One-pot synthesis of iron doped mesoporous silica catalyst for propane ammoxidation

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Iron doped hexagonal mesoporous silica (Fe-HMS) was successfully prepared by one-pot direct synthesis at ambient temperature by using dodecylamine as a templating agent. The catalyst has been characterized by various techniques including XRD, DR-UV-Vis, FTIR, SEM and N2 adsorption. The obtained material exhibits very large surface area (above 1000 m² g⁻¹), mesoporosity with narrow distribution of pores (around 4 nm) and nano-crystallites with the 20–30 nm in size. The concentration of iron in the catalyst was 0.95 wt%. The prepared material was tested in the direct ammoxidation of propane. It was shown that direct synthesis leads to better dispersion of the metal compared to the material prepared by impregnation of pure HMS silica. It is also demonstrated that this material is able to catalyze the direct ammoxidation of propane and shows quite high selectivity to acrylonitrile (52% at 11% conversion of propane).

1. Introduction

Iron doped microporous material Fe-MFI and the [Fe]-MFI represent well known and abundantly studied catalytic materials for a variety of industrial reactions, among them for example the selective oxidation of benzene to phenol,¹⁻³ the reduction of NOₓ and N₂O with ammonia or hydrocarbons⁴⁻⁹ and also the direct decomposition of N₂O.¹⁰,¹¹ Although this material reaches very interesting results in the above mentioned reactions, it also suffers from drawbacks connected to the diffusion limitations (due to its microporosity – pore size is 0.51 × 0.56 nm). In order to overcome these problems several approaches have been proposed and studied. Some of these are: (i) the synthesis of microporous materials with smaller crystal size¹² or new microporous material with increased pore size,¹³ (ii) the preparation of hierarchical materials¹⁴,¹⁵ and (iii) the preparation of mesoporous materials.¹⁶

In our previous reports we studied microporous systems containing iron species ([Fe]-MFI, [Fe]-MTW) in the direct ammoxidation of propane.¹⁷,¹⁸ These materials were recognized as active and selective catalysts for this reaction only if no strong acid sites were present in the catalyst. It was found that the fast removal of nascent acrylonitrile (ACN) from the catalysts is very important because strong acid sites bind acrylonitrile, substantially increase the residence time of this product and cause its decomposition to acetonitrile and CO₂.¹⁷ Therefore, the best MFI catalyst was Fe-silicalite, a material without strong acid sites (without aluminum in the framework). Another improvement of catalyst behavior was reached by changing the crystallite size. Superior catalytic performance was exhibited by Fe-silicalite characterized by very small size of crystals (about 0.15 μm). The acrylonitrile yield of 12% and productivity up to 9.3 molACN (kgcat h)⁻¹ at 540 °C was obtained over our best catalyst.¹⁹

This paper reports on the successful synthesis of mesoporous ferrosilicate (Fe-HMS) and its catalytic activity in direct ammoxidation of propane. Such material can benefit from two main advantages (i) as a mesoporous material it should overcome the problems connected with the diffusion, (ii) it contains low amount of strong acid sites even at high concentration of heteroatoms.²⁰ The catalytic behavior of Fe-HMS prepared by direct hydrothermal synthesis is compared with a catalyst prepared from hexagonal mesoporous silica by wet impregnation in order to elucidate the effect of iron origin and speciation. In the discussion section, the reported results are compared with our previous results obtained on microporous Fe-silicalite catalysts and discussed in a broader context.

2. Experimental

2.1 Sample preparation

Two samples of hexagonal mesoporous silica (HMS) structure with different iron content were prepared by different methods – by wet impregnation and by direct synthesis. The preparation process of HMS was described earlier.²¹ It is worth of notice that before the calcination the sample was stirred for 1 hour in ethanol in order to remove the major part of remaining DDA. Subsequently the sample was calcined in a flow of air at 540 °C.
for 8 h with heating rate 1 °C min⁻¹. The sample prepared by impregnation of HMS (denoted as Fe-HMSimp) was synthesized using an aqueous solution of Fe(NO₃)₃·9H₂O. The amount of iron precursor was calculated in order to obtain the material with ca. 0.5 wt% of iron. Impregnation of HMS proceeded below the boiling point of solvent to evaporation of solution. The impregnated sample was dried at 120 °C in air overnight and then calcined at 600 °C for 8 hours in a flow of air.

The sample prepared by direct synthesis (denoted as [Fe]-HMS) was prepared according to the procedure reported by Liu et al.²² The molar composition of this matrix solution was 1.0 SiO₂ : 0.006 Fe(NO₃)₃ : 0.27 DDA : 11 EtOH : 1 i-PrOH : 100 H₂O. The mixture was stirred for 24 hours at room temperature; the formed solid was filtered, washed thoroughly with de-ionized water, dried at ambient temperature, directly calcined in nitrogen and after reaching 500 °C calcined in air for 6 hours to obtain [Fe]-HMS.

2.2 Sample characterization and catalytic test

2.2.1 Characterization. The chemical composition of all investigated samples was determined by X-ray fluorescence spectroscopy by ElvaX (Elvatech, Ukraine) equipped with a Pd anode. The structure and crystallinity of catalysts were probed by scanning electron microscopy using a TESCAN MIRA3 FE-SEM instrument and by X-ray diffraction (D8-advance diffractometer, Bruker AXE, Germany) in the low angle 2θ range (from 0.2 to 10°) and at higher angle range (2–35°) with Cu Kα radiation (λ = 1.5406 Å).

The BET surface area and pore diameter distribution of the samples were determined according to the N₂ adsorption isotherms. Nitrogen was used as adsorptive and supplied by Linde (purity 99.999 vol.%). Sorption isotherms of nitrogen at 77 K were determined using an ASAP 2020 instrument (Micromeritics, USA) and evaluated by MicroActive Software (Micromeritics, USA). In order to attain a sufficient accuracy in the accumulation of the adsorption data, this instrument is equipped with pressure transducers covering the 133 Pa, 1.33 kPa and 133 kPa ranges. Before each sorption measurement the sample was degassed to allow a slow removal of the most of preadsorbed water at low temperatures. This was done to avoid potential structural damage of the sample due to surface tension effects and hydrothermal alternation. Starting at ambient temperature the sample was degassed at 110 °C (temperature ramp of 0.5 °C min⁻¹) until the residual pressure of 1 Pa was attained. After further heating at 110 °C for 1 hour the temperature was increased (temperature ramp of 1 °C min⁻¹) until the temperature of 250 °C was achieved. The sample was degassed at this temperature under turbomolecular pump vacuum for 8 hours. The specific surface area was calculated according to BET method. The mesopore and external surface area were determined by means of t-plot using Harkins–Jura equation for calculation of adsorbed layer thickness, pore volume and pore size distribution were determined by NL DFT approach by using the “N2 @ 77K” model for cylindrical pores and oxide surface.

3. Results and discussion

3.1 Characterization

N₂ adsorption isotherms of the impregnated Fe-HMS catalyst together with the parent HMS silica support are displayed in Fig. 1A and textural characteristics are summarized in Table 1. Both isotherms exhibit a very similar shape characterized by a gentle step around a relative pressure of 0.2 and no hysteresis loops evidencing the presence of mesopores with relatively small size. Surface area of parent silica support was 994 m² g⁻¹. Impregnation of HMS support by nitrate as a source of iron led to a much more distinct decrease of surface area (795 m² g⁻¹, i.e. by about 20 rel%), which can be caused by the presence of iron oxide clusters (evidenced by UV-Vis spectroscopy, see
below) blocking the pores. Indeed, total pore volumes of parent HMS and impregnated Fe-HMS catalyst differ by about 21 rel% in very good agreement with the decline of surface area. Area of external surface of crystallites determined from t-plots (t-plot for parent HMS is displayed in Fig. 2A, t-plot of impregnated Fe-HMS catalyst is not reported here for the sake of brevity) was found to be 55 and 50 m² g⁻¹ for parent HMS and impregnated Fe-HMS, respectively. Such relatively large values are in line with non-compact crystallites with some macroporosity evidenced by SEM (see Fig. 4). Neither parent nor impregnated Fe-HMS catalyst exhibits microporosity as is manifested by the zero intercept of the straight line fitted BET range of t-plot (t ranging from 0.35 to 0.5 nm). The texture of the catalyst prepared by direct synthesis exhibits significant differences (see Fig. 1B). Massive condensation of nitrogen, manifested by a sharp increase of the adsorbed amount at relative pressures above 0.9, indicates the presence of large mesopores/macropores or interparticle adsorption and condensation. SEM images (see Fig. 4) clearly show crystallites with size around 20–30 nm, which agrees well with the characteristics of the adsorption isotherm. Therefore, an extremely high amount of adsorbed nitrogen at relative pressures above 0.9 is caused by condensation in interparticle void space. The nanosize of crystallites is also confirmed by the determination of the external surface area from t-plot which was found to be 342 m² g⁻¹ (see Fig. 2B), whereas the total surface area determined by BET method is 1033 m² g⁻¹. The size of the pores determined from the isotherms by NL DFT method ranges from 2 to 5 nm (see Fig. 3) with the main maximum at 3.1 nm for HMS parent and impregnated sample and 4 nm for synthesized [Fe]-HMS catalyst. The pore size distribution (PSD) curve of the synthesized catalyst exhibits a narrower peak

Table 1  Chemical composition and results of physico chemical characteriztion of investigated materials

<table>
<thead>
<tr>
<th>Material</th>
<th>Fe (wt%)</th>
<th>BET (m² g⁻¹)</th>
<th>Ext (m² g⁻¹)</th>
<th>Vp (cm³ g⁻¹)</th>
<th>dp (nm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>HMS</td>
<td>—</td>
<td>994</td>
<td>55</td>
<td>0.48</td>
<td>3.1</td>
</tr>
<tr>
<td>Fe-HMSimp</td>
<td>0.54</td>
<td>795</td>
<td>50</td>
<td>0.38</td>
<td>3.0</td>
</tr>
<tr>
<td>[Fe]-HMS</td>
<td>0.95</td>
<td>1033</td>
<td>342</td>
<td>0.57</td>
<td>4.0</td>
</tr>
</tbody>
</table>

Fig. 1  (A) N₂ adsorption/desorption isotherms of parent HMS silica support (a) and impregnated Fe-HMS catalyst (b). (Fe-HMS isotherm is shifted up by 200 cm³ STP/g for sake of clarity) (B) N₂ adsorption isotherm of [Fe]-HMS.

Fig. 2  t-plots constructed by means of Harkins–Jura equation from the isotherm of parent HMS (A) and [Fe]-HMS (B).
compared with the PSD curves of silica HMS and impregnated samples. It is necessary to note that the total pore volume of synthesized [Fe]-HMS sample is also significantly larger than the total pore volume of parent silica HMS and therefore also than impregnated Fe-HMS catalyst (cf. data in Table 1).

Fig. 4 reports the scanning electron micrographs of parent HMS, of impregnated HMS and of the sample prepared by the direct synthesis [Fe]-HMS. Micrographs of HMS and Fe-HMSimp are quite similar showing spherical particles with a wide range of macropore crevices. The size of these particles is in a range from 0.5 to 1 μm. As can be seen from the micrograph, these particles form larger aggregates. The micrograph of the sample [Fe]-HMS shows that the particles are considerably smaller, being about 20–30 nm in size. The shape of the particles is markedly different from the previous ones, showing rather worm-like structure.

XRD patterns of a parent HMS and Fe-HMS samples (Fig. 5) show a broad low-angle diffraction peak at 2θ = 2–3° (100) with poorly resolved satellite at around 5°, which is typical for hexagonal lattice structure of mesoporous materials. The d_{100} distance determined from the position of the main diffraction peak by means of Bragg’s law varies between 3.2 and 3.5 nm in good agreement with data from the pore size distribution and indicates a relatively thin channel wall. The wide angle region does not show any diffraction peaks of crystalline iron oxides. Possibly, the particles of iron oxides formed upon the impregnation of HMS (see the results from UV-Vis spectroscopy) are too small, amorphous or in very low amount to be detected by X-ray diffraction.

In order to obtain more information about the presence of different iron species located in the structure of HMS of both Fe-HMS samples, the adsorption of nitric oxide was studied by means of FTIR spectroscopy. The adsorption of NO on Fe-containing systems is a very often used method for the characterization of Fe species (especially in connection with IR spectroscopy). Most authors agree that (di)nitrosyls are formed exclusively with the participation of Fe^{2+} ions because NO adsorption on oxidized samples is negligible. Therefore, samples were reduced (in 80 mbar H_{2}) at 450 °C during the night before the adsorption. It must be noted that NO adsorption on oxidized samples was measured too and only a very weak band was detected in agreement with literature. Due to negligible NO adsorption on oxidized samples, only the spectra on reduced samples are presented and discussed here.

The IR spectra of NO adsorbed on Fe-HMS samples are shown in the Fig. 6 and 7. Spectra contain a main band centered at about 1820 cm\(^{-1}\). The vibration of N=O bond is significantly shifted to lower wavenumber considering frequency of free gaseous NO (1876 cm\(^{-1}\)). It is evidence of π-back bond formation typical for metals with higher density of d-electrons and lower oxidation state. A detailed inspection of the spectra reveals distinct asymmetry of the band with a tail at high-frequency side. This is caused by the overlapping of two bands with the maxima approximately at 1820 and 1836 cm\(^{-1}\). The comparison of normalized envelope curves (see Fig. 7A) clearly shows that the tail is more intense in the case of sample [Fe]-HMS. To obtain further information, the desorption of NO from the samples under dynamic vacuum at RT and
Fig. 5  XRD patterns of HMS samples.

Fig. 6  FTIR spectra of [Fe]-HMS and Fe-HMSimp, desorption of NO with time. The first full line spectrum on the top was collected immediately after the start of desorption, the last full line spectrum at 50 min of desorption, the dotted line spectra were collected after 1 and 10 min of desorption at 100 °C.

Fig. 7  Comparison of the adsorption spectra of [Fe]-HMS and Fe-HMSimp. (A) Spectra of the samples after the start of desorption at RT, (B) spectra recorded during the desorption at 100 °C, shown spectra correspond to half of the initial value of the intensity of the desorption spectra at RT.
subsequently at 100 °C was investigated too. It is evident that the adsorption species are quite stable at RT and the intensity of the bands decreases very slowly in time (approximately 30% of NO was desorbed during the first 45 minutes under vacuum by turbomolecular pump). Not only intensity, but also shape of the band was changing during desorption. As can be seen from the spectra in Fig. 6, the band at 1820 cm$^{-1}$ decreases in intensity faster than the band at 1836 cm$^{-1}$. The spectra of the samples measured at coverage of the samples of about half of the initial value (Fig. 7B) clearly show that the band at 1836 cm$^{-1}$ dominates in the case of sample [Fe]-HMS, whereas the impregnated sample is still characterized by a band centered at 1820 cm$^{-1}$. Therefore, we can conclude that both samples differ in nature of Fe species; the synthesized sample exhibits a much higher population of species characterized by IR band at 1836 cm$^{-1}$ than the impregnated sample. At a lower coverage degree, a small and relatively broad band at about 1760 cm$^{-1}$ is detected in the spectra of [Fe]-HMS. A band with similar position and stability was observed in the spectra of NO adsorbed on Fe–Y systems and ascribed to bent species.23 It is worth noticing that spectra similar to ours were not measured against the studied HMS materials, as was observed in the case of Fe-silicalite,27 or isolated extra-framework ions.6,28 The shoulder at 35 000 cm$^{-1}$ should also correspond to isolated iron species. According to the literature it is difficult to distinguish whether these isolated Fe$^{3+}$ species are in tetrahedral or higher coordination.29 However in the literature it has been described that the absorption band at about 35 000 cm$^{-1}$ is caused by Fe$^{2+}$ in octahedral coordination.27,30 The broad absorption band below 30 000 cm$^{-1}$ is caused by small oligonuclear clusters (Fe$^{3+}$O)$^{x}$ and finally the absorption band below 20 000 cm$^{-1}$ is caused by bulky clustered Fe$_{2}$O$_{3}$ species.31

More detailed information about the dispersion and local coordination structure of iron species can be obtained from UV-Vis spectroscopy by evaluation of the absorption edge energy ($E_g$) from Tauc’s plot (the dependency of $(F(R_\infty)-h\nu)^2$ against $h\nu$, where $F(R_\infty)$ is Kubelka–Munk function, $h$ is Planck constant and $\nu$ is frequency of photon).32 $E_g$ can be determined from the Tauc’s law: $(F(R_\infty)-h\nu)^2$ is proportional to $(h\nu - E_g)$. Basically, the higher is the calculated value of the absorption band energy the higher is the dispersion of metal in the system. The calculated values of the $E_g$ of Fe catalysts together with the $E_g$ values of standard compound (diluted in the same manner as the studied HMS materials) are listed in Table 2. The $E_g$ value of [Fe]-HMS sample is 3.83 eV. Similar values have been already observed by Zhang et al. studying FeMCM-41 prepared by impregnation and by template ion exchange.33 On the basis of the correlation of the results from EXAFS and XANES measurement they suggested that the values of the band energy about 4 eV corresponds to small FeO$_x$ clusters with iron in octahedral coordination. The

![Fig. 8 UV-Vis spectra of hydrated HMS samples.](image)

| Table 2 Comparison of absorption band energy for the studied samples |
|-------------------------|----------------|
| Material                | $E_g$ (eV)   |
| Fe-HMSimp               | 2.00         |
| [Fe]-HMS                | 3.83         |
| Fe$_2$O$_3$-SiO$_2$     | 2.11         |
impregnated sample exhibits a significantly lower $E_g$ value (2.00 eV). Similar values were also found for various samples by other authors $^{33-35}$ in the case of presence oxide species. In agreement with this statement, model oxides compound (Fe$_2$O$_3$-SiO$_2$) exhibits also very low $E_g$ value (2.11 eV). Therefore, we can conclude that the catalyst prepared by direct synthesis contains mainly (besides isolated Fe cations) octahedral FeO$_x$ clusters with a small degree of polymerization, whereas the impregnated one contains substantial amount of oxide species.

### 3.2 Catalysis

The main products of the direct ammoxidation of propane were acrylonitrile (ACN), propylene, acetonitrile (AcCN), carbon oxides and ethane.

Fig. 9 shows the comparison of the catalytic performance of the Fe-HMS samples. The conversion of propane reaches very similar values; a slightly higher conversion is obtained over the sample prepared by direct synthesis (11% vs. 9% for the sample Fe-HMSimp). Consumption of oxygen is also similar (about 42% of oxygen), conversion of ammonia reaches higher values over the impregnated sample (60% vs. 51%). Selectivity to ACN varies a lot, the sample prepared by direct synthesis reached twice as high selectivity to ACN compared to the impregnated sample. Selectivity to propylene is higher in the case of the impregnated sample (33% vs. 23%), so is the selectivity to carbon oxide, the difference is twice as much (35% vs. 16%). These results clearly demonstrate that impregnation of silica does not lead to selective catalytic system. This is in accordance with the results from UV-Vis spectroscopy; the impregnated sample showed high concentration of bulky clustered Fe$_2$O$_3$ species.

Fig. 10 shows selectivity−conversion relationship for the samples [Fe]−HMS, Fe−HMSimp in comparison with Fe-silicalite samples, calcined and activated in the direct ammoxidation of propane.

![Graph showing selectivity−conversion relationship](image)

Fig. 9 Comparison of the catalytic activity of the impregnated Fe−HMS samples together with [Fe]−HMS in the direct ammoxidation of propane.

Another comparison of the catalytic activity of microporous and mesoporous systems can be given by expressing the productivity of the catalyst and turn over frequencies, see Table 3. Regarding the TOF it can be clearly seen that activated microporous material reaches the best results, however comparing the productivities we see that the mesoporous material [Fe]−HMS is able to give good results in comparison to calcined microporous Fe-silicalite. This is quite interesting mainly by taking into account the other studies from the literature dealing with N$_2$O mediated ODH of propane $^{28,36}$ where it has been stated that microporous materials reach better results than the mesoporous ones. It has been proposed that it is due to the lack of the intimate contact between potentially active Fe sites and reactant molecules $^{28}$ another assumption was that the chemical nature of iron sites is more important than the pore structure. $^{36}$

On the basis of our results from characterization techniques (UV-Vis) and from catalytic studies we tend to agree with the opinion that the nature of the iron sites is the key factor influencing the catalytic properties. Our study showed that use of the same matrix (HMS) but different method of iron incorporation leads to material with different catalytic properties. The impregnated sample contains only extra-framework Fe species with high heterogeneity covering species from
isolated Fe ions to oxidative nanoparticles. The synthesized sample could contain both framework and extra-framework species. Isolated Fe species, represented in the UV-vis spectra by absorption bands at 46 000 and 41 000 cm\(^{-1}\), can originate from isomorphously substituted iron ions in the framework of the materials or isolated extra-framework ions. However, it is not possible to distinguish between these two forms of isolated Fe cations directly from UV-Vis spectra. Framework Fe species could be distinguished by UV Raman spectroscopy\(^7\) by vibrational bands at 510 and 1090 cm\(^{-1}\) assigned to the symmetric and asymmetric Fe–O–Si stretching modes of the amorphous silica framework. On the other hand, we proved the presence of extra-framework Fe species in the synthesized sample by FT-IR spectra of adsorbed NO. Our previous studies conducted on Fe-silicates clearly showed that extraction of Fe species from the framework is essential for reaching high catalytic activity in propane ammoxidation.\(^{18,19}\) Therefore, we deduced that framework Fe atoms cannot act as an active site. We are aware that framework of amorphous mesoporous silica is more flexible than the zeolitic structure which can change the properties of tetrahedral atoms in the framework. However we believe that extra-framework species are much more active species and the catalytic performance of the catalyst is given by presence of such particles. This has been already proposed in the literature by numerous studies dealing with the investigation of the catalytic behaviour of iron doped micro- and meso-porous materials in various reactions, including hydroxylation of benzene to phenol,\(^{38,39}\) SCR or decomposition of \(\text{N}_2\text{O}\),\(^{10,41}\) ODH of propane\(^{42,43}\) and also in the direct ammoxidation of propane.\(^{17}\) Unfortunately, extra-framework Fe particles exhibit high level of heterogeneity and discovering active Fe particle is a very challenging task.

### 4. Conclusion

The direct synthesis procedure reported in this paper led to the mesoporous material with considerably larger total pore volume, surface area exceeding 1000 m\(^2\) g\(^{-1}\) and exhibiting nano-crystallites of 20–30 nm in size facilitating mass transport and diffusion of reactant and products. The direct synthesis of Fe-mesoporous silica led to better dispersion of iron in the material compared with impregnation procedures. This had the positive effect on the catalytic activity of the materials, as it has been shown, that over [Fe]-HMS quite high selectivity to acrylonitrile was achieved, meanwhile the impregnated sample gave high selectivity to carbon oxides. Catalytic activity and selectivity of the [Fe]-HMS catalyst, able to compete with Fe-MFI zeolites, was reached without any special activation (the sample was only calcined to the reaction temperature in the air), whereas the most active and selective Fe-silicalite zeolite was activated by reductive nitridation in the flow of ammonia and propane at reaction temperature for relative long time (from 5 to 15 h).

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