

DIELECTRIC RELAXATION OF WATER IN CLAY MINERALS

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Abstract—The study of confined water dynamics in clay minerals is a very important topic in aluminosilicate-surface chemistry. Aluminosilicates are among the most technologically versatile materials in industry today. Dielectric spectroscopy is a very useful method for investigating the structure and dynamics of water adsorbed on solid matrix surfaces and water in the vicinity of ions in solutions. Use of this method for the study of clay minerals has been underutilized to date, however. The main goal of the present research was to understand the relaxation mechanisms of water molecules interacting with different hydration centers in clay minerals, with a view to eventually control this interaction. Two types of natural layered aluminosilicates (clay minerals) — montmorillonite with exchangeable K^+ , Co^{2+} , and Ni^{2+} cations and kaolinite with exchangeable K^+ and Ba^{2+} cations — were examined by means of dielectric spectroscopy over wide ranges of temperature (from -121°C to $+300^\circ\text{C}$) and frequency (1 Hz–1 MHz). An analysis of the experimental data is provided in terms of four distributed relaxation processes. The low-temperature relaxation was observed only in montmorillonites and could be subdivided into two processes, each related to a specific hydration center. The cooperative behavior of water at the interface was observed in the intermediate temperature region, together with a proton percolation. The dielectric properties of ice-like and confined water structures in the layered clay minerals were compared with the dielectric response observed in porous glasses. The spatial fractal dimensions of the porous aluminosilicates were calculated by two separate methods — from an analysis of the fractality found in photomicrographs and from the dielectric response.

Key Words—Adsorbed Water, Dielectric Spectroscopy, Exchangeable Cations, Fractal Dimension, Kaolinite, Montmorillonite.

INTRODUCTION

The large specific surface area, chemical and mechanical stability, variety of structural and surface properties, large values of cation exchange capacities, *etc.*, make the clay minerals an excellent group of adsorbents (Greg and Sing, 1967; Gupta and Bhattacharyya, 2012; Kiselev, 1986; Tarasevich, 1988; Tarasevich and Ovcharenko, 1975; Volzone *et al.*, 1999). Consequently, the hydration properties of clay materials are a fundamental problem in any field of their application.

One of the most useful tools for investigating the structure and dynamics of adsorbed water on solid matrix surfaces is Dielectric Spectroscopy (DS) (Gutina *et al.*, 2003; Saltas *et al.*, 2007; Spanoudaki *et al.*, 2005; Wander and Clark, 2008). Broadband Dielectric Spectroscopy (or impedance spectroscopy) occupies a special place among the numerous modern methods used for the physical and chemical analysis of materials. By measuring the electric polarization of a dielectric material under the influence of an external time-dependent electric field at different temperatures,

investigation of relaxation processes with an extremely wide range of characteristic times (10^5 – 10^{-13} s) and monitoring different scales of molecular motions are possible (Kremer and Schöenhals, 2002; Feldman *et al.*, 2006). In particular, DS is sensitive to cooperative intermolecular interactions (Feldman *et al.*, 2006) and, thus, provides a link between the properties of the individual constituents of a complex material *via* molecular spectroscopy and the characterization of its bulk properties. Many key aspects of a material's properties can be derived from the complex dielectric functions, *e.g.* the strength and the shape parameters of molecular relaxations, the characterization of structural phase transitions, the activation energy of the relaxation process, charge mobility, *etc.* Moreover, this method is very sensitive to water dynamics in ionic aqueous solutions and can provide important information on the state of water in the vicinity of ions (Levy *et al.*, 2012).

The clay minerals represent a class of materials where the two types of water organization mentioned above can be observed simultaneously. Several attempts have been made to perform dielectric measurements of hydrated kaolinites and montmorillonites over narrow frequency and temperature ranges by changing the temperature at a fixed frequency or *vice versa* (Calvet, 1975; Hall and Rose, 1978; Hoekstra and Doyle, 1971; Lockhart, 1980a, 1980b; Mamy, 1968; Raythatha and Sen, 1986; Sposito

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and Prost, 1982). Recently, the increased interest in hydrated clay minerals has stimulated research in this field, primarily to correlate the electrical properties with their structural parameters and water content (Ishida *et al.*, 2000; Kaviratna *et al.*, 1996; Logsdon and Laird, 2002, 2004; Rotenberg *et al.*, 2005). The main aim of the present study was to examine the effect of various types of cations on the dielectric response of hydrated montmorillonite and kaolinite clay minerals over broad ranges of temperature (from -121°C to $+300^{\circ}\text{C}$) and frequency (1 Hz–1 MHz), in order to obtain detailed information on the structure and the relaxation properties of water adsorbed by different hydration centers in the minerals. Preliminary results of the work were published by Vasilyeva *et al.* (2012a, 2012b).

EXPERIMENTAL

Materials

Two types of clay minerals were considered here: Oglanlinsky montmorillonite (from Turkmenistan) and Glukhovetsky kaolinite (from Ukraine). The clay minerals were treated in the Laboratory of Disperse Systems of the Institute of Colloid and Water Chemistry (Kiev, Ukraine) according to a procedure described by Tarasevich and Ovcharenko (1975, 1980). At first, impurities of other minerals, free oxides and hydroxides of Fe and Al, as well as other impurities and inclusions were removed from the raw clay. To obtain the purest samples possible, 3–5% aqueous suspensions of the clay mineral in question were mixed thoroughly until no agglomerates were visible and then kept at quiescence for ~30 min. A method of separating the solid clay particles from the liquid was chosen to elutriate the finest fraction: the precipitate was allowed to settle and the liquid settled or carefully decanted or filtered off through a siphon (Tarasevich and Ovcharenko, 1975, 1980). For kaolinite no other treatment was performed. Additional purification and centrifugation were required for montmorillonites. The purity (homogeneity) of the samples was checked by X-ray diffraction (XRD) and infrared (IR) spectroscopy. Then cleaned natural samples were treated 7–8 times with a 1 M NaCl solution at a solid:liquid (S:L) ratio of 1:20; each time the system was shaken, kept at quiescence for ~1 h, and then decanted. Afterward, the Na-saturated minerals were washed free of excess salts using distilled water.

Table 1. Relative water content, h , in the samples studied.

Sample	h (%)
K-montmorillonite	13.7
Ni-montmorillonite	14.7
Co-montmorillonite	15.3
K-kaolinite	2.3
K,Ba-kaolinite	5.4

(resistivity not measured). Next, the clay minerals were treated 7–8 times in the same way with 1 N solutions of the required metal chlorides (KCl, BaCl₂, CoCl₂, NiCl₂) in the same S:L ratio. Finally, the samples were washed with distilled water until a negative reaction for chloride ion (with AgNO₃), dried for ~1 h at ~100°C in air, then dried for ~1 h at ~50°C in vacuum, and then stored at ambient air humidity until measured.

The relative water content, h , was determined by weighing the samples prior to and immediately after the dielectric measurements according to equation 1. The water content in a sample was calculated as a percent weight of the sample relative to the initial weight (Table 1):

$$h (\%) = 100 \times (m_{\text{initial}} - m_{\text{final}}) / m_{\text{initial}} \quad (1)$$

Methods

The elemental composition of the samples was determined by X-ray fluorescence (XRF) analysis, using an energy dispersive X-ray spectrometer EDX 800HS2 (Shimadzu, Germany) (Table 2).

The morphology of the samples was examined using a field emission scanning electron microscope (FESEM) (Merlin, Carl Zeiss, Germany). Images of the morphology of some of the samples were collected using FESEM (Figure 1).

Powder XRD measurements were performed at room temperature using a D8 Advance diffractometer (Bruker AXS, Karlsruhe, Germany) with a goniometer radius of 217.5 mm, Göbel Mirror parallel-beam optics, 2° Sollers slits, and 0.2 mm receiving slit. Profile fitting, refinement of unit-cell parameters, and crystallite-size calculations were performed using the EVA software (Bruker, AXS). The crystallite sizes of monoclinic Ni- and Co-montmorillonite obtained were 251.5 Å and 199.1 Å, respectively; the crystallite size of triclinic kaolinite was

Table 2. Elemental compositions of the four studied samples.

Sample	Mg (%)	Al (%)	Si (%)	K (%)	Ti (%)	V (%)	Fe (%)	Ni (%)	Cu (%)	Zn (%)	Ba (%)
K-montmorillonite	1.8	11.3	74.7	8.7	0.4		2.9		0.2	0.1	
Ni-montmorillonite	2.4	13.6	71.5		0.5		3.1	8.5	0.4		
K-kaolinite		35.5	57.5	2.9	1.9	0.1	1.9			0.1	
K,Ba-kaolinite		37.0	57.3	1.0	2.3		1.2		0.1	0.1	1.0

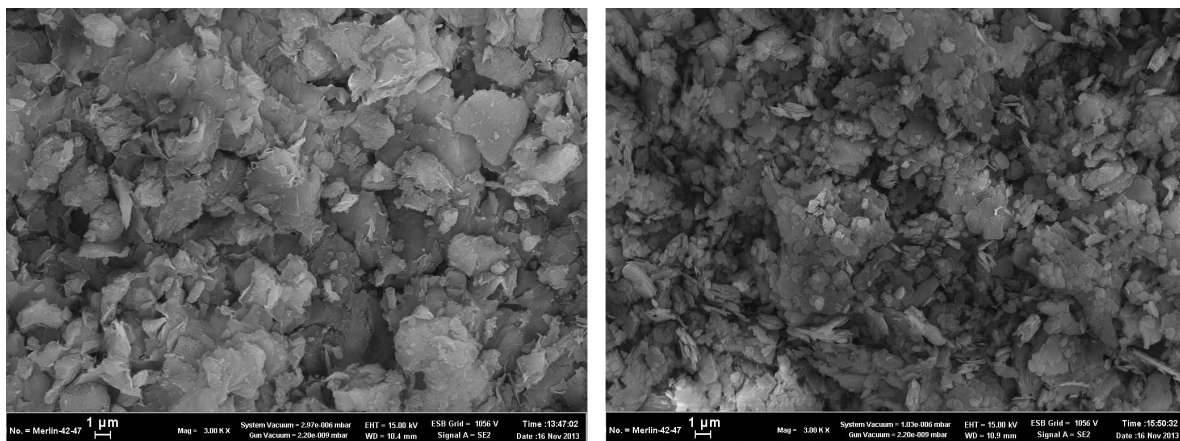


Figure 1. Electron micrographs of Ni-montmorillonite (left) and K-kaolinite (right).

382.7 Å. The kaolinite sample also contained some monoclinic sericite with a crystallite size of 409.1 Å.

Dielectric measurements in the frequency range of 1 Hz–1 MHz were performed using a BDS 80 Dielectric Spectrometer, based on an Alpha Impedance Analyzer (Novocontrol, Germany) with automatic temperature control by a Quatro Cryosystem, with a precision of 0.5°C. The Alpha High Resolution Dielectric Analyzer can measure impedances from 10^{-2} to 10^{14} Ohms with a resolution in loss factor $\tan(\delta) < 10^{-5}$ (Schaumburg, 1994; Kremer and Schoenhals, 2002). A voltage U_0 with a fixed frequency is applied to the cell which contains the material under investigation. The resulting current, I_0 , has the same frequency but a phase shift between current and voltage described by the phase angle, ϕ . The ratio of U_0 and I_0 and the phase angle ϕ is determined from the electromagnetic and the geometric properties of the sample. The formulae for the voltage $U(t)$ and current $I(t)$ in their phasor notations can be represented conveniently by the following:

$$\begin{aligned} U(t) &= U_0 \sin(\omega t) = I_m(U^* \exp(i\omega t)) \\ I(t) &= I_0 \sin(\omega t + \phi) = I_m(I^* \exp(i\omega t)) \end{aligned} \quad (2)$$

where $U^* = U' + iU'' = U_0$; $I^* = I' + iI''$; $I_0 = \sqrt{(I'^2 + I''^2)}$; $\tan(\phi) = I''/I'$ where i is the imaginary unit and ω is the cyclic frequency. For a material with a linear electromagnetic response, the measured complex impedance of the sample capacitor is

$$Z^*(\omega) = Z' + iZ'' = U^*/I^* \quad (3)$$

This is related to the complex dielectric permittivity of the sample material by:

$$\varepsilon^*(\omega) = \varepsilon'(\omega) - i\varepsilon''(\omega) = \frac{1}{i\omega C_0} \frac{1}{Z^*(\omega)} \quad (4)$$

where $\varepsilon^*(\omega)$ is the complex dielectric permittivity, $\varepsilon'(\omega)$ and $\varepsilon''(\omega)$ are real and imaginary parts, respectively, of the $\varepsilon^*(\omega)$, and C_0 is the empty cell capacity.

A parallel plate capacitor with electrodes of 16 mm diameter and a spacing of ~0.5 mm between them was used. The study was carried out using the following protocol: Each of the samples was placed in the sample cell at room temperature and the measurements were then performed by quenching the samples down to -115°C . Then the samples were measured upon heating to -58°C with 3°C intervals and then at intervals of 6°C up to $+300^\circ\text{C}$. Afterward, annealing was performed inside the cryostat at 300°C for 3 h. For montmorillonite with Co^{2+} as the exchangeable cation, the lowest temperature was -121°C . After annealing, the samples were cooled to 25°C and stored at this temperature in a nitrogen atmosphere until weighing. The montmorillonite samples with K^+ and Ni^{2+} as the exchangeable cations were also measured during the decrease in temperature down to 25°C , in order to check the reversibility of the signal. All samples were weighed prior to and immediately after the experiments. The accuracy of the complex dielectric permittivity was better than 3%.

RESULTS

The complex non-Debye dielectric behavior of the clay minerals (Figures 2 and 3) can be described in terms of several distributed relaxation processes, separated by different frequency and temperature intervals, and marked as I–IV, according to their appearance during the heating experiment.

At low temperatures from -121°C to -75°C , the spectra of montmorillonites consisted of two relaxation processes marked I.1 and I.2, respectively (Figure 4). Both of these processes are characterized by a strong temperature dependence. In the kaolinite samples measured with the same protocol, the low-temperature process I was absent.

In the mid-temperature range from -90°C to $+150^\circ\text{C}$, a relaxation process with a specific saddle-like shape and distinctive kink point was observed for mont-

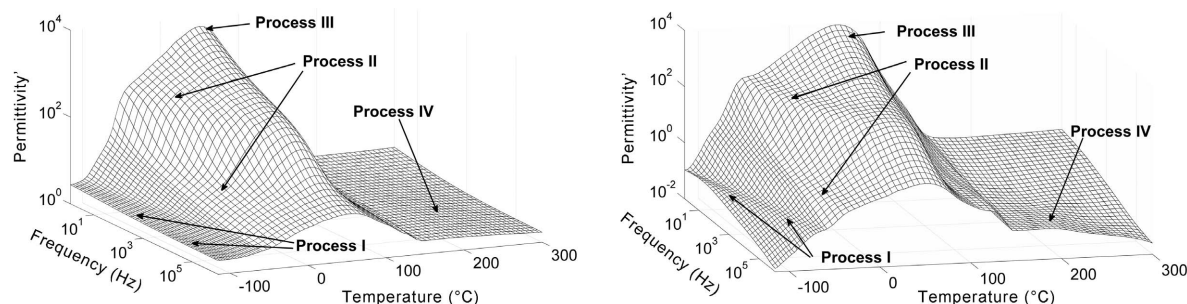


Figure 2. Typical 3D plots of the frequency and temperature dependences of the dielectric permittivity (left) and dielectric losses (right) for Ni-montmorillonite.

morillonite clay minerals at different frequency-temperature intervals simultaneously, while only one relaxation process, II, was observed in the case of kaolinite samples (Figure 5).

Relaxation process III was detected at temperatures from -50°C to $+70^{\circ}\text{C}$ for montmorillonites and from -70°C to $+40^{\circ}\text{C}$ for kaolinites. When viewed in an *ac*-conductivity presentation ($\sigma_{ac}(\omega) = (i\omega\epsilon''(\omega))$), typical *s*-like behavior of the low-frequency conductivity demonstrates the percolative nature of this process (Figure 6).

The amplitude of process III essentially decreased when the frequency increased and the maximum of the dielectric permittivity and losses vs. temperature had no temperature dependence (Figure 7) for all the clay-mineral samples.

The percolation temperature point, T_p , was determined by the maximum of the temperature dependence of the dielectric permittivity at the lowest measured frequency. The percolation temperature for montmorillonites was greater than for kaolinites (Table 3).

When the temperature rose above 100°C , the samples became more electrically conductive. The relaxation processes were overlapped in the dielectric losses by a *dc*-conductivity. Some were even screened by the conductivity (Figure 8). A detailed investigation of the high-temperature region is the subject of a separate study.

After annealing the samples, the 3D plots of the dielectric losses vs. frequency and temperature changed dramatically (Figure 9). The relaxation processes previously observed in the mid-temperature window disappeared.

Traces of the high-temperature spectrum were observed during the cooling measurement (Figures 8 and 9).

DISCUSSION

As shown above, all the samples lost some mass during the experiment. The clay-mineral samples demonstrated the absence of the dielectric relaxation processes at low- and mid-temperature levels after heating for extended periods of time at 300°C (Figures 2, right and 9). As mentioned above, clay materials can adsorb water easily at their surfaces and around exchangeable cations, suggesting that water is the origin of the dielectric relaxation in these materials in the low- and mid-temperature intervals. Note that a similar conclusion was reported for porous glass systems (Gutina *et al.*, 2003).

From a structural point of view, the surface of the broken edges of the mineral layer has primary broken Al–O and Si–O bonds, resulting in polar sites. Water molecules form hydrogen bonds with surface oxygen atoms and exchangeable cations, to create new water structures very different from those of the bulk (Schoonheydt and Johnston, 2006; Sposito and Prost, 1982). In confined spaces or near interfaces, water forms distinctively layered structures with a restriction of water molecular mobility (Behnken and Faulkner, 2011).

The minerals studied differ in their surface properties and exchangeable cations due to the differences in mineralogy and crystal structure, which lead to the variation in their affinity toward water.

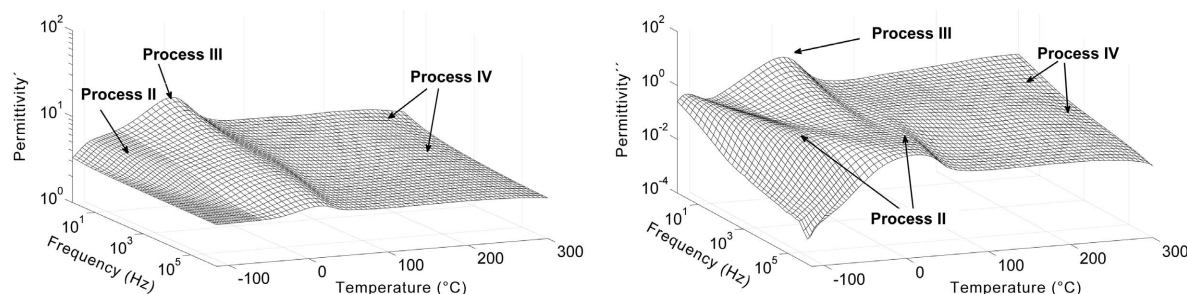


Figure 3. Typical 3D plots of the frequency and temperature dependences of the dielectric permittivity (left) and dielectric losses (right) for K-kaolinite.

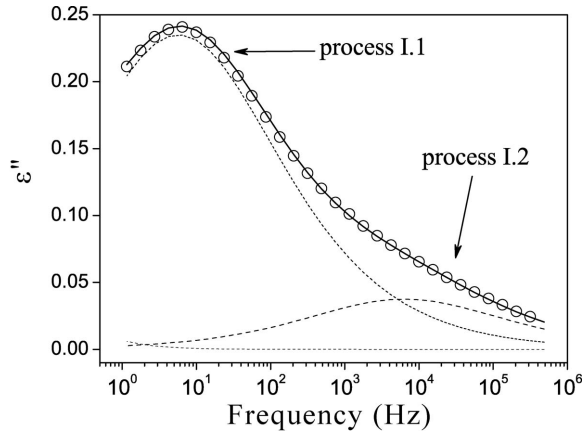


Figure 4. Measured dielectric losses vs. frequency at -121°C for Co-montmorillonite. The dashed lines mark two Cole-Cole and Jonscher terms used for fitting the experimental data (open circles). The solid line is the superposition of the terms above.

The low-temperature dielectric relaxation

In the low-temperature interval (from -121°C to -75°C) the dielectric relaxation process (I) was observed for montmorillonites only. Therefore, this process must be associated with the relaxations of interlayer water molecules which are absent from kaolinites. Moreover, the dielectric spectrum of montmorillonites consists of two low-temperature relaxation processes (Figure 4).

Whenever water interacts with another dipolar or charge entity, a broadening of its Debye dielectric relaxation spectra occurs (Puzenko *et al.*, 2010; Levy *et al.*, 2012). In most cases this broadening can be described by the phenomenological Cole-Cole (CC) spectral function (Cole and Cole, 1941):

$$\varepsilon^*(\omega) = \varepsilon_{\infty} + \frac{\Delta\varepsilon}{1 + (i\omega\tau)^{\alpha}} \quad (5)$$

where $\Delta\varepsilon = \varepsilon_s - \varepsilon_{\infty}$ is the dielectric strength of the relaxation process; ε_s and ε_{∞} are the low- and high-

Table 3. Values of the percolation temperature for the clay minerals studied.

Sample	T_p ($^{\circ}\text{C}$)
K-montmorillonite	40
Co-montmorillonite	40
Ni-montmorillonite	35
K-kaolinite	8
K,Ba-kaolinite	8

frequency limits of the complex dielectric permittivity, respectively; τ is a characteristic relaxation time; and the exponent α is referred to as a measure of symmetrical broadening in the dielectric losses relaxation peak ($0 < \alpha \leq 1$). A fruitful approach has been the application of fractional calculus to uncover a physical mechanism underlying the CC behavior in complex systems (Puzenko *et al.*, 2010; Levy *et al.*, 2012; Puzenko *et al.*, 2012). Taking into account that both low-temperature processes (I.1 and I.2) of montmorillonites are associated with the interaction of the water dipoles with the clay's mineral matrix, the quantitative analysis of the isothermal dielectric spectra uses a superposition of two CC functions, and an empirical Jonscher (J) term is convenient (Jonscher, 1996) (Figure 4):

$$\varepsilon^*(\omega) = \varepsilon_{\infty} + \sum_{j=1}^2 \frac{\Delta\varepsilon_j}{1 + (i\omega\tau_j)^{\alpha_j}} + B \cdot (i\omega)^{n-1} \quad (6)$$

where j denotes the number of the corresponding relaxation process and n is a Jonscher parameter for the high-frequency part of the respective relaxation process.

The temperature dependences of the relaxation times for the observed low-temperature processes demonstrated Arrhenius behavior (Figure 10). The activation energies of the low-temperature relaxation processes were then calculated (Table 4).

In montmorillonites the water molecules responsible for the low-temperature relaxations are located near the

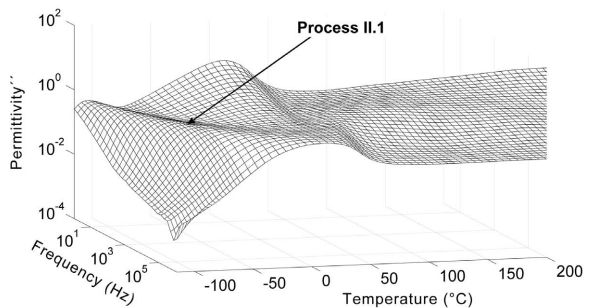
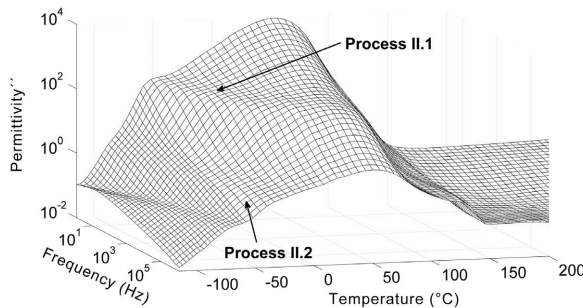


Figure 5. Typical 3D plots of the frequency and temperature behavior of the dielectric losses at a mid-temperature interval for Ni-montmorillonite (left) and K-kaolinite (right).

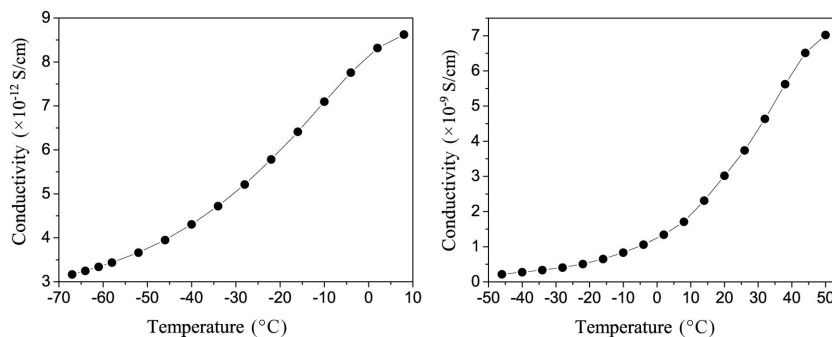


Figure 6. Temperature behavior of the low-frequency conductivity for Ni-montmorillonite (left) and K-kaolinite (right) in the mid-temperature region.

inner hydrophilic centers of interlayer channels (siloxane bonds and trapped cations) and around the interlayer cations (Salles *et al.*, 2008; Tarasevich and Ovcharenko, 1975). This is reflected by the formation of two types of structure in the low-temperature region. The slower relaxation process I.1 almost coincides with the low-temperature relaxation behavior of water adsorbed in porous silica glasses (Gutina *et al.*, 2003) (Figure 10). The activation-energy values of this process for montmorillonite samples are in the same range as that reported earlier for the porous glass materials (Gutina *et al.*, 2003; Ryabov *et al.*, 2001). The I.1 relaxation process was assumed, therefore, to correspond to ice-like water structures formed on the interior surface of interlayer channels. The second, faster, relaxation process I.2 reflects the water structures near the hydrated cations located in the interlayer channels. Thus, the nature of the hydration center plays an important role in

the water dynamics of the adsorbed water in inorganic materials. Note that the type I.2 process in the glass materials (Gutina *et al.*, 2003) was not observed, but only one type of hydration center was observed in those materials. In both processes (I.1 and I.2) the activation energy depended on the type of interlayer cations and on the water content (Table 4). The detailed dielectric investigation of the formation of hydrated clusters and their structures as a function of the exchangeable cations and water content in the montmorillonites will be reported in a future study.

The mid-temperature relaxation process

Saddle-like relaxation processes were observed in the mid-temperature range from -90°C to $+150^{\circ}\text{C}$ for all the samples studied. Quantitative analysis of the mid-temperature relaxation process was performed based on the model described by Gutina *et al.* (2003) in which the

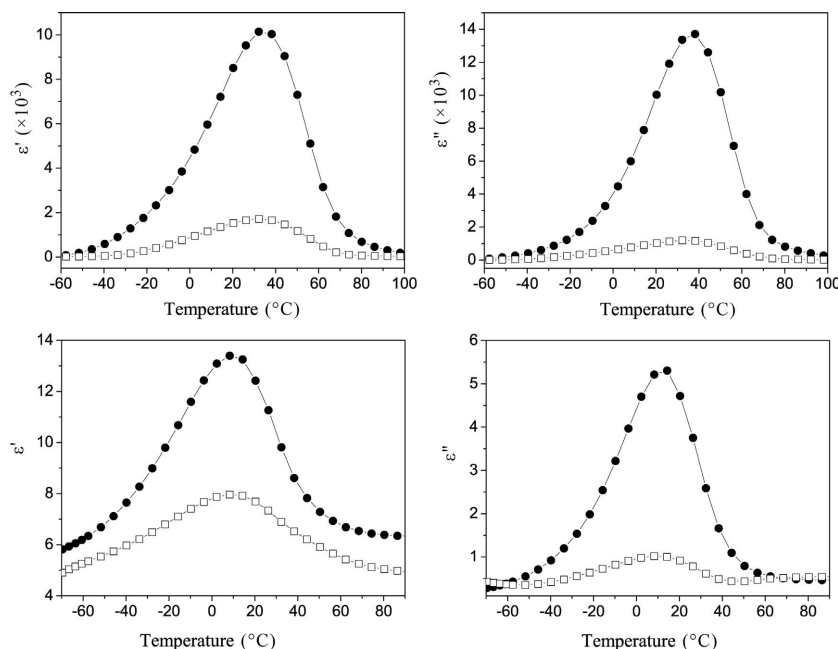


Figure 7. Typical temperature behavior of the dielectric permittivity (left) and dielectric losses (right) at percolation for Ni-montmorillonite (upper) and K-kaolinite (lower) at frequencies of 1 Hz (●) and 133 Hz (□).

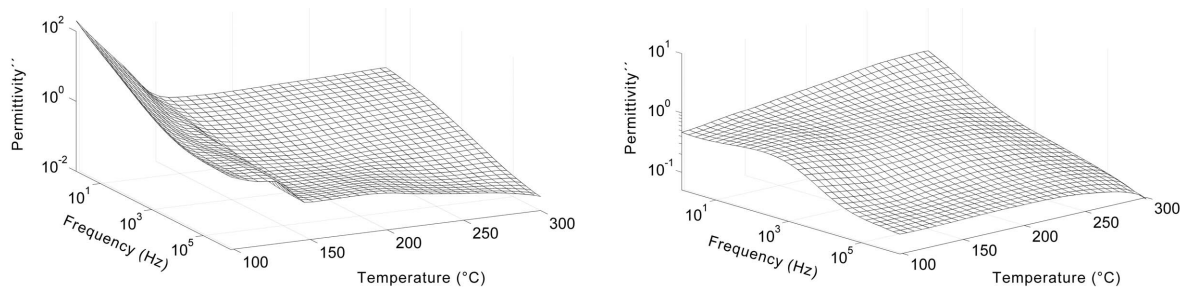


Figure 8. Typical 3D plots of the frequency and temperature dependences of the dielectric loss at high temperatures plotted for Ni-montmorillonite (left) and K-kaolinite (right).

act of relaxation was considered to be an occurrence of two simultaneous events: (1) reorientation of the molecule; and (2) the presence of a defect site in its vicinity. In the framework of the model, the temperature dependence of the relaxation time can be written in the form:

$$\tau = \tau_0 \exp \left\{ \frac{H_a}{kT} + C \exp \left(-\frac{H_d}{kT} \right) \right\} \quad (7)$$

where H_a is the height of the potential barrier of the reorientation of water molecules; H_d is the energy of the defect formation; k is Boltzmann's constant and C is a number inversely proportional to the maximum possible defect concentration, η . The pre-exponential time constant, τ_0 , can vary with temperature. The increase in temperature leads, on the one hand, to an increase in the probability of attaining sufficient energy to break the bonds between the reorienting molecule and its neighbors; or, on the other hand, it leads to a decrease in the probability of finding a defect in the vicinity of the moving molecule. The interplay of these two terms yields the appearance of the minimum observed in the temperature dependence of the relaxation time.

The temperature dependences of characteristic times corresponding to the frequency of the maximum of the relaxation loss peak (Figure 11) revealed that saddle-like process II.1 was observed in both kaolinites and montmorillonites (Figure 5), while an additional sad-

dle-like process, II.2, was found only in the Co- and Ni-montmorillonites (Figure 11).

As discussed above, two main water-adsorption centers are present in the clay-mineral structure: (1) the basal oxygen atoms and cations trapped in hexagonal holes on the surfaces, and (2) exchangeable interlayer cations. Unlike the low-temperature processes I.1 and I.2 related to the relaxation of ice-like structures, the non-monotonous saddle-like processes, labeled II.1 and II.2 in Figure 5, can be assigned to the relaxation of confined water molecules (Gutina *et al.*, 2003; Feldman *et al.*, 2006). The two types of water structures characterized by different dynamics can coexist and that was recently confirmed by differential scanning calorimetry (DSC) experiments by Kozłowski (2011, 2012). The low-temperature exothermic peaks observed in the DSC measurements show that a significant amount of adsorbed water in clay minerals remains in an unfrozen state when cooling to -70°C (Kozłowski, 2011, 2012), which confirms and follows the findings of Low *et al.* (1968a,b). The saddle-like relaxation process II.1 is clearly observed for both clay minerals (kaolinites and montmorillonites) and can be assigned to confined water between two interfaces (outer surface of powder granules on one side and air on the other). At the same time, no other saddle-like process II.2 was observed for kaolinites, which means that in the case of montmorillonites an additional source of confinement was present. The interlayer space of montmorillonite structures may have been the source of this extra second confinement. Exchangeable Co^{2+} and Ni^{2+} ions can form ion-water rigid structures in the interlayer gaps while the K^+ ion is known by its strong disordering effect on

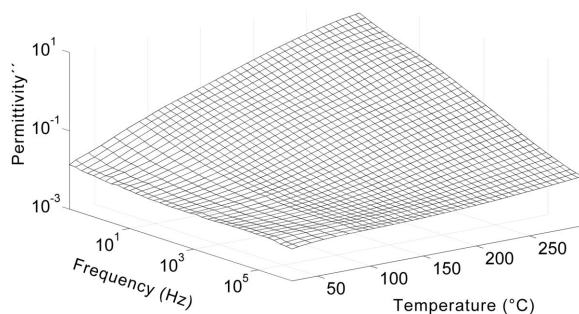


Figure 9. Typical 3D plot of the frequency and temperature dependences of the dielectric losses after annealing the sample for 3 h at 300°C plotted for Ni-montmorillonite.

Table 4. Calculated activation energies, $E_{I,1}$ and $E_{I,2}$, of the low-temperature relaxation processes (I.1 and I.2) in the montmorillonite samples.

Sample	$E_{I,1}$ (kJ/mol)	$E_{I,2}$ (kJ/mol)
K-montmorillonite	35	31
Co-montmorillonite	49	25
Ni-montmorillonite	55	34

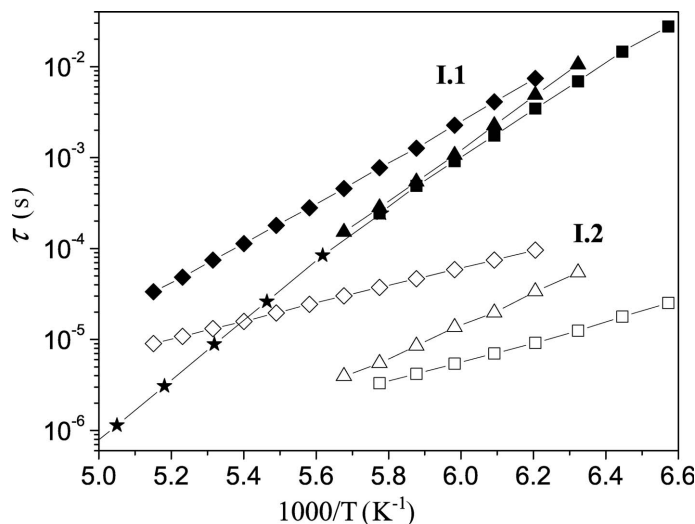


Figure 10. Temperature dependences of the relaxation times of the low-temperature processes for the montmorillonites (◆ K ◇; ■ Co □; ▲ Ni △). Filled symbols correspond to the I.1 process, open symbols to the I.2 process. Stars correspond to the temperature dependence of the relaxation time of the low-temperature process in glass (Gutina *et al.*, 2003).

unfrozen interlayer water. Indeed, the K^+ ion is characterized by hydrophobic properties ('negative hydration') at which the lifetime of water molecules to be in the immediate environment of the ion is less than that in the bulk solution (Krestov, 1991; Samoilov, 1957). Hence, the process II.2 is observed only for the Co- and Ni-montmorillonite samples.

The saddle-like process is shown (Figure 11) to depend heavily on the scale at which the relaxation takes place. Characteristic relaxation times for the process II.1 are slower than those for process II.2. The saddle-like behavior was fitted to equation 7 by means of a least-squares fit procedure. The fitting curves show good

agreement with the proposed model (Figure 11). The resultant fit parameters are listed in Table 5.

The values of the activation energies H_{a1} and H_{d1} for the process II.1 for all clay minerals under study are in good agreement with the energies of molecular re-orientation and defect formation in confined water (Gutina *et al.*, 2003) but depend on the properties of individual ions. A more detailed investigation is required and may become a subject for further study.

The percolation process

At the mid-temperature range, all of the samples studied demonstrated typical percolation behavior of the

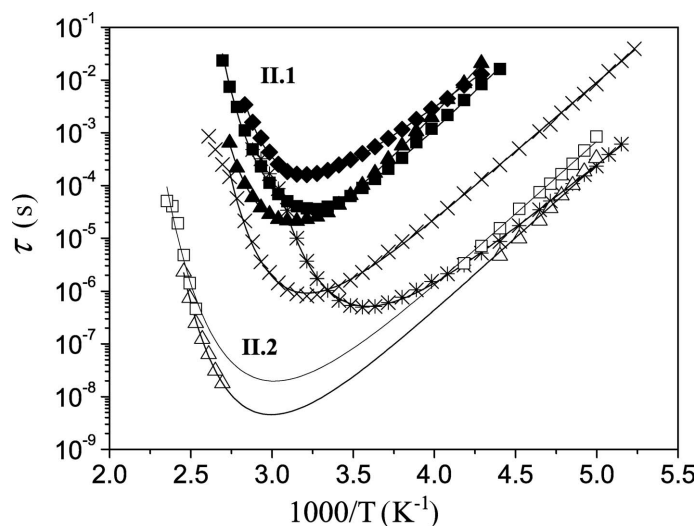


Figure 11. The temperature dependence of the characteristic times of the mid-temperature relaxation processes for montmorillonites (◆ K; □ Co ■; △ Ni ▲) and kaolinites (* K; × K, Ba). Filled symbols for the montmorillonites correspond to the II.1 mid-temperature process and the open symbols to process II.2. The solid lines are the fitting curves according to equation 7.

Table 5. Fitting parameters obtained using equation 7.

Sample Fit parameter	Process II.1				Process II.2			
	τ_0 (s)	H_{a1} (kJ/mol)	H_{d1} (kJ/mol)	η	τ_0 (s)	H_{a2} (kJ/mol)	H_{d2} (kJ/mol)	η
K-montmorillonite	2×10^{-12}	43.2	36.0	7×10^{-7}	—	—	—	—
Co-montmorillonite	1×10^{-14}	53.2	41.6	1×10^{-7}	1×10^{-17}	53.0	24.0	7×10^{-5}
Ni-montmorillonite	1×10^{-14}	55.6	42.5	1×10^{-7}	3×10^{-18}	53.2	27.7	2×10^{-5}
K-kaolinite	2×10^{-16}	46.6	26.3	6×10^{-6}	—	—	—	—
K,Ba-kaolinite	6×10^{-16}	50.3	31.6	3×10^{-6}	—	—	—	—

dielectric spectrum as mentioned in the results section above. Such a process was previously associated with the percolation of an apparent dipole moment excitation, within a developed fractal structure of connected pores in a hydrated porous medium (Gutina *et al.*, 2003; Puzenko *et al.*, 1999). Note that the fractal geometry is a mathematical tool for dealing with the complex systems that have no characteristic length scale. Scale-invariant systems are usually characterized by a non-integer (so-called fractal) dimension (Bunde and Havlin, 1995). Similar to porous silica glasses, in the clay minerals the dipole excitation can also be linked to the self-diffusion along random paths of charge carriers within the channels. A transfer of the electric excitation along the developed fractal structure of the connected pores can be described by the normalized dipole correlation function (DCF), $\Psi(t)$. Note that a transfer of the electric excitation through the porous medium can occur even in the case of closed pores which are topologically not connected one to another. However, the distance between the neighboring pores filled with the dielectric or conductive material should be small enough to provide a ‘physical’ pore coupling *via* the electric interaction. A detailed description of the relaxation mechanism associated with an excitation transfer based on a regular fractal model was introduced by Feldman *et al.* (2006), where it was applied for the cooperative relaxation of ionic micro-emulsions at percolation. Based on percolation theory and the fractal characterization of the clay minerals, the DCF at the percolation threshold was represented by an asymptotic stretched-exponential term (Gutina *et al.*, 2003; Puzenko *et al.*, 1999):

$$\Psi(t) = \exp \left[- \left(\frac{t}{\tau_p} \right)^{D_p/3} \right] \quad (8)$$

where τ_p is an effective relaxation time and D_p is a fractal dimension of the media. In order to determine the value of D_p , the relaxation law (8) can be fitted to the experimental correlation functions. For this purpose the complex dielectric permittivity data were expressed in terms of the dipole correlation function using the Laplace transform.

Dielectric permittivity data were taken at the percolation threshold temperature for all studied sam-

ples. The DCF (Figure 12) values were then fitted to equation 8 after applying the Laplace transform. The D_p values obtained are presented in Table 6.

Note that the fractal dimension discussed here is the fractal dimension of the dipole excitation transfer paths connecting the different kinds of hydration centers located on the pore surface. The calculated fractal dimension of the pathways connecting the hydration centers lies between 1 and 2 (Table 6). Due to the relatively small water-content values of all the samples, the water molecules adsorbed by the materials are bound to these centers. The paths of the excitation transfer span the fractal pore surface and ‘depict’ the backbone of clusters formed by the pores on a scale that is larger than the characteristic distance between the hydration centers. Thus, a fractal dimension of the paths D_p approximates the real surface fractal dimension in the scale interval considered.

The values of the fractal dimension, D_p , of montmorillonites are a little larger than those of kaolinites (Table 6). This fact, as well as the increase in the percolation temperature values (Table 3) for montmorillonites, can be explained by the much more developed surface structure of this type of clay mineral due to the presence of interlayer spaces that are wide compared to kaolinites.

An additional method to quantify the structural information of the clay minerals using a fractal analysis of the SEM images was applied in the present study. Fractal analysis is a powerful approach for quantifying the structure of complex systems which exhibit a measure of self-similarity. Such similarity is usually reflected at different length scales, as discussed in detail

Table 6. Fractal dimensions of the montmorillonite and kaolinite samples.

Sample	D_p	D_f
K-montmorillonite	1.74	1.85
Co-montmorillonite	1.67	1.79
Ni-montmorillonite	1.75	1.80
K-kaolinite	1.48	1.66
K,Ba-kaolinite	1.30	1.81

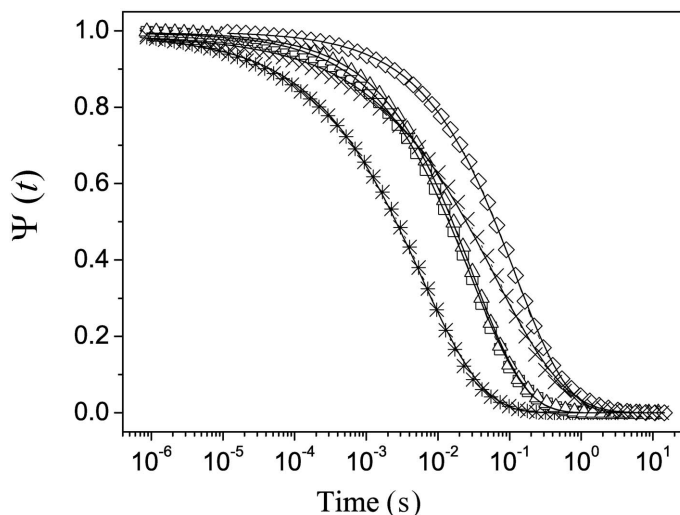


Figure 12. Semi-log plot of the dipole correlation function $\Psi(t)$ for montmorillonites (\diamond K; \square Co; Δ Ni) and kaolinites ($*$ K; \times K, Ba) at the temperatures corresponding to the percolation point. The solid line corresponds to the fit.

by Narine and Marangoni (1999) and by Tang and Marangoni (2006a, 2006b).

In an SEM image the grayscale intensity is proportional to the degree of electron scattering from the sample surface and particularly from areas of differing charge density. Consequently, an analysis of the fractality inherent in the image should reflect the underlying pathways of excitation. The fractality was calculated by counting the number of pixels, $S(N_j)$, in the image with a grayscale from 200 to 255, where the total number of pixels at level j of the recursive coarsening of the image is N_j . For simplicity, N_j was maintained as a power of 2. The detailed description of the calculations can be found in the study by Libster *et al.* (2008). The grayscale is presented below with the cutoff marked (Figure 13).

The image was coarsened by taking the average value of a moving square of four pixels, defining a reduced image. The count was then made again and the process repeated until the image size was too small. Mathematically, this is represented by the expression $N_j = 2^{2(l_{\max}-j)}$, where l_{\max} is the maximum number of recursive stages for the image. A plot was then made in double log scale of the number of counts of white pixels against the image size in pixels.

Power-law behavior was observed and the slopes were calculated from the fitting, according to the relation

$$S(N_j) \propto N_j^D \quad (9)$$

where D is the boundary fractal dimension based on the SEM image depicting the different areas of charge density. This can then be related to the spatial fractal dimension, D_f , by the relationship $D_f = 3 - D$ (Wong and Cao, 1992). The results are summarized in Table 6. The agreement that is evident between these two separate methods of obtaining the spatial fractal dimension points toward the importance of the hydration centers. The D_f values are slightly greater than D_p , as expected, because they describe in more detail the surface structure rather than only the paths connecting the hydration centers.

CONCLUSIONS

Montmorillonites with exchangeable K^+ , Co^{2+} , and Ni^{2+} cations and kaolinites with exchangeable K^+ and Ba^{2+} cations were examined by means of Broadband Dielectric Spectroscopy over wide temperature ($-121^\circ C$ to $+300^\circ C$) and frequency (1 Hz–1 MHz) ranges. The rich dielectric spectra can be described in terms of four distributed relaxation processes. Over the low-temperature interval the two Cole-Cole-type dielectric relaxation processes (I.1 and I.2) were observed for montmorillonites only, associated with the interlayer ice-like water molecule relaxations linked to the two types of hydration centers that are located at the hydrophilic centers of interlayer channels (siloxane bonds and trapped cations) and around the interlayer cations.



Figure 13. The gray scale with divisions every eight shades of gray. (The gray scale from white to black, usually varies in 255 shades of gray. For clarity in the figure each division represents the average of eight shades of gray). The cut-off used for the analysis is marked.

The saddle-like behavior of water adsorbed in the mid-temperature interval has been ascribed to the relaxation of confined water molecules (process II.1). While an additional saddle-like process (II.2) related to the interlayer space confinement was observed for Co- and Ni-montmorillonites only. The absence of the II.2 process for K-montmorillonite is explained by strong disordering effect of the K^+ ion ('negative hydration').

The third relaxation process, occurring in both clay minerals, is associated with percolation of the electric excitation (the self-diffusion of the charge carriers) within the developed fractal structure of connected pores on the outer surface of the granules. The fractal dimensions of the electric excitation paths coming along the pore surface from one hydration center to another are between 1 and 2, consistent with fractal dimension calculated from micrographs for all the samples studied.

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