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A porous, electrically conductive hexa-zirconium(IV) metal–organic framework†

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Engendering electrical conductivity in high-porosity metal–organic frameworks (MOFs) promises to unlock the full potential of MOFs for electrical energy storage, electrocatalysis, or integration of MOFs with conventional electronic materials. Here we report that a porous zirconium-node-containing MOF, NU-901, can be rendered electronically conductive by physically encapsulating C₆₀, an excellent electron acceptor, within a fraction (ca. 60%) of the diamond-shaped cavities of the MOF. The cavities are defined by node-connected tetra-phenyl-carboxylated pyrene linkers, *i.e.* species that are excellent electron donors. The bulk electrical conductivity of the MOF is shown to increase from immeasurably low to 10^{−3} S cm^{−1}, following fullerene incorporation. The observed conductivity originates from electron donor–acceptor interactions, *i.e.* charge-transfer interactions – a conclusion that is supported by density functional theory calculations and by the observation of a charge-transfer-derived band in the electronic absorption spectrum of the hybrid material. Notably, the conductive version of the MOF retains substantial nanoscale porosity and continues to display a sizable internal surface area, suggesting potential future applications that capitalize on the ability of the material to sorb molecular species.

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Introduction

Metal–organic frameworks (MOFs) are a large and growing class of porous, or potentially porous, materials. They are characterized by well-ordered (*i.e.*, crystalline) combinations of multi-topic organic ligands connected by coordination bonds to one or more metal ions, or metal-ion-containing clusters.^{1–4} The enormous potential and experimentally realized compositional variety of MOFs,^{5–8} together with their typically large internal surface areas and their molecular-scale voids, has inspired exploration of myriad candidate applications including, but not limited to, gas storage and release,^{9,10} chemical separations,^{11,12} chemical sensing,¹³ catalysis,^{14–17} energy transfer,^{18–20} drug delivery,²¹ and solar fuel production.^{22–24} Electrically conductive MOFs, while few in absolute number, constitute an intriguing and increasingly important sub-class of materials, especially in

view of the insulating character (conductivity <10^{−10} S cm^{−1}) of most MOFs. The insulating versions typically make use of redox-resistant d(0) or d(10) metal ions that are energetically poorly matched or otherwise ineffective at interacting with linker-based π – π^* systems. Thus, nodes may behave as local insulators between otherwise potentially conductive arrays of linkers (and/or *vice versa*).

Nevertheless, the demonstrated feasibility of deploying electrically conductive MOFs in field-effect transistors (FETs),²⁵ chemiresistive sensors,²⁶ electrochromic devices,^{27,28} supercapacitors^{29,30} and batteries,³¹ has motivated the search for suitably conductive MOFs. The common design rule is to provide a low energy pathway for charge transport, and reported strategies include: (a) π stacking of conjugated organic ligands,³² (b) formulation of MOFs as laterally conjugated 2D coordination compounds,³³ (c) templating the growth of conductive polymers within suitably large MOF channels,^{34,35} (d) intentionally matching energies of frontier orbitals of metal-ion nodes and organic linkers,³⁶ (e) post-synthetic modification or augmentation of MOF nodes, either with metals featuring partially filled d-shells³² or with redox couples such as ferrocenium/ferrocene,³⁷ to yield redox-hopping-type conductivity, and (f) electrochemically oxidizing or reducing the linkers of solution-immersed MOFs, again to enable redox-hopping-type conduction.^{38,39}

The availability of free space within MOFs suggests another route to eliciting electrical conductivity, namely, the incorporation

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of guest molecules (electron acceptors or donors) that are complementary, in a charge-transfer sense, to MOF nodes or linkers.^{40,41} The approach has clear parallels, for example, to the formation of conductive, organic charge-transfer salts by combining (co-crystallizing) a strong electron donor (D), such as tetrathiafulvalene (TTF), with a strong electron acceptor (A) such as tetracyanoquinodimethane (TCNQ). Generally, in order to effectively transport charges, long-range ordering of donor and acceptor components is necessary. MOFs, with their inherent crystallinity and their ability to encapsulate guest molecules, offer platforms for achieving the desired long-range ordering of electronically complementary species. In a groundbreaking study, Talin *et al.* demonstrated that infiltration of the microporous MOF $\text{Cu}_3(\text{btc})_2$ (also known as HKUST-1; btc = 1,3,5-benzene-tricarboxylate) with TCNQ boosts the MOF's electrical conductivity by a remarkable six orders of magnitude (*i.e.*, from $10^{-8} \text{ S cm}^{-1}$ to 0.07 S cm^{-1}).⁴² The basis for increase is the ligation of open Cu(II) sites by TCNQ nitrile groups such that a continuous charge-transfer pathway is formed (each TCNQ is thought to connect to four copper ions, with the overall Cu(II)trimesate unit functioning as an electron donor, and TCNQ as an acceptor).⁴³ Among other D-A pairs incorporated within MOFs to study charge transfer are: (a) TTF/naphthalenetetracarboxydiimide (NDI),^{44,45} (b) oligothiophene/[6,6]-phenyl- C_{61} -butyric acid methyl ester (PCBM),^{44,46} and (c) *N,N'*-bis(4-pyridyl)-2,6-dipyrroliidyl naphthalenediimide/methylviologen.⁴⁷ However, only in case (c) has bulk electrical conductivity been investigated (with a 35-fold increase in conductivity accompanying viologen incorporation).

Among the challenges in using guest infiltrated MOFs for device applications are: (a) limited stability of the MOF, as most of the MOFs evaluated to date feature comparatively weakly ligating nodes containing Zn(II), Cu(II) or Ni(II), and (b) substantial loss of porosity upon guest inclusion. For example, after TCNQ infiltration, the BET surface area of HKUST-1 decreases from $1844 \pm 4 \text{ m}^2 \text{ g}^{-1}$ to $214 \pm 0.5 \text{ m}^2 \text{ g}^{-1}$. Hence it is of particular interest to design stable and porous electrically conductive MOFs. Stability can be defined in many ways. For oxidative heterogeneous catalysis of gas phase reactions, thermal stability is typically paramount, while for solution-phase electrocatalysis processes such as water splitting, aqueous stability may be paramount.

These stability considerations are well satisfied by MOFs featuring Zr(IV)-oxygen(anion) coordination as the basis for node and linker connectivity.^{48–51} With this in mind, we settled on a high-surface-area MOF, NU-901,^{28,52} as a guest-accessible, electron-donating scaffold, and the iconic electron acceptor, buckminsterfullerene (C_{60}) as the complementary component for studies of MOF electrical conductivity. Notably, there are previous reports on C_{60} incorporation within the MOF channels, the earliest being a report from Chae *et al.*⁵³ To our knowledge, however, this is the first report of host-guest interaction to induce electrical conductivity in an otherwise insulating zirconium based MOF.

NU-901 consists of $\text{Zr}_6(\mu_3\text{-O})_4(\mu_3\text{-OH})_4(\text{H}_2\text{O})_4(\text{OH})_4$ nodes and tetatopic 1,3,6,8-tetrakis(*p*-benzoate)pyrene (TBAPy⁴⁻) linkers. We hypothesized that NU-901 would be an attractive candidate for encapsulation of C_{60} and electronic activation due

to: (a) the availability of diamond-shaped channels that can withstand removal of synthesis solvent and that are reasonably well size-matched to C_{60} (van der Waals diameter $\sim 7 \text{ \AA}$), (b) the modest onset potential for electrochemical oxidation of NU-901,²⁸ which suggests that its linkers may be strong enough donors to form charge-transfer complexes with easily reducible fullerenes, and (c) the arrangement of the diamond-shaped voids, as defined by the *scu* topology of the MOF, which should allow for a spatially continuous network of charge-transfer type interactions (and, therefore, electrical conductivity) in the *a-b* plane.

Results and discussion

NU-901 was synthesized by a slightly modified literature procedure.⁵² Infiltration with C_{60} was performed by soaking a microcrystalline powder sample of activated (*i.e.*, solvent-evacuated and fully porous) NU-901 in a saturated *o*-dichlorobenzene solution of C_{60} for four days at 60°C . The powder X-ray diffraction (PXRD) pattern of NU-901- C_{60} matches well with the experimentally obtained and computationally simulated patterns for pure NU-901, implying that guest uptake does not alter the material's crystallinity (Fig. 2a). Preservation of microcrystal morphology was confirmed by comparing scanning electron microscopy images of NU-901 and NU-901- C_{60} (Fig. S1†). The presence of fullerene in NU-901- C_{60} was probed by Raman spectroscopy, based on 785 nm excitation (Fig. 2d). The scattering spectrum of NU-901- C_{60} displays peaks attributable to both the MOF and the fullerene, including the $\text{A}_g(1)$ vibration of C_{60} at 492 cm^{-1} . Consistent with interactions with the fullerene, the vibrational peaks for NU-901 are slightly shifted in NU-901- C_{60} .

The porosity of both NU-901 and NU-901- C_{60} was confirmed by nitrogen adsorption-desorption measurements (Fig. 2b). Both display approximately type I isotherms, albeit with slight hysteresis presumably arising from inter-crystallite macroporosity. Brunauer-Emmett-Teller (BET) analyses of the isotherms return gravimetric surface areas of $2120 \text{ m}^2 \text{ g}^{-1}$ for guest-free NU-901 and $1550 \text{ m}^2 \text{ g}^{-1}$ for NU-901- C_{60} . DFT pore-size distribution analyses of the isotherms reveal a large peak centered at 12 \AA , together with a small peak at 27 \AA . The former is consistent with the crystallographically observed diamond-shaped pores of NU-901. The latter we speculatively attribute to pores associated with defects.

After installation of C_{60} , the incremental pore volume corresponding to the diamond shaped channels decreased from $0.54 \text{ cm}^3 \text{ g}^{-1}$ to $0.38 \text{ cm}^3 \text{ g}^{-1}$, whereas the pore volume remains unchanged for pores generated by defect sites, suggesting fullerenes primarily reside within the diamond channels of the MOF (Fig. 2c). From the decrease in gravimetric pore volume, we estimate that the fullerene incorporation to be ~ 0.6 per node (~ 0.6 per linker-defined diamond void). Notably, DFT calculations (geometry optimized) show siting in diamond pores is overwhelmingly energetically favored (by 0.8 eV) compared to siting in the smaller pores defined by pairs of nodes along the *c* axis (Fig. S8b†). The representations in the bottom of Fig. 1 are derived from the DFT geometry.⁵⁴



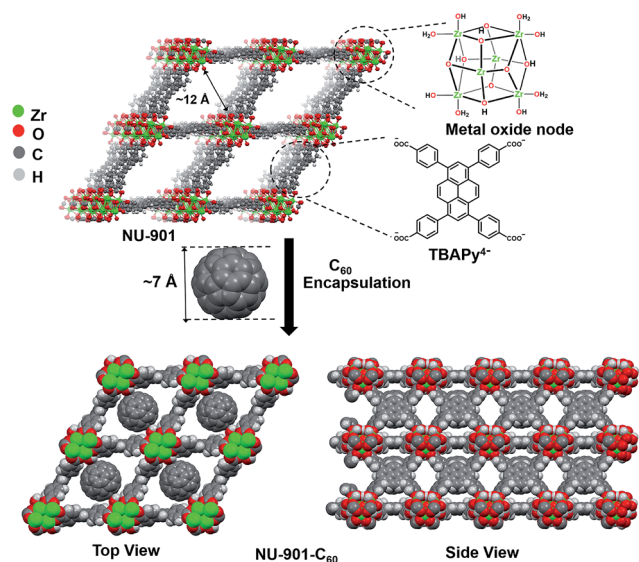


Fig. 1 Immobilization of C_{60} within diamond shaped channels of NU-901. The chemical structures of the individual components of NU-901 are shown. The diameter of C_{60} (~ 7 Å) is well suited for encapsulation in NU-901 with pore aperture of 12 Å. The top view and side view of the composite NU-901- C_{60} are shown. The top and side views are DFT-optimized structures in the limit of full occupancy of the diamond pores by the fullerene guest, i.e. 1 : 1 occupancy.

C_{60} encapsulation by NU-901 is accompanied by a change in color of the MOF from bright yellow to light brown. The visible color change is observable in corresponding diffuse reflectance UV-visible absorption spectra (Fig. S2a†). Guest-free NU-901 shows a broad absorption band beginning at 490 nm (2.53 eV) and maximizing in the UV, with no absorption beyond 490 nm. The spectrum of NU-901- C_{60} likewise displays the UV-blue band, but also presents a broad band extending from 500 nm to 700 nm (~ 1.77 eV). We attribute the new band to a donor/

acceptor (pyrene linker/ C_{60}) charge-transfer transition.⁵⁵ Consistent with the introduction of a strong electron acceptor,^{56,57} the encapsulated fullerene quenches the steady-state luminescence of the tetra-phenyl-pyrene linker by *ca.* 90%; see Fig. S2b.†

Qualitative assessments of electrical conductivity were made by first dropcasting MOF suspensions on interdigitated array electrodes (IDEs; 5 μ m gap) and then recording the current (I) transmitted between the electrodes, *via* electrode-gap-spanning MOF crystallites, in response to applied biases (V ; two-point measurements). NU-901- C_{60} samples yielded measurable currents whereas guest-free crystallites of NU-901 did not; see Fig. S7.† Unfortunately, we were unable with this approach to quantify the conductivity, σ , of NU-901- C_{60} . We therefore turned to macroscopic, two-point, I - V measurements using pressed pellets. The stability of the MOF pellets before and after C_{60} encapsulation was verified by PXRD and nitrogen adsorption isotherms (Fig. S3 and S4†). Consistent with IDE measurements, NU-901 alone displayed no measurable conductivity (where the lower limit of detection is estimated, from the magnitude of electrode noise, to be about 10^{-14} S cm^{-1}). In contrast (Fig. 3), NU-901- C_{60} shows remarkable electrical conductivity of $\sigma \sim 10^{-3}$ S cm^{-1} ,⁵⁸ i.e. eleven orders of magnitude (or more) greater than NU-901 alone, and similar to that of organic semiconductors ($\sigma > 10^{-6}$ S cm^{-1}).⁵⁹ Unlike conventional organic semiconductors, however, the MOF/fullerene material is characterized by substantial internal surface area and considerable molecular-scale porosity.

We reasoned that if TBAPy⁴⁻/ C_{60} donor/acceptor interactions are responsible for the observed electrical conductivity, increasing the strength of the MOF linker as an electron donor would increase the conductivity. We therefore replaced TBAPy⁴⁻ with a tetra-amino (phenyl-substituted) analogue.⁶⁰ PXRD (Fig. S6†) and SEM (Fig. S1†) assessments of NU-901-amino- C_{60} indicated formation of a material that is structurally similar (Fig. S5†) to NU-901- C_{60} (see (Fig. S5†)). N_2 adsorption and desorption measurements established the permanent porosity of the material, and yielded BET surface area of 980 $\text{m}^2 \text{g}^{-1}$ for



Fig. 2 (a) PXRD pattern of simulated NU-901, experimental NU-901, and experimental NU-901- C_{60} . (b) N_2 adsorption-desorption isotherms (77 K) (NU-901 and NU-901- C_{60}), (c) density functional theory (DFT) pore size distribution (NU-901- C_{60} and NU-901) and, (d) Raman spectra (NU-901, NU-901- C_{60} and C_{60}).



Fig. 3 Current (I) vs. voltage (V) plot for pressed pellets of NU-901 and NU-901- C_{60} . The scan was performed from -2 V to 2 V and the scan rate was 50 mV s^{-1} .

NU-901-amino and $370 \text{ m}^2 \text{ g}^{-1}$ for NU-901-amino- C_{60} (Fig. S6†). IDE based measurements show that C_{60} incorporation imparts electrical conductivity to NU-901-amino, with the observed currents being larger for NU-901-amino- C_{60} than for NU-901- C_{60} ,⁶¹ see Fig. S7.†

Next we turn our attention to density functional theory (the details of the methods used are in the ESI†) to gain further insight into the nature of the observed electron-donor/electron-acceptor interactions and their manifestation as electrical conductivity in the MOF/fullerene assemblies. The results of electronic density of states (DOS) calculations for NU-901, NU-901-amino, NU-901- C_{60} and NU-901-amino- C_{60} are summarized in Fig. S9.† Briefly, for the guest-free MOFs, both the valence band maximum (VBM) and conduction band minimum (CBM) are dominated by contributions from linker p orbitals (*i.e.*, those that contribute to the linkers' aromatic systems). For the C_{60} -containing materials, the VBM remains predominantly linker defined, but the CBM consists of fullerene p orbitals, consistent with donor/acceptor complex formation. The calculated CBM – VBM difference, *i.e.* the nominal bandgap, decreases by 1.2 eV with incorporation of C_{60} , a result that is qualitatively replicated in the visible-region reflectance spectra for fullerene-containing *versus* fullerene-free versions of the MOF (note that DFT methods often underestimate absolute magnitudes of bandgaps, particularly when local functionals like PBE are used).^{62,63}

Given the spatial and energetic proximity of the putative donors and acceptors and their associated bands, we also calculated charge-transfer integrals for relevant clusters cut from the NU-901- C_{60} and NU-901-amino- C_{60} periodic structures (Table S1†). Both the electron-transfer and hole-transfer integrals are larger (by 2- and 4-fold, respectively) for NU-901-amino- C_{60} compared to NU-901- C_{60} , consistent with larger IDE currents for the amino version.³⁰

Conclusions

In summary, we find that the zirconium-based MOF, NU-901, can be rendered electronically conductive by installing isolated C_{60} molecules in roughly half the linker-defined, diamond shaped pores presented by the MOF. Indeed, the experimentally evaluated conductivity increases from immeasurably small ($<10^{-14} \text{ S cm}^{-1}$) to $\sim 10^{-3} \text{ S cm}^{-1}$. The basis for the conductivity is in electron-donor (pyrene)/electron-acceptor (fullerene) interactions, as evidenced, in part, by the appearance of a visible-region charge-transfer band and by the results of DFT band structure calculations. Notably, the MOF internal surface area (gravimetric surface area) decreases by only about 27% with fullerene incorporation while the N_2 -accessible void volume (as expressed gravimetrically) decreases by only about a third. DFT computes that appending NH_2 substituents (strongly electron-donating groups) to the linker phenyl groups of NU-901 increases hole- and electron-transfer integrals, which is consistent with the increased conductivity measured experimentally. These findings point to the potential for systematic chemical tunability of the observed conductivity.

Finally, while electrocatalysis applications are beyond the scope of this initial report, it is worth considering how much conductivity is needed for a MOF to function effectively in that context. A common motif for electrochemical applications of MOFs is a thin film (a fraction of a micron to several microns thick) sited on a conventionally conductive platform – for example, a planar glassy carbon or platinum electrode. A benchmark catalytic current density for solar-driven electrochemical processes is 10 mA cm^{-2} . For a hypothetical 10 micron thick MOF film having the modest conductivity observed here, *i.e.*, $\sim 10^{-3} \text{ S cm}^{-1}$, and operating at 10 mA cm^{-2} , the energetic penalty (*i-R* drop) due to film resistivity would be just 10 mV – a value barely within the reproducibility of kinetic overpotentials for many electrocatalytic systems. We suggest that for these types of applications, conductivities in the range of $10^{-3} \text{ S cm}^{-1}$, despite falling in the semiconducting regime (phenomenologically),⁶⁴ will typically be sufficient.

Author contributions

J. T. H., O. K. F. and S. G. conceived the project, design the experiments and wrote the manuscript. S. G. performed the synthesis and experiments. D. R. and C. J. C. contributed to the theoretical studies. K. O. and C. W. C. assist in performing Raman spectroscopy and IDE measurements respectively. S. J. G., T. L. and A. A. contributed in amino linker synthesis. Y. C. assist in performing diffuse reflectance UV measurements.

Conflicts of interest

The authors declare no competing financial interests.

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Notes and references

- 1 H. Furukawa, K. E. Cordova, M. O'Keeffe and O. M. Yaghi, *Science*, 2013, **341**, 1230444.
- 2 G. Ferey, *Chem. Soc. Rev.*, 2008, **37**, 191–214.
- 3 O. K. Farha and J. T. Hupp, *Acc. Chem. Res.*, 2010, **43**, 1166–1175.
- 4 S. Horike, S. Shimomura and S. Kitagawa, *Nat. Chem.*, 2009, **1**, 695.
- 5 Y. G. Chung, J. Camp, M. Haranczyk, B. J. Sikora, W. Bury, V. Krungleviciute, T. Yildirim, O. K. Farha, D. S. Sholl and R. Q. Snurr, *Chem. Mater.*, 2014, **26**, 6185–6192.



- 6 C. E. Wilmer, M. Leaf, C. Y. Lee, O. K. Farha, B. G. Hauser, J. T. Hupp and R. Q. Snurr, *Nat. Chem.*, 2011, **4**, 83.
- 7 P. G. Boyd, Y. Lee and B. Smit, *Nat. Rev. Mater.*, 2017, **2**, 17037.
- 8 D. Nazarian, P. Ganesh and D. S. Sholl, *J. Mater. Chem. A*, 2015, **3**, 22432–22440.
- 9 M. Dincă and J. R. Long, *Angew. Chem., Int. Ed.*, 2008, **47**, 6766–6779.
- 10 O. K. Farha, A. Özgür Yazaydın, I. Eryazici, C. D. Malliakas, B. G. Hauser, M. G. Kanatzidis, S. T. Nguyen, R. Q. Snurr and J. T. Hupp, *Nat. Chem.*, 2010, **2**, 944.
- 11 J.-R. Li, R. J. Kuppler and H.-C. Zhou, *Chem. Soc. Rev.*, 2009, **38**, 1477–1504.
- 12 Z. Bao, G. Chang, H. Xing, R. Krishna, Q. Ren and B. Chen, *Energy Environ. Sci.*, 2016, **9**, 3612–3641.
- 13 L. E. Kreno, K. Leong, O. K. Farha, M. Allendorf, R. P. Van Duyne and J. T. Hupp, *Chem. Rev.*, 2012, **112**, 1105–1125.
- 14 T. Islamoglu, S. Goswami, Z. Li, A. J. Howarth, O. K. Farha and J. T. Hupp, *Acc. Chem. Res.*, 2017, **50**, 805–813.
- 15 V. Bernales, M. A. Ortuño, D. G. Truhlar, C. J. Cramer and L. Gagliardi, *ACS Cent. Sci.*, 2018, **4**, 5–19.
- 16 J. Lee, O. K. Farha, J. Roberts, K. A. Scheidt, S. T. Nguyen and J. T. Hupp, *Chem. Soc. Rev.*, 2009, **38**, 1450–1459.
- 17 J. Gascon, A. Corma, F. Kapteijn and F. X. Llabrés i Xamena, *ACS Catal.*, 2014, **4**, 361–378.
- 18 C. A. Kent, B. P. Mehl, L. Ma, J. M. Papanikolas, T. J. Meyer and W. Lin, *J. Am. Chem. Soc.*, 2010, **132**, 12767–12769.
- 19 W. A. Maza, R. Padilla and A. J. Morris, *J. Am. Chem. Soc.*, 2015, **137**, 8161–8168.
- 20 S. Goswami, L. Ma, A. B. F. Martinson, M. R. Wasielewski, O. K. Farha and J. T. Hupp, *ACS Appl. Mater. Interfaces*, 2016, **8**, 30863–30870.
- 21 P. Horcajada, T. Chalati, C. Serre, B. Gillet, C. Sebrie, T. Baati, J. F. Eubank, D. Heurtaux, P. Clayette, C. Kreuz, J.-S. Chang, Y. K. Hwang, V. Marsaud, P.-N. Bories, L. Cynober, S. Gil, G. Férey, P. Couvreur and R. Gref, *Nat. Mater.*, 2009, **9**, 172.
- 22 J.-L. Wang, C. Wang and W. Lin, *ACS Catal.*, 2012, **2**, 2630–2640.
- 23 I. Hod, M. D. Sampson, P. Deria, C. P. Kubiak, O. K. Farha and J. T. Hupp, *ACS Catal.*, 2015, **5**, 6302–6309.
- 24 N. Kornienko, Y. Zhao, C. S. Kley, C. Zhu, D. Kim, S. Lin, C. J. Chang, O. M. Yaghi and P. Yang, *J. Am. Chem. Soc.*, 2015, **137**, 14129–14135.
- 25 G. Wu, J. Huang, Y. Zang, J. He and G. Xu, *J. Am. Chem. Soc.*, 2017, **139**, 1360–1363.
- 26 M. G. Campbell, S. F. Liu, T. M. Swager and M. Dincă, *J. Am. Chem. Soc.*, 2015, **137**, 13780–13783.
- 27 K. AlKaabi, C. R. Wade and M. Dincă, *Chem*, 2016, **1**, 264–272.
- 28 C.-W. Kung, T. C. Wang, J. E. Mondloch, D. Fairen-Jimenez, D. M. Gardner, W. Bury, J. M. Klingsporn, J. C. Barnes, R. Van Duyne, J. F. Stoddart, M. R. Wasielewski, O. K. Farha and J. T. Hupp, *Chem. Mater.*, 2013, **25**, 5012–5017.
- 29 D. Sheberla, J. C. Bachman, J. S. Elias, C.-J. Sun, Y. Shao-Horn and M. Dincă, *Nat. Mater.*, 2016, **16**, 220.
- 30 K. M. Choi, H. M. Jeong, J. H. Park, Y.-B. Zhang, J. K. Kang and O. M. Yaghi, *ACS Nano*, 2014, **8**, 7451–7457.
- 31 D. Wu, Z. Guo, X. Yin, Q. Pang, B. Tu, L. Zhang, Y.-G. Wang and Q. Li, *Adv. Mater.*, 2014, **26**, 3258–3262.
- 32 S. S. Park, E. R. Hontz, L. Sun, C. H. Hendon, A. Walsh, T. Van Voorhis and M. Dincă, *J. Am. Chem. Soc.*, 2015, **137**, 1774–1777.
- 33 D. Sheberla, L. Sun, M. A. Blood-Forsythe, S. Er, C. R. Wade, C. K. Brozek, A. Aspuru-Guzik and M. Dincă, *J. Am. Chem. Soc.*, 2014, **136**, 8859–8862.
- 34 T. C. Wang, I. Hod, C. O. Audu, N. A. Vermeulen, S. T. Nguyen, O. K. Farha and J. T. Hupp, *ACS Appl. Mater. Interfaces*, 2017, **9**, 12584–12591.
- 35 B. Le Ouay, M. Boudot, T. Kitao, T. Yanagida, S. Kitagawa and T. Uemura, *J. Am. Chem. Soc.*, 2016, **138**, 10088–10091.
- 36 L. Sun, T. Miyakai, S. Seki and M. Dincă, *J. Am. Chem. Soc.*, 2013, **135**, 8185–8188.
- 37 I. Hod, W. Bury, D. M. Gardner, P. Deria, V. Roznyatovskiy, M. R. Wasielewski, O. K. Farha and J. T. Hupp, *J. Phys. Chem. Lett.*, 2015, **6**, 586–591.
- 38 S. R. Ahrenholtz, C. C. Epley and A. J. Morris, *J. Am. Chem. Soc.*, 2014, **136**, 2464–2472.
- 39 D. M. D'Alessandro, *Chem. Commun.*, 2016, **52**, 8957–8971.
- 40 M. D. Allendorf, M. E. Foster, F. Léonard, V. Stavila, P. L. Feng, F. P. Doty, K. Leong, E. Y. Ma, S. R. Johnston and A. A. Talin, *J. Phys. Chem. Lett.*, 2015, **6**, 1182–1195.
- 41 J. Liu, T. Wächter, A. Irmeler, P. G. Weidler, H. Gliemann, F. Pauly, V. Mugnaini, M. Zharnikov and C. Wöll, *ACS Appl. Mater. Interfaces*, 2015, **7**, 9824–9830.
- 42 A. A. Talin, A. Centrone, A. C. Ford, M. E. Foster, V. Stavila, P. Haney, R. A. Kinney, V. Szalai, F. El Gabaly, H. P. Yoon, F. Léonard and M. D. Allendorf, *Science*, 2014, **343**, 66–69.
- 43 X. Nie, A. Kulkarni and D. S. Sholl, *J. Phys. Chem. Lett.*, 2015, **6**, 1586–1591.
- 44 C. F. Leong, B. Chan, T. B. Faust and D. M. D'Alessandro, *Chem. Sci.*, 2014, **5**, 4724–4728.
- 45 Z. Guo, D. K. Panda, M. A. Gordillo, A. Khatun, H. Wu, W. Zhou and S. Saha, *ACS Appl. Mater. Interfaces*, 2017, **9**, 32413–32417.
- 46 K. Leong, M. E. Foster, B. M. Wong, E. D. Spoerke, D. Van Gough, J. C. Deaton and M. D. Allendorf, *J. Mater. Chem. A*, 2014, **2**, 3389–3398.
- 47 Z. Guo, D. K. Panda, K. Maity, D. Lindsey, T. G. Parker, T. E. Albrecht-Schmitt, J. L. Barreda-Esparza, P. Xiong, W. Zhou and S. Saha, *J. Mater. Chem. C*, 2016, **4**, 894–899.
- 48 V. Bon, I. Senkovska, I. A. Baburin and S. Kaskel, *Cryst. Growth Des.*, 2013, **13**, 1231–1237.
- 49 M. Kandiah, M. H. Nilsen, S. Usseglio, S. Jakobsen, U. Olsbye, M. Tilset, C. Larabi, E. A. Quadrelli, F. Bonino and K. P. Lillerud, *Chem. Mater.*, 2010, **22**, 6632–6640.
- 50 A. J. Howarth, Y. Liu, P. Li, Z. Li, T. C. Wang, J. T. Hupp and O. K. Farha, *Nat. Rev. Mater.*, 2016, **1**, 15018.
- 51 Y. Bai, Y. Dou, L.-H. Xie, W. Rutledge, J.-R. Li and H.-C. Zhou, *Chem. Soc. Rev.*, 2016, **45**, 2327–2367.
- 52 P. Deria, J. Yu, T. Smith and R. P. Balaraman, *J. Am. Chem. Soc.*, 2017, **139**, 5973–5983.



- 53 H. K. Chae, D. Y. Siberio-Pérez, J. Kim, Y. Go, M. Eddaoudi, A. J. Matzger, M. O'Keeffe and O. M. Yaghi, *Nature*, 2004, **427**, 523.
- 54 The DFT calculations also indicate a slight change pore geometry in response to complete fullerene incorporation (1 : 1 incorporation). Based on PXRD measurements, the anticipated changes are absent suggesting minor differences in framework coordinates for C₆₀ incorporation at 0.6 : 1 *versus* 1 : 1. As yet, we have been unable to boost the loading beyond 0.6 : 1. Nevertheless, it will be interesting to see (eventually) if higher loading translates not only into detectable MOF structural changes, but also significant further changes in electronic conductivity (see subsequent results and discussion).
- 55 Based on the (+/0) redox potential of TBAPy linkers in NU-901 (1.4 V *vs.* Ag/AgCl) and (−/0) redox potential of C₆₀ (−0.33 V *vs.* Ag/AgCl), the charge transfer (CT) band should appear near 1.8 eV (690 nm). The experimental band maximum is at near 550 nm and the onset is near 700 nm (1.77 eV).
- 56 D. M. Guldi, F. Spänig, D. Kreher, I. F. Perepichka, C. van der Pol, M. R. Bryce, K. Ohkubo and S. Fukuzumi, *Chem.–Eur. J.*, 2008, **14**, 250–258.
- 57 P. Dallas, G. Rogers, B. Reid, R. A. Taylor, H. Shinohara, G. A. D. Briggs and K. Porfyrakis, *Chem. Phys.*, 2016, **465–466**, 28–39.
- 58 We recognize that two-point measurements with pelletized samples do exclude effects due to electrode/sample contact resistance and do not distinguish between inter-crystallite and intra-crystallite contributions to sample resistivity. Thus, the σ value of $\sim 10^{-3}$ S cm^{−1} should be regarded as a lower-limit estimate for electrical conductivity within a single crystallite. Also obscured by pellet-based measurements is any conductivity anisotropy – specifically, differences in conductivity in the MOF *ab* plane *versus* in the *c* direction (the direction aligned with the diamond-shaped channels).
- 59 D. Hinderberger, in *EPR Spectroscopy: Applications in Chemistry and Biology*, ed. M. Drescher and G. Jeschke, Springer Berlin Heidelberg, Berlin, Heidelberg, 2012, pp. 67–89, DOI: 10.1007/128_2011_236.
- 60 T. Islamoglu, M. A. Ortuño, E. Prousaloglou, A. J. Howarth, N. A. Vermeulen, A. Atilgan, A. M. Asiri, C. J. Cramer and O. K. Farha, *Angew. Chem.*, 2018, **130**, 1967–1971.
- 61 Given uncertainties in MOF loading on the IDE platforms, we are unable to say with certainty that the conductivity of NU-901-amino-C₆₀ exceeds that of NU-901-C₆₀. Unfortunately, we lack sufficient NU-901-amino-C₆₀ to extend the study pelletized samples, *i.e.* samples that could yield quantitative conductivity data.
- 62 A. J. Garza and G. E. Scuseria, *J. Phys. Chem. Lett.*, 2016, **7**, 4165–4170.
- 63 J. P. Perdew, W. Yang, K. Burke, Z. Yang, E. K. U. Gross, M. Scheffler, G. E. Scuseria, T. M. Henderson, I. Y. Zhang, A. Ruzsinszky, H. Peng, J. Sun, E. Trushin and A. Görling, *Proc. Natl. Acad. Sci. U. S. A.*, 2017, **114**, 2801–2806.
- 64 Definitive demonstration (or not) of the putative semiconducting behavior will require variable temperature conductivity measurements. We anticipate reporting on these in due course.

