

NANOPOROUS MATERIALS: PILLARED CLAYS AND REGULAR SILICAS AS AN EXAMPLE OF SYNTHESIS AND THEIR POROSITY CHARACTERIZATION BY X-RAY DIFFRACTION

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STUDENT'S INFORMATION

General Remarks

An experimental procedure to the synthesis of Zr-PILC and MCM-41 samples is given in the following, where reagents and material are described also. A proposal for time managing is presented.

Experimental Instructions

Zr-PILC synthesis

Two grams of a montmorillonite rich clay (for instance Bentonite CAS-No-1302-78-9) in 200 mL of water are stirred for a minimum of 2 hours, but preferably overnight. 50 mL of an aqueous 0.1 mol/L Zirconium(IV) oxychloride octahydrate ($\text{ZrOCl}_2 \cdot 8\text{H}_2\text{O}$; CAS-No-13520-92-8) solution are then dropwise added to the clay dispersion (an isobaric dropping funnel can be used) under strong stirring. The mixture is then aged for 2 hours at 40 °C without stirring but under reflux, after which the solid phase is separated by centrifugation. Remove a small amount of solid for X-ray (the "Before washing" sample). Insert the solid in a dialysis tubing (example: cellulose membrane for retaining proteins of molecular weight 12000 or greater) that was previously boiled, close both sides of the tube and insert it in a container with distilled water (at least 500 mL). Keep in water preferably during a week and, occasionally, change with new water. After this time open the dialysis tube, remove the slurry and evaporate the water (use a rotary evaporator, but do not set the temperature above 60 °C). Remove a small amount of solid for X-ray (the "After washing" sample). Dry at 90-100 °C, put the solid in an alumina boat and calcined at 350 °C for 2-3 hours preferably in a open tubular oven or other type of oven that allows some degree of air to enter in it.

MCM-41 synthesis

Weight 2.642g of hexadecyltrimethylammonium bromide - CTAB (CAS-No-57-09-0) and add 132 mL of distilled water. Stir (magnetic stirring) under gentle heating until the solution becomes colourless (approximately 30 minutes). Add 11.2 mL of ammonium hydroxide solution (25% w/w), keep stirring for 2-3 minutes, add 11.4 mL of tetraethyl orthosilicate – TEOS (CAS-No-78-10-4) and maintain the stirring for 2.5 hours. Filter, wash with water preferably until the pH becomes neutral (use vacuum filtration with Büchner funnel), and dry at 90 °C until next laboratory session. Keep at least an adequate portion of the uncalcined form to obtain the X-ray diffractogram, and put the remaining solid in an alumina boat and calcined at 550 °C for 5 hours preferably in a open tubular oven or other type of oven that allows some degree of air to enter in it. For the synthesis with the mixture of hexadecyltrimethylammonium and tributylheptylammonium bromide (CAS-No-85169-31-9) the same methodology is to be used but mix 1.984 g of the former and 0.657 g of the latter.

Hazards

Tetraethyl orthosilicate – TEOS (CAS-No-78-10-4) is flammable, may cause irritation to eyes and skin and its long-term exposure may cause kidney or liver damage.

Ammonium hydroxide solution (CAS-No-1336-21-6) is corrosive, causes severe skin burns and eye damage.

Hexadecyltrimethylammonium bromide (CAS-No-57-09-0) presents acute toxicity, it causes oral skin irritation and serious eye damage and it may cause respiratory irritation.

Zirconium(IV) oxychloride octahydrate (CAS-No-13520-92-8) may causes severe skin burns and eye damage.

Tributylheptaammonium bromide (CAS-No-85169-31-9) is irritant to eyes and respiratory system and skin.

Clays (for instance Bentonite CAS-No-1302-78-9) are hazardous in case of eye contact (irritant), and of inhalation. They are slightly hazardous in case of skin contact (irritant), and of ingestion. Avoid dust formation.

Always wear suitable protective clothing, gloves and eye/face protection.

Environmental regulations should be followed for the disposal of waste.

The calcination step needs special attention and students should be advised not to remove the sample from the furnace before it cools to a temperature near ambient.

Characterization by X-ray diffraction

X-ray diffractograms are to be obtained between 2 θ degrees of 2 and 10, since this is the region where the signals for the d_{001} appear for clays^{S1,S2} and where the relevant information for MCM-41 is obtained also. In this way the X-ray apparatus can be programmed with the appropriated time per step so each diffractogram will take less than 20 minutes to obtain. Typical (for a Philips XRD model PW3710) time per step and step size are 2.5 s and 0.02 degrees, respectively.

X-ray diffractograms for the MCM-41 samples can be obtained by the usual powder method, just by depositing it in the appropriated support and compacting gently the powder. The same for the initial clay if the presence of other minerals is to be studied, cf. Figure S5, but, in this case the 2 θ degrees range to be used is from 3 to 50. For the Zr-PILC samples, and to have more accuracy in the d_{001} values, oriented mounts should be used. For this a slurry (clay plus water) is to be prepared by initially dispersing the clay in water by stirring or, preferably, by using a ultrasonic bath so the clay particles will be more efficiently dissociated. Then the slurry is concentrated, for instance by centrifugation, deposit in a glass slide (that fits the X-ray in the apparatus) and let to dry at ambient or in a oven at moderate temperature (below 50 °C). As alternative to the centrifugation, the slurry may settle and a small portion can be transferred with a Pasteur pipette to the glass slide and let to dry. Repeat the transfer/dry two or three times. Making oriented mounts is detailed described in the reference S1.

TIME MANAGEMENT

Laboratory sessions of 3 to 4 hours were considered.

Zr-PILC

First session: the theme is presented to the students, and they get acquainted with related bibliography but they also need to prepare the 0.1 M $\text{ZrOCl}_2 \cdot 8\text{H}_2\text{O}$ solution and the clay dispersion in water. Preferable, the X-ray diffractogram of the initial material should be obtained also.

Second session: the pillaring process is to be made and the pillared material is left washing in the dialysis tube until the next session (preferably a week latter). Glass slide for XRD of the “Before washing” sample can be prepared

Third session: calcination of the intercalated material. X-ray diffractograms of previous samples and preparation of the glass slide of the calcined material.

If only three working sessions are to be used, the X-ray diffractogram of the calcined sample may be obtained in student free time or the instructor may provide it since the students already made the experiment with other samples. If a fourth session is available the X-ray diffractogram of the calcined sample can then be obtained and the remaining time used to analyse the results and make a previous discussion of them with the instructor.

MCM-41

First session: presentation of the theme to the students, and get acquainted with related bibliography but, preferably, they should start the synthesis of the sample prepared with CTAB and use the 2.5 hours of stirring time to start search and reading literature. Non-calcined sample is obtained.

Second session: calcination of the material from the first synthesis and making the second synthesis (with the second template). Since the calcination will need 5 hours, plus some time to become to a temperature where it may be removed from the oven, there are mainly two options. The students (with the instructor) may use some free time before the class just to put the alumina boat in the oven and start the calcination, so it will be ready by the end of the session or the calcination would need to be started by the instructor. During the 2.5 hours of stirring time, students may progress in the literature study or they can obtain the X-ray diffractogram of the non-calcined sample from the previous session.

Third session: calcination of the material from the second synthesis. X-ray diffractograms (calcined sample from the first synthesis

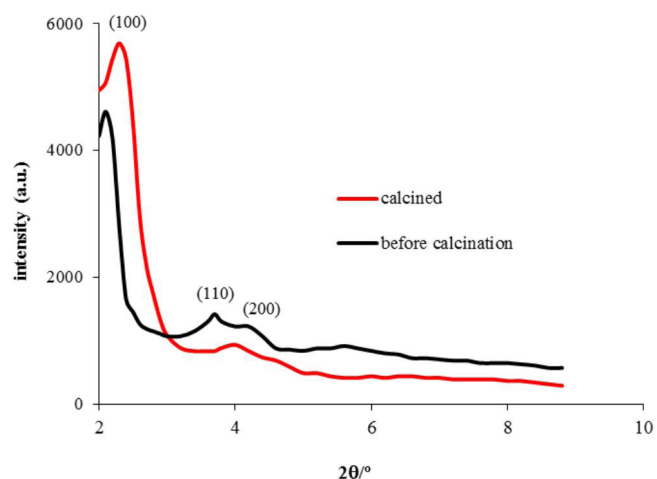


Figure 1S. X-ray diffractogram for the MCM-41 sample, prepared with cetyltrimethylammonium bromide, before and after calcination

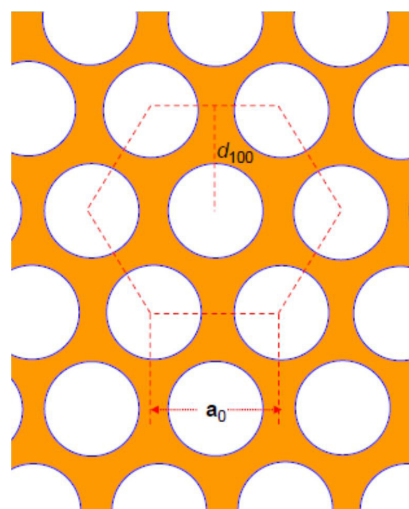


Figure 2S. Structure of the MCM-41 material (a transversal plan) with its hexagonal array of pores and the corresponding a_0 cell parameter

and non calcined sample from the second synthesis). Analysis of the diffractogram to obtain the cell parameter.

If only three working sessions are to be used the last X-ray diffractogram of the calcined sample from the second synthesis maybe obtained in student free time or the instructor may provide it since the students already made the experiment with other samples. If a fourth session is available the X-ray data can analysed and IR spectroscopy (in KBr wafers) can be used, at least for the samples before and after calcination of one of the synthesis. IR spectroscopy analysis can be made by the indexation of the major bands from the SiO_2 moiety, 1300-1000 cm^{-1} and near 800 cm^{-1} and also to show the removal of the organic alkyl ammonium template by comparing the spectra in the range 2850-3000 cm^{-1} where the organic part will absorb.

NOTES FOR THE INSTRUCTOR

Discussion

PILCs

In the case of the initial clay, the maximum of the diffraction peak is for 2θ of 7.09° , giving a value for the basal spacing (d_{001}) of 1.25 nm. This value is calculated from the basic equation of the Bragg law, defined as $n\lambda = 2d\sin\theta$, where, n is an integer number – the diffraction order, d is the distance between the planes, θ is the Bragg angle and λ is the wavelength of the X-ray (commonly from a Copper $\text{K}\alpha$ X-ray source with $\lambda = 0.15418$ nm). Students may note that the diffraction peaks for the natural clays are often broad, relatively to other minerals such as quartz for instance. This is because clays may present some variation in their chemical composition and also in the crystal sizes. The obtained d_{001} value is within the range ascribed to montmorillonite type clays.^{S1,S2} After the intercalation with the Zr-polyoxocations started (the “Before washing” sample) the diffraction peak became broader as a result of the still incomplete intercalation process. After the washing process two peaks are noticed, both due to d_{001} reflections. The peak for the lowest 2θ values (d_{001} of 1.93 nm) can be ascribed to the intercalated clay and the peak to higher 2θ values can be attributed to a portion of clay that was in fact non-intercalated, a situation that was already reported in the literature.^{S3} During calcination the Zr-polyoxocations dehydroxylate to form the oxide and that is why the d_{001} value of the final sample (1.70 nm) is lower than for the samples before calcination. To obtain the gallery heights of the pores, the dimensions of the clay layers,^{S3} 0.96 nm, needs to be subtracted from the d_{001} value, giving 0.74 nm.

MCM-41 materials

Typically, MCM-41 exhibits an XRD pattern with three or more diffraction lines which can be indexed to a hexagonal lattice as a result of the regular, hexagonal array of uniform channels.⁵⁴ For an hexagonal lattice the relation between the unit cell parameter (a_0) and the d_{100} obtained from the X-ray diffraction pattern is^{54,55} $a_0 = 2d_{100} / \sqrt{3}$ as schematically presented in Figure 2S. Pore diameters can be estimated from the unit cell parameter - a_0 , subtracting the thickness of the pore walls (cf. Figure 2S). Although some discussion exist in the literature about the thickness of the pore walls in MCM-41 type materials⁵⁵ an average value of 1 nm for the calcined materials can be admitted. The values of d_{001} and a_0 for the prepared samples, before and after calcination, are given in Table 1.

Additional and complementary activities

IR spectroscopy (in KBr wafers) can be used to study the MCM-41 samples for instance by the indexation of the major bands from the SiO_2 moiety. Examples are the strong and broad absorbance band in the $1300\text{--}1000\text{ cm}^{-1}$ region assigned to the asymmetric stretching modes, $\nu_{\text{as}}(\text{Si-O-Si})$, and the band near 800 cm^{-1} assigned to the symmetric stretching mode $\nu_{\text{s}}(\text{Si-O-Si})$.⁵⁶ Additionally, IR spectroscopy of the MCM-41 samples, before and after calcination, can also be used to show the removal of the organic alkyl ammonium template by comparing the spectra in the range $2850\text{--}3000\text{ cm}^{-1}$ where the organic part will absorb.

If nitrogen adsorption at $-196\text{ }^\circ\text{C}$ is available the characterization of porosity can be extended by the determination of the nitrogen adsorption isotherms and the evaluation of the specific surface areas and pore volumes of the materials.⁵⁷ Typical nitrogen adsorption isotherms in a MCM-41 material and in a Zr-PILC are shown in Figure 3S.

The prepared materials can be used for activities described in previous laboratory activities such as gas-chromatographic studies⁵⁸ or the adsorption of water pollutants for environmental applications.^{59,510} Both, the Zr-PILC and the MCM-41 can be used to illustrate the possibilities of porous materials as catalyst supports. For this, an inorganic complex that the students may have previously prepared, for instance in their laboratories of Inorganic Chemistry, such as a

$[\text{Cu}(\text{acac})_2]$, $[\text{Fe}(\text{acac})_3]$, Ferrocene, etc., can be selected. Various methodologies of introducing inorganic complexes in porous solids are described in the literature and can easily be adapted, with the possibilities of the characterization of the unsupported and supported complexes by spectroscopic techniques.⁵¹¹

The X-ray diffractogram of the parent clay can be used to illustrate the systematic use of the ICDD (International Center for Diffraction Data) files in the identification of the minerals when these occur in a mixture. An example is in Figure 4S.

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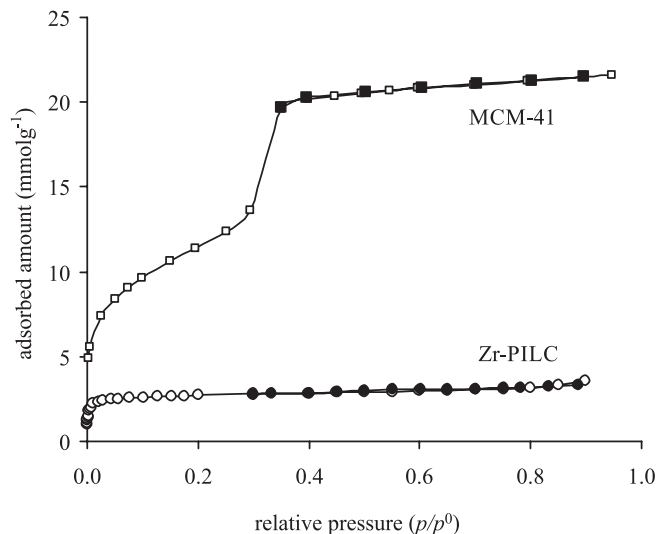
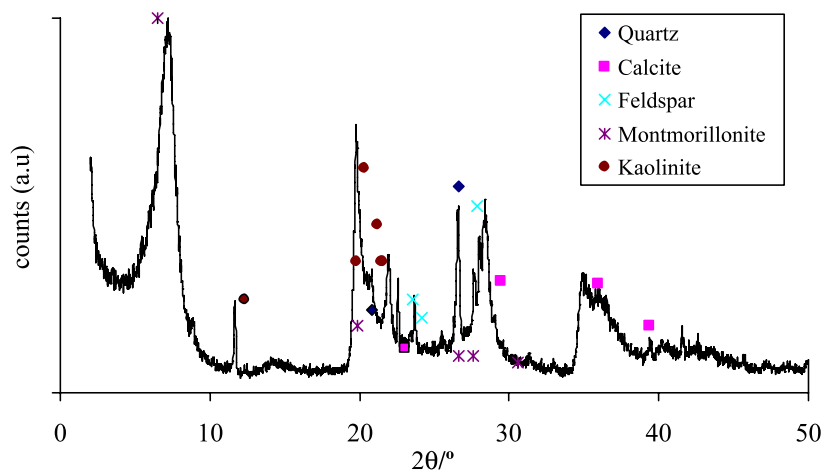


Figure 3S. Nitrogen adsorption isotherms in a mesoporous MCM-41 material and in a microporous Zr-PILC. (open points- adsorption; closed points - desorption)



Quartz		Feldspar		Calcite		Montmorillonite		Kaolinite	
39-1425		9-466		5-586		13-259		14-164	
2θ / °	relative intensity	2θ / °	relative intensity	2θ / °	relative intensity	2θ / °	relative intensity	2θ / °	relative intensity
20.85	22	27.89	100	23	12	6.494	100	12.335	100
26.662	100	23.51	25	29.4	100	19.846	18	19.81	35
36.852	8	24.13	20	35.97	14	26.668	10	20.324	60
39.456	8	24.27	16	39.402	18	27.594	10	21.208	45
50.141	14	25.36	10	43.176	18	30.591	8	21.5	35
59.948	9	13.84	20	47.49	17			21.451	35
68.323	8	22.03	16	48.513	17			24.858	80

Figure 4S. Example of identification of some minerals that, in addition to montmorillonite, can be present in starting material. The Table includes the number of the respective ICDD powder diffraction files