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Efficient amorphous platinum catalyst cluster growth on porous carbon: A combined Molecular Dynamics and experimental study

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Abstract: Amorphous platinum clusters supported on porous carbon have been envisaged for high-performance fuel cell electrodes. For this application, it is crucial to control the morphology of the Pt layer and the Pt – substrate interaction to maximize activity and stability. We thus investigate the morphology evolution during Pt cluster growth on a porous carbon substrate employing atomic scale molecular dynamics simulations. The simulations are based on the Pt-C interaction potential using parameters derived from Density Functional Theory and are found to yield a Pt cluster morphology similar to that observed in low loaded fuel cell electrodes prepared by plasma sputtering. Moreover, the simulations show amorphous Pt cluster growth in agreement with X-ray diffraction and Transmission Electron Microscopy experiments on high performance low Pt content ($10\mu\text{g}_{\text{Pt}}\text{cm}^{-2}$) loaded fuel cell electrodes and provide a fundamental insight in the cluster growth mechanism.

Keywords: Molecular dynamics; plasma sputtering deposition; cluster growth; Pt catalyst; fuel cell electrodes

1. Introduction

Physical and electronic properties of metal clusters deposited on various substrates have recently received a great deal of attention. Noble metal clusters, particularly Ru, Rh, Pd, Pt and Au,^[1] are very often used in catalytic reactions.^[2,3] In heterogeneous catalysis, reactions occur at the catalyst surface, so that a high surface-to-volume ratio leading to a large fraction of accessible atoms is required for increasing the activity of the catalyst^[4-6], and for decreasing its cost. There is thus a real interest to obtain the highest catalytic surface area from a low metal weight. Currently, a very important application of noble metals is their use for activating electrochemical reactions occurring in proton exchange membrane fuel cells (PEMFCs),^[7-9] such as the sluggish oxygen reduction reaction.^[10,11] In this latter case, Pt-based clusters are supported by nanostructured carbon substrates, among them disordered porous carbon structures^[12,13] and ordered carbon structures, such as nanotubes (CNTs)/carbon nanofibers (CNFs).^[14,15]

Lee et al. reported on the Pt cluster size distribution as a function of Pt loading for electrodeposited Pt nanocrystal on highly orientated pyrolytic graphite (HOPG) surfaces and demonstrated that the average cluster size increased as the Pt loading increased,^[16] resulting in the decrease of the active surface area. Several methods were proposed for lowering the Pt loading: by alloying Pt with foreign metal atoms,^[17,18] by forming Pt-based core-shell structures,^[19,20] by localizing the catalytic particle close to the proton conductive solid electrolyte,^[21] by modifying Pt cluster surface with a proton conductive polymer,^[22,23] etc. One of the major degradation factors of the electrical performances of PEMFC electrodes however is the agglomeration process of metal clusters under fuel cell working conditions,^[24,25] due to their mobility on the carbon supports.^[26] It is then necessary to keep the Pt clusters well attached to and dispersed on the nanostructured carbon support to avoid their agglomeration over time, which would lead to the loss of surface area and mass

activity.^[27] However, Clark and Kesmodel^[28] studied very small Pt clusters in the order of a few nm and demonstrated that their migration over the nanoporous surface leads to coalescence which in turn leads to the formation of a more stable and larger cluster. Accordingly, Lee et al.^[16] reported that the diameter distribution of mobile Pt clusters have a maximum around 14 nm. The morphology, microstructure and interaction with the support of the metal clusters are therefore properties recognized as driving the activity and stability of the catalyst. Many efforts are then directed at optimizing the catalytic performance, which in turn requires a thorough understanding of both the relationship between the nanocatalyst deposition process and the nanocatalyst activity, as well as the Pt – substrate interactions^[29, 30]. The ability to predict - and ultimately – design stable and highly active Pt clusters supported on a porous carbon substrate is thus of paramount importance. Classical reactive molecular dynamics (MD) simulations was used to investigate the dynamical and structural evolution of platinum cluster growth on a nanostructured porous carbon substrate at the atomic scale by a plasma sputtering method. Such metal deposition method has proven to be very efficient for the preparation of highly active low Pt loaded PEMFC electrodes.^[31-33]

We will demonstrate how these atomic scale simulations provide insight in the influence of the deposition conditions on the cluster growth and the resulting Pt morphology. Finally, we develop a plasma method for preparing Pt cluster deposited on carbon and provide a comparison between our simulation results and experimental results on the structural and morphological evolution of the Pt-clusters during the growth, which are leading to high performance fuel cell electrodes.

2. Experimental

The deposition chamber is equipped with two rectangular targets: one of pure carbon and one of Pt_{0.01}C_{0.99}. Deposition conditions are as follows: power 530 W, target voltage 700 V, and

target current 0.78 A. The growth rate is 4 nm.min⁻¹. The target-to-substrate distance is 6.5 cm. Targets are 45° tilted with respect to the substrate. Conventional uncatalyzed E-Tek carbon porous layers (CPL) on a carbon woven web are used as substrates. Electrodes are loaded at 0.010 mg cm⁻²_{Pt}. The Pt loading and localization profile inside the CPL was determined by performing Rutherford backscattering spectroscopy: 90% of Pt atoms were localized in the first 200 nm depth of the material.

TEM images were obtained using a JEOL JEM 2010 (equipped with a LaB6 filament and with a resolution of 0.35 nm). The size distribution was estimated from the determination of the diameter from 100 particles using ImageJ free software^[34].

The XRD measurements were performed on a Bruker D 5005 Bragg-Brentano diffractometer operated with a copper tube powered at 40 kV and 40 mA ($\text{CuK}_{\alpha 1} = 1.5406 \text{ \AA}$) with 0.06° step and a fixed acquisition time of 10 s/step. The catalytic powder scraped from a catalyzed CPL was deposited on a Kanthal foil and measurements were carried out in the 2θ range from 20° to 60°. The acquisition of the whole pattern was performed for 183 s at room temperature. The instrumental resolution function of the diffractometer was evaluated from the pattern (10-155.5°) of the NIST SRM 660 (LaB₆) line profile standard. XPS measurements were performed by using an Escalab MKII (VG scientific) set-up using the magnesium monochromatic beam (1253.6 eV) at room temperature at pressures below 8·10⁻⁹ Torr in the analysis chamber. Core level spectra were recorded at a 50 meV resolution. Binding energies were lined up with respect to the C1s peak at 284.6 eV.

The Membrane electrode assembly for fuel cell tests consisted in two 0.010 mg_{Pt}cm⁻² Pt/CPL electrodes with 5 cm² surface area mechanically pressed on a Nafion 212 membrane at 2 Nm torque in the cell hardware (Electrochem). No preliminary hot pressing process of electrodes against the Nafion 212 membrane (used as received from Electrochem. Inc.) was performed.

No Nafion solution was added in the electrode catalytic layers before use. Under our testing conditions, the pressure is manually regulated with the output valves and hydrogen and oxygen are humidified by bubbling in water at 80°C and 40°C, respectively. Measurements in a 5 cm² PEMFC using pure H₂ and O₂ gases are performed using a ECL150/MTS 150/HSA unit (Electrochem. Inc.). The polarization curves are recorded by applying increasing current values to the cell and reading the voltage stabilization (at least 30 s stabilization time for each current value).

3. Molecular Dynamics Modeling

The MD simulations are intended to mimic the Pt-cluster growth process from Pt-atom deposition on a carbon nanoporous substrate. A realistic model for this substrate is constructed by applying a constrained reverse Monte-Carlo procedure designed to reproduce a set of high resolution transmission electron microscopy pictures.^[35] The dimensions of the model substrate are 6 x 6 x 6 nm³, containing 17040 atoms, corresponding to a density of 1.57 g cm⁻³ and a porosity of 48%. Periodic boundaries are applied in the lateral directions, thus exposing the top of the structure as a free surface.

2500 platinum atoms are released towards the substrate every $\Delta t = 2$ ps. The initial position of the Pt atom is 8 Å above the substrate and with randomly chosen {x,y} coordinates. Pt atoms interact with other Pt atoms and with the nanostructured carbon substrate atoms within a cut-off distance of 8 Å. The initial velocities of the incoming Pt atoms are sampled from a Maxwell-Boltzmann distribution with three most probable mean kinetic energies $\langle E_{\text{deposit}} \rangle$ of 1 eV, 0.1 eV and 0.026 eV and with randomly chosen incident angles. The mean kinetic energy $\langle E_{\text{deposit}} \rangle$ of the incoming atoms is calculated according to a modified Thompson formula for atoms travelling through a plasma^[36,37] over a defined target-to-substrate distance. In the case of a mean kinetic energy $\langle E_{\text{deposit}} \rangle = 1$ eV, the simulation corresponds to

sputtering experiments at low argon pressure where a small amount of buffer gas only randomizes the sputtered atom velocities with little energy loss. The second case studied, $\langle E_{\text{deposit}} \rangle = 0.1$ eV, either corresponds to resistive or e-beam evaporation, which produces a vapour at the vaporization temperature of ca. 0.1 eV. Plasma sputtering deposition at a gas pressure around 2 Pa also produces a vapor with mean kinetic energy around 0.1 eV. In the last case, where $\langle E_{\text{deposit}} \rangle = 0.026$ eV, the simulation mimics a metal vapor at the temperature of 300 K which occurs when sputtered Pt atoms travel across an argon plasma at a pressure of around 5 Pa.

The temperature of the substrate in all simulations is assumed to be 300 K. The substrate temperature is controlled by applying a Berendsen heat bath, employing a relaxation time of 1.17 ps. The carbon-carbon interactions in the substrate are modeled by a Tersoff potential.^[38] The platinum-platinum interactions are modeled through an EAM potential.^[39-43] The platinum-carbon interactions are modeled using a 12-6 Lennard-Jones (LJ) potential. We used parameters that were derived by Morrow and Striolo,^[44] based on DFT calculations by Acharya et al.^[45] for the binding energy of Pt on various graphite sites. The DFT data was then used to generate a potential energy surface, to which the Lennard-Jones interaction potential was fitted. According to this parameterization, the characteristic Pt – C interaction length is $\sigma_{\text{PtC}} = 1.60$ Å and energy, $\epsilon_{\text{PtC}} = 0.1365$ eV, which are significantly different from previous sets^[46-49]. Indeed, we have chosen to use this new parametrization, as the previously used sets gave rise to very weak substrate bonding and the formation of Pt clusters which are invariably crystalline, contrary to our experiments (see below).

4. Results and discussion

The condensation coefficient of Pt onto the porous substrate is 0.83 ± 0.02 and does depend on the incoming kinetic energy.

Snapshots obtained from these simulations are shown in **Figure 1**, for the three impact energy distributions considered here.

Figure 1. Snapshots of Pt deposition on the porous carbon substrate. The green spheres represent C-atoms and the brownish spheres represent Pt-atoms.

It can be seen in **Figure 1** that simulations result in the formation of thin film-like structures for all kinetic energies investigated. The relatively higher binding energy allows the Pt deposition to closely wet the substrate morphology. Moreover, the Pt atoms are allowed to diffuse through the pores of the substrate, resulting in an overall smoother and flatter surface morphology. This result is in agreement with experimental plasma sputtering deposition of Pt on porous carbon, demonstrating the Pt diffusion into the porous carbon substrate.^[50-52]

The Pt concentration profiles for the three set of kinetic energy distribution centered at 0.26, 0.1 and 1.0 eV are plotted in Figure 2. For these three sets of calculation the concentration profile exhibits a bell shape, as in the case for thermal diffusion. When increasing the incoming kinetic energy, the concentration profile is shifted towards the in-depth of the substrate.

Figure 2. The Pt concentration profile for the three sets of kinetic energies. The values in the inset correspond to the mean kinetic energy of the sampled Maxwell-Boltzmann distribution.

Radial distribution function (RDF) plots, counting the neighbors at a given distance, for the Pt deposits are shown in **Figure 3** and detailed in **Table 1**. For the three kinetic energy distributions of the depositing Pt atoms investigated here, the RDF first peak is very broad. Moreover, the third and fourth peaks are overlapping for all energies, while the second peak, corresponding to 2nd neighbor distance expected in bulk crystalline Pt is missing. This is due to the fact that the Pt clusters are wetting the disordered surface, such that the parallel atomic planes below and above a Pt atom (where the 2nd neighbors are lying in a face centered cubic crystal), are not defined. Consequently, the corresponding peak does not appear in the

RDF. Both features are representative of disorder. On the other hand, close inspection of the snapshots in **Figure 1** reveals some atomic ordering, and specifically the formation of centered hexagons on the surface.

Figure 3. The calculated RDF of the Pt deposit grown on the porous carbon.

Table 1. RDF peak positions corresponding to Figure 3 of the deposited Pt structures grown on the porous carbon substrate displayed in **Figure 2**, and the comparison to the Pt bulk crystal values.

	1 st neighbor (Å)	2 nd neighbor (Å)	3 rd neighbor (Å)	4 th neighbor (Å)	5 th neighbor (Å)
Pt crystal value	2.7746	3.9239	4.8058	5.5492	6.79
Pt (1 eV)	2.66	-	4.79	-	-
Pt (0.1 eV)	2.65	-	4.67	-	-
Pt (0.026 eV)	2.67	-	4.85	-	-

To further resolve the the ordered/disordered state of the Pt deposits showin in Figure 1, we calculate their resulting X-ray ray diffraction (XRD) patterns. The calculated X-ray intensities vs. 2θ of the deposited Pt clusters are shown in **Figure 4** and detailed in **Table 2**.

Figure 4. The simulated XRD intensities vs. 2θ of deposited Pt structures on the model porous carbon substrate. Diamond symbols give the Pt bulk positions.

It can be seen that only two broad peaks arise for each sample, which can be attributed to Pt(111) and Pt(220) orientations. This observation is consistent with some order in the plane of the surface and the fading of 2nd and other neighbor distances.

Table 2. The 2θ position of XRD peaks for Pt crystal and simulated Pt structures (Unit: 2θ /°).

	Pt(111)	Pt(200)	Pt(220)	Pt(311)	Pt(222)	Pt(400)	Pt(331)
<i>Pt bulk crystal</i>	39.75	46.25	67.45	81.25	85.7	103.5	118.15
Pt (1 eV)	40.56	----	73.95	----	----	----	----
Pt (0.1 eV)	40.73	----	72.8	----	----	----	----
Pt (0.026 eV)	40.69	----	74.6	----	----	----	----

It has been demonstrated that it is possible to produce such amorphous Pt structure on a porous carbon by co-deposition of Pt and carbon using plasma magnetron sputtering.^[50,51] A typical TEM image of the deposited Pt clusters is shown in **Figure 5a** with the related size distribution in the inset of **Figure 5a**, and the experimental XRD pattern recorded on such magnetron sputtered electrodes in **Figure 5b**. The TEM image shows very small Pt clusters of ca. 1 nm, and the size distribution leads to an estimate of the mean size of Pt particles wetting carbon spheres of about 1.4 nm. Note, however, that the counting is complicated due to the Pt

cluster agglomeration. It is therefore likely that the bigger particles are actually agglomerated clusters rather than isolated clusters.

The XRD pattern shows no diffraction peak corresponding to the fcc structure of platinum, but only a featureless Pt background, which is consistent with the very low Pt loading ($10 \mu\text{g cm}^{-2}$) and an amorphous structure. Most apparent peaks come from polytetrafluoroethylene (PTFE) particles present in the porous carbon layer acting as substrate for the deposited Pt clusters.

Figure 5. (a) typical TEM image of Pt nanoclusters deposited by plasma sputtering on a carbon porous layer, (b) Experimental XRD pattern showing the amorphous structure of the Pt cluster deposited onto a carbon porous layer and (c) XPS spectra of the Pt 4f orbital of Pt nanoclusters deposited by plasma sputtering on a carbon porous layer (loading: $10 \mu\text{g}_{\text{Pt}} \text{cm}^{-2}$).

Moreover, this result is confirmed by XPS measurements (**Figure 4c**) showing only the presence of PtO_xH_y species (**Table 3**) in Pt clusters, which is consistent with a very small

particle size.^[53] For Pt clusters of ca. 2.5-3.0 nm, the shell Pt atoms have a +2 oxidation state, whereas the core Pt Pt atoms are metallic.^[54] The absence of metallic Pt confirmed that the size of the clusters was small enough to consider all Pt atoms as surface atoms.^[55] All these physicochemical characterizations confirm that the structure of the Pt deposit on the carbon porous layer consists of very small amorphous clusters.^[53]

Table 3. XPS catalyst characterization data. The assignment of chemical species to the binding energy is based on data from the Handbook of X-ray photoelectron spectroscopy.^[56]

	Binding energy / eV	Atomic ratio / %	Species	Concentration ratio / %
Pt 4f 7/2	72.39	39.46	Pt(OH) ₂	69.06
Pt 4f 5/2	75.74	29.6		
Pt 4f 7/2	73.80	17.68	PtO	30.94
Pt 4f 5/2	77.15	13.26		

As described in detail above, the calculated RDF and XRD patterns for the structures generated are typical for a more amorphous Pt structure. On the other hand, the experimental characterization data (TEM, XRD and XPS) confirm that Pt clusters sputtered on a porous carbon layer are very small and amorphous.

Such electrodes have been tested under H₂/O₂ fuel cell working conditions and have shown high performance,^[21,52,57] particularly concerning the Pt utilization efficiency which reach ca. 20 kW g_{Pt}⁻¹ as shown in **Figure 6**. Typical fuel cell performances of hydrogen/oxygen fed fuel cells at 80°C, with 0.35 mg cm⁻² Pt per electrode, are higher in terms of achieved power densities, ca. 1.0 to 1.2 W cm⁻² with oxygen pressure between 1.0 and 3.0 absolute bars^[58], but leads to a much lower Pt utilization efficiency of ca. 2 kW g_{Pt}⁻¹. However, the stability of such small particles could be an important issue for the system as a whole. On the other hand, such structures, consisting of strongly agglomerated Pt clusters (i.e., as if Pt clusters are supported by Pt clusters) could increase their stability towards corrosion (due to a higher chemical potential [59]) and minimize that of the support (due to less surface contact).

Figure 6. U_{cell} vs j and P vs j curves obtained at 70 °C with a membrane electrode assembly (MEA) consisting in a Nafion 212 membrane sandwiched between symmetrical 0.010 mg_{Pt}cm⁻² Pt /CPL electrodes (flow rates of 350 cm³ min⁻¹ for O₂ and 500 cm³ min⁻¹ for H₂; T_{O₂} = 40 °C, T_{H₂} = 80 °C, P_{O₂}=4 bar, P_{H₂} = 3 bar).

The power density and stability issues as well as the operating conditions of the cell may be not realistic for transport applications, where potential duty cycles and start-stop cycles may drastically deteriorate the fuel cell performances, but such low platinum loading system could be used for stationary or nomad applications.

4. Conclusion

In conclusion, plasma sputter deposition experiments have lead to high performance Pt/C fuel cell electrodes. Careful analysis of the catalytic phase allowed concluding that the catalyst consists in very small amorphous platinum clusters. For understanding this phenomenon, we have carried out molecular dynamics simulation under conditions closely matching these experiments. MD simulations demonstrated the formation of Pt clusters wetting the surface and which fit closely the surface morphology, which leads to a smooth ultrathin film. In this case, only the surface structure remains crystalline to some extent. The resulting Pt morphologies are in agreement with experiment of Pt sputtered onto fuel cell electrodes as

revealed by comparing the RDF and XRD simulated patterns to the experimental XRD, XPS and TEM results. This confirms that small amorphous Pt clusters are able to grow on carbon grains and to present high catalytic activity.

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References

- [1] T. Muroi, Role of Precious Metal Catalysts. In *Noble Metals* (Eds: Yen-Hsun Su) InTech, **2012**, pp. 301-334.
- [2] A. Chen, P. Holt-Hindle, Platinum-Based Nanostructured Materials: Synthesis, Properties, and Applications. *Chem. Rev.* **2010**, *110*, 3767.
- [3] V. Selvaraj, M. Alagar, K. S. Kumar, Synthesis and characterization of metal clusters-decorated PPY–CNT composite and their electrocatalytic oxidation of formic acid and formaldehyde for fuel cell applications. *Appl. Catal. B.* **2007**, *75*, 129.
- [4] R. Narayanan, M. A. El-Sayed, Changing Catalytic Activity During Colloidal Platinum Nanocatalysis Due to Shape Changes: Electron Transfer Reaction. *J. Am. Chem. Soc.* **2004**, *126*, 7194.
- [5] A. S. K. Hashmi, G. J. Hutchings, The First Gold-Catalyzed C-S Bond Formation: Cycloisomerization of α -Thioallenes to 2,5-Dihydrothiophenes. *Angew. Chem. Int. Ed.* **2006**, *45*, 1897.
- [6] Y. J. Xiong, B. Wiley, Y. N. Xia, Nanocrystals with Unconventional Shapes - A Class of Promising Catalysts. *Angew. Chem. Int. Ed.* **2007**, *46*, 7157.
- [7] S. S. Gupta, N. Bandyopadhyaya, J. Datta, Carbon-supported platinum catalysts for direct alcohol fuel cell anode. *Mater. Manuf. Process.* **2006**, *21*, 703.
- [8] W. Li, C. Liang, J. Qiu, W. Zhou, H. Han, Z. Wei, G. Sun, Q. Xin, Carbon nanotubes as support for cathode catalyst of a direct methanol fuel cell. *Carbon* **2002**, *40*, 787.
- [9] S. H. Joo, S. J. Choi, I. Oh, J. Kwak, Z. Liu, O. Terasaki, R. Ryoo, Ordered nanoporous arrays of carbon supporting high dispersions of platinum clusters. *Nature.* **2001**, *412*, 169.
- [10] T. R. Ralph, M. P. Hogarth, *Platinum Metal Rev.* Catalysis for Low Temperature Fuel Cells, Part I: The Cathode Challenges. **2002**, *46*, 3.

- [11] V. R. Stamenkovic, B. Fowler, B. S. Mun, G. F. Wang, P. N. Ross, C. A. Lucas, N. N. Markovic, Improved Oxygen Reduction Activity on Pt₃Ni(111) via Increased Surface Site Availability. *Science* **2007**, 315, 493.
- [12] H. A. Gasteiger, S. S. Kocha, B. Sompalli, F. T. Wagner, Activity benchmarks and requirements for Pt, Pt-alloy, and non-Pt oxygen reduction catalysts for PEMFCs. *Appl. Catal. B.* **2005**, 56, 9.
- [13] K. Y. Chan, J. Ding, J. Ren, S. Cheng, K. Y. Tsang, Supported mixed metal clusters as electrocatalysts in low temperature fuel cells. *J. Mat. Chem.* **2004**, 14, 505.
- [14] G. Che, B. B. Laskami, E. R. Fisher, C. R. Martin, Carbon nanotubule membranes for electrochemical energy storage and production. *Nature* **1998**, 393, 346.
- [15] Z. Q. Tian, S. H. Lim, C. K. Poh, Z. Tang, Z. Xia, Z. Luo, P. K. Shen, D. Chua, Y. P. Feng, Z. Shen, J. A. Lin, Highly Order –Structured Membrane Electrode Assembly with Vertically Aligned Carbon Nanotubes for Ultra-Low Pt Loading PEM Fuel Cells. *Adv. En. Mat.* **2011**, 1, 1205.
- [16] I. Lee, K. Y. Chan, D. L. Phillips, Growth of electrodeposited platinum nanocrystals studied by atomic force microscopy. *Appl. Surf. Sci.* **1998**, 136, 321.
- [17] V. R. Stamenkovic, B. S. Mun, M. Arenz, K. J. J. Mayrhofer, C. A. Lucas, G. Wang, P. N. Ross, N. M. Markovic, Trends in electrocatalysis on extended and nanoscale Pt-bimetallic alloy surfaces. *Nature Materials*, **2007**, 6, 241.
- [18] S. J. Hwang, S. K. Kim, J. G. Lee, S. C. Lee, J. H. Jang, P. Kim, T. H. Lim, Y. E. Sung, S. J. Yoo, Role of electronic perturbation in stability and activity of Pt-based alloy nanocatalysts for oxygen reduction. *J. Am. Chem. Soc.* **2012**, 134, 19508-19511.
- [19] J. Zhang, M. B. Vukmirovic, Y. Xu, M. Mavrikakis, R. R. Adzic, Controlling the Catalytic Activity of Platinum-Monolayer Electrocatalysts for Oxygen Reduction with Different Substrates. *Angew. Chem. Int. Ed.* **2005**, 44, 2132.

- [20] L. Zhang, R. Iyyamperumal, D. F. Yancey, R. M. Crooks, G. Henkelman, Design of Pt-Shell Clusters with Alloy Cores for the Oxygen Reduction Reaction. *ACS Nano*, **2013**, 7, 9168.
- [21] M. Cavarroc, A. Ennadjaoui, M. Mougnot, P. Brault, R. Escalier, Y. Tessier, J. Durand, S. Roualdès, T. Sauvage, C. Coutanceau, Performance of plasma sputtered Fuel Cell electrodes with ultra-low Pt loadings. *Electrochem. Comm.* **2009**, 11, 859.
- [22] A. C. Ferrandez, S. Baranton, J. Bigarré, P. Buvat, C. Coutanceau, Pt Particles Functionalized on the Molecular Level as New Nanocomposite Materials for Electrocatalysis. *Langmuir*, **2012**, 28, 17832.
- [23] A. C. Ferrandez, S. Baranton, J. Bigarré, P. Buvat, C. Coutanceau, Chemical Functionalization of Carbon Supported Metal Clusters by Ionic Conductive Polymer via the “Grafting From” Method. *Chem. Mater*, **2013**, 25, 3797.
- [24] R. Sellin, C. Grolleau, C. Coutanceau, J. M. Leger, S. Arrii-Clacens, S. Pronier, J. M. Clacens, Effects of Temperature and Atmosphere on Carbon-Supported Platinum Fuel Cell Catalysts. *J. Phys. Chem. C* **2009**, 113, 21735.
- [25] L. Dubau, L. Castanheira, G. Berthomé, F. Maillard, An identical-location transmission electron microscopy study on the degradation of Pt/C clusters under oxidizing, reducing and neutral atmosphere. *Electrochim. Acta* **2013**, 110, 273.
- [26] P. Urchaga, M. Weissmann, S. Baranton, T. Girardeau, C. Coutanceau, Improvement of the platinum clusters – carbon substrate interaction by insertion of a thiophenol molecular bridge. *Langmuir* **2009**, 25, 6543.
- [27] A. Tseung, S. Dhara, Loss of surface area by platinum and supported platinum black electrocatalyst. *Electrochim. Acta*. **1975**, 20, 681.
- [28] G. Clark, L. Kesmodel, Ultrahigh vacuum scanning tunneling microscopy studies of platinum on graphite, *J. Vac. Sci. Technol. B* **1993**, 11, 131.
- [29] K. Ostrikov, E. C. Neyts and M. Meyyappan, *Advances in Physics* **2013**, 62,113

- [30] E. C. Neyts, K. Ostrikov, Z. J. Han, S. Kumar, A. C. T. van Duin, A. Bogaerts, *Phys. Rev. Lett.* **2013**, 110, 065501
- [31] S. Hirano, J. Kim, S. Srinivasan, High performance proton exchange membrane fuel cells with sputter-deposited Pt layer electrodes. *Electrochim. Acta*, **1997**, 42, 1587.
- [32] A. T. Haug, R. E. White, J. Weidner, W. Huang, S. Shi, T. Stoner, N. Ranac, Increasing Proton Exchange Membrane Fuel Cell Catalyst Effectiveness Through Sputter Deposition. *J. Electrochem. Soc.* **2002**, 149, A280.
- [33] P. Brault, A. Caillard, A. L. Thomann, J. Mathias, C. Charles, R. W. Boswell, S. Escribano, J. Durand, T. Sauvage, Plasma sputtering deposition of platinum into porous fuel cell electrodes. *J. Phys. D: Appl. Phys.* **2004**, 37, 3419–3423.
- [34] Available at <http://rsbweb.nih.gov/ij/>
- [35] J. Pikunic, C. Clinard, N. Cohaut, K. E. Gubbins, J. M. Guet, R. J. M. Pellenq, I. Rannou, J. N. Rouzaud, Structural modeling of porous carbons: constrained reverse Monte Carlo method. *Langmuir*. **2003**, 19, 8565.
- [36] K. Meyer, I. K. Schuller, C. M. Falco, Thermalization of sputtered atoms, *J. Appl. Phys.* **1981**, 52, 5803.
- [37] A. Gras-Marti, J. A. Valles-Abarca, Slowing down and thermalization of sputtered particle fluxes: Energy distributions. *J. Appl. Phys.* **1983**, 54, 1071.
- [38] J. Tersoff, Empirical interatomic potential for carbon, with applications to amorphous carbon. *Phys. Rev. Lett.* **1988**, 61, 2879.
- [39] M. S. Daw, M. I. Baskes, Embedded-atom method: Derivation and application to impurities, surfaces, and other defects in metals. *Phys. Rev. B.* **1984**, 29, 6443.
- [40] S. Foiles, Application of the embedded-atom method to liquid transition metals. *Physical Review B.* **1985**, 32, 3409.
- [41] S. Foiles, M. Baskes, M. Daw, Embedded-atom-method functions for the fcc metals Cu, Ag, Au, Ni, Pd, Pt, and their alloys. *Phys. Rev. B.* **1986**, 33, 7983.

- [42] M. S. Daw, S. Foiles, M. I. Baskes, The embedded-atom method: a review of theory and applications. *Materials Science Reports* **1993**, 9, 251.
- [43] S. Foiles, M. I. Baskes, Contributions of the embedded-atom method to materials science and engineering. *Mater. Res. Bull.* **2012**, 37, 485.
- [44] B. H. Morrow, A. Striolo, Assessing how metal–carbon interactions affect the structure of supported platinum clusters. *Mol. Simulat.* **2009**, 35, 795.
- [45] C. K. Acharya, D. I. Sullivan, C. H. Turner, Characterizing the interaction of Pt and PtRu clusters with boron-doped, nitrogen-doped, and activated carbon: Density functional theory calculations and parameterization. *J. Phys. Chem. C.* **2008**, 112, 13607.
- [46] S. Y. Liem, K. Y. Chan, Simulation study of platinum adsorption on graphite using the Sutton-Chen potential. *Surf. Sci.* **1995**, 328, 119.
- [47] S. Y. Liem, K. Y. Chan, Effective pairwise potential for simulations of adsorbed platinum. *Mol. Phys.* **1995**, 86, 939.
- [48] A. Sutton, J. Chen, Long-range Finnis–Sinclair potentials. *Phil. Mag. Lett.* **1990**, 61, 139.
- [49] D. B. Graves, P. Brault, Molecular dynamics for low temperature plasma–surface interaction studies. *J. Phys. D: Appl. Phys.* **2009**, 42, 194011.
- [50] A. Caillard, P. Brault, J. Mathias, C. Charles, R. Boswell, T. Sauvage, Deposition and diffusion of platinum clusters in porous carbon assisted by plasma sputtering. *Surf. Coat. Technol.* **2005**, 200, 391.
- [51] P. Brault, C. Josserand, J. M. Bauchire, A. Caillard, C. Charles, R. W. Boswell, Anomalous diffusion mediated by atom deposition into a porous substrate. *Phys. Rev. Lett.* **2009**, 102, 045901.
- [52] Caillard, A.; Charles, C.; Boswell R.; Brault, P. Improvement of the sputtered platinum utilization in proton exchange membrane fuel cells using plasma-based carbon nanofibers. *J. Phys. D: Appl. Phys.* **2008**, 48, 185307 1.

- [53] P. Brault, A. Caillard, S. Baranton, M. Mougenot, S. Cuynet, C. Coutanceau, One-step synthesis and chemical characterization of Pt/C nanowire composites by plasma sputtering. *ChemSusChem*. **2013**, 6, 1168.
- [54] R. Sellin, J.-M. Clacens, C. Coutanceau, A thermogravimetric analysis/mass spectroscopy study of the thermal and chemical stability of carbon in the Pt/C catalytic system, *Carbon* **2010**, 48, 2244.
- [55] C. Henry C. Size Effects on Structure and Morphology of Free or Supported Clusters. In *Nanomaterials and Nanochemistry* (Eds: C. Bréchnignac, P. Houdy, M. Lahmani) Springer, Berlin, Germany **2007**, pp. 3-32.
- [56] Wagner, C. D., Riggs, W. M., Davis, L. E., Moulder, J. F., Mouilenberg, G. E. *Handbook of X-ray Photoelectron Spectroscopy*; Perkin Elmer Corporation: Eden Prairie, MN, 1978.
- [57] M. Mougenot, A. Caillard, P. Brault, S. Baranton, C. Coutanceau, High Performance Plasma Sputtered PdPt Fuel Cell Electrodes with Ultra Low Loading. *Int. J. Hydrogen Energy*. **2011**, 36, 8429.
- [58] A. J. -J. Kadjo, P. Brault , A. Caillard , C. Coutanceau , J.-P. Garnier, S. Martemianov, Improvement of proton exchange membrane fuel cell electrical performance by optimization of operating parameters and electrodes preparation, *J. Power Sources* **2007**, 172, 613-622.
- [59] J. A. S. Bett, K. Kinoshita, P. Stonehart, Crystallite growth of platinum dispersed on graphitized carbon black: II. Effect of liquid environment, *J. Catal.* **1976**, 41, 124–133.