Supplementary Materials

General Remarks

¹H-NMR and ¹³C-NMR data were recorded in CDCl₃. The chemical shifts are reported as delta (δ) units in parts per million (ppm) relative to the solvent residual peak as the internal reference. Coupling constants (*J*) for all spectra are reported in Hertz (Hz). Reactions were monitored by thin-layer chromatography on 0.25 mm E. Merck silica gel 60 plates (F254) using UV light, vanillin and *p*-anisaldehyde as visualizing agents. Potassium allyltrifluoroborate [S1] and potassium (*E*)-crotyltrifluoroborate [S2] were prepared according to the literature procedures.

References

- S1. Molander, G.A.; Figueroa, R.A. *cis*-Dihydroxylation of unsaturated potassium alkyl- and aryltrifluoroborates. *Org. Lett.* 2006, *8*, 75–78.
- S2. Batey, R.A.; Thadani, A.V.; Smil, D.V. Potassium allyl- and crotyltrifluoroborates: Stable and efficient agents for allylation and crotylation. *Tetrahedron Lett.* **1999**, *40*, 4289–4292.





















Figure S8. ¹³C-NMR spectrum (75 MHz, CDCl₃) of 3d.







Figure S10. ¹³C-NMR spectrum (75 MHz, CDCl₃) of **3e**.





Figure S13. ¹H-NMR spectrum (300 MHz, CDCl₃) of 3g.



FG. Hex.esp





Figure S15. ¹H-NMR spectrum (300 MHz, CDCl₃) of **3h**.

FG.Cin.esp ŎН 3h
 1.00
 0.99
 0.96

 6.5
 6.0
1.95 1.00 2.11 2.5 1.22 5.16 7.5 7.0 9.5 5 5.0 4 Chemical Shift (ppm) 4.5 4.0 0 9.0 8.5 3.5 3.0 2.0 0.5 5.5 1.5 1.0 8.0







3i







Figure S21. ¹H-NMR spectrum (300 MHz, CDCl₃) of 3k.



FG-p-Tol.esp











FG-30Me.esp

Figure S25. ¹H-NMR spectrum (300 MHz, CDCl₃) of **3m**.







Figure S27. ¹H-NMR spectrum (300 MHz, CDCl₃) of 3n.

FG.2-OMe.esp











Figure S31. ¹H-NMR spectrum (300 MHz, CDCl₃) of **5**.











Figure S34. ¹H-NMR spectrum (300 MHz, CDCl₃) of 7.





Figure S35. ¹³C-NMR spectrum (75 MHz, CDCl₃) of 7.