Studies in Surface Science and Catalysis 12

METAL MICROSTRUCTURES IN ZEOLITES
Preparation – Properties – Applications
Proceedings of a Workshop, Bremen, September 22–24, 1982

Editors

P.A. Jacobs
Centrum voor Oppervlaktescheikunde, Katholieke Universiteit Leuven, Leuven, Belgium

N.I. Jaeger
Forschungsgruppe Angewandte Katalyse, Universität Bremen, Bremen, F.R.G.

P. Jírů
Czechoeslovak Academy of Sciences, J. Heyrovský Institute of Physical Chemistry and Electrochemistry, Prague, Czechoslovakia

G. Schulz-Ekloff
Forschungsgruppe Angewandte Katalyse, Universität Bremen, Bremen, F.R.G.

ELSEVIER SCIENTIFIC PUBLISHING COMPANY
Amsterdam — Oxford — New York 1982
CONTENTS

Preface VII

The chemistry of ruthenium in zeolites (J.H. Lunsford) 1

EPR studies of Co(II)-bis(dimethylglyoximato)-complexes and their oxygen adducts in a zeolite X matrix
C.J. Winscom, W. Lubitz, H. Diegruber and R. Möseler) 15

In-situ-UV/VIS-microscopespectrophotometry of reactions of Co^{2+}-complexes within single crystals of faujasite-type zeolites (H. Diegruber and P.J. Plath) 23

Mossbauer spectroscopic studies of ferrous ion exchange in zeolite A (L.V.C. Rees) 33

Metal inclusion complexes of zeolite A (N. Petranović and R. Dimitrijević) 45

Quantum chemical study of the properties of Fe, Co, Ni and Cr ion-exchanged zeolites (S. Beran and P. Jiráš) 53

Cationic rhodium complexes and rhodium metal aggregates in zeolite Y (H. van Brabant, R.A. Schoonheydt and J. Pelgrims) 61

Mechanisms of formation and stabilization of metals in the pore structure of zeolites (P.A. Jacobs) 71

Reflectance spectroscopic study of Ag^{+}, Ag^{0} and Ag clusters in zeolites of the faujasite-type (L.R. Gellens and R.A. Schoonheydt) 87

Chemical evidence for charged clusters in silver zeolites (H.K. Beyer and P.A. Jacobs) 95

UV/VIS transmission spectroscopy of silver zeolites. I. Dehydration and rehydration of AgA and AgX (H.G. Karge) 103

Adsorption and decomposition of iron pentacarbonyl on Y zeolites (Th. Bein, P.A. Jacobs and F. Schmidt) 111

Stabilization and characterization of metal aggregates in zeolites. Catalytic properties in CO + H\textsubscript{2} conversion (D. Ballivet-Tkatchenko, G. Coudurier and Nguyễn Duc Chau) 123

Influence of controlled structural changes on the catalytic properties of zeolites (P. Jiráš) 137

Behaviour of Fe species in zeolite structure (B. Wichterlová, L. Kubelková, J. Nováková and P. Jiráš) 143

To the differences in properties of Ni metal particles and Cr cations in stabilized and nonstabilized zeolites (V. Patzelová, Z. Tvarůžková, K. Mach and A. Zukal) 151
Incorporation of volatile metal compounds into zeolitic frameworks (P. Fejes, I. Kiricsi and I. Hannus) 159

Characterization of metal aggregates in zeolites (P. Gallezot and G. Bergeret) 167

Nuclear magnetic resonance study of xenon adsorbed on metal-NaY zeolites (size of metal particles and chemisorption) (J. Fraissard, T. Ito, L.C. de Menorval and M.A. Springuel-Huet) 179

Characterization of metal aggregates in zeolites (F. Schmidt) 191


Investigations of the aggregation state of metals in zeolites by magnetic methods (W. Romanowski) 213

Dielectric properties of X-type zeolites containing small metallic nickel particles (J.C. Carru and D. Delafosse) 221

Modification of chemisorptive and catalytic properties of Ni° highly dispersed on zeolites of various composition (G.N. Sauvion, M.F. Guilleux, J.F. Tempere and D. Delafosse) 229

Nickel in mordenites - formation and activity of metallic complexes, clusters and particles (E.D. Garbowski, C. Mirodatos and M. Primet) 235

Dispersion of nickel and ruthenium in zeolites L, Y and mordenite (S. Narayanan) 245

Effect of the reaction medium on the metal microstructure of nickel-zeolite catalysts (N.P. Davidova, M.L. Valcheva and D.M. Shopov) 253

Activities and selectivities of reduced NiCaX faujasites in the carbon monoxide hydrogenation reaction (H. Schrubbers, G. Schulz-Ekloff and G. Wildeboer) 261

Fischer-Tropsch synthesis on polyfunctional manganese/iron-pentasil zeolite catalysts (K. Müller, W.-D. Deckwer and M. Ralek) 267

Oxidation of ethylene on silver-loaded natural zeolites (G. Bagnasco, P. Ciambelli, E. Czaran, J. Papp and G. Russo) 275

Author Index 283
ADSORPTION AND DECOMPOSITION OF IRON PENTACARBONYL ON Y ZEOLITES

Th. BEIN¹,⁴, P.A. JACOBS² and F. SCHMIDT³

¹Institut für Physikalische Chemie der Universität Hamburg, Laufgraben 24, D-2000 Hamburg 13 (F.R.G.)
²Centrum voor Oppervlaktescheikunde en Colloidale Scheikunde, Katholieke Universiteit Leuven, De Croylaan 42, B-3030 Leuven (Heverlee) (Belgium)
³Present address : ²

ABSTRACT

The adsorption isotherms of Fe(CO)₅ on NaY and HY zeolites obtained in McBain balances show micropore adsorption, the saturation at p/p₀ = 0.5 being 39 and 42 % per dry wt, respectively. IR results indicate a restricted mobility of the encaged complex. Nevertheless it can thermally be desorbed to a great extend in vacuum.

For the first time, well distinguishable decomposition phases of zeolite-adsorbed Fe(CO)₅ are found by thermogravimetric analysis. These phases are associated with species bearing 2(4) and 1/4(1) CO ligands per Fe in the case of NaY(HY). New evidence is found for the intermediate Fe(CO)₂. The slow decomposition reaction in inert atmosphere is completed already between 70 and 90°C, providing an iron content of 10.5 ± 0.5 wt %.

INTRODUCTION

With respect to the industrial importance of iron catalysts and the still not entirely understood particle size effect in catalysis, it is desirable to dispose of model catalysts with variable, narrow particle size distribution. Zeolites have proved to be suitable supports for metals (ref. 1,2) and to behave as model catalysts. Our aim is therefore to obtain Fe(0) containing zeolites with narrow particle size distributions. Reduction of Fe(II) exchanged Y-type zeolites was found to be impossible with H₂ (ref. 3,4), whereas reduction with sodium vapor resulted in highly dispersed iron metal (ref. 5-8).

With regard to the difficult procedures to be used in these methods, the decomposition of Y-zeolite adsorbed iron pentacarbonyl was chosen as an alternative.

Thermal decomposition of this complex has already been used to prepare dispersed supported iron (ref. 9), while recently it has been applied to Fe(CO)₅ loaded HY zeolite (ref. 10-12). In addition, decomposition by UV light was reported to provide a highly dispersed iron phase in the HY zeolite.

In the former studies, few quantitative details are given with regard to the parameters which govern adsorption and decomposition of the complex. In order
to arrive at a quantitative understanding of these processes, the adsorption and decomposition behaviour of iron pentacarbonyl in NaY and HY zeolite has been studied by means of gravimetric, thermogravimetric and IR-spectroscopic methods.

EXPERIMENTAL

Materials

Synthetic NaY with Si/Al = 2.46 was from Strem Chemicals. It was treated with 0.1 M NaCl solution to remove possible cation deficiencies, washed and air dried, and stored over saturated NH₄Cl solution. The NH₄Y form was obtained by conventional ion exchange. Before loading with iron carbonyl, both zeolites were degassed in situ at 450°C for about 12 hrs at 10⁻⁵ mbar, at a heating rate of 2°C/min.

Iron pentacarbonyl from Ventron (99.5 %) was cold distilled in the dark and stored over molecular sieve 5A. The zeolite samples for the McBain and IR measurements were loaded with the carbonyl as follows. The frozen carbonyl was outgassed in vacuum and allowed to warm up until the desired pressure was reached. All procedures with Fe(CO)₅ were performed in the dark, whereas the weight measurements at the McBain balance have been carried out in weak red light.

Methods

Adsorption isotherms were obtained in a McBain balance with calibrated quartz spring, with a precision of ± 0.5 %. The pressure was measured with a Bell & Howell pressure transducer BHL-4100-01, which is linear within ± 0.5 % up to 750 mbar.

Infrared spectra were taken with a Perkin Elmer 580B spectrometer from 4000 to 1200 cm⁻¹ (resolution 2 cm⁻¹) using a quartz cell with 80 mm path length and equipped with CaF windows of 3 mm thickness. The zeolite was pressed at 1 ton/cm² to selfsupporting films of ca. 5 mg/cm². All treatments were performed in situ in the IR cell.

Thermogravimetric measurements were done on a Mettler Thermoanalyzer 2 under He purge, mostly in the 10 mg range. Samples of 5 to 50 mg zeolite were outgassed by heating at 2°C/min up to 450° in a quartz oven and loaded at 20°C in a stream of dry helium containing ca. 4 mbar Fe(CO)₅. The flow rate of this stream was 2.8 1/h.

RESULTS AND DISCUSSION

1. Adsorption isotherms of Fe(CO)₅ on Y-zeolite

Comparison of the Fe(CO)₅ adsorption isotherms on NaY and HY shows rather similar behaviour (Fig. 1). The major uptake occurs at very low partial
pressures and remains almost constant up to \( p/p_0 \approx 0.5 \). At this partial pressure NaY and HY adsorb 39 and 42 mg of Fe(CO)\(_5\) per 100 mg of dry zeolite, respectively.

Desorption is reversible down to ca. \( p/p_0 = 0.1 \). After degassing for 15 hrs at \( 10^{-5} \) mbar, Fe(CO)\(_5\) loadings of 29 and 27 wt % are obtained for NaY and HY, respectively.

The adsorption behaviour can be explained in terms of nearly ideal micropore adsorption (ref. 13), the micropores being the supercages of the faujasite. The amount Fe(CO)\(_5\) adsorbed before capillary condensation occurs corresponds to 25 molecules/U.C. or 3.1 molecules per supercage. If an effective radius of 0.30 nm is assumed for the complex, the geometry of the supercages allows a maximum adsorption of three molecules per supercage.

This good agreement with the experimental results confirms the picture of completely filled supercages.

2. Infrared study of Fe(CO)\(_5\) adsorbed on zeolite Y

The Fe(CO)\(_5\) saturated zeolite wafers show no measurable transmission in the CO-stretching region. The IR spectra reported in Fig. 2 correspond therefore to samples loaded with about 10 % of the capacity obtained at saturation. The carbonyl vibrations show a rather similar pattern for both NaY and HY,
Fig. 2. (left) IR spectra of Fe(CO)$_5$/zeolite adducts at 20°C (saturated for 10%). A: Fe(CO)$_5$/NaY; B: Fe(CO)$_5$/HY; dotted lines: zeolites degassed at 450°C.

Fig. 3. (right) OH spectrum of saturated Fe(CO)$_5$/HY adduct decomposed in 600 mbar He. A: zeolite degassed at 450°C; B: HY saturated with Fe(CO)$_5$ at 20°C; C: sample B after heating at 150°C for 45 min.

respectively (Table 1a). Compared to the HY-adduct, Fe(CO)$_5$ adsorbed on NaY exhibits two additional bands at 2044 and 1945 cm$^{-1}$, all other bands only slightly being changed.

The carbonyl bands cannot be assigned definitely to particular species inside the zeolite cages. The assignment of the carbonylbands to monosubstituted species as ZO$-$Fe(CO)$_4$ (ref. 11,12) seems to be somewhat arbitrary because of the fortuitous agreement of some bands with those of complexes such as Fe(CO)$_4$P(CH$_3$)$_3$.

By $^{13}$C-nmr line-width broadening, a restricted mobility of the Fe(CO)$_5$ adsorbed in HY was found (10). This is quite reasonable since the adsorption experiments indicate complete filling of the supercages. The carbonyl bands of crystalline Fe(CO)$_5$ and of the Fe(CO)$_5$/zeolite adducts (Table 1) show fairly good agreement, although the intensities are different. This also is in line with the restricted mobility of the complex in the supercages.

In particular, the appearance of the sharp 2122 cm$^{-1}$ band can be explained by decreased site symmetry. Indeed, the intensity of the $v_1$ mode increases on
TABLE 1
IR frequencies of Fe(CO)$_5$ and its adducts (CO-stretching region).

<table>
<thead>
<tr>
<th>Adduct</th>
<th>Frequency/cm$^{-1}$</th>
<th>Ref.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Fe(CO)$_5$/NaY</td>
<td>2122w 2060s 2044s 2012s 1985s 1960sh 1945s</td>
<td>a</td>
</tr>
<tr>
<td>Fe(CO)$_5$/HY</td>
<td>2120w 2050s 2018s 1990b 1960s</td>
<td>a</td>
</tr>
<tr>
<td>Fe(CO)$_5$/HY</td>
<td>2112mw 2040s 2030sh 2010s 1985sh 1950ms</td>
<td>12</td>
</tr>
<tr>
<td>Fe(CO)$_5$ solid (-173°C)</td>
<td>2115($v_1$) 2033($v_2$) 2017sh 2003($v_6$) 1980($v_{10}$) 1956/48($^{13}C0$)</td>
<td>14</td>
</tr>
<tr>
<td>Fe(CO)$_5$ liquid (25°C)</td>
<td>[broad] 2002($v_6$) 1979($v_{10}$)</td>
<td>15</td>
</tr>
<tr>
<td>Fe(CO)$_5$ gas (25°C)</td>
<td>2034vv(T$v_6$) 2014vvs($v_{10}$)</td>
<td>16</td>
</tr>
</tbody>
</table>

*This work.

going from the gas phase to crystalline state (ref. 14).

The IR spectrum of liquid Fe(CO)$_5$ exhibits a broad and poorly structurated CO stretching band around 2000 cm$^{-1}$ (Table 1). This also is in contrast to the observed well structurated CO-bands of the Fe(CO)$_5$/zeolite adducts.

In both the NaY and the HY adduct no $v_{CO}$-bridging are shown, i.e., at 20°C no clusters with bridging CO ligands such as Fe$_2$(CO)$_9$ are generated.

The interaction of Fe(CO)$_5$ with the OH-bands of HY is illustrated in Fig. 3. Only the supercage hydroxyl groups (3645 cm$^{-1}$) disappear completely upon adsorption of the carbonyl. The band at 3550 cm$^{-1}$ broadens and increases in intensity. This can only be explained by the formation of an hydrogen bond of moderate strength between the complex and the supercage OH-groups.

3. Desorption of Fe(CO)$_5$ from Y zeolite

Heating the Fe(CO)$_5$ loaded zeolite wafers in a vacuum of $10^{-4}$ mbar results in a proportional decrease of intensity of all carbonyl bands, while no new bands appear (Fig. 4).

When the similar experiments are carried out on larger amounts of sample (100 mg), iron losses between 20 and 50 % with respect to carbonyl saturation are determined gravimetrically.

These observations must be explained by desorption of the iron complex, which also may be the reason for the different iron loadings obtained by other authors (ref. 10-12) after thermal decomposition in vacuum, with values ranging from 1 to 8 wt % Fe. The proportional decrease of all carbonyl bands (Fig. 4) also indicates that one single species is adsorbed in the zeolite cage, which seems to be the intact Fe(CO)$_5$. 
4. Decarbonylation of Fe(CO)$_5$/zeolite adducts by thermoanalysis

When in a thermobalance different amounts of zeolite are loaded with carbonyl vapor, the saturation loadings correspond to the adsorption isotherms (38 wt %). Samples are heated up to 200°C in a He stream at rates from 0.2 to 2°C/min.

a. Decomposition of Fe(CO)$_5$/NaY. Irrespective of the amount of sample and the heating rate, three distinct regions are found with respect to thermal behaviour. Two zones of slow weight loss are separated by a fast decrease in sample weight (Fig. 5A). The latter is accompanied by an endothermic DTA effect. It is striking, that the DTA effect always starts, when the sample has lost 15 wt % of its loading. The begin of the third zone is defined by the end of the DTA effect and always occurs when 26 wt % of loading are lost. Around 200°C, the sample weight becomes stable and corresponds to a loading of 10.5 %.

Begin- and end-temperature of the DTA effect are strongly dependent on the...
heating rate. Extrapolation of these temperatures to zero heating rate (ref. 17) indicates, that the same decomposition can be performed isothermally in the temperature region between 70 and 90°C. An isothermal experiment at 90°C after 11 hrs showed the break in the weight curve. Since the sample weight does not change from 200 to 400°C, the adsorbate present is considered to be metallic iron. Compared to the original carbonyl loading losses of iron must be smaller than 0.5 wt %.

This is in contrast to the results of vacuum decomposition and can be explained by an efficient hindering of the carbonyl diffusion at high pressures of inert gas.

The thermoanalytical results allow to depict carbonyl decomposition on NaY as follows:

\[
\text{NaY/Fe(CO)}_5 \xrightarrow{\text{slow}} \text{Fe(CO)}_2 \xrightarrow{\text{fast}} \text{Fe(CO)}_{1/4} \xrightarrow{\text{slow}} \text{Fe}
\]

The agreement between the measured weight loss and the one calculated according to this stoichiometry lies within 5 %.

b. Decomposition of Fe(CO)_5/HY. The thermogram of the HY/Fe(CO)_5 adduct is distinctly different from the one of the NaY adduct (Fig. 5B). First, the fast decomposition as indicated by the start of the DTA effect always appears at lower temperatures (83°C). Second, two weak, but reproducible endotherm DTA effects are observed instead of one, which correlate with the DTG minima. Above 144°C (the endpoint of the second DTA effect) no further weight loss occurs. Similar considerations as with NaY lead to the following stages of decomposition:

\[
\text{HY/Fe(CO)}_5 \xrightarrow{\text{slow}} \text{Fe(CO)}_4 \xrightarrow{\text{fast}} \text{Fe(CO)} \xrightarrow{\text{fast}} \text{Fe}
\]

Again, excellent agreement between calculated and measured values is obtained.

In previous work decomposition of Fe(CO)_5/HY in vacuum was reported to start at 25°C and to be complete at 200°C (ref. 12). The formation of Fe(CO)/HY is postulated in vacuo at 70°C (ref. 10). In these studies no reaction times were reported.

From our results it is clear that the temperature for complete decomposition is below 90°C and that decomposition is a very slow reaction.
In situ investigation of the Fe(CO)$_5$/zeolite decomposition by IR spectroscopy

Decomposition of the carbonyl adducts in a He atmosphere leads to results quite distinct from those of vacuum heating. For the Fe(CO)$_5$/NaY this is shown in Fig. 6. After heating a NaY sample (saturated with Fe(CO)$_5$) at 150°C for 10 min, a broad band around 1940 cm$^{-1}$ with a shoulder at 1860 cm$^{-1}$ is generated, replacing the 1985, 1960 and 1945 cm$^{-1}$ bands of the original adduct (Fig. 6D).

The original high frequency bands, in particular the 2044 cm$^{-1}$ vibration, strongly decrease in intensity after prolonged heating at 150°C, leaving a shoulder at the 2005 cm$^{-1}$ band and a weak band at 2070 (Fig. 6E).

The broad low frequency band changes into a vibration at 1900 cm$^{-1}$, which is the last band to survive by further heating to 200°C. Decomposition is complete after some hours at 200°C. No bands below 1800 cm$^{-1}$ are observed during decomposition.

For the Fe(CO)$_5$/HY adduct the decomposition is given in Fig. 7. Thermal treatment at 130°C of a saturated HY-adduct first leads to the generation of a new band around 1880 cm$^{-1}$ (Fig. 7D), whereas after 7 min. a relative decrease of the low frequency bands at 1960 and 1880 cm$^{-1}$ occurs (Fig. 7E). Prolonged heating at this temperature causes rapid decomposition. After 12 min. only three weak vibrations at 2065, 2030 and 2000 cm$^{-1}$ are left. Decomposition is completed after 45 min. at 150°C, and at the same time the original OH-bands are restored to about 75% of their initial intensity (Fig. 3C).

In both zeolites bands below 1900 cm$^{-1}$ are formed during the early decomposition, providing some evidence for bridged CO (ref. 18). They may be associated with Fe(CO)$_x$ species formed during the first phase of thermal decomposition. Bands at 1760 and 1790 cm$^{-1}$, which seemed to be characteristic for the Fe$_3$(CO)$_{12}$/HY adduct (ref. 12), have never been observed in the present case. The following indications exist for the intermediate formation of Fe$_3$(CO)$_{12}$:

- on the average 3 Fe(CO)$_5$ are adsorbed per supercage;
- the average stoichiometry after the first reaction step with HY is Fe(CO)$_4$;
- bands around 1880 cm$^{-1}$ which is in the region for bridged CO are also observed for Fe$_3$(CO)$_{12}$ in Ar matrix (ref. 19) or in KBr pellets and in solution (ref. 12). The previously observed bands at 1760/90 cm$^{-1}$ alternatively can be assigned to surface carbonate (ref. 20,21).

With Fe(CO)$_5$/NaY, the broad band around 1895 cm$^{-1}$ is the dominant species before reaction is complete.

In general, LFe(CO)$_x$ species exhibit a decrease of CO stretching frequency with decreasing $x$, if L is a set of the corresponding number of electron donor ligands or an inert matrix (ref. 15,22-24). The present IR results for Fe(CO)$_5$/NaY can be understood in the same way, indicating a low CO coordination.
number of the last generated intermediates.

The lower thermal stability of the HY adduct is explained by weaker $\pi$-back-bonding towards the CO ligands due to the increased electron deficiency of the iron clusters. The effect of acidity on the metal-CO bond was also observed with PdHY zeolites (ref. 25).

The HY-hydroxyl groups of the adduct are only partially restored after decomposition, indicating the consumption of protons according to

$$\text{Fe(CO)}_5 + 2\text{H}^+ \rightarrow \text{H}_2 + \text{Fe(II)} + 5\text{CO} \quad (\text{ref. 12})$$
The portion of oxidized iron, taking into account an initial proton content of 50 $H^+$/U.C., may therefore be estimated to be ca. 25 % of the iron loading.

CONCLUSION

The present work shows that the iron carbonyl at 20°C is strongly adsorbed until the supercages of the zeolites are saturated with three molecules on the average. The Fe(CO)$_3$ molecule remains intact on adsorption and is encaged in the zeolite with restricted mobility.

In HY, a hydrogen bond of moderate strength is formed with the supercage hydroxyls, which are completely involved in this process. Thermal decomposition in helium of the adducts leads to distinct CO-Fe fragments of different composition. Finally a reproducible iron loading of 10.5 ± 0.5 wt % is obtained. New evidence is found for the intermediate generation of Fe$_3$(CO)$_{12}$ during the decomposition in HY.

From the partly reversible hydroxyl interaction with the complex in HY, it is estimated that about one quart of the iron is oxidized during decomposition. Thermal decomposition is a slow reaction which goes to completion already between 70 and 90°C. It proceeds faster in case of HY due to acidic destabilization of the Fe-CO bond.

Work is in progress to determine the parameters influencing the particle size and catalytic properties of these zeolite supported iron clusters.

ACKNOWLEDGEMENTS

The technical assistance of Hugo Leeman is highly appreciated. One of us (T.B.) is indebted to the DAAD (Deutscher Akademischer Austauschdienst) and the belgian "Ministerie van Nationale Opvoeding en Nederlandse Cultuur" for a grant. P.A.J. acknowledges permanent research position as "Onderzoeksleider" from the Belgian Science Foundation (N.F.W.O.-F.N.R.S.). Financial support from the same institution and from the belgian government (Geconcerteerde Actie Catalyse, Diensten Wetenschapsbeleid) is gratefully acknowledged.

REFERENCES