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Enzymatic halocyclization of allenic alcohols and carboxylates: a biocatalytic entry to functionalized O-heterocycles†

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Chloroperoxidase from *Caldariomyces fumago* catalyzes the aerobic oxidative halocyclization of allenic alcohols and carboxylates yielding functionalized furan and pyran heterocycles as valuable synthetic scaffolds. Thanks to an oxidase-initiated redox cascade, simple halide salts – in combination with air and glucose – act as stoichiometric reagents for the *in situ* generation of reactive halonium species. Under the mild reaction conditions in an aqueous emulsion medium, the stereochemical integrity of diastereo- and enantioenriched allenes remains uncompromised and chiral dihydrofurans can be obtained via 5-*endo-trig* cyclizations with perfect axis-to-centre chirality transfer.

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Introduction

Halogenation reactions play a crucial role in organic synthesis as many target structures in agricultural and pharmaceutical chemistry feature halide substitution patterns.^{1–3} Furthermore, chlorine, bromine, and iodine atoms serve as highly important functional groups in synthetic intermediates allowing for efficient and selective carbon–carbon and carbon–heteroatom coupling reactions.^{4–6} Classically, molecular halogen has been used as reagent of choice for the introduction of halide entities both *via* radical pathways and as source of electrophilic halonium species. Due to the harmful, corrosive and toxic properties of the lighter dihalogens, organic reagents such as *N*-halosuccinimides, *N,N'*-dihalohydantoins, and tetrahalomethane have nowadays replaced elemental X₂ to some extent in certain processes.

In Nature, the major pathways for the biosynthesis of halogenated secondary metabolites rely on the enzymatic oxidative activation of chloride, bromide, and iodide as readily available source of halogen atoms.⁷ Here, halogenases^{8,9} and haloperoxidases^{10–12} generate electrophilic hypohalide species under consumption of dioxygen (and a suitable reducing agent) or hydrogen peroxide, respectively. The enzyme-bound halonium equivalents are subsequently transferred to the

acceptor molecules either by direct protein–substrate interaction or *via* liberation of free hypohalide. In recent years, the synthetic use of halogenating enzymes *ex vivo* has attracted much interest as it enables the incorporation of halide groups under very mild reaction conditions.^{13–16} Here, the substitution of high molecular weight organic halogen donors or hazardous dihalogens by simple alkali halides and air in combination with the general benefits of biocatalysis, including high selectivities, low temperatures and aqueous reaction media, can be considered a major advancement.

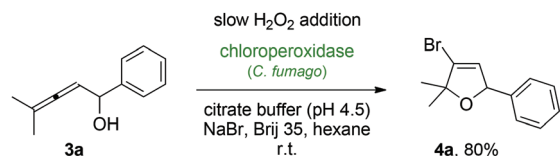
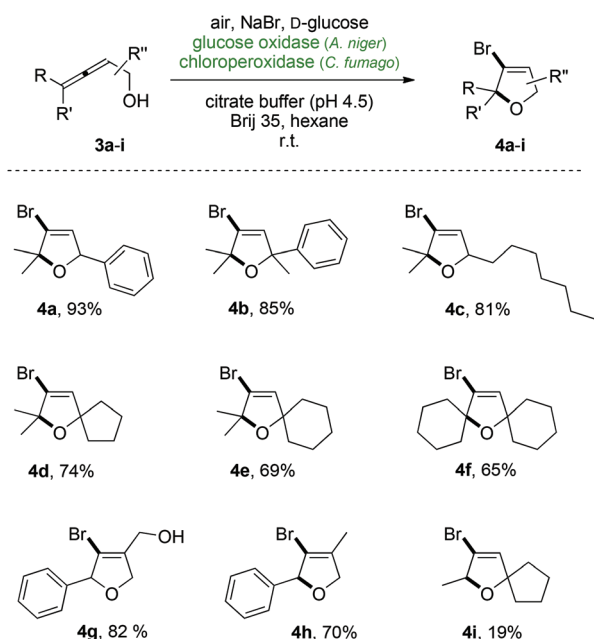
Allenes represent synthetically highly attractive building blocks that are able to engage in numerous, partly unique reactions.^{17–19} Among these many valuable transformations, suitably decorated allenes can act as acceptor for halonium ions and undergo oxidative cyclizations (both *exo-trig*^{20,21} and *endo-trig*^{22–25}) that furnish heterocyclic entities containing vinyl halide units as versatile functional handle for *e.g.* subsequent cross coupling reactions. Particularly gold catalysis proved to efficiently promote this kind of allene activation (Scheme 1, top),^{26,27} enabling the development of asymmetric halocyclizations.²⁸ Based on our recent investigations of the commercial fungal chloroperoxidase from *Caldariomyces fumago* (CPO) as catalyst in oxidative rearrangement reactions and the observation of unbeneficial halide interferences,²⁹ we became interested in the deliberate use of CPO as biological mediator for the halogenative cyclization of allenic alcohols and structurally related entities. Herein, we report on the development of a purely biocatalytic, environmentally benign,^{30–32} and high yielding protocol for the synthesis of functionalized O-heterocycles based on the halocyclization approach (Scheme 1, bottom). In our reaction design, glucose oxidase (GOx) served as catalyst for the controlled

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Scheme 2 Glucose-free halogenation via periodic addition of H₂O₂.Scheme 3 Substrate scope of the aerobic bromocyclization of α -allenols.

and **4c** in high yields. To our delight, the peroxidase-catalyzed cyclization also allowed for the preparation of spirocyclic brominated building blocks (**4d** & **4e**) and the bis-spirodihydrofuran **4f**. Furthermore, the successful halogenation of the allenic diol **3g** and allenol **3h** with high yields for the bromodihydrofurans **4g** and **4h** proved that the terminal disubstitution pattern ($R \text{ \& } R' \neq H$) is not a necessary requirement for an efficient aerobic cyclization. However, by removing the *gem*-dialkyl or aryl substituents, respectively, at the allenenes' γ -position – that would serve as stabilizing factor for potential cationic reaction intermediates – substantially reduced yields were observed (**4i**). Incubation of terminally unsubstituted allenols did not show any bromodihydrofuran formation but instead, products arising from semipinacol rearrangements were identified as minor reaction products (<10% yield).

Furthermore, a series of β -allenols was tested in order to elucidate the potential use of the oxidase/peroxidase protocol for the synthesis of halogenated pyran building blocks. The attempt to cyclize the terminally dimethylated but otherwise unsubstituted alcohol **5a** failed and the desired brominated pyran **6a** was observed only in minor concentrations (Table 2, entry 1). Geminal dimethylation, either next to the allene function (**5b**) or at the carbinol carbon (**5c**) led to slightly improved

Table 2 Bromocyclization of β -allenols^a

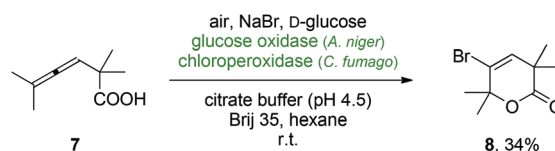
Entry	Allenol	R	R'	R''	R'''	Yield (%)
1	5a	H	H	H	H	6a , 4
2	5b	Me	Me	H	H	6b , 13
3	5c	H	H	Me	Me	6c , 13
4	5d	H	Ph	Me	Me	6d , 46

^a **5** (0.2 mmol), D-glucose (72 mg, 0.4 mmol), NaBr (62 mg, 0.6 mmol), Brij 35 (0.2 mmol), glucose oxidase (*A. niger*, 50 U), chloroperoxidase (*C. fumago*, 200 U), citrate buffer (0.1 M, pH 4.5, 10 mL), hexane (10 mL), r.t., 3–5 h. Conversions and yields were determined via ¹H NMR with dimethylsulfone as standard.

yields of the six-membered heterocycles of 13% (Table 2, entries 2 & 3). Although still lagging behind the α -allenol substrates, with the very densely substituted alcohol **5d** reasonable conversions in the halogenative 6-*endo-trig* cyclization were finally observed giving rise to the bromopyran **6d** in 46% yield.

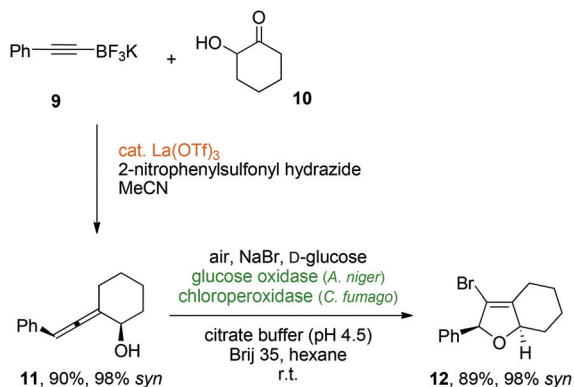
As another extension of this biocatalytic allene activation mode, we aimed to transfer the protocol to the cyclization of allenic carboxylates. To our surprise, only traces of the desired five-membered lactones were observed when 2,3-dienoic acids were incubated under the previously optimized conditions. On the other hand, bromolactonization of **3,4**-dienoate **7** resulted in the formation of δ -lactone **8** as major product (Scheme 4). Considering the low pH of the reaction medium as a prerequisite for the CPO-catalyzed halide oxidation, the presence of the low-nucleophilic free acid might attribute to the relatively sluggish conversion. The use of other haloperoxidases with activity under slightly less acidic conditions might therefore be beneficial and will be included in future studies.

In order to investigate the stereochemical integrity during the oxidative cyclization of diastereo- or enantiomerically defined allenic alcohols, Thomson's lanthanum-catalyzed traceless Petasis reaction between an alkynyltrifluoroborate (**9**) and hydroxycyclohexanone (**10**) was used to synthesize the cyclic allenol **11**,³⁶ which was obtained in excellent diastereoselectivity (98% *syn*). Gratifyingly, the protocol for the enzymatic allenol cyclization under biphasic conditions gave rise to the desired bicyclic dihydrofuran **12** in 89% yield. Most crucially, the relative *syn*-stereochemistry was quantitatively transferred from the axially chiral allene moiety to the newly



Scheme 4 Peroxidase-mediated halolactonization.

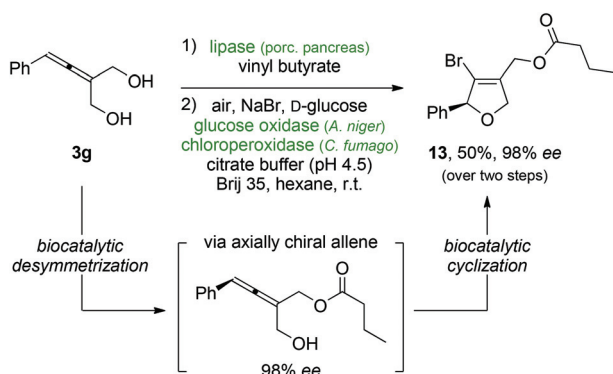




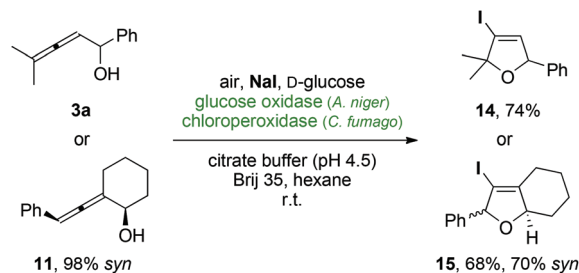
Scheme 5 Perfect chirality transfer in a metal-enzyme-coupled catalytic cascade for the synthesis of a stereochemically defined bicyclic vinyl bromide.

formed centrochiral heterocycle with a diastereomeric excess of **12** of 98% (Scheme 5).

Encouraged by these results, we turned our attention to the aspect of absolute stereochemistry and the perspective to employ multi-enzyme cascades for the preparation of enantio-enriched bromoheterocycles. On the basis of a previously described protocol for the lipase-mediated enantioselective desymmetrization of allenic diols,^{37–39} we designed a two-step sequence that allowed for the generation of optically pure, axially chiral allenols whose subsequent peroxidase-catalyzed halocyclization led to enantiomerically enriched bromodihydrofurans. The stereoselective acylation of allene **3g** by lipase from porcine pancreas with vinyl butyrate in 1,4-dioxane yielded the intermediate allenol in excellent enantiomeric excess of 98%. Also here, incubation of the axially chiral allene in the oxidase/peroxidase emulsion system resulted in smooth conversion to the expected functionalized bromide (*S*)-**13**. Subsequent basic hydrolysis of the butyrate to the alcohol (*S*)-**4g** and comparison of HPLC traces with those of a racemic reference confirmed complete conservation of the 98% optical purity during the bromoetherification (see ESI Fig. 1†) (Scheme 6).



Scheme 6 Multi enzyme cascade for the preparation of enantiomerically pure 3-bromodihydrofurans.



Scheme 7 Chloroperoxidase-triggered 5-endo-trig iodocyclizations.

Finally, the application of other halide salts was tested in order to extend the protocol towards the chlorination and iodination of allenes. In presence of sodium chloride, even after prolonged reaction times only a very low conversion was observed (<5%). In strong contrast, the corresponding iodide system provided a rapid and clean formation of the iodinated cyclization product from **3a** yielding **14** in 74% after only 1 hour (Scheme 7). Employing the diastereomerically enriched alcohol **11**, also good conversion in the iodocyclization was observed. However, due to partial epimerization of the starting material under the reaction conditions, the bicyclic iododihydrofuran **15** was obtained as mixture of diastereomers. Further studies and optimizations regarding the stereochemical integrity of axially chiral allenes in the peroxidase-mediated cyclization are hence required in order to exploit this protocol also in the synthesis of optically active iodinated O-heterocycles.

Conclusions

In this study, we disclosed that halide-oxidizing peroxidases can act as highly efficient catalysts for the oxidative 5-endo-trig cyclization of substituted α -allenols yielding brominated dihydrofuran building blocks. Hereby, the presented method avoids any kind of hazardous elemental dihalogen or high molecular weight organic halogen donors by taking advantage of an oxidase/peroxidase cascade utilizing NaBr, glucose and aerial oxygen as stoichiometric reagents in an aqueous emulsion medium. In addition to a wide substrate scope for the formation of five-membered O-heterocycles, this biocatalytic protocol can be employed to a lesser extend also for the synthesis of six-membered cyclic ethers and lactones. Of particular interest from a synthetic-organic perspective, the enzymatic bromocyclization tolerates both stereogenic carbinol centres and chiral allene axes and thus shows great potential for the chemoenzymatic stereoselective synthesis of complex heterocyclic target structure.

Experimental

General remarks

All reactions carried out under argon atmosphere were performed with dry solvents using anhydrous conditions.



Anhydrous solvents were obtained from an MBraun MB-SPS800 drying system. Commercially available reagents were used without further purification. Catalysts and cofactors were obtained from: NADH, Carbolution Chemicals GmbH; Lactoperoxidase from *Bos taurus* (LPO), 106 U mg⁻¹ (ABTS), Bio-Research Products; Peroxidase from *Armoracia rusticana* (HRP), 148 U mg⁻¹ (ABTS), Sigma; Peroxidase from *Glycine max* (SBP), 17.5 kU mL⁻¹ (ABTS), Bio-Research Products; Chloroperoxidase from *Caldariomyces fumago* (CPO), 9.9 kU mL⁻¹ (ABTS), Sigma; Glucose oxidase type II from *Aspergillus niger* (GOx), 17.3 U mg⁻¹ (HRP/ABTS), Sigma; Alcohol oxidase from *Pichia pastoris*, 1.1 kU mL⁻¹ (HRP/ABTS), Sigma; Alcohol oxidase from *Candida boidinii*, 1.0 U mg⁻¹ (HRP/ABTS), Sigma; lipase from porcine pancreas (PPL), MP Biomedicals. All products were purified either by column chromatography over silica gel (Macherey-Nagel MN-Kieselgel 60, 40–60 µm, 240–400 mesh) or by recrystallization. Reactions were monitored by thin layer chromatography (TLC) carried out on pre-coated silica gel plates (Macherey-Nagel, TLC Silica gel 60 F₂₅₄) using UV light and KMnO₄-solution or Hanessian's stain for visualization. ¹H-NMR and ¹³C-NMR spectra were recorded at room temperature on a Bruker Avance 400 instrument. Chemical shifts are reported in parts per million (ppm) calibrated using residual non-deuterated solvents as internal reference (CHCl₃ at 7.26 ppm (¹H-NMR) and 77.16 ppm (¹³C-NMR)). Infrared spectra were recorded on a Bruker Alpha ECO-ATR FT-IR-Spectrometer, absorption bands are reported in wave numbers [cm⁻¹]. High performance liquid chromatography was performed on a Waters system using a Waters 501 pump and a Waters 2487 dual detector.

Representative procedure for the enzymatic halocyclization: 3-bromo-2,2-dimethyl-5-phenyl-2,5-dihydrofuran (4a)

n-Hexane (10 mL) was added to a solution of Brij 35® (243 mg, 200 µmol) in an aqueous citrate buffer (10 mL, 0.1 M, pH 4.5) to form an emulsion by vigorous stirring. Anhydrous sodium bromide (61 mg, 600 µmol), D-(+)-glucose (72 mg, 400 µmol), alcohol **3a** (200 µmol) and chloroperoxidase from *Caldariomyces fumago* (20 µL, 200 U, 1 U µmol⁻¹) were all added to the emulsion. Glucose oxidase from *Aspergillus niger* (3 mg, 50 U, 0.25 U µmol⁻¹) was finally added to start the reaction. The mixture was stirred vigorously for 3 hours under air. Acetonitrile (10 mL) was added to the mixture and the layers were separated. The aqueous layer was extracted with *n*-pentane (3 × 30 mL). The combined organic layers were washed with brine, dried over Na₂SO₄ and concentrated under reduced pressure. Purification by flash chromatography (SiO₂, *n*-hexane/EtOAc, 9/1) provided **4a** (43 mg, 170 µmol, 85% yield) as a light yellow oil. *R*_f (cyclohexane/ethyl acetate 6/1): 0.85. ¹H-NMR (400 MHz, CDCl₃): δ [ppm] = 7.39–7.28 (m, 5H), 5.91 (d, ³J = 1.6 Hz, 1H), 5.69 (d, ³J = 1.6 Hz, 1H), 1.50 (s, 3H), 1.46 (s, 3H). ¹³C-NMR (100 MHz, CDCl₃): δ [ppm] = 140.8, 128.7, 128.3, 126.8, 126.7, 89.1, 85.3, 27.6, 26.2. FT-IR (neat, ATR): ν [cm⁻¹] = 3064 (w), 3030 (w), 1628 (m), 1454 (s), 1377 (m), 1361 (m), 1150 (m), 1047 (s), 852 (s), 760 (s), 696 (s).

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Notes and references

- 1 J. Wang, M. Sánchez-Roselló, J. L. Aceña, C. del Pozo, A. E. Sorochinsky, S. Fustero, V. A. Soloshnok and H. Liu, *Chem. Rev.*, 2014, **114**, 2432.
- 2 L. N. Herrera-Rodriguez, F. Khan, K. T. Robins and H.-P. Meyer, *Chim. Oggi Chem. Today*, 2011, **29**, 31.
- 3 L. N. Herrera-Rodriguez, H.-P. Meyer, K. T. Robins and F. Khan, *Chim. Oggi Chem. Today*, 2011, **29**, 47.
- 4 C. C. C. Johansson Seechurn, M. O. Kitching, T. J. Colacot and V. Snieckus, *Angew. Chem., Int. Ed.*, 2012, **51**, 5062, (*Angew. Chem.*, 2012, **124**, 5150).
- 5 I. P. Beletskaya and A. V. Cheprakov, *Coord. Chem. Rev.*, 2004, **248**, 2337.
- 6 D. S. Surry and S. L. Buchwald, *Angew. Chem., Int. Ed.*, 2008, **47**, 6338, (*Angew. Chem.*, 2008, **120**, 6438).
- 7 A. Butler and M. Sandy, *Nature*, 2009, **460**, 848.
- 8 D. R. M. Smith, S. Gruschow and R. J. M. Goss, *Curr. Opin. Chem. Biol.*, 2013, **17**, 276.
- 9 K.-H. van Pée and E. P. Patallo, *Appl. Microbiol. Biotechnol.*, 2006, **70**, 631.
- 10 J. M. Winter and B. S. Moore, *J. Biol. Chem.*, 2009, **284**, 18577.
- 11 M. Hofrichter and R. Ullrich, *Appl. Microbiol. Biotechnol.*, 2006, **71**, 276.
- 12 A. Butler and J. V. Walker, *Chem. Rev.*, 1993, **93**, 1937.
- 13 V. Weichold, D. Milbredt and K.-H. van Pée, *Angew. Chem., Int. Ed.*, 2016, **55**, 6374, (*Angew. Chem.*, 2016, **128**, 6482).
- 14 S. A. Shepherd, C. Karthikeyan, J. Latham, A.-W. Struck, M. L. Thompson, B. R. K. Menon, M. Q. Styles, C. Levy, D. Leys and J. Micklefield, *Chem. Sci.*, 2015, **6**, 3454.
- 15 M. Frese and N. Sewald, *Angew. Chem., Int. Ed.*, 2015, **54**, 298, (*Angew. Chem.*, 2015, **127**, 302).
- 16 E. Fernández-Fueyo, M. van Wingerden, R. Renirie, R. Wever, Y. Ni, D. Holtmann and F. Hollmann, *ChemCatChem*, 2015, **7**, 4035.
- 17 N. Krause and A. S. K. Hashmi, *Modern Allene Chemistry*, Wiley-VCH, Weinheim, 1st edn, 2004.
- 18 S. Ma, *Acc. Chem. Res.*, 2003, **36**, 701.
- 19 S. Yu and S. Ma, *Chem. Commun.*, 2011, **47**, 5384.
- 20 M. Wilking, C. G. Daniliuc and U. Hennecke, *Synlett*, 2014, 1701.
- 21 C. Jonasson, A. Horváth and J.-E. Bäckvall, *J. Am. Chem. Soc.*, 2000, **122**, 9600.
- 22 M. Hammel and J. Deska, *Synthesis*, 2012, 3789.
- 23 J. Deska and J.-E. Bäckvall, *Org. Biomol. Chem.*, 2009, **7**, 3379.
- 24 J. Li, C. Fu, G. Chen, G. Chai and S. Ma, *Adv. Synth. Catal.*, 2008, **350**, 1376.



- 25 S. Ma and S. Wu, *Chem. Commun.*, 2001, 441.
- 26 B. Gockel and N. Krause, *Eur. J. Org. Chem.*, 2010, 311.
- 27 M. Poonoth and N. Krause, *Adv. Synth. Catal.*, 2009, **351**, 117.
- 28 D. H. Milles, M. Veguillas and F. D. Toste, *Chem. Sci.*, 2013, **4**, 3427.
- 29 D. Thiel, D. Doknić and J. Deska, *Nat. Commun.*, 2014, **5**, 5278.
- 30 S. Wenda, S. Illner, A. Mell and U. Kragl, *Green Chem.*, 2011, **13**, 3007.
- 31 R. A. Sheldon, *Chem. Soc. Rev.*, 2012, **41**, 1437.
- 32 A. Sánchez-Ferrer, R. Bru and F. García-Carmona, *FEBS*, 1988, **233**, 363.
- 33 M. Moniruzzaman, N. Kamiya and M. Goto, *Langmuir*, 2009, **25**, 977.
- 34 S. Kara, J. H. Schrittwieser, S. Gagiulo, Y. Ni, H. Yanase, D. J. Opperman, W. H. van Berkel and F. Hollmann, *Adv. Synth. Catal.*, 2015, **357**, 1987.
- 35 D. van der Waals, L. E. Heim, S. Vallazza, C. Gedig, J. Deska and M. H. G. Precht, *Chem. – Eur. J.*, 2016, **22**, 11568.
- 36 D. A. Mundal, K. E. Lutz and R. J. Thomson, *J. Am. Chem. Soc.*, 2012, **134**, 5782.
- 37 C. Manzuna Sapu, J.-E. Bäckvall and J. Deska, *Angew. Chem., Int. Ed.*, 2011, **50**, 9731, (*Angew. Chem.*, 2011, **123**, 9905).
- 38 C. Manzuna Sapu and J. Deska, *Org. Biomol. Chem.*, 2013, **11**, 1376.
- 39 B. Skrobo, J. D. Rolfes and J. Deska, *Tetrahedron*, 2016, **72**, 1257.

