

Treatment Wetland effluent quality improvement by usage sorbents of various origin

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Abstract. TWs are finding as low-cost and also effective system to treat domestic wastewater. The aim of this study is to find suitable adsorbing material for application in an additional treatment unit where treatment wetlands do not provide sufficient PO_4^{3-} reduction level. Material M1 is a fine-grained by-product of thermal treatment of carbonate-siliceous rock (opoka) with high content of calcium carbonate CaCO_3 in temperature 700°C. Particular disadvantage of material M1 can be very high level of pH (11-12) and causing alkaline solution. Material M2 is lanthanum-modified bentonite (LMB) and was already used for reducing content of phosphorus compounds in lakes and other water bodies to treat effects of eutrophication process. Lanthanum-modified bentonite consists mostly SiO_2 and Al_2O_3 (~80%). The study was conducted on effluent from small (60 pe) treatment wetland (TW) in northern Poland where Hybrid Treatment Wetland (HTW) system was provided. Sorption capacity of material M1 was low and equal 0.9 mg/g. For material M2 this parameter reached value of 2.1 mg/g. Lanthanum-modified bentonite (M2) has shown better abilities to reduce phosphates from wastewater that material M1 – material containing mostly calcium oxide. Also performed study revealed that presence of others contaminations in wastewater is causing difficulties in phosphates removal.

1 Introduction

In recent years, the popularity of treatment wetlands (TWs) has been increasing as an alternative solution for rural areas and small towns [1,2]. Locating conventional (explain) wastewater treatment plants (WWTPs) in such places is not economically justified and septic tanks used in households usually leak contamination into groundwater. Thus, TWs is a low-cost and effective system to treat domestic wastewater. It has been proven that the most efficient treatment is provided by Hybrid Treatment Wetland (HTW) where the system is composed of at least two beds: subsurface vertical flow bed (SSVF) and subsurface horizontal flow bed (SSHF) [2,3]. The first stage ensures suitable conditions for the nitrification process, while the second stage ensures suitable conditions for the denitrification process, removal of organic matter and suspended solids. According to Kadlec and Wallace (2009) [4] also in the SSVF bed denitrification processes occurs, due to saturation of the bed at the bottom layers. The effect of HTW (combined VF-HF system) is focused especially on nitrogen compounds removal [2,3,5]. Unfortunately, the removal efficiency of phosphorus compounds concentration in the effluent from HTWs as well as in small WWTP below 2000 PE (person equivalent) is very often inadequate [1,6].

Eutrophication is caused by phosphorus compounds as well as nitrogen. Frequently, nutrients removal concentrates only on nitrogen, although phosphorus has

a similar effect on water bodies. Considering the current recommendations [7], which concerns about Baltic Sea protection to prevent eutrophication effects, the 70% reduction of phosphorus must be provided already for small wastewater treatment plants up to 300 PE. Baltic Sea Action Plan obligates Poland to over 40% reduction in phosphorus emission.[7].

An alternative, but recently interesting method to remove phosphorus compounds from various type of wastewater is applying materials with high sorption capacity. A few sources of adsorbents can be distinguished: (1) anthropogenic, such as AAC (autoclaved aerated concrete) [8], LECA [9], and Pollytag [10] (2) natural, such as rock opoka [11, 12] and POLONITE [13], as well as (3) by-product sources eg. fly ash [14] and blast furnace slag [15]. By-products and waste materials are adsorbents which should arouse interest due to the idea of circular economy and efficient phosphorus recovery will follow this concept, due to increasing awareness of its supply running out. It is estimated that phosphorus supplies may be exhausted within the next 100 years.

Thus, the study concerns finding a suitable adsorbing material for application in an additional treatment unit where treatment wetlands do not provide sufficient PO_4^{3-} reduction level. The aim of the performed investigation was to compare two different materials, the first one is well known material with proven effectiveness for lakes restoration but relatively expensive one, the second one is waste material with P removal potential. The idea of

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this study was to assess their applicability by kinetic equation and quality of the effluent but also to attempt fulfil the assumptions of circular economy.

The design of the experiment is multistage. In the first stage the applicability and comparison of the two materials (one well know and expansive and the second not investigated yet and the byproduct/reject material from Rockfos® production). They have been already publish some preliminary results from second stage of investigation concerning the specific features of the constructed wetlands – it was predicted that both materials should be used in posttreatment in special facility with adequate contact time and sedimentation [16,17].

2 Materials and methods

2.1. Materials

Rockfos® is produced in the process of decarbonisation (firing at high temperature ~700°C) of natural silica rock-carbonate. This material is highly reactive to phosphorus compounds as it contains large amounts of elements such as calcium, silicon, aluminium and iron (Fig. 1). According to the producer's requirements, it can be used to build P-filters to remove phosphorus and bacteriological contaminants from domestic and industrial wastewater [18].

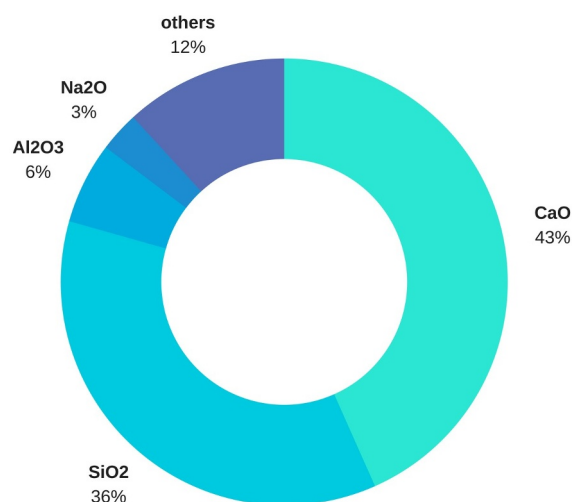


Fig. 1. Chemical composition of Rockfos®

Material M1 is a fine-grained by-product of Rockfos® production. This fraction (0-2 mm) consists mainly of calcium oxide CaO (over 80%), resulting from the decomposition of CaCO₃ during the heating process. A major disadvantage of M1 is its very high pH (11-12) making the solution alkaline [18]. Testing a waste material (byproduct) is related to the definition of circular economy.

Material M2 is a lanthanum-modified bentonite clay (LMB) developed by the Land and Water Division of Australia's CSIRO (Commonwealth Scientific and Industrial Research Organisation) and this particular

adsorbing material has already been used to reduce the phosphorus compounds content in lakes and other water bodies to treat the effects of eutrophication [19]. In the reaction of binding phosphates by lanthanum from LMB with the molar ratio 1:1, a rare-earth complex known as rhabdophane is formed [20]. Also, lanthanum concentration in the solution remains very low, due to locking La particles in the bentonite clay structure [21]. M2 consists mostly of SiO₂ and Al₂O₃ (Fig. 2). The LMB pH is close to neutral (7.0-7.5) but can be successfully used in a wide pH range from 5 to 9 [22-24].

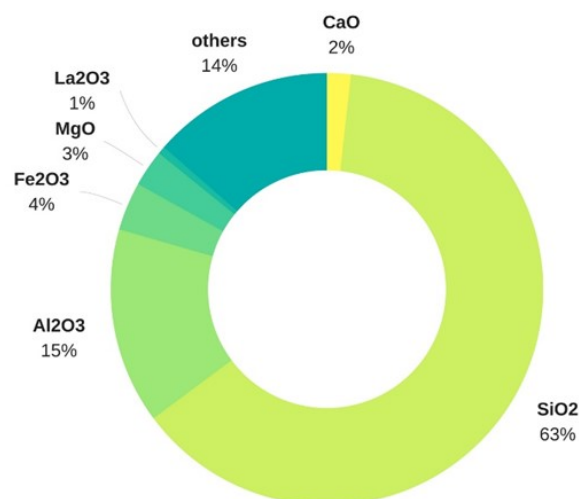


Fig. 2. Chemical composition of material M2

Several studies have been developed to prove nontoxic character of the lanthanum-modified bentonite clay and, in conclusion, to classify LMB as not hazardous [25-27].

2.2. Methods

The study was conducted on an effluent from a small wastewater treatment wetland (WTW) for 60 pe in northern Poland where Hybrid Treatment Wetland (HTW) system was provided (coordinates: 54°39'50"N; 18°07'13"E). Sedimentation and flotation processes in the mechanical part of the treatment (three-chamber sedimentation tank) are followed by a biological system (filtration, sorption, oxidation, and reduction processes). The assumed conception is based on two stages: subsurface vertical flow bed and subsurface horizontal flow bed. The natural environment and its biodiversity provide a successful process of treatment, that the effluent from HTW can be directed to the drainage system.

Samples of the wastewater for research were taken after the mechanical treatment and before the discharge to the drainage system (outflow/effluent). The basic parameters of the wastewater measured before and after HTW are presented in Table 1.

Table 1. Average physical and chemical indicator of wastewater quality in the inflow and effluent from HTW, Pomerania, Poland

parameter	unit	inflow	effluent from HTW	reduction rate
Temperature	°C	20.8	18.9	-
pH	-	7.17	7.17	-
Conductivity	µS/cm	1656	1226	-
Color	mgPt/L	392	34	91 %
Turbidity	mg/L	104.4	6.5	94 %
TSS	mg/L	28.0	9.0	68 %
PO ₄ ³⁻ -P	mg/L	26.6	4.34	84 %
COD	mg/L	329.0	20.8	94 %
NH ₄ ⁺ -N	mg/L	138.9	18.9	86 %
NO ₃ ⁻ -N	mg/L	0.75	36.1	-
NO ₂ ⁻ -N	mg/L	0.13	0.42	-

The results indicate that HTW provides relatively high efficiency of contaminants removal, although the pollution load of discharged wastewater still does not meet the Polish and European requirements Polish Regulation (Year 2014, item 1800) [28], Council Directive 91/271/EEC (European Commission, 1991) [29]. The study concentrates on removal of phosphates in TWs despite the regulations report total phosphorus. Form of phosphorus which occurs mostly in wetland are phosphates, both in organic and inorganic compounds. It has been proven that free orthophosphate is the only bioavailable form for plants in wetlands beds [30]. Moreover, the performed research was conducted on the TW effluent, thus most of organic phosphorus was removed in mechanical part of the treatment system.

The described HTW is located near Puck Bay. This specific water body is characterized with a high contamination level, as it is closed with the Hel Peninsula thus having limited connection with the Baltic Sea.

Experimental trials were carried out in steady conditions. To batch reactors filled with 1.0 L of effluent from HTW, 0.5 g of a selected adsorbing material was added and mixed for 5 minutes. Sampling was done after 5, 10, 30, 60, 120 and 300 minutes to define the materials sorption capacity and to assess their adsorption kinetics by the pseudo-second order model. The parameters such as pH, conductivity (EC), color, turbidity, and total suspended solids (TSS) were measured before and after the addition of the adsorbing materials to evaluate their influence over the wastewater quality. The concentration of PO₄³⁻ was analysed by HACH Lange cuvette tests, while others parameters were measured using HACH Lange DR 3900 spectrophotometer and WTW Multi 350i compact precision portable meter. The laboratory studies were conducted at ambient temperature.

2.3. Kinetic study – pseudo-second order model

The adsorption kinetics data analysis of both materials were presented with the pseudo-second order model, which allows predicting the sorption capacity and indicates constant parameters.

Sorption capacity of phosphates for each material was calculated according to equation (1) [31]:

$$q_e = (C_0 - C) / m \cdot V \quad (1)$$

where: q_e – sorption capacity [mg/g], V – volume of solution [L], C_0 – initial concentration of PO₄-P [mg/L]; C – final concentration of PO₄-P [mg/L], m – mass of sorption material [g].

The kinetics was investigated at solution pH. The formula to describe the adsorption kinetics is the following [23,32,33]:

$$dq/dt = k \cdot (q_e - q_t)^2 \quad (2)$$

where: q_t – adsorption capacity at time t [mg/g], q_e – adsorption capacity at equilibrium [mg/g], k – equilibrium rate constant of pseudo-second order reaction [g/mg·min].

Integrating equation (2) for the boundary condition $t = 0$ to $t = t$ and $q_t = 0$ to $q_t = q_t$, the formula becomes:

$$1/(q_e - q_t) = 1/q_e + k_t \quad (3)$$

From equation (3) linear form can be obtained:

$$t/q_t = 1/k \cdot q_e^2 + t/q_e \quad (4)$$

The analysis of a straight plot of t/q_t against time (t) allows calculating the value of parameters k , q_e and the coefficients of determination R^2 .

3 Results and discussion

The results indicate different efficiency of phosphates removal for both materials. In the performed study, the initial phosphates concentration of TW effluent was approx. 2.5 mg/L. The obtained sorption capacity of M1 was low and equal to 0.9 mg/g. The final concentration of PO₄³⁻ was close to 2.1 mg/L after 5 hours contact time which provides 17% phosphates reduction efficiency. The dose of the adsorbing material was relatively low (0.5 g), although the value of removed PO₄³⁻ was below expectations.

Various tested materials with high contents of calcium oxide presented a significant ability to remove phosphorus compounds [9, 12, 31, 34]. In the previous study performed on synthetic wastewater, the sorption capacity of this particular material reached 9.6 mg/g. The research proceeded in steady conditions with a short contact time of 1 hour. Although the described laboratory trials were conducted with an artificial P-solution (KH₂PO₄) [16], such observations may reveal a possible influence of wastewater quality and the concentration of other compounds (eg. of nitrogen or organic matter) on the removal efficiency of phosphates.

The kinetic data simulated by the pseudo-second order model fitted the experiment results well. For M1,

the coefficient of determination (R^2) was equal to 0.95 and correlation coefficient (r_{xy}) was 0.97, as shown in Figure 3. The equilibrium q_e value was 0.90 mg/g (Table 2), which was nearly the same as the sorption capacity obtained from the experimental values.

Table 2. Kinetic parameters for pseudo-second order model of phosphates adsorption

material	q_e [mg/g]	k [g/mg*min]	R^2	r_{xy}
M1	0.9	1.24	0.95	0.97
M2	2.1	0.23	0.99	0.99

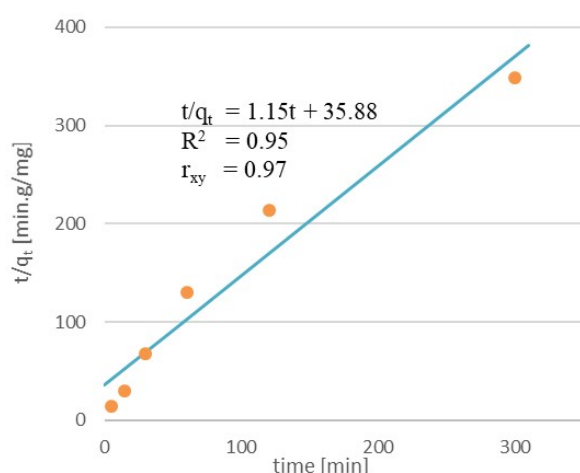


Fig. 3. Adsorption kinetics for pseudo-second order model of PO_4^{3-} onto material M1

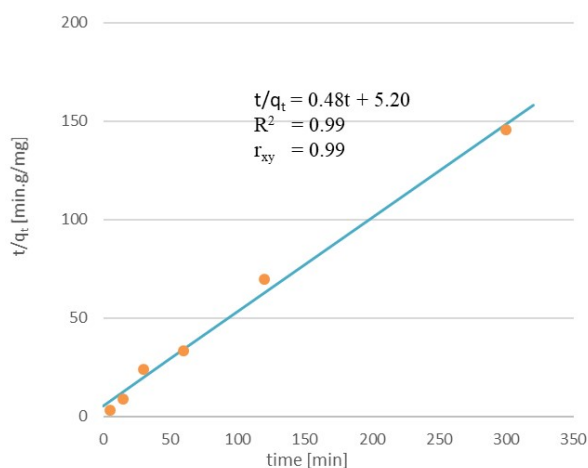


Fig. 4. Adsorption kinetics for pseudo-second order model of PO_4^{3-} onto material M2

For M2 the sorption capacity was equal to 2.06 mg/g, while the concentration of PO_4^{3-} dropped to 1.5 mg/L during the experiment. The effectiveness of phosphates removal reached 41% with the same dose and contact time. Similarly to M1, theoretical ability (obtained from research on synthetic wastewater) of M2 to reduce PO_4^{3-} was significantly higher. The study conducted by Kasprzyk et al. (2018) [16] gave the sorption capacity of M2 at the level of 9.1 mg/g. While, according to

Haghseresht et al. (2009) [23], the adsorption capacity of M2 cannot exceed 10.6 mgP/g. From the results obtained by Kurzbaum and Bar Shalom (2016) [35], the maximum sorption capacity of LMB reached 14.4 ± 4.7 mg PO_4 /g. Also Ross et al. (2008) [24] observed a decrease of phosphates reduction efficiency of M2 while using it for lake water treatment, and pointed to the presence of humic acid.

Also in the case of M2 pseudo-second order model, both the coefficient of determination (R^2) and correlation coefficient (r_{xy}) were close to 1.0 (Fig. 4) and displayed satisfactory matching of the kinetic data. The equilibrium q_e was equal to the sorption capacity calculated from the research data (Table 2).

The effect of both materials on the pH values was quite different. M1 caused a relatively high increase of pH of the solution despite a small dose of the sorption material used (0.5 g). The initial pH of the HTW effluent was close to neutral and equal to 7.2. Already after 5 minutes of the contact time, pH level reached almost 8.4 and during further sampling, pH slowly increased to the value of 8.6 after 5 hours of sedimentation (Fig. 5).

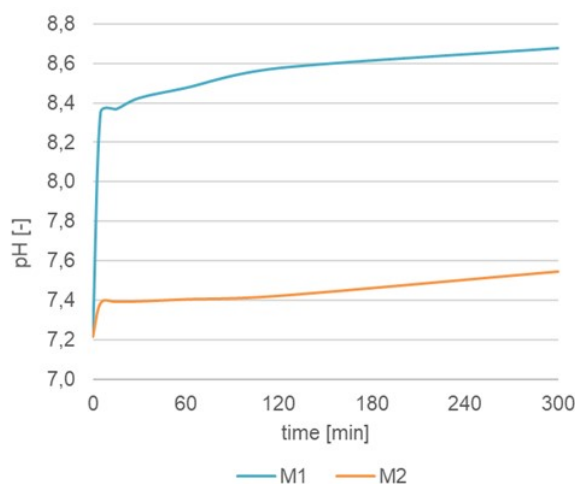


Fig. 5. Influence of adsorbing materials on pH of the effluent from HTW

This particular disadvantage of M1 is related to a significant content of calcium oxide, although its presence determines the ability to remove substantial amounts of PO_4^{3-} [36, 37]. The research described by Jucherski et al. (2017) [11], reports on the application of alkaline filtration materials for phosphorus removal, where a material with high contents of Ca (~50%) caused an increase of the solution pH to almost 13. Also, the authors pointed out that such materials may be used as the last stage in a small treatment facility on condition the effluent pH value is less than 8.5. Possible applications of M1 require adjusting pH to an acceptable level.

In the case of M2, the effect on pH was slight. An insignificant increase to the value of 7.55 was observed (Fig. 5). According to Douglas et al. (2016) [19] and Van Oosterhout and Lürling (2013) [38], M2 caused only a minor increase or had no effect on the solution

pH. The performed studies usually concentrate on how pH influences phosphorus removal efficiency using LMB, rather than how the material impacts the solution, considering pH level as one of the most important parameters of the adsorption processes [23, 24, 33]. No impact on the pH values is an extremely important advantage, although the solution pH may determine the effectiveness of M2 (with the optimal pH range of 5-7).

The initial conductivity of the HTW effluent was equal to 928 $\mu\text{S/cm}$. M1 has shown a meager increase to the value of 945 $\mu\text{S/cm}$ at the beginning of the contact time, and after 1 hour of sedimentation, the conductivity stabilized at the level of 941 $\mu\text{S/cm}$. M2 displayed almost no influence on EC (Fig. 6). Nevertheless, it should be pointed out that small doses of the reactive materials were used. The application of larger amounts of those adsorbents (especially M1) may have a negative effect on the quality of the effluent.

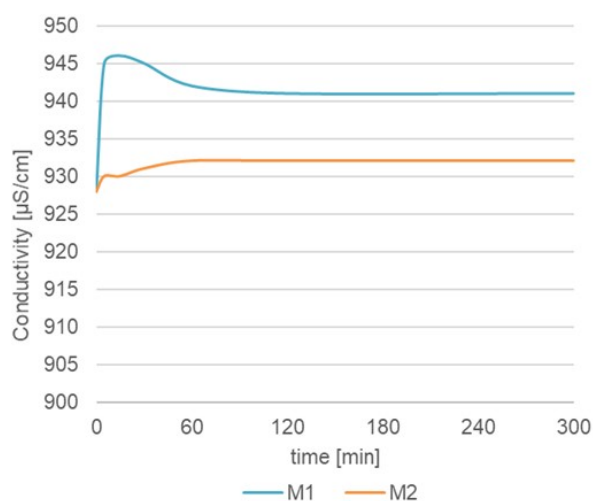


Fig. 6. Influence of adsorbing materials on EC of the effluent from HTW

Both the tested materials did not significantly affect the color and turbidity of the effluent from the HTW. Initial values were equal to 24 mgPt/L and 5.3 mg/L respectively. During the contact time, slight fluctuations of those parameters were observed. At the end of the experiment, the color and turbidity for M1 was equal to 36 mgPt/L and 7.1 mg/L, while for M2 – 38 mgPt/L and 8.1 mg/L respectively.

In the case of total suspended solids (TSS in the first taken samples (after 5 minutes of sedimentation), a high increase of the TSS values was observed for both materials (from 5 mg/L to approx. 23 mg/L). This effect could be caused by the amount of dissolving matter. It was necessary to extend the sedimentation time. After 5 hours, in both cases the values of TSS were close to initial.

In case of long term use of those materials, the maintenance costs will be entirely different due to their futures like pace of origin eg. M2 is from Australia what means costly transportation which also create big carbon food print. Assumed total daily inflow of the wastewater in Kniewo HTW was equal to 7.0 m^3/d which yields

2,500 m^3 per year. Based on the effluent phosphates concentration, the load of $\text{PO}_4\text{-P}$ can reach 10 kg/year. The sorption capacity of M1 and M2 achieved from the study was equal to 0.9 and 2.1 mg/g respectively, which concludes with the usage over 10 tons of M1 and almost 5 tons of M2.

The cost of M1 usage despite to twice the M2 requirement might be significantly lower. M1 is a by-product and waste material with no specific purpose. On the other hand, the M2 unit price may reach 2750 €/ton [39] and gives an annual maintenance cost equal to 14,000 € (Table 3). Possibility of material recovery or reuse need further analysis.

Table 3. Maintenance and cost analysis

	sorption capacity	material required	price	annual cost	costs per m^3 of treated effluent
	mg/g			€	€/m ³
M1	0.9	~ 10 tons per year	1 €/ton + transport cost	~ 700	0.3
M2	2.1	~ 5 tons per year	2750 €/ton	~ 14 000	6

4 Conclusions

The results presented in this study indicate a better ability to reduce phosphates from the HTW effluent displayed by lanthanum-modified bentonite (M2) than M1 – a material containing mostly calcium oxide. Obtained from the kinetic data, the equilibrium sorption capacity (q_e) of M1 was low and equal to 0.9 mg/g, while of M2 – 2.1 mg/g. The pseudo-second order model fitted the experiment results of both materials M1 and M2 well, due to the coefficient of determination (R^2) values equal to 0.95 and 0.99 respectively and high correlation coefficient (r_{xy}) for M1 – 0.97, for M2 – 0.99.

The performed study also revealed that the presence of other contaminants in wastewater causes difficulty in phosphates removal. Due to this fact their influence on the processes of phosphates binding should be verified and subsequently an appropriate dose of the adsorbing material for proper wastewater treatment should be established.

The influence of M1 on the pH of HTW effluent was substantial as it increased from the initial value 7.2 to 8.6. It was due to the high content of calcium oxide. As a result, the application of M1 gives an alkaline solution. The other parameters showed no significant impact on the quality of the HTW effluent, for both materials.

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