

CORRECTION

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Correction: Triamide macrocyclic chloride receptors *via* a one-pot tandem reduction–condensation–cyclization reaction

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Correction for 'Triamide macrocyclic chloride receptors *via* a one-pot tandem reduction–condensation–cyclization reaction' by Harekrushna Behera, *et al.*, *Org. Biomol. Chem.*, 2017, DOI: 10.1039/c7ob00642j.

The authors regret that some highly relevant citations reporting the synthesis of furan based triamide macrocycles were missed.

Therefore the sentence "To the best of our knowledge the only one-pot method for obtaining triamides (with a C_n symmetry) from the corresponding monomer was reported by Kilbinger and co-workers.⁴²" should be replaced by "Pyridyl based triamide macrocycles have been synthesized without the use of coupling reagents by Kilbinger and co-workers, while Chakraborty and co-workers have developed a variety of elegant furan based chiral as well as achiral triamide macrocycles using coupling reagents.⁴²"

The corrected reference 42 is shown below:

42 (a) C. Storz, M. Badoux, C. M. Hauke, T. S. Šolomek, A. Kühnle, T. Bally and A. F. Kilbinger, *J. Am. Chem. Soc.*, 2014, **136**, 12832–12835; (b) T. K. Chakraborty, S. Tapadar and S. K. Kumar, *Tetrahedron Lett.*, 2002, **43**, 1317–1320; (c) T. K. Chakraborty, S. Tapadar, T. V. Raju, J. Annapurna and H. Singh, *Synlett*, 2004, 2484–2488; (d) T. K. Chakraborty, E. Bikshapathy, R. Nagaraj, M. Vairamani, S. Kiran Kumar, and A. C. Kunwar, *J. Org. Chem.*, 2003, **68**, 6257–6263.

The Royal Society of Chemistry apologises for these errors and any consequent inconvenience to authors and readers.

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