



Cite this: *Chem. Commun.*, 2017, 53, 2555

Received 23rd December 2016,
Accepted 18th January 2017

DOI: 10.1039/c6cc10178j

rsc.li/chemcomm

6-*exo*-trig Michael addition-lactonizations for catalytic enantioselective chromenone synthesis†

Rifahath M. Neyyappadath, David B. Cordes, Alexandra M. Z. Slawin and Andrew D. Smith*

The catalytic enantioselective 6-*exo*-trig Michael addition-lactonization of enone-acid substrates to form *cis*-chromenones with high diastereo- and enantiocontrol was developed using the commercially available isothiurea tetramisole. An acidic workup proved necessary to minimize product epimerization and maximize product *er*, providing *cis*-chromenones in excellent yield, and with excellent diastereo- and enantioselectivity.

The development of catalytic processes that allow the preparation of valuable heterocyclic frameworks from readily prepared starting materials under mild conditions is of widespread importance.¹ A range of enantioselective methods that fulfil these goals has been developed.² In recent years the catalytic use of C(1)-ammonium enolates,³ particularly those using carboxylic acids as starting materials,⁴ has been popularized following the intramolecular enantioselective nucleophile-catalyzed aldol lactonization (NCAL) methodology developed by Romo for the synthesis of stereodefined β -lactones.⁵ In this area, 5-*exo*-ring closure to prepare the corresponding carbo- and heterocyclic ring systems is commonplace (Fig. 1a), and this strategy has been applied successfully for the construction of complex molecular targets.⁶ To date, only limited isolated examples of this approach for the formation of 6-membered ring systems have been developed,⁷ all of which use cinchona alkaloids as catalysts. In previous work we developed an isothiurea-catalyzed^{8,9} 5-*exo*-Michael addition-lactonization approach to 5-membered carbo- and heterocycle synthesis from enone acids (Fig. 1b).¹⁰ In this manuscript the application of this methodology for the preparation of 6-membered heterocycles is reported for the first time, allowing the synthesis of *cis*-chromenones¹¹ in up to 99 : 1 dr and 98 : 2 er (Fig. 1c).

EaStCHEM, School of Chemistry, University of St Andrews, North Haugh, St Andrews, KY16 9ST, UK. E-mail: ads10@st-andrews.ac.uk

† Electronic supplementary information (ESI) available: Experimental procedures; characterization data for novel compounds; ¹H and ¹³C NMR spectra, HPLC traces,¹² and X-ray crystallographic data files compounds 9 and 27 (CIF). CCDC 1510311 and 1510312. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c6cc10178j

a. Known: NCAL cyclizations for carbo- and heterocycle synthesis



b. Previous work: isothiurea promoted intramolecular cyclizations



c. This work: isothiurea promoted 6-*exo*-trig cyclization



Fig. 1 Summary of ammonium enolate promoted intramolecular catalytic enantioselective carbo- and heterocycle formation.

Initial model studies probed the cyclization of enone acid 1 to chromenone 2, with 1 readily prepared in three steps from 2-hydroxyphenylacetic acid.¹² Treatment of 1 with pivaloyl chloride and *i*-Pr₂NEt gave the corresponding mixed anhydride *in situ*, which was subsequently treated with isothiurea catalysts 3 to 6 and evaluated for the proposed cyclization (Table 1, entries 1–4). Achiral DHPB¹³ gave the desired *cis*-chromenone 2 in 85% yield and > 99 : 1 dr. Screening of a small range of chiral isothiureas 4–6 indicated the use of tetramisole 4 and its benzannulated counterpart, BTM 5, showed promising enantioselectivity (~ 87 : 13 er, entries 2 and 3). Subsequent optimization through variation of solvent, temperature and base¹² showed that performing the reaction at 0 °C in CHCl₃ with excess *i*-Pr₂NEt (1.5 equiv. for mixed anhydride formation, followed by an additional 2.5 equiv.) gave highest observed dr and er (entries 6–9). Lowering the catalyst loading to 5 mol% using tetramisole 4 gave 2 in 85% yield, > 99 : 1 dr and 93 : 7 er, with BTM 5 giving lower



olefination.¹² Using the 0.1 M HCl work up protocol generally high product er and dr was observed.¹⁵ Notable trends within this series showed that incorporation of halogen (4-FC₆H₄**8** and 4-ClC₆H₄**9**) substituents, as well as electron-donating (4-MeOC₆H₄**10** and 4-MeC₆H₄**11**) and 2-naphthyl substituents **15** gave the desired *cis*-chromenones in excellent enantioselectivity (97 : 3 to 98 : 2 er). Incorporation of electron-withdrawing 4-CF₃C₆H₄ or 3,5-(CF₃)₂C₆H₃ substituents was also tolerated, giving **12** with marginally reduced enantioselectivity and **13** in moderate 37% yield. Incorporation of an aliphatic enone led to decreased reactivity, requiring high catalyst loadings (20 mol%) to promote this transformation (29% isolated yield at 46% conversion), giving **14** as a single diastereoisomer in moderate 71 : 29 er.¹⁶ The relative and absolute configuration within **9** was unambiguously confirmed by X-ray crystal structure analysis,¹⁷ with the absolute configuration of all other products assigned by analogy.

The generality of this methodology was further investigated using different substituents within the aromatic tether (Table 4). Variation of the aromatic tether, incorporating substitution with electron-donating (5-Me, 4-OMe), halogen (4-F) and naphthyl groups gave *cis*-chromenones **16**–**22** with excellent enantioselectivity (95 : 5 to 98 : 2 er). Notably, incorporation of 4-OMe substituents on the aromatic tether (to give **17** and **21**) showed decreased reactivity, with the reaction taking extended reaction times (12–14 h) to reach >98% conversion, but still proceeded with excellent enantioselectivity.

Reaction scale-up and subsequent product derivatization was investigated. On a one-gram scale, complete conversion of **1** to **2** was observed using only 2.5 mol% catalyst within 6 h



Scheme 2 Product derivatization. ^a Isolated yield of major diastereoisomer (>99 : 1 dr). ^b Measured by ¹H NMR spectroscopy of crude reaction product. ^c Measured by chiral HPLC (major *cis*-diastereoisomer). ^d Starting material **2** was >99 : 1 dr and 99 : 1 er.

to give **2** in 86% isolated yield as a single diastereoisomer and 98 : 2 er.

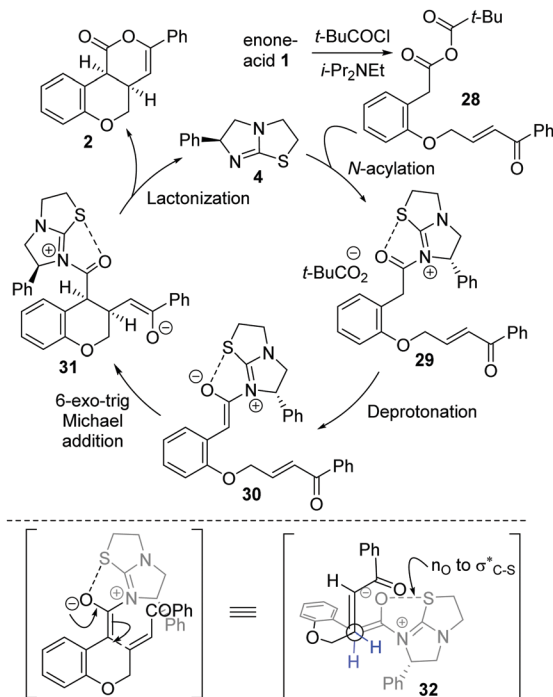
The synthetic utility of the products was then explored through a range of derivatizations (Scheme 2). Ring-opening of **2** with either methanol, morpholine or benzylamine gave the corresponding *cis*-dihydrobenzopyrans **23**–**25** in excellent yield, dr and er. Treatment of *cis*-chromenone **2** with Pd/C and H₂ (1 atm) led to hydrogenation and hydrogenolysis, giving acid **26** in excellent yield. Alternatively, treatment of a recrystallized sample of **2** (>99 : 1 er) with *m*-CPBA, followed by *p*-TSA, gave the 5-membered lactone¹⁸ **27** in excellent yield and stereo-control [96 : 4 dr, >99 : 1 er]. Recrystallization from 10% EtOAc in hexane gave **27** in >99 : 1 dr, >99 : 1 er and 82% yield. The relative and absolute configuration of **27** was confirmed by single crystal X-ray analysis.¹⁷

The mechanism of the isothiurea-catalyzed reaction, shown for the cyclization of enone-acid **1** to **2**, is postulated to proceed *via in situ* formation of mixed anhydride **28** (Scheme 3). Nucleophilic addition of isothiurea **4** to **28** gives acyl isothiuronium ion intermediate **29**, with deprotonation generating (*Z*)-ammonium enolate **30**. Subsequent intramolecular 6-*exo*-trig Michael addition to the tethered enone generates intermediate **31**, with lactonization giving *cis*-chromenone **2** and regenerating the catalyst **4**. A simplistic model to rationalize the observed diastereo- and enantiocontrol utilizes a stabilising n₀ to σ_{C-S}* interaction¹⁹ between the enolate oxygen and the sulfur of the isothiuronium ion to restrict the conformation of the (*Z*)-enolate,²⁰ forcing the stereodirecting phenyl substituent to adopt a pseudoaxial orientation to minimize 1,2-strain. Subsequent 6-*exo*-trig Michael addition occurs *anti*- to this stereodirecting group as represented by pre-transition state assembly **32**, with the two-prostereogenic centres along the developing C–C bond adopting a staggered array to minimize non-bonding interactions.

Table 4 Reaction scope: variation of aromatic tether



^a Combined isolated yield of diastereoisomers. ^b Isolated yield of major diastereoisomer (>99 : 1 dr). ^c Measured by ¹H NMR spectroscopy of crude reaction product. ^d Measured by chiral HPLC (major *cis*-diastereoisomer). ^e 12–14 h reaction time.



Scheme 3 Proposed mechanism and stereochemical rationale.

In conclusion, the catalytic enantioselective synthesis of *cis*-chromenones has been achieved using commercially available tetramisole as a catalyst. This method provides a range of *cis*-chromenone derivatives in high yield with excellent diastereo- and enantiocontrol (up to 99:1 dr and 98:2 er). On-going studies in this laboratory are focused on further applications of Lewis base organocatalysts in enantioselective catalysis.

We thank the EPSRC Centre for Doctoral Training in Critical Resource Catalysis (CRITICAT, grant code EP/L016419/1, RMNP) for funding. The European Research Council under the European Union's Seventh Framework Programme (FP7/2007-2013) ERC Grant Agreement No. 279850 is also acknowledged. ADS thanks the Royal Society for a Wolfson Research Merit Award. We also thank the EPSRC UK National Mass Spectrometry Facility at Swansea University.

Notes and references

- For select reviews see: (a) M. Álvarez-Corral, M. Muñoz-Dorado and I. Rodríguez-García, *Chem. Rev.*, 2008, **108**, 3174–3198; (b) X.-X. Guo, D.-W. Gu, Z. Wu and W. Zhang, *Chem. Rev.*, 2015, **115**, 1622–1651; (c) I. Nakamura and Y. Yamamoto, *Chem. Rev.*, 2004, **104**, 2127; (d) B. Godoi, R. F. Schumacher and G. Zeni, *Chem. Rev.*, 2011, **111**, 2937–2980.
- For select reviews see: (a) J. Yu, F. Shi and L.-Z. Gong, *Acc. Chem. Res.*, 2011, **44**, 1156–1171; (b) S. Ponra and K. C. Majumdar, *RSC Adv.*, 2016, **6**, 37784–37922; (c) L. M. Stanley and M. P. Sibi, *Chem. Rev.*, 2008, **108**, 2887–2902; (d) J. Royer, M. Bonin and L. Micouin, *Chem. Rev.*, 2004, **104**, 2311–2352.
- M. J. Gaunt and C. C. C. Johansson, *Chem. Rev.*, 2007, **107**, 5596–5605.
- L. C. Morrill and A. D. Smith, *Chem. Soc. Rev.*, 2014, **43**, 6214–6226.

- G. S. Cortez, R. L. Tennyson and D. Romo, *J. Am. Chem. Soc.*, 2001, **123**, 7945–7946.
- (a) Y. Yokota, G. S. Cortez and D. Romo, *Tetrahedron*, 2002, **58**, 7075–7080; (b) H. Henry-Riyad, C. Lee, V. C. Purohit and D. Romo, *Org. Lett.*, 2006, **8**, 4363–4366; (c) G. Ma, H. Nguyen and D. Romo, *Org. Lett.*, 2007, **9**, 2143–2146; (d) W. Zhang, A. S. Matla and D. Romo, *Org. Lett.*, 2007, **9**, 2111–2114; (e) H. Nguyen, G. Ma, T. Gladysheva, T. Fremgen and D. Romo, *J. Org. Chem.*, 2011, **76**, 2–12; (f) G. Liu and D. Romo, *Angew. Chem., Int. Ed.*, 2011, **50**, 7537–7540; (g) C. A. Leverett, V. C. Purohit, A. G. Johnson, R. L. Davis, D. J. Tantillo and D. Romo, *J. Am. Chem. Soc.*, 2012, **134**, 13348–13356.
- (a) G. S. Cortez, S. H. Oh and D. Romo, *Synthesis*, 2001, 1731–1736; (b) S. H. Oh, G. S. Cortez and D. Romo, *J. Org. Chem.*, 2005, **70**, 2835–2838; (c) K. A. Morris, K. M. Arendt, S. H. Oh and D. Romo, *Org. Lett.*, 2010, **12**, 3764–3767; (d) D. Sikriwal and D. K. Dikshit, *Tetrahedron*, 2011, **67**, 210–215.
- For reviews on isothioureia catalysis (a) J. E. Taylor, S. D. Bull and J. M. J. Williams, *Chem. Soc. Rev.*, 2012, **41**, 2109–2121; (b) J. Merad, J.-M. Pons, O. Chuzel and C. Bressy, *Eur. J. Org. Chem.*, 2016, 5589–5610.
- For seminal work on isothioureia catalysis (a) X. Li and V. B. Birman, *Org. Lett.*, 2006, **8**, 1351–1354; (b) H. Jiang, X. Li, L. Guo, E. W. Uffman and V. B. Birman, *J. Am. Chem. Soc.*, 2006, **128**, 6536–6537; (c) M. Kobayashi and S. Okamoto, *S. Tetrahedron Lett.*, 2006, **47**, 4347–4350; (d) X. Li and V. B. Birman, *Org. Lett.*, 2008, **10**, 1115–1118; (e) Y. Zhang and V. B. Birman, *Adv. Synth. Catal.*, 2009, **351**, 2525–2529; (f) C. Joannesse, C. P. Johnston, C. Concellón, C. Simal, D. Philp and A. D. Smith, *Angew. Chem., Int. Ed.*, 2009, **48**, 8914–8918.
- (a) D. Belmessieri, L. C. Morrill, C. Simal, A. M. Z. Slawin and A. D. Smith, *J. Am. Chem. Soc.*, 2011, **133**, 2714–2720; (b) D. Belmessieri, D. B. Cordes, A. M. Z. Slawin and A. D. Smith, *Org. Lett.*, 2013, **15**, 3472–3475; (c) D. Belmessieri, A. de la Houpliere, E. D. D. Calder, J. E. Taylor and A. D. Smith, *Chem. – Eur. J.*, 2014, **20**, 9762–9769; (d) D. G. Stark, P. Williamson, E. R. Gayner, S. F. Musolino, R. W. F. Kerr, J. E. Taylor, A. M. Z. Slawin, T. J. C. O'Riordan, S. A. Macgregor and A. D. Smith, *Org. Biomol. Chem.*, 2016, **14**, 8957–8965.
- For an overview see: (a) R. Pratab and V. J. Ram, *Chem. Rev.*, 2014, **114**, 10476–10526. For catalytic routes see: (b) J. Chen, J. Chen, F. Lang, X. Zhang, L. Cun, J. Zhu, J. Deng and L. Liao, *J. Am. Chem. Soc.*, 2010, **132**, 4552–4553; (c) A. M. Hardman-Baldwin, M. D. Visco, J. M. Wieting, C. Stern, S. Kondo and A. E. Mattson, *Org. Lett.*, 2016, **18**, 3766–3769.
- See ESI† for substrate synthesis, optimization and epimerization.
- V. B. Birman, X. Li and Z. Han, *Org. Lett.*, 2007, **9**, 37–40 and 9(c).
- Epimerization of the 80:20 *trans*:*cis* mixture gave exclusively *cis*-2 (>99:1 dr and 38:62 er (4aR,10bS:4aS,10bR)). See ESI† for details.
- To test the generality of the acidic work up to minimize the epimerization process, a number of reactions were quenched using either H₂O at rt or 0.1 M HCl at 0 °C. See ESI† for full information.
- After work-up a 54:46 mixture of starting material acid enone and **14** was observed in the crude reaction mixture. We hypothesize that competitive deprotonation may retard reactivity in this case.
- Crystallographic data obtained for **9** and **27** has been deposited with the Cambridge Crystallographic Data Centre and the supplementary data can be found via CCDC 1510311 and 1510312 respectively.
- Z. Fu, X. Wu and Y. R. Chi, *Org. Chem. Front.*, 2016, **3**, 145–149.
- For 1,5-S···O interactions as a control element in isothioureia catalysis see (a) M. E. Abbasov, B. M. Hudson, D. J. Tantillo and D. Romo, *J. Am. Chem. Soc.*, 2014, **136**, 4492–4495; (b) P. Liu, X. Yang, V. B. Birman and K. N. Houk, *Org. Lett.*, 2012, **14**, 3288–3291; (c) E. R. T. Robinson, D. M. Walden, C. Fallan, M. D. Greenhalgh, P. H.-Y. Cheong and A. D. Smith, *Chem. Sci.*, 2016, **7**, 6916–6927; (d) Y. Nagao, S. Miyamoto, M. Miyamoto, H. Takeshige, K. Hayashi, S. Sano, M. Shiro, K. Yamaguchi and Y. Sei, *J. Am. Chem. Soc.*, 2006, **128**, 9722–9729.
- For discussions on the origin of S···O interactions, see: (a) B. R. Beno, K.-S. Yeung, M. D. Bartberger, L. D. Pennington and N. A. Meanwell, *J. Med. Chem.*, 2015, **58**, 4383–4438; (b) X. Zhang, Z. Gong, J. Li and T. Lu, *J. Chem. Inf. Model.*, 2015, **55**, 2138–2153; (c) R. C. Reid, M.-K. Yau, R. Singh, J. Lim and D. P. Fairlie, *J. Am. Chem. Soc.*, 2014, **136**, 11914–11917.

