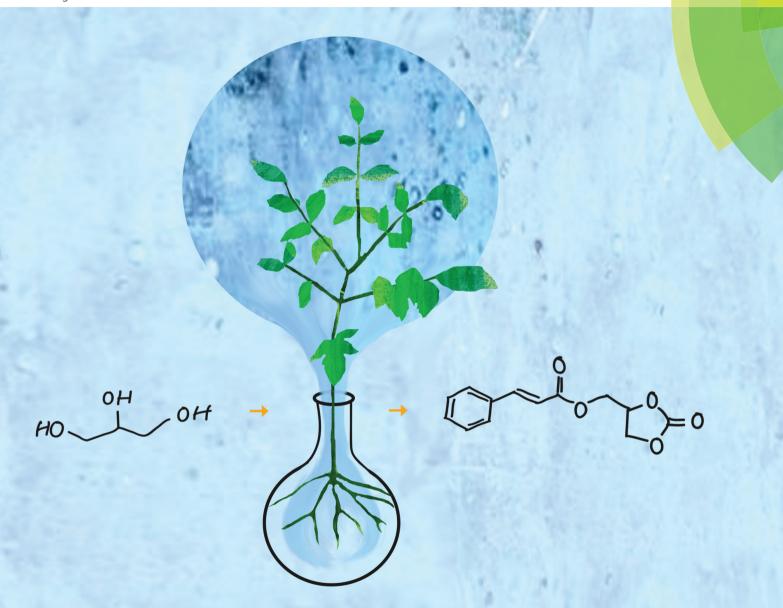
# Green Chemistry

Cutting-edge research for a greener sustainable future rsc.li/greenchem



ISSN 1463-9262



#### COMMUNICATION

H. Sundén et al.

Organocatalytic valorisation of glycerol  $\emph{via}$  a dual NHC-catalysed telescoped reaction

### **Green Chemistry**



#### COMMUNICATION

View Article Online
View Journal | View Issue



**Cite this:** *Green Chem.*, 2017, **19**, 2477

Received 13th February 2017, Accepted 16th March 2017

DOI: 10.1039/c7gc00471k

rsc.li/greenchem

## Organocatalytic valorisation of glycerol *via* a dual NHC-catalysed telescoped reaction†

A. Axelsson, A. Antoine-Michard and H. Sundén 🕩 \*

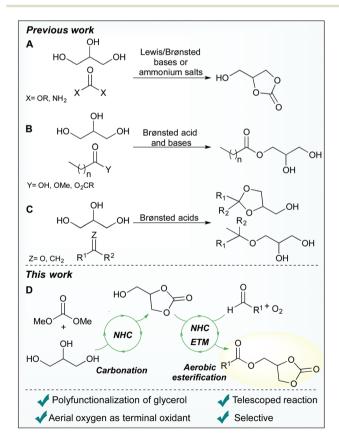
A general telescoped reaction for the NHC-catalysed carbonation and aerobic esterification of glycerol and 2-amino-2-methyl-propane-1,3-diol has been developed. The reaction provides highly functionalised glycerol derivatives in good to excellent yields (up to 95%) using low catalyst loadings and ambient conditions.

Our current dependence on fossil resources represents a major obstacle in the transition towards sustainable development.1 It is estimated that up to 95% of the carbon-containing molecules needed to sustain daily life are derived from petrochemical sources.<sup>2</sup> Hence, the valorisation of biomass into fuels, bulk and fine chemicals as well as pharmaceuticals, is of utmost importance.2-4 Moreover, in efforts to circumvent direct competition with food production the use of wastestreams instead of first generation biomass is preferred.<sup>5</sup> For this reason, glycerol is an excellent feedstock since it is obtained as a by-product from the production of biodiesel. Consequently, there is currently a large surplus of glycerol readily available.6 Conversion of glycerol into commodity and fine chemicals is typically performed using metal catalysts.<sup>7-10</sup> In contrast, organocatalytic functionalisation of glycerol is not commonly utilized. A possible explanation for this is that transition metal catalysts are typically considered to be more reactive than organocatalysts. Nevertheless, organocatalysts have many virtues such as low-cost, low-toxicity and high stability and can thus be considered as competitive alternatives to their metal counterpart. Moreover, the increasingly tough restrictions regarding metal contaminants in consumer-products further favours organocatalysis.11

Previous organocatalytic transformations of glycerol have shown that glycerol carbonate can be synthesised from glycerol and dialkyl carbonates or urea using N-heterocyclic carbenes (NHCs), quaternary ammonium salts and Brønsted-Lowry

Chemistry and Chemical Engineering, Chalmers University of Technology, Kemivägen 10, 412 96 Göteborg, Sweden. E-mail: sundenh@chalmers.se  $\dagger$  Electronic supplementary information (ESI) available: Experimental procedures and  $^1\text{H}$  NMR,  $^{13}\text{C}$  NMR and  $^{19}\text{F}$  NMR, COSY NMR, HSQC NMR, HMBC NMR, IR and HRMS data. See DOI: 10.1039/c7gc00471k

bases (Scheme 1A).<sup>12–16</sup> Additionally, acylation of glycerol is possible by transesterification of fatty methyl esters with either phosphazene- or guanidine-based catalysts, *via* Fisher esterification catalysed by sulfonic acids, and by acetate-catalysed reactions with anhydrides (Scheme 1B).<sup>17–21</sup> Both homogenous and heterogeneous sulfonic acids have also found use in the



**Scheme 1** Previous organocatalytic strategies for glycerol valorisation: transcarbonation (A), acylation (B) and acetal or ether formation (C) and the current approach combining carbonation and aerobic esterification in a telescoped reaction (D).

conversion of glycerol into a variety of ethers and acetals (Scheme 1C). 22-24

While NHC-catalysis offers a multitude of reaction paths<sup>25-28</sup> and has proven compatible with glycerol, to date, the use of NHC-catalysis in glycerol valorisation is limited to carbonation reactions. As our research interest include sustainable NHC-catalysis, we set out to merge oxidative NHC-catalysis<sup>29,30</sup> with glycerol upgrading. Instead of employing conventional methods that require stoichiometric amounts of high molecular weight oxidants, such as the Kharasch oxidant 7,31,32 an aerobic approach was envisioned, as previously reported by us and others. 33-40 More precisely, it was hypothesised that aerobic esterification of glycerol would be possible by using a biomimetic strategy comprised of electron transport mediators (ETMs), as pioneered by the Bäckvall group. 41-45 The ETMs work in concert by enabling a low energy path for electrons to flow from the substrate to oxygen (O2), circumventing the unfavourable reaction kinetics associated with direct O<sub>2</sub>-oxidations. 45 Moreover, it was predicted that a dual functionalisation of glycerol, comprising a sequential NHC-catalysed carbonation followed by an aerobic esterification, would be possible by using a telescoped reaction strategy (Scheme 1D). This approach offers several advantages including: direct valorisation of glycerol, high atom economy, improved poteconomy, 46 and the incorporation of the carbonate group as a valuable synthetic handle.12

An initial experiment with a telescoped approach that included a one-pot sequential carbonation, using dimethyl carbonate (DMC), and aerobic esterification was successful providing the unsaturated carbonate (1) in 89% yield (Table 1, entry 1). It should be noted that dimethyl carbonate is considered a benign reagent as it can be obtained from CO2, is nontoxic and biodegradable. 47 A survey of different NHC-catalysts showed that the thiazolium based catalyst 5 was inactive (entry 3), while 4 and 6 provided 1 in slightly lower yields compared to when 3 was employed (entries 2 and 4). This is due to increased carboxylic acid formation and incomplete consumption of cinnamaldehyde, respectively. The choice of base also proved important and the use of K2CO3 or Et3N in place of 1,5,7-triazabicyclo[4.4.0]dec-5-ene (TBD) resulted in no conversion of starting material (entries 5 and 6). This effect can be rationalised by the fact that the essential in situ FePc-catalysed aerobic oxidation of 8 to 7 is also base-catalysed.31 Thus, by replacing 8 with 7 the loading of TBD can be decreased from 0.5 to 0.2 equivalents with maintained yield (entry 7), albeit with prolonged reaction time (24 h). Moreover, when the reaction is performed under a nitrogen atmosphere 1 is obtained in only 5%, showcasing that aerial oxygen truly is the terminal oxidant (entry 8). Omission of the NHC-precatalyst in the carbonation step results in the incomplete conversion of glycerol with subsequent side product formation, such as, 2, the secondary- and the diester (14% determined by <sup>1</sup>H NMR) suggesting that this step is also NHC-catalyzed. Synthesis of 2 via a direct aerobic esterification of glycerol, omitting the carbonation step, was possible and 2 was obtained in 87% yield (entry 9). However, due to the formation of side products (the

**Table 1** Optimization of reaction conditions for the dual functionalisation of glycerol

Entry	Cat	ETM	ETM'	Base	Solvent	$Yield^{a}$ (%)	
						1	2
$\overline{1^b}$	3	8	FePc	TBD	MeCN	89	
$2^b$	4	8	FePc	TBD	MeCN	86	
$3^b$	5	8	FePc	TBD	MeCN	0	
$4^b$	6	8	FePc	TBD	MeCN	81	
$5^b$	3	8	FePc	$K_2CO_3$	MeCN	0	
$6^b$	3	8	FePc	Et <sub>3</sub> N	MeCN	0	
$7^c$	3	7	FePc	TBD	MeCN	88	
$8^{b,d}$	3	8	FePc	TBD	MeCN	5	
$9^e$	3	8	FePc	TBD	MeCN		87
$10^e$	3	8	FePc	TBD	Acetone		81

 $^a$  Determined by  $^1\mathrm{H}$  NMR with durene as internal standard.  $^b$  i, Glycerol (1.1 eq.), TBD (0.5 eq.), NHC (2 mol%), solvent/dimethyl carbonate 5:2. ii, Cinnamaldehyde (0.5 mmol), FePc (0.5 mol%), 8 (2 mol%).  $^c$  As in footnote b, but with 0.2 eq. TBD and 0.01 eq. of 7.  $^d$  Last step performed under a nitrogen atmosphere.  $^e$  Glycerol (4 eq.), TBD (0.5 eq.), 3 (2 mol%), cinnamaldehyde (0.5 mmol), FePc (0.5 mol%), 8 (2 mol%).

secondary ester and the diester, ca. 10%), it was not possible to obtain the pure product in a satisfactory manner. Regardless of concentration and solvent selection the amount of side products remained around 10%. For example, when acetone was used as the reaction solvent a competing aldol reaction was observed and 2 was obtained in 81% yield (entry 10).

Having successfully identified mild conditions (Table 1 entry 1) that enabled selective dual functionalisation of glycerol with aerial oxygen as the terminal oxidant, the scope of this transformation was investigated (Scheme 2). The reaction proved general, and both electron-donating (9–11, 16) and electron-withdrawing (12–15) substituents were tolerated on the cinnamaldehyde scaffold. Compound 9, isolated in 95% yield, is worth highlighting since 4-methoxycinnamates are commonly used in sunscreens and the glycerol ester has been investigated as a more benign sunscreen agent. Moreover, polyaromatics were well tolerated and anthracene ester 17 could be obtained in 76% yield. Initial reactions with aliphatic enals were sluggish. However, by using 4 mol% of the more active catalyst 4 2-hexenal yielded the corresponding ester 18 in 64% yield. Benzaldehydes proved less reactive than

**Green Chemistry** Communication

Scheme 2 The scope of the aerobic glycerol functionalisation. <sup>a</sup> (i) Glycerol (1.1 eq.), TBD, 3, MeCN/DMC 5:2. (ii) Aldehyde (0.5 mmol), FePc (0.5 mol%), **8** (2 mol%). <sup>b</sup> 1.6 grams isolated. <sup>c</sup> 4 mol% of catalyst **4** used. d 4 mol% of catalyst 3 used.

cinnamaldehydes and required 4 mol% (instead of 2 mol%) of catalyst 3 to react efficiently. With the increase in catalyst loading both electron rich and electron poor aromatic as well as heteroaromatic aldehydes worked well, affording the corresponding products (19-25) in moderate to good yields (51-79%).

For instance, piperonal derived ester 23 could be obtained in 79% yield. Moreover, this reaction is scalable and α,β-unsaturated ester 1 could be obtained in 87% yield and 1.6 grams.

Encouraged by the successful functionalisation of glycerol, the compatibility of the developed system with the aminoalcohol 2-amino-2-methyl-1,3-propanediol (26) was investigated (Scheme 3). The approach was successful and both electron rich and electron poor enals could be transformed into the corresponding 2-oxooxazolidine esters in excellent yields. For instance, cinnamaldehyde and 4-chlorocinnamaldehyde both reacted well and provided the corresponding α,β-unsaturated esters in 92% and 91% yield respectively (27 and 30).

Ring opening of glycerol carbonate and its derivatives have obtained a lot of attention, for example, in sustainable synthesis of pharmaceuticals and polymers. 9,49,50 Consequently, the herein developed procedure could generate several new glycerol carbonate building blocks for use as green monomers or for further manipulation. With this in mind, the nucleophilic ring opening of the 2-oxo-1,3-dioxolan group with piperidine was investigated (Scheme 4). Inspired by reports from Kleij et al., 51,52 it was possible to selectively activate the less electrophilic carbonate in the presence of the unsaturated ester using 5 mol% of TBD, vielding acyclic carbamate 31 in 83% vield and good regioselectivity (Scheme 4). Usage of higher loadings of TBD or other solvents (MeCN, THF) leads to formation of the amide as a side product.

The proposed catalytic cycle starts with addition of the NHC (I) to aldehyde II forming the Breslow intermediate (III) (Scheme 5). Intermediate III is oxidized by the coupled system of ETMs to the acyl azolium IV. Glycerol carbonate (V), formed by NHC/TBD catalysis in the first step of the telescoped reac-

Scheme 3 Selective modification of aminopolyol 26 by telescoped carbamate formation and aerobic esterification. (i) 26 (1.0 eq.), 3, TBD, MeCN/DMC 5: 2. (ii) Aldehyde (0.5 mmol), FePc, 8, isolated yields.

Scheme 4 Organocatalytic ring opening of 1 to form acyclic carbamate 31. 1 (0.2 mmol), piperidine (1.2 eq.), isolated yield, regioisomeric ratio determined by <sup>1</sup>H NMR of crude material, major isomer separable by flash chromatography.

Communication **Green Chemistry** 

Scheme 5 Proposed catalytic cycle.

tion, adds as a nucleophile to IV forming the product (VI) and regenerating the NHC.

In summary, a telescoped protocol for the NHC-catalysed carbonation and aerobic esterification of glycerol has been developed. The presented method gives access to highly functionalised glycerol derivatives from sustainable resources such as glycerol, dimethyl carbonate and aerial oxygen. The reaction occurs under ambient conditions, is high vielding, scalable and has a broad scope that tolerates (hetero)aromatic aldehydes, aromatic and aliphatic  $\alpha,\beta$ -unsaturated aldehydes. The method could also be further extended to amino-polyols, affording 2-oxooxazolidine esters in excellent yields. Lastly, chemoselective activation of the less electrophilic carbonate was demonstrated, showcasing the use of the obtained products. These results demonstrate the possibility to extend the role of NHC-catalysis in glycerol valorisation beyond carbonation reactions, enabling access to new sustainable building blocks for chemical synthesis.

Funding from the Swedish Research Council (VR and Formas) are gratefully acknowledged. Furthermore, we thank the C. F. Lundström foundation for supporting the running costs of this project.

#### References

- 1 G. G. Brundtland, Our common future, The World Commision on Environmental Development, Oxford, 1987.
- 2 C. H. Christensen, J. Rass-Hansen, C. C. Marsden, E. Taarning and K. Egeblad, ChemSusChem, 2008, 1, 283-
- 3 R. A. Sheldon, Green Chem., 2014, 16, 950-963.
- 4 A. Corma, S. Iborra and A. Velty, Chem. Rev., 2007, 107, 2411-2502.
- 5 J. H. Clark, J. Chem. Technol. Biotechnol., 2007, 82, 603-609.
- 6 M. Pagliaro and M. Rossi, The Future of Glycerol, Royal Society of Chemistry, Cambridge. U.K, 2nd edn, 2010.
- 7 C.-H. Zhou, J. N. Beltramini, Y.-X. Fan and G. Q. Lu, Chem. Soc. Rev., 2008, 37, 527-549.

- 8 A. Said Stålsmeden, J. L. Belmonte Vázquez, K. van Weerdenburg, R. Rae, P.-O. Norrby and N. Kann, ACS Sustainable Chem. Eng., 2016, 4, 5730-5736.
- 9 A. M. Truscello, C. Gambarotti, M. Lauria, S. Auricchio, G. Leonardi, S. U. Shisodia and A. Citterio, Green Chem., 2013, 15, 625-628.
- 10 J. Bensemhoun and S. Condon, Green Chem., 2012, 14, 2595-2599.
- 11 G. Fiorani, W. Guo and A. W. Kleij, Green Chem., 2015, 17, 1375-1389.
- 12 M. O. Sonnati, S. Amigoni, E. P. Taffin de Givenchy, T. Darmanin, O. Choulet and F. Guittard, Green Chem., 2013, 15, 283-306.
- 13 J. A. Stewart, R. Drexel, B. Arstad, E. Reubsaet, B. M. Weckhuysen and P. C. A. Bruijnincx, Green Chem., 2016, 18, 1605-1618.
- 14 P. U. Naik, L. Petitjean, K. Refes, M. Picquet and L. Plasseraud, Adv. Synth. Catal., 2009, 351, 1753-1756.
- 15 H. Mutlu, J. Ruiz, S. C. Solleder and M. A. R. Meier, Green Chem., 2012, 14, 1728-1735.
- 16 J. R. Ochoa-Gomez, O. Gomez-Jimenez-Aberasturi, C. Ramirez-Lopez and B. Maestro-Madurga, Green Chem., 2012, 14, 3368-3376.
- 17 G. Kharchafi, F. Jerome, J.-P. Douliez and J. Barrault, Green Chem., 2006, 8, 710-716.
- 18 F. Jerome, G. Kharchafi, I. Adam and J. Barrault, Green Chem., 2004, 6, 72-74.
- 19 J. n. Pérez-Pariente, I. Díaz, F. Mohino and E. Sastre, Appl. Catal., A, 2003, 254, 173-188.
- 20 X. Liu, H. Ma, Y. Wu, C. Wang, M. Yang, P. Yan and U. Welz-Biermann, Green Chem., 2011, 13, 697-701.
- 21 B. Ren, M. Rahm, X. Zhang, Y. Zhou and H. Dong, J. Org. Chem., 2014, 79, 8134-8142.
- 22 K. Klepáčová, D. Mravec and M. Bajus, Appl. Catal., A, 2005, 294, 141-147.
- 23 J. Deutsch, A. Martin and H. Lieske, J. Catal., 2007, 245, 428-435.
- 24 A. Behr, J. Eilting, K. Irawadi, J. Leschinski and F. Lindner, Green Chem., 2008, 10, 13-30.
- 25 D. Enders, O. Niemeier and A. Henseler, Chem. Rev., 2007, 107, 5606-5655.
- 26 N. Marion, S. Díez-González and S. P. Nolan, Angew. Chem., Int. Ed., 2007, 46, 2988-3000.
- 27 X. Bugaut and F. Glorius, Chem. Soc. Rev., 2012, 41, 3511-3522.
- 28 D. M. Flanigan, F. Romanov-Michailidis, N. A. White and T. Rovis, Chem. Rev., 2015, 115, 9307-9387.
- 29 C. E. I. Knappke, A. Imami and A. J. von Wangelin, ChemCatChem, 2012, 4, 937-941.
- 30 S. De Sarkar, A. Biswas, R. C. Samanta and A. Studer, Chem. - Eur. J., 2013, 19, 4664-4678.
- 31 M. S. Kharasch and B. S. Joshi, J. Org. Chem., 1957, 22, 1439-1443.
- 32 S. D. Sarkar, S. Grimme and A. Studer, J. Am. Chem. Soc., 2010, 132, 1190-1191.

33 R. S. Reddy, J. N. Rosa, L. F. Veiros, S. Caddick and P. M. P. Gois, *Org. Biomol. Chem.*, 2011, 9, 3126–3129.

**Green Chemistry** 

- 34 E. G. Delany, C.-L. Fagan, S. Gundala, A. Mari, T. Broja, K. Zeitler and S. J. Connon, *Chem. Commun.*, 2013, 49, 6510–6512.
- 35 E. G. Delany, C.-L. Fagan, S. Gundala, K. Zeitler and S. J. Connon, *Chem. Commun.*, 2013, **49**, 6513–6515.
- 36 J. Zhao, C. Mück-Lichtenfeld and A. Studer, *Adv. Synth. Catal.*, 2013, 355, 1098–1106.
- 37 L. Ta, A. Axelsson and H. Sunden, *Green Chem.*, 2016, **18**, 686–690.
- 38 A. Axelsson, E. Hammarvid, L. Ta and H. Sunden, *Chem. Commun.*, 2016, 52, 11571–11574.
- 39 D. Xie, D. Shen, Q. Chen, J. Zhou, X. Zeng and G. Zhong, *J. Org. Chem.*, 2016, **81**, 6136–6141.
- 40 A. Axelsson, L. Ta and H. Sunden, *Synlett*, 2017, DOI: 10.1055/s-0036-1588395.
- 41 J. E. Bäckvall, A. K. Awasthi and Z. D. Renko, *J. Am. Chem. Soc.*, 1987, **109**, 4750–4752.
- 42 J.-E. Bäckvall, R. B. Hopkins, H. Grennberg, M. Mader and A. K. Awasthi, *J. Am. Chem. Soc.*, 1990, **112**, 5160–5166.

- 43 J. Piera, K. Närhi and J.-E. Bäckvall, Angew. Chem., Int. Ed., 2006, 45, 6914–6917.
- 44 J. Wöltinger, J.-E. Bäckvall and Á. Zsigmond, *Chem. Eur. I.*, 1999, 5, 1460–1467.
- 45 J. Piera and J.-E. Bäckvall, Angew. Chem., Int. Ed., 2008, 47, 3506–3523.
- 46 Y. Hayashi, Chem. Sci., 2016, 7, 866-880.
- 47 P. Tundo and M. Selva, Acc. Chem. Res., 2002, 35, 706-716.
- 48 R. A. Holser, T. R. Mitchell, R. E. Harry-O'kuru, S. F. Vaughn, E. Walter and D. Himmelsbach, *J. Am. Oil Chem. Soc.*, 2008, **85**, 347–351.
- 49 C. Duval, N. Kébir, R. Jauseau and F. Burel, *J. Polym. Sci.*, *Part A: Polym. Chem.*, 2016, 54, 758–764.
- 50 G. Rokicki, P. Rakoczy, P. Parzuchowski and M. Sobiecki, *Green Chem.*, 2005, 7, 529–539.
- 51 W. Guo, J. Gónzalez-Fabra, N. A. G. Bandeira, C. Bo and A. W. Kleij, *Angew. Chem., Int. Ed.*, 2015, 54, 11686–11690.
- 52 S. Sopeña, V. Laserna, W. Guo, E. Martin, E. C. Escudero-Adán and A. W. Kleij, *Adv. Synth. Catal.*, 2016, 358, 2172–2178.