

Halocyclization of *o*-(Alkynyl)styrenes. Synthesis of 3-Halo-1*H*-indenes

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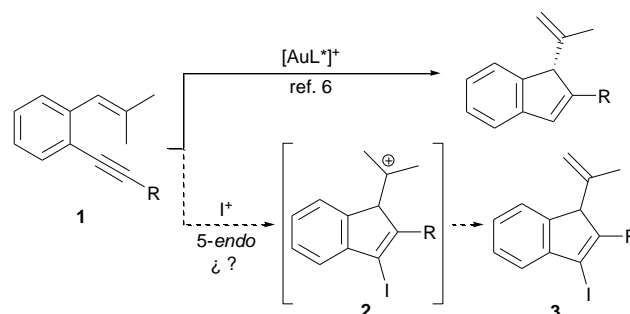
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o-(Alkynyl)styrenes undergo halocarbocyclization processes via a 5-*endo-dig* ring closure. By this strategy an efficient synthesis of 3-halo-1*H*-indene derivatives has been developed.

10 The electrophilic cyclization of heteroatomic nucleophiles with alkynes, mainly with iodine-containing electrophiles, is a useful method for the synthesis of functionalized heterocyclic compounds.¹ However, the analogous carbocyclization reactions, i.e. the internal nucleophile is a carbon-based moiety, have been much less developed. Few examples have been reported involving the use of arenes² and malonates³ as nucleophilic partners. Remarkably, the use of olefins⁴ for triggering this type of cyclization reactions has remained unknown until very recently, when Kirsch and co-workers have established that 1,5-enynes can undergo a iodonium-induced carbocyclization via 6-*endo-dig* processes.⁵ On the other hand, we have recently described the (enantioselective) synthesis of 1*H*-indene derivatives by an unprecedented gold(I)-catalyzed 5-*endo-dig* cyclization of *ortho*-(alkynyl)styrenes **1**.⁶ In this context, and considering that the presence of a halogen atom at the C-3 position of the final indene derivative could be of interest for further functionalization, we wondered about the feasibility of synthesizing 3-halo-1*H*-indenes⁷ from the same *o*-(alkynyl)styrenes **1** as proposed in Scheme 1.

Gratifyingly, we found that 3-iodoindene **3a** (R = Ph) was selectively formed and isolated in high yield when **1a** (R = Ph) was treated with an excess of NIS in CH₂Cl₂ at room temperature for 24 h, in the absence of any catalyst. This result shows that a direct halocyclization has occurred, probably through the formation of a stabilized carbocation **2** that undergoes proton elimination to afford the functionalized indene moiety (Scheme 1). It should be noted that in contrast to those examples recently reported by Kirsch and co-workers that proceed through a 6-*endo* cyclization process,⁵ our reaction implies an unprecedented 5-*endo* halocyclization reaction of an enyne derivative.⁸ The stability of the tertiary carbocation intermediate **2** could be the key for the success of the proposed 5-*endo* cyclization.

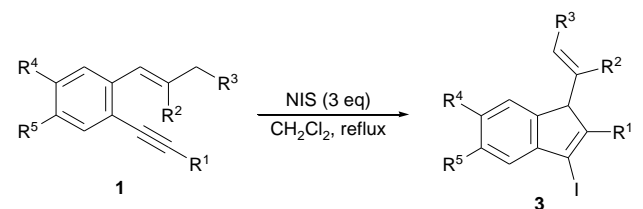
As shown in Table 1, a variety of *o*-(alkynyl)styrenes **1** are useful substrates for this iodocyclization. Reactions were performed at reflux in few hours under an air atmosphere. Different substituents at the alkyne (R¹), including functionalized-aromatic, heteroaromatic, (functionalized)-alkyl, and heteroatomic ones are well tolerated (Table 1, entries 1–6). Regarding the alkene, besides two methyl groups, R² and R³ could be part of a cyclic alkyl moiety (Table 1, entries 8 and 9) as well as two different groups such



Scheme 1 Our proposal for the synthesis of 3-iodoindenes

as methyl and phenyl (Table 1, entries 7 and 14). However, terminal alkynes (R¹ = H) proved to be unreactive under the reaction conditions, whereas substrates lacking a substituent at the β-position of the styrene moiety (R² = H) gave rise to a complex mixture of products.¹⁰ In addition, substrates bearing electron-withdrawing groups as well as electron-donating groups at the benzenoid moiety efficiently underwent the iodocyclization reaction (Table 1, entries 10–14).

Table 1 Synthesis of 3-iodo-1*H*-indenes **3** by iodocyclization of *o*-(alkynyl)styrenes **1**

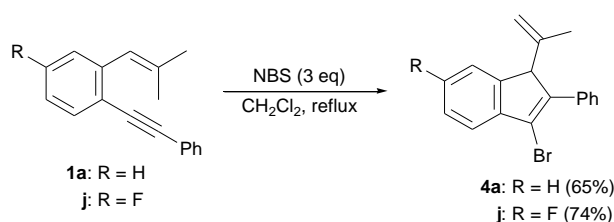


Entry	1	R ¹	R ²	R ³	R ⁴	R ⁵	t/h	3	Yield (%) ^a
1	1a	Ph	Me	H	H	H	3	3a	85
2	1b	4-MeOC ₆ H ₄	Me	H	H	H	1	3b	92(71) ^c
3	1c	3-Th ^b	Me	H	H	H	2.5	3c	87
4	1d	<i>n</i> Bu	Me	H	H	H	2	3d	81
5	1e	(CH ₂) ₃ CN	Me	H	H	H	15	3e	71
6	1f	SPh	Me	H	H	H	1	3f	80
7	1g	Ph	Ph	H	H	H	4.5	3g	81
8	1h	Ph	–(CH ₂) ₃ –	H	H	H	2	3h	71
9	1i	Ph	–(CH ₂) ₄ –	H	H	H	3	3i	88
10	1j	Ph	Me	H	F	H	15	3j	79
11	1k	Ph	Me	H	Br	H	7	3k	84
12	1l	Ph	Me	H	–OCH ₂ O–	H	1	3l	78(67) ^c
13	1m	<i>n</i> Bu	Me	H	Br	H	16	3m	82
14	1n	<i>n</i> Bu	Ph	H	Br	H	36	3n	55 ^d

^a Isolated yield after column chromatography referred to starting material **1**. ^b 3-Thienyl. ^c Carried out with I₂/K₃PO₄ (3 equiv) at rt. ^d ~80% conversion.

The possibility of using molecular iodine as the electrophilic reagent was also demonstrated by the isolation of iodoindenes **3b** and **3l**, from treatment of substrates **1b** and **1l** with I₂ and base instead of NIS (Table 1, entries 2 and 12). However, the yields were slightly lower and, moreover, in some cases, such as the reaction of **1a**, some side-products were formed along with the desired compound.

Interestingly, the same halocyclization reaction was observed when NBS was used. In this case, 3-bromo-1*H*-indenes **4a** and **4j** were isolated in good yields (Scheme 2).



Scheme 2 Bromonium-mediated synthesis of 3-bromo-1*H*-indenes **4**

We next decided to check the possibility of introducing further functionality on the C-1 substituent of the indene core by performing the iodocyclization in the presence of external oxygen nucleophiles. After some optimization, we found that although the competitive elimination reaction that affords iodoindenes **3** could not be completely suppressed, 1-alkoxy-substituted 3-iodo-1*H*-indene derivatives **5** could be isolated in useful yields and in pure form by using a large excess of MeOH (Table 2). In these cases, reactions are faster and could be performed at room temperature in 1–4 h. Better selectivities to **5** and therefore yields were obtained with substrates such as **1j** and **1k** bearing electron-withdrawing substituents at the aromatic nucleus (see entries 5 and 6 vs 1), and with **1g** (entry 4) that possess a phenyl group at the alkene. In addition, we have also demonstrated that I₂/K₃PO₄ could be used as iodonium source in the alkoxyiodocyclization reaction of **1a** (Table 2, entry 1), although with slightly lower selectivity.

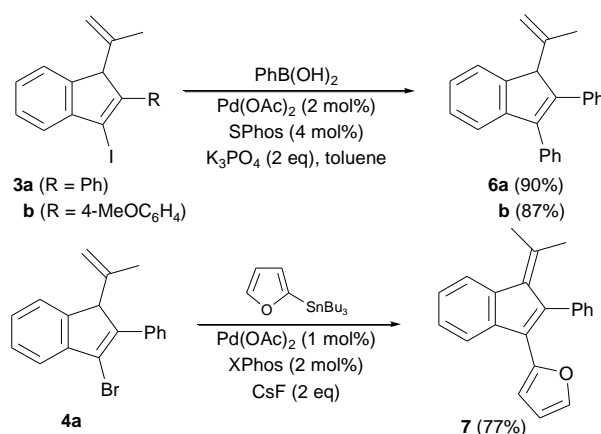
Table 2 Alkoxyiodocyclization of *o*-(alkynyl)styrenes **1**

Entry	1	R ¹	R ²	R ³	<i>t</i> /h	5	Yield (%) ^a
1	1a	Ph	H	H	2	5a	66(58) ^b
2	1b	4-MeOC ₆ H ₄	H	H	1	5b	70
3	1f	SPh	H	H	1	5f	58
4	1g^c	Ph	H	Ph	3	5g	81 ^d
5	1j	Ph	F	H	2	5j	78
6	1k	Ph	Br	H	4	5k	72

^a Isolated yield after column chromatography referred to starting material **1**. ^b Carried out with I₂/K₃PO₄ as electrophilic source. ^c Used as a ca. 2:1 mixture of geometrical isomers. ^d Obtained as a ca. 2:1 mixture of diastereoisomers.

Having in mind that 2,3-diarylindenes are known to possess biological activity,¹¹ we decided to employ the

synthesized 3-haloindenes as precursors for the introduction of an aromatic substituent at the C-3 of the indene moiety by Pd-catalyzed cross coupling reactions. In this way, 2,3-diarylindene derivatives **6** were obtained in high yield from **3a** and **3b** through a Suzuki coupling¹² with phenylboronic acid and Pd(OAc)₂/SPhos as catalytic system (Scheme 3). Interestingly, when bromoindene **4a** underwent a Stille coupling¹³ with 2-(tributylstannyl)furan, the functionalized benzofulvene¹⁴ derivative **7** was obtained, in which further isomerization of the propenyl group at the C-1 position of the indene scaffold has occurred (Scheme 3).



Scheme 3 Synthetic applications of 3-haloindenes **3** and **4**

In summary, we have reported the direct halocyclization of *o*-(alkynyl)styrenes to give 3-halo-1*H*-indene derivatives in good to high yields under mild conditions. These results together with those recently reported by Kirsch and coworkers represent the first examples of electrophilic cyclizations of alkynes promoted by halonium ions where the nucleophilic counterpart is an alkene. In particular, our results represent the first examples of this type of halocyclization via a 5-*endo-dig* ring closure mechanism. The presence of a halogen in the final products allows further functionalization at the C-3 position of the indene through conventional palladium-catalyzed cross-coupling reactions. Current efforts in our lab are devoted to the development of an enantioselective version of this reaction and further applications of the obtained haloindenes in the synthesis of complex molecules.

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Notes and references

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† Electronic Supplementary Information (ESI) available: Experimental procedures, characterization data, and NMR spectra. See DOI: 10.1039/b000000x/

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