



# Catalytic activity of maghemite supported palladium catalyst in nitrobenzene hydrogenation

Viktória Hajdu<sup>1</sup> · Ádám Prekob<sup>1</sup> · Gábor Muránszky<sup>1</sup> · István Kocserha<sup>2</sup> · Zoltán Kónya<sup>3</sup> · Béla Fiser<sup>1,4</sup> · Béla Viskolcz<sup>1</sup> · László Vanyorek<sup>1</sup> 

Received: 29 October 2019 / Accepted: 31 December 2019 / Published online: 9 January 2020  
© The Author(s) 2020

## Abstract

A maghemite supported palladium catalyst was prepared and tested in nitrobenzene hydrogenation. The catalyst support was made by a newly developed combined technique, where sonochemical treatment and combustion have been used. As a first step, maghemite nanoparticles were synthesized. Iron(II) citrate was treated in polyethylene glycol by high-intensity ultrasound cavitation to get a homogeneous dispersion, then the product was combusted. The produced powder contained maghemite nanoparticles with 21.8 nm average particle size. In the second step of catalyst preparation, the magnetic nanoparticles were dispersed in the ethanolic solution of palladium(II) nitrate. The necessary energy for the reduction of Pd<sup>2+</sup> ions was achieved in the “hot spots” by acoustic cavitation, thus catalytically active palladium was formed. The prepared maghemite supported Pd catalyst have been tested in nitrobenzene hydrogenation at three different temperatures (283 K, 293 K and 303 K) and constant pressure (20 bar). At 293 K and 303 K, the conversion and selectivity of nitrobenzene was above 99% and 96%, respectively. However, the selectivity was only 73% at 273 K because the intermediate species (azoxybenzene and nitrosobenzene) have not been transformed to aniline. All in all, the prepared catalyst is successfully applied in nitrobenzene hydrogenation and easily separable from the reaction media.

**Keywords** Magnetic catalyst · Selectivity · Nitrobenzene · Aniline

## Introduction

Several different complex catalysts have been successfully applied in the hydrogenation of nitro groups, such as carbon (C), silica (SiO<sub>2</sub>) or alumina (Al<sub>2</sub>O<sub>3</sub>) supported Pd, Pt, Ru, Rh, Ni, Fe or bimetallic systems [1–13]. The easy handling and

✉ László Vanyorek  
kemviki@uni-miskolc.hu

Extended author information available on the last page of the article

separability are very important properties for the catalysts. These can be improved by introducing magnetic features (e.g. magnetic catalyst supports), which will allow the easy and efficient removal of the catalysts after the reactions. For this reason, magnetic systems have been widely used in various applications. Magnetite ( $\text{Fe}_3\text{O}_4$ )/silica composite catalyst was used for esterification of palmitic acid with methanol [14]. Pd, Pr–Cu and  $\text{Pr}_6\text{O}_{11}$  decorated  $\text{Fe}_3\text{O}_4/\text{SiO}_2$  catalyzed the reduction of 2,4-dinitrophenylhydrazine, 4-nitrophenol and chromium(VI) ions, Mizoroki–Heck coupling reaction, and the catalytic ozonation of acetochlor [15–17]. Magnetite/carbon support was applied in Suzuki–Miyaura cross-coupling of 4-iodotoluene and phenylboronic acid and aniline synthesis by palladium [18, 19]. By magnetite/alumina supported Pd catalyst the hydrogenation of nitrate in water and 4-nitrophenol can be achieved [20, 21]. Magnetic iron oxides can be combined with different layered double hydroxides ( $\text{Fe}_3\text{O}_4$ -LDH), complex magnesium silicates ( $\text{Fe}_3\text{O}_4$ -sepiolite) and hydroxyapatite ( $\gamma$ - $\text{Fe}_2\text{O}_3$ -HAP) to use as a support for Pd and these catalytic systems can be applied to catalyze the Heck reaction between iodobenzene and styrene, and the reduction of nitroarenes and nitrobenzene [22–24]. Magnetite itself is also a promising catalyst support as it was proved by the applicability of Ag/ $\text{Fe}_3\text{O}_4$ , Ag–Ni/ $\text{Fe}_3\text{O}_4$ , Pd/ $\text{Fe}_3\text{O}_4$  and Rh/ $\text{Fe}_3\text{O}_4$  systems in the synthesis of 3,4-dihydropyrimidinones, 2,4-dihydropyrano[2,3-*c*]pyrazoles, and the hydrogenation of soybean oil and nitroarenes [25–28]. The two main components of the catalysts mentioned above are the support and the catalytically active metal. The catalysts are prepared through several steps, including the activation of metal, within which metal (e.g. palladium ions) ions or their complex ions are reduced to the catalytically active form (e.g.  $\text{Pd}^0$ ). In the case of Pd ions, the activation (reduction) can be done on the supports in aqueous solution by molecular hydrogen (6 atm, 75 °C) or by using  $\text{NaBH}_4$  in ethanol but the ethylene glycol is also efficient [21, 24, 29].

In our work, a simplified reduction step was applied during the catalyst production (palladium(II) nitrate to  $\text{Pd}^0$ ) by applying alcohol and acoustic cavitation. The high energy of the ultrasonic treatment in liquids generates acoustic cavitation, which leads to the formation of micro vapor-bubbles. The collapse of the formed bubbles leads to „hot spots” where intense local heating (~ 5000 K), high pressure (~ 1000 atm), enormous heating and cooling rates (> 109 K/s) and liquid jet streams (~ 400 km/h) appear in a small volume [30]. The energy in the „hot spots” can cover the needs of the reduction of metal ions to metals in the presence of a reducing agent [31–36]. By using of ultrasonic cavitation, palladium nanoparticles were deposited on the surface of maghemite in methanol phase. Owing to the magnetic properties of the maghemite, this is a remarkable catalyst support in liquid phase hydrogenation because the catalyst easily separated from the reaction media by magnetic field.

## Experiment

### Materials

Iron(III) citrate hydrate ( $\text{FeC}_6\text{H}_5\text{O}_7 \cdot \text{H}_2\text{O}$ , PanReac AppliChem) as precursor and polyethylene glycol (PEG400, Sigma Aldrich) were applied for the synthesis of

maghemite. Palladium(II) nitrate dihydrate ( $\text{Pd}(\text{NO}_3)_2 \cdot 2\text{H}_2\text{O}$ , Merck) and absolute ethanol (VWR) was used to synthesize catalytically active palladium.

### Application of maghemite supported palladium catalyst

Maghemite nanoparticles, as catalyst supports were synthesized by a combustion method. 3.5 g iron(III) citrate hydrate was dispersed in 20 g polyethylene glycol (PEG 400, Sigma Aldrich) by using a Hielscher Ultrasound tip homogenizer. The iron precursor containing dispersion was heated up and burned at 500 °C in a calcining furnace for two hours.

The before-synthesized maghemite was applied for catalyst preparation by using a Hielscher Ultrasound tip homogenizer (UIP1000hDT). The palladium precursor (0.125 g) was solved in 50 ml abs. ethanol, and 1.00 g maghemite was added to the solution. The ethanolic dispersion was sonicated by using the homogenizer (115 W, 19.43 kHz) for 2 min. Then, the catalyst was removed from the dispersion with a Nd magnet, washed with ethanol, and dried at 105 °C overnight.

### Characterization techniques of the nanoparticles

Maghemite and palladium nanoparticles were examined by using high-resolution transmission electron microscopy (HRTEM, FEI Technai G2 electron microscope, 200 kV). The samples were prepared by dropping their aqueous suspension on 300 mesh copper grids (Ted Pella Inc.). The diameters of the nanoparticles were measured on the HRTEM images, based on the original scale bar by using the ImageJ software. X-ray diffraction (XRD) measurements were used to identify and quantify the crystalline phases, by applying a Rigaku Miniflex II diffractometer with Cu  $K_\alpha$  radiation source (30 kV, 15 mA). The palladium content was determined with a Varian 720 ES inductively coupled optical emission spectrometer (ICP-OES), by using a Merck Certipur ICP multi-element standard IV.

### Catalytic tests

The catalytic hydrogenation was carried out in a Büchi Uster Picoclave reactor, in a 200 ml stainless steel vessel with heating jacket. The hydrogen pressure was 20 bar and the reactions were carried at 283 K, 293 K and 323 K. Sampling took place after the beginning of hydrogenation at 5, 10, 15, 20, 30, 60, 120, 180, and 240 min. The initial concentration of nitrobenzene was 0.125 mol  $\text{dm}^{-3}$  in methanol. The total amount of the solution was 150 ml and 0.2 g catalyst was used during each test. Aniline formation was followed by applying Agilent 7890A gas chromatograph coupled with Agilent 5975C Mass Selective detector. Analytical standards (aniline, nitrobenzene, nitrosobenzene, azoxybenzene, dicyclohexylamine, o-toluidine, cyclohexylamine and n-methylaniline) were provided by Dr. Ehrenstorfer and Sigma Aldrich. The efficiency of the catalytic hydrogenation was compared by calculating the conversion (X%) of nitrobenzene based on the following equation (Eq. 1):

$$X\% = \frac{\text{consumed } n_{\text{nitrobenzene}}}{\text{initial } n_{\text{nitrobenzene}}} \times 100. \quad (1)$$

The selectivity (S%) of the catalyst was calculated as follows (Eq. 2):

$$S\% = \frac{n_{\text{aniline}}}{n_{\text{nitrobenzene}}} \times 100 \quad (2)$$

re  $n_{\text{aniline}}$  and  $n_{\text{nitrobenzene}}$  are the corresponding chemical amounts of the compounds.

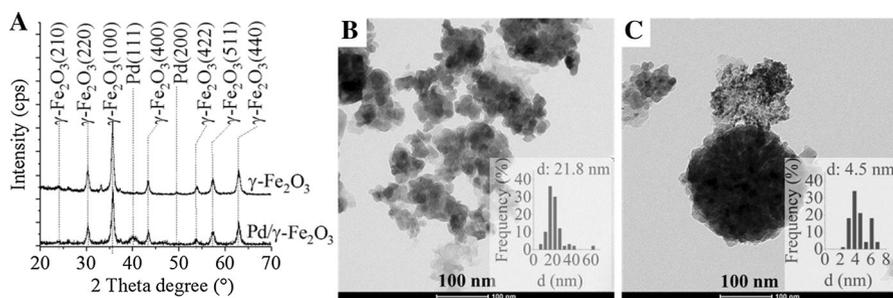
By assuming that the process is a first order reaction, based on the initial and measured nitrobenzene concentrations ( $c_0$  and  $c_k$ , mol/dm<sup>3</sup>), the reaction rate constant (k) was calculated at different temperatures by non-linear regression (Fig. 4) according to the following (Eq. 3):

$$c_k = c_0 * e^{-k*t} \quad (3)$$

## Results and discussion

### Surface morphology and phase composition of the catalyst

The reduction of palladium ions to elemental Pd have been confirmed by XRD measurements (Fig. 1a). Reflections at 40° and 48.8° 2θ degrees were identified on the XRD pattern, which are attributed to the Pd(111) and Pd(200) phases (Fig. 1a, red line). Other reflections were also identified such as the peaks at 24.1°, 30.3°, 35.7°, 43.3°, 54°, 57.3° and 63° 2θ degrees, which are assigned to the presence of (210), (220), (100), (400), (422), (511) and (400) planes of maghemite ( $\gamma\text{-Fe}_2\text{O}_3$ ) crystalline phase. The average size of the maghemite particles was found to be 21.8 nm (Fig. 1b). Palladium nanoparticles were deposited onto the surface of the maghemite crystals. The palladium deposition onto the surface of the maghemite led to the aggregation of the magnetic particles, the size of the nanocomposite aggregates are between 70–200 nm. The palladium particles on the maghemite aggregates



**Fig. 1** XRD pattern of the maghemite (blue line) and Pd/maghemite catalyst (red line) (a) HRTEM image and size distribution of maghemite (b) and Pd/maghemite (c)

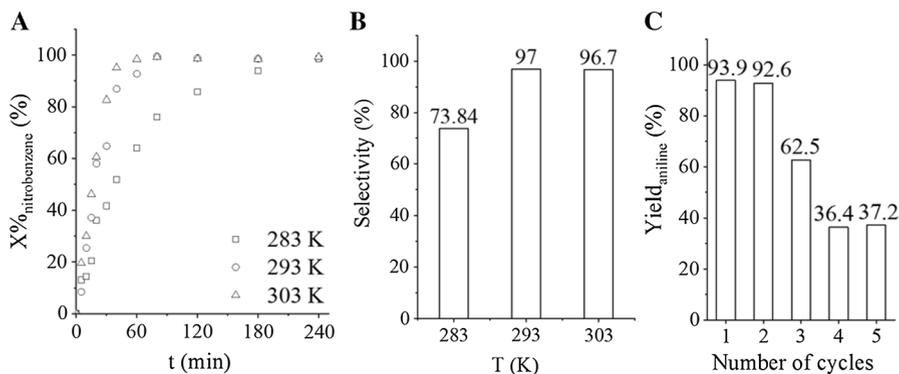
are smaller than 8 nm (the average size is 4.5 nm) (Fig. 1c). The formation of palladium nanoparticles can be explained by the reduction effect of the appearing  $\cdot\text{CH}_2\text{R}$  radicals. These reactive species generated by the ultrasonic treatment through the reaction of  $\cdot\text{OH}$  radicals and ethanol [37, 38].

### Catalytic activity of the magnetic Pd catalyst

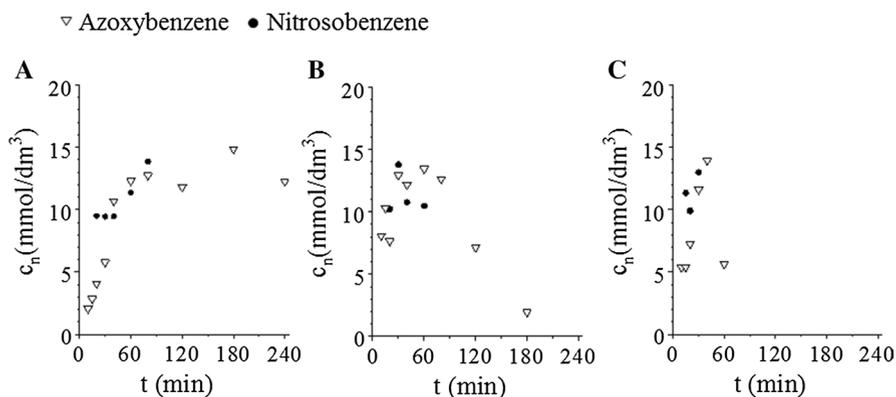
The prepared magnetic Pd catalyst was tested in nitrobenzene hydrogenation. The maximum conversions were reached after 80 min at 293 K and 303 K (Fig. 2a). At 283 K the reaction is slower, but the total amount of nitrobenzene was transformed to aniline. The aniline selectivity was high, 97% and 96.7% at 293 K and 303 K, respectively (Fig. 2b). The catalytic activity was tested through five cycles at 303 K and 20 bar hydrogen pressure, while the reaction time was 80 min. The catalyst was not regenerated between the cycles, only washed with methanol. The activity started to decrease from the third cycle, which indicates that the regeneration of the catalyst is necessary (Fig. 2c).

The selectivity was lower, only 73.8%, at 283 K which can be explained by the low reaction rate, and the persistence of the intermediates which are not converted to aniline (Fig. 3a). At 283 K, azoxybenzene and nitrosobenzene have been detected during the reactions, which indicates that, the hydrogenation process follows the Haber mechanism [38–42]. At higher temperatures the intermediates transformed to aniline (Fig. 3b and c). The catalyst was very selective towards the formation of aniline, by-products have not been detected. All in all, the prepared maghemite supported palladium catalyst at 303 K reaction temperature and 20 bar hydrogen pressure can be applied effectively for aniline synthesis.

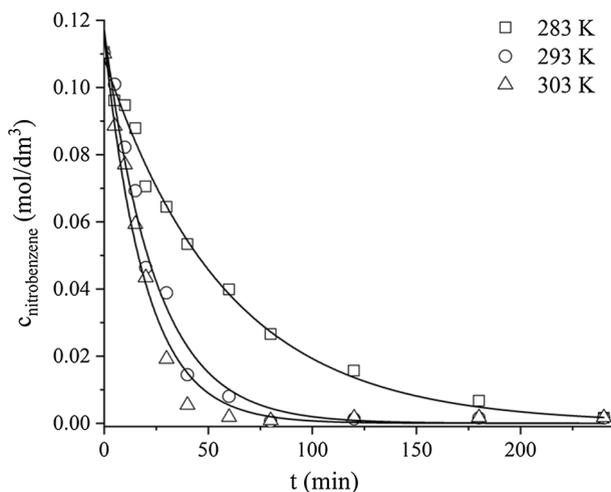
The reaction rate constants ( $k$ ) at different temperatures were calculated based on the measured nitrobenzene concentrations by using non-linear regression [43] (Fig. 4; Table 1).



**Fig. 2** Conversion of nitrobenzene vs time of hydrogenation (a) and aniline selectivity (b) at various temperatures (283, 293 and 303 K). Aniline yield vs number of cycles at 20 bar pressure and 303 K, after 80 min of hydrogenation



**Fig. 3** Concentration of the intermediates vs time of hydrogenation, at 283 K (a), 293 K (b) and 303 K(c) at 20 bar pressure



**Fig. 4** Concentration of nitrobenzene vs time of hydrogenation

**Table 1** Reaction rate constants of nitrobenzene hydrogenation

Temperature (K)	283	293	303
Reaction rate constant ( $s^{-1}$ )	$1.73 \times 10^{-2}$	$4.09 \times 10^{-2}$	$5.10 \times 10^{-2}$
SD	$7.76 \times 10^{-4}$	$2.99 \times 10^{-3}$	$3.83 \times 10^{-3}$

## Conclusion

Maghemite supported palladium catalyst was prepared. The maghemite catalyst support was made by a newly developed combined technique, where sonochemical

treatment and combustion have been used. This procedure leads to nanoparticles with smaller crystalline size (21.8 nm) and high adsorption capability. The catalyst is in an active form immediately after the production of the Pd/maghemite nanocomposite, as the sonochemical treatment initiated the involvement of the dispersion media in the reduction of palladium ions to elemental palladium particles ( $\text{Pd}^0$ ). In this sense, the catalyst does not require further post-treatments, and it does not need to be reduced under a hydrogen atmosphere, therefore the catalyst preparation method is simplified. The synthesized magnetic catalyst was efficiently applied in nitrobenzene hydrogenation at 293 K and 303 K and the conversion was more than 99% in both case. The catalyst was selective towards aniline, and the selectivity was 97.0% and 96.7% at 293 K and 303 K, respectively. By-products were not detected during the reaction. All in all, a simple method has been designed for magnetic catalyst production. The achieved catalytic system is easily separable from the reaction media, thanks to its magnetic property and successfully applicable in nitrobenzene hydrogenation.

**Acknowledgements** Open access funding provided by University of Miskolc (ME). This research was supported by the European Union and the Hungarian State, co-financed by the European Regional Development Fund in the framework of the GINOP-2.3.4–15-2016–00004 project, aimed to promote the cooperation between the higher education and the industry. The EFOP-3.6.1-16-2016-00014 project is also gratefully acknowledged due to support our work.

## Compliance with ethical standards

**Conflict of interest** On behalf of all authors, the corresponding author states that there is no conflict of interest.

**Open Access** This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article's Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article's Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit <http://creativecommons.org/licenses/by/4.0/>.

## References

1. Zhao F, Zhang R, Chatterjee M et al (2004) Hydrogenation of nitrobenzene with supported transition metal catalysts in supercritical carbon dioxide. *Adv Synth Catal* 346:661–668. <https://doi.org/10.1002/adsc.200303230>
2. Srikanth CS, Kumar VP, Viswanadham B et al (2015) Vapor phase hydrogenation of nitrobenzene to aniline over carbon supported ruthenium catalysts. *J Nanosci Nanotechnol* 15:5403–5409. <https://doi.org/10.1166/jnn.2015.9872>
3. Zhao Y, Li C-H, Yu Z-X et al (2007) Effect of microstructures of Pt catalysts supported on carbon nanotubes (CNTs) and activated carbon (AC) for nitrobenzene hydrogenation. *Mater Chem Phys* 103:225–229. <https://doi.org/10.1016/j.matchemphys.2007.02.045>
4. Wu S, Wen G, Zhong B et al (2014) Reduction of nitrobenzene catalyzed by carbon materials. *Chin J Catal* 35:914–921. [https://doi.org/10.1016/S1872-2067\(14\)60102-9](https://doi.org/10.1016/S1872-2067(14)60102-9)

- Chen P, Yang F, Kostka A, Xia W (2014) Interaction of cobalt nanoparticles with oxygen- and nitrogen-functionalized carbon nanotubes and impact on nitrobenzene hydrogenation catalysis. *ACS Catal* 4:1478–1486. <https://doi.org/10.1021/cs500173t>
- Collins DJ, Smith AD, Davis BH (1982) Hydrogenation of nitrobenzene over a nickel boride catalyst. *Ind Eng Chem Prod Res Dev* 21:279–281. <https://doi.org/10.1021/i300006a016>
- Hatziantoniou V, Andersson B, Schoon NH (1986) Mass transfer and selectivity in liquid-phase hydrogenation of nitro compounds in a monolithic catalyst reactor with segmented gas-liquid flow. *Ind Eng Chem Process Des Dev* 25:964–970. <https://doi.org/10.1021/i200035a021>
- Liao H-G, Xiao Y-J, Zhang H-K et al (2012) Hydrogenation of nitrocyclohexane to cyclohexanone oxime over Pd/CNT catalyst under mild conditions. *Catal Commun* 19:80–84. <https://doi.org/10.1016/J.CATCOM.2011.12.027>
- Li C-H, Yu Z-X, Yao K-F et al (2005) Nitrobenzene hydrogenation with carbon nanotube-supported platinum catalyst under mild conditions. *J Mol Catal A* 226:101–105. <https://doi.org/10.1016/j.molcata.2004.09.046>
- Dong B, Li Y, Ning X et al (2017) Trace iron impurities deactivate palladium supported on nitrogen-doped carbon nanotubes for nitrobenzene hydrogenation. *Appl Catal A* 545:54–63. <https://doi.org/10.1016/j.apcata.2017.07.035>
- Hao L, Li H, Hu Y et al (2014) Carbon nanotube-supported bimetallic Pt–Fe catalysts for nitrobenzene hydrogenation. *Micro Nano Lett* 9:97–99. <https://doi.org/10.1049/mnl.2013.0624>
- Zhao F, Ikushima Y, Arai M (2004) Hydrogenation of nitrobenzene with supported platinum catalysts in supercritical carbon dioxide: effects of pressure, solvent, and metal particle size. *J Catal* 224:479–483. <https://doi.org/10.1016/j.jcat.2004.01.003>
- Kim Y, Ma R, Reddy DA, Kim TK (2015) Liquid-phase pulsed laser ablation synthesis of graphitized carbon-encapsulated palladium core–shell nanospheres for catalytic reduction of nitrobenzene to aniline. *Appl Surf Sci* 357:2112–2120. <https://doi.org/10.1016/J.APSUSC.2015.09.193>
- dos Santos-Durndell VC, Peruzzolo TM, Ucoski GM et al (2018) Magnetically recyclable nanocatalysts based on magnetite: an environmentally friendly and recyclable catalyst for esterification reactions. *Biofuel Res J* 5:806–812. <https://doi.org/10.18331/BRJ2018.5.2.4>
- Nasrollahzadeh M, Issaabadi Z, Safari R (2019) Synthesis, characterization and application of Fe<sub>3</sub>O<sub>4</sub>@SiO<sub>2</sub> nanoparticles supported palladium(II) complex as a magnetically catalyst for the reduction of 2,4-dinitrophenylhydrazine, 4-nitrophenol and chromium(VI): a combined theoretical (DFT) and experimental study. *Sep Purif Technol* 209:136–144. <https://doi.org/10.1016/j.seppur.2018.07.022>
- Yavari I, Mobaraki A, Hosseinzadeh Z, Sakhaee N (2019) Copper-catalyzed Mizoroki-Heck coupling reaction using an efficient and magnetically reusable Fe<sub>3</sub>O<sub>4</sub>@SiO<sub>2</sub>@PrNCu catalyst. *J Organomet Chem* 897:236–246. <https://doi.org/10.1016/J.JORGANOCHEM.2019.06.029>
- Wang J, Lou Y, Zhuang X et al (2018) Magnetic Pr<sub>6</sub>O<sub>11</sub>/SiO<sub>2</sub>@Fe<sub>3</sub>O<sub>4</sub> particles as the heterogeneous catalyst for the catalytic ozonation of acetochlor: Performance and aquatic toxicity. *Sep Purif Technol* 197:63–69. <https://doi.org/10.1016/j.seppur.2017.12.052>
- Desmecht A, Pennetreau F, L'hoost A et al (2019) Preparation of magnetically recoverable carbon nanotube-supported Pd(II) catalyst. *Catal Today* 334:24–29. <https://doi.org/10.1016/j.catto.2019.02.057>
- Mei N, Liu B (2016) Pd nanoparticles supported on Fe<sub>3</sub>O<sub>4</sub>@C: an effective heterogeneous catalyst for the transfer hydrogenation of nitro compounds into amines. *Int J Hydrogen Energy* 41:17960–17966. <https://doi.org/10.1016/j.ijhydene.2016.07.229>
- Rahimi E, Sajednia G, Baghdadi M, Karbassi A (2018) Catalytic chemical reduction of nitrate from simulated groundwater using hydrogen radical produced on the surface of palladium catalyst supported on the magnetic alumina nanoparticles. *J Environ Chem Eng* 6:5249–5258. <https://doi.org/10.1016/j.jece.2018.08.026>
- Gil J, Ferreira LF, Silva VC et al (2019) Facile fabrication of functionalized core-shell Fe<sub>3</sub>O<sub>4</sub>@SiO<sub>2</sub>@Pd microspheres by urea-assisted hydrothermal route and their application in the reduction of nitro compounds. *Environ Nanotechnol Monit Manag* 11:100220. <https://doi.org/10.1016/j.enmm.2019.100220>
- Wang G, Ling Y, Wheeler DA et al (2011) Facile synthesis of highly photoactive  $\alpha$ -Fe<sub>2</sub>O<sub>3</sub>-based films for water oxidation. *Nano Lett* 11:3503–3509. <https://doi.org/10.1021/nl202316j>
- Ghonchepour E, Islami MR, Bananezhad B et al (2019) Synthesis of recoverable palladium composite as an efficient catalyst for the reduction of nitroarene compounds and Suzuki cross-coupling

- reactions using sepiolite clay and magnetic nanoparticles ( $\text{Fe}_3\text{O}_4$ @sepiolite- $\text{Pd}^{2+}$ ). *Comptes Rendus Chim* 22:84–95. <https://doi.org/10.1016/j.crci.2018.07.008>
24. Jiang L, Zhang Z (2016) Efficient transfer hydrogenation of nitro compounds over a magnetic palladium catalyst. *Int J Hydrogen Energy* 41:22983–22990. <https://doi.org/10.1016/j.ijhydene.2016.09.182>
  25. Mirhashemi F, Ali Amrollahi M (2019) Decoration of  $\beta$ -CD on  $\text{Fe}_3\text{O}_4$ @Ag core-shell nanoparticles as a new magnetically recoverable and reusable catalyst for the synthesis of 3,4-dihydropyrimidinones and 2,4-dihydropyranol[2,3-*c*]pyrazoles in  $\text{H}_2\text{O}$ . *Inorganica Chim Acta* 486:568–575. <https://doi.org/10.1016/j.ica.2018.11.009>
  26. Wang L, Wang W, Wang Y et al (2018) Structural characteristics of a Ni–Ag magnetic catalyst and its properties in soybean oil hydrogenation. *Food Bioprod Process* 109:139–147. <https://doi.org/10.1016/j.fbp.2018.03.008>
  27. Kandathil V, Koley TS, Manjunatha K et al (2018) A new magnetically recyclable heterogeneous palladium(II) as a green catalyst for Suzuki-Miyaura cross-coupling and reduction of nitroarenes in aqueous medium at room temperature. *Inorganica Chim Acta* 478:195–210. <https://doi.org/10.1016/j.ica.2018.04.015>
  28. Shaikh MN, Helal A, Kalanthoden AN et al (2019) Sub-nanometric Rh decorated magnetic nanoparticles as reusable catalysts for nitroarene reduction in water. *Catal Commun* 119:134–138. <https://doi.org/10.1016/j.catcom.2018.09.002>
  29. Park J, Bae S (2019) Highly efficient and magnetically recyclable Pd catalyst supported by iron-rich fly ash@fly ash-derived  $\text{SiO}_2$  for reduction of *p*-nitrophenol. *J Hazard Mater* 371:72–82. <https://doi.org/10.1016/j.jhazmat.2019.02.105>
  30. Suslick KS (2000) Sonochemistry. In: Suslick KS (ed) *Kirk-Othmer encyclopedia of chemical technology*. Wiley, Hoboken.
  31. Qiu X-F, Zhu J-J (2003) Synthesis of palladium nanoparticles by a sonochemical method. *Chinese J Inorg Chem* 19:766–770
  32. Yu Y, Zhang QY, Li XG (2003) Reduction process of transition metal ions by zinc powder to prepare transition metal nanopowder. *Acta Phys Chim Sin* 19:436–440. <https://doi.org/10.3866/PKU.WHXB20030512>
  33. Qiu X-F, Zhu J-J, Chen H-Y (2003) Controllable synthesis of nanocrystalline gold assembled whiskery structures via sonochemical route. *J Cryst Growth* 257:378–383. [https://doi.org/10.1016/S0022-0248\(03\)01467-2](https://doi.org/10.1016/S0022-0248(03)01467-2)
  34. Wu S-H, Chen D-H (2003) Synthesis and characterization of nickel nanoparticles by hydrazine reduction in ethylene glycol. *J Colloid Interface Sci* 259:282–286. [https://doi.org/10.1016/S0021-9797\(02\)00135-2](https://doi.org/10.1016/S0021-9797(02)00135-2)
  35. Kan C, Cai W, Li C et al (2003) Ultrasonic synthesis and optical properties of Au/Pd bimetallic nanoparticles in ethylene glycol. *J Phys D* 36:1609–1614. <https://doi.org/10.1088/0022-3727/36/13/328>
  36. Li Q, Li H, Pol VG et al (2003) Sonochemical synthesis, structural and magnetic properties of air-stable Fe/Co alloy nanoparticles. *New J Chem* 27:1194. <https://doi.org/10.1039/b302136j>
  37. Okitsu K, Bandow H, Maeda Y, Nagata Y (1996) Sonochemical preparation of ultrafine palladium particles. *Chem Mater* 8:315–317. <https://doi.org/10.1021/cm950285s>
  38. Turáková M, Salmi T, Eränen K et al (2015) Liquid phase hydrogenation of nitrobenzene. *Appl Catal A* 499:66–76. <https://doi.org/10.1016/j.apcata.2015.04.002>
  39. Leipzig B (1898) Über stufenweise Reduktion des Nitrobenzols mit begrenztem Kathodenpotential. *Z Elektrochem Elektrochem* 4:506–514. <https://doi.org/10.1002/bbpc.18980042204>
  40. Qu R, Macino M, Iqbal S et al (2018) Supported bimetallic AuPd nanoparticles as a catalyst for the selective hydrogenation of nitroarenes. *Nanomaterials* 8:690. <https://doi.org/10.3390/nano8090690>
  41. Peureux J, Torres M, Mozzanega H et al (1995) Nitrobenzene liquid-phase hydrogenation in a membrane reactor. *Catal Today* 25:409–415. [https://doi.org/10.1016/0920-5861\(95\)00128-3](https://doi.org/10.1016/0920-5861(95)00128-3)
  42. Easterday R, Sanchez-Felix O, Losovyj Y et al (2015) Design of ruthenium/iron oxide nanoparticle mixtures for hydrogenation of nitrobenzene. *Catal Sci Technol* 5:1902–1910. <https://doi.org/10.1039/C4CY01277A>
  43. Lente G (2015) *Deterministic kinetics in chemistry and systems biology*. Springer, Cham

## Affiliations

Viktória Hajdu<sup>1</sup> · Ádám Prekob<sup>1</sup> · Gábor Muránszky<sup>1</sup> · István Kocserha<sup>2</sup> · Zoltán Kónya<sup>3</sup> · Béla Fiser<sup>1,4</sup> · Béla Viskolcz<sup>1</sup> · László Vanyorek<sup>1</sup> 

Viktória Hajdu  
kempviki@uni-miskolc.hu

Ádám Prekob  
kempadam@uni-miskolc.hu

Gábor Muránszky  
kemmug@uni-miskolc.hu

István Kocserha  
istvan.kocserha@uni-miskolc.hu

Zoltán Kónya  
konya@chem.u-szeged.hu

Béla Fiser  
kempfiser@uni-miskolc.hu

Béla Viskolcz  
bela.viskolcz@uni-miskolc.hu

- <sup>1</sup> Institute of Chemistry, University of Miskolc, 3515 Miskolc-Egyetemváros, Hungary
- <sup>2</sup> Institute of Ceramics and Polymer Engineering, University of Miskolc, 3515 Miskolc-Egyetemváros, Hungary
- <sup>3</sup> Department of Applied and Environmental Chemistry, University of Szeged, Rerrich Béla sq. 1, 6720 Szeged, Hungary
- <sup>4</sup> Ferenc Rákóczi II. Transcarpathian Hungarian Institute, Beregszász, Transcarpathia 90200, Ukraine