The combination of zinc bromide with tosyl chloride promoted sulfonylation, allowing temperature-controlled regioselective sulfonylation of carbohydrates which possess both primary and secondary hydroxy groups.

A sulfonate ester is one of the most useful intermediates in organic chemistry and sulfonylation of alcohols by use of tosyl chloride is one of the most prevalent reactions. It is possible to selectively tosylate primary over secondary alcohols in pyridine at 0 °C. It is also possible to tosylate nonprotected hydroxy functionalities regioselectively by use of organotin compounds. However, application of these methods to relatively large molecules possessing a number of primary and secondary hydroxy groups is not always straightforward, as discrimination of the hydroxy groups is difficult. In the case of the preparation of hepta-6-O-sulfonylated β-cyclodextrin which is considered a suitable intermediate for hepta-6-functionalized derivatives, a significant amount of the 2-O-sulfonylated derivative was also produced. In the synthetic study to improve the yield of 1, we found a unique role of zinc bromide to promote sulfonylation and here we will describe the results.

Steric crowding on the primary side of a cyclodextrin during the later stages of sulfonylation to prepare the derivative may enable the reaction of the less reactive secondary hydroxy groups before the completion of 6-O-sulfonylation. The effect of metal halides as additives was studied because either selective activation or protection of hydroxy groups by coordination of metal ion was expected. The results are shown in Figure 1. Metal ions such as Mg2+ and Ca2+ suppressed the sulfonylation reaction. In contrast, zinc halides facilitated the reaction. Among them ZnBr2 showed the highest acceleration effect, although the ratio of the tosylate and the additionally 2-O-sulfonylated derivative was the same as that observed on a reaction without the additive. The lack of selectivity suggests that ZnBr2 promotes tosylation on both primary and secondary hydroxy groups. A possible role of ZnBr2 is considered to be coordination to an oxygen atom of the sulfonyl moiety making the halogen a better leaving group. Coordination of ZnBr2 to tosyl chloride is suggested by the fact that the solubility of a mixture of ZnBr2 and tosyl chloride, in a molar ratio of 1:1.5, in pyridine is greater than the metal halide itself. Use of a mixture of zinc bromide and tosyl chloride allows the reaction to proceed more efficiently.
of cycloextrin (1.5 times the concentration of the glucose residues) reduced the reaction time by a factor of twelve. While ZnBr$_2$ promoted the sulfonylation at both primary and secondary hydroxy groups, it was found that low reaction temperature effectively suppressed the undesirable 2-O-sulfonylation. On lowering the reaction temperature to below –20 °C the yield of the desired product 1 increased to more than 70% in the absence of ZnBr$_2$. The HPLC chromatogram was much simpler with fewer peaks. However, the reaction required 25 mole equivalents of tosyl chloride and took more than 30 h to obtain the maximal yield of 1. Use of ZnBr$_2$ enabled reaction to be carried out at lower reaction temperature, in a shorter time, and with less sulfonyl chloride, namely at –40 °C, 6 for 4 h, and in the presence of ZnBr$_2$ and tosyl chloride, ten and fifteen mole equivalents, respectively, to give the desired product 1 in 84% yield. Through the use of twenty mole equivalents of tosyl chloride at –40 °C the tosylate 1 (83%) was obtained within 1 h.

On tosylation of methyl α-D-glucoside in pyridine, similar effects of ZnBr$_2$ were demonstrated. The sulfonylation at r.t. in the absence of ZnBr$_2$, using ten mole equivalents of tosyl chloride took 30 min to consume the glucoside affording mono- and ditosylate (57 and 43%, respectively). Reaction at –20 °C using both ZnBr$_2$ and tosyl chloride (two and three mole equivalents, respectively) converted the glucoside in 20 min to the mono- and disulfonates in the yield of 64 and 36 %, respectively. The latter reaction needed less tosyl chloride to consume the glucoside. Taking into account differences between the two conditions above in reaction temperature, amount of sulfonyl chloride, and reaction time, addition of ZnBr$_2$ increased the efficiency of conversion of the glucoside to sulfonates by a factor of eighty. Lowering the reaction temperature from r.t. to –20 °C resulted in preferential sulfonylation on primary hydroxy group, as observed in the case of cyclodextrin, although the effect of temperature was smaller than that of cyclodextrin. It is probable that macrocyclic structure of the cyclodextrin makes its secondary hydroxy groups more crowded than that of the glucoside, leading to a greater difference in reactivity between the primary and secondary hydroxy groups.

In conclusion, zinc bromide promoted tosylation such that less sulfonyl chloride was required. Further, the additive enabled low temperature reactions, which gave selective reaction on primary hydroxy groups, to be carried out efficiently. Although the use of zinc halides as Lewis acids is known, this is the first example of the influence of zinc bromide on the reactivity of sulfonyl chloride with hydroxy groups, to the best of our knowledge, in the literature. The results suggest the possibility of very rapid ("instant") sulfonylation and the use of much lower temperatures for sulfonylation in an effort to increase specificity. Further studies including the application of this process to other alcohols including carbohydrates and cyclitols such as mannitol, are underway in our laboratory.

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References and Notes
4 The reaction using ZnI$_2$ gave a mixture of tosylates and products with lower UV absorbance whose sulfonyl groups were presumably replaced by iodides. 5
6 Reaction mixture in pyridine froze at –50 °C.