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Fabrication of Robust Spatially-Resolved Photochromic Patterns on Cellulose Papers by Covalent Printing for Anti-Counterfeiting Applications

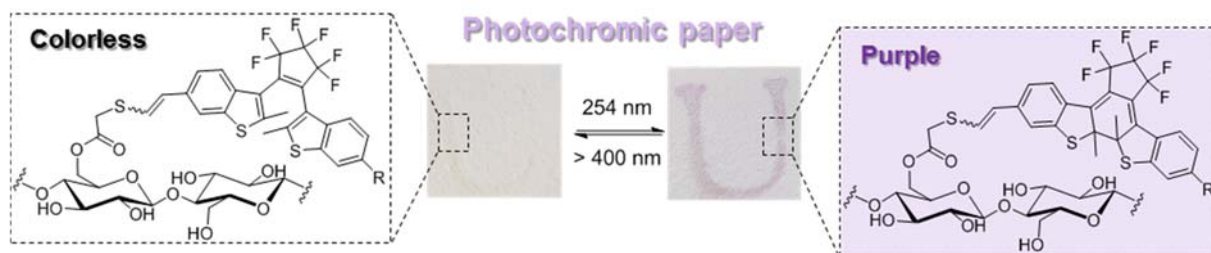
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■ ABSTRACT GRAPHIC



■ ABSTRACT

Despite its millennial age, cellulose paper remains the preferred material for domestic and professional printings, covering applications from simple office paperwork to fiducial solutions such as bills, passports, and head letters. The creation of robust photochromic patterns on cellulose papers for anti-counterfeiting applications is an important and still partially unaddressed challenge. In this contribution, we report the covalent printing of dibenzothienylethenes as photochromic compounds through a spatially-controlled light mediated thiol-X ligations. Photophysical and theoretical studies provide evidences for a reversible photochromism behavior, not affected significantly by the polar environment of the cellulose matrix, and demonstrates a high fatigue resistance over 18 successive write-erase cycles. The strong coloration-discoloration switch can be easily followed by a direct naked-eye readout.

■ KEYWORDS

Cellulose paper; Photochromic pattern; Thiol-yne reaction; Covalent printing; Dibenzothienylethene, Photoswitch

■ INTRODUCTION

While electronic devices flood our everyday life, paper prints paradoxically remain a modern media of communication and storage in both domestic and professional usages. Indeed, despite its millennial age, cellulose paper is more than ever a modern material as its low cost, disposability, stability, recyclability, and biocompatibility, are all major benefits in the development of sustainable societies.

The fabrication of functional papers, expanding the traditional usages of cellulose paper, is a particularly active field of research with promising applications in area as diverse as medicine,¹⁻⁴ catalysis,⁵⁻⁹ analytical,¹⁰⁻¹³ and environmental sciences.¹⁴⁻¹⁶ In this framework, the incorporation of photochromic dyes within the cellulose matrix allows the fabrication of photochromic papers as potential anti-counterfeiting devices or rewritable materials.¹⁷ Creating photochromic patterns on cellulose papers for anti-counterfeiting applications is an ambitious objective needing important requirements: (i) a spatial control of the cellulose functionalization; (ii) a reversible photochromism, not affected by the cellulose matrix; (iii) a robust patterning, insensitive to washing processes; and (iv) a good contrast for a naked-eye readout.

The vast majority of published works reports the preparation of photochromic papers using merocyanine dyes as photoactive compounds without any spatial control of the cellulose functionalization, excluding any molecularly imprinted patterns.¹⁸⁻²⁵ Moreover, when covalently anchored to the material surface, the photochromism of merocyanine dyes is frequently affected by the polar environment of the cellulose matrix as the colored open form is strongly stabilized within the hydrogen bond network of cellulose.¹⁹⁻²⁰ The irreversibility of the photochromism can be prevented through the impregnation of merocyanine dyes with a polymer carrier within the cellulose matrix,^{18, 22, 26} but the creation of photochromic patterns through a spatial control of the functionalization is still not demonstrated through these studies.

Diarylethenes are well-known photochromic compounds, derived from stilbene, which exist under a poorly conjugated open-form, absorbing in the UV region, and a strongly conjugated closed-form presenting a red-shifted absorption, typically in the 500-600 nm

domain.²⁷⁻²⁸ Upon UV-light irradiation the open-ring isomer undergoes a 6π -electrocyclization to furnish the closed-ring isomer which can be back converted upon exposure to visible light. Most diarylethenes are thermally irreversible (P-type) compounds, although, thermally unstable (T-type) diarylethenes have been reported as well.²⁸ Koga et al. prepared a remarkable photochromic paper by impregnation of cellulose fibers with a solution of diarylethenes in ionic liquids.²⁹ The absence of charges in both the open- and closed-ring isomers prevents the polar environment of cellulose to excessively stabilize one form with respect to the other and induce an unwanted irreversible photochromism behavior. Again, such strategy excludes any creation of photochromic patterns as paper fibers are all impregnated with the photochromic solution and physisorbed dyes could suffer from a limited stability as they are susceptible to be washed off from the material upon contact with liquids.³⁰

Therefore, robust photochromic papers remain to be developed through an original strategy allowing a spatial control of the cellulose functionalization. In this work, we report the first strategy allowing the creation of robust photochromic patterns on cellulose paper through our concept of covalent printing,³¹⁻³² which employs spatially-controlled thiol-X ligations. Our strategy exploits the photophysical properties of dibenzothienylethenes (DBE)³³ and features: (i) an efficient and fast coloration-discoloration process upon appropriate light irradiation; (ii) a thermal stability of the DBE isomers as shown by both theory and experiment; (iii) a good color contrast; and (iv) a high fatigue resistance, allowing many cycles to be performed without significant loss of performances.

■ EXPERIMENTAL SECTION

General remarks. All commercial solvents and reagents were used as received from Sigma-Aldrich, Fischer Scientific Ltd or Alfa Aesar. Whatman grade 6 filter paper was used as the cellulose source. ^1H and ^{13}C NMR spectra were recorded at 300 or 400 MHz and 75 or 100 MHz, respectively. Proton chemical shifts were internally referenced to the residual proton resonance in CDCl_3 (7.26 ppm). Carbon chemical shifts were internally referenced to the deuterated solvent signals in CDCl_3 (77.16 ppm). Melting points were recorded on a Stuart Scientific 7SMP3 apparatus. FT-IR spectra were recorded on a Bruker Tensor 27 spectrometer by the ATR technique. HRMS spectra were recorded on a Xevo G2-XS Qtof. UV-visible absorption spectra were recorded using a Shimadzu UV-2501 PC

spectrophotometer for all compounds before the grafting onto cellulose and a Perkin Elmer lambda 1050 was used for the study of papers.

General procedure for the pre-treatment of cellulose filters. A piece of paper (ca. 140 mg) was immersed in an aqueous solution of NaOH (40 mL, 10 wt%) and the mixture was shaken for 24 h on an orbital agitator. The paper was washed 6 times with EtOH and stored in EtOH.

Synthesis of dithiodiglycolic acid. A round-bottom flask was charged with 1% H₂O₂ aqueous solution (120 mL), tetrabutylammonium iodide (200 mg, 0.54 mmol) and thioglycolic acid (5 g, 54.3 mmol). The resulting mixture was stirred for 1 h. The crude product was extracted with ethyl acetate (3 × 50 mL). The combined organic layers were washed with water (5 × 40 mL), dried with MgSO₄ and concentrated under vacuum to give the corresponding product as a red powder (9.88 g, 99%) which was used without further purification.

Grafting of dithiodiglycolic acid onto cellulose paper. A pre-treated cellulose paper was washed 3 times with anhydrous toluene and placed in a flame-dried round-bottom flask under argon with dry toluene (20 mL). A solution of BF₃·Et₂O in toluene (1% in 1.1 mL) and dithiodiglycolic acid (471.9 mg, 2.59 mmol) was added to the reaction mixture. The resulting solution was stirred for 24 h at 110 °C on an orbital agitator, protected from light and under argon. Then, the paper was washed successively with EtOH, MeOH, acetone and CH₂Cl₂, dried under vacuum and stored under argon.

2-Methylbenzothiophene (1). A four necked flask was charged with benzothiophene (5 g, 37.26 mmol) in dry THF (40 mL) and cooled to −78 °C. *n*-BuLi (25.6 mL, 40.99 mmol, 1.6 M in hexane) was added slowly and the mixture was stirred for 1 h. Then, methyl iodine (2.6 mL, 40.99 mmol) was added and the solution was stirred for 40 minutes at −78 °C. The resulting mixture was quenched with cold water (100 mL), extracted with ether (2 × 50 mL), washed with brine (100 mL) and concentrated to give **1** as a white powder (5.46 g, 99%). mp 50–51 °C [Lit.³⁴ 51–52 °C]. IR (ATR) ν 1430, 1196, 1134, 1016, 865, 828, 743, 721, 562, 488 cm^{−1}. ¹H NMR (CDCl₃, 400 MHz) δ 7.76 (d, 1H, *J* = 7.9 Hz), 7.66 (d, 1H, *J* = 7.4 Hz), 7.34–7.23 (m, 2H), 6.98 (s, 1H), 2.60 (s, 3H). ¹³C NMR (CDCl₃, 100 MHz) δ 141.0, 140.6, 139.9, 124.2, 123.5, 122.7, 122.1, 121.7, 16.3. HRMS (ASAP⁺) *m/z* [M + H]⁺ Calcd for C₉H₉S 149.0425; Found 149.0427.

3-Bromo-2-methylbenzothiophene. In a round-bottom flask, a solution of 2-methylbenzothiophene (5.45 g, 36.82 mmol) in AcOH/CHCl₃ (1/1, 100 mL) was stirred with

N-bromosuccinimide (7.21 g, 40.49 mmol). The reaction mixture was stirred for 17 h. Then, a part of chloroform and acetic acid was evaporated (*ca.* 80 mL) and a saturated aqueous Na₂S₂O₃ solution (60 mL) was added and stirred for 5 min. The resulting mixture was extracted with CH₂Cl₂ and washed with a saturated aqueous NaHCO₃ solution and H₂O. The organic layer was dried over MgSO₄ and concentrated under vacuum. Purification by flash chromatography (pentane) afforded the titled compound as a white solid (7.83 g, 95%). mp 42–43 °C [Lit.³⁵ 43–45 °C]. IR (ATR) ν 1432, 1300, 1252, 916, 748, 719 cm⁻¹. ¹H NMR (CDCl₃, 300 MHz) δ 7.75–7.70 (m, 2H), 7.44–7.31 (m, 2H), 2.57 (s, 3H). ¹³C NMR (CDCl₃, 75 MHz) δ 138.6, 137.3, 135.4, 125.0, 124.9, 122.7, 122.3, 106.7, 15.65. HRMS (ASAP⁺) *m/z* [M]⁺ Calcd for C₉H₇SBr 225.9452; Found 225.9455.

3,3'-(Perfluorocyclopent-1-ene-1,2-diyl)bis(2-methylbenzothiophene) (4). To a solution of 3-bromo-2-methylbenzothiophene (6.23 g, 27.7 mmol) in dry THF (60 mL) under N₂ atmosphere, *n*-BuLi (19.04 mL, 30.5 mmol, 1.6 M in hexane) was added dropwise at -78 °C. The mixture was stirred for 30 minutes and octafluorocyclopentene (1.66 mL, 12.47 mmol) was added dropwise to the mixture with a cooled syringe. The resulting mixture was stirred for 2 h at -78 °C, warmed to room temperature and quenched with cold water (100 mL). The organic phase was extracted with ether (3 × 50 mL) and the organic layers were collected, washed with brine (2 × 50 mL), dried over MgSO₄, filtered and evaporated under vacuum. The resulting mixture was purified by flash column chromatography (100% hexane) to give **4** as a yellow solid (3.62 g, 62%) as mixture of parallel (p) and antiparallel (ap) isomers in a ratio of 36:64. mp 153–154 °C [Lit.³⁶ 154–156 °C]. IR (ATR) ν 1437, 1337, 1274, 1102, 962, 755, 735, 574, 565 cm⁻¹. ¹H NMR (300 MHz, CDCl₃) δ 7.72–7.58 (m, 4H), 7.42–7.16 (m, 4H), 2.52 (s, 6H × 0.36, p), 2.25 (s, 6H × 0.64, ap). ¹³C NMR (CDCl₃, 75 MHz) δ 142.8, 142.3, 138.5, 138.3, 124.8, 124.6, 124.5, 122.2, 122.3, 122.0, 119.3, 15.29. HRMS (ASAP⁺) *m/z* [M + H]⁺ Calcd for C₂₃H₁₅F₆S₂ 469.0519; Found 469.0514.

Dibenzothienylethenes (5) and (6). A solution of 3,3'-(perfluorocyclopent-1-ene-1,2-diyl)bis(2-methylbenzothiophene) **4** (2.65 g, 5.66 mmol) in acetic acid (200 mL) was treated with water (9.4 mL), sulfuric acid (4 mL), iodine (1.72 g, 6.79 mmol) and orthoperiodic acid (554 mg, 2.43 mmol) at room temperature. The mixture was stirred for 3 h at 70 °C and the solution was poured into water (500 mL). The organic phase was extracted with AcOEt (2 × 100 mL). The organic extracts were washed with water (100 mL), saturated aqueous NaHCO₃ solution (100 mL) and saturated aqueous Na₂S₂O₃ solution (100 mL). The resulting solution was dried over MgSO₄, filtered and concentrated. The residue was purified flash

chromatography (100% hexane) to afford a white solid (2.77 g, 72%) as a mixture of **5/6** in a ratio of 33/66 which was directly used in the next step.

Dibenzothienylethenes (7) and (8). Under nitrogen atmosphere, a solution of a mixture of **5/6** in a ratio of 33/67 (760 mg, 0.374 mmol **5** and 0.747 mmol **6**), Pd(PPh₃)₂Cl₂ (137 mg, 0.195 mmol), CuI (411 mg, 2.16 mmol) and trimethylsilylacetylene (1.42 g, 2.16 mmol) in Et₃N (11 mL) was heated at 80 °C for 15 h. The crude mixture was hydrolyzed with water (10 mL). Et₃N was removed under vacuum and the solution was extracted with CH₂Cl₂ (3 × 50 mL). The organic extracts were washed with brine (50 mL), dried with MgSO₄ and concentrated under reduced pressure. The residue was purified with short cake of celite followed by a flash chromatography on silica gel (100% hexane) to afford **7** (200 mg, 95%, based on **5**) and **8** (449 mg, 91%, based on **6**) as yellow solids.

Dibenzothienylethene 7 was isolated as a mixture of parallel (p) and antiparallel (ap) isomers in a ratio of 38/62. mp 121–122 °C. IR (ATR) ν 1272, 1250, 1139, 1110, 897, 841, 756 cm⁻¹. ¹H NMR (300 MHz, CDCl₃) δ 7.83–7.19 (m, 7H), 2.49 (s, 6H × 0.38, p), 2.23 (s, 3H × 0.62, ap), 2.22 (s, 3H × 0.62, ap), 0.30 (s, 9H × 0.62, ap), 0.27 (s, 9H × 0.38, p). ¹³C NMR (CDCl₃, 75 MHz) δ 144.6, 144.1, 142.8, 142.4, 138.4, 138.3, 138.2, 128.5, 128.3, 125.9, 125.7, 124.8, 124.6, 122.3, 122.0, 121.9, 119.5, 119.2, 104.9, 95.2, 94.9, 88.2, 86.1, 15.4, 15.2, 0.1. HRMS (ES⁻) m/z [M - H]⁻ Calcd for C₂₈H₂₁F₆S₂Si 563.0758; Found 563.0748.

Dibenzothienylethene 8 was isolated as a mixture of parallel (p) and antiparallel (ap) isomers in a ratio of 41/59. mp 198–199 °C ; IR (ATR) ν 1462, 1343, 1271, 1146, 1106, 840, 820 cm⁻¹. ¹H NMR (300 MHz, CDCl₃) δ 7.81 (s, 2H × 0.59, ap), 7.73 (s, 2H × 0.41, p), 7.53 (d, 2H × 0.59, J = 8.3 Hz, ap), 7.47–7.38 (m, 2H), 7.24 (s, 2H × 0.41, p), 2.46 (s, 6H × 0.41, p), 2.18 (s, 6H × 0.59, ap), 0.26 (s, 18H × 0.59, ap), 0.24 (m, 18H × 0.41, p). ¹³C NMR (CDCl₃, 75 MHz) δ 144.6, 144.1, 138.1, 128.4, 128.4, 125.9, 125.8, 121.9, 121.6, 119.5, 119.3, 104.8, 95.2, 15.4, 0.1. HRMS (ES⁺) m/z [M + Na]⁺ Calcd for C₃₃H₃₀F₆NaS₂Si₂ 683.1129; Found 683.1145.

Dibenzothienylethene (9(o)). A solution of **7** (100 mg, 0.177 mmol) in a mixture of CH₂Cl₂ (20 mL) and MeOH (10 mL) was treated with K₂CO₃ (600 mg, 4.35 mmol) under nitrogen. The resulting mixture was stirred for 1 h at room temperature and quenched with NH₄Cl (50 mL). The organic phase was extracted with CH₂Cl₂ (3 × 50 mL) and washed with water (2 × 20 mL). The resulting solution was dried over MgSO₄, filtered and concentrated to give **9(o)** as a yellow solid (83.6 mg, 96%) as a mixture of parallel (p) and antiparallel (ap) isomers in a ratio of 37/63. mp: 163–164 °C. IR (ATR) ν 3302, 2108, 1345, 1273, 1101, 965, 725, 646 cm⁻¹. ¹H NMR (300 MHz, CDCl₃) δ 7.88–7.12 (m, 7H), 3.12 (s, 1H × 0.63, ap), 3.06

(s, 1H \times 0.37, p), 2.49 (s, 6H \times 0.37, p), 2.23 (s, 3H \times 0.63, ap), 2.21 (s, 3H \times 0.63, ap). ^{13}C NMR (CDCl_3 , 75 MHz) δ 144.8, 144.2, 142.9, 142.4, 138.5, 138.4, 138.2, 138.1, 128.6, 128.4, 126.1, 125.9, 124.8, 124.7, 124.6, 122.3, 122.1, 122.1, 119.4, 119.2, 118.4, 83.5, 77.9, 15.4, 15.3. HRMS (ASAP $^+$) m/z $[\text{M}]^+$ Calcd for $\text{C}_{25}\text{H}_{14}\text{F}_6\text{S}_2$ 492.0441; Found 492.0435.

Dibenzothienylethene (10(o)). A solution of **8** (231 mg, 0.35 mmol) in a mixture of CH_2Cl_2 (20 mL) and MeOH (10 mL) was treated with K_2CO_3 (1.38 g, 10 mmol) under nitrogen. The resulting mixture was stirred for 1 h at room temperature and quenched with saturated aqueous NH_4Cl solution (50 mL). The organic phase was extracted with CH_2Cl_2 (3 \times 50 mL) and washed with water (2 \times 50 mL). The organic extracts were dried over MgSO_4 , filtered and concentrated under vacuum to give **10(o)** as a yellow solid (179 mg, 99%) as mixture of parallel (p) and antiparallel (ap) isomers in a ratio of 40/60. mp: 194–195 $^\circ\text{C}$. IR (ATR) ν 3302, 2111, 1461, 1336, 1269, 1106, 965, 876, 816 cm^{-1} . ^1H NMR (300 MHz, CDCl_3) δ 7.84 (s, 2H \times 0.6, ap), 7.75 (s, 2H \times 0.4, p), 7.57 (d, 2H \times 0.6, J = 8.3 Hz, ap), 7.48 (m, 2H), 7.30 (m, 2H \times 0.4, p), 3.12 (s, 2H \times 0.6, ap), 3.08 (s, 2H \times 0.4, p), 2.48 (s, 6H \times 0.4, p), 2.20 (s, 6H \times 0.6, ap). ^{13}C NMR (CDCl_3 , 75 MHz) δ 144.8, 138.4, 138.2, 128.6, 128.5, 126.2, 126.0, 122.0, 121.8, 119.3, 118.5, 83.4, 77.9, 15.4. HRMS (ASAP $^+$) m/z $[\text{M}]^+$ Calcd for $\text{C}_{27}\text{H}_{14}\text{F}_6\text{S}_2$ 516.0441; Found 516.0432.

Dibenzothienylethene (11). A solution of **8** (300 mg, 0.454 mmol) in CH_2Cl_2 (15 mL) under inert atmosphere was treated with *m*-CPBA (673 mg, 2.73 mmol, 70 % in water) at 0 $^\circ\text{C}$ and the resulting mixture was stirred for 48 h at 25 $^\circ\text{C}$. The organic phase was extracted with CH_2Cl_2 (3 \times 50 mL) and washed with water (2 \times 10 mL). The organic extracts were dried over MgSO_4 , filtered and concentrated under vacuum to give **11** as a yellow solid (329 mg, 89%) as a mixture of parallel (p) and antiparallel (ap) isomers in a ratio of 52/48. mp 151–153 $^\circ\text{C}$. IR (ATR) ν 2158, 1697, 1317, 1145, 842 cm^{-1} . ^1H NMR (300 MHz, CDCl_3) δ 7.81 (s, 1H), 7.77 (s, 1H), 7.63 (d, J = 7.7 Hz, 1H), 7.47 (d, J = 7.7 Hz, 1H), 7.04 (m, 2H), 2.18 (s, 3H), 2.03 (s, 3H), 0.25 (s, 18H); ^{13}C NMR (CDCl_3 , 75 MHz) δ 145.0, 144.3, 137.0, 136.8, 135.5, 128.6, 126.7, 126.5, 126.2, 123.6, 122.2, 122.1, 102.0, 100.9, 9.1, -0.22. HRMS (ES $^-$) m/z $[\text{M} - \text{H}]^-$ Calcd for $\text{C}_{33}\text{H}_{29}\text{O}_4\text{F}_6\text{S}_2\text{Si}_2$ 723.0950; Found 723.0969.

Covalent printing on cellulose paper with dibenzothienylethenes 9(o) - Cell-DBE-1. A piece of **Cell-Dis** (12 mg), wedged between a photomask and an aluminium plate (2 x 2.5 cm), was immersed in the solution of dibenzothienylethenes **9(o)** (135 mg, 0.27 mmol) and 2,2-dimethoxy-2-phenylacetophenone (DMPA, 5 mg, 0.02 mmol) in DMSO (6 mL). The solution, under vigorous stirring, was irradiated with a Xenon lamp (50 W) for 150 minutes.

The paper was washed successively with EtOH, MeOH, acetone and CH₂Cl₂, dried under vacuum and stored under argon.

Covalent printing on cellulose paper with dibenzothienylethenes **10(o) - Cell-DBE-2.**

A piece of **Cell-Dis** (12 mg), wedged between a photomask and an aluminium plate (2 x 2.5 cm), was immersed in the solution of dibenzothienylethenes **10(o)** (140 mg, 0.27 mmol) and 2,2-dimethoxy-2-phenylacetophenone (DMPA, 5 mg, 0.02 mmol) in DMSO (6 mL). The solution, under vigorous stirring, was irradiated with a Xenon lamp (50 W) for 150 minutes. The paper was washed successively with EtOH, MeOH, acetone and CH₂Cl₂, dried under vacuum and stored under argon.

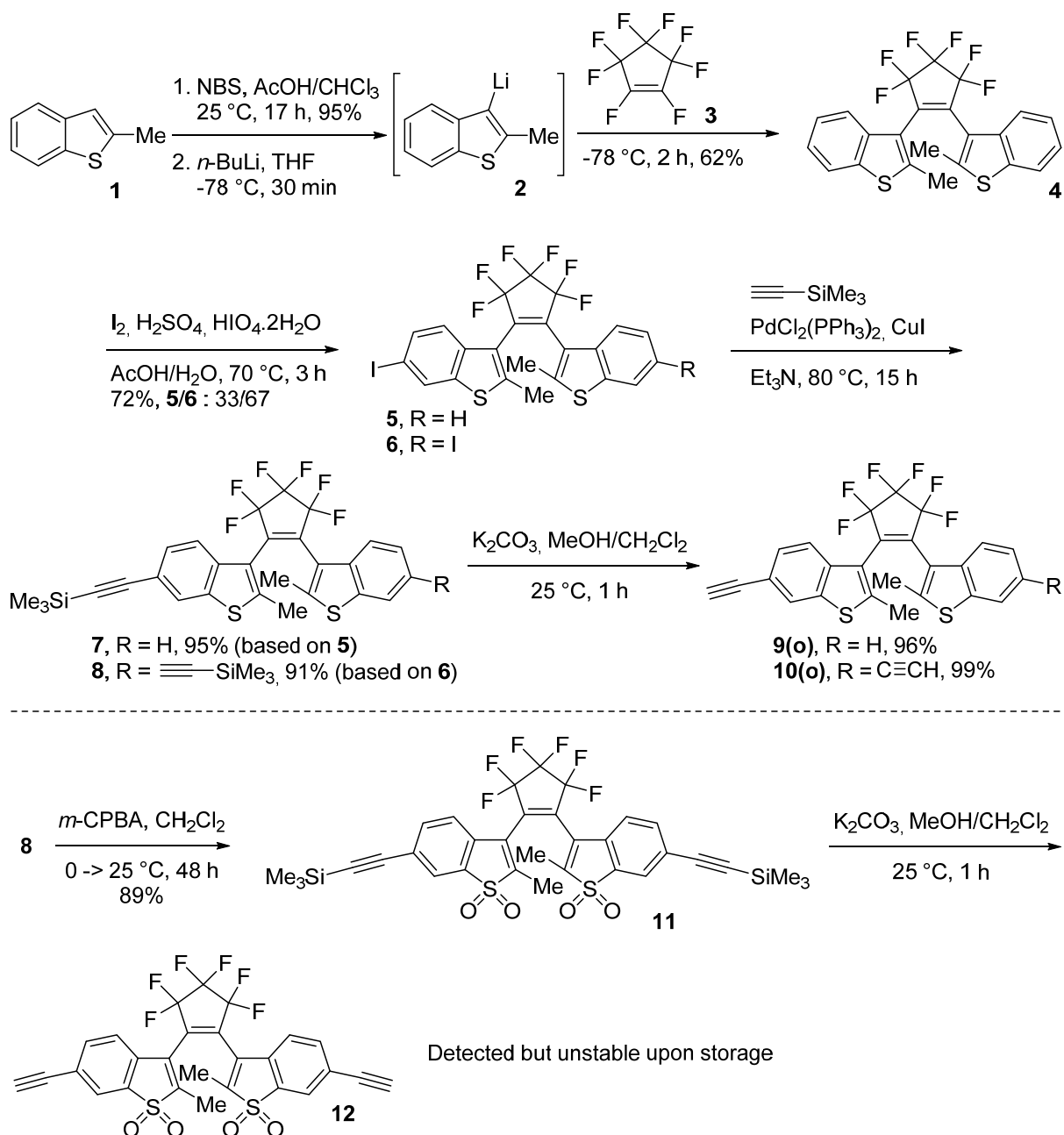
Computational details. All theoretical calculations have been performed with the Gaussian 16 program.³⁷ The ground-state geometries of the different dyes have been optimized at the PBE0/6-31G(d) level,³⁸ taking into account dispersion effects through the use of the D3-BJ approach.³⁹ These geometry optimization were made using a *tight* convergence threshold and the so-called *ultrafine* DFT integration grid. Next, the minimum nature of all optimized geometries was confirmed by a frequency calculation that returned no imaginary frequency. The excited-state were explored using TD-DFT with the same PBE0 functional but a diffuse containing atomic basis set, namely 6-31+G(d). The thermal transition state between the open and closed forms was determined at the PBE0/6-31G(d) level using the broken-symmetry (BS-DFT) approach and it was verified that the final vibrational mode indeed corresponds to the cyclization process. During all calculations, solvent effects (CH₂Cl₂) were accounted for using the Polarizable Continuum Model (PCM),⁴⁰ in its linear-response *non-equilibrium* form for the TD-DFT part of the calculation.

■ RESULTS AND DISCUSSION

Synthesis of dibenzothienylethenes. We recently reported the concept of covalent printing onto cellulose paper.³¹ The covalent printing strategy uses photoresponsive papers functionalized with dithiodiglycolic ester which upon light irradiation allows thiol-X ligations of chromophoric inks with a powerful spatio-temporal control. The application of the covalent printing strategy to the fabrication of photochromic papers required the synthesis of dibenzothienylethenes bearing reactive alkyne groups for thiol-yne ligations onto the surface of cellulose paper.

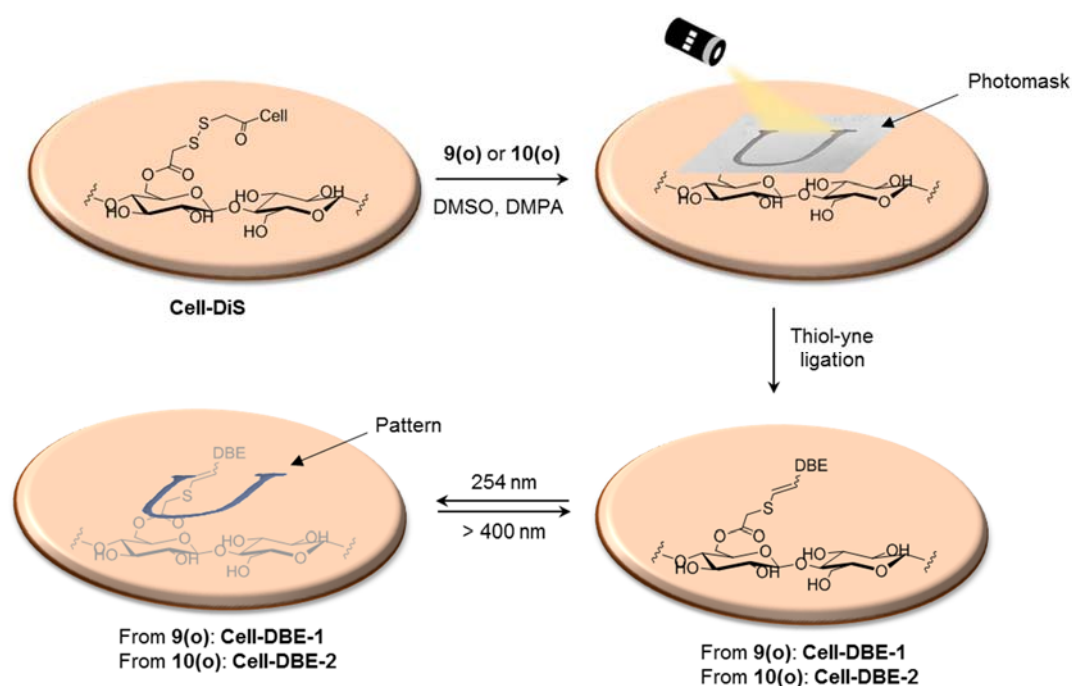
The dibenzothienylethene backbone **4** was prepared in two steps from commercially available 2-methylbenzothiophene **1** by adaptation of literature procedures (Scheme 1).⁴¹

Bromination of 2-methylbenzothiophene **1** with NBS followed by lithium-halogen exchange with *n*-Buli furnished the corresponding lithiated benzothiophene derivative **2** which reacted with octafluorocyclopentene **3** at $-78\text{ }^{\circ}\text{C}$ to give 1,2-bis-(2-methylbenzo[*b*]thiophen-3-yl)perfluorocyclopentene **4** in 62% yield over two steps. Iodination of the benzothiophene units furnished an inseparable mixture of mono- and diiodinated dibenzothienylethenes **5/6**, in a ratio of 1/2, respectively, and 72% yield. The subsequent introduction of alkyne functional groups through a Sonogashira reaction with trimethylsilylacetylene was carried out on the iodinated mixture **5/6**, furnishing the corresponding dibenzothienylethenes **7** and **8** which were separated by flash chromatography. The cleavage of the trimethylsilyl protecting group on **7** and **8** with K_2CO_3 in $\text{CH}_2\text{Cl}_2/\text{MeOH}$ proceeded smoothly to give the expected terminal alkynes **9(o)** and **10(o)** which were immediately engaged for the covalent printing of cellulose paper due to their rapid degradation upon storage. We also envisaged the preparation of the sulfonyl derivative **12** since it has been reported that the oxidation of benzothienyl moieties significantly increase the fluorescence quantum yield of the closed form, as compared to the corresponding unoxidized dibenzothienylethenes.⁴² The oxidized photochromic compound **12** was tentatively prepared in two steps by sulfur oxidation of **8** with *m*-CPBA, furnishing **11** as intermediate, followed by trimethylsilyl cleavage. Unfortunately, the oxidized derivative **12** proved to be unstable and could not be obtained in pure form for further covalent printing on cellulose paper.



SCHEME 1. Preparation of dibenzothienylethenes bearing alkyne functional groups

The covalent printing of **9(o)** and **10(o)** on photoactive cellulose paper (**Cell-Dis**), through a thiol-yne ligation, was carried out following an adaptation of our recently developed protocol.³¹ **Cell-Dis**, prepared by grafting of dithiodiglycolic acid on mercerized cellulose paper, was inserted in a sample holder consisting of two aluminum plates; the top plate being machined with the letter “U”. Upon immersion in a solution of **9(o)** or **10(o)** in DMSO, also containing 2,2-dimethoxy-2-phenylacetophenone (DMPA) as radical initiator, **Cell-Dis** was irradiated with a Xenon lamp (60 W) to print the letter “U” (Scheme 2).



SCHEME 2. General strategy for the covalent printing of photochromic compounds **9(o)** and **10(o)**.

Material characterization. The covalent printing of **9(o)** and **10(o)** onto **Cell-DiS** was followed by Fourier-transform infrared spectroscopy (FTIR) and X-ray photoelectron spectroscopy (XPS) analyses (Figure 1). The appearance of a new band at *ca.* 1645 cm^{-1} on the FTIR spectra of **Cell-DBE-1** and **Cell-DBE-2** characteristic of C=C stretching frequencies of the dibenzothienylethene backbone, successfully confirmed the grafting. The band at *ca.* 1725 cm^{-1} accounts for the C=O stretching vibration of the ester groups confirming that the covalent printing conditions did not affect the thiomethylglycolate function. The remaining bands between 400 to 1500 cm^{-1} are characteristic of the finger print of the cellulose backbone while the bands in the region of 3100-3400 cm^{-1} are attributed to the stretching of O-H bonds. The XPS spectra of **Cell-DBE-1** and **Cell-DBE-2** also attested the successful grafting of **9(o)** and **10(o)**, respectively, through the presence of F1s peaks at *ca.* 688 eV.

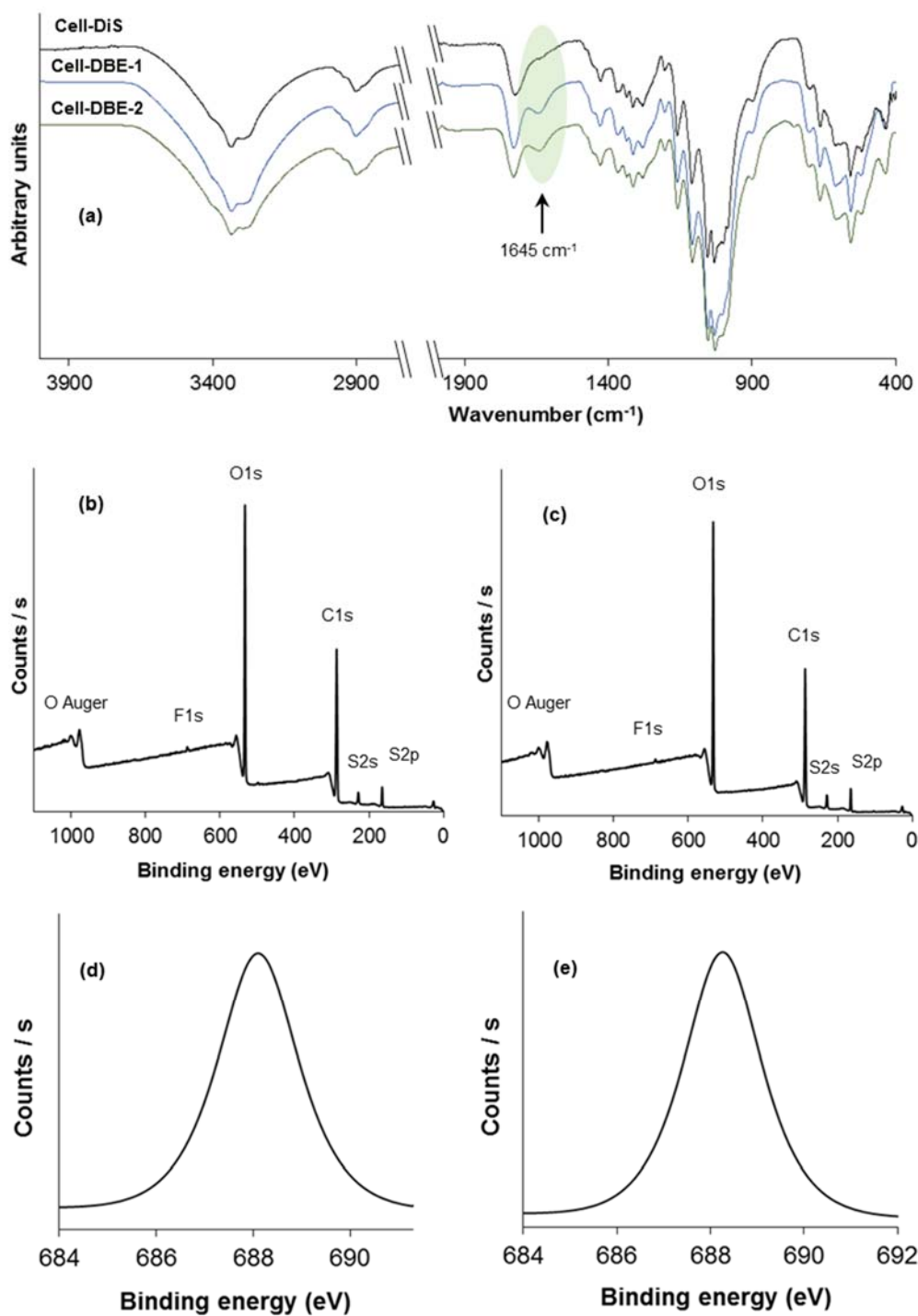


FIGURE 1. (a) FTIR spectra of **Cell-DiS**, **Cell-DBE-1** and **Cell-DBE-2**. (b) Survey scan spectra of **Cell-DBE-1**. (c) Survey scan spectra of **Cell-DBE-2**. (d) High-resolution F1s spectrum of **Cell-DBE-1**. (e) High-resolution F1s spectrum of **Cell-DBE-2**.

The thermal stability of **Cell-DBE-1** and **Cell-DBE-2** were assessed using thermogravimetric analysis (TGA) experiments under N₂ atmosphere (Figure 2) and

compared to pristine cellulose paper. The mass loss of ca. 4% below 200 °C for the three samples is attributed to the desorption of volatiles such as residual solvents and moisture. **Cell-DBE-1** and **Cell-DBE-2** display very similar stability profiles and are stable until 250 °C; Cell showing a slightly increased stability, up to 300. The sudden mass loss at 260 °C for **Cell-DBE-1** and **Cell-DBE-2** can be explained by the elimination of the grafted photochromic moieties. This result suggests that **Cell-DBE-1** and **Cell-DBE-2** are stable in a large range of temperatures from 25 to 250 °C.

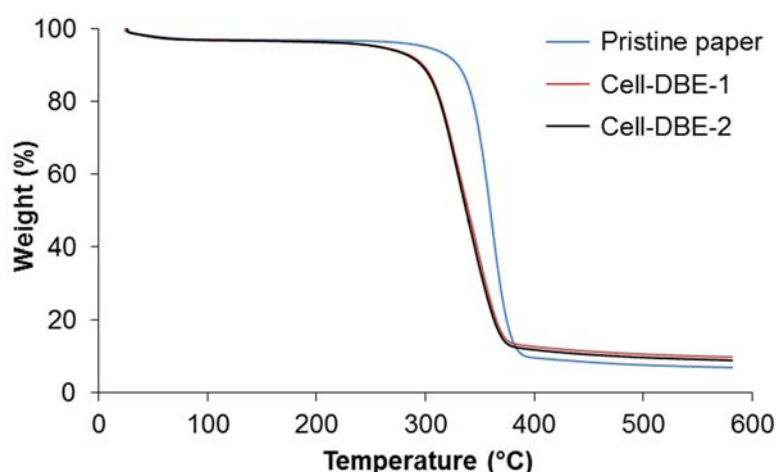


Figure 2. Thermogravimetric analysis of **Cell**, **Cell-DBE-1** and **Cell-DBE-2**.

The atomic composition of the patterned surface was determined by XPS and the degree of substitution (DS) was calculated to be *ca.* 1% for both paper. This value means that 1 glucose unit is functionalized with photochromic compounds every 100 glucose units. As the degree of substitution of **Cell-DiS** by dithiodiglycolic acid units was *ca.* 14%, the yield of the covalent printing was calculated to be *ca.* 7%. While the DS of the photochromic papers and the associated covalent printing yields could be hastily considered as unsatisfactory, the absorbance of the closed form at very long-wavelength allowed to follow the photochromic behavior of the patterned papers by naked eyes (*vide infra*). Moreover, a high concentration of the dyes onto the surface of cellulose paper might generate detrimental photochromic effects due to cross interactions.

Photophysical properties. In order to evaluate the influence of the cellulose matrix on the photophysical properties of dibenzothienylethenes **9(o)** and **10(o)**, we initially recorded their absorption spectra in solution for comparison with **Cell-DBE-1** and **Cell-DBE-2** (Figure 3). We also used theoretical chemistry to model their properties (*vide infra*).

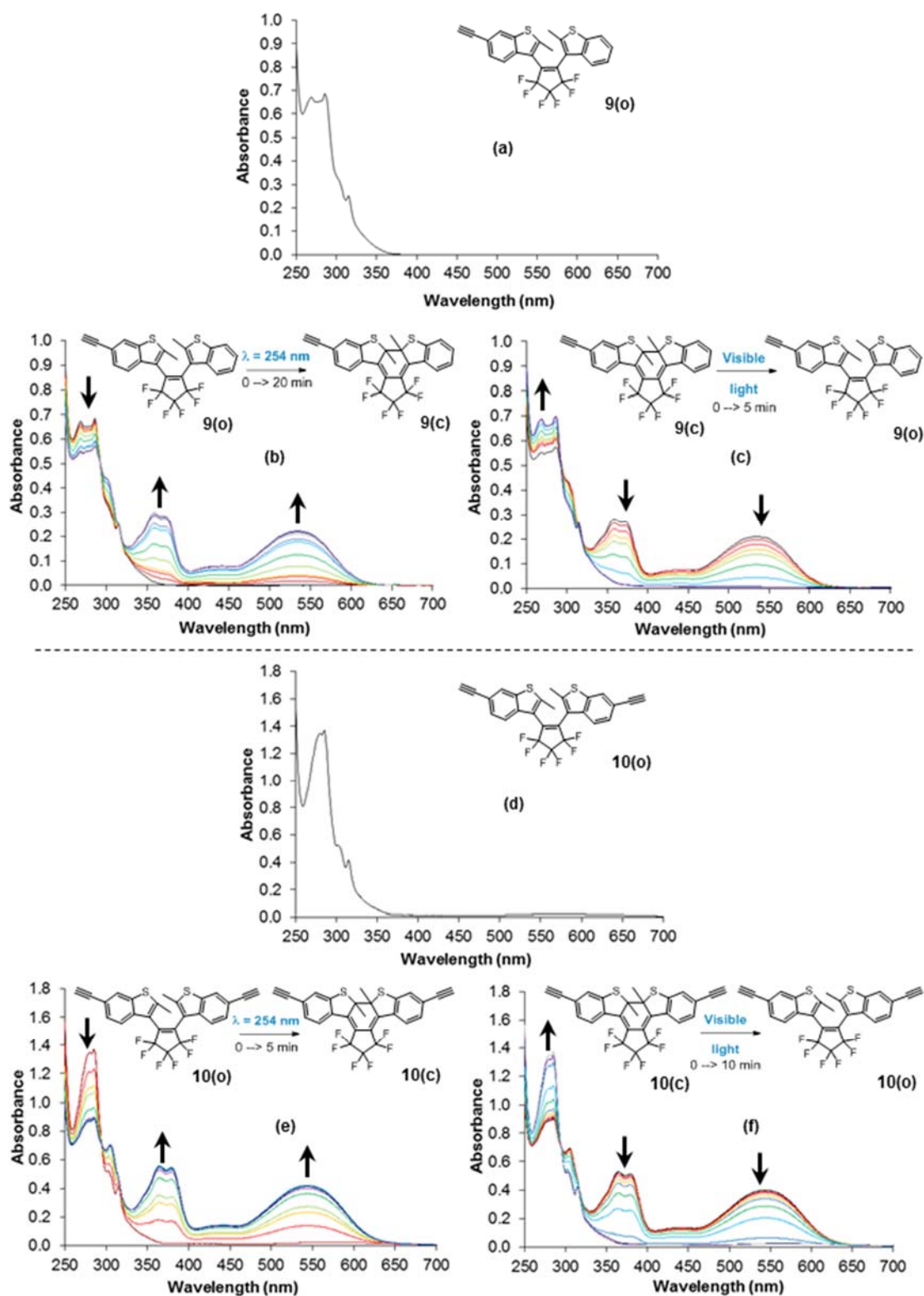


FIGURE 3. (a) Absorption spectra of **9(o)** in the ground state. Time evolution of the absorbance of **9(o)** (b) under irradiation at 254 nm, (c) under visible light. (d) Absorption spectra of **10(o)** in the ground state. Time evolution of the absorbance of **10(o)** (e) under irradiation at 254 nm, (f) under visible light. All data obtained in CH_2Cl_2 at 4×10^{-5} M and 25 °C.

The absorption spectra of **9(o)** and **10(o)**, recorded in CH₂Cl₂ (4×10^{-5} M) at 25 °C, only show absorption bands in the UV region (< 350 nm), as expected. Upon light irradiation of **9(o)** and **10(o)** at 254 nm, the corresponding colorless solutions turn deep pink and purple, respectively. Such color change indicates the successful 6π -electrocyclization of the ring-open isomers **9(o)** and **10(o)** to the corresponding closed forms **9(c)** and **10(c)**, respectively. The photocyclization, upon irradiation at 254 nm, is evidenced by the gradual decay of bands in the UV region ($\lambda < 300$ nm) together with the appearance of strong absorption bands at 359, 371 and 535 nm for the closed-ring isomer **9(c)**, and 364, 379 and 546 nm for the closed-ring isomer **10(c)**. These results show that the presence of a second alkyne function in **10(c)** with respect to **9(c)** redshifts absorption bands due to the increased conjugation (see also calculations below). The photostationary state is reached after 20 minutes of irradiation for **9(c)** and 5 min for **10(c)**. The cycloreversion gradually occurs under daylight (standard lightning in the laboratory) as indicated by the decay of the intensities of the bands in the region of 340-550 nm and the complete bleaching of the solution. The photostationary state is obtained after 5 min for **9(o)** and 10 min for **10(o)** under daylight. We also confirmed that **9(o)** and **10(o)** are both P-type photochromic compounds as thermal relaxation is not observed upon prolonged heating in the dark (refluxing CH₂Cl₂ for 2 hours).

We have performed Density Functional Theory (DFT) calculations to characterize dyes **9(o)**, **10(o)**, **9(c)**, and **10(c)** (see Experimental Section for computational details). For the open-ring structures, the relative free energies of the parallel and anti-parallel conformers are nearly equal (difference < 0.5 kcal.mol⁻¹) indicating that, consistently with experiment, there is therefore only half of the DBE that can undergo photochromism, a usual outcome in diarylethene switches. The closed-ring forms **9(c)** and **10(c)** are respectively less stable by 7.7 and 8.0 kcal.mol⁻¹ as compared to the open **9(o)** and **10(o)**, whereas the thermal **10(c)**-to-**10(o)** barrier is as large as 36 kcal.mol⁻¹, consistent with the observed P-type photochromism. We also carried out TD-DFT calculations to probe the nature of the excited-states. For the hallmark visible absorption of the closed forms, TD-DFT returns transitions at 567 nm ($f=0.36$) and 584 nm ($f=0.47$) for **9(c)** and **10(c)**, respectively. These values are in good agreement with the experimental ones (see Figure 3), the redshift upon addition of a second ethynyl group being reproduced. As expected, these transitions are attributed to a HOMO-LUMO transition. These orbitals are displayed on Figure 4. As can be seen, both orbitals are significantly delocalized on the ethynyl segment(s), consistently with an increase of delocalization and hence a redshift of the absorption spectra.

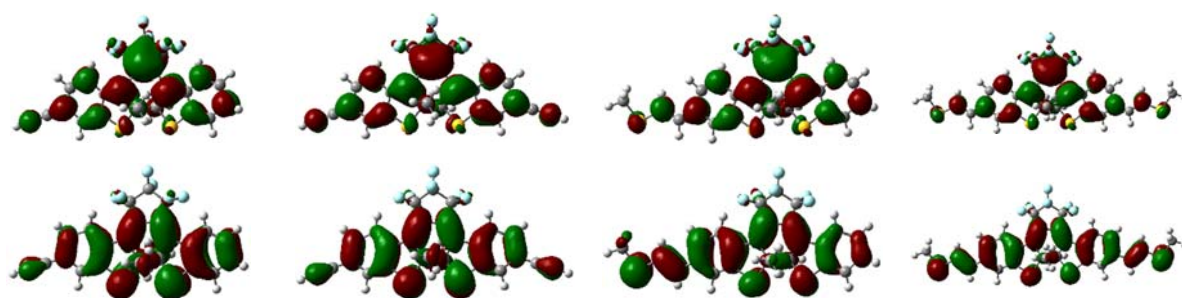


FIGURE 4. Representation of the PCM (CH_2Cl_2)-PBE0-D3^{BJ}/6-31+G(d) frontier orbitals (bottom: HOMO, top: LUMO) for dyes **9(c)** (leftmost), **10(c)** (center left) and the dyes corresponding to DBE anchored onto cellulose **Cell-DBE-1 (9'(c))**, center right) and **Cell-DBE-2 (10'(c))**, rightmost).

Reflective UV-Vis spectroscopy was used to follow the photochromic behavior of **Cell-DBE-1** and **Cell-DBE-2** (Figure 5). Due to the presence of whitening agents, absorbing between 250 to 350 nm, in commercially available cellulose paper, we restricted the absorbance recording at wavelengths >350 nm to follow the characteristic bands of the ring-closed isomers at *ca.* 540 nm. The time evolution of the absorption in the 400-750 nm window was followed upon irradiation at 254 nm. While the ground-state absorption spectra of **Cell-DBE-1** and **Cell-DBE-2** does not show any band between 400 and 750 nm, a new band centered at 534 and 548 nm for **Cell-DBE-1** and **Cell-DBE-2**, respectively, appear after 10 seconds of irradiation. These measurements show that the λ_{max} of the absorption bands at *ca.* 540 nm are similar to the one of **9(o)** and **10(o)** as the band was blue-shifted by 1 nm on **Cell-BDE-1** and red-shifted by 2 nm on **Cell-DBE-2**, respectively. We have also modeled at the TD-DFT level the properties of the DBE presenting a form equivalent to the one after grafting, that is, the DBE in which we replaced the ethynyl segment(s) by 2-methylthiovinyl moiety(ies), and we denote this species **9'** and **10'**. The thermodynamic properties were rather comparable to the one of the original switches with nearly equal free energies for the parallel and anti-parallel conformers and a closed form less stable by 6.4 and 4.9 kcal.mol⁻¹ than the open form for **9'** and **10'**, respectively. The thermal **10'(c)**-to-**10'(o)** barrier is as large as 37 kcal.mol⁻¹, very close to the value obtained for **10** (*vide supra*). The PCM (CH_2Cl_2)-TD-DFT calculations predict that **9'(c)** and **10'(c)** have their lowest transition at 581 nm ($f=0.66$) and 617 nm ($f=1.07$) indicating significant bathochromic and hyperchromic shifts compared to **9(c)** and **10(c)** in the same conditions. The strong increase of the oscillator strength is a positive news as it means that even a small ratio of grafted photochrome could be detected. The predicted bathochromic displacements are a logical consequence of the stronger π -

conjugation in double than triple bonds (see the extended HOMO and LUMO of **9'(c)** and **10'(c)** in Figure 4), but remains unseen experimentally when grafting the dyes on the paper, which hints that cellulose induces a small hypsochromic shift compared to solution. In any case, these theoretical and experimental results suggest that the conjugated π -electron systems were not strongly affected by the grafting, highlighting the high value of our thiol-X ligation for preserving the photophysical properties of photochromic compounds. Comparing **Cell-DBE-1** and **Cell-DBE-2** was also informative as the bi-functional photochromic compound grafted to **Cell-DBE-2** redshifts the absorption band by 12 nm with respect to **Cell-BDE-1** (534 vs. 548 nm). The photostationary state was reached after 20 and 60 minutes for **Cell-BDE-1** and **Cell-DBE-2**, respectively. The reversible photochromic switches were evidenced upon subsequent exposure of the patterned paper to daylight. Then band centered at ca. 540 nm completely disappeared after 40 min for **Cell-BDE-1** and 60 min **Cell-DBE-2**.

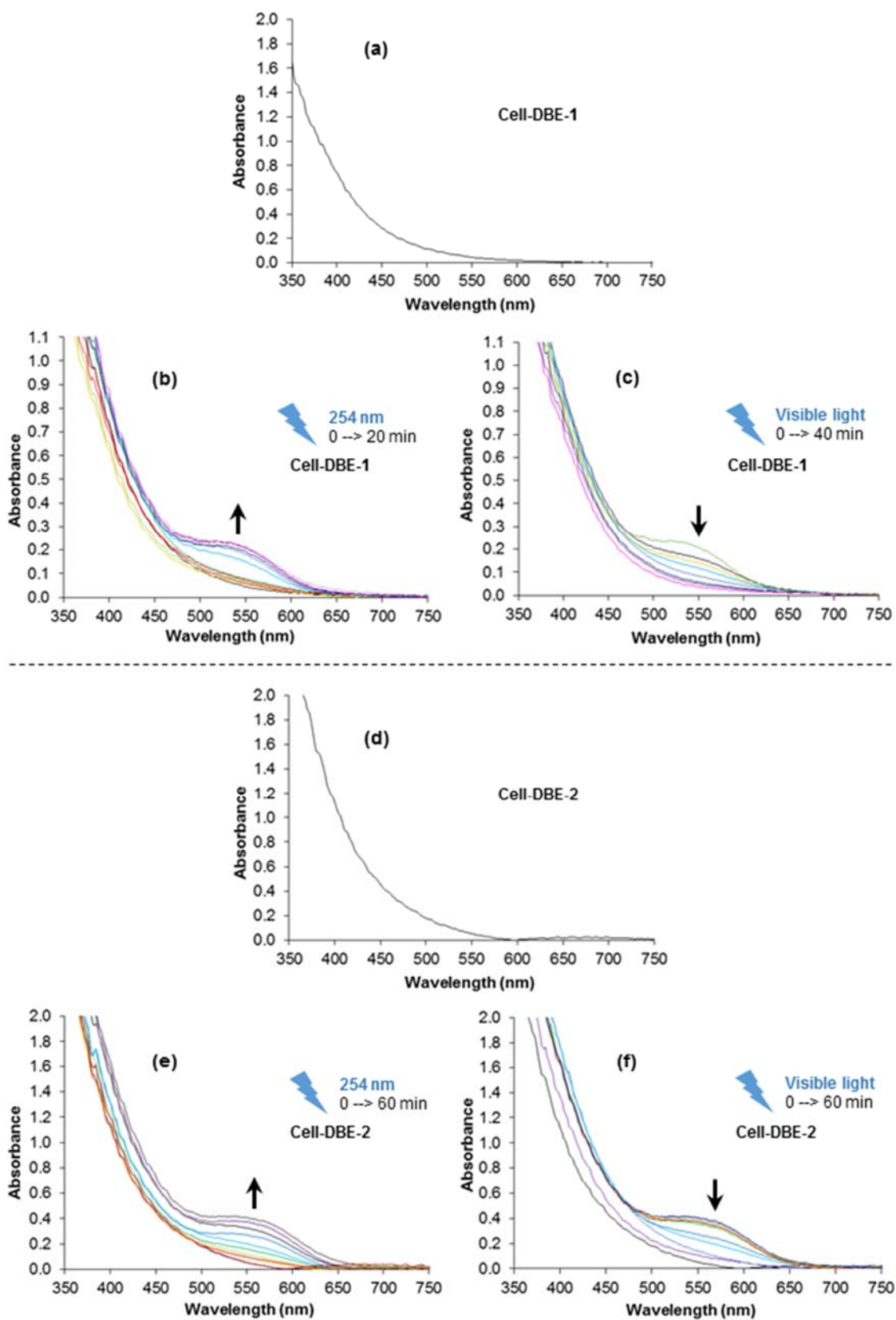


FIGURE 5. (a) Absorbance spectra of **Cell-DBE-1** in the ground state. Time evolution of the absorbance of **Cell-DBE-1** (b) under irradiation at 254 nm, (c) under visible light. (d)

Absorption spectra of **Cell-DBE-2** in the ground state. Time evolution of the absorbance of **Cell-DBE-2** (e) under irradiation at 254 nm, (f) under visible light.

The photochromic behavior of our patterned paper is illustrated on Figure 6. Pictures of patterned papers **Cell-BDE-1** and **Cell-DBE-2** under daylight (Figures 6a and 6c, leftmost) and after irradiation at 254 nm (Figures 6a and 6c, rightmost) show the strong coloration-discoloration switch. An excellent color contrast and a good resolution of the printed letter “U” can be observed for both papers. The precision covalent printing allows patterns with line thickness of *ca.* 250–500 μM . Increasing the precision to line thickness $<200\text{ }\mu\text{M}$ would not be relevant for cellulosic material due to a high heterogeneity of the surface composed of fibers with a diameter usually ranging from 10 to 100 μm . The grafting mostly occurs on the surface of the paper as the photochromic patterns are not visible on the opposite side of **Cell-BDE-1** and **Cell-DBE-2**. The repeatability of the photochromic behavior was studied by applying 18 successive write-erase cycles and neither significant color change nor color intensity were observed. Indeed, as initially expected, both the photophysical and fatigue resistance studies demonstrate that the photochromic behaviors of grafted dibenzothienylethenes are much less affected by the polar environment of the cellulose matrix than merocyanine switches.^{20, 43} The absence of charges in both the open- and closed forms of dibenzothienylethenes most probably prevents any negative interaction with cellulose that could produce unwanted irreversible photochromism behavior.

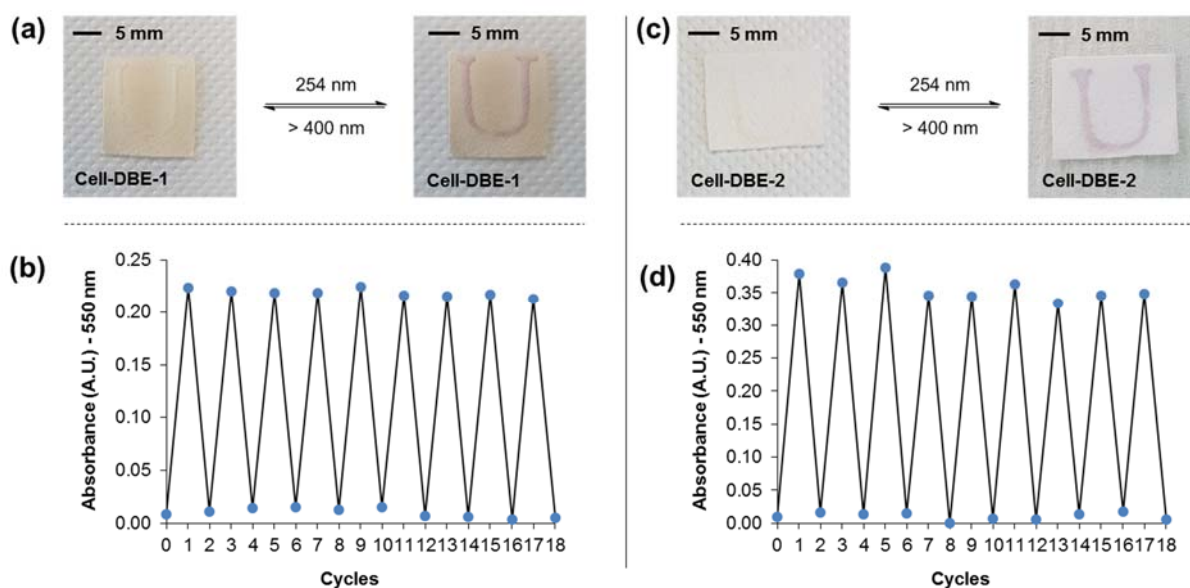


FIGURE 6. Photochromic behavior of patterned papers **Cell-BDE-1** (a) and **Cell-DBE-2** (c). Variation of absorbance intensity of **Cell-BDE-1** (b) and **Cell-DBE-2** (d) under UV (254 nm) and daylight cycles.

In order to compare the robustness of our covalent printing strategy with a simple adsorption process, we prepared the photochromic paper **Cell-DBE-A** which consists of the photochromic compound **8** adsorbed onto pristine cellulose paper (Figure 7). The high instability of **9(o)** and **10(o)** under the dry state precludes any adsorption test with these photochromic compounds on cellulose. While the photochromic behavior of **8** is visually preserved upon adsorption, a single washing of **Cell-DBE-A** in CH_2Cl_2 completely washed-off the circular pattern, validating the better robustness of our covalent printing with respect to adsorption processes. The same washing process carried out on **Cell-DBE-2** (Figure 7d-f) does not affect the pattern, confirming that a covalent printing is much more robust than a simple adsorption.

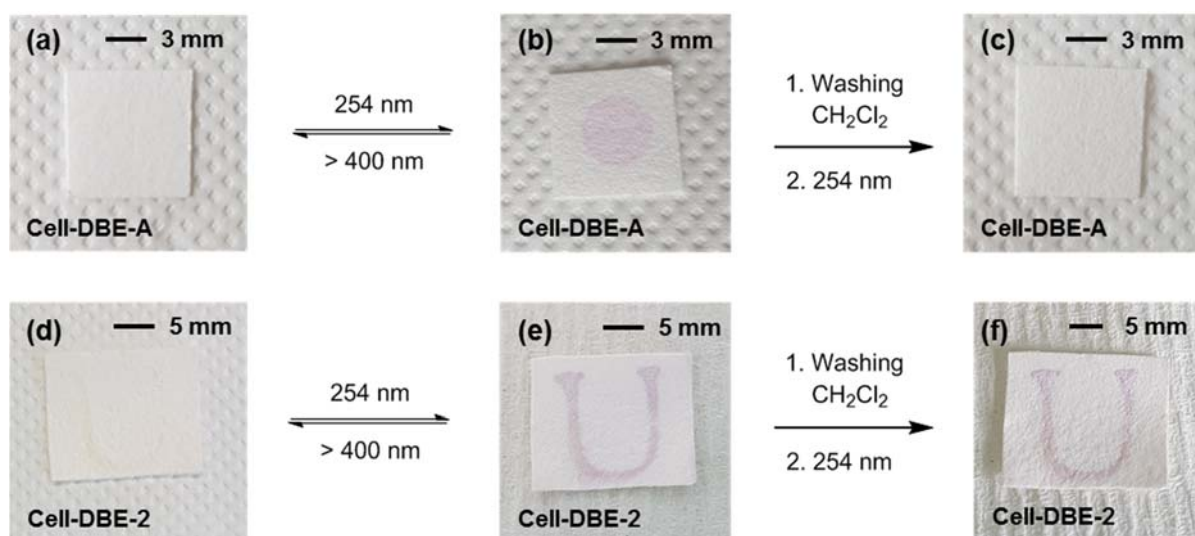


FIGURE 7. Photochromic behavior of patterned papers **Cell-BDE-A** (a) under visible light, (b) after irradiation at 254 nm for 1 min, (c) after a single washing in CH_2Cl_2 followed by an irradiation at 254 nm for 1 min and **Cell-BDE-2** (d) under visible light, (e) after irradiation at 254 nm for 1 min, (f) after a single washing in CH_2Cl_2 followed by an irradiation at 254 nm for 1 min.

■ CONCLUSION

We have developed the first covalent patterning of cellulose paper with photochromic compounds. The strategy stands on the spatially-controlled light-mediated thiol-X ligation of dibenzothiénylenes (DBE) as photochromic compounds. The photophysical and theoretical studies are consistent with a reversible photochromic behavior of a thermally-stable DBE, not significantly affected by the hydrogen bond network of the cellulose matrix, and demonstrates the high fatigue resistance of the photochromic material over 18 successive write-erase cycles without decrease of the absorbance intensity. The robust covalent patterns are insensitive to successive washings in sharp contrast with patterns created through a non-covalent physisorbed process. We trust that the creation of robust molecularly imprinted photochromic patterns on cellulose paper is a promising technology that will find applications as anti-counterfeiting device in fiducial documents.

■ ASSOCIATED CONTENT

The supporting Information is available free of charge on the ACS Publication website at DOI: 10.1021/acsapi.XXXX.

NMR spectra of compounds: **1**, 3-bromo-2-methylbenzothiophene, **4**, **9**, **7**, **8**, **9(o)**, **10(o)** and **11** (PDF)

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Notes

The authors declare no competing financial interest.

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