

# Vinylene-Linked Covalent Organic Frameworks by Base-Catalyzed Aldol Condensation

Amitava Acharjya, Pradip Pachfule, Jérôme Roeser, Franz-Josef Schmitt, and Arne Thomas\*

**Abstract:** Two 2D covalent organic frameworks (COFs) linked by vinylene ( $-\text{CH}=\text{CH}-$ ) groups (V-COF-1 and V-COF-2) are synthesized by exploiting the electron deficient nature of the aromatic *s*-triazine unit of  $C_3$ -symmetric 2,4,6-trimethyl-*s*-triazine (TMT). The acidic terminal methyl hydrogens of TMT can easily be abstracted by a base, resulting in a stabilized carbanion, which further undergoes aldol condensation with multitopic aryl aldehydes to be reticulated into extended crystalline frameworks (V-COFs). Both V-COF-1 (with terephthalaldehyde (TA)) and V-COF-2 (with 1,3,5-tris(*p*-formylphenyl)benzene (TFPB)) are polycrystalline and exhibit permanent porosity and BET surface areas of  $1341 \text{ m}^2 \text{ g}^{-1}$  and  $627 \text{ m}^2 \text{ g}^{-1}$ , respectively. Owing to the close proximity ( $3.52 \text{ \AA}$ ) of the pre-organized vinylene linkages within adjacent 2D layers stacked in eclipsed fashion, [2+2] photo-cycloaddition in V-COF-1 formed covalent crosslinks between the COF layers.

The discovery of covalent organic frameworks (COFs) demonstrated that organic building blocks could be reticulated via strong covalent bonds into well-ordered 2D and 3D extended crystalline frameworks.<sup>[1–6]</sup> These ordered crystalline materials feature defined porosities and functionalities and have consequently garnered increasing attention in various applications such as gas storage and separation,<sup>[7,8]</sup> energy storage,<sup>[9,10]</sup> photovoltaics,<sup>[9,11]</sup> opto-electronics,<sup>[12,13]</sup> proton and ion conduction,<sup>[14–19]</sup> and heterogeneous catalysis.<sup>[20–26]</sup> The formation of such extended crystalline frameworks is generally achieved by implementing reversible condensation reactions to polymerize rigid building blocks. Careful control over the rate of reversible bond formation allow the construction of the desired crystalline framework by applying the principles of reticular chemistry.<sup>[27–29]</sup> Until recently, linkages (that is, bonds formed to reticulate the building blocks) reported for COF synthesis were mostly based on reversible B–O,<sup>[1,2,16,30,31]</sup> B–N,<sup>[32]</sup> Si–O,<sup>[33,34]</sup> and C–

N bond formation.<sup>[35–39]</sup> The reversible nature of those linkages affected by the insertion–deletion of water molecules, allow the formation of the desired extended crystalline frameworks, but was also identified to be detrimental in terms of chemical stability, especially in aqueous media.<sup>[28,40]</sup> Indeed, although thermally robust, many COFs are prone to hydrolysis and structural collapse in aqueous/acidic media or even in contact with moisture. Several strategies, including keto–enol tautomerization,<sup>[35]</sup> interlayer stacking optimization,<sup>[36]</sup> or post-functionalization<sup>[41,42]</sup> have been applied to circumvent these limitations, yielding COFs featuring higher chemical and thermal stability.

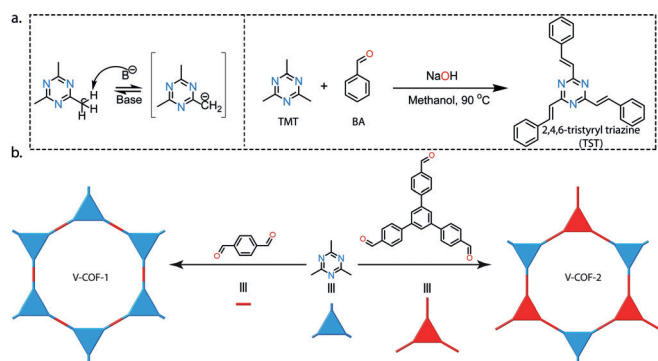
In this context, recently reported 2D COFs based on cyanovinylene ( $-\text{CH}=\text{C}(\text{CN})-$ ) linkages hold the promise for the generation of chemically stable COFs as the cyanovinylene linkage is significantly less prone to hydrolysis compared to boroxine, boronate ester, or imine linkages that are mostly used for the synthesis of COFs.<sup>[10,43]</sup> The challenge to crystallize a COF by reversible  $-\text{CH}=\text{C}(\text{CN})-$  bond formation was overcome by implementing reversible Knoevenagel condensation of 1,4-phenylenediacetonitrile (PDAN) with multitopic aldehyde-functionalized building blocks. Such COFs featuring fully conjugated backbones are also promising materials in terms of optoelectronic or photovoltaic applications.<sup>[44]</sup>

Herein, we report the synthesis of two purely vinylene ( $-\text{CH}=\text{CH}-$ ) linked 2D COFs (V-COF-1 and V-COF-2) achieved by base-catalyzed reversible aldol condensation, exploiting the highly electron deficient *s*-triazine core of 2,4,6-trimethyl-*s*-triazine (TMT). The methyl protons of TMT are acidic and undergo base-catalyzed aldol condensation with benzaldehyde to yield 2,4,6-tristyryl triazine quantitatively (Scheme 1a).<sup>[45,46]</sup> V-COF-1 and V-COF-2 were obtained by extending this model reaction to terephthalaldehyde (TA) and 1,3,5-tris(4-formyl)phenyl benzene (TFPB) (Scheme 1b). During our investigation on the base-catalyzed pathway, Yaghi et al. reported the acid-catalyzed synthesis of a biphenyl analogue of V-COF-1, which however could not be synthesized after extensive trials employing basic conditions.<sup>[47]</sup> On the other hand, we were so far also unable to synthesize crystalline V-COF-1 and -2 using the reported acid-catalyzed pathway, showing that the choice of reaction conditions can be crucial to obtain V-COFs with varying backbone structure. V-COF-1 and V-COF-2, possess surface areas ( $\text{SA}_{\text{BET}}$ ) of  $1341 \text{ m}^2 \text{ g}^{-1}$  and  $627 \text{ m}^2 \text{ g}^{-1}$ , respectively. Although chemically and thermally stable, we identified that V-COF-1 is photo-responsive, as the pre-organized vinylene groups of adjacent 2D layers offer a platform for [2+2] cycloaddition upon UV/Vis irradiation fulfilling Schmidt's criteria for photocyclization.<sup>[48]</sup>

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Supporting information and the ORCID identification number(s) for the author(s) of this article can be found under:  
<https://doi.org/10.1002/anie.201905886>.

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**Scheme 1.** V-COF synthesis. a) Illustration of the base-catalyzed aldol condensation of 2,4,6-trimethyl-s-triazine (TMT) and benzaldehyde (BA) to yield 2,4,6-tristyryl-s-triazine (TST). b) Representation of reticulation of the crystalline V-COFs by condensation of 2,4,6-trimethyl-s-triazine with ditopic or tritopic aldehydes, respectively.

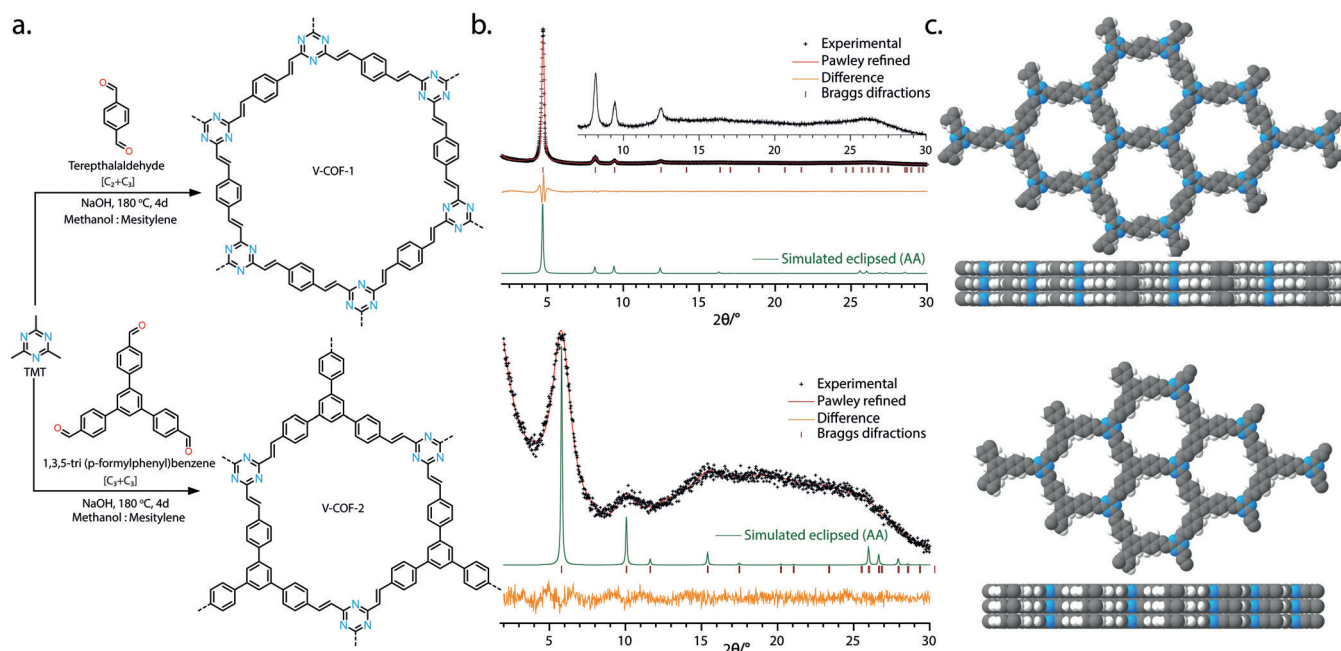
Syntheses of V-COF-1 and V-COF-2 were performed by solvothermal condensation of TMT with TA and TFPB, in 7:1 and 1:1 (methanol/mesitylene) solvent mixtures, respectively, for 4 days at 180 °C in presence of NaOH as base (Figure 1 a). Both the COFs were collected as light-yellow polycrystalline solids by filtration and further washed with methanol, water, and acetone before being dried under vacuum at 100 °C overnight.

Crystallinity of V-COF-1 and V-COF-2 was assessed by powder X-ray diffraction (PXRD) analyses. The experimental PXRD pattern of V-COF-1 and V-COF-2 confirmed the formation of a crystalline framework with no evidence of remaining starting material (Supporting Information, Figure S2). Given the connectivity of the building blocks, several

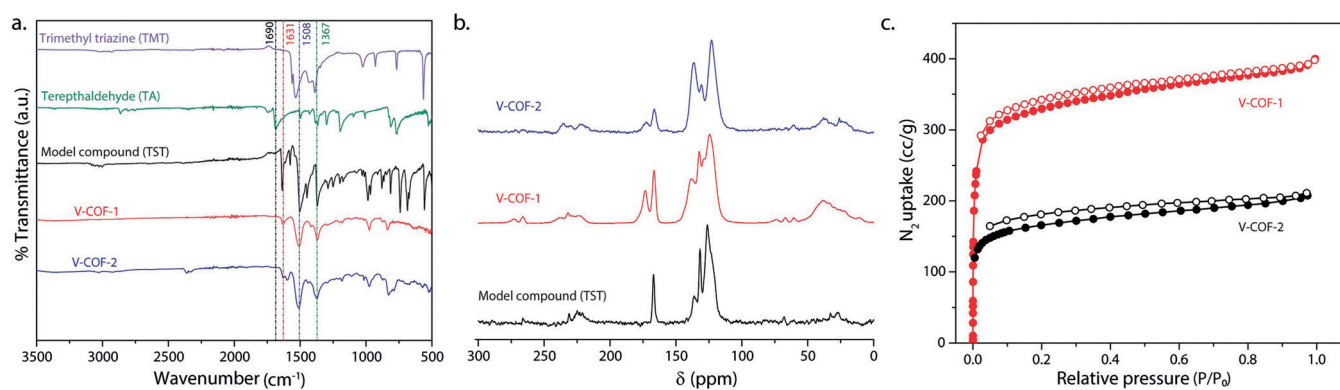
models with different stacking modes were generated in hexagonal 2D nets with **hcb** topology (Supporting Information, Section S4). A good match was found between the experimentally obtained PXRD patterns and the calculated patterns of a fully eclipsed model (Figure 1; Supporting Information, Figures S3, S5). The final lattice parameters were extracted after Pawley refinement and V-COF-1 was found to crystallize in a hexagonal unit cell ( $P6/m$ ,  $a=b=21.6934$  Å,  $c=3.5232$  Å,  $R_p=2.66\%$ ;  $R_{wp}=3.40\%$ ) with similar unit cell parameters as the isoreticular LZU-1 constructed from benzene nodes and imine linkages.<sup>[49]</sup> V-COF-2 was poorly crystalline as evidenced by the broad diffraction peaks observed experimentally, but was found to crystallize in a similar hexagonal unit cell (space group:  $P\bar{6}$ ,  $a=b=17.5632$  Å,  $c=3.4544$  Å,  $R_p=2.82\%$ ;  $R_{wp}=3.58\%$ ) (Figure 1; Supporting Information, Figures S4, S6).

Fourier-transform infrared (FTIR) spectroscopy analyses were applied to assess the structural integrity of the crystalline framework (Figure 2 a). Complete disappearance of  $-C=O-$  stretching frequency of starting aldehyde monomers at  $1690\text{ cm}^{-1}$  and appearance of a new band at  $1631\text{ cm}^{-1}$  attributed to  $-C=C-$  stretching indicated complete condensation of the starting building blocks and the successful formation of the vinylenic linkage in V-COF-1 and V-COF-2.

$^{13}\text{C}$  cross-polarization magic-angle-spinning (CP-MAS) NMR spectroscopy confirmed the full conversion of starting monomers as no residual carbonyl resonance located at  $\delta \approx 190$  ppm could be detected. The two expected signals for the vinylenic ( $-C=C-$ ) carbons at about 138.1 ppm and 132.1 ppm could be unambiguously assigned within the aromatic carbon signals (Figure 2 b; Supporting Information, Figure S7). V-COF-1 and V-COF-2 surprisingly exhibited two



**Figure 1.** Synthesis and structural characterization of V-COFs. a) Synthetic conditions for the reticulation of V-COF-1 and V-COF-2. b) PXRD patterns of V-COF-1 and V-COF-2: experimental patterns (black), Pawley refined profile curves (red), Bragg diffractions (brown), difference (orange), simulated patterns for the eclipsed layer stackings (green), and c) structural models of both V-COF-1 and V-COF-2 with their respective stacking patterns.



**Figure 2.** Structural characterization of V-COFs. a) FTIR analyses of V-COFs compared with the starting materials (TMT and TA) and the model compound (TST). b)  $^{13}\text{C}$  CP-MAS NMR spectra of V-COF-1 and V-COF-2 and model compound (TST). c)  $\text{N}_2$  sorption measurements of V-COF-1 and V-COF-2 powders.

distinct signals for the aromatic triazine carbon atoms at about 172.9 and about 166.4 ppm in the  $^{13}\text{C}$  CP-MAS NMR spectra, respectively, while just one peak, as expected, was observed for the molecular model compounds. Splitting of the triazine signals has been also observed in the solid-state  $^{13}\text{C}$  NMR spectrum of melamine and was attributed to different H-bonding environment leading to two chemically inequivalent carbon atoms.<sup>[50,51]</sup> However, this explanation could be ruled out here, since there is no electron-deficient hydrogen donor within the building blocks. Instead, the second triazine environment is probably caused by a [2+2] cycloaddition between vinylene groups.<sup>[52–54]</sup> This is corroborated by the presence of a broad signal in the aliphatic region ( $\delta \approx 38.1$  ppm) indicating the formation of cyclobutane moieties in the COFs (see below).

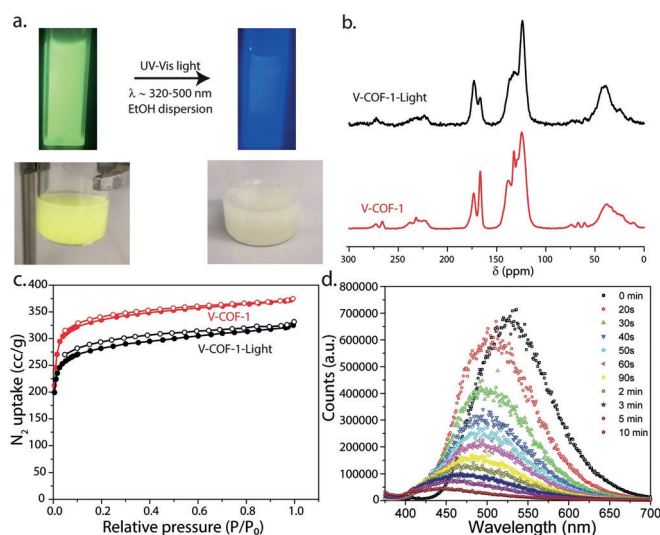
The permanent porosity of V-COF-1 and V-COF-2 was evaluated by low-pressure nitrogen ( $\text{N}_2$ ) and argon (Ar) sorption studies on evacuated samples, at 77 K and 87 K, respectively. A steep gas uptake in the low relative pressure range ( $p/p_0 < 0.05$ ) of the  $\text{N}_2$  adsorption branch shows the microporosity of the samples (Figure 2c). The Brunauer–Emmett–Teller surface areas ( $\text{SA}_{\text{BET}}$ ) calculated from the  $\text{N}_2$  adsorption were found to be  $1341 \text{ m}^2 \text{ g}^{-1}$  and  $627 \text{ m}^2 \text{ g}^{-1}$ , respectively. The  $\text{SA}_{\text{BET}}$  of V-COF-1 is almost twice as high as the  $\text{SA}_{\text{BET}}$  reported for the isorecticular LZU-1 COF ( $729 \text{ m}^2 \text{ g}^{-1}$ ).<sup>[49,55]</sup> The corresponding pore size distribution of 1.69 nm was derived by fitting the Ar adsorption branch data at 87 K with a cylindrical quenched-solid density functional theory (QSDFT) model, and matched well with the theoretically calculated pore size of 1.6 nm of the proposed eclipsed structural model (Supporting Information, Figure S8). The pore size distribution of V-COF-2 was found to be centered at 0.82 nm, derived by fitting the  $\text{N}_2$  adsorption branch data at 77 K with the QSDFT model and again matched well with theoretically calculated pore size of 0.8 nm (Supporting Information, Figure S9).

Architectural stability of the V-COF-1 and V-COF-2 was investigated by thermogravimetric analysis (TGA) under  $\text{N}_2$  atmosphere (Supporting Information, Figures S10, S11). After an initial weight loss of about 2 wt% owing to desorption of adsorbed solvents below  $100^\circ\text{C}$ , the frame-

works remained stable until  $400^\circ\text{C}$ . Chemical stability of crystalline V-COF-1 and V-COF-2 was further investigated in harsh acidic and basic conditions (Supporting Information, Figures S12, S13). Both V-COF-1 and V-COF-2 were found to be stable in concentrated acidic and basic medium for at least 4 days. V-COF-1 was also found to be stable in common organic solvents such as THF, acetone, and DMF, for 4 days, with no signs of decomposition (Supporting Information, Figure S14).

Considering the high crystallinity and conjugated structure, we investigated the optical properties of V-COF-1. The diffuse-reflectance UV/Vis (UV-DRS) spectrum of V-COF-1 shows a distinct red-shift in absorbance with respect to the monomers (TA and TMT), indicating a higher degree of conjugation (Supporting Information, Figure S15). V-COF-1 shows an absorbance edge at about 410 nm, whereas the absorbance tail is extended up to about 535 nm. Interestingly, V-COF-1 was found to be photosensitive, as the light-yellow color of pristine COF powders slowly faded with time upon sunlight irradiation, indicating a structural change within V-COF-1 (Figure 3a). To further elucidate the effect of light irradiation, V-COF-1 powder was irradiated directly with UV/Vis light ( $\lambda \approx 320\text{--}500$  nm) considering that the maximum absorbance of the COF lies within this wavelength range (Supporting Information, Section S10).  $^{13}\text{C}$  CP-MAS NMR spectra of the irradiated (48 h) sample revealed a cycloaddition reaction between the COF layers. Thus, the vinylene carbon signal ( $\delta \approx 138.1, 132.4$  ppm) intensity significantly decreased and the ratio of the triazine carbon signals (at  $\delta \approx 172.9$  and  $\delta \approx 166.4$ ) reversed compared to the pristine COF (Figure 3b). Finally, a broad signal ( $\delta \approx 38.1$  ppm) at the aliphatic region of the  $^{13}\text{C}$  CP-MAS NMR spectra proved the formation of cyclobutane moieties.

Solid-state supramolecular photochemistry offers an efficient way to engineer new functional polymers by taking advantage of the preorganization of reactive moieties in the crystalline matrix.<sup>[56–58]</sup> In general, different strategies such as hydrogen bonding interaction,  $\pi \cdots \pi$  stacking, and metal–ligand interactions have been explored to control the topochemical arrangement of the reactive modules. Efficient  $\pi \cdots \pi$  stacking of 2D layers within 2D COFs offer a suitable

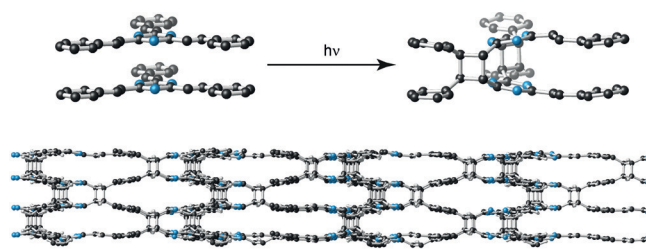


**Figure 3.** Effect of light irradiation on V-COF-1. a) Photographs of V-COF-1 dispersions, and b)  $^{13}\text{C}$  CP-MAS NMR spectra and c)  $\text{N}_2$  sorption isotherms of V-COF-1 before (V-COF-1) and after light irradiation (V-COF-1-Light). d) Fluorescence emission quenching upon irradiation of light with time, monitored by fluorescence emission spectroscopy.

platform for the preorganization of photoreactive moieties in a particular fashion to undergo photochemical reactions.<sup>[59]</sup> Topochemical [2+2] photocycloaddition in the solid state requires parallel orientation of two double bonds within a distance of 3.5–4.2 Å, entailed by Schmidt's criteria.<sup>[48]</sup> Thus, preorganization of the vinylene based templates are necessary for controlled synthesis of new 2D or 3D polymers. Schlüter et al. reported the synthesis of a 2D polymer by topochemical [2+2] photocycloaddition of a trivinylene monomer, whereas numerous vinylene building units inside metal organic frameworks (MOFs) or coordination polymers are also reported to undergo [2+2] cycloaddition.<sup>[60,61]</sup> Considering the high crystallinity and eclipsed stacking of 2D layers, V-COF-1 offers a suitable platform for [2+2] photo-cycloaddition within the vinylene linkages of adjacent layers. Still, this is the first report where parallel oriented vinylene moieties of adjacent layers undergo [2+2] cycloaddition within the 2D layers of a covalent organic framework.

The effect of cyclobutane formation from the vinylene linkages was also observed in the fluorescence emission spectrum of the V-COF-1 powders measured over time with light irradiation (Figure 3d). The pristine COF powders dispersed in ethanol was found to be highly fluorescent upon excitation at  $\lambda \approx 365$  nm. With light irradiation over time, a fast quenching of fluorescence emission together with the blue-shift of the emission maximum was observed and fluorescence emission was completely quenched after 10 minutes. This can be expected, as the degree of conjugation decreases upon formation of cyclobutane from conjugated vinylenes.<sup>[62]</sup>

Crystallinity of V-COF-1 was affected upon the [2+2] photo-cycloaddition, as  $\text{sp}^2$ -hybridized vinylene carbons convert into  $\text{sp}^3$ -cyclobutane species, causing severe strain within the crystalline lattice (Figure 4). To compensate the strain



**Figure 4.** Illustration of the [2+2] photocycloaddition between the layers and possible structural changes in the V-COF-1 lattice (side view). Note that the partial cycloaddition of vinylenes is represented, corroborating the  $^{13}\text{C}$  CP-MAS NMR spectrum of V-COF-1 after light irradiation.

within the 2D layers, V-COF-1 framework renders amorphous (Supporting Information, Figure S17). Such a change in crystallinity upon [2+2] photocycloaddition within crystalline lattices has been widely investigated in vinylene-based monomer single crystals, sometimes with a photosalt effect.<sup>[63]</sup> Notably, the framework remained porous even after exposure to UV/Vis irradiation for long time (48 h) with a  $\text{S}_{\text{A,BET}}$  of  $1093 \text{ m}^2 \text{ g}^{-1}$  (Figure 3c).

In summary, we report the successful crystallization of two new vinylene-linked 2D COFs (V-COF-1 and V-COF-2), achieved by base-catalyzed aldol condensation of TMT with ditopic terephthalaldehyde (TA) and tritopic 1,3,5-tris(p-formylphenyl)benzene (TFPB). This work demonstrates a successful strategy to exploit the highly electron-deficient s-triazine core for the crystallization of 2D vinylene-linked COFs at base-catalyzed conditions. Both COFs were found to be chemically stable under concentrated acidic as well as in basic conditions. [2+2] photocycloaddition within the columnar  $\pi \cdots \pi$  stacked 2D layers of a COF was observed for the first time upon UV/Vis light irradiation on V-COF-1 powders. Loss of crystallinity of V-COF-1 was observed after light irradiation owing to formation of highly strained cyclobutane rings within the 2D layers, as confirmed by a  $^{13}\text{C}$  CP-MAS NMR measurements. However, the framework remains porous with a  $\text{S}_{\text{A,BET}}$  of  $1093 \text{ m}^2 \text{ g}^{-1}$  even after exposure to UV/Vis irradiation for long time (48 h). Therefore, our focus is now dedicated towards the cycloreversion of the cyclobutanes, which could lead to a photo-switchable crystalline framework and could be implemented as a rewritable memory storage device. However, such a task is extremely difficult to achieve in the solid polycrystalline state, as the [2+2] cycloreversion from cyclobutane to parent vinylenes can follow different pathways. Thus, suitable conditions to photoswitch the frameworks reversibly are currently under investigation.

## Acknowledgements

This work is funded by the “Deutsche Forschungsgemeinschaft (DFG, German Research Foundation) within the project TH1463/12-1 and under Germany's Excellence Strategy—EXC 2008/1 (UniSysCat) -390540038”. A.A. thanks the BIG-NSE and Unicat for doctoral fellowship. We thank Christina Eichenauer and Maria Unterweger for their assis-

tance. Dr. Matthias Trunk, Nicolas Chaoui, and Dr. Johannes Schmidt are acknowledged for the fruitful discussions.

### Conflict of interest

The authors declare no conflict of interest.

**Keywords:** covalent organic frameworks · [2+2] cycloaddition ·  $\pi$ – $\pi$  stacking · porous polymers · vinylenes links

**How to cite:** *Angew. Chem. Int. Ed.* **2019**, *58*, 14865–14870  
*Angew. Chem.* **2019**, *131*, 15007–15012

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Manuscript received: May 12, 2019  
Revised manuscript received: June 25, 2019  
Accepted manuscript online: July 24, 2019  
Version of record online: September 9, 2019