Assessment of the location of Pt nanoparticles in Pt/zeolite Y/Al2O3 composite catalysts

Citation for published version (APA):

Document license:
CC BY-NC

DOI:
10.1002/cctc.201901617

Document status and date:
Published: 18/01/2020

Document Version:
Publisher’s PDF, also known as Version of Record (includes final page, issue and volume numbers)

Please check the document version of this publication:

• A submitted manuscript is the version of the article upon submission and before peer-review. There can be important differences between the submitted version and the official published version of record. People interested in the research are advised to contact the author for the final version of the publication, or visit the DOI to the publisher’s website.
• The final author version and the galley proof are versions of the publication after peer review.
• The final published version features the final layout of the paper including the volume, issue and page numbers.

Link to publication

General rights
Copyright and moral rights for the publications made accessible in the public portal are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights.

• Users may download and print one copy of any publication from the public portal for the purpose of private study or research.
• You may not further distribute the material or use it for any profit-making activity or commercial gain
• You may freely distribute the URL identifying the publication in the public portal.

If the publication is distributed under the terms of Article 25fa of the Dutch Copyright Act, indicated by the “Taverne” license above, please follow below link for the End User Agreement:
www.tue.nl/taverne

Take down policy
If you believe that this document breaches copyright please contact us at:
openaccess@tue.nl
providing details and we will investigate your claim.

Download date: 03. Apr. 2020
Assessment of the Location of Pt Nanoparticles in Pt/zeolite Y/γ-Al₂O₃ Composite Catalysts


The location of Pt nanoparticles was studied in Pt/zeolite Y/γ-Al₂O₃ composite catalysts prepared by H₂PtCl₆·6H₂O (CPA) or Pt (NH₃)₂(NO₃)₂ (PTA) as Pt precursors. The aim of this study is to validate findings from Transmission Electron Microscopy (TEM) by using characterization techniques that sample larger amounts of catalyst per measurement. Quantitative X-ray Photoelectron Spectroscopy (XPS) showed that the catalyst prepared with CPA led to a significantly higher Pt/Al atomic ratio than the catalyst prepared with PTA confirming that the 1-2 nm sized Pt nanoparticles in the former catalyst were located on the open and mesoporous γ-Al₂O₃ component, whereas they were located in the micropores of zeolite Y in the latter. By using infrared spectroscopy, a shift in the absorption band maximum of CO chemisorbed on Pt nanoparticles was observed, which can be attributed to a difference in electronic properties depending on the support of the Pt nanoparticles. Finally, model hydrogenation experiments were performed using β-phenylcinnamaldehyde, a reactant molecule with low diffusivity in zeolite Y micropores, resulting in a 5 times higher activity for the catalyst prepared by CPA compared to PTA. The combined use of these characterization techniques allow us to draw more robust conclusions on the ability to control the location of Pt nanoparticles by using either CPA or PTA as precursors in zeolite/γ-Al₂O₃ composite catalyst materials.

Introduction

The structural properties of supported metal catalysts, e.g.: metal loading, metal nanoparticle size and their distribution across the catalyst support, as well as the support acidity and porosity, strongly affect their selectivity, activity and stability in catalytic reactions.[1–3] Bifunctional catalysts with zeolite acid sites and metal sites are a special class of solid catalysts that are used for the conversion of crude oil fractions or renewable hydrocarbon feedstocks into fuels and chemicals.[4–6] For optimal catalytic performance of these catalysts, a close intimacy between the metal and acid sites is essential.[7–9] When metal nanoparticles, responsible for dehydrogenation-dehydrogenation, are located inside zeolite micropores, a ‘closest’ intimacy with the zeolite acid sites, responsible for isomerization and cracking, is ensured. On the other hand, locating metal nanoparticles inside zeolite micropores can limit their accessibility for larger hydrocarbon reactants that have low diffusivity through such small pores.[10,11]

Due to the strong impact of metal nanoparticle location on the catalytic performance, various synthetic approaches were developed over the last decades to improve control over location of metal nanoparticles within (micro)porous supports.[12–14] For example, Kim et al. were able to control the location of Pt nanoparticles by using [Pt(NH₃)₂Cl₂]²⁺ (aq) to ion-exchange with zeolite protons to obtain Pt nanoparticles inside ZSM-5 micropores, whereas impregnation with previously synthesized colloidal particles (~1.4 nm in diameter) was used to deposit Pt nanoparticles on the outer surface of the ZSM-5 crystallites (micropore width: 0.56 nm).[15] Several papers report methods to locate metal clusters within zeolite micropores by adding stabilized metal colloids or metal complexes before complete crystallization of the zeolites.[16–19] Alternatively, applying specific thermal treatments have been used to control the distribution of metal nanoparticles in porous supports.[20]

In these and many other studies of supported metal catalysts, TEM is one of the most commonly used techniques to investigate the location of metal nanoparticles on the support.[21–24] However, the severe limitations of TEM are the low amount of catalyst that is sampled per measurement, and the projection of a catalyst’s three dimensional structure in a two dimensional image. The latter issue can be overcome by the use of Electron tomography (or 3D TEM) that has been successfully applied to establish the location of metal particles within e.g. zeolite micropores.[22,24–26] Aside from TEM, several other characterization techniques have been sporadically employed to try to directly or indirectly determine the location

[a] J. Oenema, Dr. J. Zečević, Prof. K. P. de Jong
Inorganic Chemistry and Catalysis
Debye Institute for Nanomaterials Science
Utrecht University
Universiteitsweg 99
Utrecht 3584 CG (The Netherlands)
E-mail: J.Zecevic@uu.nl
K.P.dejong@uu.nl

[b] Dr. J. P. Hofmann, Prof. E. J. M. Hensen
Laboratory for Inorganic Materials and Catalysis
Department of Chemical Engineering and Chemistry
Eindhoven University of Technology
P.O. Box 513
Eindhoven 5600 MB (The Netherlands)

Supporting information for this article is available on the WWW under https://doi.org/10.1002/cctc.201901617
©2019 The Authors. Published by Wiley-VCH Verlag GmbH & Co. KGaA, Germany. This is an open access article under the terms of the Creative Commons Attribution Non-Commercial License, which permits use, distribution and reproduction in any medium, provided the original work is properly cited and is not used for commercial purposes.
of the metal nanoparticles on a support, including catalysis,[16–19,27–29] CO infrared spectroscopy,[18,40] SAXS,[23] XAS,[33] and XPS.[27,28,33]

Zeolite based catalysts used in industrial processes are typically shaped as millimeter sized bodies using a porous metal oxide binder, and may consist of additional components to further improve catalyst performance.[22–3] While academic catalyst research focuses often on structurally-simple model catalysts, a number of recent publications focuses on more practical catalysts and the interplay of all the structural features that determines the catalytic performance.[34,35] In a recent study, industrially relevant zeolite Y/γ-Al₂O₃ extrudates were used as support for Pt nanoparticles, located either inside zeolite Y or on the γ-Al₂O₃ binder.[35] The resulting bifunctional catalysts were investigated by using a heavy hydrocarbon feedstock, and remarkable beneficial effects on product selectivity were observed for the catalyst with Pt nanoparticles on the γ-Al₂O₃ binder compared to the catalysts with Pt nanoparticles inside zeolite Y. TEM was used to investigate the location of Pt nanoparticles in the zeolite Y/γ-Al₂O₃ extrudates and to overcome the challenge of overlapping features in two dimensional TEM images, authors used ultramicroscopy to section the catalysts into 70 nm thick slices. Besides that, an electron tomography study was used to study the location of Pt nanoparticles. However, both of these approaches suffer from sampling issues, with an estimated amount of sample being imaged in the order of 10⁻¹⁴ g. In studies of supported metal catalysts, synthesis-originated structural heterogeneities, such as: variation in local metal loading or metal nanoparticle size and location are regularly observed, either at the nanoscale between individual support particles[24,36] up to the microscale between individual support grains.[37] It is therefore desirable to validate results of characterization techniques, especially those that sample low amounts of catalysts per measurement, with complementary characterization techniques. In this study, the location of Pt nanoparticles in Pt/zeolite Y/γ-Al₂O₃ catalysts is assessed using a combination of commonly used laboratory techniques (quantitative XPS, CO infrared spectroscopy and catalysis) that sample significantly more catalyst than can be achieved by TEM analysis (Table 1, SI section 1.1, Table S1). The presented methods for assessment of the location of metal nanoparticles could be adopted for a wider range of catalysts, in particular catalyst with bimodal porosities, for example, to study the migration of metal nanoparticles during catalysis.[38]

**Results and Discussion**

Pt/zeolite Y/γ-Al₂O₃ catalysts were prepared with the aim to generate Pt nanoparticles onto either the zeolite Y or the γ-Al₂O₃ component of a zeolite Y/γ-Al₂O₃ composite support (50/50 wt.), using a previously reported methodology.[25] H₃PtCl₆·6H₂O or Pt(NH₃)₄(NO₃)₂ precursors dissolved in water yield the anionic [PtCl₆]⁶⁻ (aq) complex or the cationic [Pt(NH₃)₄]²⁺ (aq) complex, respectively, whereafter these solutions were added to an aqueous suspension of the zeolite Y/γ-Al₂O₃ composite support. In acidic conditions (pH~3.0), [PtCl₆]⁶⁻ (aq) adsorbs on the positively charged γ-Al₂O₃ surface (pzc~8.5) support due to electrostatic attraction, while interactions with the zeolite Y component are minimal.[39,40] On the other hand, the [Pt(NH₃)₄]²⁺ (aq) complex ion-exchanges with protons of the zeolite Y component, which is rather independent of pH.[41] The ion-exchange was performed at mildly acidic conditions (pH~5.0) whereby γ-Al₂O₃ is slightly positively charged to restrict adsorption of the cationic Pt complex. After Pt deposition, samples were dried and calcined/reduced to convert the adsorbed Pt complexes into Pt nanoparticles of 1–2 nm in diameter. The properties of the prepared catalysts are provided in Table 2.

High Angle Annular Dark Field Scanning Transmission Electron Microscopy (HAADF-STEM) analysis of 70 nm thick micromtombed sections of the samples provided local information about the size and location of Pt nanoparticles. It was observed that in the Pt–Y/A–Cl catalyst Pt nanoparticles were mostly present on γ-Al₂O₃ (Figure 1a), whose highly irregular porous structure of aggregated ~10 nm long platelets can be easily distinguished from large (200–1000 nm) mesoporous zeolite crystals. In Pt–Y/A–NH₃ (Figure 1b), Pt nanoparticles were almost exclusively present in the zeolite Y component. Previous research using electron tomography has shown that Pt nanoparticles in zeolite Y are located inside zeolite crystalline domains even though their size exceeds that of the zeolite micropores.[24] To further aid distinguishing the zeolite Y from the γ-Al₂O₃, HAADF-STEM was combined with EDX elemental mapping as can be observed in Figure S1.

---

**Table 1.** Employed characterization techniques and the respective amount of catalyst that is sampled per measurement, estimated using the parameters employed in this study.

<table>
<thead>
<tr>
<th>Characterization technique</th>
<th>Amount [g]</th>
</tr>
</thead>
<tbody>
<tr>
<td>TEM</td>
<td>~ 10⁻¹⁴</td>
</tr>
<tr>
<td>Quantitative XPS</td>
<td>~ 10⁻¹</td>
</tr>
<tr>
<td>CO infrared spectroscopy</td>
<td>~ 10⁻²</td>
</tr>
<tr>
<td>Catalysis</td>
<td>5·10⁻²</td>
</tr>
</tbody>
</table>

**Table 2.** Properties of the catalysts used for XPS and HAADF-STEM measurements. Pt weight loadings were determined by ICP whereas Pt nanoparticle sizes were determined from HAADF-STEM images.

<table>
<thead>
<tr>
<th>Sample</th>
<th>Pt precursor</th>
<th>Pt adsorption mechanism</th>
<th>Pt wt. loading [%]</th>
<th>Pt nanoparticle size [nm]</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pt–Y/A–NH₃</td>
<td>Pt(NH₃)₄(NO₃)₂</td>
<td>Ion-exchange</td>
<td>0.5[a]</td>
<td>1.4 ± 0.4</td>
</tr>
<tr>
<td>Pt–Y/A–Cl</td>
<td>H₃PtCl₆·6H₂O</td>
<td>Strong electrostatic adsorption</td>
<td>0.5[a]</td>
<td>1.8 ± 0.4</td>
</tr>
</tbody>
</table>

[a] Pt wt. loading indicated corresponds to catalysts characterized with HAADF-STEM and quantitative XPS. Catalysts wt. loadings used for CO infrared spectroscopy and catalysis are between 0.3–0.4 wt.% Pt.
In both composite catalysts, a small number of Pt nanoparticles close to the interface between γ-Al₂O₃ and zeolite Y appeared to be located on the ‘opposite’ component, as can be evidenced in Figure S2. In such cases, electron tomography could be used to determine the exact location of individual Pt nanoparticles. However, considering the low amount of catalyst sampled in a TEM measurement, it is desirable to investigate the severity of these heterogeneities (Pt nanoparticles located on the ‘opposite’ component) within larger amounts of the Pt/zeolite Y/γ-Al₂O₃ composite catalysts.

Quantitative XPS

XPS can provide quantitative information about atomic composition of the outer few nanometres of solid samples. It has been used by Wang et al. to determine if Pd nanoparticles were identified by their different morphology and are indicated in the images with red and green, respectively. 1–2 nm Pt nanoparticles can be identified as bright white dots, almost exclusively present in the γ-Al₂O₃ for Pt–Y/A–Cl and almost exclusively present in zeolite Y for Pt–Y/A–NH₃.

Because the thickness of the EMFP in Al₂O₃ (kinetic energy: 1410–1415 eV), this implies that every part of the sample has equal probability of being sampled and thus, for 1–2 nm sized Pt nanoparticles supported on γ-Al₂O₃, the XPS signal is an average of the entire sample. The zeolite Y crystallites are between 200 and 1000 nm in size while the EMFP for zeolite Y is approximately 4 nm (kinetic energy: 1410–1415 eV). As a consequence, only the outer layer of zeolite Y crystallites is studied, with a thickness determined by the EMFP, whereas the part below this layer does not contribute to the XPS signal (Figure 2). For more details about the XPS measurements and the approximation of EMFP in the samples, we refer to SI section 1.2. To assess the location of Pt nanoparticles in Pt–Y/A–Cl and Pt–Y/A–NH₃, samples were used with similar Pt weight loading and Pt particle size (Table S2).

XPS spectra of the fitted Pt 4f core lines can be observed in Figure 3, while for the full regional spectra we refer to Figure S5.

Table 3 shows the surface atomic ratios obtained by XPS, obtained after correction with atomic sensitivity factors, while bulk atomic ratios based on ICP analysis for Pt and manufacturers specifications for Si and Al. Furthermore, physical mixtures of Pt–Y–NH₃ with γ-Al₂O₃ (Pt–Y–NH₃ + A) and Pt–A–Cl

<table>
<thead>
<tr>
<th>Sample</th>
<th>Bulk Si/Al [at/at]</th>
<th>Pt/Al [at/at]</th>
<th>Surface (XPS) Si/Al [at/at]</th>
<th>Pt/Al [at/at]</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pt–Y/A–NH₃</td>
<td>0.80</td>
<td>0.0024</td>
<td>0.18</td>
<td>0.0003</td>
</tr>
<tr>
<td>Pt–Y/A–Cl</td>
<td>0.80</td>
<td>0.0025</td>
<td>0.18</td>
<td>0.0021</td>
</tr>
<tr>
<td>Pt–Y–NH₃ + A</td>
<td>0.78</td>
<td>0.0024</td>
<td>0.79</td>
<td>0.0001</td>
</tr>
<tr>
<td>Pt–A–Cl + Y</td>
<td>0.69</td>
<td>0.0025</td>
<td>0.83</td>
<td>0.0009</td>
</tr>
</tbody>
</table>

Figure 1. HAADF-STEM images of microtomed sections of Pt–Y/A–Cl (a) and Pt–Y/A–NH₃ (b). The γ-Al₂O₃ and zeolite Y components of the samples can be identified by their different morphology and are indicated in the images with red and green, respectively. 1–2 nm Pt nanoparticles can be identified as bright white dots, almost exclusively present in the γ-Al₂O₃ for Pt–Y/A–Cl and almost exclusively present in zeolite Y for Pt–Y/A–NH₃.
with zeolite Y (Pt–A–Cl + Y) were measured, that have similar bulk composition to the studied Pt/zeolite Y/γ-Al₂O₃ catalysts. For other relevant reference samples that were used for the fitting and standard deviations in the XPS measurements, we refer to Table S2.

The Si/Al surface ratios obtained by XPS were lower for the Pt/zeolite Y/γ-Al₂O₃ composite catalysts (Pt–Y/A–Cl and Pt–Y/A–NH₃) than their bulk Si/Al ratios, that can be attributed to coating of zeolite Y crystallites by γ-Al₂O₃ platelets thus attenuating the XPS signal of Si, as was also evidenced by TEM imaging (Figure S6). Another factor that could contribute to the higher aluminum concentration as observed by XPS, is the preferential breaking at void-rich γ-Al₂O₃ domains during the grinding prior XPS measurements, whereas the more dense zeolite Y crystals stay intact.

The surface Pt/Al ratio reported in Table 3 for Pt–Y/A–Cl is by a factor 7 higher than for Pt–Y/A–NH₃, while a similar difference in Pt/Al ratio is observed for physical mixtures. For Pt–Y/A–NH₃ and Pt–Y–NH₃, we expect that the majority of Pt nanoparticles was located in zeolite Y ‘underneath’ the outer surface layer invisible to the XPS signal, whereas for Pt–Y/A–Cl the Pt nanoparticles are located on the γ-Al₂O₃ component where they can contribute to the XPS signal. When the zeolite Y crystallites are approximated as cubes with side
length L (Figure 2), we anticipate that the difference in the XPS Pt/Al ratio depending on Pt nanoparticle location inside versus outside the zeolite crystals can be described by equation (1):

$$\left(\frac{I_{Pt}}{I_{Al}}\right)_{Pt \text{ in zeolite } Y} = \left(\frac{I_{Pt}}{I_{Al}}\right)_{Pt \text{ on zeolite } Y} \times \left(1 - \frac{2\gamma_{Pt/Al \text{ Y}}}{L}\right)$$  (1)

Wherein \(I_{Pt}\) and \(I_{Al}\) correspond to the Pt and Al peak areas, respectively, after correction for atomic sensitivity factors. \(\gamma_{Pt/Al \text{ Y}}\) is the EMFP of Pt 4f/Al 2p photoelectrons in zeolite Y. Assuming a uniform distribution of crystallites between 200–1000 nm and an EMFP of 4 nm, this should correspond to a difference of a factor \(\sim 50\), that is significantly larger difference than is observed here. A complicating factor for the studied composite catalysts, is that Pt nanoparticles outside zeolite Y are located on the \(\gamma\)-AlO\(_3\) binder, with a significantly higher specific surface area of 314 m\(^2\)/g compared to the external surface of zeolite Y crystallites of \(\sim 8\) m\(^2\)/g\(^{[46]}\) leading to a lower surface coverage of Pt nanoparticles. This results in a lower Pt/Al surface ratio as observed by XPS and, consequently, a smaller difference between the Pt/zeolite Y/\(\gamma\)-AlO\(_3\) composite catalysts based on the location of Pt nanoparticles.\(^{[46]}\) Due to the structural complexity of the composite catalysts, the differences in XPS Pt/Al ratio cannot be accurately described by the Pt nanoparticle inside versus outside zeolite Y approximation.

The quantification of the spectra of the Pt/zeolite Y/\(\gamma\)-AlO\(_3\) composite catalysts was challenging, due to the low intensity of the Pt 4f signals and overlap in the Pt 4f and Al 2p core levels, while the isolated Pt 4d core levels could not be used because of a too low signal/noise ratio. Nevertheless, regarding the standard deviations in Pt/Al ratios indicated in Table S2, there is a significant difference in the XPS Pt/Al ratio between Pt–Y/A–Cl and Pt–Y/A–NH\(_3\), that is also observed for the physical mixture reference samples. The uncertainties in the measurements as a consequence of low Pt weight loading and the overlap in Al 2p and Pt 4f do therefore not affect the outcome of this study.

The accurate quantification of XPS spectra, proved to be a useful tool that is able to differentiate the location of Pt nanoparticles in Pt/zeolite Y/\(\gamma\)-AlO\(_3\) composite catalysts. A further quantification of XPS signals calls for a more elaborate study (i.e. the intensity of the Pt 4f signal could be significantly enhanced in resonant photoemission experiments at a synchrotron) and modelling efforts. Alternatively, another metal could be used that does not overlap strongly with core lines of the support.

**CO Infrared Spectroscopy**

Fourier-transform infrared spectroscopy (FTIR), transmission IR spectra combined with CO chemisorption can provide information on the electronic properties of Pt nanoparticles. Typically, uniformly sized supported Pt particles, results in a CO absorption band between 2100–2000 cm\(^{-1}\).\(^{[46]}\) Due to metal-support interactions, the electronic properties of Pt particles (1–2 nm) are affected by the support and therefore CO infrared spectroscopy can be used to determine Pt nanoparticle location.\(^{[46,48]}\) Stakhiev et al. previously studied metal-support interactions using CO-FTIR to determine the location of Pt metal clusters in K–L zeolite and was able to distinguish between Pt particles inside or outside zeolite crystallites, whereas Liu et al. was able to describe differences in electronic properties between Pt/MCM-22 catalysts with different Pt dispersion and Pt nanoparticle location.\(^{[16,20]}\)

Here we have used samples with 1–2 nm Pt nanoparticles on a composite support and used references with Pt nanoparticles supported on either \(\gamma\)-AlO\(_3\) (Pt–A–Cl) or zeolite Y (Pt–Y/A–NH\(_3\)) to obtain the CO band maximum corresponding to Pt nanoparticles on these supports. The CO surface coverage of Pt nanoparticles and temperature influences the degree of dipolar coupling between CO molecules.\(^{[46,49]}\) All samples had similar Pt nanoparticle sizes that resulted in a similar CO coverage, although Pt nanoparticles located inside zeolite micropores could suffer from geometrical constraints making it more difficult for CO molecules to access the Pt sites. Figure 4 shows the CO FT-IR spectra of chemisorbed CO on the samples after subtraction of a background spectrum taken prior to CO adsorption. Both the Pt–Y/A–Cl and Pt–A–Cl samples, displayed in red in Figure 4 have a band maximum around \(\sim 2060\) cm\(^{-1}\) that has been reported for CO coordinated to Pt particles that do not have extended crystal faces on Pt/AlO\(_3\) at room temperature.\(^{[50]}\)

This is a clear indication that for Pt–Y/A–Cl monodisperse Pt nanoparticles are located on the \(\gamma\)-AlO\(_3\) binder. The band maximum of linearly coordinated CO on Pt–Y/A–NH\(_3\) is present at 2077 cm\(^{-1}\) shown in green in Figure 4b, in accordance with earlier studies reporting band maxima of 2083 cm\(^{-1}\) for Pt nanoparticles of \(<1\) nm supported on H–Y zeolite.\(^{[51]}\) The band maximum of Pt–Y/A–NH\(_3\) is present at 2070 cm\(^{-1}\), that is lower than for Pt–Y/NH\(_3\) but significantly higher than for Pt–A–Cl. This could indicate that Pt nanoparticles could be mainly located in the zeolite Y component with a small fraction located on the \(\gamma\)-AlO\(_3\) and the CO band maximum forms therefore an average of these populations of Pt nanoparticles. The downside of CO infrared spectroscopy is that differences between band maxima were small and a large overlap existed between relatively broad peaks. Overall, CO infrared spectroscopy data indicates Pt nanoparticles were located on \(\gamma\)-AlO\(_3\) for Pt–Y/A–Cl and predominantly located inside zeolite Y micropores for Pt–Y/A–NH\(_3\).

**Catalysis**

In a model reaction using 50 mg of catalyst, the catalytic activity of Pt/zeolite Y/\(\gamma\)-AlO\(_3\) composite catalysts is determined for hydrogenation of \(\beta\)-phenylcinnamaldehyde (RPCMA, 0.81 × 1.0 nm).\(^{[26]}\) Due to its relatively large size, it has low diffusivity in zeolite Y micropores (0.74 × 0.74 nm)\(^{[52]}\) and therefore can hardly reach Pt nanoparticles located inside the zeolite Y. If Pt nanoparticles are located on ‘accessible’ locations such as the zeolite Y mesopores or the external surface or the \(\gamma\)-AlO\(_3\) binder, this would lead to fast conversion of the reactant. Similar approaches have been reported, either based on gas phase or liquid phase reactions, to determine the location of metal nanoparticles in microporous catalysts.\(^{[16,19,27,28]}\)
Catalytic experiments were performed in a batch reactor loaded with equal amounts of BPCMA and catalyst. Figure 5 shows rate constants calculated from concentration profiles obtained by GC analysis of samples taken during the catalytic experiments. The references are used to study the catalytic activity of catalysts for which the location of Pt nanoparticles is known. For the concentration profiles that were used to calculate rate constants by fitting, we refer to Figure S7 and Table S3 and for product identification we refer to Figure S8,9.

The catalytic activity of Pt–Y/A–NH$_3$ and Pt–Y–NH$_3$ in this reaction was low, as could be expected for catalysts with Pt nanoparticles located inside zeolite micropores. Pt–Y/A–NH$_3$ had a factor 1.5 higher activity with respect to Pt–Y–NH$_3$ that can be contributed to a small fraction of Pt nanoparticles that are located on the γ-Al$_2$O$_3$ binder. Pt–Y/A–Cl on the other hand, had a factor 5 higher catalytic activity compared to Pt–Y/A–NH$_3$ that forms a strong indication that Pt nanoparticles are located on the γ-Al$_2$O$_3$ component for the former sample. The catalytic activity of Pt–A–Cl in the same reaction was also studied and was, as the activity of Pt–Y/A–Cl, significantly higher than Pt–Y/A–NH$_3$ and Pt–Y–NH$_3$. The difference in activity between Pt–Y/A–Cl and Pt–A–Cl could be attributed to Pt nanoparticles located in the zeolite, but more likely it is caused by undesired side reactions due to the reactivity of BPCMA on Brønsted acid sites. In the supporting information data is provided of a catalytic test performed with a mixture of Pt–A–Cl and zeolite Y powder, having a catalytic activity very similar to Pt–Y/A–NH$_3$. Overall, the catalytic experiments performed here proved to be a very useful tool in assessment of the Pt nanoparticle location in the Pt/zeolite Y/γ-Al$_2$O$_3$ composite catalysts.

**Conclusions**

The location of metal nanoparticles may have great impact on the performance of supported metal catalysts, but for structurally complex, practical catalysts used in industrial processes and consisting of multiple components, assessment of the location of metal particles presents great challenges.[25,53,54]

TEM has proven to be an accurate technique for determining the size and the location of metal nanoparticles in such catalysts, but a limitation is the amount of catalyst that can be sampled per measurement. In this work we explored quantitative XPS, CO infrared spectroscopy and catalysis as techniques that can offer information on metal location, while being able to sample larger amounts of catalysts. Quantitative XPS analysis of catalysts with equal Pt loading, equally sized Pt nanoparticles, but with Pt nanoparticles located either inside zeolite Y crystals or on the γ-Al$_2$O$_3$ binder of Pt/zeolite Y/γ-Al$_2$O$_3$ catalysts, lead to a significant differences in Pt/Al atomic ratios as a consequence of a different...
diffusivity in zeolite Y, large differences in activity were observed using the method described above, but using γ-Al2O3 in powder form as support. The γ-Al2O3 was obtained by calcination of HMPA pseudo-boehmite at 550 °C for 2 hours, using a ramp of 5 °C/min. Prior to the reduction the powder was pressed into a pellet, crushed and sieved into a sieve fraction of 0.2–0.5 mm in diameter, while gas flows were kept constant.

Elemental analysis. Pt elemental analysis was performed at Kolbe Mikroanalytisches Laboratorium, Oberhausen using an ICP-optical emission spectrometer (Perkin Elmer) after sample dissolution according to standard in-house procedures.

Transmission Electron Microscopy. High-Angle Annular Dark-Field Scanning Transmission Electron Microscopy (HAADF-STEM) imaging was performed on an FEI Talos F200X transmission electron microscope, equipped with a high-brightness field emission gun (X-PEG) and operated at 200 kV. For these analyses, catalysts were embedded in Epoxy resin, left to cure in air overnight at 60 °C, and cut to 70 nm sections using a Reichert-Jung Ultracut E ultramicrotome with Diatome Ultra 35° diamond knife. Sections were deposited on carbon-coated copper TEM grids.

X-ray Photoelectron Spectroscopy. XPS was performed using a Thermo Scientific K-Alpha spectrometer equipped with a monochromated Al Kα (hν = 1486.6 eV) X-ray source. The peak binding energies (BE) were calibrated against the sp2 C 1s peak of adventitious carbon at 284.6 eV. Sample charging was compensated by low energy e− and Ar+ ion flooding during measurements. Quantitative analysis of XPS data was performed using Casa XPS software based on high resolution regional spectra covering the Al 2 s, Si 2p, Pt 4f core levels in the BE range of 60–130 eV. The ratio between the Al 2p peak and Al 2 s peak was obtained from a reference Al2O3 sample and used during the fitting of the Pt containing samples to determine the area of the Al 2p in the overlapping Pt 4f/Al 2p region. The remaining area was then fitted to two components with asymmetric (metallic) line shape for Pt 4f7/2 and Pt 4f5/2 core levels, whereas the line shape and the FWHM were kept identical for both peaks of the spin-orbit split doublet (splitting was fixed to 3.35 eV). Atomic ratios were computed using atomic sensitivity factors. Values were averaged for two locations of the sample using a spot size of 400 μm. More information about the fitting of XPS spectra can be found in SI section 1.2.

Pt nanoparticle location. By using CO infrared spectroscopy, a systematic shift in CO band maximum between catalysts was observed depending on the location of Pt nanoparticles, while by performing model reactions using a reactant molecule with low diffusivity in zeolite Y, large differences in activity were observed between catalysts with Pt nanoparticle located either inside zeolite Y or on the more accessible γ-Al2O3 binder. Therefore, the results of this study show that Pt/zeolite Y/γ-Al2O3 composite catalysts can be prepared with great control over Pt nanoparticle location and without significant variations within samples, simply by using either CPA or PTA as Pt precursor during catalyst synthesis.

Considering the techniques used in this study (quantitative XPS, CO infrared spectroscopy, catalysis) are more common laboratory techniques compared to TEM, the methods used in this study, and combinations thereof, can be a valuable asset in screening practical catalysts. Furthermore, they can be used to ensure that the results obtained from TEM imaging are not compromised by local variations in structure.

Experimental Section

Catalyst Synthesis. Cylindrical extrudates consisting of 50 wt.% of mesoporous zeolite Y (Zeolyst CBV 760) and 50 wt.% of γ-Al2O3 (designated as Y/A) were obtained from Shell Projects and Technology. Mesoporous zeolite Y was purchased from Zeolyst (Si/Al = 30, CBV760) and HMPA pseudo-boehmite as precursor for γ-Al2O3 was obtained from Shell Projects and Technology. Extrudates were crushed and sieved to particles of 0.2–0.5 mm in diameter. Textural analysis of the Y/A support, as well as γ-Al2O3 and zeolite Y was performed using N2 physisorption at liquid nitrogen temperature on a Micromeritics TriStar II Plus, after drying the samples at 300 °C for 10 hours in an N2 flow. The isotherms and textural properties can be found in Figure S10–12 and Table S4.

Pt–Y/A–NH3 and Pt–Y–NH3 catalysts. 1 g of 0.2–0.5 mm Y/A particles were suspended in 300 ml Milli-Q water at room temperature and stirred for 1 hour, after which the pH of the suspension was 6.4. ~50 ml of aqueous solution containing 8.8 mg of Pt (NH3)2(NO3)2 (Sigma-Aldrich, 99.995% purity) was added dropwise to the suspension (aiming at nominal Pt loading of 0.5 wt.%), after which the pH dropped to 6.0. After addition of the Pt precursor, the suspension was stirred for another 3 hours, after which the pH dropped to 5.6. The suspension was filtered and washed with Milli-Q water and dried in air overnight at 120 °C (1 h, 5 °C/min). Pt–Y–NH3 reference catalyst was prepared using the method described above, but using only zeolite Y (CBV 760) in powder form as support. Prior to the calcination/reduction the powder was pressed into a pellet, crushed and sieved into a sieve fraction of 0.2–0.5 mm in diameter while gas flows were kept constant.

Pt–Y/A–Cl and Pt–A–Cl catalysts. A suspension containing 1 g of 0.2–0.5 mm Y/A particles in 300 ml Milli-Q water was stirred for 1 hour at room temperature, after which the pH of the suspension was lowered from 5.5 to 3.0 by adding a few drops of a 1 M HCl solution. ~30 ml of aqueous solution containing 13.5 mg of H2PtCl6·6H2O (Sigma-Aldrich, ~36 wt.% Pt) was added dropwise to the suspension (aiming at nominal Pt loading of 0.5 wt.%). After 3 hours of stirring (pH increased to 3.8), the suspension was filtered and washed with Milli-Q water, and dried in air overnight at 120 °C. Dried catalyst precursor was reduced in a flow of H2 for 3 hours at 500 °C, with a ramp of 5 °C/min (GHSV ~3300 h−1). Pt–A–Cl reference catalyst was prepared using the method described above, but using γ-Al2O3 in powder form as support. The γ-Al2O3 was obtained by calcination of HMPA pseudo-boehmite at 550 °C for 2 hours, using a ramp of 5 °C/min. Prior to the reduction the powder was pressed into a pellet, crushed and sieved into a sieve fraction of 0.2–0.5 mm in diameter, while gas flows were kept constant.

CO infrared spectroscopy. CO infrared spectroscopy experiments were performed in a Perkin Elmer 2000 FTIR using a transmission in-situ cell with CaF2 windows. Spectra were recorded with a resolution of 4 cm−1 by coaddition of 25 scans. For each measurement, a self-supporting pellet was made of ~10 mg of sample. The pellet was dried overnight at 120 °C (p < 10−5 mbar). Thereafter, the sample was reduced in a flow of H2 (Linde Gas, quality 5.0) at ~1 bar whilst heating up to 300 °C (1 h, 5 °C/min) followed by evacuation of the cell for 1 h. Then, the temperature was lowered to 50 °C and a reference spectrum was measured in vacuum (p < 10−1 mbar). A 10% CO/He gas mixture (Linde Gas, 99.998% purity) of 200 mbar was introduced in the cell for 30 min followed by evacuation for 30 min. After evacuation of gaseous CO (p < 10−5 mbar), a spectrum was measured of the sample with remaining chemisorbed CO. The final spectra were obtained by subtraction of the reference spectrum from the spectrum of the sample with chemisorbed CO.

Catalytic experiments. Hydrogenation of β-phenylcinamaldehyde (BPCMA, Sigma-Aldrich) was performed in stainless steel autoclaves with a total volume of 15 ml. 50 mg of Pt–Y/A catalysts (or 25 mg of Pt–A–Cl or Pt–Y–NH3 reference catalysts) with a sieve fraction of 0.2–0.5 mm, was suspended in 7 ml of 2-propanol (Merck, ≥ 99.5%) containing 200 mg of dissolved BPCMA and 100 μl of n-tetradecane.
Conflict of Interest

The authors declare no conflict of interest.

Keywords: Bifunctional catalysts  ·  Metal nanoparticle location  ·  Quantitative XPS  ·  CO infrared spectroscopy  ·  Catalysis

Acknowledgements

Mark Meijerink and Lars van der Wal (both UU) are acknowledged for TEM analysis of the microtomed samples. Nikos Nikolopoulos (UU) is acknowledged for performing N2 physisorption measurements. Fouad Saulimani (UU) is acknowledged for providing the training for CO infrared spectroscopy measurements and Pascal Wijten (UU) is acknowledged for the training for the autoclave setup and GC. Shell Projects and Technology is acknowledged for supplying Zeolite Y/γ-Al2O3 exudates and HMHPA pseudo-boehmite.