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Chemoselective homologative preparation of trisubstituted alkenyl halides from carbonyls and carbenoids†

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The chemoselective synthesis of trisubstituted alkenyl halides (Cl, Br, F, I) starting from ketones and aldehydes and lithium halocarbenoids is reported. Upon forming the corresponding tetrahedral intermediate adduct, followed by the addition of thionyl chloride, a selective E2-type elimination is triggered, furnishing the targeted motifs. The transformation takes place under full chemocontrol: various sensitive functionalities (e.g. ester, nitrile, nitro, or halogen groups) can be placed on the starting materials, thus documenting a wide reaction scope, as well as the application of the technique to biologically active substances.

The interest in the alkenyl halide moiety encompasses distinct areas of organic synthesis; in fact, it represents a versatile motif well suited for undergoing further elaboration through conceptually different regimes.¹ As a consequence of the unique electronic environment imparted by the halogen, the material can be advantageously employed in C–C or C–heteroatom bond formation (either *via* polar metalation or transition-metal-catalyzed sequences – Scheme 1).² Moreover, it is expressed in some natural products (mainly in the form of vinyl-type chlorides) and can also be present in medicinally relevant structures or agrochemicals.³

The desired substitution pattern—and consequently the stereochemistry—of the alkenyl cluster dictate the rationale for selecting the most suitable preparative route. Accordingly, the hydrohalogenation of alkynes offers a valuable strategy, which is also tuneable in both *syn*- and *anti*-Markovnikov-type additions when organic halides replace pure mineral acids (HX – Scheme 1, path 1a).⁴ Although various transition-metal-catalyzed approaches have

been developed, procedures leading to alkenyl fragments featuring the four halogens have rarely been described.⁵ Nevertheless, previously considered elusive internal alkynes can nowadays be used as competent starting materials.^{5c,6} More traditional methods are based on the conversion of carbonyl groups into vinyl halides, well illustrated by the so-called Barton synthesis, which unfortunately shows high substrate sensitivity and may require the preparation of capricious intermediate species (e.g. hydrazones – Scheme 1, path 1b left).⁷ In this context, although Prati's modification addresses these difficulties, it still remains challenging for common materials such as benzophenones.⁸ Alternatively, vinylic-type nucleophilic substitutions conducted on functionalized alkenes in transition-metal-catalyzed reactions have been also reported (Scheme 1, path 1b right).⁹ From a specular perspective, the same carbonyl moiety constitutes a placeholder for a haloalkene through a formal homologative event. Wittig-like methodologies with α -halosubstituted phosphorous ylides can furnish either *E*- or *Z*-olefins,¹⁰ but due to the highly basic conditions required, undesirable elimination to alkynes or reduction to methylene units could affect the chemoselectivity (Scheme 1, path 2a).¹¹ In general, gaining remarkable control during these processes is somewhat counterbalanced by the restriction of the protocol to more electrophilic carbonyls (aldehydes). In this sense, the venerable Takai haloolefination furnishes disubstituted alkenes starting from aldehydes and a haloform in the presence of problematic Cr(II) salts (Scheme 1, path 2b).¹² However, engaging ketones proved to be more challenging and usually needed the employment of bimetallic species (e.g. Mg–TiCl₄) to generate an adequately reactive nucleophile (Scheme 1, path 2c).¹³ Significantly, the inconvenient manipulability of fluoroform (bp –82 °C)¹⁴ precluded the adoption of these protocols for the preparation of alkenyl fluorides.

Cognizant of the excellent reactivity of lithiated halo-methanes towards carbonyls (ketones and aldehydes) generating tetrahedral nucleophilic addition intermediates,¹⁵ we

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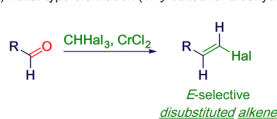
Alkenyl halides in synthesis

- Versatile motifs in C-C and C-Heteroatom bond formations
- Modifiable for polar and/or TM-catalytic operations
- Valuable motif in med chem and natural product chemistry

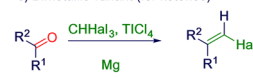
a) Hydrohalogenation of alkynes (tuneable stereochemical control)



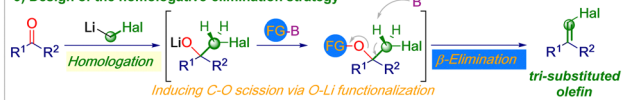
a) Wittig-like halo-olefinations



c) Bimetallic variant (for ketones)



3) Design of the homologative-elimination strategy



Scheme 1 General context of the presented work.

4-Bromoacetophenone (**1**) was selected as a suitable model compound susceptible to base-mediated enolate formation in the reaction with the basic carbenoid LiCH_2Cl (Table 1). The constitutive presence of the exchangeable bromine atom would permit assessment of the chemocontrol over the transformation in the presence of the lithiated carbanion-type species. Based on our previous studies on homologative/deoxygenative sequences,¹⁶ 1.4 equiv. of carbenoid-generated from 1.5 equiv. of ClCH_2I and 1.4 equiv. of MeLi-LiBr at -78°C in THF-we could verify the complete conversion (upon acidic quenching) of the ketone into chlorohydrin **1b** (90% yield). With this confirmation in hand, we studied the subsequent E2-type elimination run on the lithiated chlorohydrin **1a** triggered by an external agent. Thus, by slowly adding SOCl_2 (1.5 equiv.) at -78°C and stirring for 10 min followed by 30 min at ambient temperature, the desired *E*-chloroalkene (**2**) was obtained in 86% yield and *E/Z* ratio $> 99:1$, as deduced from NOESY

Reaction scheme for the synthesis of **2** from **1**:

Starting material **1** (a ketone with a methyl group and a substituent R^1) reacts with $ClCH_2I$ (1.5 equiv), $MeLi-LiBr$ (1.4 equiv), THF, $-78^\circ C$, 0.5 h, and $[R^1 = 4-BrC_6H_4]$ to form intermediate **1a** (a lithium enolate).

Intermediate **1a** reacts with SO_2 to form intermediate **1c** (a sulfonate salt).

Intermediate **1c** reacts with H_3O^+ to form intermediate **1b** (a sulfonic acid, 90% yield).

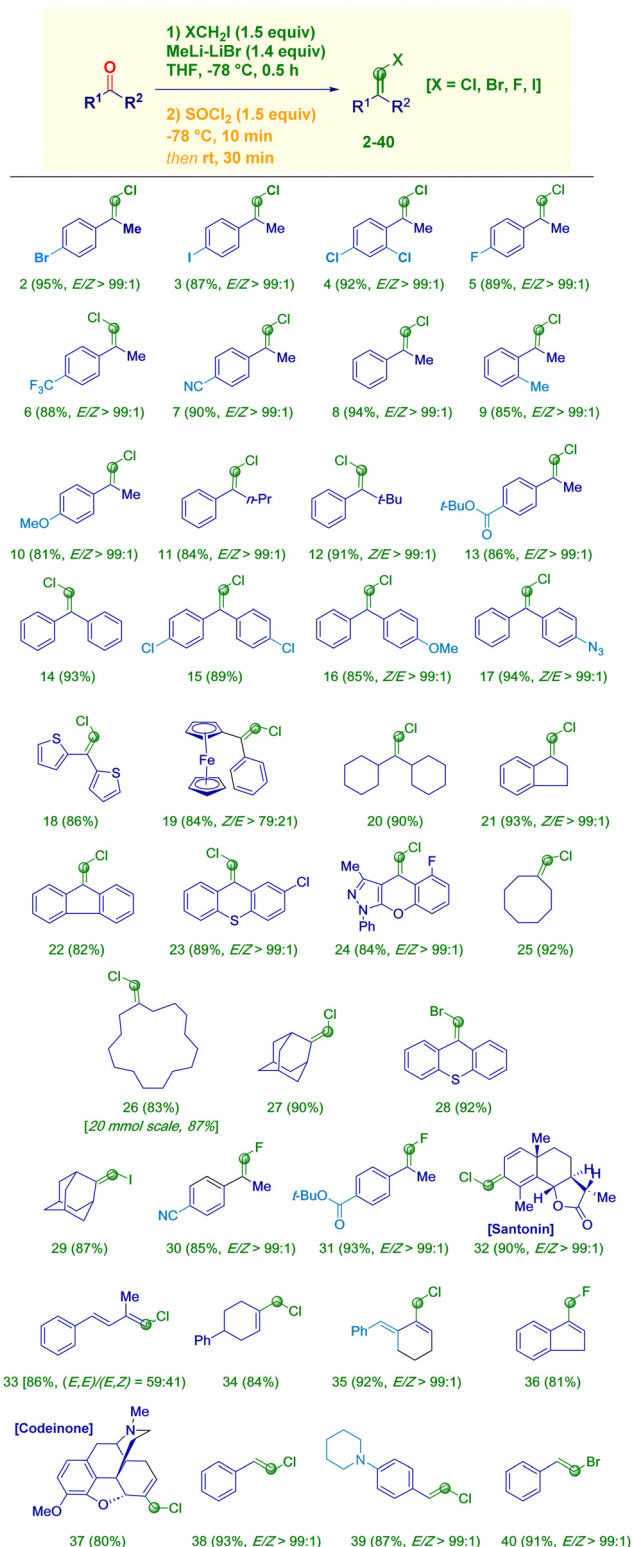
Intermediate **1b** is then converted to the final product **2** (a ketone with a methyl group and a substituent R^1), with SO_2 (gas) as a byproduct.

Entry	Eliminating agent (equiv.)	Temperature [°C]	Reaction time ^a (h)	Yield of 2 ^b (%)	E/Z ratio of 2
1	SOCl ₂ (1.5)	−78 to rt	0.5	95	>99:1
2	SOCl ₂ (1.5)	0 to rt	0.5	67	93:7
3 ^c	SOCl ₂ (1.5)	−78	2	—	—
4 ^d	SOCl ₂ (1.5)	−78 to rt	0.5	75	98:2
5 ^e	SOCl ₂ (1.5)	−78 to rt	0.5	58	98:2
6	SOBr ₂ (1.5)	−78 to rt	0.5	82	>99:1
7	(COCl) ₂ (1.5)	−78 to rt	0.5	63	98:2
8	POCl ₃ (1.5)	−78 to rt	0.5	54	99:1
9	POBr ₃ (1.5)	−78 to rt	0.5	62	99:1
10	PCl ₃ (1.5)	−78 to rt	0.5	48	>99:1

^a Reaction time refers to the stirring of the mixture after removing the cooling bath. ^b Isolated yield. ^c See Scheme 3 for isolation of chlorosulfite intermediates. ^d DIPEA (1.3 equiv.) was added after concluding the addition of SOCl₂. ^e Pyridine (1.3 equiv.) was added after concluding the addition of SOCl₂.

experiments (entry 1). Some points merit mention: (a) Adding SOCl_2 to the lithiated intermediate cooled at 0°C had a detrimental effect, since the formation of impurities was observed in the $^1\text{H-NMR}$ of the reaction crude and the E/Z ratio sensitively diminished (entry 2). (b) Continuing stirring at -78°C did not induce any elimination and the corresponding chlorosulfite intermediates could be isolated (entry 3 - *vide infra*). (c) The addition of bases such as DIPEA or pyridine, although it enabled the formation of the alkene, was not comparable to the process carried out with SOCl_2 alone, suggesting the chloride ion (released during the formation of the chlorosulfite ester) to be the active base for boosting elimination on intermediate **1c** (entries 4 and 5). (d) Replacing SOCl_2 with SOBr_2 reduced the chemical yield (entry 6), as did the use of analogous $(\text{COCl})_2$ and phosphorous-based electrophiles (POCl_3 , POBr_3 , PCl_3 , entries 7–10). Plausibly, the release of gaseous SO_2 at the end of the eliminative sequence accounts for the high efficiency of the transformation.

Having demonstrated that SOCl_2 induces an E2-type elimination conducted on the nucleophilic addition intermediates (alkoxides)–without furnishing any ($\text{OH} \rightarrow \text{Cl}$) substitution products¹⁷–we then studied the scope of the method (Scheme 2). Various aryl-substituted acetophenones were all amenable substrates, giving the corresponding trisubstituted chloro-alkenes in very high chemical yield and *E*-selectivity. Not only could the whole series of halogens (2–5)–including the trifluoromethyl analogue (6)–be conveniently placed on the aromatic ring but substrates decorated with nitrile (7), ether (10) or ester (13) functionalities also worked equally well, furnishing in all cases halo-olefins in *E/Z* ratio > 99 : 1. The simple acetophenone (8) and the more sterically hindered *o*-methyl analogues (9) underwent the transformation with comparable efficiency. While using the (linear) propiophenone



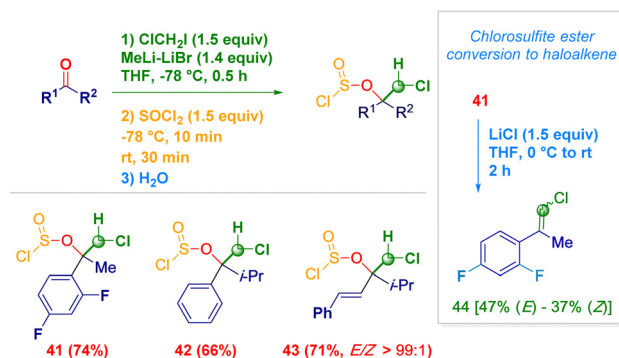
Scheme 2 Scope of the sequential carbonyl homologation/elimination en route to haloalkenes.

gave compound **11** in E/Z ratio > 99:1, switching to the highly sterically demanding *t*-butyl-phenyl ketone (**12**) reversed the ratio in favour of the *Z*-isomer. The employment of di(hetero)arylketones [benzophenones (**14–17**) the bis-thienyl-

(**18**), the ferrocenyl-(**19**) and the bis-cyclohexyl (**20**) analogues] guaranteed access to (mainly) (*Z*)-halo-alkenes in comparable yields. Extending the protocol to bi- and tri-cyclic systems was possible, as observed in the cases of 1-indanone (**21**), 9*H*-fluoren-9-one (**22**), 9*H*-thioxanthene-9-one (**23**) and (the more elaborate) fluoro-substituted chromeno[2,3-*c*]pyrazol-4(1*H*)-one (**24**). Pure carbocyclic analogues, including octan-1-one (**25**) and the expanded 15-membered ketone (**26**) gave the trisubstituted alkenes in high yield (also when scaling up to 20 mmol), as well as, the sterically hindered adamant-1-one (**27**). The chemo-selective methodology was not restricted to the use of LiCH₂Cl as the C1-donor but was also applicable to the preparation of bromo-(**28**) iodo-(**29**) and fluoro-(**30–31**) olefins with LiCH₂Br,¹⁸ LiCH₂I¹⁹ and LiCH₂F,²⁰ respectively, showing remarkable synthetic versatility. Conjugated haloalkenes could be easily prepared in the cases of both cyclic (**32**) and acyclic (**33**) materials. It is interesting to highlight the success of the transformation conducted on the anthelmintic natural product Santonin (**32**) whose constitutive lactone moieties were inert under the reaction conditions. As a consequence of the higher thermodynamic stability of endocyclic double bonds (compared to exocyclic ones),²¹ vinyl-allyl halide isomerization could take place, yielding structures **34–37** during the usual work-up. Again, the application of the method to the narcotic drug codeinone (**37**) further illustrates the significance of the protocol for the elaboration of medicinally relevant substances. Finally, switching to aromatic aldehydes as electrophilic partners for carbenoids—*coeteris paribus*—provided a smooth route to β-haloalkenes (**38–40**), validating the protocol for obtaining (*E*)-haloalkenes.²²

As mentioned in the optimization study (Table 1), by keeping the temperature at –78 °C after the addition of SOCl₂, it was possible to unambiguously demonstrate the genesis of chlorosulfite esters as the pertinent intermediates (Scheme 3). Nevertheless, subsequent treatment with a solution of LiCl in THF at room temperature yielded the corresponding halo-alkene **44**, albeit with poorer stereocontrol (presumably due to conducting the reaction at high temperature).

In summary, we have reported an effective synthesis for trisubstituted halo-alkenes from ketones through a sequential homologation with lithium halocarbenoids (LiCH₂Cl, LiCH₂Br,



Scheme 3 Trapping chlorosulfite esters for mechanistic analysis.



LiCH₂I and LiCH₂F-generated from dihalomethanes and MeLi–LiBr under Barbier-type conditions at –78 °C), followed by an E2-type elimination triggered on the tetrahedral intermediate with thionyl chloride. The protocol exhibits remarkable chemocontrol, as indicated by reacting ketones presenting a wide range of chemical functionalities (e.g. halogen, azido, nitrile, ferrocenyl, ether, ester, lactone, conjugated olefin, or amine), which, in principle, may interfere with the carbenoids during the homologative event. Not only are simple alkyl-aryl ketones amenable for the process, but (hetero)aryl-(hetero)aryl, (alkyl)-(alkyl) including macrocycles and a series of biologically active substrates featuring the ketone group could also be equally employed. The application of the method to aromatic aldehydes furnishes β-halostyrenes, whereas aliphatic analogues are currently under investigation and the results will be reported in due course.

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Data availability

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

Conflicts of interest

There are no conflicts to declare.

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