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# Dendritic structured palladium complexes: magnetically retrievable, highly efficient heterogeneous nanocatalyst for Suzuki and Heck cross-coupling reactions†

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The recyclable nanomagnetic Pd-complex PAMAM G<sub>0</sub>-Pd@γ-Fe<sub>2</sub>O<sub>3</sub> is reported for catalytic C–C cross-coupling reactions of challenging substrates. Mainly, a great variety of aryl chlorides can be used as substrates for Suzuki–Miyaura and Mizoroki–Heck reactions under mild reaction conditions (60–90 °C) and low catalyst loading (<1 mol% Pd) in aqueous media. The presence of numerous polar groups in the polymer matrix increases the solubility of the catalyst in water, thus facilitating its operation in aqueous environments. The immobilization of the catalyst on the surface of a magnetic platform allows its effective recovery and reuse without significant loss of catalytic activity for at least six cycles with total leaching of <1% palladium metal, meeting the requirements for acceptable metal residues in the pharmaceutical industry.

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## 1. Introduction

The development of new protocols and methods according to green chemistry standards to meet the growing demands of human society for energy, health, and agricultural purposes remains a pressing challenge for chemistry and the related sciences.<sup>1,2</sup> Arguably, catalysis stands at the forefront of such efforts, providing innovation by the development of new chemical transformations. Nevertheless, making catalysis feasible for large-scale applications by improving catalytic activity and selectivity, and especially in the case of transition metal catalysis, obviating metal leaching is equally essential even for well-established transformations.<sup>3</sup>

In this context, palladium-catalyzed cross-couplings have been manifold proven to be of particular significance for synthesizing medicinally relevant compounds even though palladium is one of the scarcest metals in the world.<sup>4–6</sup> Efficient recycling of such species is economically desirable; moreover, the considerable toxicity of palladium mandates its removal below 10 ppm in pharmaceuticals.<sup>7</sup> Separating homogeneous palladium catalysts from products is challenging; consequently,

heterogenization of palladium catalysts is a viable approach to overcome these difficulties. However, typical heterogeneous palladium catalysts such as Pd/C, being most successful for hydrogenations,<sup>8</sup> are less efficient in cross-coupling reactions, especially with industrially relevant aryl chlorides.<sup>9</sup> Such substrates remain challenging with any catalyst despite some impressive advances in designing molecular palladium complexes with specialized ligands.<sup>10–17</sup>

Considering large-scale and recycling, many efforts have been made to develop heterogeneous Pd-catalysts,<sup>10–14</sup> including molecular palladium–ligand complexes deposited onto various solid supports.<sup>15–19</sup> Among them, iron oxide nanoparticles have emerged as an attractive platform for metal catalysts<sup>20</sup> and especially for palladium complexes to arrive at eco-friendly and robust heterogeneous catalysts for Suzuki–Miyaura and Heck–Mizoroki C–C cross-coupling reactions.<sup>21–24</sup> Such catalysts can be easily manipulated and retrieved by external magnets avoiding cumbersome centrifugation or filtration processes.<sup>25–27</sup> Moreover, iron oxide nanoparticles have a high surface-to-volume ratio, which allows generating a large number of active sites that are available to promote catalytic reactions.<sup>28,29</sup>

Polyamidoamine (PAMAM) dendrimers have a particularly well-defined structure favorable for the encapsulation of metal nanoparticles.<sup>30</sup> The nanoparticle size and the metal loading can be controlled by the dendrimer generation, whereas the solubility in common solvents can be adjusted by the nature of the dendritic end group.<sup>31</sup> Despite these attractive properties, dendrimers are often difficult to recycle and to reuse; thus, attaching them to magnetic supports holds the promise to

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combine tunability of the dendritic core to generate a suitable environment of catalytically active moieties with magnetic properties beneficial for recovery and reuse.<sup>32</sup>

Previously, various homogeneous and heterogeneous systems of palladium nanoparticles encapsulated in modified PAMAM dendrimers of generation G<sub>2</sub>–G<sub>6</sub> have been reported for Suzuki–Miyaura and Mizoroki–Heck cross-coupling reaction.<sup>33–36</sup> For example, PAMAM-G<sub>3</sub> grafted on functionalized multiwall carbon nanotubes (MWCNTs-NH<sub>2</sub>) showed good activity for aryl chlorides in Heck reactions employing *N*-methyl-2-pyrrolidone (NMP) as a solvent,<sup>33</sup> and the Fe<sub>3</sub>O<sub>4</sub>-G<sub>4</sub>-PAMAM-Pd(0) could convert aryl chlorides in DMF as a solvent in Suzuki–Miyaura couplings.<sup>34</sup> Moving towards environmentally more benign solvent systems, the G<sub>6</sub>-OH(Pd<sub>74</sub>Cu<sub>73</sub>) PAMAM<sup>34</sup> dendrimer or Fe<sub>3</sub>O<sub>4</sub>@SiO<sub>2</sub>-PAMAM-Pd<sup>36</sup> were competent catalysts for aryl chlorides in Suzuki–Miyaura reactions in a mixture of EtOH/H<sub>2</sub>O. Similar to the PAMAM structure, polypropylene polybenzyl isocyanate (PPI) dendrimers modified by diphosphines as coordination site for Pd(II) complex have been employed successfully in Suzuki–Miyaura reactions with aryl chlorides in a THF/H<sub>2</sub>O solvent mixture.<sup>37</sup> More Pd-PAMAM-based systems were reported, but these only catalyze the coupling of aryl iodides and bromides effectively and show little activity at best for aryl chlorides.<sup>38–42</sup>

We report here the synthesis of a hybrid material, consisting of a superparamagnetic  $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> core decorated with zero-generation PAMAM dendrimer G<sub>0</sub> in which palladium nanoparticles were encapsulated (Fig. 1). This catalytic system shows unusual high activity for Suzuki–Miyaura and Heck coupling reactions, most notably with a wide variety of aryl chlorides as substrates.

## 2. Experimental section

The preparation of PAMAM G<sub>0</sub>-Pd@ $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> nanocatalyst is described in detail in the experimental section (ESI, Section 1. Schemes 1–3†). Briefly, at first  $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> nanocore was prepared as the platform for the immobilization, using Fe(II) and Fe(III) chloride salts. Then the iron oxide nanoparticles were functionalized with (3-chloropropyl)triethoxysilane followed by the attachment of the zero-generation of polyamidoamine dendrimer (PAMAM) to the  $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> MNPs through the siloxy linker. Subsequently, PAMAM G<sub>0</sub>@ $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> was treated with palladium dichloride, which produced the PAMAM G<sub>0</sub>-Pd@ $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> (ESI, Schemes 1–3†). The physicochemical properties of PAMAM G<sub>0</sub>-Pd@ $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> were characterized by FT-IR, TEM, XRD, FE-SEM,

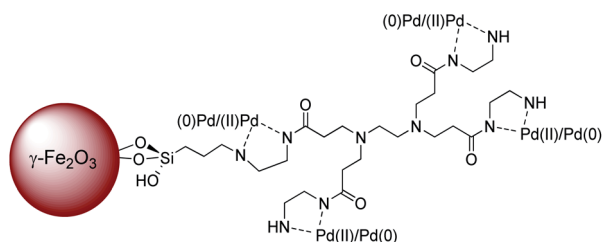


Fig. 1 The proposed structure of PAMAM G<sub>0</sub>-Pd@ $\gamma$ -Fe<sub>2</sub>O<sub>3</sub>.

ICP, TGA/DTG, VSM, MAP, XPS, EDS, and LC-MS analyzes (see the ESI, Fig. S1–S16†).

### 2.1 General procedure for the Suzuki–Miyaura cross-coupling reaction (9a–r)

PAMAM G<sub>0</sub>-Pd@ $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> (0.47 mol%, 0.007 g) was added to a mixture of K<sub>2</sub>CO<sub>3</sub> (2 mmol), phenylboronic acid (1.2 mmol), and aryl halide (1.0 mmol) in water (2 mL), under stirring and heated at 60 °C (90 °C when we used aryl chlorides). The progress of reaction was monitored by TLC until completion (15 min), then the nanocatalyst was separated by an external magnet. The reaction mixture was extracted with ethyl acetate (3 × 5 mL), and the combined organic layers were dried over anhydrous Na<sub>2</sub>SO<sub>4</sub> and concentrated. The residue was purified on silica to give rise to the pure products 9a–r.

### 2.2 General procedure for the Mizoroki–Heck cross-coupling reaction (12a–r)

PAMAM G<sub>0</sub>-Pd@ $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> (0.61 mol%, 0.009 g) was added to a mixture of K<sub>2</sub>CO<sub>3</sub> (2 mmol), alkene (1.5 mmol), and aryl halide (1.0 mmol) in water (2 mL), under stirring and heated at 80 °C (90 °C when we used aryl chlorides). The reaction progress was screened by TLC at different time intervals. After the completion of the reaction, the nanocatalyst was separated by an external magnet. The reaction mixture was extracted with ethyl acetate (3 × 5 mL), and the combined organic layers were dried over anhydrous Na<sub>2</sub>SO<sub>4</sub> and concentrated. The residue was purified on silica to give rise to the pure products 12a–r.

## 3. Results and discussion

### 3.1 Synthesis and characterization of the PAMAM G<sub>0</sub>-Pd@ $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> MNPs

The successful synthesis of PAMAM G<sub>0</sub>-Pd@ $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> was established by comparing its XRD pattern with those of  $\gamma$ -Fe<sub>2</sub>O<sub>3</sub>. The XRD pattern of  $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> (Fig. 2) confirms the cubic magnetite crystal structure according to the peaks at 15, 23.1, 26.9, 30.3, 35.8, 43.6, 50, 54.8, 57.3, 63.2, 71.1, and 74.4° (2 $\theta$ ), corresponding to the (110), (210), (211), (220), (311), (400), (421), (422), (511), (440), (620) and (533) miller indices respectively

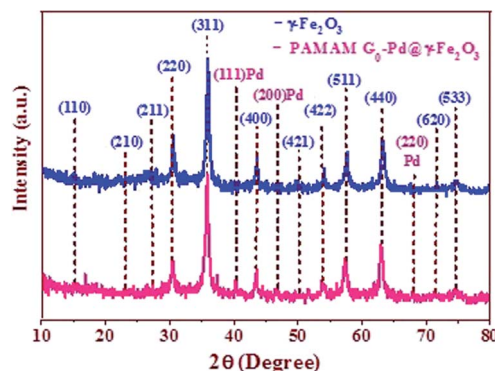


Fig. 2 XRD patterns of  $\gamma$ -Fe<sub>2</sub>O<sub>3</sub>, and PAMAM G<sub>0</sub>-Pd@ $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> complex.



(JCPDS card no. 39-1346).<sup>43,44</sup> The unit cell is assigned as the space group  $P4_132$  with the crystallographic parameters  $a = b = c = 8.3515 \text{ \AA}$ , and  $\alpha = \beta = \gamma = 90^\circ$ . The observed diffraction peaks of PAMAM  $G_0$ -Pd@ $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> at 40.1, 46.7 and 68.1° ( $2\theta$ ) are attributed to the standard crystallographic pattern of Pd NPs corresponding to the (111), (200), and (220) miller indices (JCPDS Card no. 65-2867, Fig. 2).<sup>45,46</sup> The decrease of the XRD peak intensity of PAMAM  $G_0$ -Pd@ $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> can be attributed to the formation of palladium complexes on the surface of  $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> MNPs being stabilized by various functional groups present in the PAMAM dendrimer acting as ligands.

The FE-SEM images of  $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> NPs,  $\gamma$ -Fe<sub>2</sub>O<sub>3</sub>-CPTES and supported catalyst were shown in Fig. S10.† According to the FE-SEM images, the  $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> and  $\gamma$ -Fe<sub>2</sub>O<sub>3</sub>-CPTES NPs have a uniform spherical morphology (Fig. S10A†). Also, the particles of the supported catalyst on the surface of  $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> NPs have a homogeneous spherical morphology and uniform size distribution. This clearly shows the synergistic effect of organic and inorganic diverse ingredients in a homogeneous network of PAMAM  $G_0$ -Pd@ $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> (6) (Fig. S10B†).

The TEM image of PAMAM  $G_0$ -Pd@ $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> is in line with the proposed core-shell structure placing palladium nanoparticles at the surface of magnetic Fe<sub>2</sub>O<sub>3</sub> nanoparticles (Fig. 3). Energy-dispersive X-ray (EDX) choosing a point at random on the surface of PAMAM  $G_0$ -Pd@ $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> NPs (Fig. 4) confirmed the presence of Pd, N, O, Si, Fe, Cl, and C, in 8.10, 7.74, 21.03, 2.45, 43.79, 2.69, and 14.20 wt% respectively.

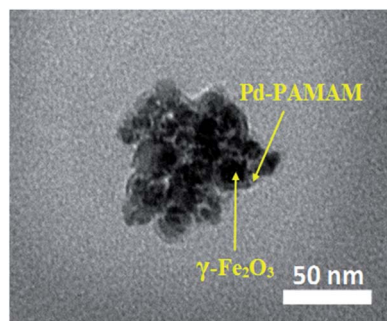


Fig. 3 TEM images of the PAMAM  $G_0$ -Pd@ $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> complex.

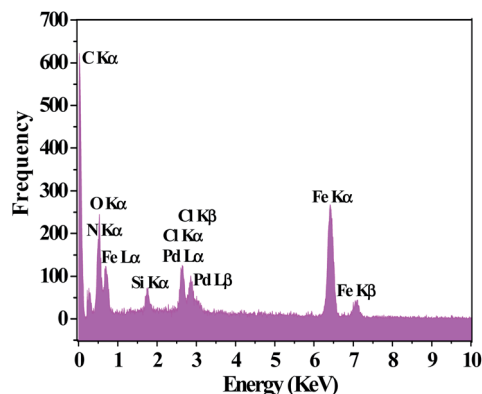


Fig. 4 EDX analysis of the PAMAM  $G_0$ -Pd@ $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> complex.

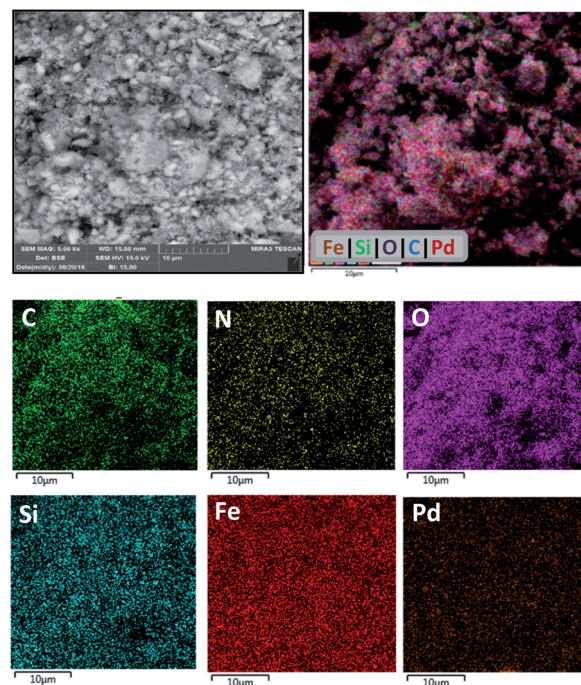


Fig. 5 EDS elemental mappings of the PAMAM  $G_0$ -Pd@ $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> complex.

Elemental mapping analysis by energy dispersive spectroscopy (EDS) of PAMAM  $G_0$ -Pd@ $\gamma$ -Fe<sub>2</sub>O<sub>3</sub> (Fig. 5) indicates that Pd NPs are distributed almost uniformly and without aggregation

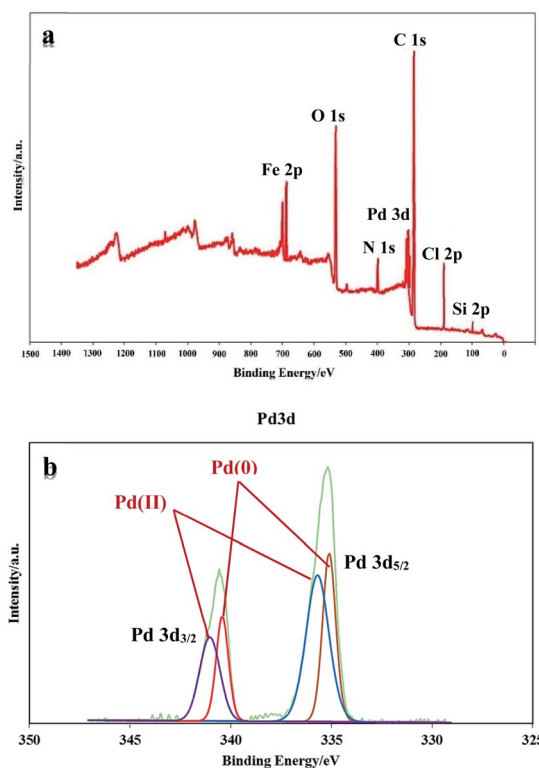


Fig. 6 XPS patterns of PAMAM  $G_0$ -Pd@ $\gamma$ -Fe<sub>2</sub>O<sub>3</sub>: overall elemental survey spectrum (a), and, Pd3d (b).

Table 1 Suzuki–Miyaura cross-coupling reactions of different aryl halides with phenylboronic acid catalyzed by PAMAM G<sub>0</sub>-Pd@γ-Fe<sub>2</sub>O<sub>3</sub><sup>a</sup>

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Table 1 (Contd.)


					
Prod.	X	Yield	<i>t</i> (min)	TON	TOF
<b>9r</b>	I	85	80	180	135
<sup>a</sup> Reaction conditions: phenylboronic acid (1.2 mmol), aryl halide (1.00 mmol), K <sub>2</sub> CO <sub>3</sub> (Et <sub>3</sub> N when we used the aryl chlorides) (2.00 mmol), catalyst (0.007 g, 0.47 mol% of Pd), H <sub>2</sub> O (2.0 mL), 60 °C (90 °C when we used the aryl chlorides); all yields are isolated.					

Table 2 Mizoroki–Heck cross-coupling reactions of different aryl halides with styrene catalyzed by PAMAM G<sub>0</sub>-Pd@γ-Fe<sub>2</sub>O<sub>3</sub><sup>a</sup>

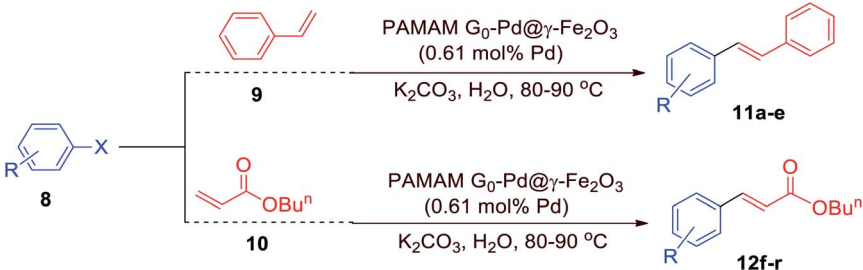
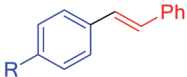
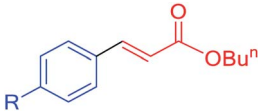
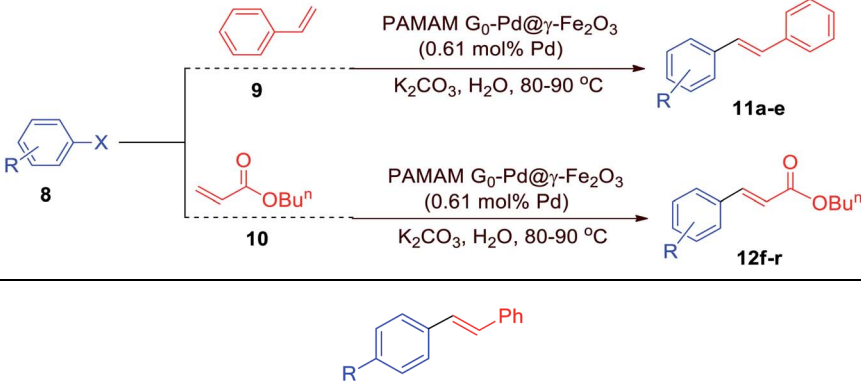
						
Prod.	X	R	Yield	<i>t</i> (min)	TON	TOF
<b>12a</b>	I	Me	75	180	122	41
<b>12a</b>	Br	Me	65	180	106	35
<b>12a</b>	Cl	Me	55	220	90	24
<b>12b</b>	I	OMe	90	90	148	98
<b>12c</b>	I	H	95	60	155	150
<b>12c</b>	Br	H	95	80	155	117
<b>12c</b>	Cl	H	90	120	147	73
<b>12d</b>	I	CN	85	60	139	139
<b>12e</b>	I	NO <sub>2</sub>	95	45	155	206
<b>12e</b>	Br	NO <sub>2</sub>	95	50	155	186
<b>12e</b>	Cl	NO <sub>2</sub>	90	60	147	147
						
<b>12f</b>	I	OH	90	120	147	73
<b>12g</b>	I	NH <sub>2</sub>	90	120	147	73
<b>12h</b>	I	OMe	92	500	150	181
<b>12h</b>	Cl	OMe	65	300	106	21
<b>12i</b>	I	Me	80	120	131	65
<b>12i</b>	Br	Me	80	140	131	56
<b>12i</b>	Cl	Me	70	150	114	45
<b>12j</b>	I	H	95	50	155	187
<b>12j</b>	Br	H	90	75	147	117
<b>12j</b>	Cl	H	80	100	131	78
<b>12k</b>	I	Cl	85	55	139	153
<b>12k</b>	Br	Cl	70	60	114	114
						



Table 2 (Contd.)



Prod.	X	R	Yield	<i>t</i> (min)	TON	TOF
12l	I	Br	75	70	122	105
12m	I	CO <sub>2</sub> Et	95	40	131	65
12m	Br	CO <sub>2</sub> Et	93	50	152	183
12m	Cl	CO <sub>2</sub> Et	90	80	147	110
12n	I	COMe	92	70	150	129
12o	Br	CN	92	50	150	181
12o	Cl	CN	85	120	139	69
12p	I	NO <sub>2</sub>	95	30	155	77
12q	Br	Me	65	130	106	49
12q	Cl	Me	50	240	81	20
12r	I	NO <sub>2</sub>	92	70	150	129

<sup>a</sup> Reaction conditions: styrene or *n*-butyl acrylate (1.5 mmol), aryl halide (1.00 mmol), K<sub>2</sub>CO<sub>3</sub> (Et<sub>3</sub>N when we used the aryl chlorides) (2.00 mmol), catalyst (0.009 g, 0.61 mol% of Pd), H<sub>2</sub>O (2.00 mL), 80 °C (90 °C when we used aryl chlorides); all yields are isolated.

on the catalyst surface along with the other elements (C, N, O, Si, Fe) present.

XPS analysis was performed to investigate the chemical state of the surface of PAMAM G<sub>0</sub>-Pd@γ-Fe<sub>2</sub>O<sub>3</sub> (Fig. 6a). In the XPS elemental survey of PAMAM G<sub>0</sub>-Pd@γ-Fe<sub>2</sub>O<sub>3</sub>, the peaks corresponding to carbon, oxygen, nitrogen, chlorine, silicon, palladium and iron are clearly observed (Fig. 6a). Also, Pd(0)/(II) existence in the catalyst structure were confirmed through XPS analysis through studying binding energy range of 335.15 to 340.97 eV (Fig. 6b). As shown in Fig. 6b, the peaks located at 335.15 (3d<sub>5/2</sub>) and 340.35 eV (3d<sub>3/2</sub>) correspond to Pd(II) species, and also the peaks appeared at 336.11 (3d<sub>5/2</sub>) and 340.97 eV (3d<sub>3/2</sub>) are attributed Pd(0) (Fig. 6b). According to XPS results, the ratio of Pd(0)/Pd(II) is equal to 0.40 (in atomic number).

### 3.2 Evaluation of the catalytic activity of PAMAM G<sub>0</sub>-Pd@γ-Fe<sub>2</sub>O<sub>3</sub> MNPs in cross-coupling reactions

Taking the Suzuki-Miyaura C-C coupling reaction between iodobenzene and phenylboronic acid as the model reaction,

PAMAM G<sub>0</sub>-Pd@γ-Fe<sub>2</sub>O<sub>3</sub> was evaluated varying solvent, temperature, base, and catalyst loading (see ESI for details, Fig. S17†), suggesting that the best results were achieved with polar solvents. Being green and safe, we were pleased to see that the reaction proceeded exceptionally well in water, giving rise to 95% yield with a catalyst loading of 0.47 mol% within 15 minutes of reaction time at 60 °C.

With the optimized reaction conditions in hand, we evaluated the substrate scope of PAMAM G<sub>0</sub>-Pd@γ-Fe<sub>2</sub>O<sub>3</sub> for Suzuki-Miyaura coupling reactions (Table 1). A wide variety of electron-deficient or electron-rich aryl halides or phenylboronic acids were amenable, and functional groups that could be potentially sensitive under the basic, aqueous reaction conditions employed (CHO, CN, NH<sub>2</sub>) were tolerated well. As expected, the reactivity decreased from aryl iodides *via* aryl bromides to aryl chlorides, but even for the latter high yields could be obtained by raising the reaction temperature to 90 °C and prolonging the reaction time 100–220 min.

Analogously, the catalytic activity of PAMAM G<sub>0</sub>-Pd@γ-Fe<sub>2</sub>O<sub>3</sub> was evaluated for Mizoroki-Heck C-C coupling reactions.



Screening of conditions for the reaction of iodobenzene with styrene (see ESI for details, Fig. S18†) also revealed for this transformation water to be a suitable choice in combination with  $K_2CO_3$  at a reaction temperature of 80 °C, giving rise to 1,2-diphenylethylene in 95% yield within 60 minutes at a catalyst loading of 0.61 mol%.

Evaluating the substrate scope under the optimized reaction conditions demonstrated that a great variety of electronically and sterically differentiated aryl halides reacted smoothly in the coupling with styrene or *n*-butyl acrylate (Table 2). Noteworthy, electron-rich and electron-poor aryl chlorides could be converted, requiring only slightly longer reaction times and higher temperatures (90 °C).

Control experiments confirmed for both, the Suzuki–Miyaura as well as the Mizoroki–Heck reactions that no coupling is observed with the parent materials PAMAM  $G_0@ \gamma\text{-Fe}_2\text{O}_3$  and PAMAM $G_0$ , but also that  $PdCl_2$  as a stand-alone catalyst does not promote the transformations. The high activity and efficiency of the PAMAM  $G_0\text{-Pd}@ \gamma\text{-Fe}_2\text{O}_3$  catalyst compares well to other catalysts examined for comparison (see Fig. S19†).

### 3.3 Recyclability study

The recyclability, activity, and stability of PAMAM  $G_0\text{-Pd}@ \gamma\text{-Fe}_2\text{O}_3$  catalyst were examined using the hot-filtration method

taking the Mizoroki–Heck reaction between iodobenzene and styrene (Table 2, entry 5) as a representative example.

After the catalyst was magnetically removed from the reaction medium (28 min, 50% yield of 1,1-diphenylethylene monitored by GC), the reaction, by and large, did not proceed further (53% yield after 90 min, Fig. S20†), indicating that little to no catalytically active species were present outside the support.

ICP analysis of the Mizoroki–Heck model reaction solution was performed to determine the amount of palladium leaching for six consecutive catalytic runs (Fig. S21†).

Base on the results, Pd catalyst, showed high stability, with little leaching of Pd (total less than 1%), while consistent yields (95–93%) of 1,1-diphenylethylene were obtained for each cycle (see ESI for details, Table S1†).

Likewise, recyclability of the PAMAM  $G_0\text{-Pd}@ \gamma\text{-Fe}_2\text{O}_3$  complex was evaluated for the Suzuki–Miyaura reaction between iodobenzene and phenylboronic acid for 6 sequential runs (see ESI for details, Table S1†), giving rise to 93–95% yield in each run. The comparison of TEM and SEM images of fresh and the used catalyst after 6 runs in the Mizoroki–Heck reaction did not show significant changes in size and morphology (Fig. 7a and b). In addition, VSM measurements (Fig. 7c) and FT-IR spectra (Fig. 7e) further confirmed that the catalyst did not change in its structure and magnetic properties. Also, evaluation and comparison leaching, some of hydrophilic/

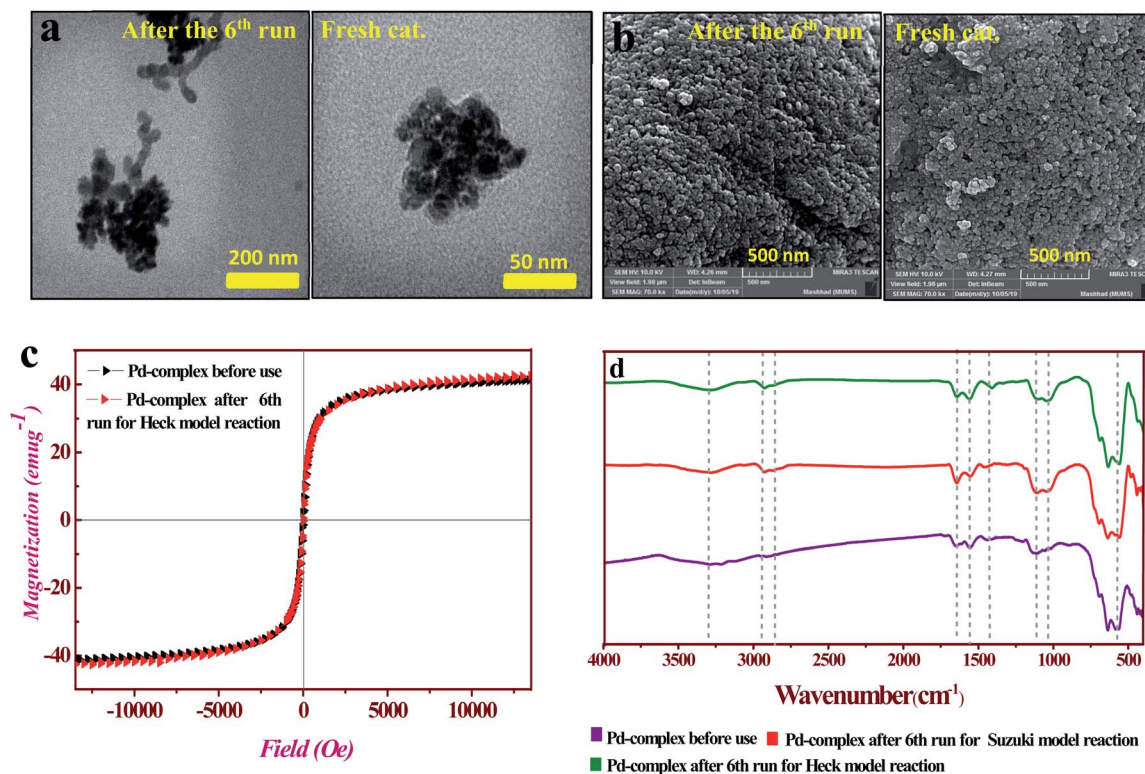


Fig. 7 (a) TEM, and (b) FE-SEM, images recovered of the PAMAM  $G_0\text{-Pd}@ \gamma\text{-Fe}_2\text{O}_3$  complex after the 6<sup>th</sup> run for the model reaction of Mizoroki–Heck; (c) magnetization curves of PAMAM  $G_0\text{-Pd}@ \gamma\text{-Fe}_2\text{O}_3$  before use and PAMAM  $G_0\text{-Pd}@ \gamma\text{-Fe}_2\text{O}_3$  after the 6<sup>th</sup> run for the model reaction of Mizoroki–Heck, at 300 K; (d) FT-IR spectra of PAMAM  $G_0\text{-Pd}@ \gamma\text{-Fe}_2\text{O}_3$  before use; FT-IR spectra of PAMAM  $G_0\text{-Pd}@ \gamma\text{-Fe}_2\text{O}_3$  after 6<sup>th</sup> run for the model reactions of Suzuki–Miyaura and Mizoroki–Heck cross-couplings.

amphiphilic catalysts effective in c-c cross coupling reactions with PAMAM G<sub>0</sub>-Pd@γ-Fe<sub>2</sub>O<sub>3</sub> catalyst were performed to better understand the activity of the PAMAM G<sub>0</sub>-Pd@γ-Fe<sub>2</sub>O<sub>3</sub> (see Table 3. ESI† for found more details).

## 4. Summary and conclusions

In conclusion, we developed a recyclable magnetic nano Pd-complex, formulated as PAMAM G<sub>0</sub>-Pd@γ-Fe<sub>2</sub>O<sub>3</sub> MNPs, which proved to be an efficient catalyst for the Suzuki-Miyaura and Mizoroki-Heck coupling. Notable features are its high activity in water, which allows the employment of aryl chloride at moderately forcing conditions (90 °C), facile recyclability and reuse, as well as little palladium leaching (<1% in a total of 6 runs).

## Conflicts of interest

All the authors, declare that they haven't any of conflicts of interest.

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