Physico-chemical studies of sorption materials based on biomass waste

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Abstract. This article discusses the possibility of using cheap raw materials for the treatment of wastewater from heavy metal ions using residual biomass and agricultural waste (sunflower husk, millet, buckwheat). Residual biomass is formed after the extraction of valuable components from microalgae and duckweed. The authors proposed to modify the sorption material using heat treatment and the introduction of additives, such as chitosan and thermally expanded graphite. Chitosan allows you to get sorption materials in the form of granules, which are convenient to use. The physicochemical properties of sorbents were studied. The sorption capacity for the obtained materials was from 5.0 to 32.0 mg/g.

1 Introduction

The problem of wastewater treatment is relevant for the whole world. The greatest attention is paid to cheap water purification methods. The manufacture of sorption materials from waste allows us to solve two problems at once: water purification and waste disposal. Many research papers have been published in which it is proposed to use waste from various industries as sorption materials [1,2,3].

In the works of Finnish scientists studied the modification of peat and sawdust with hydrochloric and citric acid [4].

Turkish scientists have studied the use of tea waste as a biosorbent. The optimum pH and temperature were selected for the extraction of copper and nickel ions, which were 5 and 50 °C, respectively. The samples used for treatment of wastewater to remove the heavy metal ions have been obtained by the following procedure: pre-waste of tea was washed and dried in an oven at a temperature of 105 °C. The sorption capacity for nickel ions of 10.8 mg/g and for copper ions of 14.9 mg/g was achieved [5-6].

Known work [7-8], in which it is proposed to use as a sorbent feathers of Dromaius novaehollandiae (DNF) were collected from poultry and chitosan. As a result of this technology, it is possible to obtain a biopolymer composite having a sorption capacity 70.42 mg/g.

Coffee grounds are a cheap sorption material for the treatment of wastewater from lead and fluoride [9]. The exhausted coffee grounds gathered from industrial wastes have been acid-activated and examined for their adsorption capacity. The maximum cure efficiency of nickel (II) ions was found at pH 5.0. The adsorption equations were well correlated by the Langmuir equations and the Freundlich isotherm. The maximum sorption capacity was 254.3 mg/g for a chitosan-based biosorbent coated with a silicon layer [10].

Marine plants are also widely used to extract heavy metal ions of zinc and copper [11,12]. The use of living algae is not advisable, therefore it is proposed to use dry algae as a sorbent. The walls of algae cells contain mainly cellulose and proteins associated with polysaccharides. Cell walls play a key role in the removal of heavy metals from water. In different types of algae, the composition of the cell wall is different, hence the ability to bind metal ions.

As a rule, cell walls are rich in functional groups such as carboxyl, amines, hydroxyls, phosphates, imidazoles and sulfates. These groups are negatively charged and attract positively charged metal ions. However, at low pH, they are partially protonated and lose some strength to absorb positive ions. In [13], the

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microalga strain Chlorella Sorokinianian was used, biomass was grown in cultivators, then dried in a drying cabinet and used to extract gallium ions. The sorption capacity was 38.5 mg/g.

The study [14] directed the study of the biosorption of lead, cadmium, copper and arsenic ions using natural algae. The biosorption of these metals is based on an ion exchange mechanism, accompanied by the release of light metals such as calcium, magnesium and sodium. The pH value, initial concentrations and extraction temperature were studied.

To improve the sorption properties, it is proposed to carry out the modification of sorption materials [15,16]. The dried algae U. fasciata to be carbonized is impregnated with solution of chloride salts such as calcium chloride for 24 h [17]. Accordingly, sufficient quantities were soaked well with 10 % chloride solution of 5 L capacity, respectively, so that the solution get well adsorbed for a period of 24 h. At the end of 24 h the excess solution was decanted off and air dried. Then, the materials were placed in muffle furnace carbonized at 400 °C. The dried materials were powdered and activated in a muffle furnace kept at 800 °C for a period of 10 min. After activation, the carbon was washed sufficiently with 4 N HCl to remove the cations. Then, the materials were washed with plenty of water to remove excess acid, dried and powdered.

2 Materials and methods

The object of the study was the residual biomass of Lemma minor duckweed and Chlorella sorokiniana microalgal, which is formed after the extraction of valuable components, such as lipids, pigments, pectin substances [18].

We have previously studied the problems of the cultivation of the biomass of microalgae Chlorella sorokiniana and duckweed Lemma minor as a source of valuable components. Residual biomass, which is formed as waste, requires further utilization. It is proposed to use the residual biomass as a sorption material for the purification of wastewater from heavy metal ions (zinc, cadmium, lead). The material is loose and inconvenient during operation, therefore, to increase the sorption capacity and shape, it is proposed to add chitosan.

TEG and millet husk are widely known as sorbents for the extraction of heavy metals. This article compares the sorption properties of sorbents of different composition.

Granules of 3 types were obtained by the technology described below:

Sample No1 – thermally expanded graphite (TEG) (5 g). Residual microalgae biomass (RMB) (5 g) – chitosan (100 ml);

Chitosan 40 g was dissolved in 3% acetic acid 960 ml. The mixture is stirred for 4-5 h 3% ours until complete dissolution of chitosan, add RMB (5 g), further, the process of preparation is the same as that of the sample No1.

Sample No2 – RMB (5 g) – chitosan (100 ml). Chitosan 40 g was dissolved in 3% acetic acid 960 ml. The mixture is stirred for 4-5 h 3% ours until complete dissolution of chitosan, add RMB (5 g), further, the process of preparation is the same as that of the sample No1.

Sample No3- RMB (5 g) – chitosan (100 ml) – millet husk (5g). Chitosan 40 g was dissolved in 3% acetic acid 960 ml. The mixture is stirred for 4-5 h 3% ours until complete dissolution of chitosan, add RMB (5 g) and millet husk (5 g), further, the process of preparation is the same as that of the sample No1.

The manufacture of samples No 2, 3 differs only in the composition of the added additives.

Sample No 0- Residual microalgal biomass without modification.

Chitosan- is an amino saccharide, a polysaccharide derivative. macromolecules consist of randomly-linked β-(1–4) D-glucosamine units and N-acetyl-D-glucosamine (Figure 1).

Residual biomass and agricultural waste were previously thermally treated in a special steel cell with limited oxygen access. The residual biomass was burned at a temperature of 400 °C for 20 minutes and the waste of agriculture at a temperature of 300 °C for 20 minutes.

Structure of adsorbents by mean of scanning electron microscope Zeiss Leo 1530.

Specific surface of the granulated sorbents are study using NOVA 4000e. Surface area & Pore Size Analyzer, brand Quantachrome.

![Fig. 1. Structural formula of chitosan.](image)

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\[
\frac{p/p_0}{a(1 - p/p_0)} = \frac{1}{a_nC} + \frac{(C - 1)p/p_0}{a_nC}
\]

where \(p/p_0\) is the ratio of the pressure in the system to the condensation pressure, \(a\) is the amount of adsorption, \(a_n\) -is the volume of the monolayer on the surface of the adsorbent, \(C\) -is the ratio of the adsorption equilibrium constants in the first layer and the condensation constant.

The residual concentration of heavy metal ions was determined by the voltammetric method (Russian standart 14.1.2:4.222-06) on the device TA-Lab. The accuracy of the measurement procedure is 5%. To calculate the sorption capacity, model solutions were prepared with a concentration of 1 g/l. The sorption capacity was calculated as the difference between the initial and equilibrium concentrations, taking into account the mass of the sorbent and the volume of the
solution. The accuracy of the measurement procedure is 5%.

To modify the residual biomass after the extraction of valuable components, heat treatment is used, which allows to obtain a material with high sorption properties

3 Results and Discussion

To determine the sorption capacity, sorption was carried out for 20 minutes, at the rate of 2 g of sorption material per 1 liter of a model solution containing heavy metal ions (zinc, cadmium, lead). The mixture was stirred using a laboratory shaker. The calculation of the sorption capacity was carried out according to the formula (1) and the results are presented in table 1.

\[ q = \frac{Co - Ce}{m} \cdot V \]  

Where q is the sorption capacity, Co and Ce are the initial and final concentrations of heavy metals in the solution, V(L) is the volume of solution and m (g) is the waste amount.

From table 1 it can be seen that the sorption material No. 3 has the highest sorption capacity; its specific surface is 2.720 m²/g. The specific surface is not large, but the layered structure suggests that the extraction of metal ions occurs due to chemisorption and physical sorption. From microstructural studies, it follows that the samples have a layered structure characteristic of an additive made of chitosan. The sorption properties are also due to surface irregularities (cracks).

Table 1. Sorption properties of granular materials.

| Type                  | q, mg/g | Pb²⁺ | Cd²⁺ | Zn²⁺ |
|-----------------------|---------|------|------|------|
| Residual microalgal biomass Sample No 0 | 13.0    | 10.0 | 5.0  |
| Sample No 1           | 22.0    | 17.0 | 16.0 |
| Sample No 2           | 28.5    | 16.0 | 26.3 |
| Sample No 3           | 32.0    | 22.4 | 27.0 |

Sample No. 1 illustrates the worm-like structure and packed layers of expanded graphite. As a result of thermal expansion, furrows or open channels can serve as pores and facilitate the adsorption of metals in them.

At the same time, the presence of chitosan in the analyzed adsorbents shows a porous and rough structure, which can provide a higher surface area and higher adsorption properties. As its chemical analogue (cellulose), chitosan has a similar fiber morphology with distinct crystallites and holes in the shape of domes.

Figures 2 show micrographs of an adsorbent from millet husks of irregular shape. The surface structure appears to be heterogeneous, which can be explained by the presence of lignin in this adsorbent. On its outer surface there are cavities that can be called irregular and inhomogeneous. Some of the pore holes and cracks may contribute to the flow of model solutions into the pores.

Fig. 2. Surface morphology sample No1 (a) and sample No2 (b), sample No3 (c).

For a comparative analysis of the porosity of the samples, the specific surface area was determined by the BET method. (table2).

Table 2. The specific surface of the granulated sorbents.

| Granular sorbent              | S sp, m²/g |
|-------------------------------|------------|
| thermally expanded graphite   | 1.093      |
| Residual microalgal biomass – |            |
| chitosan                      | 1.599      |
| Residual microalgal biomass – |            |
| chitosan- millet husk         | 2.720      |

From table 2 it can be seen that the specific surface values are not large, and the highest value for sample No 3. It is possible that sorption processes occur due to chemical sorption. The specific surface area is consistent with the value of sorption capacity for all samples.

4 Conclusions

1. It is proposed to use as a sorption material the residual biomass obtained after the extraction of valuable components from microalgae Chlorella sorokiniana and Lemma minor duckweed.
2. Granulated materials were obtained with the addition of thermally expanded graphite, chitosan and agricultural waste.
3. The sorption properties of the materials obtained were studied, the values of the sorption capacity for lead, cadmium and zinc ions were calculated.

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