

Photoredox Propargylation of Aldehydes Catalytic in Titanium

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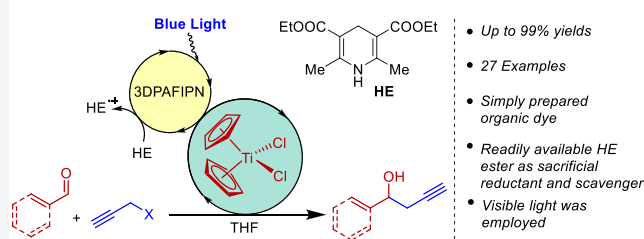
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ABSTRACT: A practical and effective photoredox propargylation of aldehydes promoted by 10 mol % of $[\text{Cp}_2\text{TiCl}_2]$ is presented. No stoichiometric metals or scavengers are used for the process. A catalytic amount of the cheap and simply prepared organic dye 3DPAFIPN is used as the reductant for titanium. The reaction displayed a broad scope, and no traces of allenyl isomers were detected for simple propargyl bromide, whereas mixtures of propargyl and allenyl isomers were observed for substituted propargyl bromides.

Photoredox propargylation of aldehydes mediated by titanium



In just a few years, photoredox catalysis has reached an extraordinary level of advancement, introducing new and exciting methodologies in organic chemistry.¹ Besides electron transfer, many interesting reactions can also be promoted by photocatalytic methodologies using energy transfer (EnT) to reach a reactive transition state in molecules or complexes.² Now, dual photoredox catalysis,³ that is the combination of metal-promoted processes with photoredox cycles, is in continuous development.⁴ From the application of synergistic dual photoredox catalysis to cross-coupling reactions, metal-catalyzed processes were addressed to radical to polar cross-over reactions (RPC),⁵ with the aim of developing important C–C bond-forming reactions.⁶ In this context, allylation methodologies were introduced with chromium,⁷ nickel,⁸ and titanium.⁹ Particularly, the earth-abundant titanium can give important advantages in terms of sustainability and eco-friendliness of the process,¹⁰ and its use in combination with photoredox catalysis was first explored by Gansäuer.¹¹ In addition, the interesting photophysical properties of titanium complexes make the further exploration of their chemistry in the excited state even more intriguing.¹² We recently have reported the allylation reaction of aldehydes, employing 10 mol % of the inexpensive $[\text{Cp}_2\text{TiCl}_2]$, in the presence of an organic dye, 3DPAFIPN,¹³ and Hantzsch's ester as the stoichiometric reductant.⁹ Quite recently, Glorius has reported an interesting carbonyl propargylation¹⁴ via dual chromium/photoredox catalysis, taking advantage of a catalytic radical three components coupling of 1,3-enynes, aldehydes, and radical precursors, in the presence of CrCl_3 and visible light.¹⁵ However, the direct use of propargylic halides or alkynes for the generation of propargyl chromium species is still not reported by means of the emerging dual photoredox catalytic system.

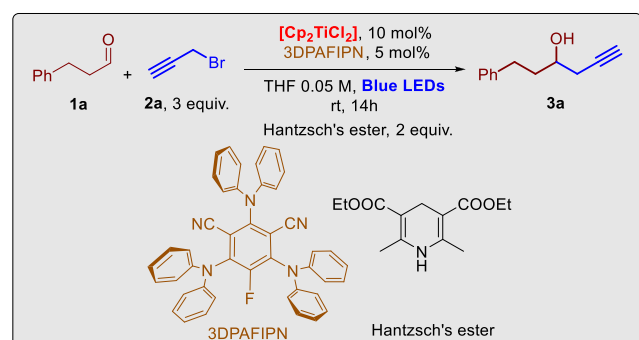
Preliminarily, we have also mentioned that the propargylation reaction of aldehydes was also accessible by the dual photoredox-mediated catalysis with titanium,^{9,16} and herein we illustrate the potentiality, cleanness, and simplicity of the propargylation reaction conducted under photoredox conditions by the use of propargyl bromide.

Starting by employing hydrocinnamic aldehyde **1a**, we have optimized the model propargylation reaction using propargyl bromide, and in general, it was found that the reaction was promoted by the organic dyes 3DPAFIPN, affording the product **3a** in good yields. 4CzIPN¹⁷ and 3CzCIIPN¹⁸ were also tested in the model reactions and gave inferior results (Table 1, entries 3 and 4, respectively). All reactions were performed with the cheap and commercially available $[\text{Cp}_2\text{TiCl}_2]$. As reported in our allylation reaction,^{9a} the use of THF, with a substrate concentration of 0.05 M, was important to allow the desired transformation due to the strong overlap in the absorbance of the photocatalyst and the titanium complex. The optimized conditions are in line with the photophysical observation because the concentration of the red titanium complex ($\epsilon_{455\text{nm}} \approx 250 \text{ M}^{-1} \text{ cm}^{-1}$) allows a significant absorption of the blue photons by 3DPAFIPN ($\epsilon_{455\text{nm}} \approx 2900 \text{ M}^{-1} \text{ cm}^{-1}$) to promote the photoinduced electron transfer. Hantzsch's ester was found to be the best choice as the stoichiometric reductant since various sacrificial reductants (i.e., different amines) were proved to be not suited for the propargylation reaction. This is in part related to the

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Table 1. Propargylation Reaction of Hydrocinnamaldehyde and Variation of Some Reaction Parameters

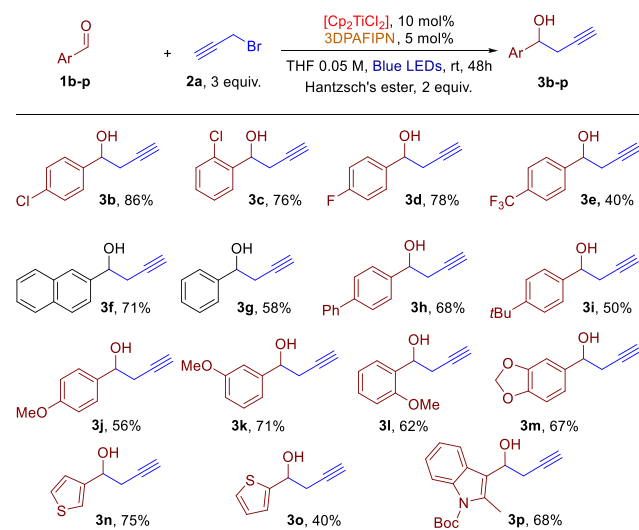
entry ^a	deviations from standard conditions	yield (%) ^b
1 ^c	none	>99(98)
2	1.0 mmol scale, 48 h, 3 mol % of 3DPAFIPN	>99(93)
3	4CzIPN instead of 3DPAFIPN	62
4	3CzClIPN instead of 3DPAFIPN	57
5	no Light	0
6	no Hantzsch's ester	0
7	no [Cp ₂ TiCl ₂]	0
8	no degassed solvent	75
9	no photocatalyst, with irradiation at 456 nm	0
10	in the presence of TEMPO (1 equiv)	10
11	DMF instead of THF	traces
12	CH ₃ CN instead of THF	55

^aReactions performed on a 0.1 mmol scale irradiating with a 40W Kessil lamp, 456 nm. ^bDetermined by ¹H NMR analysis. Values in parentheses represent the yield after chromatographic purification. ^cReaction performed on a 0.2 mmol scale.

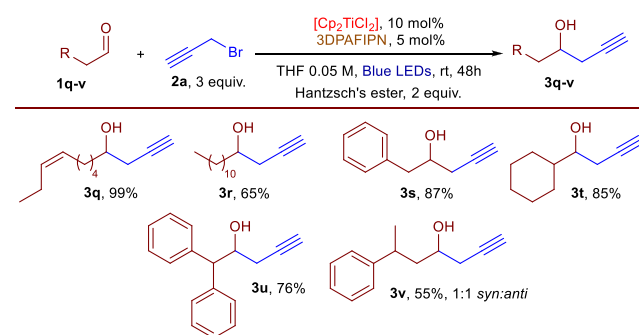
lack of stability of [Cp₂TiCl₂] in the presence of different amines under irradiation.¹⁶ The propargylation reaction was quite sensitive to traces of oxygen and water, and low yields were isolated, performing the reaction in the presence of oxygen (Table 1, entry 8). It is worth noting that the photocatalyst does not decompose during the photoreaction and can be easily recovered at the end of the reaction by flash chromatography.

It was possible to scale the reaction up to 1.0 mmol, increasing the reaction time to 48 h without observing an appreciable decrease of the yield (Table 1, entry 2).

The selected reaction conditions were employed to test the scope of the reaction with aromatic and aliphatic aldehydes. As evident by the data reported in Scheme 1, aromatic and heteroaromatic aldehydes are suitable substrates for the reaction. Yields are, in general, from good to moderate. The presence of electron-withdrawing groups on the aromatic ring reduced the yield of the reaction. Sterical hindrance in the *ortho*-position does not hamper the reactivity, with either activating or deactivating groups. Although the oxidized product of Hantzsch's ester (the corresponding protonated pyridinium) is strongly acidic and could favor undesired reaction pathways involving the indole ring, indole substrates are reactive in the propargylation reaction, giving much better yields compared to what was observed for the allylation reaction.^{9a} Various substituted thiophene carboxaldehydes are suitable substrates for the transformation. As already noted in the allylation reaction, no additives or scavengers are required for the release of the desired homopropargylic alcohol with the concomitant restoration of the titanium complex from

Scheme 1. Dual Photoredox Propargylation of Aromatic Aldehydes

homopropargylic alkoxide. As a matter of fact, the protonated pyridine derivative obtained by the oxidation of the Hantzsch's ester behaves as a scavenger for the reaction, enabling the desired turnover of the employed titanium complex. As was observed for the allylation reaction, also in the photoredox propargylation reaction, 10 mol % of the [Cp₂TiCl₂] complex gave the optimal catalyst concentration. In all reported examples with aromatic aldehydes, no traces of the possible allenic product were detected by ¹H NMR analysis of the crude reaction mixture. The reaction is also quite effective with aliphatic aldehydes (3q–v), and excellent results were obtained (Scheme 2). Branched aliphatic aldehydes were

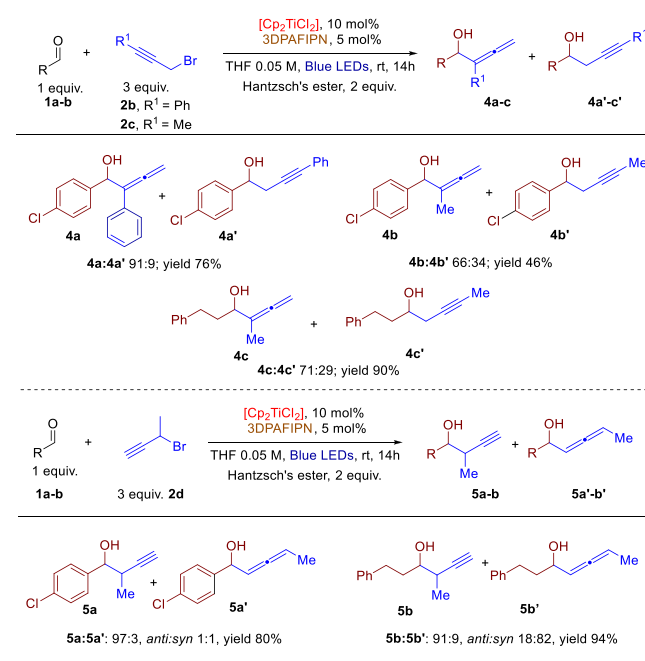
Scheme 2. Dual Photoredox Propargylation of Aliphatic Aldehydes

found to be reactive substrates, furnishing the respective homopropargylic alcohols in good yields. In addition, aliphatic aldehydes with acidic protons, whose propargylation products can suffer from undesired water elimination pathways, are suitable substrates. No modifications in the conditions were made to perform the reaction with aliphatic aldehydes with respect to aromatic substrates.

In general, the reactions ran smoothly without any significant inconveniences. Also, with aliphatic aldehydes, the reactions favored the propargylic derivatives in all examined cases.

We have briefly investigated the outcome of the reaction in the case of different propargylic bromides (Scheme 3).

Scheme 3. Dual Photoredox Propargylation of Aliphatic Aldehydes



Interestingly, the presence of aliphatic or aromatic substituents on the propargylic moiety favors the allenylic product, probably due to the major sterical hindrance of the allenylic titanium intermediate, compared to the propargylic. The synthesis and structural characterizations of allenyl titanocene(IV) and propargyl titanocene(IV) were reported in literature.¹⁹ These compounds are involved in fast metal-lotropic allenyl-propargyl equilibria in solution prior to the electrophilic quenching,²⁰ as confirmed by DFT calculations. Therefore, the reactivity of differently substituted propargylic halides are controlled by the different energy barriers in transition states relative to the reaction of the propargyl and allenyl titanium(IV) precursors with carbonyls via S_E2 mechanism and not by the metalotropic equilibria.¹⁹

The reaction with secondary propargyl bromides gave unsatisfactory simple diastereoselection; in this case, the result could be imputable to the absence of control in the formation of the allenyl titanium intermediate. We have conducted the Stern–Volmer analysis of the reaction, similarity to our previous study of allylation, by simply varying the concentration of propargyl bromide added to the 3DPAFIPN. As illustrated in Figure S3B (see Supporting Information), in air-equilibrated solution, the emission intensity of 3DPAFIPN is barely decreasing upon increasing the concentration of propargyl bromide, thus highlighting a slow diffusional quenching ($k_q 1.0 \times 10^8 \text{ M}^{-1} \text{ s}^{-1}$).

The same conclusions can be drawn by observing the negligible changes in the emission lifetime in the presence of propargyl bromide at concentrations up to ca. 0.13 M (Figure S3C). In degassed solutions, the long-excited state lifetime of pristine 3DPAFIPN (172 μs) is decreasing to 41 μs upon the addition of propargyl bromide at high concentrations (ca. 0.11 M). The estimated quenching constant is 3 orders of magnitude lower than that determined for $[\text{Cp}_2\text{TiCl}_2]$ in the same experimental conditions ($k_q \approx 10^5$ and $5.2 \times 10^8 \text{ M}^{-1} \text{ s}^{-1}$, respectively; see Figure S4B), pointing out that a photo-induced electron transfer is more likely to occur to the latter.

We have already reported⁹ that Hantzsch's ester and $[\text{Cp}_2\text{TiCl}_2]$ are effective quenchers of the photocatalyst. It is also worth mentioning that we have also established that the pyridine salt formed in the reaction is not a quencher at any concentration. In the case of propargylation, the absence of quenching with propargylic bromide suggests the mechanism illustrated in Figure 1. The oxidative quenching of

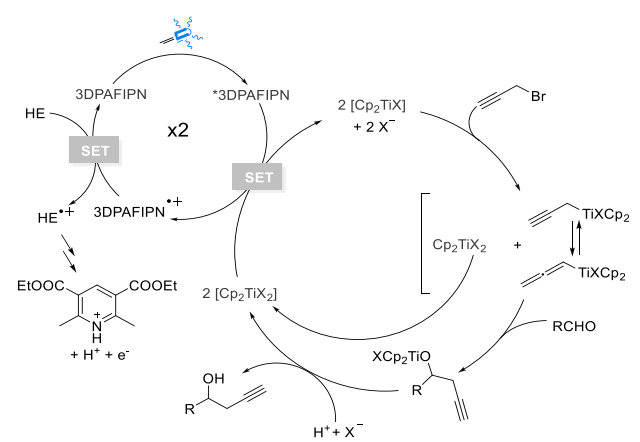


Figure 1. Proposed catalytic cycle.

*3DPAFIPN is responsible for the formation of $[\text{Cp}_2\text{TiCl}]$ and 3DPAFIPN $^{\bullet+}$. The latter is a strong oxidant ($E(\text{PC}^{\bullet+}/\text{PC}) = +1.30$ vs SCE),¹³ and the photoredox cycle is closed by the Hantzsch's ester ($E(\text{HE}^{\bullet+}/\text{HE}) = +1.0$ vs SCE), which reduces the 3DPAFIPN $^{\bullet+}$ back to 3DPAFIPN. The reaction produces the strong reductant $\text{HE}^{\bullet+}$ that can participate in further electron transfer events,²¹ generating the rearomatized Hantzsch's ester, in the form of its pyridinium salt. Furthermore, the oxidative quenching of the photocatalyst in its excited state by the titanium complex $[\text{Cp}_2\text{TiCl}_2]$ generates the $[\text{Cp}_2\text{TiCl}]$ species. A radical-mediated addition²² of $[\text{Cp}_2\text{TiCl}]$ to the propargyl bromide gave the corresponding allenylic/propargylic titanium reagents. Subsequent reaction of the allenylic species with aldehydes gave the titanium-alkoxy derivatives that are transformed into the corresponding alcohols by acidic protons available from the oxidized Hantzsch's ester pyridinium salt. In fact, the latter is an acidic compound, and it features a low pK_a compared to other reagents used as scavengers in the catalytic redox reaction promoted by titanium chemistry (such as collidine-HCl).²³

In summary, we have described a direct photoredox propargylation reaction mediated by a cheap and not toxic titanium complex that affords the desired homopropargylic alcohols in good yields with both aromatic and aliphatic aldehydes. Further studies about metal photoredox-mediated reactions²⁴ are in progress in our laboratory.

EXPERIMENTAL SECTION

General Methods and Materials. ^1H NMR, ^{13}C NMR, and ^{19}F NMR spectra were recorded on a Varian Mercury 400 spectrometer. The residual protic signal of the solvent for the ^1H and $^{13}\text{CDCl}_3$ signals for ^{13}C were used as references for spectra recorded in CDCl_3 (7.26 and 77.16 ppm, respectively). The trifluoroacetic acid signal (−76.55 ppm) was used as a reference for ^{19}F NMR spectra. Chemical shifts are reported in parts per million (ppm) of the δ scale relative to TMS for ^1H and ^{13}C NMR spectra and CFCl_3 for ^{19}F NMR spectra. Data are reported as follows: chemical shift, multiplicity (s = singlet, d = doublet, t = triplet, dd = doublet of doublet, ddd =

doublet of doublet of doublet, td = triplet of doublet, m = multiplet), coupling constants (Hz). Chromatographic purifications were performed with 240–400 mesh silica gel. HPLC-MS analyses were performed on an Agilent Technologies HP1100 instrument coupled with an Agilent Technologies MSD1100 single-quadrupole mass spectrometer using a Phenomenex Gemini C18 3 μm (100 mm \times 3 mm) column; mass spectrometric detection was performed as follows: in full-scan mode from m/z 50 to 2500, scan time 0.1 s in positive ion mode, ESI spray voltage 4500 V, nitrogen gas 35 psi, drying gas flow rate 11.5 mL min^{-1} , and fragmentor voltage 30 V. HRMS was performed on a Waters Xevo G2-XS QToF, ESI⁺, cone voltage 40 V, Capillary 3KV, with a source temperature of 120 $^{\circ}\text{C}$. All reactions were set up under an argon atmosphere in oven-dried glassware (borosilicate) using standard Schlenk techniques. The reaction mixture was irradiated with a Kessil PR160L@456 nm. Lamp technical specifications are available on the manufacturer's web site.²⁵ The reaction vessel was placed 10 cm approximately from the lamp, and the temperature was maintained at room temperature by cooling with a PR160 ring w/fan kit.²⁶ 3DPAFIPN,¹³ 4CzIPN,¹³ 3CzClIPN,¹³ and diethyl 2,6-dimethyl-1,4-dihydropyridine-3,5-dicarboxylate (Hantzsch's Ester)²⁷ were prepared following literature procedures.

Standard Procedure for Photoredox Titanium-Catalyzed Propargylation of Aldehydes. All reactions were performed on a 0.2 mmol scale of aldehyde. A dry 10 mL Schlenk tube, equipped with a Rotaflo stopcock, magnetic stirring bar, and argon supply tube, was first charged under argon with the organic photocatalyst 2,4,6-tris(diphenylamino)-5-fluoroisophthalonitrile 3DPAFIPN (5 mol %, 0.010 mmol, 6.4 mg), [Cp_2TiCl_2] catalyst (10 mol %, 0.02 mmol, 5.0 mg), and diethyl 1,4-dihydro-2,6-dimethyl-3,5-pyridinedicarboxylate, i.e. Hantzsch's ester (2 equiv, 0.4 mmol, 100 mg). Inhibitor-free dry THF (4 mL, in order to obtain a 0.05 M solution of aldehyde) was then added, the reaction mixture was further subjected to a freeze–pump–thaw procedure (three cycles), and the vessel was refilled with argon. Then, propargyl bromide derivative 2a–d (0.6 mmol, 3 equiv) and the substrate 1a–v (0.2 mmol) were added. The reaction was irradiated under vigorous stirring for 14 h. After that, the reaction mixture was quenched with H_2O (approximately 5 mL) and extracted with AcOEt (4 \times 5 mL). The combined organic layers were dried over anhydrous Na_2SO_4 , and the solvent was removed under reduced pressure. The crude was subject of flash column chromatography (SiO_2) to afford the products 3a–v, 4a–c, and 5a–b in the stated yields.

Procedure for 1 mmol Scale. A dry 50 mL Schlenk tube, equipped with a magnetic stirring bar and argon supply tube, was first charged under argon with the organic photocatalyst 3DPAFIPN 2,4,6-tris(diphenylamino)-5-fluoroisophthalonitrile (3 mol %, 0.03 mmol, 19 mg), [Cp_2TiCl_2] catalyst (8 mol %, 0.08 mmol, 0.020 g), and diethyl 1,4-dihydro-2,6-dimethyl-3,5-pyridinedicarboxylate, namely, Hantzsch's ester (2 equiv, 2 mmol, 0.500 g). Inhibitor-free dry THF (20 mL in order to obtain a 0.05 M solution of aldehyde) was then added, the reaction mixture was further subjected to a freeze–pump–thaw procedure (four cycles), and the vessel refilled with argon. Then, propargyl bromide 2a (80% v/v in toluene, 3 mmol, 3 equiv, 0.280 mL) and the substrate 1a (1 mmol, 0.134 g, 0.132 mL) were added. The reaction was irradiated under vigorous stirring for 48 h. After that, the solvent was removed under reduced pressure. The crude was subjected to flash column chromatography (SiO_2) to afford the products 3a in 93% yield (0.93 mmol, 0.162 g).

1-Phenylhex-5-yn-3-ol (3a): brown oil, 98% (0.195 mmol, 34 mg). The general procedure was applied using 1a (0.2 mmol, 26 μL) previously distilled and 2a (solution 80% v/v in toluene, 0.6 mmol, 56 μL , 3 equiv). The title compound was isolated by flash column chromatography (100% DCM). Spectroscopic data were in agreement with those reported in literature.²⁸

1-(4-Chlorophenyl)but-3-yn-1-ol (3b): brown oil, 86% (0.17 mmol, 31 mg). The general procedure was applied using 1b (0.2 mmol, 28 mg) and 2a (solution 80% v/v in toluene, 0.6 mmol, 56 μL , 3 equiv). The title compound was isolated by flash column

chromatography (100% DCM). Spectroscopic data were in agreement with those reported in literature.²⁹

1-(2-Chlorophenyl)but-3-yn-1-ol (3c): brown oil, 76% (0.15 mmol, 27 mg). The general procedure was applied using 1c (0.2 mmol, 22.4 μL) previously distilled and 2a (solution 80% v/v in toluene, 0.6 mmol, 56 μL , 3 equiv). The title compound was isolated by flash column chromatography (100% DCM). Spectroscopic data were in agreement with those reported in literature.²⁸

1-(4-Fluorophenyl)but-3-yn-1-ol (3d): brown oil, 78% (0.16 mmol, 22.3 mg). The general procedure was applied using 1d (0.2 mmol, 22 μL) previously distilled and 2a (solution 80% v/v in toluene, 0.6 mmol, 56 μL , 3 equiv). The title compound was isolated by flash column chromatography (100% DCM). Spectroscopic data were in agreement with those reported in literature.³⁰

1-(4-(Trifluoromethyl)phenyl)but-3-yn-1-ol (3e): brown oil, 40% (0.08 mmol, 18 mg). The general procedure was applied using 1e (0.2 mmol, 28 μL) previously distilled and 2a (solution 80% v/v in toluene, 0.6 mmol, 56 μL , 3 equiv). The title compound was isolated by flash column chromatography (100% DCM). Spectroscopic data were in agreement with those reported in literature.³¹

1-(Naphthalen-2-yl)but-3-yn-1-ol (3f): brown oil, 71% (0.14 mmol, 28 mg). The general procedure was applied using 1f (0.2 mmol, 32 mg) and 2a (solution 80% v/v in toluene, 0.6 mmol, 56 μL , 3 equiv). The title compound was isolated by flash column chromatography (100% DCM). Spectroscopic data were in agreement with those reported in literature.⁸

1-Phenylbut-3-yn-1-ol (3g): brown oil, 58% (0.12 mmol, 18 mg). The general procedure was applied using 1g (0.2 mmol, 20.4 μL) previously distilled and (solution 80% v/v in toluene, 0.6 mmol, 56 μL , 3 equiv). The title compound was isolated by flash column chromatography (100% DCM). Spectroscopic data were in agreement with those reported in literature.³²

1-([1,1'-Biphenyl]-4-yl)but-3-yn-1-ol (3h): brown oil, 68% (0.14 mmol, 30.2 mg). The general procedure was applied using 1h (0.2 mmol, 36 mg) and 2a (solution 80% v/v in toluene, 0.6 mmol, 56 μL , 3 equiv). The title compound was isolated by flash column chromatography (100% DCM). Spectroscopic data were in agreement with those reported in literature.³³

1-(4-(tert-Butyl)phenyl)but-3-yn-1-ol (3i): brown oil, 50% (0.1 mmol, 20.2 mg). The general procedure was applied using 1i (0.2 mmol, 32 μL) previously distilled and 2a (solution 80% v/v in toluene, 0.6 mmol, 56 μL , 3 equiv). The title compound was isolated by flash column chromatography (100% DCM). Spectroscopic data were in agreement with those reported in literature.³⁴

1-(4-Methoxyphenyl)but-3-yn-1-ol (3j): brown oil, 56% (0.11 mmol, 19 mg). The general procedure was applied using 1j (0.2 mmol, 26 μL) and 2a (solution 80% v/v in toluene, 0.6 mmol, 56 μL , 3 equiv). The title compound was isolated by flash column chromatography (100% DCM). Spectroscopic data were in agreement with those reported in literature.³⁵

1-(3-Methoxyphenyl)but-3-yn-1-ol (3k): brown oil, 71% (0.14 mmol, 25 mg). The general procedure was applied using 1k (0.2 mmol, 26 μL) and 2a (solution 80% v/v in toluene, 0.6 mmol, 56 μL , 3 equiv). The title compound was isolated by flash column chromatography (100% DCM). Spectroscopic data were in agreement with those reported in literature.²⁹

1-(2-Methoxyphenyl)but-3-yn-1-ol (3l): brown oil, 62% (0.12 mmol, 22 mg). The general procedure was applied using 1l (0.2 mmol, 26 μL) and 2a (solution 80% v/v in toluene, 0.6 mmol, 56 μL , 3 equiv). The title compound was isolated by flash column chromatography (100% DCM). Spectroscopic data were in agreement with those reported in literature.³¹

1-(Benzo[d][1,3]dioxol-5-yl)but-3-yn-1-ol (3m): brown oil, 67% (0.13 mmol, 25 mg). The general procedure was applied using 1m (0.2 mmol, 38 mg) and 2a (solution 80% v/v in toluene, 0.6 mmol, 56 μL , 3 equiv). The title compound was isolated by flash column chromatography (100% DCM). Spectroscopic data were in agreement with those reported in literature.²⁹

1-(Thiophen-3-yl)but-3-yn-1-ol (3n): brown oil, 75% (0.15 mmol, 23 mg). The general procedure was applied using 1n (0.2

mmol, 18 μL) and **2a** (solution 80% v/v in toluene, 0.6 mmol, 56 μL , 3 equiv). The title compound was isolated by flash column chromatography (100% DCM). Spectroscopic data were in agreement with those reported in literature.³⁶

1-(Thiophen-2-yl)but-3-yn-1-ol (3o): brown oil, 40% (0.08 mmol, 12 mg). The general procedure was applied using **1o** (0.2 mmol, 18 μL) and **2a** (solution 80% v/v in toluene, 0.6 mmol, 56 μL , 3 equiv). The title compound was isolated by flash column chromatography (100% DCM). Spectroscopic data were in agreement with those reported in literature.³²

tert-Butyl 3-(1-Hydroxybut-3-yn-1-yl)-2-methyl-1H-indole-1-carboxylate (3p): brown oil, 68% (0.14 mmol, 41 mg). The general procedure was applied using **1p** (0.2 mmol, 52 mg) and **2a** (solution 80% v/v in toluene, 0.6 mmol, 56 μL , 3 equiv). The title compound was isolated by flash column chromatography (100% DCM). ¹H NMR (401 MHz, CDCl₃): δ 8.10 (d, J = 8.0 Hz, 1H), 7.77 (d, J = 7.6 Hz, 1H), 7.30–7.16 (m, 2H overlapped with the residual peak of the solvent), 5.22 (dd, J = 7.9, 6.2 Hz, 1H), 2.96 (ddd, J = 16.7, 8.2, 2.4 Hz, 1H), 2.68 (ddd, J = 16.7, 5.8, 2.4 Hz, 1H), 2.60 (s, 3H), 2.35 (s, 1H), 2.06 (t, J = 2.3 Hz, 1H), 1.68 (s, 9H). ¹³C{¹H} NMR (101 MHz, CDCl₃): δ 150.6, 136.0, 134.2, 127.3, 123.5, 122.4, 119.4, 118.4, 115.4, 83.9, 80.9, 70.5, 67.1, 28.2, 27.4, 14.2. HRMS (ESI/Q-TOF): m/z [M-Boc-H₂O + H]⁺ calcd for C₁₃H₁₁N, 181.0891; found, 182.0961. HRMS (ESI/Q-TOF): m/z [M + Na]⁺ calcd for C₁₈H₂₁NO₃Na, 322.1419, found, 322.1409.

(Z)-Dodec-9-en-1-yn-4-ol (3q): yellow oil, 99% (0.2 mmol, 36 mg). The general procedure was applied using **1q** (0.2 mmol, 34 μL) and **2a** (solution 80% v/v in toluene, 0.6 mmol, 56 μL , 3 equiv). The title compound was isolated by flash column chromatography (100% DCM). ¹H NMR (401 MHz, CDCl₃): δ 5.50–5.19 (m, 2H), 3.80–3.71 (m, 1H), 2.43 (ddd, J = 16.7, 4.7, 2.6 Hz, 1H), 2.31 (ddd, J = 16.7, 6.7, 2.6 Hz, 1H), 2.11–1.98 (m, 5H), 1.97–1.85 (m, 1H), 1.57–1.51 (m, 2H), 1.44–1.35 (m, 3H), 1.25 (m, 1H), 0.95 (td, J = 7.5, 2.2 Hz, 3H). ¹³C{¹H} NMR (101 MHz, CDCl₃): δ 131.8, 128.8, 80.8, 70.7, 69.8, 36.1, 29.6, 27.3, 26.9, 25.2, 20.5, 14.3. ESI-MS: m/z 198.2 [M + NH₄]⁺. HRMS (ESI/Q-TOF): m/z [M + NH₄]⁺ calcd for C₁₂H₂₄NO, 198.1858; found, 198.1853.

Pentadec-1-yn-4-ol (3r): brown oil, 65% (0.13 mmol, 25 mg). The general procedure was applied using **1r** (0.2 mmol, 32 μL) and **2a** (solution 80% v/v in toluene, 0.6 mmol, 56 μL , 3 equiv). The title compound was isolated by flash column chromatography (100% DCM). Spectroscopic data were in agreement with those reported in literature.³⁷

1-Phenylpent-4-yn-2-ol (3s): brown oil, 87% (0.17 mmol, 28 mg). The general procedure was applied using **1s** (0.2 mmol, 23 μL) and **2a** (solution 80% v/v in toluene, 0.6 mmol, 56 μL , 3 equiv). The title compound was isolated by flash column chromatography (100% DCM). Spectroscopic data were in agreement with those reported in literature.³⁸

1-Cyclohexylbut-3-yn-1-ol (3t): brown oil, 85% (0.17 mmol, 26 mg). The general procedure was applied using **1t** (0.2 mmol, 22 μL) and **2a** (solution 80% v/v in toluene, 0.6 mmol, 56 μL , 3 equiv). The title compound was isolated by flash column chromatography (100% DCM). Spectroscopic data were in agreement with those reported in literature.²⁹

1,1-Diphenylpent-4-yn-2-ol (3u): brown oil, 76% (0.15 mmol, 36 mg). The general procedure was applied using **1u** (0.2 mmol, 39.2 mg) and **2a** (solution 80% v/v in toluene, 0.6 mmol, 56 μL , 3 equiv). The title compound was isolated by flash column chromatography (100% DCM). ¹H NMR (401 MHz, CDCl₃): δ 7.41 (dd, J = 8.2, 1.0 Hz, 2H), 7.37–7.18 (m, 8H), 4.54 (ddd, J = 8.7, 6.3, 4.5 Hz, 1H), 4.13 (d, J = 8.7 Hz, 1H), 2.46 (ddd, J = 16.9, 4.3, 2.7 Hz, 1H), 2.30 (ddd, J = 16.9, 6.4, 2.7 Hz, 1H), 2.17–2.08 (m, 1H), 2.04 (s, 1H). ¹³C{¹H} NMR (101 MHz, CDCl₃): δ 141.7, 140.6, 128.8 (3C), 128.7 (2C), 128.6 (2C), 128.2, 126.8, 126.9, 80.6, 71.7, 71.1, 56.9, 25.3. HRMS (ESI/Q-TOF) m/z : [M-H₂O + H]⁺ calcd for C₁₇H₁₅N, 219.1174, found, 219.1164.

6-Phenylhept-1-yn-4-ol (3v): brown oil, 55% (0.11 mmol, 20 mg) as a *syn/anti* mixture, dr of 1:1. The general procedure was applied using **1v** (0.2 mmol, 29.6 mg) and **2a** (solution 80% v/v in

toluene, 0.6 mmol, 56 μL , 3 equiv). The title compound was isolated by flash column chromatography (100% DCM). In analogy with the reported data in literature for similar compounds,³⁹ it was possible to distinguish the *syn* and the *anti* isomer. The diastereoisomeric ratio was calculated, considering the ¹H NMR spectrum of the reaction crude, comparing the integral of the signal at 3.77 ppm for the *syn* product and at 3.49 for the *anti* product. ³v_{syn} ¹H NMR (401 MHz, CDCl₃): δ 7.33–7.18 (m, 5H, overlapped with the residual peak of the solvent and peaks related to the other isomer aromatic protons), 3.77 (s, 1H), 3.09–2.97 (m, 1H), 2.52–2.39 (m, 1H), 2.05 (t, J = 12.0, 2.6 Hz, 2H), 1.93–1.83 (m, 3H), 1.33–1.29 (m, 3H). ¹³C NMR (101 MHz, CDCl₃): δ 146.9, 128.6 (2H), 127.1 (2H), 126.2, 80.8, 70.9, 68.0, 44.6, 36.5, 27.9, 23.1. ³v_{anti} ¹H NMR (401 MHz, CDCl₃): δ 7.33–7.18 (m, 5H, overlapped with the residual peak of the solvent and peaks related to the other isomer aromatic protons), 3.49 (s, 1H), 2.96–2.87 (m, 1H), 2.37–2.20 (m, 1H), 2.02 (t, J = 12.0, 2.6 Hz, 2H), 1.82–1.74 (m, 3H), 1.28–1.24 (m, 3H). ¹³C{¹H} NMR (101 MHz, CDCl₃): δ 146.3, 128.5 (2H), 126.8 (2H), 126.1, 80.6, 70.8, 67.7, 44.5, 36.4, 27.4, 22.0. HRMS (ESI/Q-TOF) m/z : [M-H₂O + H]⁺ calcd for C₁₃H₁₅, 171.1174; found, 171.1160.

1-(4-Chlorophenyl)-2-phenylbuta-2,3-dien-1-ol and 1-(4-Chlorophenyl)-4-phenylbut-3-yn-1-ol (4a–4a'): brown oil, 76% (0.14 mmol, 36 mg) as a mixture regioisomers **4a/4a'** (91:9). The general procedure was applied using **1b** (0.2 mmol, 28 mg) and **2b** (0.6 mmol, 117 mg, 3 equiv). The ratio of regioisomer was calculated considering the ¹H NMR spectrum of the reaction crude and comparing the integral of the signal at 5.71 ppm for **4a** and at 4.91 ppm for **4a'**. The title compound was isolated by flash column chromatography (100% DCM). Spectroscopic data were in agreement with those reported in literature.⁴⁰

1-(4-Chlorophenyl)-2-methylbuta-2,3-dien-1-ol and 1-(4-chlorophenyl)pent-3-yn-1-ol (4b–4b'): brown oil, 46% (0.1 mmol, 18 mg) as a mixture of regioisomers **4b/4b'** (66:34). The general procedure was applied using **1b** (0.2 mmol, 28 mg) and **2c** (0.6 mmol, 52 μL , 3 equiv). The regioisomeric ratio was calculated considering the ¹H NMR spectrum of the reaction crude and comparing the integral of the signal at 5.10 ppm for **4b** and at 4.77 ppm for **4b'**. The title compound was isolated by flash column chromatography (100% DCM). Spectroscopic data were in agreement with those reported in literature.⁴¹

4-Methyl-1-phenylhexa-4,5-dien-3-ol and 1-phenylhept-5-yn-3-ol (4c–4c'): brown oil, 90% (0.18 mmol, 34 mg) as a mixture of regioisomers **4c/4c'** (71:29). The general procedure was applied using **1v** (0.2 mmol, 26 μL) and **2c** (0.6 mmol, 52 μL , 3 equiv). The regioisomeric ratio was calculated considering the ¹H NMR spectrum of the reaction crude and comparing the integral of the signal at 4.81 ppm related for **4c** and at 4.72 ppm for **4c'**. The title compound was isolated by flash column chromatography (100% DCM). Spectroscopic data were in agreement with those reported in literature.⁴²

1-(4-Chlorophenyl)-2-methylbut-3-yn-1-ol and 1-(4-chlorophenyl)penta-2,3-dien-1-ol (5a–5a'): brown oil, 80% (0.16 mmol, 31 mg) as a mixture of regioisomers and diastereoisomers **5a/5a'** of 97:3, **5a_{syn}/5a_{anti}** dr of 1:1. The regioisomeric ratio was calculated considering the ¹H NMR spectrum of the reaction crude and comparing the integrals of the signal at 4.71 ppm for **5a** and at 5.29 ppm for **5a'**. The diastereoisomeric ratio was calculated considering the ¹H NMR spectrum of the reaction crude comparing the integral of the signal at 4.71 ppm for the product **5a_{syn}** and at 4.51 ppm for the product **5a_{anti}**. The general procedure was applied using **1b** (0.2 mmol, 28 mg) and **2d** (0.6 mmol, 79 mg, 3 equiv). The title compound was isolated by flash column chromatography (100% DCM). Spectroscopic data were in agreement with those reported in literature.⁴³

4-Methyl-1-phenylhex-5-yn-3-ol and 1-phenylhepta-4,5-dien-3-ol (5b–5b'): brown oil, 94% (0.19 mmol, 35 mg) as a mixture of regioisomers and diastereoisomers of **5b/5b'** of 92:8, **5b_{syn}/5b_{anti}** dr of 18:82. The regioisomeric ratio was calculated, considering the ¹H NMR spectrum of the reaction crude, comparing the integral of the signal at 3.49 ppm for **5b** and at 4.80 ppm for **5b'**. The diastereoisomeric ratio was calculated, considering the ¹H NMR

spectrum of the reaction crude, comparing the integral of the signal at 3.54 ppm related to the product **5b_{syn}** and the multiplet (1H) at 3.49 ppm related to the product **5b_{anti}**. The general procedure was applied using **1a** (0.2 mmol, 26 μ L) and **2d** (0.6 mmol, 79 mg, 3 equiv). The title compound was isolated by flash column chromatography (100% DCM). Spectroscopic data were in agreement with those reported in literature.⁴⁴

Photophysical and Mechanistic Studies. All photophysical analyses were carried out in air-equilibrated tetrahydrofuran at 298 K unless otherwise specified. UV–vis absorption spectra were recorded with a PerkinElmer λ 40 spectrophotometer using quartz cells with an optical path length of 1.0 cm. Degassed solutions are obtained by means of repeated pump–freeze–thaw cycles (ca. 4×10^{-6} mbar) in sealed quartz cuvettes. Luminescence spectra were performed with a PerkinElmer LS-50, a Varian Cary Eclipse, or an Edinburgh FLS920 spectrofluorimeter equipped with a Hamamatsu R928 phototube. Lifetimes shorter than 10 μ s were measured by the above-mentioned Edinburgh FLS920 spectrofluorimeter equipped with a TCC900 card for data acquisition in time-correlated single-photon counting experiments (0.5 ns time resolution). The estimated experimental errors are 2 nm on the band maximum, 5% on the molar absorption coefficient, and luminescence lifetime.

■ ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/acs.joc.1c00521>.

Photos of reaction setup, copies of NMR spectra, and photophysical study results (PDF)

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