

A Facile and Efficient Method for the Debromination of *vic*-Dibromides to Alkenes with BiCl₃/Indium System

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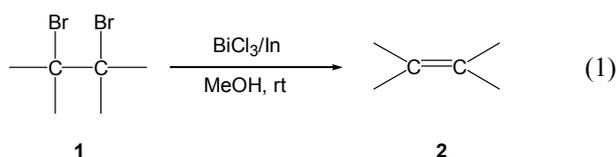
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The *vic*-dibromide-alkene functional group interconversion represents a useful step in the double bond protection-deprotection strategy.¹ To date, several protocols have been utilized for the debromination of *vic*-dibromides to alkenes, but some of these methods are associated with limitations regarding chemo-selectivity, low yields, prolonged reaction times, and harsh reaction conditions.² Consequently an efficient and mild procedure for debromination of *vic*-dibromides continues to be developed. The chemical reactivity of BiCl₃/M system (M = Zn, Fe, Al, Mg) has been the subject of considerable interest and the reducing ability of these systems has been extensively studied.³ Because of the close resemblance of indium to zinc in several respects, including first ionization, we reasoned that BiCl₃/indium system could serve as a protocol for the debromination of *vic*-dibromides. In recent years indium metal has drawn an increasing attention for its unique properties such as low toxicity and high stability in water and air compared with other metals.⁴ Bismuth(III) chloride is inexpensive, relatively non-toxic, fairly water insensitive, and environmentally benign reagent, which have been used as mild Lewis acid catalyst for synthetic transformations. In continuation to our interest in exploring the utility of metal-metal salt system in organic synthesis,⁵ we wish to report here an efficient and mild method for debromination of *vic*-dibromides **1** to alkenes **2** with BiCl₃/indium system in methanol at room temperature. The reaction can be generalized as in eq. 1.⁶ The new reagent system was generated by the addition of indium powder to a stirred solution of bismuth chloride in methanol under sonication.⁷ To gauge the scope and limitations of this methodology, various substituted *vic*-dibromides were subjected to this protocol at room temperature. We have found that *vic*-dibromides on treatment with BiCl₃/indium system in methanol solution are smoothly converted into the corresponding alkenes in high yields.



The high yields of the debromination products demonstrate the efficiency of this new method. To ensure the role of indium, a controlled experiment was carried out using *vic*-dibromides

with BiCl₃ without indium metal, which failed to yield any desired product. A 2 : 1 ratio of indium and BiCl₃ was the best ratio in terms of yield and reaction time. And methanol has been found to be the most suitable solvent for the reaction.

Table 1. Reductive debromination of *vic*-dibromides with BiCl₃/indium system

entry	substrate	product	time (hr)	yield (%) ^a
1			0.5	95
2			0.5	91
3			0.5	94
4			1.5	92
5			1.5	89
6			1.0	86
7			2.0	86
8			0.5	94
9			1.0	90
10			2.0	84
11			1.0	89

^aIsolated yields. The products are commercially available and were characterized by comparison of their spectral data with authentic samples.

The result of the reaction is summarized in Table 1, where we examined a series of functionally and structurally various *vic*-dibromides. Clearly a broad range of functional group (ester, carboxyl, aldehyde, methoxy, chloro, and ketone) was tolerated under the reaction conditions and only *trans* olefins were obtained. It was worth commenting that the sensitive carbonyl group remains intact without any further reduction under the reaction condition (entries 5-10). In comparison with other procedures, the present procedure reduced *vic*-dibromides in higher yields and showed a good chemoselectivity under mild conditions. The notable advantages of this methodology are mild reaction conditions, simple manipulation, high yields, and tolerance of various functional groups. No over-reduction of the produced alkene was observed with any substrate.⁸ Some of the *vic*-dibromides in the table are commercially available, or could be prepared conveniently from the corresponding precursors through known olefin halogenation reactions.⁹ Although the reaction mechanism is still not clearly understood at this time, the reaction can be envisaged to proceed in two stages. In the first bismuth(III) chloride is probably reduced by indium to form low-valent bismuth species, which in the subsequent step would debrominate *vic*-dibromides **1** to give the corresponding alkenes **2** through a SET (single electron transfer) process. The reducing property exhibited by metal-metal salt combinations proceeds through transfer of one electron from the metal surface to the substrate. In such combinations elementary metal part needs to be more electropositive than the metal part of the salt. The involvement of zero valent bismuth resulting from the transmetallation of bismuth chloride with indium metal may be presumed.¹⁰ We have been able to demonstrate the utility of BiCl₃/indium system for effecting chemoselective debromination of *vic*-dibromides.

In conclusion, we have demonstrated that BiCl₃/indium system mediates an efficient and mild conversion of *vic*-dibromides to the corresponding alkenes. Although the scope and limitations were not fully established, the present method could be a practical alternative to the conventional method. Further work on the application of BiCl₃/indium system is in progress.

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6. A typical procedure for the debromination of *vic*-dibromides is as follows: Indium powder (230 mg, 2.0 mmol) and bismuth(III) chloride (315 mg, 1.0 mmol) were mixed in methanol (4 mL). The resulting mixture was stirred at room temperature for 0.5 hr under sonication producing a solution of the low-valent bismuth-indium complex. 1,2-Dibromoethylbenzene (264 mg, 1.0 mmol) was then added to this solution and the reaction mixture was stirred for 30 min at room temperature. The solvent was removed under reduced pressure and the residue was extracted with ether, washed with brine, and dried over anhydrous Na₂SO₄. The crude product was purified by silica gel column chromatography (hexane:ethyl acetate = 10:1) to afford styrene (99 mg, 95%). All of the products are commercially available compounds, and their identification was based on spectral comparison with authentic samples.
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