в наноконкомпозитной системе комплексов $\text{Fe}_3\text{O}_4/\text{ГЦ}/\text{HER2}$ в клетках MDA-MB-231 наблюдается значительное повышение уровня «свободного железа», что способствует образованию активных форм кислорода и вызывает оксидативный стресс (реакция Фентона). Последствиями оксидативного стресса являются индукция апоптоза, усиление процессов пероксидного окисления липидов и структурно-функциональная перестройка биологических мембран. Показана перспективность дальнейших исследований МЖ $\text{Fe}_3\text{O}_4@\text{ГЦ}/\text{Ол.На}/\text{ПЭГ}/\text{HER2}+\text{ФР}$ с целью создания на их основе магнитоуправляемого лекарственного средства для использования в адресной противоопухолевой терапии.

Ключевые слова: гемцитабин, наноразмерный однодоменный магнетит, наноконкомпозиты ядро-оболочка, магнитные жидкости, антитело HER2, адресная противоопухолевая терапия

REFERENCES

- Plentz R.R., Malek N.P. Systemic therapy of cholangiocarcinoma. *Visc. Med.* 2016. **32**(6): 427.
- Jain A., Kwong L.N., Javle M. Genomic profiling of biliary tract cancers and implications for clinical practice. *Curr. Treat. Options Oncol.* 2016. **17**(11): 58.
- Internet resource. Gemcitabine. Recent clinical trials of II and III stages in metastatic pancreatic cancer. 2017. [in Russian]. <https://www.meir-health.ru>
- Gutorov S.L. Gemcitabine and pemetrexed: recent results in chemotherapy of solid tumors. *Farmateka.* 2005. **21**: 16. [in Russian].
- Arias J.L., Reddy L.H., Couvreur P. Fe_3O_4 /chitosan nanocomposite for magnetic drug targeting to cancer. *J. Mater. Chem.* 2012. **22**(15): 7622.
- Popescu R.C., Andronescu E., Vasile B.S., Truscă R., Boldeiu A., Mogoantă L., Mogosanu G.D., Temelie M., Radu M., Grumezescu A.M., Savu D. Fabrication and cytotoxicity of gemcitabine-functionalized magnetite nanoparticles. *Molecules.* 2017. **22**(7): 1080.
- Iglesias G.R., Reyes-Ortega F., Checa Fernandez B.L., Delgado Á.V. Hyperthermia-triggered gemcitabine release from polymer-coated magnetite nanoparticles. *Polymers.* 2018. **10**(3): 269.
- Petranovska A.L., Abramov M.V., Opanashchuk N.M., Turanska S.P., Kussyak N.V., Gorbyk P.P. Synthesis and properties of magnetically sensitive nanocomposites based on magnetite and gemcitabine. *Him. Fiz. Tehnol. Poverhni.* 2018. **9**(4): 353.
- Petranovska A.L., Abramov M.V., Opanashchuk N.M., Turanska S.P., Gorbyk P.P., Kussyak N.V., Kussyak A.P., Lukyanova N.Yu., Chekhun V.F. Magnetically sensitive nanocomposites and magnetic liquids based on magnetite, gemcitabine, and antibody Her2. *Him. Fiz. Tehnol. Poverhni.* 2019. **10**(4): 419.
- Gorbyk P.P. Medico-biological nanocomposites with nanorobot functions: state of investigations, development, and prospects of practical introduction. *Him. Fiz. Tehnol. Poverhni.* 2020. **11**(1): 128. [in Ukrainian].
- Abramov M.V., Petranovska A.L., Kussyak N.V., Kussyak A.P., Opanashchuk N.M., Turanska S.P., Gorbyk P.P., Luk'yanova N.Yu., Chekhun V.F. Synthesis and properties of magnetosensitive nanocomposites and ferrofluids based on magnetite, gemcitabine and HER2 antibody. *Funct. Mater.* 2020. **27**(2): 1.
- Levy L., Sahoo Y., Earl B.J. Synthesis and characterization of multifunctional nanoclinics for biological applications. *Chem. Mater.* 2002. **14**: 3715.
- Shpak A.P., Gorbyk P.P. *Nanomaterials and Supramolecular Structures: Physics, Chemistry, and Applications.* (Netherlands: Springer, 2009).
- Gorbyk P.P., Turov V.V. *Nanomaterials and Nanocomposites in Medicine, Biology, Ecology.* (Kyiv: Naukova Dumka, 2011). [in Russian].
- Patent UA 99211. Gorbyk P.P., Petranovska A.L., Turelyk M.P., Turanska S.P., Vasylieva O.A., Chekhun V.F., Luk'yanova N.Yu., Shpak A.P., Korduban O.M. Nanocapsule with nanorobot functions. 2012.
- Gorbyk P.P., Chekhun V.F. Nanocomposites of medicobiologic destination: reality and perspectives for oncology. *Funct. Mater.* 2012. **19**(2): 145.

17. Gorbyk P.P. Nanocomposites with functions of medico-biological nanorobots: synthesis, properties, application. *Nanosystems, Nanomaterials, Nanotechnologies*. 2013. **11**(2): 323. [in Ukrainian].
18. Gorbyk P.P., Lerman L.B., Petranovska A.L., Turanska S.P. Magnetosensitive Nanocomposites with Functions of Medico-Biological Nanorobots: Synthesis and Properties. In: *Advances in Semiconductor Research: Physics of Nanosystems, Spintronics and Technological Applications*. (NY: Nova Science Publishers, 2014).
19. Gorbyk P.P., Lerman L.B., Petranovska A.L., Turanska S.P., Pylypchuk I.V. Magnetosensitive Nanocomposites with Hierarchical Nanoarchitecture as Biomedical Nanorobots: Synthesis, Properties, and Application. In: *Fabrication and Self-Assembly of Nanobiomaterials, Applications of Nanobiomaterials*. (Elsevier, 2016).
20. Pylypchuk I.V., Abramov M.V., Petranovska A.L., Turanska S.P., Budnyak T.M., Kusiak N.V., Gorbyk P.P. Multifunctional magnetic nanocomposites on the base of magnetite and hydroxyapatite for oncology applications. In: *Nanochemistry, Biotechnology, Nanomaterials, and Their Applications (NANO 2017)*. Selected Proc. 5th Int. Conf. "Nanotechnology and Nanomaterials" (Aug. 23–26, 2017, Chernivtsi, Ukraine). P. 35.
21. Abramov M.V., Kusiak A.P., Kaminskiy O.M., Turanska S.P., Petranovska A.L., Kusiak N.V., Gorbyk P.P. Magnetosensitive Nanocomposites Based on Cisplatin and Doxorubicin for Application in Oncology. In: *Horizons in World Physics*. (Nova Publishers, 2017). **293**: 1.
22. Uvarova I.V., Gorbyk P.P., Gorobets' S.V., Ivashchenko O.A., Ulianchenko N.V. *Nanomaterials of Medical Destination*. (Kyiv: Naukova Dumka, 2014). [in Ukrainian].
23. Gorobets' S.V., Gorobets' O.Yu., Gorbyk P.P., Uvarova I.V. *Functional Bio- and Nanomaterials of Medical Destination*. (Kyiv: Kondor, 2018). [in Ukrainian].
24. Kusiak A.P., Petranovska A.L., Gorbyk P.P. Adsorption of Pb^{2+} cations from blood plasma by nanocomposites based on magnetite. *Surface*. 2016. **8**(23): 179. [in Ukrainian].
25. Abramov M.V., Turanska S.P., Gorbyk P.P. Magnetic properties of nanocomposites of superparamagnetic core-shell type. *Metalo fizyka i Novitni Tekhnologiyi*. 2018. **40**(4): 423. [in Ukrainian].
26. Abramov M.V., Turanska S.P., Gorbyk P.P. Magnetic properties of fluids based on polyfunctional nanocomposites of superparamagnetic core-multilevel shell type. *Metalo fizyka i Novitni Tekhnologiyi*. 2018. **40**(10): 1283. [in Ukrainian].
27. Karaman O.M., Fedosova N.I., Voyeikova I.M., Cheremshenko N.L., Ivanchenko A.V., Savtsova Z.D. Prospects of application of lectins for diagnostics and treatment of malignant neoplasms. *Onkologiya*. 2018. **20**(1): 10. [in Ukrainian].
28. Turanska S.P., Opanashchuk N.M., Petranovska A.L., Kusiak N.V., Tarasiuk B.I., Gorobets' S.V., Turov V.V., Gorbyk P.P., Abramov M.V. Synthesis, properties of nanocomposites based on gemcitabine, and application in oncotherapy. *Poverkhnia*. 2019. **11**(26): 577. [in Ukrainian].
29. Moiseenko V.M. Probabilities of monoclonal antibodies in treatment of malignant tumors. *Practical Oncology*. 2002. **3**(4): 253. [in Russian].
30. Tan M., Yu D. Molecular mechanisms of erbB2-mediated breast cancer chemoresistance. *Adv. Exp. Med. Biol.* 2007. **608**: 119.
31. Santin A.D., Bellone S., Roman J.J., McKenney J.K., Pecorelli S. Trastuzumab treatment in patients with advanced or recurrent endometrial carcinoma overexpressing HER2/neu. *Int. J. Gynaecol. Obstet.* 2008. **102**(2): 128.
32. Borisenko N.V., Bogatyrev V.M., Dubrovin I.V., Abramov N.V., Gayevaya M.V., Gorbyk P.P. Synthesis and Properties of Magnetosensitive Nanocomposites Based on Oxides of Iron and Silicon. In: *Physics and Chemistry of Nanomaterials and Supramolecular Structures*. (Kyiv: Naukova Dumka, 2007). **1**: 394. [in Russian].
33. Mornet S., Vasseur S., Grasset F., Veverka P., Goglio G., Demourgues A., Portier J., Pollert E., Duguet E. Magnetic nanoparticle design for medical applications. *Prog. Solid State Chem.* 2006. **34**(2–4): 237.
34. Petranovska A.L., Abramov N.V., Turanska S.P., Gorbyk P.P., Kaminskiy A.N., Kusiak N.V. Adsorption of cis-dichlorodiammineplatinum by nanostructures based on single-domain magnetite. *J. Nanostruct. Chem.* 2015. **5**: 275.
35. Abramov N.V., Turanska S.P., Kusiak A.P., Petranovska A.L., Gorbyk P.P. Synthesis and properties of magnetite/hydroxyapatite/doxorubicin nanocomposites and magnetic fluids based on them. *J. Nanostruct. Chem.* 2016. **6**: 223.
36. Freshni R.Ya. *Animal Cell Culture: Practical Manual*. (Moscow: BINOM, Laboratoriya znania, 2010). [in Russian].
37. Luk'yanova N.Yu. Doctoral (Biol.) Thesis. (Kyiv, 2015). [in Ukrainian].

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