

SOLID-STATE SYNTHESIS AND PHYSICO-CHEMICAL CHARACTERIZATION OF MODIFIED SMECTITES USING NATURAL CLAYS FROM BURKINA FASO

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ABSTRACT. Solid-state intercalation was applied to prepare organo-smectites using cationic surfactants and natural clays containing smectite at various surfactant/smectite ratios. The surfactants enter the interlayers of smectites causing a swelling of the clays. The used clays were collected in Siétougou and Diabari villages located in the eastern part of Burkina Faso. The solid-state intercalation was successful for all four surfactants applied, *n*-dodecyltrimethylammonium bromide, (*n*-C₁₂H₂₅(CH₃)₃NBr), *n*-tetradecyltrimethylammonium bromide, (*n*-C₁₄H₂₉(CH₃)₃NBr), *n*-hexadecyltrimethylammonium bromide (*n*-C₁₆H₃₃(CH₃)₃NBr) and di-*n*-dodecylidimethylammonium bromide ((*n*-C₁₂H₂₅)₂(CH₃)₂NBr) at different levels of the cation exchange capacity (CEC). The synthesized organo-smectites were characterized regarding relative density, structural and textural properties. XRPD data showed a systematic increase of the basal spacing of the unit cell of the smectite up to >38 Å at 2.0-CEC loading of the surfactant. This increase indicates that the surfactants penetrated into the smectite interlayers with the surfactants being arranged parallel to the layers at low concentrations and almost perpendicular at high ones. FTIR spectra of the organo-smectites showed a decrease in the intensities of the water bands at around 1630 cm⁻¹ and 3400 cm⁻¹, and new specific bands close to 2920 cm⁻¹ and 2850 cm⁻¹ were assigned to the asymmetric and symmetric stretching of CH₂ groups of the surfactants, respectively.

KEY WORDS: Smectite, Organo-smectite, Surfactant, Intercalation

INTRODUCTION

Smectite clay minerals are abundant in nature, and they are known to have swelling properties due to their ability to take up water and swell. Surface modification and intercalation of clay minerals have received significant attention as it allows to prepare materials with properties suitable in a wide range of applications at low cost. Several routes to modify clays have been developed to be used in applications as adsorption, ion exchange with inorganic and organic cations, acidification, calcination, and re-aggregation of smectites [1, 2]. Generally, the reported scientific papers in this field describe the preparation of the organoclays at laboratory scale using different techniques and chemical conditions, different type of clays, and a range of surfactants as reviewed by Paiva *et al.* [2]. For modification, e.g. clays quaternary ammonium salts are the most frequently used compounds, but *n*-alkyl-pyrrolidones, maleic anhydride, biomolecules, polymeric quaternary alkylammonium, alkyl-imidazolium and phosphonium salts have also been used for this purpose [2-6]. The areas of application of modified clays are wide including nanocomposites, adsorbents of organic and inorganic pollutants in soil, water, air, etc. [2, 7-9]. Clays and chemically modified clays have extensively been used as adsorbents in environmental systems. The most commonly used clay types in such applications are smectites due to their high CEC, swelling properties, large surface areas, and good adsorption and absorption properties [2, 8, 10, 11].

Intercalation of cationic surfactants into the 2:1 sheets, e.g. smectites changes their surface properties from hydrophilic to hydrophobic [12, 13] because of the replacement of hydrated

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metal cations between the sheets by cationic surfactants. This is seen through a decrease in the intensities of the water bands in the infrared region at around 1630 cm^{-1} and 3400 cm^{-1} [12, 13] and new bands related to the surfactants. Transmission and scanning electron microscopy analyses (TEM and SEM, respectively) have been applied to study the macrostructural features of modified clays [14-16]. The TEM analyses showed the packing density of modified clays within the interlayer space, and SEM micrographs showed the difference in surface morphology of untreated and modified clays [14-16]. Thermogravimetry (TG) and differential scanning calorimetry (DSC) have been applied to get information on the thermal behavior of modified clays [17, 18]. The result of the modification of clays is seen as swelling, thus, the basal spacing between the sheets has increased [2, 14, 19]. As a result, organoclays exhibited better adsorption capacity, e.g. organic contaminants [2, 20, 21]. According to literature, the behavior and properties of organoclays are strongly related to their structure and the chemical properties of the intercalated compound between the sheets [1, 2, 14, 22]. Numerous previous studies have shown that the d_{001} spacing of the modified clays is strongly related to the length and number of the alkyl chains and the packing density of the intercalated surfactants of the modified clays [22, 23, 24].

Modified clays are normally prepared in aqueous solution by cation exchange or solid-state reactions. Organic molecules can be intercalated in dried clay minerals by solid-state reactions without use of solvents and this makes the preparation procedure environmentally friendly and more suitable for industrialization [2, 25]. Solid-state intercalation is therefore to be preferred for modified clays [25, 26]. However, so far it has been less employed than procedures in aqueous solution [2, 12, 14]. The first solid-state intercalation of alkylammonium cations into clay minerals was reported by Ogawa *et al.* [25, 27]. Although solid-state intercalation reactions showed several advantages, so far a limited number of studies are reported on the synthesis of modified clays by solid-state reactions [2, 27, 28]. The use of intercalation reactions of organic compounds into clays and clay minerals using solid-state reactions has been described elsewhere [2, 26].

In the present work, the solid-state intercalation method was applied to the preparation of organoclays because of simple handling and relatively low cost [25, 26, 28]. Although different types of clays from several regions and suppliers have been used to prepare modified clays, further studies using solid-state intercalation technique to introduce organic compounds into clays and clay minerals are relevant from both scientific and applied point of view. Since the chemical composition of smectite clays varies from one deposit to another, more studies are needed to assess their usefulness as cheap and effective materials with sorption capacity regarding removal of organic contaminants in different environmental applications. To the best of our knowledge, although large deposits of smectite clays, which constitute an important mineral resource, are located in the Eastern part of the country, they have not yet been used for organoclays preparation. These deposits of smectite clays being largely available for the synthesis of modified clays, by intercalating cationic surfactants between the raw clay sheets, a wide range of applications could be considered at low cost in the country.

Four cationic surfactants $n\text{-C}_{12}\text{H}_{25}(\text{CH}_3)_3\text{N}^+$, $n\text{-C}_{14}\text{H}_{29}(\text{CH}_3)_3\text{N}^+$, $n\text{-C}_{16}\text{H}_{33}(\text{CH}_3)_3\text{N}^+$ and $(n\text{-C}_{12}\text{H}_{25})_2(\text{CH}_3)_2\text{N}^+$ ions as bromide salts have been used in solid-state intercalation reactions with two natural clays of smectite type collected in the Eastern part of Burkina Faso. Clays modified with quaternary alkylammoniums derivatives such as *n*-hexadecyltrimethylammonium and *n*-dodecyltrimethyl-ammonium bromides have been reported in a number of studies [12, 21, 24, 29, 30, 31], those intercalated with *n*-tetradecyltrimethylammonium and di-*n*-dodecyldimethylammonium bromides have only been the subject of a limited number of studies [21, 23, 24, 29, 32].

The present work was undertaken to investigate the solid-state intercalation of three alkyltrimethylammonium and one di-alkyldimethylammonium cations into two natural smectite clays collected from large deposits located in the Eastern part of Burkina Faso, and the

subsequent changes of structure and physicochemical properties. The relationship between the d_{001} basal spacing of the modified clays and the length, the number of long alkyl chains in the quaternary alkylammonium ions has been one of the main objectives of this study. Furthermore, the interlayer configuration of intercalated surfactants in the two smectite clays has been elucidated. The variation of the relative density of the organoclays depending on different loading of surfactants has been determined, as this has been scarcely studied [33].

EXPERIMENTAL

Materials

The natural clays used in this study are from Siétougou and Diabari villages, denoted AH and DI, respectively, located in the Eastern region of Burkina Faso at the following coordinates, AH: latitude = 11°54'43.2'', longitude = 00°38'22.7'', and DI: latitude = 11°51'19.9'', longitude = 00°25'08.07''.

The used surfactants *n*-dodecyltrimethylammonium bromide ($n\text{-C}_{12}\text{H}_{25}(\text{CH}_3)_3\text{NBr}$, (C_{12})), *n*-tetradecyltrimethylammonium bromide ($n\text{-C}_{14}\text{H}_{29}(\text{CH}_3)_3\text{NBr}$, (C_{14})), *n*-hexadecyltrimethylammonium bromide ($n\text{-C}_{16}\text{H}_{33}(\text{CH}_3)_3\text{NBr}$, (C_{16})) and di-*n*-dodecyldimethylammonium bromide ($(n\text{-C}_{12}\text{H}_{25})_2(\text{CH}_3)_2\text{NBr}$, (2C_{12})), were purchased from Sigma-Aldrich ($\geq 98\%$ purity). Silver nitrate (AgNO_3 , Sigma-Aldrich, $\geq 99.8\%$ purity) was used as purchased to test the presence of the bromide ions in the washing water after treatment of clays with the surfactants.

Organosmectites synthesis

The synthesis of organoclays was undertaken by the following procedure: 4 g of dried clay (powder size, $\leq 80\ \mu\text{m}$) and a well-defined amount of surfactant, which represents a level of 0.5 CEC, 1.0 CEC, 1.5 CEC and 2.0 CEC of the used clay, was placed in an agate mortar. The mixture was ground for 10-15 min to get a homogeneous powder. The treated clays were washed with distilled water to get rid of the bromide ions as checked by the addition of an aqueous solution of silver nitrate, dried at room temperature, and further dried in oven at 100 °C for 4 hours. The dried modified clays were ground in an agate mortar and stored in closed bottles. The modified clays were labelled as surfactant loading-type of surfactant-original location of the clay, e.g. 0.5 CEC- C_{12} -AH.

Characterization techniques

Cation exchange capacity was determined by adsorption of copper ethylenediamine $\text{Cu}_2(\text{en})_2^{2+}$ complex [34]. The CEC values were determined at 34.2 meq/100 g and 25.4 meq/100 g for the raw clays from AH and DI, respectively.

The relative densities of raw and modified clays are calculated by dividing the mass of the samples by the mass of water in a calibrated 5.0 mL beaker, weighed in the same way as possible.

The elemental composition of raw clays was determined by using an ICP-OES (Inductively Coupled Plasma Optical Emission Spectroscopy) Optima 7300V HF Perkin Elmer instrument. A well-defined quantity of the sample was mixed with hydrochloric acid and nitric acid in a microwave oven for complete dissolution of the sample and the elementary composition was determined.

X-ray powder diffraction (XRPD) patterns of the samples were recorded between 2 and 62° with a step size of 0.2° in scattering angle, 2θ , by a θ - θ goniometer as described elsewhere [35] using $\text{Mo}(\text{K}_{\alpha 1})$ radiation, $\lambda = 0.71073\ \text{\AA}$.

Fourier transform infrared spectroscopy (FT-IR) was used to characterize the samples in this study. The raw clays, organoclays and pure surfactants were mixed with dry KBr and pressed

into discs with 2 mg of sample and 200 mg KBr in each tablet. The data were recorded on a Perkin Elmer Spectrum 100 FT-IR spectrometer over the spectral range of 400 cm^{-1} - 4000 cm^{-1} .

RESULTS AND DISCUSSION

X-ray diffraction of raw clays

The XRD diffractograms of the AH and DI raw clays are shown in Figure 1.

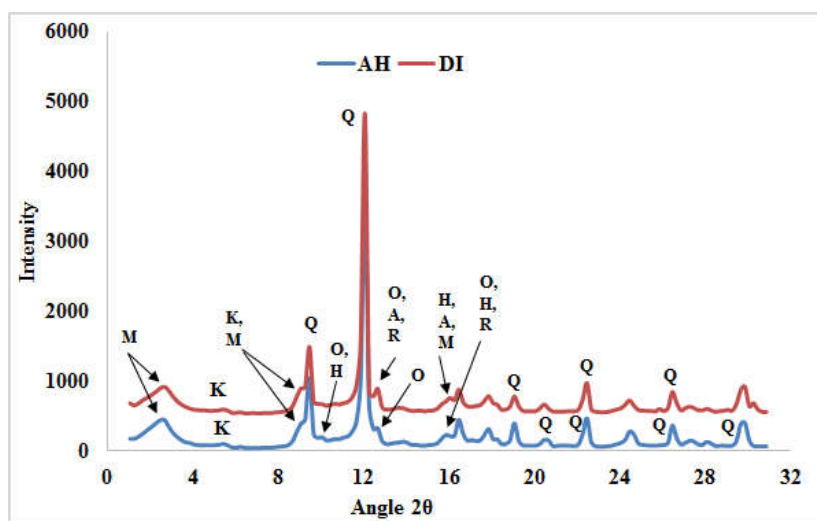


Figure 1. X-ray powder diffraction patterns of AH and DI raw clays: Q = quartz, M = Montmorillonite, K = Kaolinite, H = Hematite, R = Rutile, O = orthoclase and A = Anorthite.

The diffractograms show that the raw clays contain montmorillonite, kaolinite, quartz, anorthite, orthoclase, hematite and rutile. According to literature studies, the d_{001} basal spacing values reported for Na-montmorillonite are the following: 12.0 Å and 11.78 Å, Cubuk *et al.* [30]; 12.0 Å, Caglar *et al.* [31]; 12.96 Å, Tetsuka *et al.* [36]. In the case of Ca-montmorillonite, the d_{001} values reported are the following: 15 Å, He *et al.* [29]; 15.72 Å, Tetsuka *et al.* [36]; 15.4 Å, Zhang *et al.* [37]. In the present study, the d_{001} basal spacing of the smectite phase is seen at 15.3 Å (2.266° denoted M in Figure 1). This d_{001} basal spacing value (15.3 Å) is closer to the d_{001} basal spacing of Ca-montmorillonite than the one corresponding to Na-montmorillonite, indicating that the AH and DI raw clays contain Ca-montmorillonite as the dominant smectite mineral clay. AH and DI clays contain different amounts of montmorillonite with a higher content in AH according to the cation exchange capacity and the XRPD patterns, see above. The presence of montmorillonite, which belongs to the smectite clay group, in the raw clays is advantageous for the intercalation of surfactants as smectite clays have a high cation exchange capacity and large surface areas, which confer strong adsorption/absorption properties. Generally in the literature, Na-montmorillonite, Ca-montmorillonite and bentonite are the mineral clays that are used for the preparation of organoclays. Compared to the existing literature, our study focuses on raw clays containing Ca-montmorillonite, kaolinite, orthoclase and anorthite as mineral phases, with Ca-montmorillonite being the major phase.

Chemical analysis of raw clays

The elemental composition of the raw clays was determined and the oxide content is summarized in Table 1. The high values of the $\text{SiO}_2/\text{Al}_2\text{O}_3$ ratio for each raw sample (5.27 for AH and 5.70 for DI) can be explained by the presence of bentonite (montmorillonite) which has a ratio at around 3.78 [38] and the presence of a highly free silica.

Table 1. Chemical composition (expressed as mass %) of the AH and DI clay samples.

AH clay								
Oxide	Al_2O_3	SiO_2	Fe_2O_3	CaO	NiO	P_2O_5	MgO	SrO
Mass %	12.7	67.1	8.54	1.12	0.03	0.04	0.96	0.01
Oxide	Cr_2O_3	K_2O	SO_3	MnO	TiO_2	ZrO_2	Na_2O	$\text{SiO}_2/\text{Al}_2\text{O}_3$
Mass %	0.04	0.61	0.02	0.16	1.61	0.12	0.18	5.27
DI clay								
Oxide	Al_2O_3	SiO_2	Fe_2O_3	CaO	NiO	P_2O_5	MgO	SrO
Mass %	12.4	70.4	5.26	1.40	0.02	0.04	0.87	0.02
Oxide	Cr_2O_3	K_2O	SO_3	MnO	TiO_2	ZrO_2	Na_2O	$\text{SiO}_2/\text{Al}_2\text{O}_3$
Mass %	0.02	0.77	0.02	0.13	0.93	0.10	0.59	5.70

Relative density measurements

The relative densities of the samples are summarized in Table 2. These results show a systematic decrease of the relative density with increasing level of surfactant loading which means that the clays swell with the intercalation of the surfactant. The relative density of the organoclays also decreases with the length and number of the long alkyl chains of the surfactants. According to Yevtushenko *et al.*, there is an inverse relationship between porosity and the relative density of soil, which means that the decrease of the relative density causes an increase in soil porosity [39]. Although porosity measurements were not carried out, the decrease of the relative density of the organoclays could indicate that their porosities are higher than the ones of the corresponding untreated samples [39]. The swelling and increased hydrophobicity is certainly the reason for the difficulty to grind the organoclays with high surfactant loading level. Therefore, a simple way to confirm the intercalation of the surfactant molecules into the clays is relative density measurements.

Table 2. Relative densities of the raw and modified clays.

Surfactant loading level	Surfactants							
	C_{12}		C_{14}		C_{16}		2C_{12}	
	AH	DI	AH	DI	AH	DI	AH	DI
0.0 CEC	1.55	1.50	1.55	1.50	1.55	1.50	1.55	1.50
0.5 CEC	1.46	1.46	1.38	1.39	1.23	1.36	1.46	1.47
1.0 CEC	1.37	1.28	1.26	1.25	1.20	1.20	1.18	1.18
1.5 CEC	1.34	1.27	1.28	1.19	1.15	1.13	1.09	1.20
2.0 CEC	1.35	1.29	1.27	1.20	1.14	1.19	1.09	1.11

X-ray diffraction of the organoclays

The XRPD diffractograms of the DI and AH raw and modified clays prepared at different loading levels (0.5 CEC, 1.0 CEC, 1.5 CEC and 2.0 CEC) using the C₁₂, C₁₄, C₁₆ and 2C₁₂ surfactants are shown in Figure 2.

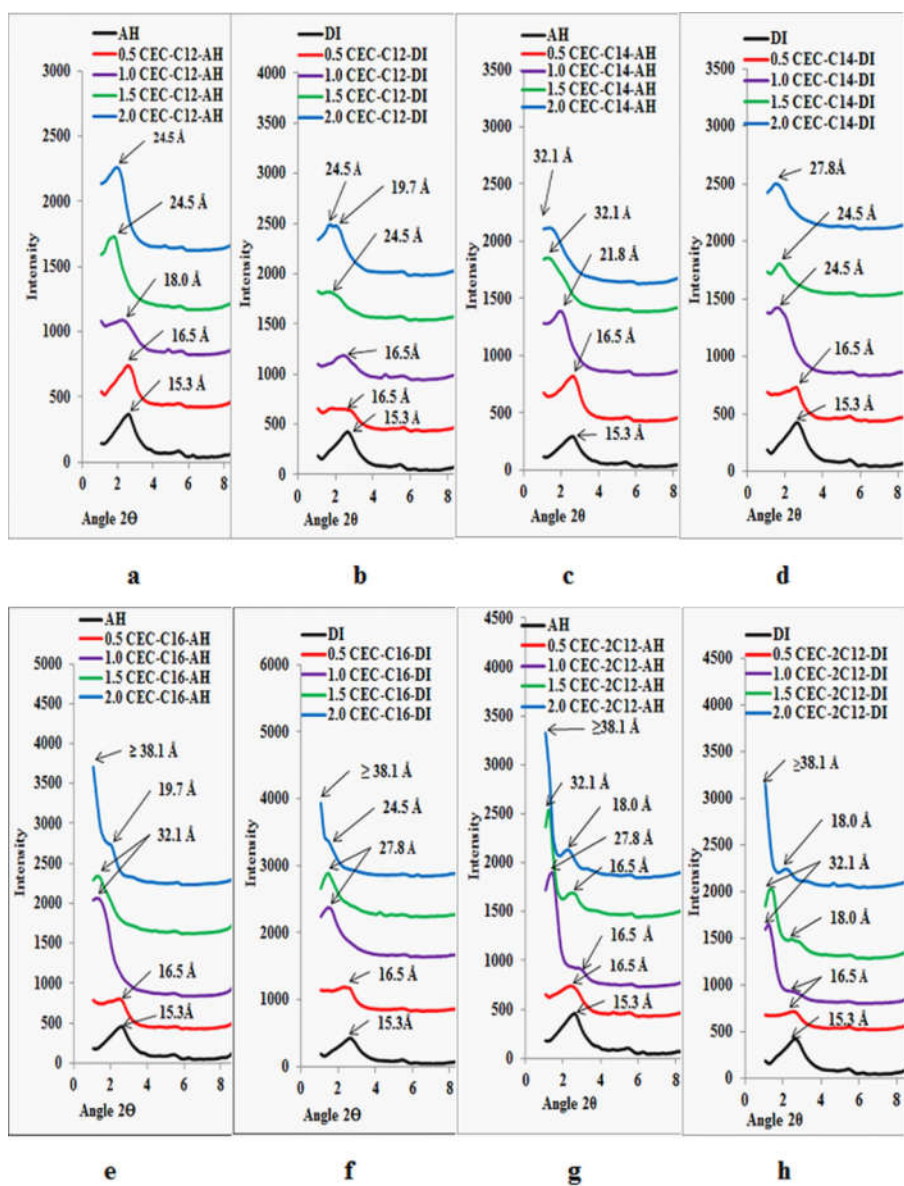


Figure 2. X-ray powder diffraction patterns of AH, DI clays and C₁₂-AH, C₁₂-DI, C₁₄-AH, C₁₄-DI, C₁₆-AH, C₁₆-DI, 2C₁₂-AH and 2C₁₂-DI organoclays.

The basal spacing information of the modified clays obtained from the XRPD experiments proves the intercalation of the organic surfactant into the clay layers. According to the XRPD patterns of the raw and modified clays, the basal spacing increases with increasing loading of surfactant. This increase shows that the surfactant molecules are intercalated between the layers of the mineral clays and that increased loading increases the basal spacing, d_{001} .

The XRPD patterns show also an increasing d_{001} value with increasing length of the long alkyl chain (C_{12} , C_{14} and C_{16}) and number of long alkyl chains (C_{12} and $2C_{12}$), Figure 2. Furthermore, the increase in the d_{001} values seems to take place in steps with preferred intervals, Figure 2.

The value of the basal spacing d_{001} increases with increasing length of long alkyl chains. The d_{001} value of 2.0 CEC- C_{12} -AH and 2.0 CEC- C_{12} -DI (25 Å) is smaller than the d_{001} values of 2.0 CEC- C_{14} -AH (32 Å) and 2.0 CEC- C_{14} -DI (28 Å), which are also smaller than the d_{001} values of 2.0 CEC- C_{16} -AH and 2.0 CEC- C_{16} -DI (≥ 38 Å). According to Park [24], at 0.5 CEC level loading, the d_{001} value increases marginally with the length of the surfactant: C_{12} (14.1 Å), C_{14} (14.3 Å) and C_{16} (14.4 Å) at a XRD step size of 0.0167° for 2θ . In principle the same observation is made in this study with the same d_{001} spacing for all applied surfactants at 0.5 CEC loading, Figure 2. This can be explained by the low loading level and the long alkyl chains that are oriented parallel to the sheets, Figures 3a and 3b. With increasing loading, length and number of long alkyl chains in the quaternary ammonium ions the d_{001} spacing increases, Figure 2. This means that the orientation of the intercalated quaternary ammonium ions changes from being parallel with the sheets to become more and more upright with increasing loading, Figure 3.

At 1.0 CEC loading the increase in d_{001} spacing for C_{12} and C_{14} is marginal compared to the 0.5 CEC loading, but significant for C_{16} and $2C_{12}$, Figure 2. At 1.5 CEC loading the d_{001} spacing for C_{12} and C_{14} increase and reach the same values as for the 1.0 CEC loading of C_{16} and $2C_{12}$. However, the latter keep the same d_{001} spacing at 1.5 CEC as at 1.0 CEC loading. At 2.0 CEC loading the C_{12} and C_{14} surfactants maintain the loading at 1.5 CEC, while for the C_{16} and $2C_{12}$ surfactants a further increase in the d_{001} spacing is observed, Figure 2. These preferred d_{001} spacing's are most likely connected to certain preferred orientations and stacking of the long alkyl chain(s) of the surfactants even though this study cannot give any detailed models for the orientation and stacking of the long alkyl chains of the surfactants applied. Figure 3 shows tentative orientations and stacking of the surfactants with increasing CEC loading. At full intercalation it can be expected that the structure of the surfactants resembles their structure in solid-state. He *et al.* found that there is an increase when the long alkyl chain number in the quaternary alkylammonium ions increases from one to two in the case of C_{12} and C_{16} surfactants [29], as also found in the present study for the C_{12} surfactant. Indeed, the d_{001} value of 2.0 CEC- C_{12} -AH and 2.0 CEC- C_{12} -DI (24.5 Å) is smaller than the d_{001} value of 2.0 CEC- $2C_{12}$ -AH and 2.0 CEC- $2C_{12}$ -DI (≥ 38 Å), Figure 2. This effect of the level of loading on the d_{001} basal spacing values is in agreement with the previous studies [23, 24, 29, 37, 40, 41]. Cubuk *et al.* and Caglar *et al.* [30, 31] reported that the basal spacing (d_{001}) values of modified clays, prepared with C_{12} and C_{16} surfactants added between 0.5 CEC and 3.0 CEC increased for each successive level loading (XRD step size 0.026° for 2θ). However, in the present work we found that the basal spacing remains the same in some cases for two successive level loadings. The step size in the present experiment, 0.2° in 2θ , causes a lower resolution than in previous experiments. Nevertheless, the resolution is sufficient for a basic understanding of the systematic changes in the expansion of the modified clays along the c -axis of the montmorillonite unit cell with increasing loading level, length and number of long alkyl chains, of the surfactant.

The XRPD diffractograms indicate that in the case of the C_{16} and $2C_{12}$ surfactants, from 1.0 CEC to 2.0 CEC level loading, the d_{001} basal spacing has two values, Figure 2. This result is in agreement with the fact that C_{16} and $2C_{12}$ surfactants have more than one molecular arrangement as described by Park [24] and Cubuk *et al.* [30] for Ca-montmorillonite interlayers. This may

depend on different availability for loading of different particles. The peaks related to d_{001} spacing are shifted to lower Bragg angles for all modified clay samples with increased surfactant loading. This variation in the intercalation spacing is systematic with length, number of long alkyl chains and level of surfactant loading applied for the Ca-montmorillonite clays, which is in agreement with the previous studies [23, 24, 30, 42]. The increase of the d_{001} basal spacing at the surfactant intercalation also sustains the decrease of the relative density.

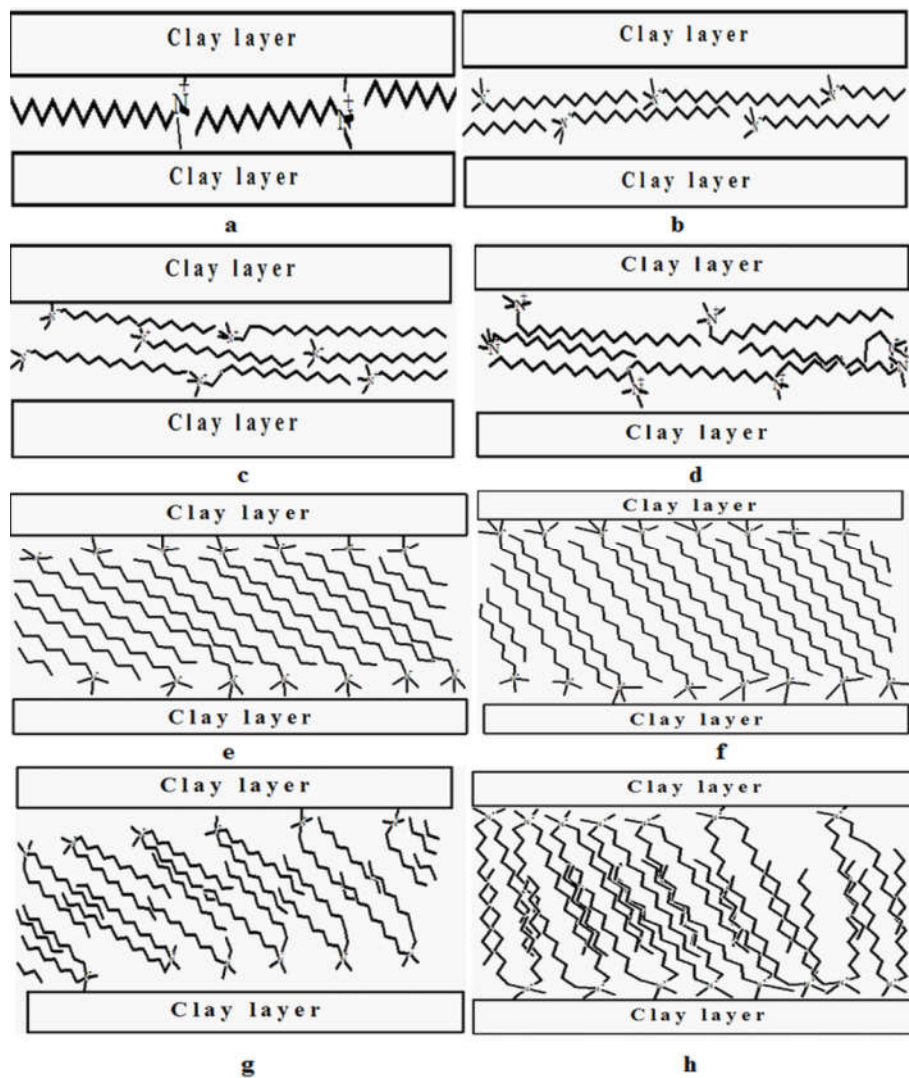


Figure 3. Surfactant molecules orientations into the interlayer space: arrangements (a) monolayer, (b) bilayers, and (c, d) pseudo-trilayers and (e, f and g, h) paraffin-type for one and two long alkyl chains, respectively [45].

The orientation of the surfactants depends most likely on the quantity and the arrangement and orientation of the surfactant in the interlayer. The structure of the adsorbed surfactant layer in the interlayers of smectite is most likely also affected by the charge distribution and chemical composition of the smectite.

Considering the interlayer expansions and the molecular dimensions of the surfactants, tentative arrangements of the surfactant molecules between the 2:1 sheets of the smectite have been distinguished. The expansion of the smectite evaluated by deducting the thickness of the smectite layer structure (tetrahedral-octahedral-tetrahedral) 2:1 unit (9.7 Å) [30, 31, 43] from basal spacing of 0.5 CEC organoclays samples is 6.8 Å. The molecular dimensions are approximately 3.8 Å in width and 18.0 Å in length for $C_{12}H_{25}(CH_3)_3N^+$, 4.0 Å in width and 20.8 Å in length for $C_{14}H_{29}(CH_3)_3N^+$ and 4.2 Å in width and 23.5 Å in length for $C_{16}H_{33}(CH_3)_3N^+$ when the long alkyl chain for each surfactant is parallel to the plan of smectite clay [30, 31, 43]. Regarding the interlayer expansion of 6.8 Å for all modified clays at 0.5 CEC loading of AH and DI smectites and the molecular sizes of C_{12} , C_{14} and C_{16} , it can be concluded that the surfactant molecules are located as a lateral monolayer arrangement. The interlayer distance and basal spacing of modified smectites at 0.5 CEC surfactant loading in this work are in agreement with previous investigations showing that C_{12} , C_{14} and C_{16} cations are inserted laterally in a monolayer arrangement [29, 30, 31, 43], Figure 3a.

The expansion of the interlayers when increasing the surfactant loading from 1.0 CEC to 2.0 CEC levels are 8.3 Å, 10.0 Å, 12.1 Å, 14.8 Å, 18.1 Å, 22.4 Å and more than 28.3 Å depending on the raw clay and/or the surfactant, Figures 3. Considering the molecular size of the surfactant, the expansion values, 8.3 Å and 10.0 Å, correspond to a bilayer arrangement of surfactant molecules with an angle which depends on the interaction between the surfactant molecules and the clay layers, the chemical composition of the clay and the structure of the surfactant. The d_{001} basal spacing values corresponding to this arrangement are 18.0 Å and 19.7 Å. For 12.1 Å and 14.8 Å as expansion values, the molecular orientation of the carbon alkyl chain is contained in an oblique plan and the arrangement of the surfactant molecule reaches a pseudo-trilayer arrangement. In case the expansions are 18.1 Å, 22.4 Å and more than 28.4 Å, the alkyl chain arrangement is paraffin-type and approximate to the alternate antiparallel packing between the clay layer as found in the crystal structures in solid bromide salts of the surfactants C_{12} , C_{14} and C_{16} [44]. The d_{001} basal spacing values corresponding to the paraffin-type arrangement are 27.8 Å, 32.1 Å and more than 38 Å. Therefore, the angle between the clay layers and the plane of the surfactant molecules approaches 90°. These arrangements of the long alkyl chains in the organoclays are similar to those proposed by Lagaly *et al.* [45].

A general decrease in intensity of the diffraction peaks with increasing loading of surfactants, Figure 3, strongly indicates decreasing crystallinity of organoclays with increase length of the c-axis due to the surfactant loading.

Fourier transform infrared (FTIR) spectroscopy of organoclays

The FTIR spectra of raw clays and modified clays prepared at 0.5 CEC, 1.0 CEC, 1.5 CEC and 2.0 CEC level loadings of C_{12} , C_{14} , C_{16} and $2C_{12}$ surfactants, and pure surfactants are shown in Figures 4-7.

In all these FTIR spectra, the H-O-H bending bands of water molecules adsorbed on raw and modified clays appear at ca. 1600 cm^{-1} . The peak at 3622 cm^{-1} for the raw clays is assigned to OH stretching vibrations of the structural hydroxyl group, and the band around 3420 cm^{-1} is assigned to water molecules adsorbed between the layers of the clay. Madejová *et al.* assigned the peaks at 3694 cm^{-1} and 3620 cm^{-1} to AlMg-OH and Al₂-OH vibrations, respectively. Those bands are normally assigned to stretching vibrations of hydroxyl groups coordinated to octahedral cations in montmorillonite or kaolinite [46-48]. Moreover the absorption peaks seen

at 3620 cm^{-1} on the raw clays and organoclays are typical for smectites with high amount of Al in the octahedra [46].

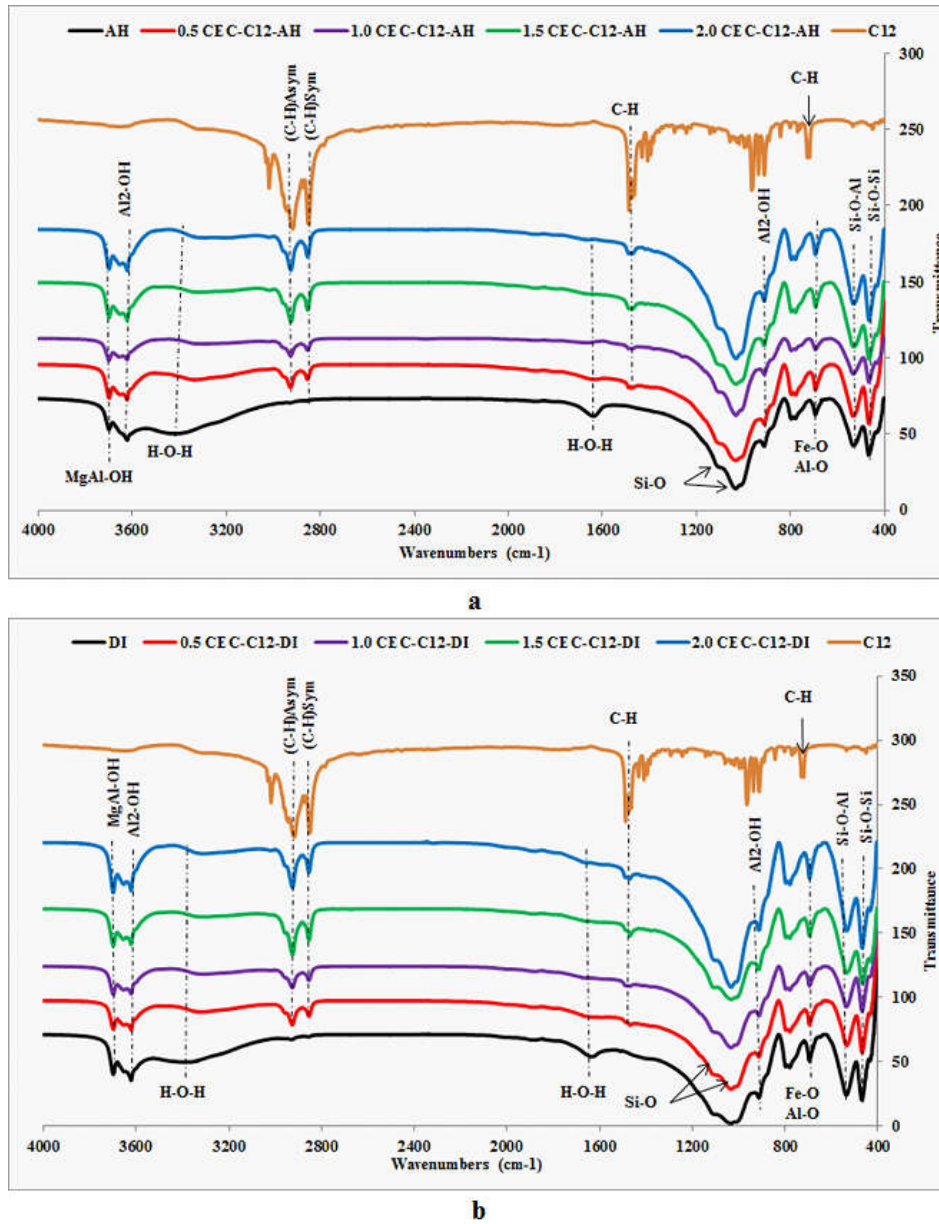


Figure 4. FTIR spectra of AH and DI raw clays, and C_{12} -AH and C_{12} -DI organoclays.

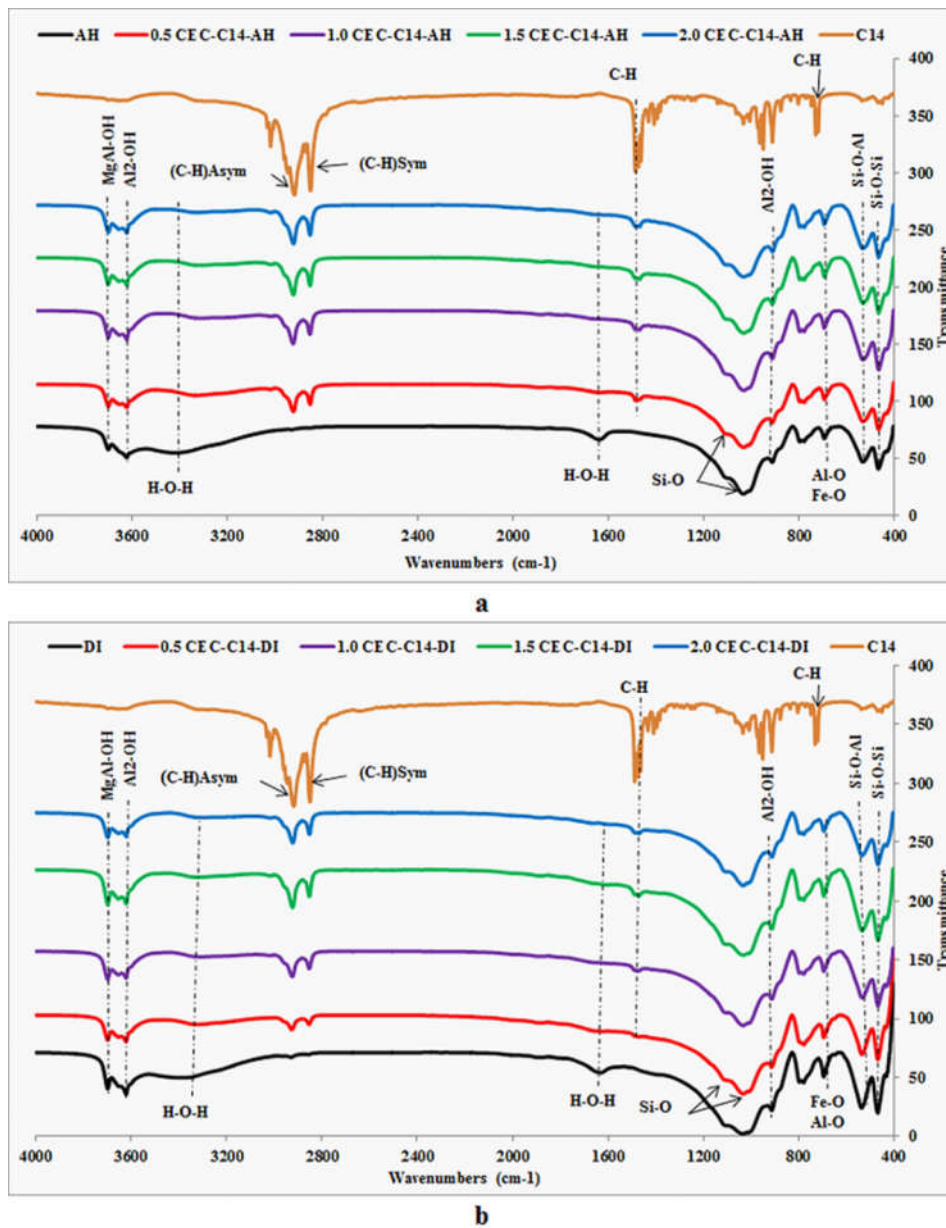


Figure 5. FTIR spectra of AH and DI raw clays, and C₁₄-AH and C₁₄-DI organoclays.

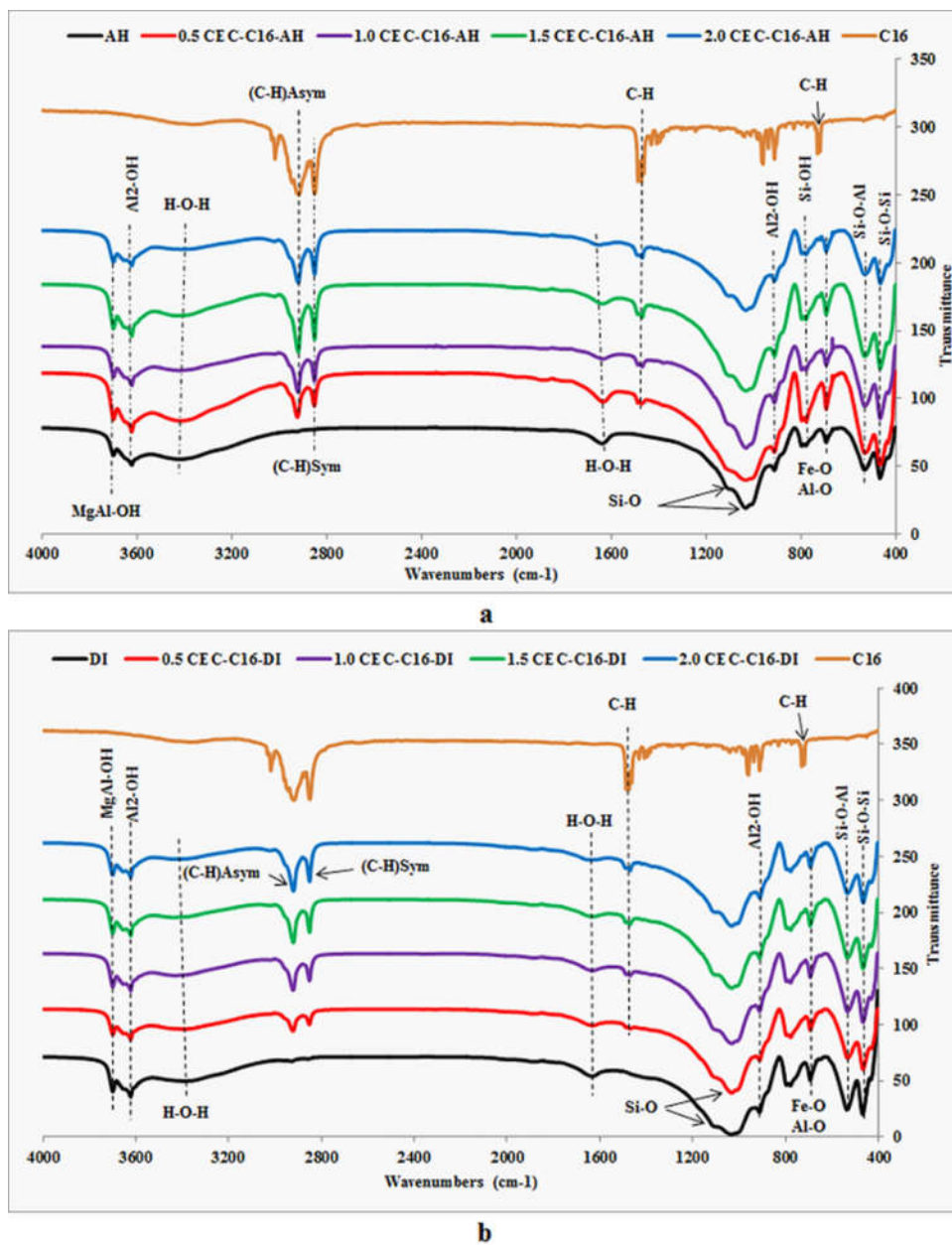


Figure 6. FTIR spectra of AH and DI raw clays, and C₁₆-AH and C₁₆-DI organoclays.

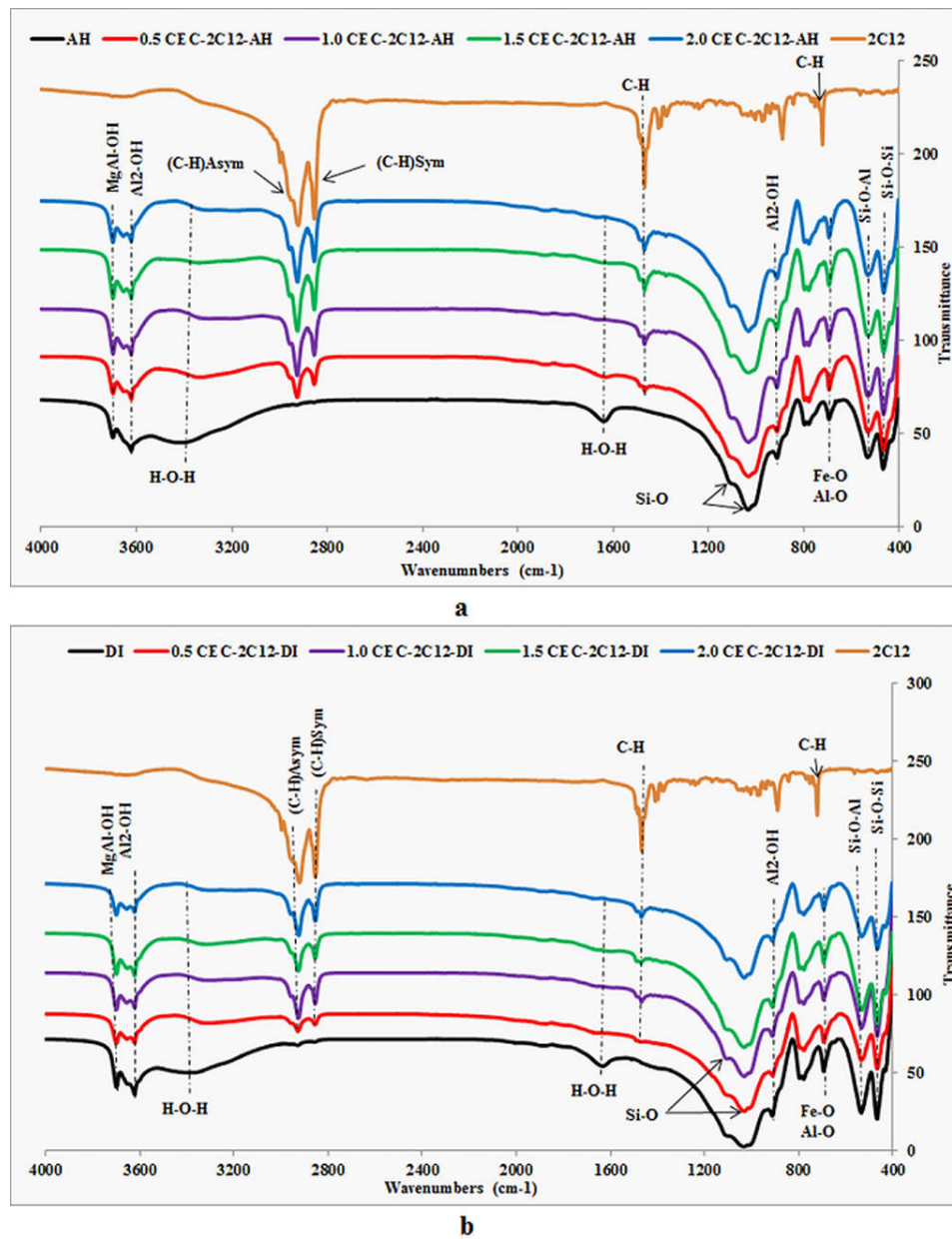


Figure 7. FTIR spectra of AH and DI raw clays, and $2C_{12}$ -AH and $2C_{12}$ -DI organoclays.

All FTIR spectra contain bands attributed to Si-O stretching at 1100 cm^{-1} and Si-O in-plane bending at 470 cm^{-1} [47]. The absorption peaks at 913 cm^{-1} and 536 cm^{-1} are assigned to Al_2OH bending vibration and Al-O-Si deformations of raw clays, respectively [47, 48]. The peaks observed around 694 cm^{-1} are assigned to Fe-O or Al-O out of plane bending vibrations [47].

The CH₂ asymmetric stretching bands are slightly shifted to lower wavenumbers upon intercalation of the C₁₂, C₁₄, C₁₆ and 2C₁₂ surfactants from 2927 cm⁻¹ to 2920 cm⁻¹ depending on AH or DI raw clays, and surfactant loading, 0.5 CEC to 2.0 CEC level. Furthermore, the wavenumber of the symmetric CH₂ stretching mode is shifted from 2855 cm⁻¹ to 2850 cm⁻¹ seen in the modified clays, Figures 4-7. When the surfactant loading level increases, the wavenumbers of asymmetric and symmetric CH₂ stretching bands of surfactants in the modified clays approach the wavenumbers of the pure surfactant. This further indicates that the alkyl chains are packed in a similar way as in the pure surfactant when the intercalation approaches saturation. According to Kamitori *et al.* [44], Silva *et al.* [49] and Campanelli and Scaramuzza [50], the molecular arrangements of C₁₂, C₁₄ and C₁₆ alkyl chains are as parallel as possible within a layer and antiparallel in alternate layers in the pure surfactant. The infrared spectroscopy data support that arrangement of alkyl chains in the organoclays changes gradually from being parallel with clay sheets at low surfactant loading to standing in a similar or identical way as the pure surfactants at high loading as proposed in previous studies [12, 13, 24].

The wavenumbers of asymmetric and symmetric stretching modes of CH₂ of the pure surfactant molecules and the modified clays are summarized in the Table 3. The frequency and the intensity of the asymmetric and symmetric stretching bands of CH₂ change with the length and the number of long alkyl chains as well as with level of loading. However, the CH₂ scissoring and rocking vibrations seen at 1473 cm⁻¹ and 731 cm⁻¹, respectively [30], seem to be independent of the length of alkyl chain and level of loading.

Moreover, the additional weak band, which is seen between 3670 cm⁻¹ and 3700 cm⁻¹ on the raw clay spectra, increases with surfactants molecules intercalation. This is related to the insertion of the surfactants molecules into the Ca-montmorillonite interlayers. It is observed that the band intensities at 3420 cm⁻¹ and 1635 cm⁻¹ decreased progressively with the surfactant loading level due to the intercalation of the quaternary alkylammonium cations and this causes that the hydrophilic properties of raw clays change to hydrophobic ones [30, 31, 46]. The FTIR spectra support the model of intercalation of surfactants between the sheets in clays as described above, Figure 3.

Table 3. Wavenumbers variation of CH₂ asymmetric and symmetric stretching mode in AH and DI modified clays.

AH clay								
Surfactants and wavenumbers (cm ⁻¹) of asymmetric and symmetric CH ₂ stretching								
Loading level	C ₁₂		C ₁₄		C ₁₆		2C ₁₂	
	CH ₂ asym	CH ₂ sym	CH ₂ asym	CH ₂ sym	CH ₂ asym	CH ₂ sym	CH ₂ asym	CH ₂ sym
0.5 CEC	2926	2855	2923	2853	2923	2852	2925	2854
1.0 CEC	2926	2854	2923	2852	2921	2851	2924	2854
1.5 CEC	2925	2854	2922	2852	2920	2851	2924	2853
2.0 CEC	2925	2854	2921	2851	2920	2850	2921	2853
DI clay								
Surfactants and wavenumbers (cm ⁻¹) of asymmetric and symmetric CH ₂ stretching								
Loading level	C ₁₂		C ₁₄		C ₁₆		2C ₁₂	
	CH ₂ asym	CH ₂ sym	CH ₂ asym	CH ₂ sym	CH ₂ asym	CH ₂ sym	CH ₂ asym	CH ₂ sym
0.5 CEC	2926	2855	2926	2854	2923	2852	2927	2855
1.0 CEC	2926	2855	2924	2853	2922	2851	2925	2855
1.5 CEC	2924	2854	2923	2852	2921	2851	2925	2854
2.0 CEC	2924	2854	2922	2852	2920	2851	2924	2854
Pure surfactants	2918	2850	2918	2849	2919	2849	2921	2853

CONCLUSION

Two natural clays from Burkina Faso containing Ca-montmorillonite as main clay type were successfully intercalated by cationic alkyltrimethylammoniums ($n\text{-C}_{12}\text{H}_{25}(\text{CH}_3)_3\text{N}^+$, $n\text{-C}_{14}\text{H}_{29}(\text{CH}_3)_3\text{N}^+$ and $n\text{-C}_{16}\text{H}_{33}(\text{CH}_3)_3\text{N}^+$) and di-alkyldimethylammonium ($(n\text{-C}_{12}\text{H}_{25})_2(\text{CH}_3)_2\text{N}^+$) using solid-state reaction methodology. Relative density measurements show that the relative density of the organoclays decreases with increasing surfactant loading. Thus, the natural clays swell substantially at treatment with surfactants. The increase of the d_{001} spacing shown by XRPD proved the swelling of layers by intercalation of the surfactants. The value of basal spacing reaches 25 Å for modified clays treated with C_{12} , 32 Å for those treated with C_{14} , and more than 38 Å for those treated with C_{16} and 2C_{12} at 2.0 CEC loading level of the surfactant. The IR spectra showed the CH_2 asymmetric and symmetric stretching bands at around 2920 cm^{-1} and 2850 cm^{-1} , respectively, proving successful intercalation. FTIR spectra showed that the wavenumbers of asymmetric and symmetric stretching vibrations of the surfactants in organoclays decrease and approach the values observed for the pure surfactants with increasing surfactant level. The intercalation of the quaternary alkylammonium cations into the clays interlayers causes that the hydrophilic properties of raw clays change to hydrophobic and organophilic. The solid-state intercalation of alkyltrimethylammonium and di-alkyldimethylammonium cations is an effective and simple method to prepare useful modified clays for environmental applications.

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