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COMMUNICATION

Facile formation of ZIF-8 thin film on ZnO nanorods

Cite this: DOI: 10.1039/x0xx00000x

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Accepted 00th January 2015

DOI: 10.1039/x0xx00000x

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In this work, thin films of the well-known metal organic framework ZIF-8 were formed on zinc oxide nanorods through the reaction with 2-methyl imidazole solution (Hmim). Deposition of a thin film of the linker solution allows the underlying zinc oxide nanorod morphology to be preserved, resulting in the facile and precise formation of a nanostructured hybrid materials in short reaction times and under conventional heating. The effect of various synthesis parameters on the morphology of the resulting thin film is reported.

Metal-organic frameworks (MOFs) are porous solid materials comprising metal ions or clusters bound to organic linkers. Their surprising tuneability has led to their applications in various fields.^{1, 2} The combination of metal-organic frameworks with other functional materials results in the formation of multi-functional hybrid materials with combined properties or synergistic effects of the individual components.^{3, 4} Such combination allows for the formation of materials with novel properties with applications in fields such as sensing,⁵ catalysis,⁶ and drug delivery.⁷

One of the most often studied MOFs is ZIF-8. Known for its remarkable stability⁸ and ease of synthesis,⁹ this MOF has been used in a variety of areas such as sensing,^{10, 11} catalysis,^{12, 13} drug encapsulation,¹⁴ and separation.¹⁵ However, ZIF-8 thin films are notoriously difficult to grow due to MOF's inherent nucleation mechanism¹⁶ and are therefore often prepared through methods such as secondary nucleation,¹⁷ surface functionalization¹⁸ or by the use of harsh and time consuming solvothermal methods.¹⁹ Recently, zinc oxide was shown to aid the formation of ZIF-8 thin films by acting both as the nucleation site as well as the source of zinc ions.¹⁷ Such development allows for the one-step formation of ZIF-8 membranes and composites, combining the properties of ZIF-8 and zinc oxide.

Zinc oxide is a semi-conductor with a wide bandgap of ~3.3 eV and a high exciton binding energy of 60 meV at room temperature.²⁰ In recent years, zinc oxide has attracted much attention due to its biocompatibility, high electron mobility, chemical stability, and ease

of synthesis.²¹⁻²³ One of the most interesting aspects of zinc oxide is the possibility to form a large variety of nanostructures.²⁴ Whereas the morphology of zinc oxide has been investigated alongside synthesis and application of metal nanoparticles,²⁵ quantum dots,²⁶ and enzymes,²⁷ the exploitation of the large variety of zinc oxide morphologies alongside synthesis and application of MOFs is relatively unexplored.

Even though the use of zinc oxide is shown to facilitate the formation of ZIF-8 thin films, solvothermal methods are often still required to form ZIF-8 films.^{28, 29} A recent trend has been the use of solvent-free methods, relying instead on molten linkers for the formation of ZIF-8 as well as other MOFs.^{30, 31} Whereas before, the linker was dissolved in a solvent, here the powdered linker is simply placed in contact with a metal oxide and heated to its melting point so as to form a MOF. This method is often performed using metal oxide powders, with the aim of complete conversion of the oxide to the desired MOF.^{28, 29, 32, 33} Ameloot *et al.* have recently used this method to form coatings of ZIF-8 on zinc oxide nanostructures.³⁴ The main advantages of using a molten linker for the formation of MOF thin films are the lack of solvent, the ability to re-use unreacted linker and shorter synthesis time with a full surface coverage achieved in less than an hour in contrary to other methods that rely on a longer synthesis time of a day or more.^{32, 35} Furthermore, this method is green with water being the only side-product. However, the thin films synthesised by Ameloot *et al.* were not totally uniform, with some crystals being as large as 1 μm in diameter, making this method unsuitable for the coating of finer structures in the nanoscale regime.

Here, a new method is developed for the synthesis of ZIF-8 thin films on electrochemically grown zinc oxide nanorod arrays. This method combines the advantages of zinc oxide electrodeposition with using a molten linker for the facile formation of ZIF-8 thin films. Deposition of a thin film of the linker solution instead of using a linker powder allows more control over the reaction while preserving the underlying delicate nanorod morphology.

Zinc oxide nanoarrays were prepared using an amperometric method based on literature.³⁶ The optimization of the

electrochemical deposition of well-defined zinc oxide nanorods can be found in the ESI†. By casting 200 μL Hmim linker solution onto the zinc oxide nanorod array and allowing the solvent to evaporate, a thin uniform linker film was formed. After heating this sample at 150 $^{\circ}\text{C}$ in an oven, ZIF-8 is formed on zinc oxide nanoarrays. The effects of heating rate and nanorod morphology are given in the ESI†. Figure 1 shows the effect of using different amounts of linker and reveals a significant change between the use of a thin film of linker solution as opposed to a pure linker powder. Addition of a pure linker powder results in a MOF film on top of the rods with inhomogeneous crystal sizes, consistent with previous reports.³⁴ However, the drop casting of a thin film of linker solution allows more control over the linker amount which results in an even distribution of linker while the underlying nanorod morphology is maintained, indicating the growth of a thin MOF-film.

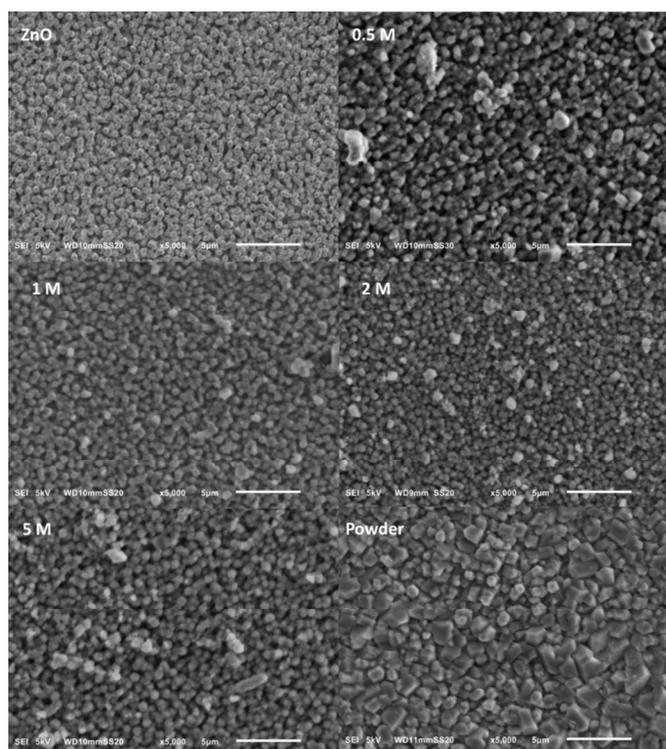


Figure 1. SEM images showing the ZIF-8 morphology for different concentrations of Hmim linker and comparison with the use of pure Hmim linker powder (20 minutes reaction time, scale bar represents 5 μm).

The effect of reaction time on ZIF-8 weight percentage, as calculated from XRD analysis (see ESI†),³⁷ is shown in Figure 2. An exponential increase in weight percentage is observed with reaction time, in good agreement with kinetic studies on MOF formation.³⁸ This behaviour has been observed before when using zinc oxide^{34, 35} as well as zinc ions in a solution of linker^{16, 39, 40} and has been attributed to the rapid growth of formed ZIF-8 crystal nuclei.¹⁶ The weight percentage seems to stabilize after 20 minutes, indicating no more zinc oxide is consumed after this time.

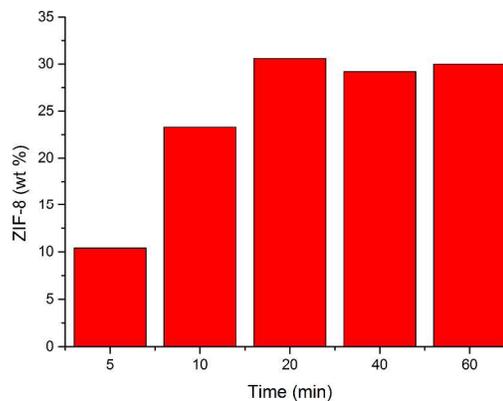


Figure 2. ZIF-8 weight percentage as a function of the reaction time using a 2 M linker solution in methanol.

The corresponding SEM images shown in Figure 3 indicate that the nanorod morphology is maintained throughout the synthesis and appears to become more defined with increasing the reaction time. Unfortunately, an increase in reaction time seems to also affect the growth of larger crystals on top. As almost no change in weight percentage of ZIF-8 is observed, the formation of these larger crystals can be largely attributed to Ostwald ripening.^{16, 34}

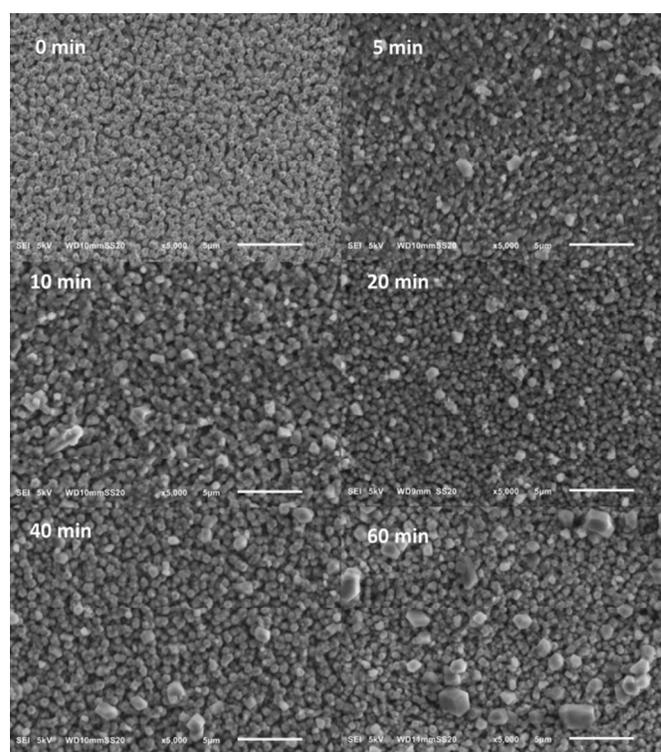


Figure 3. SEM images showing the effect of reaction time on ZIF-8 morphology using a 2 M Hmim linker solution in methanol.

A more detailed examination of the ZnO nanorods before and after coating is given in figure S14. ZnO nanorods are formed with a diameter of approximately 350 nm. After ZIF-8 formation, the rods diameter increases to approximately 520 nm.

Zinc oxide possesses a polar top plane and apolar side planes,²⁴ implying that the polarity of the solvent used to deposit the thin linker layer could affect the coverage of the nanorods. It is known that DMF has a larger surface tension and a more apolar nature

compared to methanol. Thus DMF would allow more linker to be retained onto the substrate and accordingly a more thorough coverage of the zinc oxide nanorods. As can be seen in Figure 4, the effect of reaction time when using DMF differs from methanol. An increase in the ZIF-8 weight percentage at shorter synthesis times is observed in both cases. However, this increase is much less significant when using DMF and could indicate different crystal growth kinetics.³⁹

The differences in reaction progression are evident in Figure 5. While the use of methanol preserves the nanorod morphology throughout the reaction, this is not the case for DMF as large crystals of ZIF-8 coat the surface. The change in crystal morphology appears to be similar to previous observations,^{34, 40} as small cubic crystals seem to grow in size and become more truncated. The increase in crystal size, coupled with a decrease in the amount of crystals, indicate Ostwald ripening.¹⁶ However, a steady increase in ZIF-8 weight percentage indicates that crystal growth at the expense of zinc oxide still occurs up to 40 minutes reaction time. Ameloot *et al.* reported that an increase in the film thickness of a flat zinc oxide layer resulted in the formation of larger ZIF-8 crystals on top of the layer.³⁴ The use of DMF could result in more linker being retained resulting in a much faster reaction and hence the growth of larger crystals.³⁹ Furthermore, the incomplete removal of DMF must be taken into account. It is known that DMF itself can decompose thermally, yielding formate and dimethylamine which can act as deprotonating agents and increase the rate of ZIF-8 formation. Furthermore, DMF can also interact with both linker and metal. However, it is worth mentioning that DMF decomposition is very slow with 3.3% decomposition after 33 h at 100 °C and the interactions of DMF mentioned above are known to slow down nucleation and would result in the opposite of what is observed here.³⁹

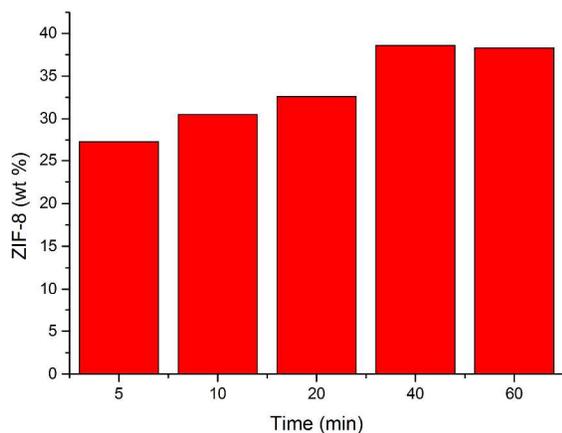


Figure 4. ZIF-8 weight percentage as a function of reaction time using a 2 M Hmim linker solution in DMF.

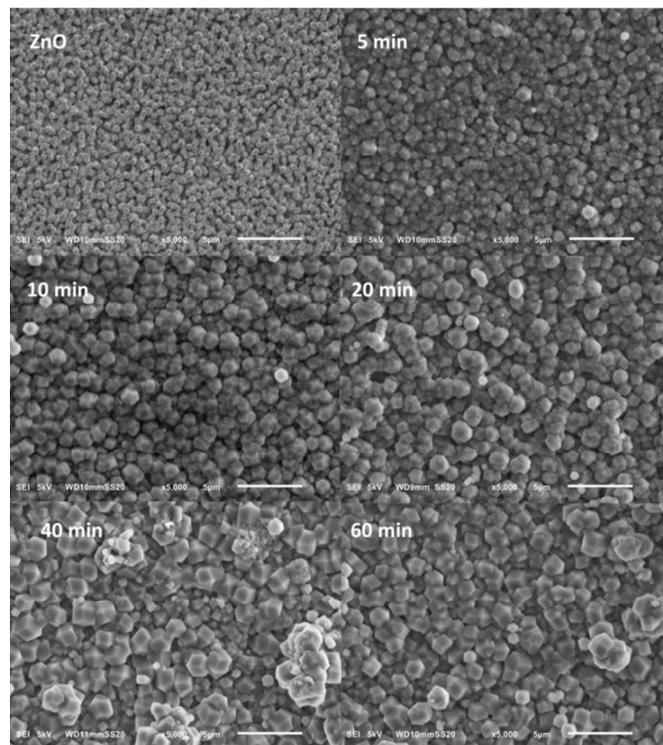


Figure 5. SEM images showing the effect of the reaction time on ZIF-8 morphology using a 2 M Hmim linker solution in DMF.

Conclusion

A new approach was suggested for the formation of a ZIF-8 thin layer on nanostructured zinc oxide. By casting a thin film of the linker solution, the morphology of the underlying zinc oxide was preserved and more control over the reaction was achieved. This method allows for the formation of nanostructured thin films in a fast and facile manner, while minimizing the use of both solvents and linker. Furthermore, an increase in the synthesis time results in the formation of more defined ZIF-8 coated rods. However, the growth of large crystals on top of the nanorod array is also observed to increase and is thought to be due to Ostwald ripening.

Much is still unknown regarding the crystallisation kinetics inherent to this method. Understanding the kinetics would not only lead to more insight into this method but could also allow for ways to reduce the growth of larger crystals on the surface of the rods and hence improve their function.

The technique presented in this work is versatile and can be applied to other metal oxide nanostructures and metal organic frameworks. This type of hybrid materials can be used in a variety of applications such as membrane technology, sensing and catalysis. The next step should hence focus on the investigation of their properties for various potential applications. In view of the chemical stability, high porosity and sieving properties of ZIF-8, assessment of porosity is an important factor. Finally, ZIF-8/ZnO nanohybrids prepared on transparent FTO substrates provide the possibilities into their use in photochemistry. Thus, the assessment of photochemical properties of this material is therefore of great interest.

Acknowledgements

G.C. and A.D. thank the European Research Council under the

European Union's Seventh Framework Programme (FP/2007-2013)/ERC Grant Agreement n. 335746, CrystEng-MOF-MMM for financial support.

Notes and references

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† Electronic Supplementary Information (ESI) available: [The experimentals, details for the synthesis of ZnO nanorods, and formation of ZIF-8 can be found in ESI]. See DOI: 10.1039/c000000x/

- L. E. Kreno, K. Leong, O. K. Farha, M. Allendorf, R. P. Van Duyne and J. T. Hupp, *Chemical Reviews*, 2011, **112**, 1105-1125.
- B. Seoane, J. Coronas, I. Gascon, M. E. Benavides, O. Karvan, J. Caro, F. Kapteijn and J. Gascon, *Chemical Society Reviews*, 2015, DOI: 10.1039/C4CS00437J.
- Q.-L. Zhu and Q. Xu, *Chemical Society Reviews*, 2014, **43**, 5468-5512.
- H. Al-Kutubi, J. Gascon, E. J. R. Sudhölter and L. Rassaei, *ChemElectroChem*, 2015, DOI: 10.1002/celc.201402429, n/a-n/a.
- X. Lin, G. Gao, L. Zheng, Y. Chi and G. Chen, *Analytical chemistry*, 2013, **86**, 1223-1228.
- K. F. Babu, M. A. Kulandainathan, I. Katsounaros, L. Rassaei, A. D. Burrows, P. R. Raithby and F. Marken, *Electrochemistry Communications*, 2010, **12**, 632-635.
- F. Ke, Y.-P. Yuan, L.-G. Qiu, Y.-H. Shen, A.-J. Xie, J.-F. Zhu, X.-Y. Tian and L.-D. Zhang, *Journal of Materials Chemistry*, 2011, **21**, 3843-3848.
- K. S. Park, Z. Ni, A. P. Côté, J. Y. Choi, R. Huang, F. J. Uribe-Romo, H. K. Chae, M. O'Keeffe and O. M. Yaghi, *Proceedings of the National Academy of Sciences*, 2006, **103**, 10186-10191.
- Y. Pan, Y. Liu, G. Zeng, L. Zhao and Z. Lai, *Chemical Communications*, 2011, **47**, 2071-2073.
- G. Lu and J. T. Hupp, *Journal of the American Chemical Society*, 2010, **132**, 7832-7833.
- S. Liu, Z. Xiang, Z. Hu, X. Zheng and D. Cao, *Journal of Materials Chemistry*, 2011, **21**, 6649-6653.
- U. P. Tran, K. K. Le and N. T. Phan, *ACS Catalysis*, 2011, **1**, 120-127.
- C. M. Miralda, E. E. Macias, M. Zhu, P. Ratnasamy and M. A. Carreon, *ACS Catalysis*, 2011, **2**, 180-183.
- C.-Y. Sun, C. Qin, X.-L. Wang, G.-S. Yang, K.-Z. Shao, Y.-Q. Lan, Z.-M. Su, P. Huang, C.-G. Wang and E.-B. Wang, *Dalton Transactions*, 2012, **41**, 6906-6909.
- H. Bux, C. Chmelik, R. Krishna and J. Caro, *Journal of Membrane Science*, 2011, **369**, 284-289.
- S. R. Venna, J. B. Jasinski and M. A. Carreon, *Journal of the American Chemical Society*, 2010, **132**, 18030-18033.
- M. Shah, H. T. Kwon, V. Tran, S. Sachdeva and H.-K. Jeong, *Microporous and Mesoporous Materials*, 2013, **165**, 63-69.
- O. Shekhah and M. Eddaoudi, *Chemical Communications*, 2013, **49**, 10079-10081.
- J. Yao and H. Wang, *Chemical Society Reviews*, 2014, **43**, 4470-4493.
- C. Jagadish and S. J. Pearton, *Zinc oxide bulk, thin films and nanostructures: processing, properties, and applications*, Elsevier, 2011.
- Z. Li, R. Yang, M. Yu, F. Bai, C. Li and Z. L. Wang, *The Journal of Physical Chemistry C*, 2008, **112**, 20114-20117.
- Z. Fan and J. G. Lu, *Journal of nanoscience and nanotechnology*, 2005, **5**, 1561-1573.
- L. Rassaei, R. Jaber, S. E. Flower, K. J. Edler, R. G. Compton, T. D. James and F. Marken, *Electrochimica Acta*, 2010, **55**, 7909-7915.
- Z. L. Wang, *Materials Today*, 2004, **7**, 26-33.
- N. P. Herring, K. AbouZeid, M. B. Mohamed, J. Pinsk and M. S. El-Shall, *Langmuir*, 2011, **27**, 15146-15154.
- H. Chen, W. Li, H. Liu and L. Zhu, *Electrochemistry Communications*, 2011, **13**, 331-334.
- S. K. Arya, S. Saha, J. E. Ramirez-Vick, V. Gupta, S. Bhansali and S. P. Singh, *Analytica chimica acta*, 2012, **737**, 1-21.
- Y. Yue, B. Guo, Z.-A. Qiao, P. F. Fulvio, J. Chen, A. J. Binder, C. Tian and S. Dai, *Microporous and Mesoporous Materials*, 2014, **198**, 139-143.
- M. Drobek, M. Bechelany, C. Vallicari, A. Abou Chaaya, C. Charmette, C. Salvador-Levehang, P. Miele and A. Julbe, *Journal of Membrane Science*, 2015, **475**, 39-46.
- M. Lanchas, S. Arcediano, A. T. Aguayo, G. Beobide, O. Castillo, J. Cepeda, D. Vallejo-Sánchez and A. Luque, *RSC Advances*, 2014, **4**, 60409-60412.
- J.-B. Lin, R.-B. Lin, X.-N. Cheng, J.-P. Zhang and X.-M. Chen, *Chemical Communications*, 2011, **47**, 9185-9187.
- Y. Yue, Z.-A. Qiao, X. Li, A. J. Binder, E. Formo, Z. Pan, C. Tian, Z. Bi and S. Dai, *Crystal Growth & Design*, 2013, **13**, 1002-1005.
- E. Zanchetta, L. Malfatti, R. Ricco, M. J. Styles, F. Lisi, C. J. Coghlan, C. J. Doonan, A. J. Hill, G. Brusatin and P. Falcaro, *Chemistry of Materials*, 2014, DOI: 10.1021/cm502882a.
- I. Stassen, N. Campagnol, J. Fransaer, P. Vereecken, D. De Vos and R. Ameloot, *CrystEngComm*, 2013, **15**, 9308-9311.
- W.-w. Zhan, Q. Kuang, J.-z. Zhou, X.-j. Kong, Z.-x. Xie and L.-s. Zheng, *Journal of the American Chemical Society*, 2013, **135**, 1926-1933.
- M. Skompska and K. Zarebska, *Electrochimica Acta*, 2014, **127**, 467-488.
- J. Martín-Ramos, A. Cambeses, A. López-Galindo, J. Scarrow and J. Díaz-Hernández, *Pathways for quantitative analysis by X-Ray diffraction*, INTECH Open Access Publisher, 2012.
- M. Goesten, E. Stavitski, E. A. Pidko, C. Gücüyener, B. Boshuizen, S. N. Ehrlich, E. J. M. Hensen, F. Kapteijn and J. Gascon, *Chemistry – A European Journal*, 2013, **19**, 7809-7816.
- P. Y. Moh, M. Brenda, M. W. Anderson and M. P. Attfield, *CrystEngComm*, 2013, **15**, 9672-9678.
- J. Cravillon, C. A. Schroder, H. Bux, A. Rothkirch, J. Caro and M. Wiebcke, *CrystEngComm*, 2012, **14**, 492-498.