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Thermodynamic properties of mixed-layers illite-smectite by calorimetric methods. Acquisition of the enthalpies of mixing of illite and smectite layers.

H. Gailhanou^{1*}, P. Blanc¹, J. Rogez², G. Mikaelian², H. Kawaji³, J. Olives⁴, V. Montouillout⁵, J-M. Grenèche⁶,
P. Vieillard⁷, E.C. Gaucher¹, C.I. Fialips⁸, B. Made⁸

¹BRGM, 3 av. C. Guillemin, BP6009, 45060 Orléans, France

²IM2NP, F.S.T. Saint-Jérôme, Case 251, Aix Marseille Université, 13397 Marseille Cedex 20, France

³Tokyo Institute of Technology, 4259 Nagatsuta-cho, Midori-ku, Yokohama, 226-8503 Japan

⁴CINaM-CNRS, Aix-Marseille Univ., Campus de Luminy, Case 913, 13288 Marseille Cedex 9, France

⁵CEMHTI-CNRS-UPR-3079, 1D av. de la Recherche Scientifique, 45071 Orléans, France

⁶IMMM-UMR-CNRS-6283, Le Mans Université, 72085 Le Mans cedex, France

⁷CNRS-IC2MP-UMR-7285 Hydrasa, 5 av. Albert Turpain, 86073 POITIERS-Cedex, France

⁸ANDRA, 92298 Châtenay-Malabry, France.

Corresponding author:

Hélène GAILHANOU

BRGM, 3 av. Claude Guillemin BP6009

45060 Orléans, France.

*E-mail: h.gailhanou@brgm.fr

Abstract --- The stability of illite-smectite interstratified with respect to discrete illite and smectite minerals was investigated by measuring thermodynamic properties. The standard thermodynamic properties of the illite-smectite ISCz-1 (G, H, S, Cp, V) were determined between 298.15K and 375K by using calorimetric methods. Moreover, the enthalpies of mixing between illite and smectite layers were measured at 298.15K by acid solution calorimetry from a complete serie of illite-smectite interstratified minerals (Shinzan area, Japan). The measured values were slightly negative, with a minimum value at -3.7 kJ.mol^{-1} , and contribute to the stability of the mixed-layer with respect to a mechanical mixture of illite and smectite. Besides, the model from Blanc et al. (2015) was implemented to estimate the thermodynamic properties of the interstratified illite-smectite ISCz-1 mineral. The predicted values were consistent with the experimental ones. However, the estimates were slightly improved by considering the thermodynamic properties of the mixture of illite and smectite components and then adding the terms of energies of mixing. This could be confirm by setting up the stability domains of ISCz-1 and the corresponding illite and smectite end-members, according to Meunier and Velde (1989) determination of the smectite to illite reaction pathways.

Key words: illite, smectite, enthalpies, heat capacities, entropies, Gibbs free energies, calorimetry, enthalpies of mixing

Running title: Thermodynamic properties of illite-smectite by calorimetry

1. INTRODUCTION

Illite/smectite mixed layer clay minerals (I/S) are widely present in natural environment and occur in various contexts, notably during diagenetic processes in sedimentary basins (Perry and Hower, 1970; Hower et al., 1976, among others), or resulting from the hydrothermal alteration of volcanic or plutonic rocks (Steiner, 1968; McDowell and Elders, 1980; Inoue and Utada., 1983, among others). Understanding the processes associated with the transformation of smectite into illite/smectite and consequences on the mechanical and the geochemical behaviour of the clayey formation is of great importance in some major applications fields, such as oil exploration or nuclear waste disposal. For instance, this transformation is an indicator of possible oil reserves for a given reservoir. It was and is still widely used as a geothermometer for the early diagenesis stage (possibly combined with other tools like the Na/K or Si geothermometer or stable isotope analyses) in order to assess the maturity, in terms of P-T-X reaction pathway, of a given reservoir (Velde and Vasseur, 1992). In addition, the smectite-to-illite conversion may generate overpressures in sedimentary basins due to the dehydration of smectite and the precipitation of secondary minerals (Powers, 1967; Colton-Bratley, 1987; Osborne and Swarbrick, 1997) that can be problematic for the functioning of oil wells. Besides, according to Gaucher and Blanc (2006) review, the smectite transformation to illite-smectite and then illite could be possibly expected at cement/clay interfaces, after long period of reaction times, within the framework of deep disposal. Alkaline disturbance will generate mineralogical transformations at the cement/clay interface, in particular of bentonite, implying a decrease of smectite content. This latter will affect the swelling properties and the retention capacities of clayey barriers. Such reactions have to be integrated in the geochemical calculations used for performance assessment.

The thermodynamic stability of illite/smectite remains an unsolved issue. Numerous studies based on mineralogical characterization of clays in natural systems (Inoue et al., 2005; Ylagan et al., 2000; Murakami et al., 2005; Claret et al., 2004; McCarty et al., 2008; Lanson et al., 2009, among others) and on laboratory experiments (Ferrage et al., 2011; Yates and Rosenberg, 1996, 1997; Rosenberg et al., 1990; Whitney and Northrop, 1988; Whitney and Velde, 1993, among others) were performed in order to better understand the

reaction mechanisms of the smectite illitization during hydrothermal alteration and diagenetic processes. Several authors (Whitney and Northrop, 1988; Whitney and Velde, 1993; Rosenberg et al., 1990; Yates and Rosenberg, 1996; Ferrage et al., 2011 and references therein) showed the coexistence of several phases (smectite, illite and one or more intermediate illite/smectite phases) during the transformation of smectite to illite. Collecting the chemical analyses of natural I/S samples, Meunier and Velde (1989) have described the interstratified minerals by a ternary solid-solution model with one illite and two montmorillonite end-members. Based on this approach, Blanc et al. (1997) developed a thermodynamic model allowing the calculation of the energies of mixing of illite and smectite layers depending on the degree of ordering of the stacking sequences. The resulting Gibbs free energies of mixing are then slightly negative, suggesting the stability of illite-smectite mixed-layers with respect to the mechanical mixture of pure illite and smectite layers.

The present work focuses on the measurement of thermodynamic properties for I/S minerals with two main objectives:

- to acquire a complete set of thermodynamic properties (G, H, S, Cp) for an illite-smectite mineral (ISCz-1), based on calorimetric methods (Gailhanou et al., 2012, 2013), which could be used to supply the thermodynamic databases (e.g., ThermoChimie (Giffaut et al., 2014) and Thermoddem (Blanc et al., 2012)) for geochemical modelling calculations. Until now, the lack of thermodynamic data for illite/smectite mixed layers is a limitation to properly model the smectite-to-illite conversion;

- to acquire the enthalpies of mixing of illite-smectite layers for a series of interstratified illite-smectite (hydrothermal alteration minerals, coming from Shinzan Area; Inoue and Utada, 1983; Inoue et al., 1987).

Besides, the present work allows checking the reliability of the predictive model from Blanc et al. (2015) for the thermodynamic properties of interstratified illite-smectite. Based on the properties measured here, the influence of the thermodynamic parameters of mixing can be discussed, including the configurational entropy associated with cationic disordering in crystallographic sites of clay minerals and specific interactions between smectite and illite layers, within the I/S stacking sequence.

2. MATERIALS AND METHODS

2.1. Clay samples

The ISCz-1 sample is a mixed-layer illite-smectite, originating from Slovakia. It was provided by the Source Clays Repository of the Clay Minerals Society (Chipera and Bish, 2001).

After sieving the ISCz-1 sample at a size $< 20 \mu\text{m}$, the $< 2 \mu\text{m}$ fraction was separated by centrifugation. This latter was then twice dispersed in CaCl_2 0.5 M solution during 24h and centrifuged. Therefore, the sample was washed with a CaCl_2 0.001 M solution, by using dialysis bag for 3 days. In the present study, Ca^{2+} was preferred to Na^+ for the saturation of the smectitic interlayers, to facilitate the washing of the clay sample by avoiding osmotic swelling of smectite. Eventually, the sample was dried at 40°C then 60°C for several days. In the following, the sample referenced ISCz-1 designates this treated sample.

A series of fourteen illite-smectite mixed-layers samples coming from the Shinzan area, northeast Japan, has also been used for this study. This series, well described in literature (Inoue et al., 1987; Inoue and Utada, 1983), is noticeable as it consists in a complete mixed-layers series with chemical compositions varying from montmorillonite to illite end-members. The series results from the hydrothermal alteration of dioctahedral micas/smectites. The continuous conversion of smectite into illite has been described in Inoue et al. (1987). The main transformation mechanism suggested by the authors was based on the dissolution of smectite and the crystallization of illite. A solid-state transformation mechanism was also shown from HRTEM observations (Amouric and Olives, 1991; Olives et al., 2000).

2.2. Analysis and characterization of the samples

2.2.1. ISCz-1 sample

Chemical analyses for Si, Al, Ti, Fe(total), Mn, Ca, Mg, K, Na and P were performed by X-ray fluorescence spectrometry. The results of the analysis are presented in Table 1; the relative uncertainties are estimated at

±2 %. The amounts of total carbon and total sulfur were determined by infra-red spectroscopy after burning the samples at 900 °C in an oxygen atmosphere. The amount of Fe²⁺ was determined by ⁵⁷Fe Mössbauer spectrometry, as described further in the text.

The chemical composition of the ISCz-1 sample (Ca-saturated) differs slightly from the previous Na-saturated ISCz-1 sample used for heat capacity measurements in Gailhanou et al. (2007) (Table 1). Both samples were supplied by the Clay Minerals Society with several years of intervals. The following study of the thermodynamic properties of the present ISCz-1 sample is complementary to the previous work from Gailhanou et al. (2007).

Table 1 Chemical analyses of the sample ISCz-1 (< 2µm clay fraction). (a) X-ray fluorescence; (b) ⁵⁷Fe Mössbauer spectrometry; (c) weight loss by heating up to 1000°C; (d) infra-red spectroscopy.

	ISCz-1 (this study) wt. %	Na-sat. ISCz-1 (Gailhanou et al., 2007) wt. %	Analytical technique
SiO ₂	53.8	52.6	(a)
Al ₂ O ₃	24.4	26.7	(a)
CaO	1.1	< 0.1	(a)
Fe ₂ O ₃	1.39	0.97	(a), (b)
FeO	0.17	0.30	(b)
K ₂ O	5.07	5.96	(a)
MgO	2.2	2.1	(a)
Na ₂ O	< 0.2	1.0	(a)
TiO ₂	0.13	0.09	(a)
P ₂ O ₅	< 0.05	0.13	(a)
Loss on ignition	11.9	10.0	(c)
C total	0.07	0.04	(d)
S total	< 0.01	0.06	(d)
Total	100.25	99.95	

The cationic exchange capacity (CEC) of the sample was measured after saturating the sample with a chloride cobalt hexammine solution, according to the method described in Gaucher et al. (2009). The total CEC was measured by colorimetry and the exchangeable cations (Na⁺, Ca²⁺, K⁺, Mg²⁺) were analyzed by ionic chromatography. The results are provided in Table 2. The obtained CEC value is consistent with the one measured for the ISCz-1 sample in Gailhanou et al. (2007), at 39.0 meq/100 g.

Table 2. CEC measured for ISCz-1 sample (< 2 µm clay fraction), in meq/100 g of dry sample.

Sample	Na	Ca	Mg	K	Sum of exch. Cations	CEC (colorimetry)
ISCz-1	0.4	39.7	< 0.5	3.8	43.9	44.1

Moreover, XRD analyses were carried out on the ISCz-1 sample, on powder and on oriented deposit, using the method described in Gailhanou et al. (2007). For the powder sample, both an identification of the nature of impurities and a semi-quantification were performed by using Modpattern (Guyonnet et al., 2009) with an internal corundum standard (Figure 1). In the present case, only rough estimates of the proportions of the impurities could be obtained by modelling, due to the lack of a reference pattern for Ca-saturated illite-smectite.

Moreover, the modelling of the XRD pattern on the glycolated, oriented deposit was performed by using the Arquant code (Blanc et al., 2007) and allowed the identification and the quantification of a small amount of kaolinite. Finally, the amounts of impurities in the purified sample estimated by XRD modeling are 4.2, 3.8 and 0.45 wt.% of quartz, K-feldspar and kaolinite, respectively. Moreover, resulting from the modelling of the XRD pattern on the glycolated deposit and in agreement with Chipera and Bish (2001) results, the interstratified ISCz-1 was ordered with a degree of interstratification ordering $R = 1$ (Reichweite index) and the illite layers content was estimated at 70 %.

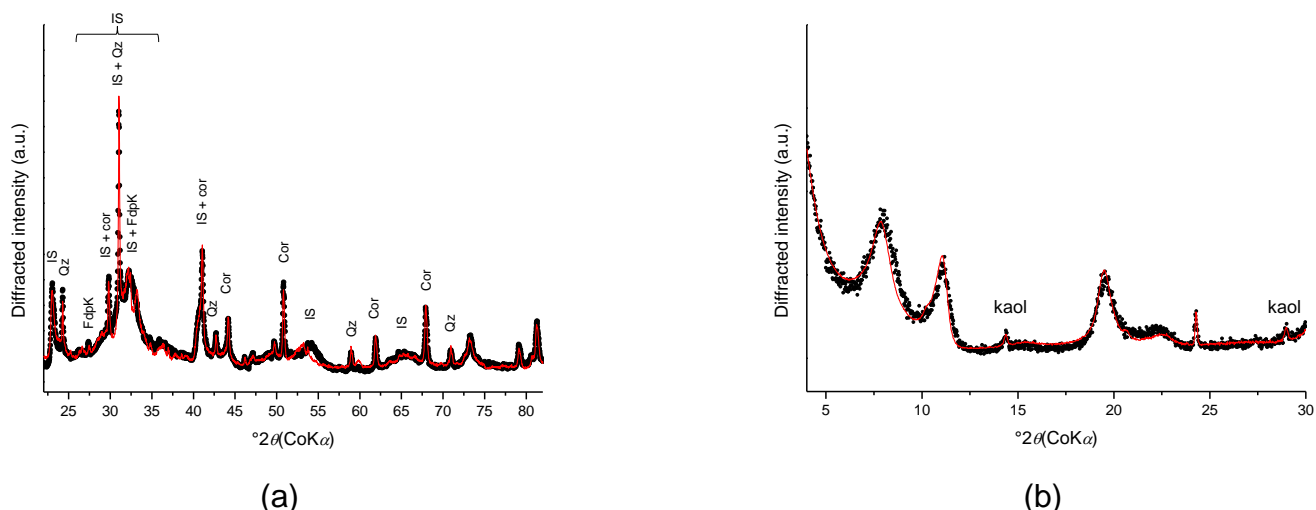


Figure 1. Modelled XRD patterns obtained for ISCz-1 sample (< 2 μm , Ca-saturated) on (a) powder sample and (b) glycolated oriented deposit. IS = mixed-layer ISCz-1; Qtz = quartz; Cor = corundum (internal standard, 9.19 wt.%); FdpK = K-feldspar; Kaol = Kaolinite (in black, experimental; in red, modelling).

Besides, some investigations were performed by using ^{29}Si and ^{27}Al NMR in order to complete the characterization of the silicates and aluminosilicates impurities. A ^{29}Si NMR spectrum was acquired with a Bruker Advance 300 MHz ($B_0 = 7.0\text{ T}$) spectrometer, operating at 79.5 MHz frequency for ^{29}Si (Figure 2(a)). It revealed notably, in addition to illite-smectite component (between -80 and -100 ppm), the presence of quartz (-107.52 ppm) and the absence of other silica impurity (cristobalite or amorphous silica, notably). The amount of quartz was estimated to 5.4 wt.% in the ISCz-1 sample. This result is in good agreement with the XRD estimate at 4.2 wt.%.

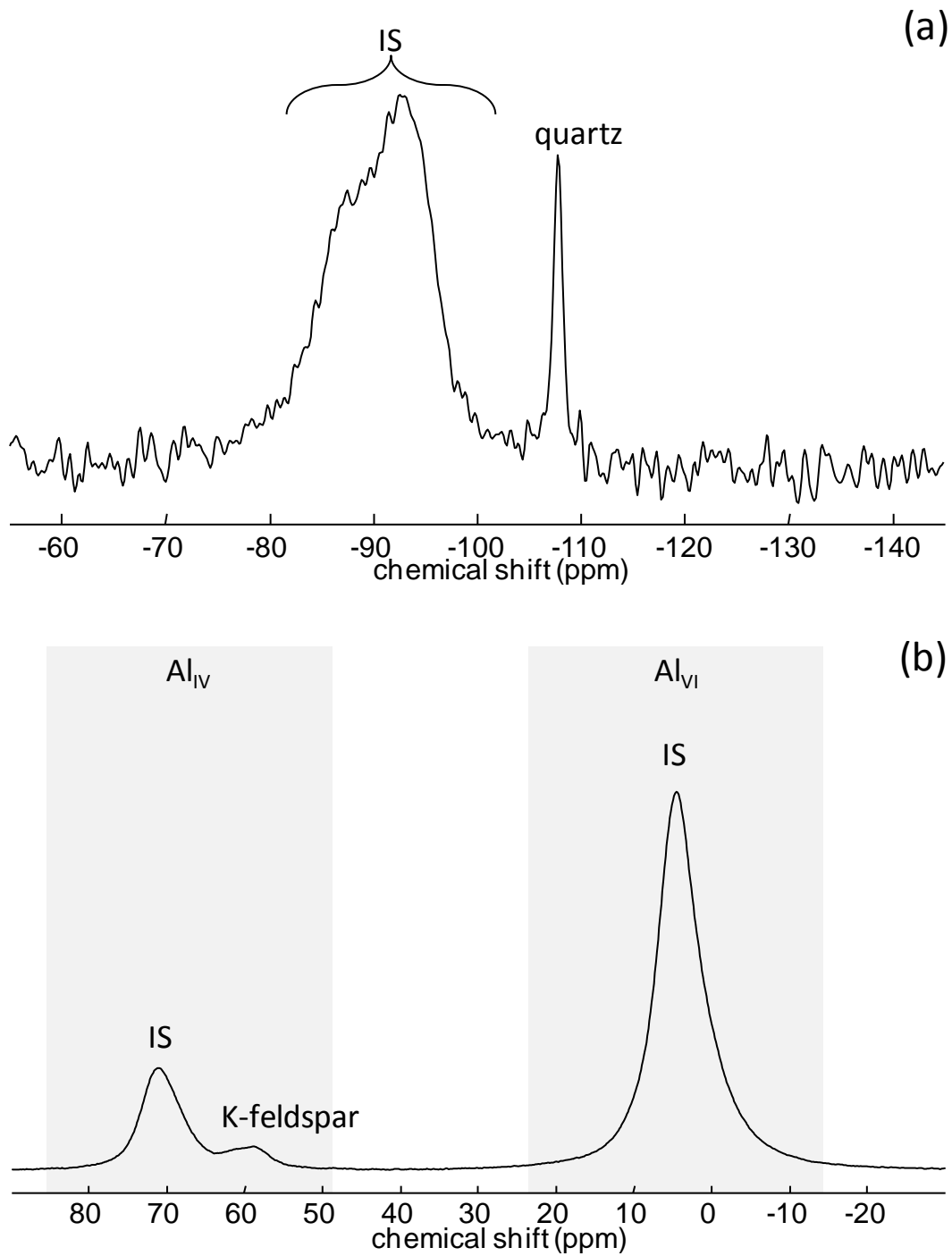


Figure 2. Identification and quantification of impurities in ISCz-1 sample (< 2 μm clay fraction) by NMR: (a) ^{29}Si and (b) ^{27}Al NMR spectra.

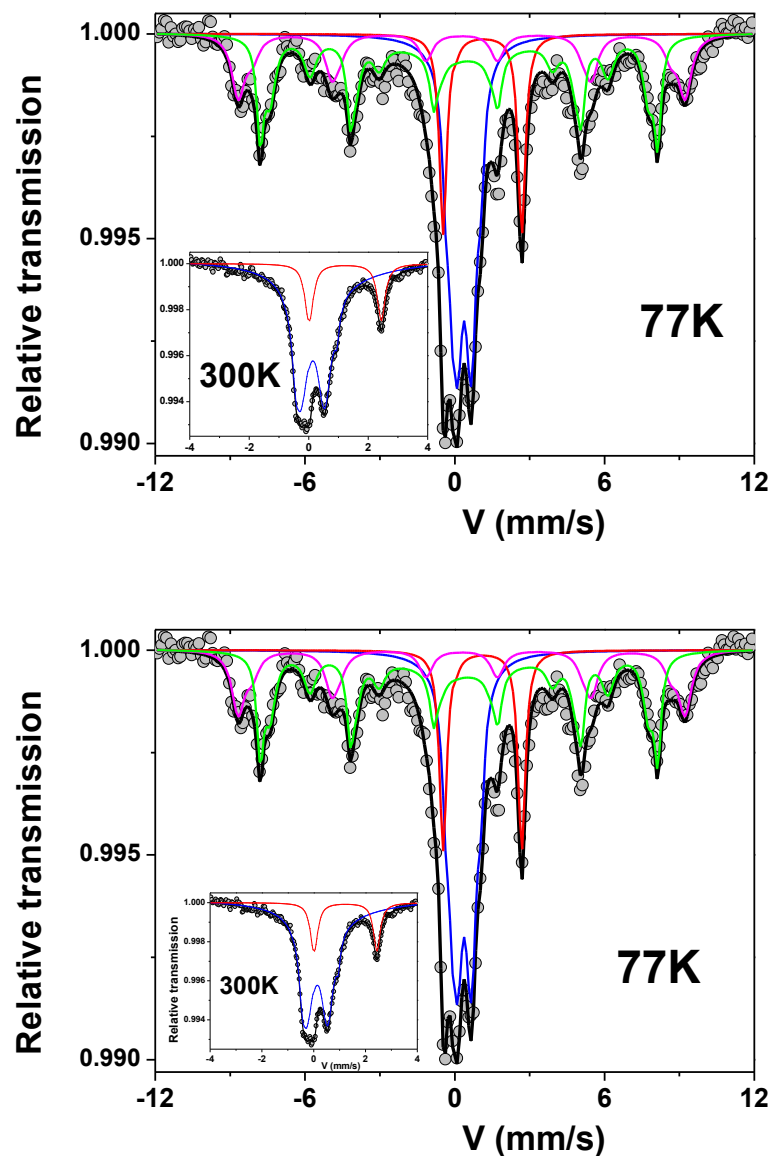
For its part, the ^{27}Al NMR spectrum (Figure 2 (b)) acquired with a Bruker Advance 750 MHz spectrometer ($B_0 = 17.7 \text{ T}$, $\nu_{\text{RF}}(^{27}\text{Al}) = 195.5 \text{ MHz}$) exhibits three main signals for one six-fold coordinated aluminum

($\delta_{\text{CG}} = 4.6$ ppm) and two four-fold coordinated aluminum ($\delta_{\text{CG}} = 59.4$ and 71.1 ppm). Both $\delta_{\text{CG}} = 4.6$ and 71.1 ppm sites are commonly attributed to Al^{VI} and Al^{IV} in illite-smectite (Cuadros and Linares, 1996; Garg and Skibsted, 2016), while the signal at $\delta_{\text{CG}} = 59.4$ ppm is compatible with Al^{IV} in K-feldspar, detected by XRD (Cuadros and Linares, 1996). The spectrum was modeled using the Dmfit software (Massiot et al., 2002). The component at $\delta_{\text{CG}} = 59.4$ ppm corresponds to 2.9 at % of the aluminum of the sample. The resulting proportion of K-feldspar impurity in the ISCz-1 sample was then estimated to 3.8 wt.%, in agreement with the value obtained by XRD modeling.

The proportions of ferric and ferrous ions in the clay mineral and, if present, the amounts of ferrous or ferric impurities can be determined by ^{57}Fe Mössbauer spectrometry. These analyses were performed using a conventional constant acceleration Mossbauer spectrometer, in transmission geometry, equipped with a $^{57}\text{Co}(\text{Rh})$ source. The isomer shift values were corrected according to the calibration of the velocity scale made from $\alpha\text{-Fe}$ at 300 K. The amount of Fe in the sample was about 5 mg/cm^2 . The spectra obtained at 300 and 77 K were fitted using a discrete number of independent quadrupolar doublets and magnetic sextets composed of lorentzian lines where the line width at half-height Γ (mm.s^{-1}), the isomer shift δ (mm.s^{-1}) and the quadrupole splitting ΔE_{Q} (mm.s^{-1}) were refined using a least-square fitting procedure.

The ^{57}Fe Mössbauer spectra for the ISCz-1 sample ($< 2 \mu\text{m}$) are displayed in Figure 3. The 300K spectrum (inset Fig. 3) exhibits a quadrupolar broadened asymmetrical feature and a single line which can be described by means of two components clearly attributed to ferric and ferrous species. But one can see that the baseline remains distorted (as confirmed by a spectrum obtained at 12 mm/s, not shown here): this feature results from the presence of superparamagnetic Fe species, i.e. the presence of ultra-fine Fe-containing grains. Consequently, the hyperfine structure was investigated at 77K to get blocked magnetic states of some Fe species, as confirmed in Fig.3. The description requires unambiguously several magnetic sextets and quadrupolar doublets but a single broadened line was added to better describe the baseline. The refined values of Mössbauer parameters extracted from spectral modeling are listed in Table 3. The proportion of Fe in the clay mineral was estimated to 42 at.% of the total Fe content and the ratio $\text{Fe}^{3+}/(\text{Fe}^{2+} +$

210 Fe^{3+}) to 72 % (Table 5). It is important to note the good agreement between the Fe^{2+} contents estimated at
 211 300K and 77K, assuming that the proportions are proportional to the relative absorption areas, i.e. assuming
 212 the same values of the Lamb-Mössbauer recoilless factors. The modeling allowed to distinguish three
 213 magnetic phases, goethite ($\alpha\text{-Fe}^{3+}\text{O}(\text{OH})$) as fine grains (magnetic sextets) and ultra-fine superparamagnetic
 214 grains (single line) and probably bernalite (ideally $\text{Fe}^{3+}(\text{OH})_3$, McCammon et al., 1995) or bernalite/hematite
 215 mixture.
 216



217
 218
 219 Figure 3. ^{57}Fe Mössbauer spectra for sample ISCz-1 (< 2 μm clay fraction). The 77K spectrum and its decomposition
 220 into 4 components: in red, for Fe^{2+} in clay; in dark blue, for Fe^{3+} in clay; in green, magnetic component for Fe^{3+} in

goethite; in pink, magnetic component for Fe³⁺ in bernalite. Inset represents the 300K spectrum with blue and red components attributed to ferric and ferrous species, respectively (see text).

Table 3. ⁵⁷Fe Mössbauer parameters at 77 K for the ISCz-1 sample (< 2 µm clay fraction)

Mineral	I.S. (mm/s)	2ε/ΔE _Q (mm/s)	B _{hf} (T)	%	attributions
	± 0.02	± 0.01		± 2	
Illite-smectite	0.47	0.57		27	Fe ³⁺ associated with clay minerals (structural and/or sorbed ions)
	0.42	1.32		4	Fe ³⁺ associated with clay minerals
Illite-smectite	1.30	3.06		12	Fe ²⁺ associated with clay minerals
Goethite	0.46	-0.27	49.2	28	Fe ³⁺ in goethite
	0.47	-0.27	46.7		
	0.47	-0.27	36.8		
Bernalite	0.40	0.00	51.8	17	Fe ³⁺ in bernalite
	0.40	0.00	55.3		
Goethite (ultra-fine grains)	0.46	0.00		12	Fe ³⁺ in goethite

I.S. = Isomer shift value relative to that of the α-Fe at 300 K; 2ε: quadrupolar shift; ΔE_Q = quadrupolar splitting value; B_{hf} = magnetic field; % = ratio of each component.

The 2σ standard errors for I.S., 2ε, ΔE_Q, B_{hf} and % are, respectively, ± 0.02, ± 0.02, ± 0.01, ± 0.5 and ± 2.

Thus, from these results and considering the total amount of Fe₂O₃ in the sample (Table 1), the amounts of goethite and bernalite were estimated to 0.70 wt.% and 0.36 wt.%, respectively.

From the preceding results, the final amounts of impurities were calculated and are reported in Table 4. The mean structural formula of illite-smectite was eventually calculated (Table 4) by subtracting the contributions of impurities, obtained at ambient R.H., to the global chemical analyses from Table 1.

Table 4. Amounts of impurities in the ISCz-1 sample, mean structural formula and molar mass of illite-smectite.

Impurity	Amounts at ambient R.H. (wt.%)	Amounts in dry sample (wt.%)	Analytical mean
Kaolinite	0.45	0.49	XRD
Quartz	5.40	5.85	²⁹ Si NMR
K-feldspar (orthoclase)	3.80	4.12	²⁷ Al NMR
goethite	0.70	0.76	⁵⁷ Fe Mössbauer
bernalite	0.36	0.39	⁵⁷ Fe Mössbauer
Sample	Mean structural formula		Molar mass (g.mol ⁻¹)
ISCz-1 (this work)	(Ca _{0.092} K _{0.440})(Si _{3.562} Al _{0.438})(Al _{1.721} Mg _{0.255} Fe ³⁺ _{0.029} Fe ²⁺ _{0.011} Ti _{0.008})O ₁₀ (OH) ₂		382.01
ISCz-1 (Gailhanou, 2007)	(Na _{0.134} K _{0.530})(Si _{3.565} Al _{0.435})(Al _{1.717} Mg _{0.218} Fe ³⁺ _{0.050} Fe ²⁺ _{0.017})O ₁₀ (OH) ₂		385.04

Finally, the mineralogical and chemical compositions of the present ISCz-1 sample were compared to those obtained for the previous Na-saturated ISCz-1 sample, which had been used for heat capacity measurements in Gailhanou et al. (2007). Both samples were supplied by the Clay Minerals Society with several years of intervals. The amounts of impurities are significantly higher in the present ISCz-1 sample than in the previous one, which contained about 0.75 wt.% of quartz and 1.5 wt.% of kaolinite. Moreover, the mean structural formula of the Ca-saturated ISCz-1 sample (this study) exhibits a slightly lower total layer charge (0.62 eq./O₁₀(OH)₂) compared with the one previously obtained for the Na-saturated ISCz-1 sample, at 0.66 eq./O₁₀(OH)₂. The decrease of the layer charge is probably correlated to the increase of the total CEC, respectively 0.44 and 0.39 eq.kg⁻¹, for the present Ca-saturated ISCz-1 sample and the previous Na-saturated ISCz-1 sample, both indicating an amount of smectite layers slightly higher in the present case than for the sample analyzed previously.

2.2.2. Illite-smectite series from the Shinzan Area

The illite/smectite samples covers all the range of compositions from 0 % to 100 % of smectite (Table 5) as determined by X-ray diffraction from ethylene-glycol solvated samples (Inoue et al., 1987). According to these authors, the samples containing more than 50 % of smectite layers were randomly interstratified (Reichweite index R = 0), while the degrees of interstratification ordering increased progressively to R = 1, R = 2 and R ≥ 3 with increasing illite contents.

The chemical compositions of all the studied samples were determined by atomic absorption spectroscopy (Perbost et al., 2002). The mean structural formulae for the illite/smectite samples are given in Table 6.

Table 5. Illite/smectite interstratified minerals from the Shinzan hydrothermal alteration area in Japan.

Sample	Drill Hole	Depth (m)	% smectite layers
A (smectite)	Ws-2	183,5	100 % (R = 0)
B	Ws-4	329,5	85 % (R = 0)
C	Ws-4	383,0	73 % (R = 0)
D	Ws-2	377,8	70 % (R = 0)
E	Ws-2	407,4	68 % (R = 0)
F	Ws-4	392,8	50 % (R = 0)
G	Ws-2	423,2	40 % (R = 1)
H	Ws-10	199,9	35 % (R = 1)
I	Ws-4	440,6	30 % (R = 1)
J	Ws-5	240,1	20 % (R = 2)
K	Ws-7	123,6	15 % (R ≥ 3)
L	Ws-7	115,5	15 % (R ≥ 3)

M	Ws-7	197,0	5 % (R ≥ 3)
N (illite)	Ws-8	420,9	0 %

Table 6. Mean structural formulae of illite/smectite mixed-layers from the Shinzan area, on the basis of 12 oxygen atoms.

	A	B	C	D	E	F	G	H	I	J	K	L	M	N
Si	3.886	3.822	3.786	3.714	3.692	3.728	3.586	3.524	3.560	3.483	3.542	3.473	3.397	3.329
Al	1.685	1.658	1.722	1.847	1.908	1.738	2.079	2.059	2.036	2.268	2.101	2.214	2.316	2.401
Fe ³⁺	0.124	0.116	0.096	0.086	0.069	0.130	0.078	0.070	0.096	0.033	0.017	0.029	0.033	0.065
Fe ²⁺	0.020	0.035	0.025	0.025	0.042	0.037	0.019	0.038	0.028	0.040	0.017	0.023	0.029	0.055
Mg	0.314	0.397	0.390	0.340	0.304	0.388	0.258	0.314	0.301	0.190	0.332	0.264	0.231	0.171
Ti	0.012	0.006	0.009	0.020	0.011	0.005	0.003	0.045	0.009	0.008	0.015	0.018	0.003	0.005
Mn	0.000	0.000	0.000	0.000	0.000	0.001	0.000	0.000	0.000	0.001	0.002	0.003	0.007	0.001
Ca	0.072	0.063	0.049	0.022	0.025	0.038	0.029	0.027	0.026	0.023	0.013	0.009	0.022	0.001
Na	0.162	0.223	0.211	0.258	0.226	0.152	0.208	0.101	0.103	0.004	0.004	0.004	0.005	0.011
K	0.008	0.148	0.222	0.235	0.292	0.388	0.353	0.481	0.516	0.621	0.686	0.702	0.769	0.802

N.B. : Fe³⁺ and Ti were assumed to be in the octahedral sheets.

A statistical analysis of the chemical compositions of the I/S minerals was performed using the Principal Component Analysis method (XLSTAT®). The main trend for the chemical dispersion is described by the eigenvector **F**₁ and corresponds to the substitution Si → KAl occurring during the transformation of montmorillonite to illite (Figure 4). A second trend, less important, seems to emerge along the **F**₂ axis. Regarding the composition of the eigenvector **F**₂, it represents the chemical dispersions for the elements Si, Mg and Al and shows a correlation between Si and Mg, whereas these latter are anti-correlated with Al. Then, the dispersion of compositions along **F**₂ axis is probably associated with the Tschermak substitution Al^{IV}Al^{VI} → Si^{IV}Mg, corresponding to muscovite → celadonite. Finally, the main chemical dispersion occurs along the montmorillonite N – illite A axis, as expected. Nevertheless, because some compositions deviate from this axis, it is also necessary to take into account a supplementary dimension, along the **F**₂ axis, in order to describe all the compositions of the I/S minerals. In the following, we propose to describe the compositions of these illite-smectite series by a ternary solid-solution, composed of illite N, montmorillonite A and interstratified F end-members. The interstratified F was selected as it exhibits the stronger celadonite-type component along **F**₂ (Figure 4).

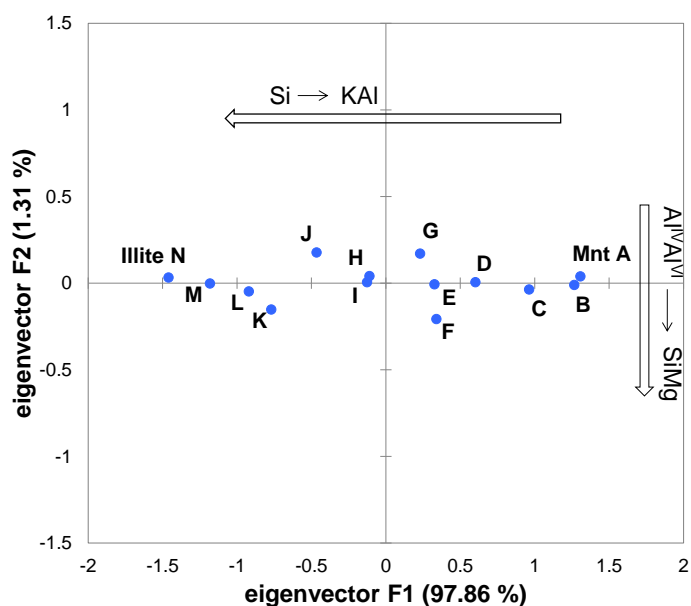


Figure 4. Statistical treatment by PCA of the 14 compositions of the I/S from Shinzan area. The chemical trends shown by arrows were depicted from the components of the main eigenvectors F_1 and F_2 (see text). Mnt A = Montmorillonite A.

2.2.3. Hydration state of samples for calorimetric measurements

The calorimetric measurements were performed on dehydrated ISCz-1 sample. The dehydrated state was obtained by heating the sample at 150°C for 20 hours. The sample was then stored in a glove box under dry Ar atmosphere ($H_2O < 4$ ppm).

For the interstratified samples from the Shinzan hydrothermal area, the enthalpies of dissolution were acquired by Perbost et al. (2002). The samples were stored at room relative humidity before the measurements. Their hydration states were not determined. Nevertheless, slight variations in the relative humidities, and therefore in the amounts of water adsorbed for a given sample, were not expected to affect significantly the enthalpy of dissolution of the sample, because the contribution of the enthalpy of dilution is negligible compared to the total measured enthalpy of dissolution and its associated uncertainty.

2.3. Calorimetric methods

2.3.1. Heat capacity measurements

For the dehydrated ISCz-1 sample, the heat capacities were measured by heat-pulse calorimetry between 2 and 330 K, using a commercial apparatus (PPMS model 6000, Quantum Design Inc.; Dachs and Bertoldi

(2005)). Measurements were performed twice in the temperature range, by cooling the sample from 50 K to 2 K and heating from 50 K to 330 K, respectively. They were carried out on two pressed pellets, weighing 4.16 and 14.93 mg respectively in order to check the reproducibility of the measurements. The relative uncertainties of the C_p values were estimated at $\pm 2\%$ for $T > 50$ K and $\pm 5\%$ for lower temperatures, according to the results from Dachs and Benisek (2011) obtained for several standard reference powder samples encapsulated in Al-pans.

Moreover, the heat capacities of the dehydrated Ca-saturated ISCz-1 sample were also measured by using differential scanning calorimetry (DSC), between 300 and 520 K. The apparatus was a Calvet DSC111 from Setaram. The amount of dehydrated sample was 140.03 mg. The experimental procedure used for the C_p^0 measurements was described in Gailhanou et al. (2012).

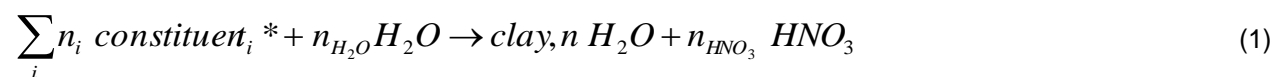
2.3.2. Acid solution calorimetry at 298.15 K

The enthalpies of dissolution of the dehydrated ISCz-1 sample and of the fourteen illite/smectite samples from the Shinzan area were measured by isothermal dissolution calorimetry at 298.15 K in HF-HNO₃ solutions. The apparatus was a highly sensitive Tian-Calvet calorimeter, the Calsol calorimeter (Ganteaume et al., 1991). The calorimeter was calibrated by dissolving THAM (tris amino-2 hydroxymethyl-2 propanediol-1,3) crystal in a solution of hydrochloric acid (0.12 M), in the same conditions as for the experiments of Ganteaume et al. (1991). The experimental procedure for the measurements was described in details in Gailhanou et al. (2012). The acid solution compositions were HF(13M)-HNO₃(2M) for the ISCz-1 sample, and HF(3M)-HNO₃(2M) for the illite/smectite samples from the Shinzan area. A higher HF concentration was necessary in the case of the ISCz-1 sample compared with the other samples, because of the presence of a quite high amount of quartz (5.85 wt.%) in the ISCz-1 sample. The compositions of the acid solutions were optimized to obtain the complete dissolution of the samples after stirring at 540 rpm during about 3 hours. For the series of I/S samples from Shinzan area, about 20 mg to 30 mg of samples were dissolved in 50 mL of acid solution. For the dehydrated ISCz-1 sample, the amount of sample was about 40 mg for 50 mL of acid solution. The weighing of this latter sample and the filling of the sample vessel were carried out in a glove box under dry argon atmosphere in order to prevent from the hydration of the sample.

2.3.3. Standard enthalpy of formation of a clay mineral from constituents at 298.15 K

The experimental method described in Gailhanou et al. (2012, 2013) was used for the determination of the enthalpy of formation of the illite/smectite in the ISCz-1 sample. It was based on a set of dissolution reactions performed in order to obtain the enthalpy of formation of the clay mineral from secondary references, i.e. constituents chosen among oxides, hydroxides and nitrates, at 298.15 K. The reactions involved were (i) the dissolution of the clay sample, containing the clay mineral and impurities, (ii) the dissolution of a mixture composed of secondary references and of impurities with the same composition of those present in the sample and (iii) dilution type reactions. The masses of constituents (except H₂O), and impurities are calculated to present the elemental stoichiometry of 40 mg of anhydrous clay sample. The impurities were added to the mixture of constituents in the same amounts than those present in the clay sample. The weighings were realized in air atmosphere for a better stability of the balance, with a maximum deviation of 10 µg. Thus, about 46 mg of a mechanical mixture of constituents and impurities was made for each dissolution experiment to accurately determine the amounts of each of these compounds present in the mixture. Four or five measurements of enthalpies of dissolution were performed on each clay sample and constituents and impurities mixtures. Relative uncertainties for enthalpy values are lower than 0.5%. For a given sample, slight variations in the final solution composition existed due to the slight variations in sample mass. However, no dependency of the enthalpy of dissolution (per unit mass) on these variations of solution composition was observed.

The reaction of formation of a clay mineral from the constituents is expressed by (Gailhanou et al., 2013):



The superscript (*) refers to all constituents except H₂O, n_i is the amount of constituent i (in mol/mol of clay), n_{H_2O} is the amount of constituent H₂O, obtained indirectly from the balance on O and H elements, n is the amount of water adsorbed in the clay, and n_{HNO_3} is the amount of HNO₃, calculated from the balance on nitrates.

Therefore, the enthalpy of formation of the clay mineral from constituents $\Delta H_{f/constit}^0$ expressed in kJ/mol $O_{10}(OH)_2$ may be calculated according to the following formula described in Gailhanou et al. (2013):

$$\Delta H_{f/constit}^0 = \left(\frac{M_{min}}{\eta} + (n - n_{H_2O}) M_{H_2O} + n_{HNO_3} M_{HNO_3} \right) \Delta H_{diss,mixt} - \left(\frac{M_{min}}{\eta} + n M_{H_2O} \right) \Delta H_{diss,sample} + n_{H_2O} M_{H_2O} \Delta H_{dil} - n_{HNO_3} M_{HNO_3} \Delta H_{sol} \quad (2)$$

where M_{min} is the molar mass of the dehydrated clay mineral, η is the mass fraction of dehydrated clay in the dehydrated sample, $\Delta H_{diss,mixt}$ is the enthalpy of dissolution of the constituents and impurities mixture (in J/g), and $\Delta H_{diss,sample}$ the enthalpy of dissolution of the clay sample (in J/g), ΔH_{dil} and ΔH_{sol} expressed in J/mol are respectively, the enthalpy of dilution of the HF-HNO₃ solution and the integral enthalpy of mixing of pure HNO₃ in HF-HNO₃ solution.

Finally, the standard enthalpy of formation ΔH_f^0 of the illite/smectite mineral was determined by considering $\Delta H_{f/constit}^0$ and the standard enthalpies of formation of the chosen secondary references.

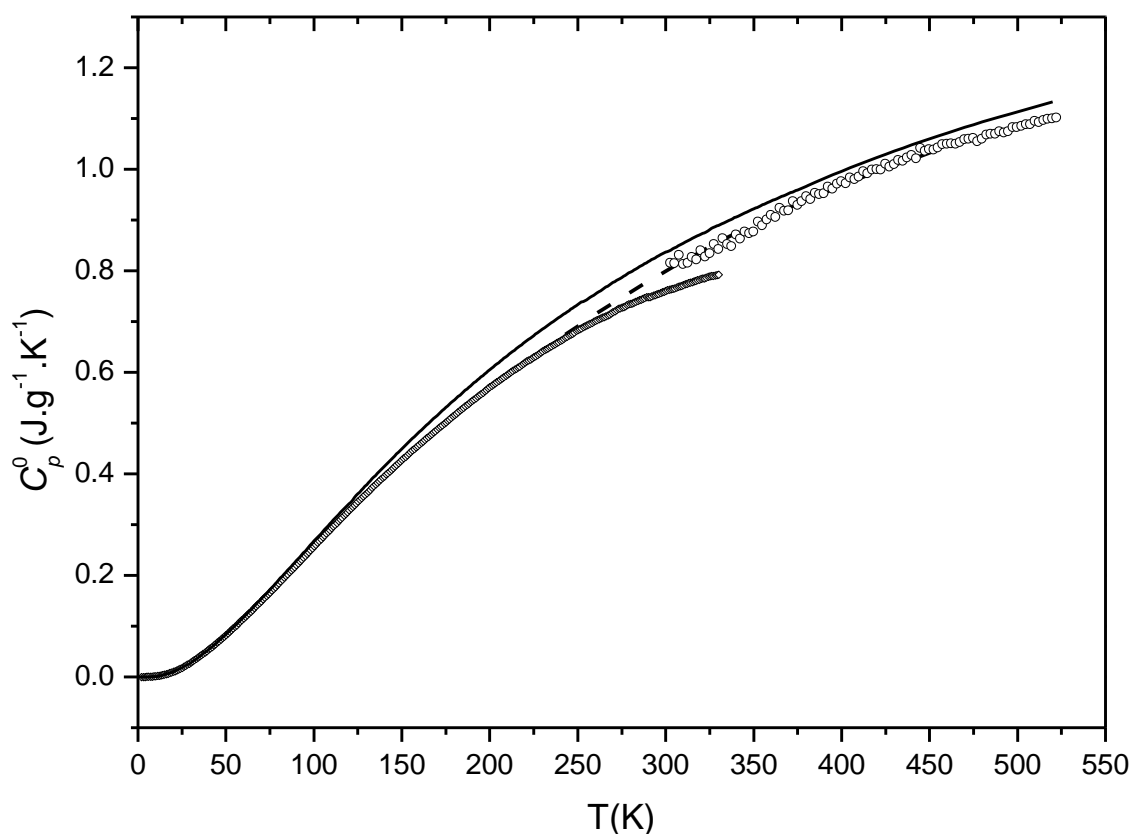
3. RESULTS

3.1. Thermodynamic properties of anhydrous illite-smectite ISCz-1

3.1.1. Heat capacities

The heat capacities measured for the Ca saturated ISCz-1 sample are plotted between 2 and 330 K for PPMS data and between 300 and 520 K for DSC data (Figure 5). All the raw data are given in Electronic Annex EA-1. A significant discrepancy between the measurements performed by both techniques appears in the common temperature range 300 – 330K. By comparison with the Cp data obtained previously for the Na-saturated ISCz-1 sample (Gailhanou et al., 2007), the Cp values measured by DSC increase similarly, whereas the Cp values from PPMS tend to drastically discard from the general trend, between about 230 K and 330 K. This comparison with the Na-saturated ISCz-1 sample reveals an inaccuracy of the Cp values measured by PPMS at the highest temperature range. This could possibly be due to the low thermal conductivity of the sample. Indeed, Kennedy et al. (2007) tested the reliability of PPMS technique on several

371 phases and noticed that the C_p data could be precise but inaccurate at high temperature ($T > 150$ K) in the
 372 particular case of poor thermal conductor phases. Therefore, the heat capacities were interpolated between
 373 200 and 520 °K using a least-square fit method and the polynomial function :
 374 $C_p(T) = 2.7070 - 3.1540 \cdot 10^{-4}T + 1.2571 \cdot 10^{-4}T^{-2} - 33.827T^{-0.5}$. Relative errors for the least square fit
 375 method are lower than 2 % at $T > 300$ K.



376
 377 Figure 5. Heat capacities measured for ISCz-1(Ca) sample (diamonds : by PPMS; white circles: by DSC; dashed line:
 378 interpolated data) and ISCz-1(Na) sample (line : data from Gailhanou et al., 2007), between 2 K and 520 K.

379
 380 The heat capacities of the clay sample are linearly interpolated at every degree between 0 and 520 °K. The
 381 heat capacities of the interstratified mineral are then calculated from the heat capacities of the sample by
 382 subtracting the contributions of the impurities (Table 4) according to the method described in Gailhanou et al.
 383 (2007). The heat capacity data for the impurities are taken from the literature (see references in EA-2).
 384 Because of the lack of thermodynamic data for the bernalite ($\text{Fe}(\text{OH})_3$) mineral, this latter was assimilated to

goethite (FeOOH), for the correction calculations. Therefore, the amount of goethite considered in the following was adjusted to 0.98 wt.%, regarding the total amount of Fe in goethite and bernalite. Besides, the heat capacity data for goethite provided by Majzlan et al. (2003) are limited to the temperature range 0 – 375 °K, due to the occurrence of the magnetic Neel transition at about 375 °K. Consequently, the correction of the heat capacities of the sample is performed between 0 and 375 °K. The heat capacities of the interstratified mineral, $C_{p,m}^0$ expressed in J.mol⁻¹.K⁻¹, are given in Table 7 for this temperature range.

3.1.2. Heat contents

The heat contents of the mineral, $H^0(T) - H^0(298.15\text{ K})$, are calculated at any temperature by integration of the $C_{p,m}^0$ function, according to

$$H^0(T) - H^0(298.15\text{ K}) = \int_{298.15}^T C_{p,m}^0 dT \quad (3)$$

using the trapezoid method at every degree (Table 7).

3.1.3. Entropy

The third-law entropy of the interstratified mineral, S^0 , is expressed by

$$S^0 = \int_0^T \frac{C_{p,m}^0}{T} dT + S^{0(resid.)} \quad (4)$$

with $S^{0(cal)} = \int_0^T \frac{C_{p,m}^0}{T} dT$ the calorimetric entropy obtained directly from the heat capacity measurements, and $S^{0(resid.)}$, the residual entropy. This latter term is not available from the calorimetric measurements and requires additional structural information about the mineral (Ulbrich and Waldbaum, 1976). The chemical site configurational entropy, $S^{0(conf)}$, associated with cationic disorder in the tetrahedral, octahedral and interlayer crystallographic sites, is a significant part of the residual entropy. It is assessed according to the method described in Blanc et al. (2015). The residual entropy may also include a magnetic term $S^{0(mag)}$ associated with the disordering of magnetic spins of transition metals in the clay minerals, occurring at very low temperatures (e.g. $T < 10\text{ K}$ for iron-bearing chlorites, Townsend et al., 1986). Indeed, for clay minerals, recent studies (Gailhanou et al., 2009) have demonstrated that the magnetic contribution is difficult to detect

and to measure directly. A maximum value of the magnetic entropy is assessed by using the method described by Ulbrich and Waldbaum (1976), based on the maximum number of spin configurations according to:

$$S^{0(mag)} = R \sum_i x_i \ln(2S_i + 1) \quad (5)$$

where R is the gas constant, x_i the amount of element i and S_i its spin number (for Fe^{3+} , $S = 5/2$; for Fe^{2+} , $S = 2$). Finally, in the case of the illite-smectite ISCz-1 mineral, the configurational entropy $S^{0(conf)}$ is estimated at $23.19 \text{ J.mol}^{-1}.\text{K}^{-1}$ and the maximum magnetic entropy, at $0.65 \text{ J.mol}^{-1}.\text{K}^{-1}$. The calorimetric entropy data are given in Table 7.

Table 7. Molar heat capacities and derived thermodynamic functions of the dehydrated Ca-saturated illite-smectite ISCz-1 mineral (corrected for impurities) at selected temperatures.

T	$C_{p,m}^0(T)$	$S^{(0)cal}$	$H^0(T) - H^0(298.15)$
K	$\text{J.mol}^{-1}.\text{K}^{-1}$	$\text{J.mol}^{-1}.\text{K}^{-1}$	kJ.mol^{-1}
0	0.00	0.00	-45.596
10	0.65	0.42	-45.594
20	3.81	1.65	-45.574
30	10.61	4.37	-45.505
40	19.85	8.64	-45.354
50	30.53	14.20	-45.103
60	43.01	20.85	-44.736
70	55.96	28.44	-44.242
80	69.39	36.78	-43.616
90	83.08	45.75	-42.853
100	97.33	55.24	-41.951
110	111.25	65.17	-40.908
120	124.93	75.43	-39.728
130	138.00	85.95	-38.413
140	150.81	96.66	-36.968
150	163.30	107.49	-35.397
160	175.05	118.41	-33.704
170	186.45	129.37	-31.896
180	197.55	140.34	-29.976
190	208.35	151.31	-27.946
200	216.66	162.25	-25.812
210	226.42	173.06	-23.597
220	236.19	183.82	-21.284
230	245.88	194.53	-18.873
240	255.41	205.20	-16.367
250	264.72	215.81	-13.766
260	273.81	226.38	-11.073
270	282.64	236.88	-8.291
280	291.20	247.31	-5.421
290	299.49	257.67	-2.468
298.15	306.05	266.06	0.000
300	307.51	267.96	0.568
310	315.23	278.17	3.682

320	322.66	288.30	6.871
330	329.83	298.34	10.134
340	336.73	308.29	13.467
350	343.37	318.14	16.868
360	349.78	327.91	20.334
370	355.95	337.58	23.862
375	358.95	342.37	25.650

3.1.4. Standard enthalpy of formation

Enthalpies of dissolution of the Ca-saturated ISCz-1 sample and of the constituents* mixture (superscript * means: except H₂O constituent) are given in Table 8. Uncertainties associated with the mean enthalpies are calculated from measured values using Student's *t*-distribution with a 95 percent-confidence interval.

Table 8. Enthalpies of dissolution of the dehydrated ISCz-1 sample and of the corresponding constituents and impurities mixtures

		Mass (mg)	ΔH_{diss} (J/g)	Mean ΔH_{diss} (J/g)
ISCz-1 in HF(13M)- HNO ₃ (2M)	Dehydrated sample ISCz-1	39.86	-2256.60	-2257.82 (± 1.19)
		40.01	-2257.62	
		40.00	-2258.31	
		40.20	-2258.74	
	Mixture of constituents* + impurities	46.58	-1845.16	-1848.08 (± 3.28)
		46.55	-1849.41	
		46.55	-1850.65	
		46.57	-1845.16	
		46.55	-1850.00	
Note: Uncertainties between brackets are calculated using Student's <i>t</i> -distribution with a 95% confidence interval				

The enthalpy of dilution ΔH_{dil} and the solution enthalpy ΔH_{sol} associated with the mixing of pure HNO₃ in HF(13M) - HNO₃(2M) were determined in Gailhanou et al. (2012). Their values were respectively - 1.57 (± 0.30) kJ.mol⁻¹ H₂O and -27.7 (± 26) kJ.mol⁻¹ HNO₃.

Afterwards, the enthalpy of formation of the mineral from constituents was then calculated using equation (2) (Table 9).

Table 9. Reaction of formation of illite-smectite ISCz-1 from the secondary reference constituents and associated enthalpy $\Delta H_{f/constit.}^{\circ}$ in kJ.mol⁻¹ (at 1 bar and 298.15 K).

Reactions of formation of mixed-layer mineral	$\Delta H_{f/constit.}^{\circ}$ (kJ.mol ⁻¹)
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Illite/smectite ISCz-1 (n = 0)	3.562 SiO _{2,quartz} + 2.159 Al(OH) _{3,gibbsite} + 0.255 Mg(OH) ₂ + 0.029 FeOOH _{goethite} + 0.011 FeO + 0.008 TiO _{2,rutile} + 0.440 KNO ₃ + 0.092 Ca(OH) ₂ = (Ca _{0.092} K _{0.440})(Si _{3.562} Al _{0.438})(Al _{1.721} Mg _{0.255} Fe _{0.029} ³⁺ Fe _{0.011} ²⁺ Ti _{0.008})O ₁₀ (OH) ₂ +0.440 HNO ₃ +2.380 H ₂ O	62.36 (±3.49)
Notes: $M_{min}(ISCz-1) = 382.007 \text{ g.mol}^{-1}$; $\eta(ISCz-1) = 0.88456$; $M_{H_2O} = 18.02 \text{ g.mol}^{-1}$; $M_{HNO_3} = 63.01 \text{ g.mol}^{-1}$; $n_{H_2O} = -2.380$; $n = 0$; $n_{HNO_3} = 0.440$. Standard uncertainties given between brackets correspond to one standard deviation of the means.		

Eventually, the standard enthalpy of formation of the illite-smectite ΔH_f^0 was obtained from the enthalpy of formation of the mineral from the secondary reference constituents and from the standard enthalpies of formation of these reference constituents (Table 10). These latter values are selected consistently with the Thermochimie database (Giffaut et al., 2014), at 298.15 K, and are tabulated in Electronic Annex EA-3.

The uncertainty associated with ΔH_f^0 was standard deviation of the mean, calculated using the error propagation method (Ellison et al., 2000):

$$\Delta(\Delta H_f^0) = \sqrt{[\Delta(\Delta H_{f/constit.}^0)]^2 + \sum_i \nu_i^2 [\Delta(\Delta H_f^{constit.i})]^2} \quad (3)$$

Where ν_i is the stoichiometric coefficient associated with the constituent i , and $\Delta H_f^{constit.i}$ is the standard enthalpy of formation of the constituent i . The uncertainties associated with $\Delta H_{f/constit.}^0$ and $\Delta H_f^{constit.i}$ correspond to one standard deviation of the mean.

3.1.5. Standard Gibbs free energy of formation

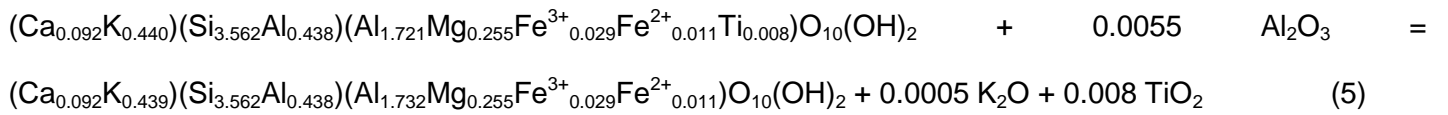
The standard Gibbs free energy of formation ΔG_f^0 is expressed by:

$$\Delta G_f^0(T) = \Delta H_f^0(T) - T\Delta S_f^0(T) \quad (4)$$

where $\Delta S_f^0(T)$ is the standard entropy of formation of the interstratified mineral calculated considering the reaction of formation of the mineral from the elements in their stable thermodynamic form in standard conditions. The value for the ISCz-1 mineral was obtained at 298.15 K from the third-law entropy of the mineral S^0 (Table 10) and from the entropies of the elements at 298.15 K. These latter data were taken from Cox et al. (1989) for Si_(cr), Al_(cr), Mg_(cr), Ti_(cr), K_(cr), Ca_(cr), O_{2(g)}, H_{2(g)} and from Chase (1998) for Fe_(cr).

3.2. Synthesis of the thermodynamic properties for anhydrous illite-smectite ISCz-1

The complete set of thermodynamic properties of the interstratified ISCz-1 mineral at 1 bar and 298.15 K is given in Table 10. Since its structural formula contains a small amount of TiO₂, a simplified structural formula is also proposed in order to ease the introduction of the phase into geochemical databases. The simplification is based on fictive solid-solid reactions with properties of reaction equal to 0. In the case of the interstratified ISCz-1, the following fictive reaction was considered :



The thermodynamic data for alumina- α , rutile and K₂O are taken from the Thermoddem database (<http://thermoddem.brgm.fr>, Blanc et al. (2012)).

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475 Table 10. Thermodynamic properties of the dehydrated illite-smectite ISCz-1 mineral, at 1 bar and 298.15 K.

T = 298.15 K									A	B*10 ⁻³	C*10 ⁻⁵	T range for C _p functions
Mineral	ΔG_f^0 (kJ mol ⁻¹)	ΔH_f^0 (kJ mol ⁻¹)	$S^{(calc)}$ (J mol ⁻¹ K ⁻¹)	$S^{(conf)}$ (J mol ⁻¹ K ⁻¹)	$S^{(mag)}$ (J mol ⁻¹ K ⁻¹)	S^0 (J mol ⁻¹ K ⁻¹)	V^0 (cm ³ mol ⁻¹)	C _p (298.15 K) (J mol ⁻¹ K ⁻¹)	J/(mol K)	J/(mol K ²)	(J K /mol)	
Illite/smectite ISCz-1	-5417.38 (± 7.60)	-5785.74 (± 7.46)	266.06	29.53	0.58	296.17	137.13	306.05	228.34	438.31	-47.04	298 – 375 K
Properties recalculated for a simplified structural formula*												
Illite/smectite ISCz-1*	-5418.62 (± 7.60)	-5787.22 (± 7.46)	265.89	28.89	0.58	295.36	137.13	306.00	236.14	422.80	-49.92	298 – 375 K

Notes: Between 298 and 375 K, Cp(T) functions are expressed according to $C_p(T) = A + B \cdot 10^{-3} T + C \cdot 10^{-5} T^2$, where A, B and C are Maier-Kelley coefficients, obtained by fitting the Cp curves.

Standard uncertainties associated with ΔH_f^0 and ΔG_f^0 correspond to one standard deviation of the means (see text for details).

[⊥] Molar volume V^0 is obtained from values of unit-cell parameters a, b and c determined from XRD patterns, according to the method described in Gailhanou et al. (2012): for the present dehydrated ISCz-1 sample, $a = 5.20 \text{ \AA}$, $b = 9.01 \text{ \AA}$ and $c = 10.12 \text{ \AA}$.

[‡] Simplified formula is introduced to ease the predictive calculations and their implementation into databases. The simplification process is detailed in the text.

Illite/smectite ISCz-1: (Ca_{0.092}K_{0.440})(Si_{3.562}Al_{0.438})(Al_{1.721}Mg_{0.255}Fe³⁺_{0.029}Fe²⁺_{0.011}Ti_{0.008})O₁₀(OH)₂

Illite/smectite ISCz-1*: (Ca_{0.092}K_{0.439})(Si_{3.562}Al_{0.438})(Al_{1.732}Mg_{0.255}Fe³⁺_{0.029}Fe²⁺_{0.011})O₁₀(OH)₂

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3.3. Enthalpies of mixing for mixed-layers from Shinzan area samples

The analysis of the chemical dispersions among the illite-smectite compositions (Figure 4) shows that all of these one can be described by a ternary solid solution, with illite N, montmorillonite A and illite-smectite F end members. Although the chemical dispersions occur mostly along the illite – smectite join, the ternary solid solution is preferred to the binary illite-smectite solid solution for accuracy reasons. Since the enthalpies of mixing are expected to be very low, and consequently difficult to determine accurately regarding the associated uncertainties, it is preferable to perform the measurements for all the samples without adding a supplementary source of uncertainty induced by the consideration of the binary solid solution. In the following, the enthalpies of mixing for illite-smectite compositions from the Shinzan area samples are determined by using isothermal dissolution calorimetry at 298.15 K.

3.3.1. Enthalpies of dissolution at 298.15 K

The enthalpies of dissolution of all the samples in HF(3M)-HNO₃(2M) at 25 °C were measured by Perbost et al. (2002). The data are given in

Table 11. Uncertainties associated with the mean enthalpies are calculated from measured values using Student's *t*-distribution with a 95 percent-confidence interval. The relative uncertainties are lower than 1%, except for the sample E, at 2.4%.

Table 11. Mean values of the enthalpies of dissolution of the I/S samples from Shinzan area, measured by Perbost et al. (2002) in HF(3M)-HNO3(2M) solution at 25 °C. Mnt A = Montmorillonite A. Uncertainties calculated using Student's t -distribution with a 95 percent-confidence interval.

Sample	Mean value of ΔH_{diss} (kJ.mol ⁻¹)
Mnt A	-846.80 (± 7.64)
B	-844.03 (± 1.45)
C	-859.94
D	-847.84 (± 2.95)
E	-842.04 (± 20.2)
F	-840.77 (± 2.52)
G	-859.50
H	-848.02 (± 6.14)
I	-860.21
J	-866.77
K	-854.33
L	-858.14
M	-863.52
Illite N	-863.57 (± 6.39)

The opposite values of the enthalpies of dissolution are plotted in Figure 6a, in the 3-D space (\mathbf{F}_1 , \mathbf{F}_2 , $-\Delta H_{\text{diss}}$). The values were then modelled by a polynomial surface, by using the least-square fit method. The polynomial function is expressed by:

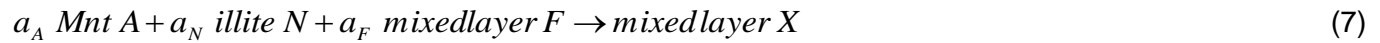
$$P(z_1, z_2) = 852.124 - 6.029 z_1 + 41.159 z_2 + 1.938 z_1^2 - 4.027 z_1 z_2 + 45.824 z_2^2 \quad (6)$$

where z_1 and z_2 correspond to the coordinates of the I/S samples in the ($\mathbf{F}_1, \mathbf{F}_2$) plane.

The discrepancies between the measured and modelled values are lower than 1%, which corresponds to the relative uncertainties for the measured enthalpies of dissolution. The modelled values are then considered to represent correctly the measured enthalpies.

3.3.2. Enthalpies of mixing at 298.15 K

The illite-smectite compositions may be expressed by a linear combination of the compositions of the three end-members A, F and N. The barycentric coordinates (a_A, a_F, a_N) for all the I/S samples are given in Table 12. With respect to these end-members, the enthalpy of mixing for an interstratified X from the Shinzan series is the enthalpy of the reaction of formation:



which practically represents the enthalpy of mixing of illite and smectite (montmorillonite) layers. The enthalpy of mixing is then obtained from the combination of the reactions of dissolution of the minerals and is expressed by:

$$\Delta H_m(\text{mixedlayer X}) = a_A \Delta H_{\text{diss}}(A) + a_N \Delta H_{\text{diss}}(N) + a_F \Delta H_{\text{diss}}(F) - \Delta H_{\text{diss}}(\text{mixedlayer X}) + \Delta H_{\text{mix},\text{sol}} \quad (8)$$

where $\Delta H_{\text{mix},\text{sol}}$ is the enthalpy of mixing of solutions resulting from the dissolution reactions of the minerals, which is neglected in the following calculations. The enthalpies of mixing are then calculated for all the illite-smectite compositions, from the modelled values of the enthalpies of dissolution (

Table 12), and are plotted in the ($F_1, F_2, \Delta H_m$) space (Figure 6b).

Table 12. Enthalpies of mixing of the I/S minerals from the Shinzan area, with respect to the three end-members Mnt A, illite N and interstratified F. a_A , a_N and a_F are the barycentric coordinates of the I/S minerals, associated with the corresponding end-members.

Samples	a_A	a_N	a_F	ΔH_m (kJ.mol ⁻¹)
Mnt A	1.00	0.00	0.00	0.00
B	0.85	-0.06	0.20	0.20
C	0.68	0.02	0.30	-0.99

D	0.66	0.21	0.13	-2.64
E	0.53	0.29	0.18	-3.18
F	0.00	0.00	1.00	0.00
G	0.97	0.58	-0.55	-2.98
H	0.50	0.52	-0.02	-3.70
I	0.40	0.47	0.12	-3.62
J	0.74	0.85	-0.59	-2.41
K	-0.25	0.48	0.77	-1.47
L	-0.02	0.69	0.33	-2.19
M	0.01	0.85	0.14	-1.39
Illite N	0.00	1.00	0.00	0.00

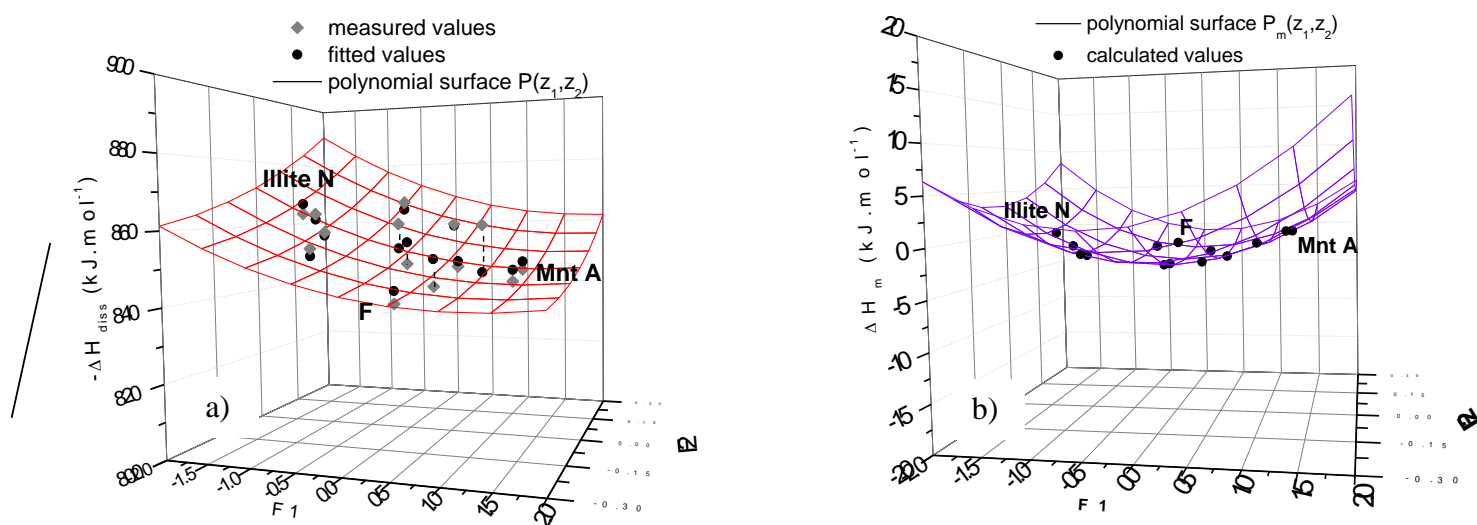


Figure 6. Representations of (a) the polynomial surface $P(z_1, z_2)$ obtained by fitting the experimental values of $-\Delta H_{\text{diss}}$ for the I/S samples and (b) the enthalpies of mixing of the illite and smectite layers (per mol $\text{O}_{10}(\text{OH})_2$) for the I/S minerals and the modelling polynomial surface $P_m(z_1, z_2)$.

The resulting enthalpies of mixing are slightly negative, with a minimum value at about -3.7 kJ.mol^{-1} for the H and I samples. According to XRD modelling (Inoue and Utada, 1983), these latter are first order R1 type and should contain about 30-35 % of smectite layers. A model predicting ΔH_m was proposed by Blanc et al. (1997), resulting in $\Delta H_m = -3.6 \text{ kJ.mol}^{-1}$ for illite/smectite displaying such composition and ordering types.

4. DISCUSSION: comparison with estimates and stability domains

The standard thermodynamic properties of the illite-smectite ISCz-1 obtained experimentally are compared with estimated values calculated from Blanc et al. (2015) model that includes a consistent set of models for predicting the standard thermodynamic properties (H, S, Cp, V) for phyllosilicates. First, the model from Blanc et al. (2015) is merely applied by considering the whole structural formula of the illite-smectite. The simplified structural formula, $(\text{Ca}_{0.092}\text{K}_{0.439})(\text{Si}_{3.562}\text{Al}_{0.438})(\text{Al}_{1.732}\text{Mg}_{0.255}\text{Fe}^{3+}_{0.029}\text{Fe}^{2+}_{0.011})\text{O}_{10}(\text{OH})_2$, is considered in order to ease the predictive calculations. Results are given in Table 13. Besides, a second approach is carried out, predicting separately the thermodynamic parameters of illite and smectite components and combining them according to the required composition of illite-smectite. This approach aims to differentiate the chemistry of the illite and smectite layers in the interstratified, consistently with the structure of the mineral illite-smectite.

The method for calculating the chemical compositions of illite and smectite end-members relies on the work of Meunier and Velde (1989), based on the graphical representation of $\text{M}^+-4\text{Si}-\text{R}^{2+}$ ternary system for natural sample compositions. Plotting ISCz-1 coordinates in such a diagram (Figure 7) reveals that illite-smectite ISCz-1 can be considered as a binary solid solution with a high-charge illite ($0.87/\text{O}_{10}(\text{OH})_2$), namely IllitePP, and a low-charge montmorillonite (0.21), namely MontmorillonitePP, end-members. Meunier and Velde (1989) have proposed relations between illite and montmorillonite theoretical end-members. The compositions of the end-members are deduced from their coordinates in the $\text{M}^+-4\text{Si}-\text{R}^{2+}$ triangle, which provide chemical constraints, and from a calculation based on a least-square method to adjust the proportions of illite and smectite components. These latter were determined considering 70% of illite, consistently with the proportions obtained from XRD. Finally, the thermodynamic properties of the illite and smectite end-members are calculated by using the model from Blanc et al. (2015) for each end-members, then considering a linear combination of both. The set of data is completed by considering the mixing enthalpy measured for the I/S sample with 70% of illite layers (sample I, Table 5), -3.6 kJ.mol^{-1} .

Regarding the results (reported in Table 13) , the prediction obtained by considering the mixture of illite and smectite components falls closest to the experimental values except for heat capacity and volume. For these latter properties, both estimate method provide similar results and, on the whole, the method based on a mixture of layers is slightly improving the accuracy of the predictions for the properties of illite-smectite ISCz-1 mineral.

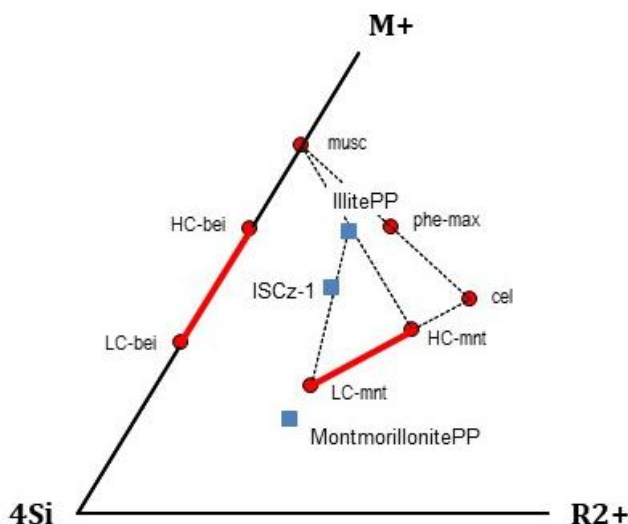


Figure 7. Composition of clay minerals in the $M^+-4Si-R^{2+}$ system (M^+ = layer charge of ideal mica, $4Si$ = Si content of the tetrahedral sheet, R^{2+} = amount of bivalent cations in the octahedral position). Musc = Muscovite, phe-max = Phengite, cel = celadonite, IllitePP = illite end-member (layer charge 0.87), HC-mnt = high charge montmorillonite, LC-mnt = low charge montmorillonite, beid = beidellite. IllitePP and MontmorillonitePP are the predicted compositions of the illite and smectite components in the mixed-layer ISCz-1.

Table 13. Estimated thermodynamic properties of ISCz-1, at 1 bar and 298.15K, according to the model from Blanc et al. (2015). Data are compared with the experimental values obtained for the simplified structural formula, given in Table 11.

Predicted	ΔG_f^0 kJ.mol ⁻¹	ΔH_f^0 kJ.mol ⁻¹	$S^{0(tot)}$ J.mol ⁻¹ .K ⁻¹	$C_p^0(298.15\text{ K})$ J.mol ⁻¹ .K ⁻¹	V cm ³ .mol ⁻¹
Mean composition	-5412.63	-5779.31	301.78	317.63	137.94
$\Delta(\text{exp-pred})$	-5.99	-7.91	-6.41	-11.58	-0.81
$\Delta(\%)$	-0.11	-0.13	-2.17	-3.78	-0.59
Mix illite/smectite layers*	-5415.19	-5785.52	290.03	317.79	138.00
Mixing terms	-3.6	-3.6			
Final estimate	-5418.89	-5789.22	290.03	317.79	138.00
$\Delta(\text{exp-pred})$	0.17	1.90	5.33	-11.74	-0.87
$\Delta(\%)$	0.00	0.03	1.80	-3.80	-0.63
*illite component : $K_{0.683}Ca_{0.084}(Si_{3.319}Al_{0.681})(Al_{1.777}Mg_{0.234}Fe^{2+}_{0.016})O_{10}(OH)_2$					
smectite component : $Ca_{0.107}Si_4(Al_{1.652}Mg_{0.292}Fe^{3+}_{0.081}Fe^{2+}_{0.003})O_{10}(OH)_2$					

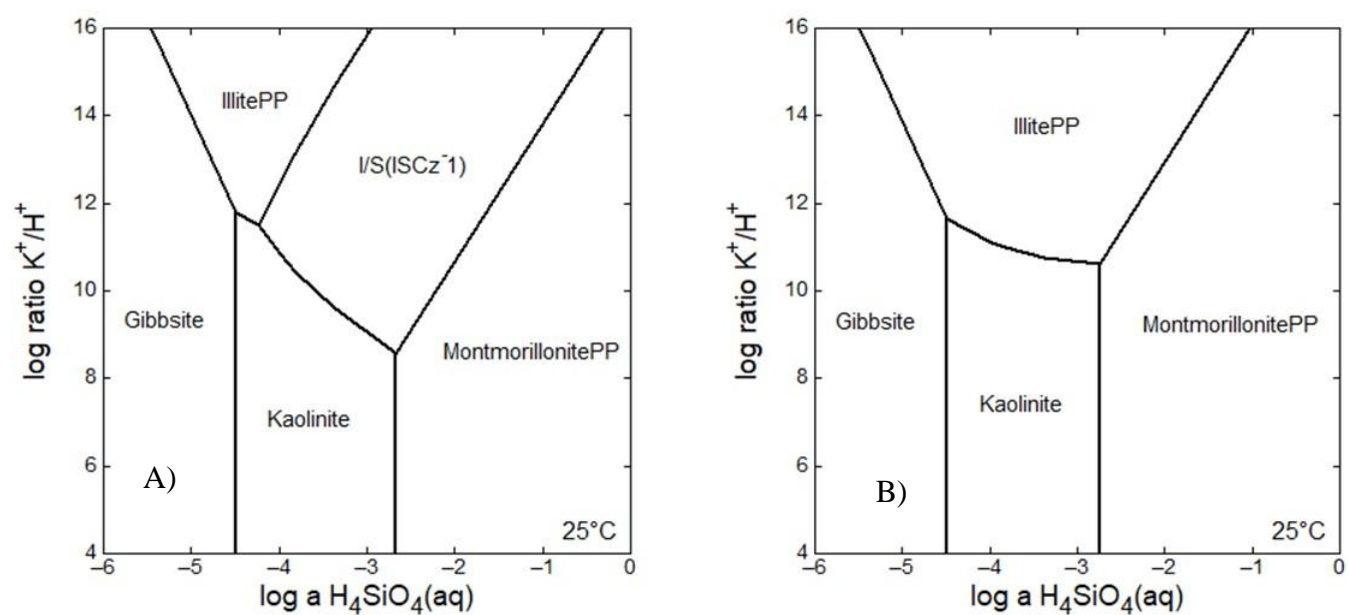
As stated by Blanc et al. (1997), the mixing of layers introduces an additional entropy term, which is not measurable directly by calorimetry, for instance. Blanc et al. (1997) have proposed a predicting model based on nearest neighbor interaction up to the fourth neighbor. In a simpler approach, we can first state for an ideal solid solution between illite and smectite end members, which would result, for ISCz-1 with 70 % illite, in $\Delta S_m = 5.38 \text{ J.mol}^{-1}.\text{K}^{-1}$.

In Figure 8, the stability fields for ISCz-1, Illite and Montmorillonite end-members are drawn at 25°C, considering two different sets of thermodynamic parameters for ISCz-1:

- a set including ΔH_m for ISCz-1 properties of formation (Figure 8.A)
- a set excluding ΔH_m for ISCz-1 properties of formation (Figure 8.B).

Comparing Fig. 8.A and Fig. 8.B enlighten the importance of mixing terms. Indeed, ISCz-1 stability field just disappears when the mixing terms are not integrated to the thermodynamics function, whereas it lies as expected between illite and montmorillonite end members when the mixing terms are added to the ISCz-1 formation properties. On the whole, in the calculation process followed here to predict the thermodynamic properties of illite/smectite, we consider a mixture of layers, with the composition of the montmorillonite component being itself ranging between high and low charge montmorillonite, consistently with Meunier and Velde (1989). As can be seen from Fig. 8 a simple mixture is not sufficient and additional and mixing terms are necessary to result in a significant stability, ranging as expected between illite and montmorillonite end members.

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Figure 8 – Stability fields for I/S ISCz-1 mineral: A) including enthalpy of mixing and B) without enthalpy of mixing.

5. SUMMARY AND CONCLUSIONS

In early diagenesis and underground storage studies, the transformation of smectite into illite has been recognized for long to involve the formation of interstratified illite/smectite as intermediate product (Srodon and Eberl, 1984, Meunier and Velde, 1989). However the lack of thermodynamic properties prevented it to be implemented into geochemical calculations, so far. This study presents the first calorimetric direct measurements of such properties.

The measurements were carried out onto a purified, Ca-saturated ISCz-1 sample from Slovakia, provided by the Source Clays Repository of the Clay Minerals Society. ΔH°_f was obtained from acid solution calorimetry experiments, S° and C_p were measured by PPMS and DSC techniques. In addition, a specific mixing enthalpy was measured by acid solution calorimetry, considering samples from the Shinzan hydrothermal series (Inoue and Utada, 1983). For the whole set of measured thermodynamic properties, the comparison with previously published estimation method (Blanc et al., 2015) was favorable, provided the estimate is calculated distinguishing the respective contribution of illite and smectite layers. The respective compositions of these latter are derived from composition relations established by Meunier and Velde (1989).

Among the concluding points, comparing the different estimating methods brings out that the energetic terms for layer mixing are necessary for the mineral stability, even though their magnitude is small with respect to formation terms, i.e. even falling within the uncertainty obtained for the formation terms. This implies that further improvements and advances could be made by considering weak energies terms like hydration or exchange reaction contributions (Gailhanou et al., 2017).

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