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CORRECTION

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Correction: Rapid synthesis of internal peptidyl α-ketoamides by on resin oxidation for the construction of rhomboid protease inhibitors

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Correction for 'Rapid synthesis of internal peptidyl α-ketoamides by on resin oxidation for the construction of rhomboid protease inhibitors' by Tim Van Kersavond *et al.*, *RSC Adv.*, 2021, **11**, 4196–4199, DOI: 10.1039/D0RA10614C.

The authors regret that an incorrect version of Fig. 1 was presented in the original manuscript. The R-group in compound **7** was incorrectly indicated as $(CH_2)_5Ph$. The corrected version of the figure with the R group as $(CH_2)_4Ph$ is shown below.

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

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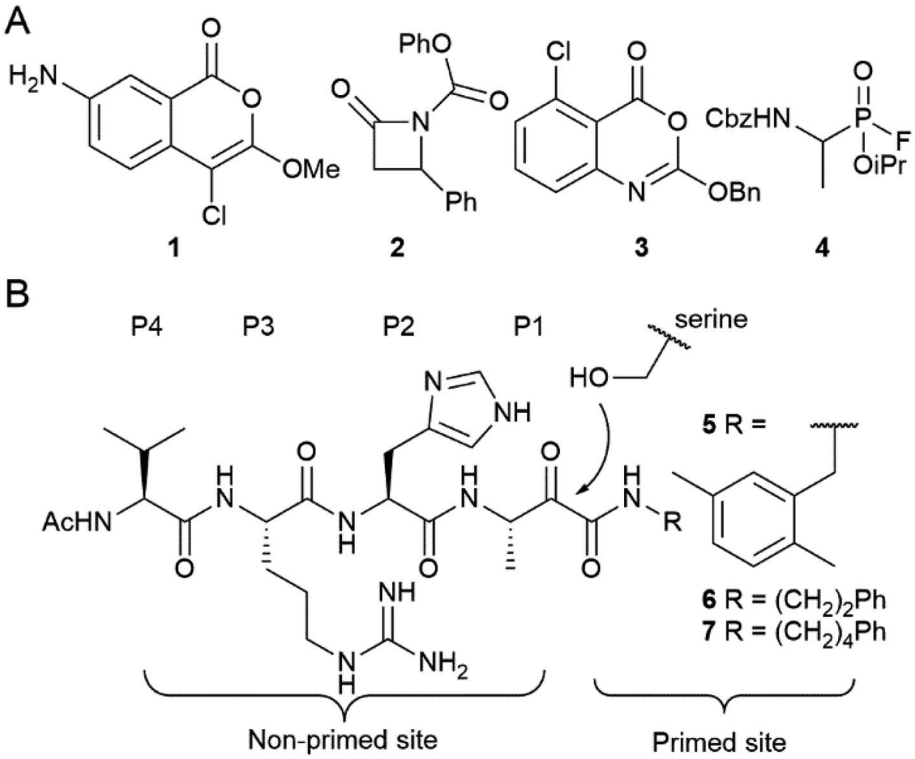

Fig. 1 Examples of rhomboid inhibitors. (A) 4-Chloro-isocoumarins (**1**), β-lactams (**2**), benzoxazinones (**3**) and fluorophosphonates (**4**). (B) α-Ketoamide rhomboid inhibitors (**5**–**7**). The peptidic element in the non-primed site is indicated with the P1–P4 position according to the Schechter and Berger protease substrate nomenclature.[1]

References

1 I. Schechter and A. Berger, *Biochem. Biophys. Res. Commun.*, 1967, **27**, 157–162.