


 Cite this: *RSC Adv.*, 2025, 15, 31806

Tandem synthesis of dihydronaphthalen-1(2*H*)-one derivatives *via* aldol condensation–Diels–Alder–aromatization sequence of reactions

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A new series of dihydronaphthalen-1(2*H*)-one derivatives were synthesized in high yields starting from commercially available 3,5,5-trimethylcyclohex-2-en-1-one **1a**, aromatic aldehydes **2**, and diethyl acetylenedicarboxylate. Reaction of **1a** with the aldehydes produced the respective dienones **3**, which could cycloadd to dialkyl acetylenedicarboxylate, either stepwise or *in situ*, under aqueous/organocatalyzed (DMAP) conditions. The respective adducts **4**, were produced efficiently *via* a Diels–Alder–double bond isomerization–oxidative aromatization sequence and were characterized based on the analysis of their ¹H and ¹³C NMR spectra.

 Received 1st July 2025
 Accepted 25th August 2025

DOI: 10.1039/d5ra04673d

rsc.li/rsc-advances

Introduction

4-Dimethylaminopyridine (DMAP) is an amine with improved basicity¹ and nucleophilicity,² suitable to accelerate various organic transformations conveniently.³ As a catalyst, DMAP is used in the synthesis of α,β -unsaturated δ -lactones,⁴ Buchwald–Hartwig C–N coupling,⁵ *N*-vinylation,⁶ [4 + 2] cycloadditions,⁷ and many other synthetic reactions.⁸ DMAP is also conveniently used in domino carbonylation–electrocyclization synthesis of imidazodipyridines⁹ and cascade synthesis of α -pyrones.¹⁰ In addition to usual homogeneous uses, DMAP can be used in the form of ionic liquids¹¹ or immobilized derivatives.¹² The advantages of DMAP in organic chemistry are not limited to its regular basic or nucleophilic properties and are additionally highlighted due to its uses in biological,¹³ medicinal,^{14,15} and nanoparticle disciplines,¹⁶ while DMAP has shown several applications in chiral catalysis¹⁷ and asymmetric synthesis¹⁸ and has got involved in synthetic protocols as a reactant¹⁹ or the active part of starting materials.²⁰ Some illustrative cases are highlighted in Fig. 1.

Cyclohexenone derivatives are a group of usually inexpensive commercially available reactants for various convenient transformations to more complex products,^{21–23} including the aromatization process to their respective phenol equivalents.²⁴ Among this group of compounds, the 3-methyl substituted derivative **1** would be a suitable candidate for both aromatization to the respective phenol and chain extension *via* aldol condensation with aldehydes²⁵ or coupling reactions with

vinyllogous species²⁶ at the allylic methyl position. The resulting dienes have the advantages for further synthetic manipulations through cycloaddition reactions with acetylenic dienophiles to access the respective polysubstituted dihydronaphthalen-1-one (A) or naphthol analogues (B), which in turn are useful synthons to access natural^{27,28} or synthetic²⁹ structures (Fig. 2).

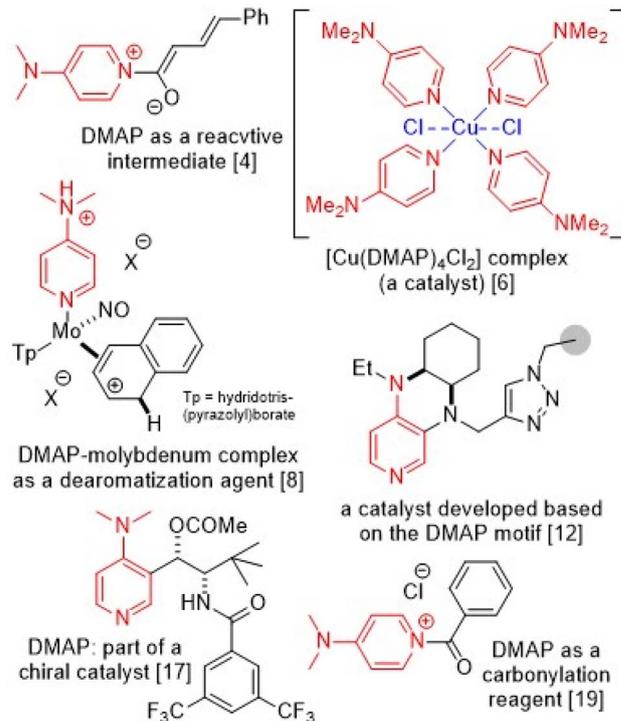


Fig. 1 Some important synthetic roles of DMAP.

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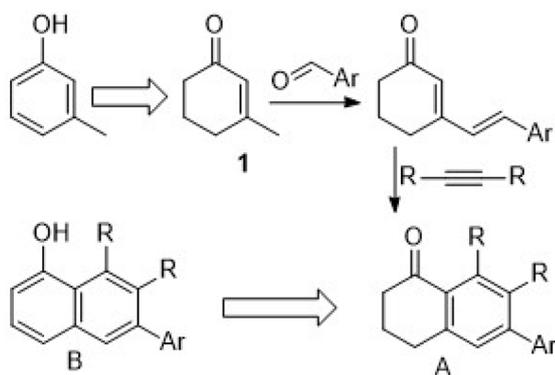



Fig. 2 3-Methylcyclohexenone route to aromatic DA adducts.

The [4 + 2] Diels–Alder (DA) cycloaddition is a key reaction in synthetic organic chemistry,³⁰ due to efficient formation of a cyclohexene ring with up to four stereogenic centres with predictable selectivity.³¹ Incorporation of this reaction with other important organic transformations into tandem processes multiplies the efficiency of the DA cycloaddition so that many synthetic applications have arisen in recent decades as a result of DA-Hantzsch,³² Biginelli-DA,³³ Heck-DA,³⁴ Knoevenagel-DA,³⁵ aldol condensation-DA,³⁶ and other sequential reactions.^{37,38}

We are interested in the study of DA reactions and have communicated our findings on the synthesis and cycloaddition reactions of the styrylcyclohexene diene system in recent years.^{39,40} Based on this background, we were persuaded to

extend our study to embark on direct synthesis of the dihydronaphthalen skeleton starting from 3,5,5-trimethylcyclohex-2-en-1-one (**1a**, isophorone). We hereby disclose the synthesis of various derivatives of **4** through a tandem aldol-condensation-Diels–Alder-rearrangement-aromatization sequence of reactions under aqueous/DMAP conditions, as depicted in Scheme 1 for the reaction of **1a** with benzaldehyde and the following cycloaddition step of **3a** with diethyl acetylenedicarboxylate (DEAD).

Results and discussion

We first optimized the conditions for both steps. While a mixture of **1a** and benzaldehyde **2a** in aqueous DMAP gave low quantities of **3a** at room temperature (entry 1), treatment of the same mixture at 60 °C produced 84% of the product after 3 h (entry 2). In contrast, the yield diminished in the absence of either DMAP (entry 3) or water (entry 4). This was also the case when water was replaced with an alcohol (entries 5–6), toluene (entry 7), MeCN (entry 8), or THF (entry 9) as the medium of the reaction.⁴¹ Alternatively, separation of **3a** and its treatment with DEAD in the same H₂O/DMAP medium gave **4a** in much higher quantities at reflux (entry 10).⁴² Treatment of the same mixture at lower temperatures was less productive (entries 11 and 12) (Table 1).

Having the two sets of optimized conditions, we next evaluated the generality of the method by using other derivatives of **2** (Table 2). Thus, in addition to unsubstituted aromatic aldehydes (entries 1 and 2), the H₂O/DMAP mediated condensation of various aldehydes bearing electron releasing (entries 3 and 4)

Scheme 1 Typical stepwise reaction pathway for the conversion of **1** to **3** and **3** to **4**.Table 1 Three-component optimization of the synthesis of **3a**

Entry	Reactants	Conditions ^a	Temperature (°C)	Time (h)	Product	Yield ^b (%)
1	1a + 2a	DMAP, H ₂ O	rt	3	3a	17
2	1a + 2a	DMAP, H ₂ O	60	3	3a	84
3	1a + 2a	H ₂ O	60	3	3a	>5
4	1a + 2a	DMAP	60	3	3a	24
5	1a + 2a	DMAP, EtOH	60	3	3a	10
6	1a + 2a	DMAP, MeOH	60	3	3a	12
7	1a + 2a	DMAP, toluene	60	3	3a	>5
8	1a + 2a	DMAP, MeCN	60	3	3a	12
9	1a + 2a	DMAP, THF	60	3	3a	>5
10	3a + DEAD	DMAP, H ₂ O	100	48	4a	81
11	3a + DEAD	DMAP, H ₂ O	80	48	4a	38
12	3a + DEAD	DMAP, H ₂ O	60	48	4a	22

^a 10 mol% DMAP was used in all reactions. ^b Isolated yields.



Table 2 Stepwise synthesis of various derivatives of 3 and 4

Entry	Conditions ^a	Reactants	Ar	Product	Yield ^{bb} (%)
1	DMAP, H ₂ O, 60 °C	1a + 2a	C ₆ H ₅	3a	84
2	DMAP, H ₂ O, 60 °C	1a + 2b	2-Naphthyl	3b	84
3	DMAP, H ₂ O, 60 °C	1a + 2c	4-MeC ₆ H ₄	3c	81
4	DMAP, H ₂ O, 60 °C	1a + 2d	4-MeOC ₆ H ₄	3d	87
5	DMAP, H ₂ O, 60 °C	1a + 2e	3-MeOC ₆ H ₄	3e	78
6	DMAP, H ₂ O, 60 °C	1a + 2f	4-FC ₆ H ₄	3f	82
7	DMAP, H ₂ O, 60 °C	1a + 2g	4-BrC ₆ H ₄	3g	80
8	DMAP, H ₂ O, 60 °C	1a + 2h	4-CF ₃ C ₆ H ₄	3h	81
9	DMAP, H ₂ O, 60 °C	1a + 2i	3-O ₂ NC ₆ H ₄	3i	80
10	DMAP, H ₂ O, 60 °C	1a + 2j	4-ClC ₆ H ₄	3j	83
11	DMAP, H ₂ O, 60 °C	1a + 2k	2,6-Cl ₂ C ₆ H ₃	3k	75
12	DMAP, H ₂ O, 60 °C	1a + 2l	2-Thienyl	3l	79
13	DMAP, H ₂ O, reflux	3a + DMAD	C ₆ H ₅	4a	85
14	DMAP, H ₂ O, reflux	3a + DEAD	C ₆ H ₅	4a'	83
15	DMAP, H ₂ O, reflux	3b + DEAD	2-Naphthyl	4b	78
16	DMAP, H ₂ O, reflux	3c + DEAD	4-MeC ₆ H ₄	4c	83
17	DMAP, H ₂ O, reflux	3d + DEAD	4-MeOC ₆ H ₄	4d	86
18	DMAP, H ₂ O, reflux	3e + DEAD	3-MeOC ₆ H ₄	4e	77
19	DMAP, H ₂ O, reflux	3f + DMAD	4-FC ₆ H ₄	4f	81
20	DMAP, H ₂ O, reflux	3g + DEAD	4-BrC ₆ H ₄	4g	78
21	DMAP, H ₂ O, reflux	3h + DEAD	4-CF ₃ C ₆ H ₄	4h	82
22	DMAP, H ₂ O, reflux	3i + DEAD	3-O ₂ NC ₆ H ₄	4i	80
23	DMAP, H ₂ O, reflux	3j + DEAD	4-ClC ₆ H ₄	4j	81
24	DMAP, H ₂ O, reflux	3k + DEAD	2,6-Cl ₂ C ₆ H ₃	4k	75
25	DMAP, H ₂ O, reflux	3l + DEAD	2-Thienyl	4l	80

^a 10 mol% DMAP was used in all reactions. ^b Isolated yields.

or electron withdrawing (entries 5–11) and heteroaromatic groups (entry 12) with 1a gave the respective aldol condensation products in high yields. Alternatively, isolation of 3a–l and their separate treatment with DEAD (entries 14–18 and 20–25) or

dimethyl acetylenedicarboxylate (DMAD) (entries 13 and 19) in the same aqueous DMAP mixtures yielded 4a–l efficiently.

We then evaluated the feasibility of conducting both steps in a one-pot procedure (Table 3). For this, we examined the

Table 3 One-pot synthesis of derivatives of 4



Entry	Reactants ^a	Ar	R	Product	Yield ^b (%)
1	1a + 2a + DMAD	C ₆ H ₅	Me	4a	82
2	1a + 2a + DEAD	C ₆ H ₅	Et	4a'	80
3	1a + 2b + DEAD	2-Naphthyl	Et	4b	80
4	1a + 2c + DEAD	4-MeC ₆ H ₄	Et	4c	77
5	1a + 2d + DEAD	4-MeOC ₆ H ₄	Et	4d	78
6	1a + 2e + DEAD	3-MeOC ₆ H ₄	Et	4e	80
7	1a + 2a + DMAD	4-FC ₆ H ₄	Me	4f	86
8	1a + 2f + DEAD	4-BrC ₆ H ₄	Et	4g	80
9	1a + 2g + DEAD	4-CF ₃ C ₆ H ₄	Et	4h	80
10	1a + 2a + DMAD	3-O ₂ NC ₆ H ₄	Et	4i	75
11	1a + 2a + DMAD	4-ClC ₆ H ₄	Et	4j	85
12	1a + 2a + DMAD	2,6-Cl ₂ C ₆ H ₃	Et	4k	81
13	1a + 2a + DMAD	2-Thienyl	Et	4l	80

^a 10 mol% DMAP was used in all reactions. ^b Isolated yields.



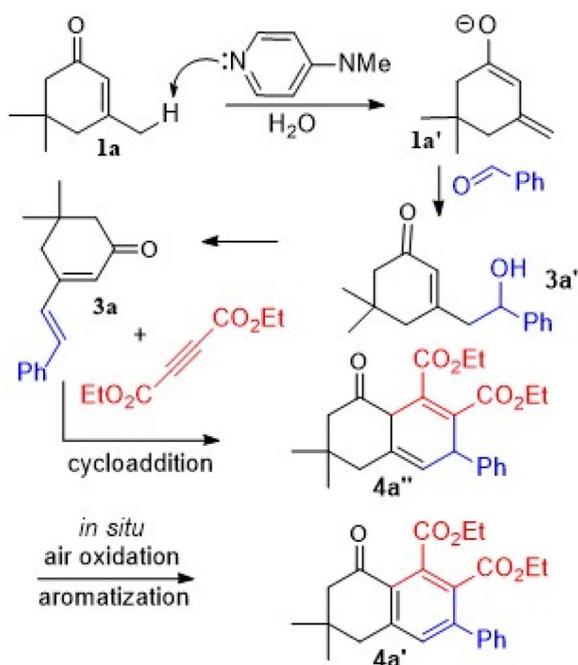


Fig. 3 The proposed mechanism.

conditions for the reaction of **1a** with benzaldehyde and DEAD in a water/DMAP mixture, where initial warming of the mixture of **1a** and **2a** to 60 °C (1 h) and delayed addition of the dienophile DMAD to the mixture and switching the conditions to refluxing temperature produced **4a** in 82% yield within 48 h (entry 1). Successful reactions of other aldehydes with either DMAD or DEAD illustrated the generality of the process by producing their respective adducts in high yields (entries 2–13).

Based on the results obtained here and in view of the basicity of DMAP as an organocatalyst in aqueous media,⁴³ a mechanism was proposed for the one-pot combination of the reactants, as shown in Fig. 3 for the reaction of benzaldehyde with **1a** and DEAD. Primarily, the DMAP preferably removed the γ acidic proton (Me group) to give the dienolate intermediate (**1a'**). In continuation, the intermediate attacked the aldehyde to complete the aldol step, giving **3a'** and then **3a**. Cycloaddition of **3a** with DEAD followed by *in situ* aromatization of **4a''** produced the final product **4a'**.⁴⁴

Experimental

General

All reagents were commercially available and used as received. Progress of the reactions was monitored by TLC using silica gel coated plates and EtOAc/petroleum ether mixture as the eluent. Melting points are uncorrected and obtained by Buchi Melting Point 530 apparatus. ¹H NMR (300 MHz or 600 MHz) and ¹³C NMR (75 MHz or 150 MHz) spectra are obtained on a FT-NMR Bruker Ultra Shield™ (or Bruker DRX-600) instrument as CDCl₃ solutions, and the chemical shifts are expressed as δ units with Me₄Si as the internal standard. Chemical ionization (CI) HRMS data are obtained (with MeOH as the ionization

source) using Agilent technologies 6530 Q-TOF-LC-MS. The identity of the known products was confirmed by the comparison of their ¹H NMR and ¹³C NMR spectra with those of authentic compounds reported in the literature.^{45,46} All new products were fully characterized based on their spectral data.

Typical synthesis of **3a**

A mixture of **1a** (300 μ L, 2.0 mmol), benzaldehyde **2a** (284 μ L, 2.0 mmol), and DMAP (25 mg, 10 mol%) in water (1.0 mL) was stirred at 60 °C for 3 h. After completion of the reaction (monitored by TLC using EtOAc/hexanes (1 : 4) as the eluent), the mixture was extracted with EtOAc (3 \times 5 mL), washed with brine, dried over anhydrous Na₂SO₄, and concentrated under reduced pressure. Product **3a** (380 mg, 84%) was obtained by column chromatography fractionation of the residue using (EtOAc/hexanes, 1 : 4, v/v).

Typical stepwise synthesis of **4a'**

A mixture of **3a** (1.0 mmol, 226 mg) and DEAD (255 mg, 1.5 mmol), and DMAP (13 mg, 10 mol%) in H₂O (2.0 mL) was refluxed for 48 h, until TLC showed completion of the process. The product was extracted from the reaction mixture with EtOAc (5.0 mL), washed with brine, dried over anhydrous Na₂SO₄, and concentrated under reduced pressure. Product **4a'** (327 mg, 83%) was obtained by column chromatography fractionation of the residue using (EtOAc/hexanes, 1 : 4, v/v).

Typical one-pot synthesis of **4a'**

A mixture of **1a** (150 μ L, 1.0 mmol), benzaldehyde **2a** (142 μ L, 1.0 mmol), and DMAP (13 mg, 10 mol%) in water (1.0 mL) was stirred at 60 °C for 3 h. At this point, DEAD (255 mg, 1.5 mmol) was added to the mixture and the mixture was refluxed for another 36 h. The product was extracted from the reaction mixture with EtOAc (5.0 mL), washed with brine, dried over anhydrous Na₂SO₄, and concentrated under reduced pressure. Product **4a'** (316 mg, 80%) was obtained by column chromatography fractionation of the residue using (EtOAc/hexanes, 1 : 4, v/v).

Spectral data of new products

(E)-5,5-dimethyl-3-(4-(trifluoromethyl)styryl)cyclohex-2-en-1-one 3h. Mp 103–105 °C; IR (KBr): 1324, 1586, 2161, 1655, 2933, 9961 cm⁻¹; ¹H NMR (300 MHz, CDCl₃) δ 1.12 (s, 6H), 2.33 (s, 2H), 2.49 (s, 2H), 6.12 (s, 1H), 6.99 (s, 2H), 7.59 (d, *J* = 8.0 Hz, 2H), 7.63 (d, *J* = 8.0 Hz, 2H); ¹³C NMR (75 MHz, CDCl₃) δ 28.4, 33.3, 38.9, 51.3, 125.7 (q, *J* = 28 Hz), 127.2, 128.2, 130.3, 130.7, 131.9, 133.0, 139.4, 153.7, 200.1; MS (70 eV) *m/z* (%), 294 (M⁺), 142, 170, 262, 277; anal. calcd for C₁₇H₁₇F₃O: C, 69.38; H, 5.82. Found: C, 69.20; H, 5.79.

(E)-3-(2,6-dichlorostyryl)-5,5-dimethylcyclohex-2-en-1-one 3k. Mp 69–71 °C; IR (KBr): 1587, 1616, 1657, 2930, 2947 cm⁻¹; ¹H NMR (300 MHz, CDCl₃) δ 1.12 (s, 6H), 2.32 (s, 2H), 2.50 (s, 2H), 6.06 (s, 1H), 6.94 (d, *J* = 16.5 Hz, 1H), 7.01 (d, *J* = 16.5 Hz, 1H), 7.13 (t, *J* = 8.0 Hz, 1H), 7.33 (d, *J* = 8.0 Hz, 2H); ¹³C NMR (75 MHz, CDCl₃) δ 28.4, 33.2, 38.5, 51.3, 128.2, 128.4, 128.6, 128.9,



133.3, 134.4, 137.7, 153.7, 200.1; MS (70 eV) m/z (%), 294 (M^+), 176, 204, 260; anal. calcd for $C_{16}H_{16}Cl_2O$: C, 65.10; H, 5.46. Found: C, 64.87; H, 5.37.

Dimethyl 6,6-dimethyl-8-oxo-3-phenyl-5,6,7,8-tetrahydronaphthalene-1,2-dicarboxylate (4a). Mp 151–153 °C; IR (KBr): 2949, 1739, 1694, 1588 cm^{-1} ; 1H NMR (600 MHz, $CDCl_3$): 1.03 (s, 6H), 2.58 (s, 2H), 2.93 (s, 2H), 3.62 (s, 3H), 3.97 (s, 3H), 7.22 (d, $J = 8.0$ Hz, 2H), 7.29 (s, 1H), 7.56 (d, $J = 8.0$ Hz, 2H); ^{13}C NMR (150 MHz, $CDCl_3$): 28.1, 33.7, 43.8, 52.5, 52.6, 53.0, 122.9, 128.5, 129.6, 129.7, 131.7, 132.0, 134.2, 138.2, 144.5, 145.4, 167.4, 169.0, 196.3; ESI-HRMS (MeOH) (m/z): calcd for $[C_{22}H_{22}O_5 + H]^+$: 367.1540, found: 367.1545.

Diethyl 7,7-dimethyl-5-oxo-5,6,7,8-tetrahydro-[2,2'-binaphthalene]-3,4-dicarboxylate (4b). Mp 124–125 °C; IR (KBr): 2969, 2255, 1726, 1241 cm^{-1} ; 1H NMR (600 MHz, $CDCl_3$): 0.50 (t, $J = 7.5$ Hz, 3H), 1.14 (s, 3H), 1.17 (s, 3H), 1.41 (t, $J = 7.5$ Hz, 3H), 2.62 (q, $J = 7.5$ Hz, 2H), 2.96 (q, $J = 7.5$ Hz, 2H), 2.93 (s, 2H) 4.46 (q, $J = 7.5$ Hz, 2H), 7.34 (dd, $J = 8.0$ Hz, 1H), 7.36 (s, 1H); 7.45 (dd, $J = 8.0, 8.0$ Hz, 1H), 7.50 (dd, $J = 8.0, 8.0$ Hz, 2H), 7.59 (d, $J = 8.0$ Hz, 1H), 7.89 (d, $J = 8.0$ Hz, 1H), 7.91 (d, $J = 8.0$ Hz, 1H); ^{13}C NMR (150 MHz, $CDCl_3$): 13.0, 13.9, 28.0, 28.3, 33.8, 43.8, 52.7, 61.1, 61.9, 125.0, 125.7, 126.1, 126.3, 126.4, 128.2, 128.4, 128.7, 131.3, 131.5, 133.2, 133.3, 134.4, 137.4, 144.9, 145.0, 166.4, 168.7196.4; ESI-HRMS (MeOH) (m/z): calcd for $[C_{28}H_{28}O_5 + H]^+$: 445.2009, found: 445.2010.

Diethyl 3-(4-fluorophenyl)-6,6-dimethyl-8-oxo-5,6,7,8-tetrahydronaphthalene-1,2-dicarboxylate (4f). Mp 93–95 °C; IR (KBr): 3062, 1729, 1233 cm^{-1} ; 1H NMR (600 MHz, $CDCl_3$): 1.02 (t, $J = 7.0$ Hz, 3H), 1.11 (s, 6H), 1.40 (t, $J = 7.0$ Hz, 3H), 2.58 (s, 2H), 2.93 (s, 2H), 4.07 (q, $J = 7.0$ Hz, 2H), 4.44 (q, $J = 7.0$ Hz, 2H), 7.11–7.14 (m, 2H), 7.28 (s, 1H), 7.33–7.35 (m, 2H); ^{13}C NMR (150 MHz, $CDCl_3$): 13.6, 13.9, 28.1, 33.8, 43.8, 52.6, 61.8, 62.0, 115.5 (d, $J = 21.0$ Hz), 128.3, 128.8 (d, $J = 9.0$ Hz), 130.3, 131.9, 134.2, 135.5, 144.8 (d, $J = 84.0$ Hz), 144.4 (d, $J = 247.5$ Hz), 145.1 (d, $J = 211.5$ Hz), 196.3; ESI-HRMS (MeOH) (m/z): calcd for $[C_{24}H_{25}FO_5 + H]^+$: 413.1759, found: 413.1759.

Diethyl 3-(4-bromophenyl)-6,6-dimethyl-8-oxo-5,6,7,8-tetrahydronaphthalene-1,2-dicarboxylate (4g). Mp 133–135 °C; IR (KBr): 2981, 1737, 1591 cm^{-1} ; 1H NMR (300 MHz, $CDCl_3$): 0.94 (t, $J = 7.0$ Hz, 3H), 1.04 (s, 6H), 1.31 (t, $J = 7.0$ Hz, 3H), 2.49 (s, 2H), 2.84 (s, 2H), 4.00 (q, $J = 7.0$ Hz, 2H), 4.35 (q, $J = 7.0$ Hz, 2H), 7.14 (d, $J = 7.5$ Hz, 2H), 7.18 (s, 1H), 7.45 (d, $J = 7.5$ Hz, 2H); ^{13}C NMR (75 MHz, $CDCl_3$): 13.6, 13.9, 28.1, 33.8, 43.8, 52.6, 61.8, 62.0, 122.7, 128.5, 129.7, 130.1, 131.6, 131.7, 134.3, 138.4, 144.4, 145.1, 166.9, 167.4, 167.1, 196.2; ESI-HRMS (MeOH) (m/z): calcd for $[C_{24}H_{25}BrO_5 + H]^+$: 473.0958, found: 473.0964.

Diethyl 6,6-dimethyl-8-oxo-3-(4-(trifluoromethyl)phenyl)-5,6,7,8-tetrahydronaphthalene-1,2-dicarboxylate (4h). IR (KBr): 2971, 1728, 1592 cm^{-1} ; 1H NMR (300 MHz, $CDCl_3$): 0.96 (t, $J = 7.0$ Hz, 3H), 1.12 (s, 6H), 1.40 (t, $J = 7.0$ Hz, 3H), 2.58 (s, 2H), 2.93 (s, 2H), 4.04 (q, $J = 7.0$ Hz, 2H), 4.44 (q, $J = 7.0$ Hz, 2H), 7.30 (s, 1H), 7.46 (d, $J = 7.5$ Hz, 2H), 7.69 (d, $J = 7.5$ Hz, 2H); ^{13}C NMR (75 MHz, $CDCl_3$): 13.2, 13.6, 27.8, 33.5, 43.6, 52.3, 61.5, 61.8, 122.0 (q, $J = 28.0$ Hz), 125.1, 128.2, 128.5, 129.8, 130.0, 130.4, 131.6, 134.2, 142.9, 143.8, 145.1, 167.1 (d, $J = 125.0$ Hz), 195.9; ESI-HRMS (MeOH) (m/z): calcd for $[C_{25}H_{25}F_3O_5 + H]^+$: 463.1727, found: 463.1732.

Diethyl 6,6-dimethyl-3-(3-nitrophenyl)-8-oxo-5,6,7,8-tetrahydronaphthalene-1,2-dicarboxylate (4i). Mp 140–141 °C; IR (KBr): 2949, 1738, 1587, 1237 cm^{-1} ; 1H NMR (300 MHz, $CDCl_3$): 1.03 (t, $J = 7.0$ Hz, 3H), 1.12 (s, 6H), 1.39 (t, $J = 7.0$ Hz, 3H), 2.58 (s, 2H), 2.94 (s, 2H), 4.08 (q, $J = 7.0$ Hz, 2H), 4.43 (q, $J = 7.0$ Hz, 2H), 7.32 (s, 1H), 7.60 (dd, $J = 7.5, 7.5$ Hz, 1H), 7.67 (dt, $J = 1.5, 7.5$ Hz, 1H), 8.20–8.30 (m, 2H); ^{13}C NMR (75 MHz, $CDCl_3$): 13.6, 13.9, 28.1, 33.8, 43.8, 52.5, 61.9, 62.1, 123.0, 123.1, 129.1, 129.4, 129.9, 132.0, 134.1, 134.6, 141.0, 142.9, 145.5, 148.1, 166.3, 168.1, 196.1; ESI-HRMS (MeOH) (m/z): calcd for $[C_{24}H_{25}NO_7 + H]^+$: 440.1704, found: 440.1709.

Diethyl 3-(2,6-dichlorophenyl)-6,6-dimethyl-8-oxo-5,6,7,8-tetrahydronaphthalene-1,2-dicarboxylate (4k). Mp 182–185 °C; IR (KBr): 2972, 1729, 1593 cm^{-1} ; 1H NMR (300 MHz, $CDCl_3$): 0.95 (t, $J = 7.0$ Hz, 3H), 1.13 (s, 6H), 1.41 (t, $J = 7.0$ Hz, 3H), 2.60 (s, 2H), 2.94 (s, 2H), 4.05 (q, $J = 7.0$ Hz, 2H), 4.45 (q, $J = 7.0$ Hz, 2H), 7.16 (s, 1H), 7.28 (dd, $J = 7.5, 7.5$ Hz, 1H), 7.39 (dd, $J = 1.5, 7.5$ Hz, 2H); ^{13}C NMR (75 MHz, $CDCl_3$): 13.4, 13.9, 28.1, 33.8, 43.8, 52.7, 61.4, 61.9, 127.1, 129.2, 129.4, 129.5, 132.6, 134.1, 135.4, 138.0, 141.5, 145.9, 165.0, 168.5, 196.4; ESI-HRMS (MeOH) (m/z): calcd for $[C_{24}H_{24}Cl_2O_5 + H]^+$: 463.1074, found: 463.1079.

Conclusions

In summary, we introduced a novel three-component method for the synthesis of a series of polysubstituted dihydronaphthalene derivatives. Both the diene formation and the cycloaddition steps were performed sequentially in the same pot using all three required reactants. The process was easy to operate by using an aqueous DMAP medium with no extra catalyst or additive needed. Additionally, a single product was obtained for each of the reactions in a good overall yield.

Author contributions

M. S. Abaee conceived and designed the work. Y. L. Nosood and E. Akbarzadeh performed the experiments and collected data. M. M. Mojtahedi reviewed the draft and performed the literature survey. A. Al-Harrasi conducted some of the analyses. All authors analyzed the data, discussed the results, and reviewed the manuscript.

Conflicts of interest

There are no conflicts to declare.

Data availability

The data supporting this article have been included as part of the SI. Supplementary information: Spectra of new compounds (1H NMR, ^{13}C NMR). See DOI: <https://doi.org/10.1039/d5ra04673d>.

Acknowledgements

We would like to thank CCERCI for financial support of this work (Grant number 1402-102).



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