

Urban Wastewater Treatment by Adsorption of Organic Matters on Modified Bentonite by (Iron-Aluminum)

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Abstract

In this research, the natural bentonite clay (from Maghnia, western Algeria) was purified (Na⁺⁻montmorillonite, CEC = 91 meq/100 g), noted (puri.bent) and modified with mixed hydroxy-Fe-Al (FeAl-PILC). The purified bentonite clay and FeAl-PILC were heated at 383 K for 2 hr and characterized by the chemical analyses data, XRD, and N₂ adsorption to 77 K techniques. Puri.bent and FeAl-PILC were applied to fix the organic matter (OM) present in urban wastewater from the city of Sidi Bel-Abbes (western Algeria). The adsorption of organic matter was followed by spectro-photometry at 470 nm, and the adsorption data were a good fit with Freundlich isotherm for puri.bent but for FeA-IPILC, were well fit by Elovitch isotherm model. The maximum adsorption capacity (q_m) was 571.6 mg/g for puri.bent and 1120.69 mg/g for FeAl-PILC. The degree of OM removal was 67% for puri.bent and 97% for FeAl-PILC. FeAl-PILC can be considered as a promising adsorbent for the removal of OM from wastewater.

Keywords

Adsorption, Urban Wastewater, Bentonite Clay, Isotherm Models

1. Introduction

In Algeria, recycled wastewater effluent was an important source of irrigation water, about 500 million m³ of wastewater effluent was produced per year [1]. Wastewater treatment plants (STEP) of Sidi Bel-Abbes city, lo-

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cated on the Mekerra River (western Sidi Bel-Abbes) at the exit of the city have the capacity for treating 300,000 Eq/H, purifying the liquid part of the sewage and attempt to remove most pollutants, excess nutrients and pathogens from wastewater [2]. Wastewater effluent treated discharged into river and the leftover solids and semi solids were filtered from wastewater making up the "sewage sludge some" [2] [3].

Wastewater effluent undergoes five major processes: primary treatment, secondary treatment, disinfection and finally sludge treatment, before the treated wastewater, disinfected and discharged into Mekerra River. However, treated wastewater contains higher concentrations of suspended and dissolved organic and inorganic matter was only minimally removed from the effluent [4] [5]. Adsorption on solid substrates, such as clay or activated carbon, was one of the methods, which have been used for removing OM from wastewater.

Bentonite clay which was available in large quantities can be used as an adsorbent particularly FeAl-PILC, for the removal of many aromatic organic pollutants from wastewater [6]-[10], heavy toxic metals [11] [12], and colors [13].

This study aimed the removal of the organics matters from urban wastewater treatment plant (STEP) from Sidi Bel-Abbes city by purified bentonite clay and FeAl-PILC.

2. Materials and Methods

2.1. Purified Bentonite Clay

The natural bentonite (from Maghnia, west Algeria) was fractionated by sedimentation to obtain the <2 μ m montmorillonite rich faction. The purification was prepared in laboratory [2], the carbonates were removed by sodium acetate/chloridric acid, iron oxides by sodium thiosulfate/sodium chloridric and organic materials by hydrogen peroxide (30% vol). To ensure complete transformation into the sodium from all samples, they were washed several times with 0.5 M NaCl. The important physico-chemical properties reported in Table 1 with the SiO₂/Al₂O₃ ratio equal 2.63 indicated that the major mineral was the montmorillonite.

2.2. Preparation of Solution of Hydroxyl-(FeAl) Inter-Layered

The pillaring solutions of hydroxy-Al and hydroxy-Fe were prepared separately according to the procedure previously reported in literature [14]. 0.207 M AlCl₃ was hydrolysed with 0.207 M NaOH from in solutions, with OH/Al ratio of 2.5 and a pH of 4.3 - 4.5. The final concentration of the Al-solution was 0.09 M, the hydroxyl-Al solution was aged at room temperature during 6 days.

For hydroxyl-Fe solution, by slowly adding a 0.1 M solution aqueous of 0.1 M NaOH to a solution of 0.1 M FeCl₃ to obtain final acidic ratios OH/Fe³⁺ = 2.5, pH of the final solution ([Fe³⁺] = 0.045 M) was close 1, *i.e.* The resulting solution was aged for 30 days at room temperature.

2.3. Modified Bentonite by Mixed Hydroxyl FeAl (FeAl-PILC)

The mixing solution hydroxy-Fe and hydroxy-Al (50% - 50%, in % atomic) was slowly added under vigorous

Table 1. Chemical-physical properties of bentonite clay.					
Chemical	Composition (%	mass)			
Element	SEM	EDS	Physical Characteristics		
SiO ₂	57.96	52.51	Loss on ignition (%)	12.19	
Al_2O_3	22.05	21.25	Density	1	
Fe_2O_3	2.83	2.25	Granulometry (µm)	2 to 1	
CaO	8.49	15.0	Capacity swelling Cg	8.27	
Cl ₂ O	1.138	1.25	pH for 10 g/L	10.7	
MgO	2.37	2.37	Conductance (μS) for 0.5 g/L	66.4	
Na ₂ O	1.91	1.25	CEC (meq/g)	0.90	
K_2O	2.84	1.25			
TiO ₂	0.12	2.50			

stirring to purified bentonite clay taking M^{3+} ($M^{+3} = Fe^{3+}$, Al^{3+})/clay ratio equal 0.065 and allowed to age overnight. Then, the product was collected by filtration and washed with deionized doubly distilled water until free of chlorides, as indicated by the AgNO₃ test. The solid was dried overnight at 110°C, ground and sieved.

2.4. Characterisation Studies

The clay mineralogy were analyzed by, X-ray Diffraction (XRD), the solids were calcined at 110°C for overnight, and measured by Philips PW 1729 powder-diffractometer, with CoK α radiation at 35 Kv, 30 mA, passing a Fe filter. To precisely confirm the presence of the (0 0 1) reflection peak in each sample, the XRD patterns in the 2θ range 2° - 14° were also taken at steps of 0.02° with scanning rate 0.05°/min. Specific surface areas—BET, was determined by adsorption of nitrogen at 77 K, using a Micromeritics ASAP 2010 instrument. The samples were previously outgassed by treatment at 120°C for 2 h, under flow of nitrogen. The FTIR spectra were obtained using a Mattson-1000 FTIR spectrometer. The samples were prepared by mixing 1 mg of sample with 100 mg of KBr from pellets.

2.5. Adsorption Equilibrium Experiment

An mass *m* (g) of adsorbents was put into a 50 mL reactor, into which OM of urban wastewater with different initial diluted concentrations, ranging from 3.16, 1.92, 3.72 and 4.53, for periods (03/06/2012, 08/09/2012, 13/04/2013, 19/12/2013), respectively, **Table 2** were added to reach a final volume of 50 ml. These dilutions were such that the initial concentration of OM was 126.4 mg/l for four periods, or a dilution average of 3.33 times. At T = 25°C and P = 1 atm the suspension was stirred for 30 min, time of adsorption equilibrium (Z. Meçabih *et al.*, 2006). After, by decanting for 2 h, two-thirds of the supernatant liquid was removed and filtrated through a membrane (1.2 μ m) fiber glass filter. The concentration of the organic matter (OM) in 30 ml the supernatant was titrated by KMnO₄, 10⁻³ M. This titration was followed by sepctrophotometric (Speckol 10) at 470 nm (λ_{max} of OM absorption) [2]. All experiments were carried out in triplicates (test 3 × 4 periods) and the average values were reported in **Figure 1(a)**, **Figure 1(b)**, show the variation of the optical density of MO as a function of solution volume KMnO₄ 10⁻³ M.

3. Results and Discussion

3.1. Characterisation of Bentonite Clay and FeAl-PILC

3.1.1. XRD Analysis

The basal spacing d_{001} of two samples after calcinations at 383 K, and the specific surface areas were presented in **Table 3**. As seen in **Figure 2**, the d_{001} position of FeAl-PILC was 14.50 Å in comparison to Na⁺-montmorillonite which was 12.79 Å. The increase in basal spacing was due to pillaring [15]. The increase of the interlayer distance (d_{001}) after pillaring with hydroxyl FeAl-PILC was typical [10].

Table 2. Urban wastewater characteristics of Sidi Bel-Abbes city samples.						
	Period					
Characteristic	1: removal of	2: removal of	3: removal of	4: removal of		
	03/06/2012	08/09/2012	13/04/2013	19/12/2013		
MES (mg/L)	675	417	798	939		
MO (mg/L)	398.9	243.1	469.8	572.8		
DCO (mg·O ₂ /L)	338	320	410	998.65		
$DBO_5 (mg \cdot O_2/L)$	179.25	160.6	210.87	530		
Nitrogenize (mg/L)	76.2	32.96	54.12	83		
Nitrate-nitrite (mg/L)	13-0.08	15-0.11	7-1.43	12-0.35		
Phosphate total (mg/L)	19.66	25.80	42.15	47.09		
Temperature (°C)	25	25	26	25		
PH	7	7	7.5	7.5		



solution volume, KMnO₄ of: (a) Purified bentonite; (b) FeAl-PILC. \Box 10 mg, O 15 mg, Δ 20 mg, ∇ 25 mg, \Diamond 30 mg.



Figure 2. Basal spacing (d_{001}) of: (a) Purified bentonite; (b) FeAl-PILC at 383 K.

3.1.2. BET Analysis

The specific surface area, calculated from the N_2 adsorption isotherms by the use of the Brunauer-Emmett-Teller (BET) equation, increased from 100.20 $\text{m}^2 \cdot \text{g}^{-1}$ for puri.bent to 277.16 $\text{m}^2 \cdot \text{g}^{-1}$ for FeAl-PILC Table 3. This increase after pillaring with FeAl-PILC, suggest also the increasing of the micropores [7].

3.1.3. FTIR Analysis

The FTIR spectra of purified bentonite (puri.bent) and modified clay FeAl-PILC samples were presented in Figure 3. The region of interest for determining structural composition was the frequency region between (1400 -400) cm⁻¹ which was characteristic of metal bonded silica [16] [17]. The broad absorption bands observed at 3446 - 3628 cm⁻¹ represent the fundamental stretching vibrations of different -OH groups present in Mg-OH-Al (3640 cm⁻¹), Al-OH-Al (3620 cm⁻¹) and Fe-OH-Al (885 cm⁻¹) units in the octahedral layer [17]. The most intensive band between 115 - 1020 cm⁻¹ was related to antisymmetric stretching of the \equiv Si-O-Si \equiv bridge. The deformation mode was placed at 798 cm⁻¹. The dolomite and quartz were identified by infrared spectroscopy as the main admixtures of clay raw material that was used as the source of montmorillonite. For montmorillonite this band ought to be evident at 1100 cm⁻¹ according to Besson et al., 1987 [17]. The bands at 525, 620 and 917 cm⁻¹ were assigned to Al-O-Si deformation, Si-O stretching, and Al-Al-OH deformation, respectively [16].



Figure 3. FT-IR spectra of: (a) Purified bentonite; (b) FeAl-PILC.

These bands slightly shift to high wavenumber in the FT-IR spectra of (Fe-Al)-pillared bentonite at 627.96, 790.22 and 919.76 cm⁻¹ respectively. The bands at 1036.51 cm⁻¹ assigns to Si-O-Si stretching vibrations and band at 472.96 cm⁻¹ assigns to the Si-O bending and M-O stretching vibrations. These bands slightly shift to high wavenumber in IR spectra of Fe-Al pillared bentonite, which can be indirect evidence of the incorporation of Fe³⁺ into Al-polycation, because the Fe-O bond is longer than the Al-O bond, so that the substitution of ions Al³⁺ by Fe³⁺ should be accompanied by a general shift to high wavenumber. Bands Si-O-Si, Si-O, and M-O were the most sensitive to these substitutions.

3.2. Adsorption of Organic Matters from Urban Wastewater

Table 2 illustrates the average values of the various parameters determined ($T = 22^{\circ}C \pm 2^{\circ}C$) on the sample of urban wastewater. The concentration of OM was determined by calcination at ($625^{\circ}C \pm 5^{\circ}C$), for periods (1 - 4), 59.1, 58.3, 58.9, respectively with 61.0% of suspended solids (TS). In the period 4 (**Table 2**), the wastewater treatment plant (STEP) of Sidi Bel-Abbes city's, received the effluent flood and wineries waste very charged in organic matters OM. The values of the chemical oxygen demand (COD) indicated that the urban wastewater of Sidi Bel-Abbes city's, industrial quality.

The BOD₅/COD, around 0.52 indicates that the organic matters in the sample of urban wastewater can be biodegradable. Than more, pH around 7 was favourable for biological treatment, and the effluent of urban wastewater as for irrigation were feasible.

3.3. Isotherm Models

All isotherms were obtained at pH 7 with initial OM concentrations 126.4 mg/l with different masses of clay ranging from 10 - 30 mg. The experimental data were fitted to Freundlich and Elovitch models using Equations (1) and (2), respectively. The parameters of the Freundlich and Elovitch models and correlation coefficient (R^2) values for each clay type were presented in **Table 4**. The data were adequately fitted by the two models which were consistent with reported results [2]. However, the Langmuir equation was not obeyed by OM adsorption data for both puri.bent and FeAl-PILC.

$$q = \frac{x}{m} = K_F \times C^{\frac{1}{n}}$$
(1)

$$\frac{q}{q_m} = K_E \times \exp\frac{q}{q_m} \tag{2}$$

where *x* and *m* (g) were masses of adsorbed and adsorbent respectively, *C* was the equilibrium concentration of the adsorbent (mg·L⁻¹). *q* (mg·g⁻¹) and q_m (mg·g⁻¹) were capacity adsorption and maximum capacity adsorption respectively which were related to θ : recovery rate adsorption sites Equation (3). K_F and *n* were Freundlich constants; and K_E was Elovich constant (L·g⁻¹). *n* was related to energy of adsorption Q_0 Equation (4) constant of digital distribution energy of sorptive sites, *R* universal gas constant and *T* absolute temperature.

$$\theta = \frac{q}{q_m} \tag{3}$$

$$n = \frac{Q_0}{RT} \tag{4}$$

The residual concentrations at equilibrium of the MO, diluted urban wastewater treated by both Na^+ -montmorillonite and FeAl-PILC clay, were shown in Figure 4. It was found that the concentration of M.O decreased with an in-

Table 4. Estimated isotherm parameters for OM adsorption.							
	Samples	Freundlich			Elovitch		
		п	$K_F(\mathbf{L}\cdot\mathbf{g}/\mathbf{L})$	R^2	$K_E (L \cdot g/L)$	$q_m (\mathrm{mg} \;\mathrm{OM/g})$	R^2
	Puri.bent	1.09	70.91	0.98	-	-	-
	FeAl-PILC	-	-	-	0.099	378.79	0.99



crease in adsorbent mass. This was expected, for an adsorbent mass of 25 mg for puri.bent and 10 mg for FeAl-PILC clay by 15.01 cm³ of raw urban wastewater were sufficient to remove most of the OM from the raw urban wastewater, fixation rates of MO per unit mass of the adsorbent were shown in **Table 5**, FeAl-PILC clay showed an excellent potential for the removal of organic matter from wastewater with a rate 97%.

Organic matters adsorption data were very well described by the Freundlich equation for puri.bent (Figure 5), however, the adsorption data of the FeAl-PILC was good fitted to the Elovitch equation (Figure 6).

The value of the Freundlich parameter k_F and *n* were 70.91 and 1.09 respectively, the magnitudes of k_F and *n* showed easy sorption of OM from the diluted wastewater and indicate favourable adsorption. The intercept k_F value was an indication of the adsorption capacity of the adsorbent, which reflects the number of sorptive sites. The slope 1/n indicates the effect of concentration on the adsorption capacity and represents adsorption intensity, parameter characterizing energetic heterogeneity of the adsorption surface, *i.e.*, surface with non-energetically equivalent sites. As seen from **Table 4**, *n* value > 1 indicated favourable adsorption can be explained by assumption of multilayer adsorption phenomena, energy of digital distribution of sorptive sites Q_0 equal 2669.9 J·g⁻¹, Equation (1) and Equation (2). This value corresponding to the conditions the residual concentration at equilibrium of MO was 34 mg·L⁻¹ or 90 mg (*x*) of organic matter adsorbed on 25 mg (*m*) of clay (**Figure 4**, **Figure 5**). Freundlich isotherm fitted well with the correlation coefficient (R^2) of 0.98.

The adsorption data for FeAl-PILC clay were also evaluated according to the Elovitch isotherm Equation (2), this model also assumes a multilayer behavior for the adsorption of OM onto FeAl-PILC, with recovery rate adsorption sites 0.91 and fixation rate of OM equal 97% (Table 5). The value of Elovitch constant ($K_E = 0.099 \text{ L/g}$) and the maximum adsorption capacity ($q_m = 378.79 \text{ mg} \cdot \text{MO} \cdot \text{g}^{-1}$) were listed in Table 4 and the plot of this isotherm is shown in Figure 6. The correlation coefficient (R^2) of 0.99 obtained showed that adsorption of OM also followed Elovitch isotherm.

The Freundlich isotherm was more widely used but provides no information on the maximum adsorption capacity (q_m) for puri.bent but a polynomial regression of order 2 (**Figure 7**) indicates that was, 571 mg·MO·g⁻¹ and 1120 mg·MO·g⁻¹ for FeAl-PILC. Therefore the adsorptions of organic matters were on monolayer for puri.bent and, on multilayer for FeAl-PILC.

4. Conclusions

In this work, hydroxy-Fe and hydroxy-Al, mixed have been used as pillaring species to modify the Algerian natural bentonite (Na⁺-montmorillonite, C E C = 91 meq/100 g) for adsorption of organic matters from the effluent from wastewater treatment plant (STEP) of Sidi Bel-Abbes city. The study results demonstrated that increase in basal spacing (d₀₀₁), and specific surface area occurs in the presence of the mixed hydroxy-FeAl (OH/M = 2.5), as mentioned in the literature. Experimental results were in good agreement with Freundlich and

Samples	% OM fixation	Mass (g)	Mass (g)/1 L of raw urban wastewater	
		50 ml of diluted urban wastewater/15.01 ml of raw urban wastewater	Hydroxyl (50/50%, % atom)	Clay
Pruri.bent	67.16	25	0.00	1.67
FeAl-PILC	97	10	3.75	0.6





Figure 5. Linear plot of Freundlich isotherm for purified bentonite.







Figure 7. Adsorption capacity (q) for: ■ purified bentonite; ● FeAl-PILC.

Elovitch isotherm models, and had showed a good fitting to the experimental data. Adsorption of OM on puribent obeys Freundlich model with good correlation.

 $(R^2 \approx 0.98)$ and the value of Freundlich parameter, n = 1.09 confirmed that the adsorption was favourable. But the adsorption on FeAl-PILC was fitted to the Elovitch equation, with R^2 value ≈ 0.99 . The adsorptions of organic matters were on monolayer for puri.bent and, on multilayer for FeAl-PILC. The maximum adsorption capacity (q_m) was determined by the polynomial regression of order 2 of the capacity adsorption (q) as a function of the mass of the adsorbent, at 571.6 for puri.bent and 1120.69 mg·g⁻¹ for FeAl-PLC. It was evident that this last sample due to the fixation rate of OM equals 97%.

Finally as a result this bentonite modified clay, by the mixed hydroxy-iron and aluminum, was the best performance which could be successfully applied in the urban wastewater treatments.

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