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One-step flashlight processing of MOF thin films for non-linear light absorption†

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Although flashlight processing is an energy-efficient and fast technological approach for one-step patterning of polymers, metals, and dielectrics for diverse applications, it has not yet been implemented for hybrid materials such as metal–organic frameworks (MOFs). Herein, we report a one-step and large-scale patterning of different MOF thin films using a flash xenon lamp and a given mask in real time with $\text{cm}^2 \text{s}^{-1}$ speed. Structural and morphological analyses confirmed that the resulting patterns consisted of carbon and metal derivatives. We demonstrated that the resulting patterns acted as visible and near-infrared light absorbers with nonlinear absorption coefficients of 10^2 to 10^4 cm GW^{-1} , exceeding those of most of the metal films, initial MOFs, and metal–carbon hybrids. Thus, these results open up a universal one-step approach to pattern arbitrary MOFs for nonlinear optical applications in desired ways.

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Introduction

Metal–organic frameworks (MOFs), composed of inorganic and organic building blocks linked together *via* coordination bonds, possess tunable porosity and composition.^{1,2} These have made MOFs promising materials for gas separation/sorption, catalysis, and energy applications.^{3–6} Followed by the conventional synthetic approaches to fabricate MOFs⁷ for diverse applications, their patterning (by post-processing or structuring) endows them with new functionalities. For instance, the self-assembly of MOF particles,^{8,9} MOF patterning,¹⁰ shaping,¹¹ pyrolysis, and scaling from nanocrystal to thin-film (TF) forms¹² enable tuning the sorption and catalytic properties and extending the optical and electronic applications. Moreover, MOF patterning provides simple integration with optoelectronic devices.^{13,14} For optical and electronic applications, patterns greatly influence nonlinear optics and the corresponding planar/microscale devices as the hybrid structure of MOFs provides efficient modulation, nonlinear absorption, and high conversion of light.^{15–21}

Laser processing^{21–24} and lithography^{25–27} are considered as scalable and industry-oriented approaches to pattern MOFs.

They make it possible to pattern the MOFs (and other materials)^{28–30} to create planar components for optics, electronics, medicine, and sensors. However, despite their high level of scalability and pattern resolution (up to 50 nm for MOFs),²⁵ these approaches are still energy-consuming and require relatively long patterning times (with rare exceptions).^{28–30}

Herein, we report a one-step process to pattern MOF TFs through flashlight. Although flashlight approach has been proven as an energy-efficient and fast technological approach to pattern polymers, metals, and dielectrics,³¹ it has not yet been implemented for hybrid materials such as MOFs. Using a flash xenon lamp and a given mask, we developed arbitrary images on different MOF TFs in real time with $\text{cm}^2 \text{s}^{-1}$ speed in this study. Structural and morphological analyses confirmed that the resulting patterns consisted of carbon and metal derivatives. We also demonstrated that the resulting patterns acted as visible and near-infrared light absorbers: Z-scan technique revealed that patterns of varying thicknesses demonstrated the saturation absorption process with up to 10^4 cm GW^{-1} nonlinear coefficient, which exceeded the values of most of the metal films, initial MOFs, and metal–carbon hybrids. Thus, these results open up a universal one-step approach for fast and large-scale optical patterning of arbitrary MOFs for nonlinear optics in desired ways.

Results and discussion

As model MOFs, we selected HKUST-1,³² consisting of Cu ions, coordinated together with a 1,3,5-benzenetricarboxylate (BTC)

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ligand, its bimetallic forms (containing Cu–Ni, Cu–Co, and Cu–Fe ions),³³ and ZIF-8 (see ESI†). A relatively simple process of nucleation and growth of these MOFs³⁴ makes them suitable for TF deposition.³⁵ In this work, 220–280 °C decomposition temperature³⁶ of HKUST-1 should simplify the flashlight processing and reduce the energy consumption. As a result, the decomposition of HKUST-1 and ZIF-8 MOFs (depending on the environmental conditions)³⁶ generally provides carbon, metal, and metal oxide derivatives.^{21,37}

Initial TFs of HKUST-1 were deposited by a spin-coating method (see Experimental section and Video S1 (ESI†)); for its bimetallic forms and ZIF-8, see the ESI†). In detail, two automated microfluidic pumps were filled with a solution of Cu(NO₃)₂ and BTC in ethanol (1.5 mmol and 3 mmol per 100 mL of ethanol, respectively), releasing up to 700 drops of solution one by one onto Menzel glass, mounted on a rotating and heated holder (300 rpm, 100 °C, see Video S1, ESI†). This spin-coating setup have allowed us to fabricate MOF TFs recently with a surface roughness up to 200 nm.³⁸ The structure of obtained HKUST-1 TFs was confirmed by powder X-ray diffraction (PXRD, Fig. S1, ESI†), elemental analysis (Fig. S2 and S3, ESI†), and Raman analysis (Fig. 3b), while the thickness of the TFs (app. 30 μm) was determined by scanning electron microscopy (SEM). The structure of initial bimetallic HKUSTs and ZIF-8 TFs was also confirmed by PXRD, elemental, and Raman analyses (Fig. S9, S10, S12, and S14, ESI†).

Then, we performed flashlight processing of the TFs (see Experimental section and Video S2, ESI†). For this, each TF was mounted under a light source (pulsed xenon lamp, Fig. 1A) at a distance of 2 cm, while 0.5 mm-thick brass mask of the desired pattern was placed on the top of the TF. The obtained patterns (logo and variable width stripes, Fig. 1B–H) on HKUST-1 TF can be characterized by a metallic shine. It should also be noted that the spatial resolution (with up to 70 μm, Fig. S21, ESI†) and the quality of the flashlight patterning in Fig. 1A are limited by white light diffraction at the aperture,³⁹ heat transfer, and the geometry of the mask itself (smoothed boundaries of the patterns appeared, when the width of the aperture is less than the mask thickness). Concerning the contribution of heat transfer, the transition from isolators (such as glass) to heat-conducting surfaces can reduce the gradient of the pattern boundary (Fig. S21, ESI†), thereby additionally improving the spatial resolution.

The comparison of the optical properties of initial HKUST-1 TF and its pattern (Fig. 2) revealed that the flashlight process caused radical changes in both transmission (2-fold decrease of transmission for the pattern, Fig. 2A) and reflection spectra. Here, the latter is similar to the spectrum of light reflection from pure copper (Fig. 2B).

In a series of test experiments with HKUST-1 TF of a fixed thickness (app. 30 μm), we discovered the threshold values of the surface energy of flashlight, required to initiate the patterning. At the surface energy values less than 1.5 J cm⁻², no changes of HKUST-1 TF could be observed, regardless of the number of the flashlight pulses. However, we detected the characteristic changes (*i.e.*, the appearance of a metallic sheen)

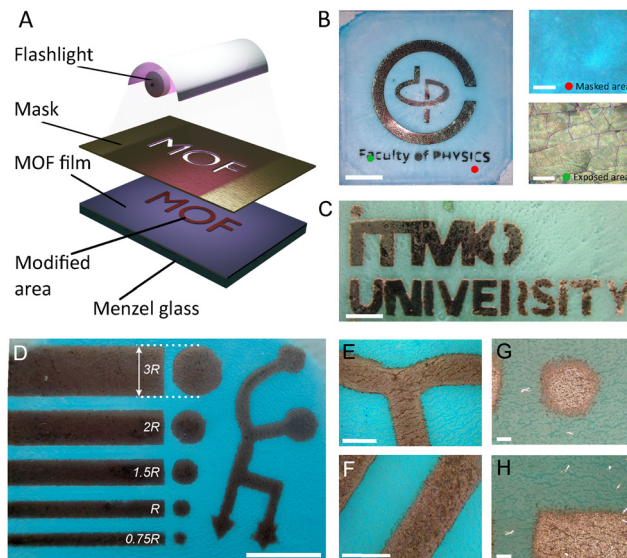


Fig. 1 (A) Scheme of flashlight processing of MOF TF (Video S2, ESI†). (B) Pattern on HKUST-1 TF after a series of 10 pulses with a surface energy density of 2.7 J cm⁻², and the optical images of the characteristic masked (upper image) and exposed (lower image) areas. Scale bars = 1 cm (left), 250 μm (right). (C) Optical image of the pattern obtained after a series of 50 pulses with a surface energy density of 1.8 J cm⁻². Scale bar = 500 μm. (D) Optical image of the arbitrary pattern with the fixed size of elements ($R = 1$ mm). Scale bar = 500 μm. (E)–(H) Magnified optical images of an arbitrary pattern from (D). Scale bars = 1 mm (E) and (F) and 200 μm (G) and (H).

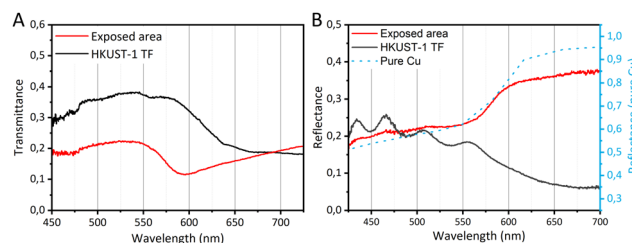


Fig. 2 (A) Transmittance and (B) reflectance spectra for the initial (black curve) and patterned (red curve) HKUST-1 TF.

when operating at 1.8 J cm⁻² even at a single flashlight pulse. An increase in the number of the flashlight pulses (irradiation dose) did not lead to additional changes in this case. However, the thickness inhomogeneities of the initial HKUST-1 TF resulted in different degrees of patterning by a single flashlight pulse (three distinguished areas in Fig. 3A). Using a confocal Raman spectrometer in the non-destructive mode (see the Experimental section), we analysed these areas (Fig. 3A): The first unmodified area of HKUST-1 TF did not absorb enough energy to initiate the visual changes (Fig. 3B, red curve correlating well with the Raman spectrum of initial HKUST-1).²¹ The second area demonstrated the metallic shine, probably associated with broken coordination bonds and partial diffusion/nucleation of Cu.⁴⁰ Indeed, Raman spectra revealed the decreased intensity of the coordination bond peak (270 cm⁻¹),²¹ and the appearance of halo in the region of 1200–1700 cm⁻¹



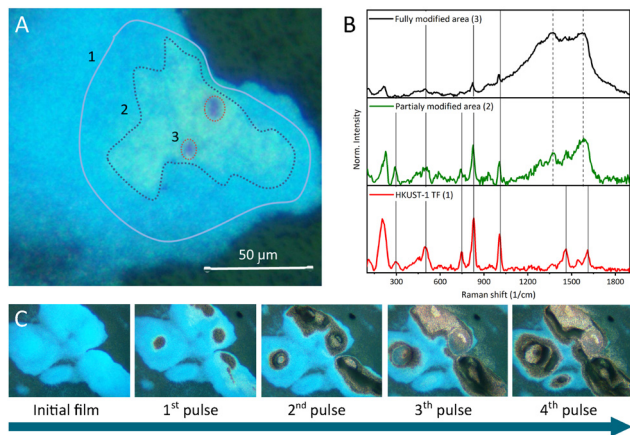


Fig. 3 (A) Characteristic stages of the patterning of HKUST-1 TF by single pulse and a set of pulses (C) at a fixed surface energy of 1.8 J cm^{-2} with the corresponding Raman spectra (B).

(Fig. 3B, green curve), which indirectly confirmed the amorphization (Fig. S5, ESI[†]) and carbonization⁴¹ of MOFs. The third area in Fig. 3A corresponds to a complete MOF decomposition, characterized by the vanished Raman vibration of organic ligands, coordination bond breaking,²¹ and appearance of carbon Raman peaks (Fig. 3B, black curve).⁴¹ Elemental analysis (Fig. S2–S7, ESI[†]) also confirmed the carbonization and appearance of Cu after flashlight patterning of HKUST-1 TF. Herein, the morphological analysis of the patterns (Fig. 4A) revealed the heterogeneity of the surface of the patterned TF with a well-defined macroscale granular structure (partially detached from the surface due to the

thermal stresses, Fig. 4B). The cross-section SEM micrographs determined the thickness of the patterned area varying from 0.1 to $2 \mu\text{m}$ (Fig. 4D and Fig. S4, ESI[†]), while the elemental analysis along the pattern (Fig. 4H–K) revealed a decrease in carbon (due to the thinning of the pattern itself and partial evaporation of carbon). Nevertheless, the patterned area still contained carbon (Fig. 3B) and agglomerated Cu-based particles (Fig. 4F–G and Fig. S6, S7, ESI[†]). Similar results were obtained for the patterned bimetallic HKUST-1 and ZIF-8 (Fig. S11 and S13, ESI[†]).

The transition from a single flashlight pulse mode to a set of pulses at a fixed surface energy of 1.8 J cm^{-2} is shown in Fig. 3C. It can be observed that the first pulse led to a local change in Fig. 3A, corresponding to the appearance of carbon (black area) and Cu (responsible for the metallic shine). Then, subsequent pulses only increase the probability of absorption of their energy and expand the surface of patterned area of the TF. This process repeats until the whole TF becomes completely patterned and the flashlight pulses does not cause any changes.

It is important that the number of pulses (30 pulses with 1.8 J cm^{-2} in our case), needed for the complete patterning of the TF, depends on the pulse energy (above the threshold) and the TF thickness, which determine the level of absorbed light. Thus, the flashlight patterning can be considered as adaptive and tunable through TF thickness and the number/energy of the flashlight pulses.

We should also mention that at a high level of pulse energy for patterning, the surface roughness and crystalline orientations of MOF TFs vanished its significance. Thus, the flashlight patterning can be applicable for diverse MOF TFs (with different thermal stabilities) obtained by spin coating (HKUST-1 and

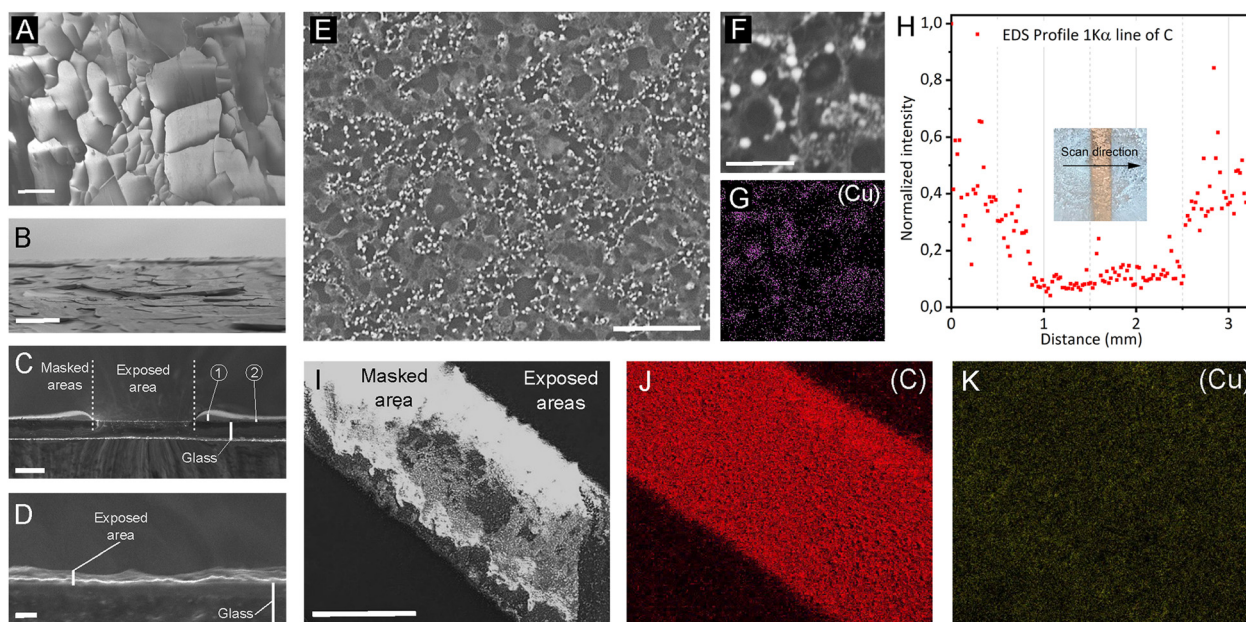


Fig. 4 SEM micrographs and elemental analysis of the patterned area of HKUST-1 TF: (A) top view, (B) angled view and (C) and (D) cross-section view. Scale bars = 200 μm, 100 μm, 200 μm and 2.5 μm, respectively. (E) Top-view SEM micrograph of the exposed area. Scale bar = 10 μm. (F) SEM micrograph and (G) the corresponding EDS map of Cu particle agglomerates. Scale bar = 2 μm. (H) EDS scan of carbon distribution across the exposed line. (I) SEM micrograph and the corresponding EDS map of (J) carbon and (K) copper distribution on the exposed and unexposed areas of HKUST-1 TF. Scale bar = 500 μm.



Table 1 Overview of the optical approaches of top-down MOF patterning. For non-optical approaches, see Table S3 (ESI)

Approach	Source	Resolution	Speed/ Exposure time	MOF	Product of patterning	Substrate	Ref.
Laser metallurgy	Pulsed fiber laser (1064 nm central wavelength, 20 kHz pulse repetition rate, 80–100 ns pulse duration)	55–380 μm	15 $\text{mm}^2 \text{s}^{-1}$	MOFs based on Fe, Co, Ni, Zn, Cd, In, Pb and Bi Ce-MOF	Metal nanoparticles covered with graphene shell CeO ₂ nanoclusters with Pt nanoparticles	Glass	23
			70 $\text{mm} \text{s}^{-1}$				42
Laser ablation	Pulsed Femto-, pico- and nano-second lasers			HKUST-1, ZIF-8, ZIF-67, UiO-66	Carbon, metal and metal oxide nanocrystals, MOF nanocrystals, amorphous MOF		21
Laser nanoforging	Pulsed fiber laser (1064 nm central wavelength, 1–10 Hz pulse repetition rate, 5 ns pulse duration)	150 nm		ZIF-8	ZIF-8 thin film	Metal foil, PET	43
Laser carbonization	Pulsed CO ₂ laser (10.6 μm central wavelength, 9–10 W integral power) CO ₂ laser (10.6 μm central wavelength, 2.4–7.8 W integral power)	150 μm		ZIF-67 (Co)	Porous graphene	LIMDG polyamide	44
			80–270 $\text{mm} \text{s}^{-1}$	Fe (MIL), Ni-BDC, HKUST-1	Metal nanoparticles with an amorphous carbon	Metal foil, carbon paper, carbon cloth	45
Laser interference patterning	CaN laser diode (450 nm central wavelength, 5.5 W integral power) Continuous diode laser (532 nm central wavelength, 40 mW integral power)	50 μm 5 μm		ZIF-67 (Co)	Co/N doped amorphous carbon		46
				HKUST-1	Copper oxide and amorphous carbon	Glass	47
Nonlinear lithography	Femtosecond laser (850–1300 nm wavelength, 150 fs pulse duration, 1 MHz pulse repetition rate) Femtosecond laser (780 nm central wavelength, 80 MHz pulse repetition rate, 100 fs pulse duration)	400 nm	150 fs, and 30–120 s	Er-BTC	Amorphous area inside a crystal volume	MOF crystal on Menzel glass	48
			700–800 nm	250 ms	MOF-5	Metallic patterns inside a crystal volume	
UV-lithography	Hg lamp		min	ZIF-8	Etched ZIF-8	Glass, Si, Au, Ag, Pt, Pd	50
Flashlight	Pulsed Xe lamp	70–120 μm	140 $\mu\text{s}/\text{cm}^2 \text{s}^{-1}$	HKUST-1, bimetallic HKUST-1 (Cu–Co, Cu–Ni, Cu–Co), ZIF-8	Metal oxide nanoparticle agglomerations with an amorphous carbon	Glass	This work

ZIF-8, Fig. S12, ESI[†]) and other methods (for example, dripping of bimetallic HKUST-1, Fig. S8, ESI[†]). Despite the increased thickness of bimetallic HKUST-1 TFs (up to 0.2 mm), the flashlight patterning (at 1.4 to 7 Hz repetition rate, 30 pulses) during 30 s was also successful (Fig. S11 and Fig. S20, ESI[†]).

In accordance with the capabilities of the flashlight approach,³¹ the pulse repetition rate can be increased up to 30 Hz, thereby reducing the patterning time. The latter is important for the transition of the Lab approach to industry: the laser processing and lithography of various materials^{25–30} and MOFs (Table 1 and Table S3, ESI[†])^{42–50} scale from $\text{cm}^2 \text{h}^{-1}$ to $\text{cm}^2 \text{s}^{-1}$ speed, while we reached the desired patterning speed up to $\text{cm}^2 \text{s}^{-1}$ (see Video S2, ESI[†]). The next feature of the Lab approach is spatial resolution (70 μm in our case, Fig. S21, ESI[†]): In contrast to laser processing and lithography of MOFs,^{21–26} demonstrating up to 50 nm resolution (Table 1 and Table S3, ESI[†]), the flashlight is limited by the white light

diffraction at the aperture³⁹ and heat transfer process, and can potentially reach several hundred nanometers.⁴⁸

Concerning the application,^{21–23} the large-scale MOF derivatives and patterns possess diverse functionalities from catalysis to energy and optics. This is due to the presence of metal open sites, residual porosity, and carbon nature. It is important that this set can additionally provide efficient absorption of light over a wide spectral range for nonlinear light absorbers^{17,18} and coatings. To prove this, we performed Z-scan measurements with an open aperture for the patterns on HKUST-1, bimetallic, and ZIF-8 TFs (see the Experimental section and ESI[†]).⁵¹ For this, Thorlabs AC lenses were used to build a telescope system at wavelengths of 1030 nm (254-50-AB-ML) and 515 nm (Mitutoyo NIR 10 \times NA 0.26), while Ophir PD10-C was used to detect the light in both cases. As the laser pulses, 515 nm and 1030 nm femtosecond light was utilized (Avesta Antaus fs laser, 6 Hz pulse repetition rate). The patterned TFs were then mounted on the linear



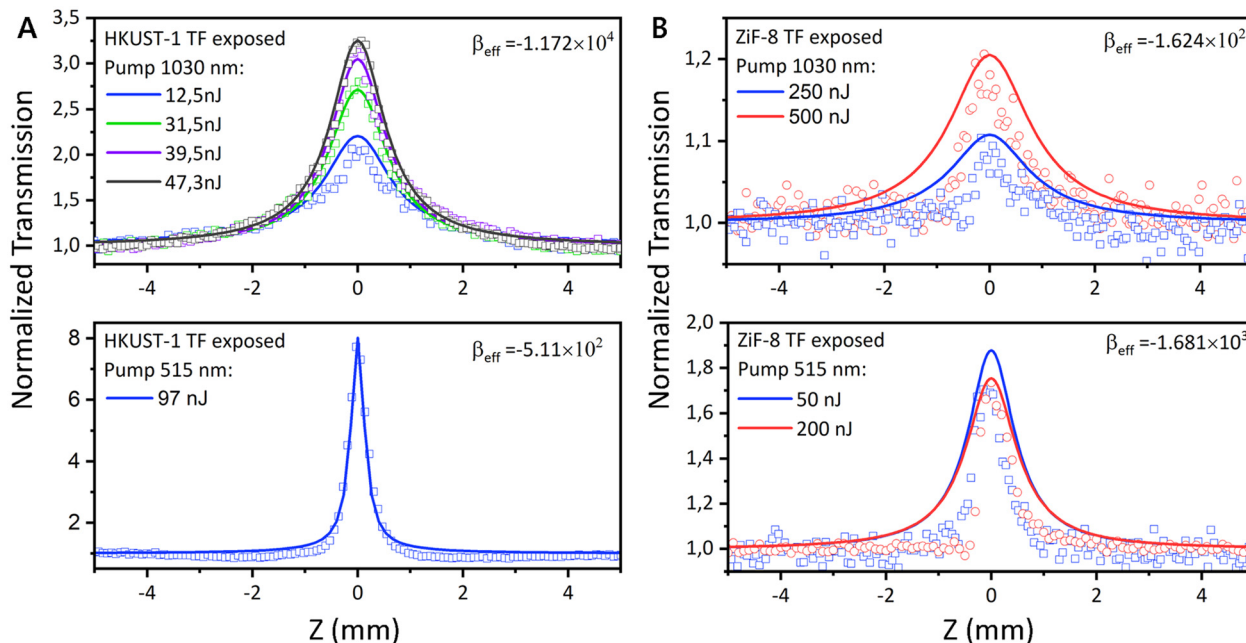


Fig. 5 Open-aperture Z-scan curves for the pattern on HKUST-1 TF from Fig. 1B at 1030 nm and 515 nm wavelengths (A), and the pattern on ZIF-8 TF (B) at 1030 nm and 515 nm wavelengths with varying laser pulse energies (12.5 to 500 nJ).

translation stage and moved between two confocally aligned objectives, while the incoming and transmitted energies of the laser pulses were detected.

Fig. 5 and Fig. S22 (ESI[†]) show the results of Z-scan. It can be observed that 50 to 100 nm-thick (Fig. S4, ESI[†]) patterns demonstrated a well-defined saturation absorption (SA) process.⁵² The curves shown in Fig. 5 were also used to estimate the nonlinear coefficient of light absorption β_{eff} (for the one-photon process, see details in the ESI[†]).^{51,52} We revealed that β_{eff} varied from 10^2 to 10^4 cm GW⁻¹ (Table S4, ESI[†]) depending on the wavelength and composition of the initial MOF. The complex morphology and hybrid nature of the patterned TF provided a nonlinear coefficient β_{eff} higher than that of common metals, carbon-based TFs (see Table S4, ESI[†]), and most initial MOFs (with some exceptions), which is intriguing.^{53–55}

Additionally, after Z-scan experiments, we noticed no damage of the patterned TF. Therefore, the observed nonlinear optical effect, in combination with high laser damage stability of the patterns (up to TW cm⁻²), outlines their potential application for large-scale and nanometer-thick semi-transparent patterns for passive modulation devices and photonics in general.^{56–58}

Conclusions

We report a one-step process to pattern MOF thin films through flashlight processing. Using a flash xenon lamp and a given mask, we developed arbitrary images on different MOF TFs in real time with cm² s⁻¹ speed and 70 μm spatial resolution. Structural and morphological analyses confirmed that the resulting patterns consist of carbon and metal derivatives. We also demonstrated that the resulting patterns act as visible

and near-infrared light absorbers: Z-scan revealed that the patterns demonstrated the saturation absorption process with a nonlinear absorption coefficient up to 10^4 cm GW⁻¹, which exceeds that of most metal films, initial MOFs, and metal-carbon hybrids. Thus, these results, open up a universal one-step approach to optically pattern MOFs (Table 1) for nonlinear optics in the desired ways.

Experimental section

HKUST-1 thin film synthesis

Thin films were synthesized on the self-made spin coating setup (see Video S1, ESI[†]) with two microfluidic pumps and a two-dimensional positioner system. The Menzel glass, used as a wafer, has been previously sonified in isopropanol. Two microfluidic pumps were filled with Cu(NO₃)₂ and BTC (1,3,5-benzenetricarboxylate) solutions in ethanol (1.5 mmol and 3 mmol per 100 mL of ethanol, respectively). A rotating sample holder (300 rpm) has been previously heated up to 100 °C. Drops were released from pumps one by one – two drops per each of 700 cycles, while drop counting was controlled by an optical pair feedback system. Each cycle takes approximately 10 s, which means that the deposition of one film takes two hours, without any control required from the setup operator. By the end of the synthesis procedure, films were dried in air on the sample holder heated up to 70 °C for 15 min.

Powder X-ray diffraction

The PXRD analysis of the film was performed using a Rigaku SmartLab 3 X-ray diffractometer (Tokyo, Japan) with a 2 kW characteristic CuK_α ($K_{\alpha 1}$ $\lambda = 1.54059$ Å, angular range



$2\theta = 5^\circ - 45^\circ$) X-ray radiation source and the Bragg–Brentano goniometer geometry. The measurements were performed with an angular resolution of 0.01° at a scanning speed of $0.7^\circ \text{ min}^{-1}$. The analysis confirmed the HKUST-1 structure (Fig. S1, ESI†) based on the merging of diffraction peaks with the etalon samples from the Cambridge Crystallographic Data Centre (CCDC numb. 112954).

Microscopic imaging

Microscopic images of HKUST-1 thin films were acquired using a Karl Zeiss AxioImager A2m in a bright-field regime.

Scanning electron microscopy and elemental analysis

SEM micrographs were obtained using a Carl Zeiss Neon 40 EsB CrossBeam system, being a combination of a field emission electron-optical column GEMINI and a field emission ion column with a gallium source, focused on a single point of the sample. The Neon 40 system uses three types of signal detectors: a secondary electron detector (In-lens SE) built into the column to study the surface topography with high resolution; a detector of reflected electrons with the possibility of energy filtering (EsB); and an Everhart-Thornley secondary electron detector (SE). For obtaining the images of the patterns, the system was operated in a high-resolution field emission SEM mode, in combination with elemental analysis. The sample was mounted on a translation stage inside the chamber (stage positioning: $X/Y = 152 \text{ mm}$, $Z = 43 \text{ mm}$, tilt $-15^\circ - +65^\circ$, rotation 360°).

Flashlight processing

For flashlight processing of MOF thin films, we designed the setup based on a pulsed Xe lamp (emission spectrum in Fig. S17, ESI†). The condenser battery charged for the estimate value ignited the discharge of the Xe lamp, thus creating the high-temperature plasma channel, irradiating the sample with the high-intensity full-spectrum pulsed light. In order to achieve the optimal distribution of the irradiance field, the reflector was used. The MOF thin film was placed at a distance of 2 cm from the light source, while the amount of energy put in the discharge ($0.51 \pm 0.04 \text{ J cm}^{-2}$ and more, see Table S2, ESI†) was regulated *via* the initial voltage ($1.00 \pm 0.05 \text{ kV}$) on the condenser battery. The time needed for cm^2 flashlight patterning varied from 1 s to 2 min, depending on the number of pulses (for more technical details, see the detailed description of the setup in the ESI† and Video S2).

Confocal transmission and reflection spectroscopy

Confocal transmission and reflection measurements were performed on a self-made confocal optical setup. A halogen lamp (Avantes Hal-s mini) was used as the light source. For transmission measurements, the incoming light was focused *via* the bottom objective (Mitutoyo M Plan Apo x50, NA = 0.55) and transmitted light was collected with the top one (Mitutoyo M Plan Apo x50, NA = 0.55). For the reflection, the top objective was used both for focusing the incoming radiation on the TF, and for collecting the reflected light. In both cases, the

collected signal was transferred to a Horiba Labram spectrometer with 150 g mm^{-1} diffraction grating and a water-cooling ANDOR CCD detector.

Confocal Raman spectroscopy

Raman scattering measurements were implemented under excitation by a 632.8 nm He-Ne continuous laser radiation in the reflection mode through $100\times/0.9 \text{ NA}$ objective. To prevent the TF from unintended overheating, the integral power of He-Ne was limited with a natural density filter by 10% (from 25 mW to 2.5 mW). The collected signal was transferred to a Horiba Labram Spectrometer with 1800 g mm^{-1} diffraction gratings and a water-cooling ANDOR CCD detector.

Z-scan measurements

The optical setup for Z-scan analysis is based on the fs pulsed ANTAUS fiber laser (Avesta projects) with the main harmonic wavelength of 1030 nm and a pulse duration of 270 fs. The main laser is also occupied with the second and third harmonic generators, which provides the conversion of the main wavelength into 515 nm. The pulse repetition rate was adjusted using the laser operational software, while the energy per pulse was regulated with the polarization attenuator (Glan-Taylor prism with rotating half wavelength plate). A telescope system was mounted with two long-working distance objectives, $10\times/0.26\text{NA}$ Mitutoyo Plan Apo NIR. The pulse energy measurements of the incoming and the transmitted pulses were conducted using two Ophir Photonics PD10C detectors. The vertical translation of the sample (along the Z axis) was provided using a Standa Motorized Stage 8MT177-100.

Author contributions

V. A. M. and N. A. Z. conceptualized the work. L. V., R. C., and M. M. performed the flashlight processing. Y. K. performed optical spectroscopy. S. A. P. performed z-scan experiments. S. S. R. and A. V. L. performed MOF thin film synthesis. S. A. S. performed bimetallic HKUST-1 synthesis. M. V. T. performed structural analysis. All the authors participated in writing the text.

Data availability

The data supporting this article have been included as part of the ESI.†

Conflicts of interest

There are no conflicts to declare.

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