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Borate esters as convenient reagents for direct amidation of carboxylic acids and transamidation of primary amides†

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Simple borates serve as effective promoters for amide bond formation with a variety of carboxylic acids and amines. With trimethyl or tris(2,2,2-trifluoroethyl) borate, amides are obtained in good to excellent yield and high purity after a simple work-up procedure. Tris(2,2,2-trifluoroethyl) borate can also be used for the straightforward conversion of primary amides to secondary amides *via* transamidation.

The amide linkage¹ is prevalent in nature² and is extensively used in synthetic chemistry.³ A vast number of approaches for amide bond formation exist,^{3–7} many of which are problematic with regard to cost and atom-efficiency,⁸ as well as recyclability and functional group tolerance.⁹ Recently, boric acid and arylboronic acids were shown to be able to mediate direct amide coupling of carboxylic acids and amines.¹⁰ Although they can be used in catalytic amounts (1–20 mol%), effective removal of water is essential, either by heating under azeotropic reflux in high-boiling point solvents (*e.g.*, PhMe, xylene),^{10b–f} or by conducting the reaction at lower temperatures (rt–50 °C), in the presence of molecular sieves.^{10g} However, high dilution conditions and prolonged reaction times (24–48 h) are necessary in this latter case. The use of stoichiometric boron reagents for amidation has also been reported (*e.g.* BF₃·OEt₂,^{11a} catecholborane,^{11b} BH₃·NMe₃,^{12a} and BH₃·THF^{12b}), but such reagents require strictly anhydrous reaction conditions and, in some cases, an excess of either the carboxylic acid or amine in order to obtain a good conversion.¹² Herein, we demonstrate the application of simple borate esters to direct carboxamidation under convenient reaction conditions. Additionally, we show that tris(2,2,2-trifluoroethyl) borate can be used to activate amides towards transamidation.¹³ These reactions show good functional group tolerance and do not require anhydrous reaction conditions.

During the course of our work on the development of new boron reagents for organic synthesis,¹⁴ we examined a range of compounds as mediators for amidation reactions, and observed that B(OMe)₃¹⁵ can act as an effective reagent for direct carboxamidation. Activation of the carboxylic acid presumably occurs

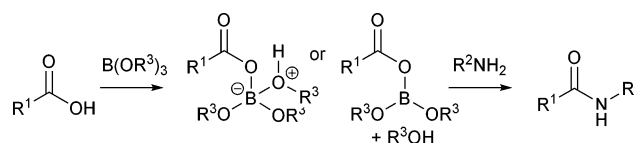
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† Electronic supplementary information (ESI) available: Experimental procedures, spectral data and copies of ¹H and ¹³C NMR spectra for all compounds. See DOI: 10.1039/c0ob01069c

Table 1 Optimisation of Borate-Promoted Direct Carboxamidation under Microwave Conditions^a

Entry	Reagent	Solvent	Conversion (%) ^b
1	None	MeCN	2
2	B(OMe) ₃	MeOH	0
3	B(OMe) ₃	MTBE	4
4	B(OMe) ₃	PhMe	12
5	B(OMe) ₃	THF	20
6	B(OMe) ₃	DMSO	27
7	B(OMe) ₃	B(OMe) ₃ ^c	30
8	B(OMe)₃	MeCN	35 (35)^e
9	B(OMe) ₃	MeCN	6 ^d
10	B(OMe) ₃	MeCN	14 ^e
11	B(OMe) ₃	MeCN	19 ^f
12	B(O ⁱ Pr) ₃	MeCN	9
13	B(OSiMe ₃) ₃	MeCN	9
14	Si(OMe) ₄	MeCN	22
15	B(OCH₂CF₃)₃	MeCN	63 (63)^e

via in situ generation of a three or four-coordinate boron species (Scheme 1).^{10b} Through subsequent reaction optimisation under microwave conditions (Table 1), we identified the best solvent: acetonitrile (entries 2–8); as well as the most effective reagent: tris(2,2,2-trifluoroethyl) borate (entry 15). The background conversion to amide in the absence of any reagent was very low (entry 1).^{10a} The reaction was much more effective in polar aprotic solvents (entries 5, 6, 8) than when conducted neat (entry 7),¹⁵ even though a smaller quantity of reagent was employed. The presence of ROH compounds, including the by-products of the reaction (water, MeOH, *etc.*), significantly reduces the conversion (entries 9–11), presumably *via* inhibition of the reagent.



Scheme 1 Borate-mediated direct amidation.

Table 2 Borate-promoted direct amide formations

Entry	Product	Isolated yield (%)		
		B(OCH ₂ CF ₃) ₃ ^a	B(OMe) ₃ ^a	Thermal ^b
1		91 (74) ^c	92 (66) ^c	18
2		70	73	<1
3		70	51	7
4		76	44	5
5		14 (50) ^d	2	0
6		27 (71) ^d	12	<1
7		92	45	9
8		82	51	6
9		66	66	6
10		61	36	3
11		70	Quant	9
12		71	17	8
13		95	11	6

Table 2 (Contd.)

Entry	Product	Isolated yield (%)		
		B(OCH ₂ CF ₃) ₃ ^a	B(OMe) ₃ ^a	Thermal ^b
14		72	4	0
	PMP = <i>p</i> -MeOC ₆ H ₄ Ar = <i>p</i> -CF ₃ C ₆ H ₄			
15 ^c		94	60	5
16		81 ^f	49 ^g	7

^a Reaction conditions: acid (1 equiv), amine (1 equiv), borate (2 equiv), 0.5 M MeCN, 80 °C, 15 h. ^b Reaction conditions: acid (1 equiv), amine (1 equiv), 0.5 M MeCN, 80 °C, 15 h. ^c 1 equiv of borate used. ^d Carried out at 100 °C in a carousel tube. ^e From amine-HCl salt, with ^tPr₂N⁺Et⁻ (1 equiv). ^f 88% ee. ^g >99% ee.¹⁶

A brief study indicated that thermal conditions were more effective than microwave heating, so we then compared the reactivity of B(OMe)₃ and B(OCH₂CF₃)₃ with a range of acids/amines in acetonitrile at 80 °C. The reactions were conducted in the absence of any additional dehydrating agents or water removal apparatus (Table 2), and good to excellent conversions were obtained in all cases. Although thermally promoted carboxamidation was observed,^{10a} it remained at background levels. In the case of unreactive systems such as pivalic and benzoic acids, yields improved significantly on raising the temperature to 100 °C (entries 5 and 6).

The amides were obtained in high purity after a simple aqueous work-up, and B(OCH₂CF₃)₃ proved to be the optimal reagent in nearly all cases. Both α - and β -substituted acids, as well as α -substituted amines, gave higher yields with this more electron-deficient reagent (entries 3–5, 8, 15–16). B(OCH₂CF₃)₃ was particularly effective for unsaturated carboxylic acids (entries 12–14), and the acylation of an aniline could also be successfully achieved (entry 14).¹⁷ However, it should be noted that B(OMe)₃ was effective for the formation of several amides (e.g. entries 1–2, 11), providing an extremely economical method for accessing these systems. In contrast to other boron reagents and catalysts, anhydrous reaction conditions (dry solvents, inert atmosphere) are not required. The use of acetonitrile as solvent is also practically useful as it enables polar substrates to be coupled effectively (entries 7, 11, 15–16). Notably, highly polar amines such as ethanolamine (entry 7) and tryptamine (entry 11) can be acylated without protection. An amine salt could be used directly in the coupling reactions in the presence of Hünig's base (entry 15), and a Boc-protected amino acid was coupled with only low levels of racemisation (entry 16). The acid-labile Boc group is not cleaved under the reaction conditions, despite the presence of the Lewis acidic boron reagent. A range of other functional groups including

Table 3 Tris(2,2,2-trifluoroethyl) borate-promoted transamidation

Entry	Product	Isolated yield (%)
1		73
2		63
3		82
4		62

alkenes, cyclopropanes, indoles, hydroxyl groups and esters were also well tolerated (entries 7, 9–11, 15).

Given the fact that these borate ester reagents had proved highly effective for the activation of carboxylic acids, we were keen to explore their potential for activating other related systems. Although esters did not undergo amidation (Table 2, entry 15),⁷ B(OCH₂CF₃)₃ was observed to activate primary amides (Table 3). This boron-mediated transamidation reaction gave good yields of secondary amides, and shows very good functional group tolerance (entries 3–4). Although a number of different procedures for transamidation have been reported,¹³ there are few methods available for the transamidation of primary amides without a separate pre-activation step.^{13e–13f} In contrast to these other reports, this method is experimentally simple,^{13f} and requires only equimolar quantities of amine/amide.^{13e} No transamidation was observed in the absence of B(OCH₂CF₃)₃, or in the presence of B(OMe)₃.

In summary, we have demonstrated that simple borates are practical reagents for direct amide bond formation under both thermal and microwave conditions. Unlike many other coupling methods, this approach exhibits good functional group tolerance and purification is extremely straightforward. Tris(2,2,2-trifluoroethyl) borate was also shown to activate amides toward transamidation, providing a convenient and practical method for the direct conversion of primary amides to secondary amides. Further work on the development and application of other boron-centered reagents is ongoing and will be reported in due course.

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