Supplementary Material

The synthesis of N^{α} -protected amino hydroxamic acid from N^{α} -protected amino acids employing versatile chlorinating agent CPI-Cl

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Figure 1. ¹H NMR spectrum of compound 2a



Figure 2. ¹³ C NMR spectrum of compound 2a



Figure 3. ¹H NMR spectrum of compound 2b



Figure 4. ¹³C NMR spectrum of compound 2b



Figure 5. ¹H NMR spectrum of compound 2c



Figure 6. ¹³C NMR spectrum of compound 2c



Figure 7. ¹H NMR spectrum of compound 2d



Figure 8. ¹³C NMR spectrum of compound 2d



Figure 9. ¹H NMR spectrum of compound 2e



Figure 10. ¹³ C NMR spectrum of compound 2e



Figure 11. ¹H NMR spectrum of compound 2g



Figure 12. ¹³ C NMR spectrum of compound 2g



Figure 13. ¹H NMR spectrum of compound 2h



Figure 14. ¹³ C NMR spectrum of compound 2h



Figure 15. ¹H NMR spectrum of compound 2i



Figure 16. ¹³ C NMR spectrum of compound 2i



Figure 17. ¹H NMR spectrum of compound 2j



Figure 18. ¹³ C NMR spectrum of compound 2j



Figure 19. ¹H NMR spectrum of compound 2k



Figure 20. ¹³C NMR spectrum of compound 2k



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Figure 21. ¹H NMR spectrum of compound 2k*



Figure 22. ¹³C NMR spectrum of compound 2k*



Figure 23. ¹H NMR spectrum of compound 2I



Figure 24. ¹³C NMR spectrum of compound 2I



Figure 25. RP-HPLC profile of 2K



Figure 26. RP-HPLC profile of 2k*



Figure 27. RP-HPLC profile of equimolar mixture of 2k and 2k*



The RP-HPLC analysis of epimers was carried out using an Agilent instrument (method: gradient 0.1% TFA wateracetonitrile (0–100%) in 20 min; VWD at λ 254 nm; flow rate: 1.0 mL/min; column: Agilent Eclipse, XDB-C18, pore size 5 μ m, diameter × length = 4.6 × 150 nm)