



**Cite this:** *Green Chem.*, 2025, **27**, 4995

Received 27th January 2025,  
Accepted 8th April 2025

DOI: 10.1039/d5gc00499c

[rsc.li/greenchem](https://rsc.li/greenchem)

## Supramolecular assisted *O*-acylation of carbohydrates†

Soumyadip Dey,  Debabrata Giri,  Adrita Nandy  and Abhijit Sau  \*

The acylation of hydroxy groups serves as one of the most employed protecting group strategies in carbohydrate chemistry. Here, we present a base-free, supramolecular assisted approach for the *O*-acylation of carbohydrates under mild conditions, using 18-crown-

6 in combination with a catalytic amount of potassium fluoride. This sustainable and useful method successfully converted various functional groups containing carbohydrate hydroxy groups into *O*-acyl, *O*-benzoyl, and *O*-propionyl derivatives with up to 99% yields.

### Green foundation

1. In this manuscript, we describe an efficient and green method *via* supramolecular assisted *O*-acylation of carbohydrate alcohols. This approach removes the use of hazardous and toxic pyridine and its derivatives, which are harmful to the ecosystem. Moreover, it is efficient and eco-friendly for acylation, including aliphatic and phenolic hydroxy groups under neat conditions, without using any complicated setups or inert gasses.
2. This method requires very cheap and easily available chemicals: potassium fluoride (KF) and 18-crown-6. Our cost-effective acylation method achieves a very low E-factor (0.6) compared to other existing acylation techniques, which makes it more efficient and sustainable for synthesizing acylating substrates.
3. The long reaction time and the requirement of 40 °C for effective acylation are minor drawbacks of this synthetic method. However, we have confidence that this work provides a significant proof of concept, establishing supramolecular assisted acylation under mild conditions.

Alcohols,<sup>1</sup> amines,<sup>2</sup> phenols,<sup>3</sup> and thiols<sup>4</sup> are often acylated for functional group transformations in organic synthesis.<sup>5–9</sup> In carbohydrate chemistry, *O*-acetylation is one of the most commonly used techniques for protecting hydroxyl groups.<sup>10–12</sup> Moreover, per-*O*-acetylated sugars are affordable and practically valuable intermediates for the synthesis of natural glycosides, glycoconjugates, and other oligosaccharides.<sup>13</sup> Additionally, acylation is also useful in structural elucidations of various carbohydrate moieties containing natural products by converting them into their per-*O*-acetates.<sup>14</sup> Apart from acetyl protection, benzoyl, pivaloyl, and chloroacetyl are also used in acyl-protecting groups during carbohydrate transformation. Generally, these common acyl protected sugars are synthesized with acetic anhydride or acetyl chloride, benzoyl chloride, pivaloyl chloride, and chloroacetyl chloride, respectively, in the presence of a base.<sup>15</sup> Acetylation of sugar alcohols has been frequently accomplished with a significant excess of

acetic anhydride in the presence of pyridine, acting as the base and solvent, despite its toxicity and disagreeable odour.<sup>16</sup> In addition, different pyridine derivatives, like 4-(1-pyrrolidino)pyridine and 4-(*N,N*-dimethylaminopyridine) (DMAP), are largely used as co-catalysts to accelerate the reaction rate.<sup>17,18</sup> Base-catalyzed acylation<sup>19</sup> is often performed in the presence of imidazole or its derivatives.<sup>13,20,21</sup> However, the preferred synthetic procedure for stereoselective synthesis of anomeric  $\beta$ -glycosyl acetates is Wolfrom's anhydrous sodium acetate method.<sup>22,23</sup> In the literature, acid-catalyzed acylation has been reported using different Lewis acids such as In(OTf)<sub>3</sub>,<sup>24</sup> InCl<sub>3</sub>,<sup>25</sup> ZnCl<sub>2</sub>,<sup>26</sup> FeCl<sub>3</sub>,<sup>27</sup> Fe<sub>2</sub>(SO<sub>4</sub>)<sub>3</sub>,<sup>28</sup> iodine,<sup>29,30</sup> Cu(OTf)<sub>2</sub>,<sup>31</sup> Sc(OTf)<sub>3</sub>,<sup>32</sup> BiOCl–SOCl<sub>2</sub>,<sup>33</sup> CoCl<sub>2</sub>,<sup>34</sup> BiCl<sub>3</sub>,<sup>35</sup> LiClO<sub>4</sub>,<sup>36</sup> *etc.* Perchloric acid,<sup>37</sup> sulfuric acid,<sup>38</sup> sulfamic acid,<sup>10</sup> and HClO<sub>4</sub>–SiO<sub>2</sub><sup>39</sup> have also been used in combination with acetic anhydride in various acetylations.<sup>40</sup> In addition to these classical methods, various heterogeneous catalysts have been employed for acetylation, such as zeolites,<sup>41</sup> montmorillonite K-10,<sup>14</sup> and Nafion-H.<sup>42</sup> Although different protocols, such as microwave irradiation conditions<sup>25,26,43,44</sup> ionic liquids,<sup>45</sup> and carbene-mediated<sup>46–49</sup> selective acylation<sup>50</sup> of saccharides, have also been investigated, these methods often face several challenges,

Department of Chemistry, Indian Institute of Technology Hyderabad, Kandi, 502284 Sangareddy, Telangana, India. E-mail: [asau@chy.iith.ac.in](mailto:asau@chy.iith.ac.in)

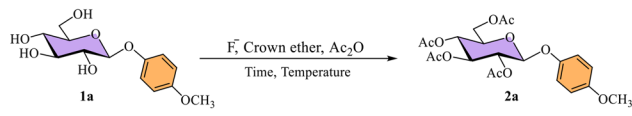
† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d5gc00499c>



including excessive use of acylating reagents, excessive environmental toxicity, moisture sensitivity, preparation of catalysts, incompatibility with acid-sensitive groups, high costs, handling the large amount of toxic pyridine and isomerization of unprotected reducing carbohydrate derivatives into their furanose and pyranose forms. To overcome these limitations, a quick, environmentally friendly, and clean reaction protocol that minimizes the need for traditional workup and purification processes is still desired for acetylation of carbohydrates. We sought a mild, efficient, non-toxic, green, and sustainable catalytic method for acylation. On the other hand, crown ethers have gained significant importance in the field of organic synthesis in recent years for numerous applications such as metal ion separations, phase transfer catalysis, metal-organic frameworks, and electron transfer reactions.<sup>51,52</sup> Various types of crown ethers,<sup>53</sup> such as 12-crown-4, 15-crown-5, 18-crown-6, *etc.*, have been developed, and their own central hole size correlate with the diameters of different cations. The exceptional chemical property of excellent capacity to encapsulate metal ions from their salts with stability and high specificity makes crown ethers unique and significant in their applications.<sup>54</sup> Furthermore, the strong interaction ability of crown ethers with metal cations has inspired the development of catalytic properties within the reaction mixture by activating the metal salts through enhancing their solubility.<sup>55</sup> Several efforts have been made in the past for the acylation of carbohydrates, but still, there is a high demand for a less hazardous, cheaper, solvent-free new method for the synthesis of acyl-protected sugars that can avoid the extreme toxicity of pyridine. Keeping in mind the exciting properties of crown ethers, we envisioned exploring the catalytic ability of a metal salt-crown ether combined system for acylation in sugar molecules (Scheme 1).

In the preliminary screening, we started with four free hydroxyl groups containing the substrate 4-methoxyphenyl- $\beta$ -D-glucopyranoside (**1a**) for the acylation reaction. During the first set of reactions, compound **1a** was treated with 0.4 equiv. (0.1 equiv. per -OH) of potassium fluoride and 4.5 equiv. of acetic anhydride in acetonitrile, resulting in a trace amount of spot-on TLC at a temperature of 50 °C (Table 1, entry 1). After

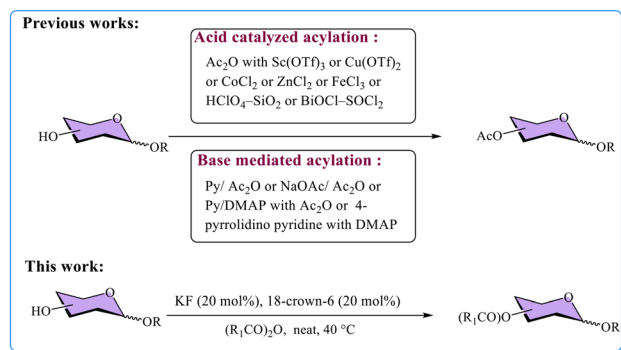
**Table 1** Optimisation of the acylation reaction<sup>a</sup>

					
Entry	Reagent (equiv. per -OH)	Temp (°C)	Solvent	Time (hours)	Yield <sup>b</sup> (%)
1	KF (0.1)	50 °C	MeCN	12 h	Trace
2	KF (0.1), 18-C-6 (0.1)	25 °C	Neat	12 h	43
3	KF (0.2), 18-C-6 (0.2)	25 °C	Neat	12 h	68
4	KF (0.2), 18-C-6 (0.2)	25 °C	DCM	12 h	71
5	<b>KF (0.2), 18-C-6 (0.2)</b>	<b>40 °C</b>	<b>Neat</b>	<b>12 h</b>	<b>94</b>
6	NaF (0.2)	50 °C	MeCN	12 h	5
7	NaF (0.2), 15-C-5 (0.2)	25 °C	Neat	12 h	43
8	NaF (0.2), 15-C-5 (0.2)	40 °C	Neat	12 h	58
9	CsF (0.2)	50 °C	MeCN	6 h	8
10	KF (0.2), 18-C-6 (0.2)	70 °C	Neat	6 h	92

<sup>a</sup> Reaction conditions: substrate **1a** (0.34 mmol) and acetic anhydride (1.61 mmol). <sup>b</sup> Isolated yield.

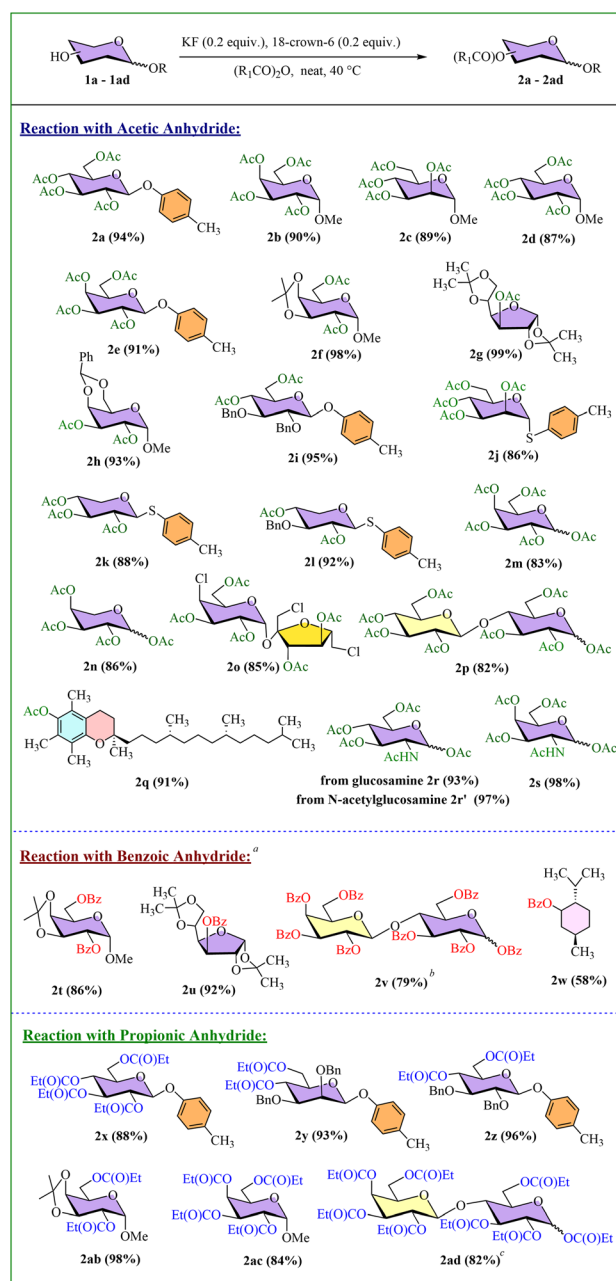
that, the use of a combination of 0.1 equiv. potassium fluoride and crown ether (0.1 equiv. per -OH) made a noticeable improvement in the formation of the desired product peracetylated 4-methoxyphenyl- $\beta$ -D-glucopyranoside (**2a**), yielding 43% under neat conditions after 12 hours (Table 1, entry 2). Next, increasing the loading of both potassium fluoride and crown ether to 0.8 equiv. (0.2 equiv. per -OH) at 25 °C under solvent free and DCM conditions after 12 hours resulted in good yields of our desired per-acetylated product **2a**, which were 68% and 71%, respectively (Table 1, entries 3 and 4). However, the yield was enhanced to 94% by increasing the temperature to 40 °C, while maintaining the same reagent and catalyst loading under neat conditions (Table 1, entry 5). Subsequently, we also explored the catalytic amount of other metal fluorides like sodium fluoride (NaF) and cesium fluoride (CsF), which resulted in 5% and 8% of desired compound **2a**, respectively, at 40 °C (Table 1, entries 6 and 9). Moreover, reacting compound **1a** with a combination of 0.8 equiv. of NaF and 15-crown-5 (0.2 equiv. per -OH) afforded up to 58% of peracetylated 4-methoxyphenyl- $\beta$ -D-glucopyranoside (**2a**) (Table 1, entries 7 and 8). In addition, we noticed that at 70 °C within 6 hours all starting materials were consumed and resulted in a similar isolated yield (Table 1, entry 10). After this extensive screening, we concluded that using catalytic amounts of potassium fluoride and 18-crown-6 (0.2 equiv. per -OH) with acetic anhydride (1.15 equiv. per -OH) at 40 °C under neat conditions is the best condition for the acylation reaction.

Optimizing the standard reaction conditions, we further explored the substrate scope with various sugar molecules as well as by employing different acylating anhydrides (Scheme 2). Initially, we focused on the acylation reaction of methyl- $\alpha$ -D-galactopyranoside (**1b**), methyl- $\alpha$ -D-mannopyranoside (**1c**), and methyl- $\alpha$ -D-glucopyranoside (**1d**) with acetic anhydride under the optimized reaction conditions and obtained the corresponding peracetylated products (**2b**), (**2c**), and (**2d**) with excellent yields of 90%, 89%, and 87%, respect-



**Scheme 1** Previous studies and this work on O-acylation of carbohydrates.





**Scheme 2** Substrate scope of acylation with various anhydrides.

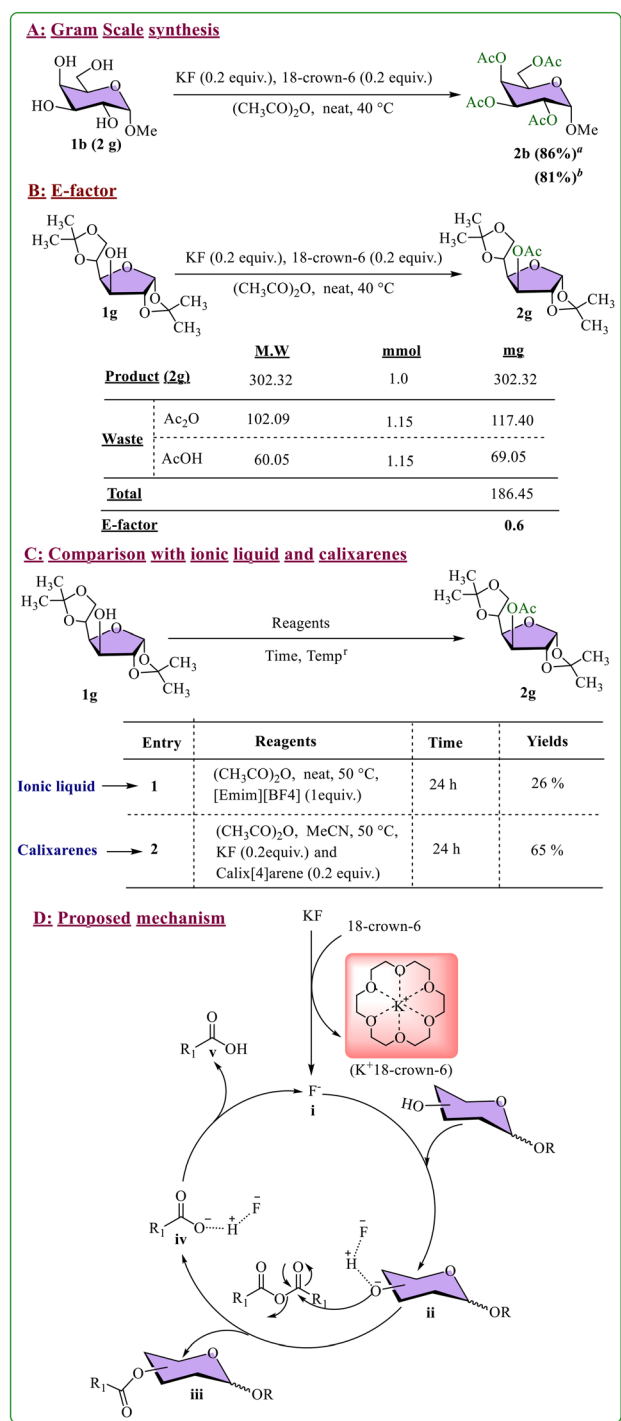
<sup>a</sup>Reaction conditions: acetonitrile was used as a solvent, <sup>b</sup>reaction was performed at 60 °C and 12 equiv. of Bz<sub>2</sub>O was used, and <sup>c</sup>reaction was performed at 60 °C for 20 h.

Next, acylation of 4-methoxyphenyl-β-D-galactopyranoside (**1e**) also resulted in an excellent yield of the desired peracetylated compound **2e** (91%). After that, we aimed for acylation of different protected sugars such as methyl 3,4-*O*-isopropylidene-α-D-galactopyranoside (**1f**) and 3-hydroxy-1,2:5,6-di-*O*-isopropylidene-α-D-glucopyranoside (**1g**) with acetic anhydride and observed excellent yields of desired acetylated products **2f** (98%) and **2g** (99%), respectively. Similarly, methyl 4,6-*O*-benzylidene-α-D-galactopyranoside (**1h**) was successfully converted to the acetylated product (**2h**) with 93% yield. These results

suggest that acid-sensitive isopropylidene and benzylidene rings can tolerate the reaction conditions easily. Then, we obtained an excellent yield of the compound *p*-methoxyphenyl 2,3-di-*O*-benzyl-4,6-*O*-acetyl-β-D-glucopyranoside (**2i**, 95%) under the same reaction conditions. Besides, different thioglycosides, *p*-methoxyphenyl 1-thio-α-D-mannopyranoside (**1j**), *p*-methoxyphenyl 1-thio-β-D-xylopyranoside (**1k**) and *p*-methoxyphenyl 3-*O*-benzyl-1-thio-β-D-xylopyranoside (**1l**) were smoothly converted to the respective acylated products with good yields (**2j**, 86%; **2k**, 88%; and **2l**, 92%). Next, we attempted the acetylation of free sugar molecules such as galactose (**1m**) and arabinose (**1n**), and interestingly, observed impressive yields in both cases (83% and 86%, respectively). In addition, disaccharides sucralose (**1o**) and cellobiose (**1p**) were also smoothly converted to the per-acetylated form with good yields (**2o**, 85% and **2p**, 82%) under the optimal reaction conditions. Furthermore, we also extended the acylation of the phenolic hydroxyl group of the essential natural product tocopherol (**1q**) and obtained an excellent yield (**2q**, 91%). Additionally, we expanded the substrate scope towards amino sugars such as glucosamine (**1u**), N-acetylglucosamine (**1v**) and galactosamine (**1w**) and obtained per-acetylated products **2r**, **2r'** and **2s** with excellent yields (93%, 97% and 98%, respectively, Scheme 2). To showcase the utility of this green approach, we further investigated acylation with different acylating reagents. Hence, compound **1f** was reacted with 0.4 equiv. of benzoic anhydride in acetonitrile, resulting in the benzoylated product (**2t**) with a good yield (86%) under standard reaction conditions. Then, we executed a reaction of compound **1g** with benzoic anhydride, which resulted in the corresponding benzoyl-protected product **2u** with excellent yield (92%). In addition, disaccharide lactose and aliphatic alcohol 1-menthol were easily converted to their corresponding benzoylated products **2v** and **2w** with good yields – 79% and 58%, respectively. We also observed good to excellent yields of 82% to 98% while extending the acylation scope with propionic anhydride and different sugar molecules, including disaccharide lactose (Scheme 2; **2x–2ad**).

After exploring the substrate scope, we planned gram-scale synthesis of per-acetylated carbohydrates under the optimized reaction conditions. For this purpose, compound **1b** was easily converted to per-acetylated product **2b** with a good yield (86% in a 2 g scale). Moreover, we recovered the crown ether from the reaction mixture following the reported work-up procedure<sup>54</sup> (details in the ESI†) and utilized it in the further acylation reaction that provided a similar yield to the desired per-acetylated product (81%). A commonly used indicator for the quantitative assessment of a synthetic reaction method's environmental impact is called the E-factor.<sup>56,57</sup> The calculated E-factor of our supramolecular assisted acylation reaction was 0.6 (Scheme 3B). In addition, we checked the reaction in an ionic liquid ([Emim][BF<sub>4</sub>]) and calixarene (Calix[4]arene) and noticed 26% and 65% yields of compound **2g**, respectively (Scheme 3C). The most probable mechanistic pathway is displayed in Scheme 3, where initially, crown ether encapsulates the potassium ion (K<sup>+</sup>) in the cavity, making free fluoride ions (F<sup>−</sup>) more nucleophilic in the reaction mixture. Then, F<sup>−</sup>,





**Scheme 3** (A) Gram-scale synthesis: <sup>a</sup>fresh catalyst loading and <sup>b</sup>reused catalyst. (B) Calculation of the E-factor, (C) comparison with an ionic liquid and calixarenes and (D) proposed mechanism of supramolecular assisted acylation.

which abstracts the proton of sugar alcohols to form intermediate (ii), further reacts with the anhydride to yield the desired acylated product (iii). Next, R<sub>1</sub>COO<sup>−</sup> protonates in the medium to generate fluoride ions again to continue the catalytic cycle (Scheme 3D).

## Conclusions

In conclusion, we have developed an efficient and practical method for acylation by using catalytic amounts of potassium fluoride (KF) and 18-crown-6 under solvent-free conditions, avoiding toxic pyridine and its derivatives. This supramolecular assisted approach was successfully applied to a wide range of carbohydrates and aliphatic alcohols, providing straightforward access to their acetyl, benzoyl, and propionyl derivatives. Importantly, functional groups commonly used for protecting and manipulating carbohydrates remained unaffected under the reaction conditions. Moreover, these mild reaction conditions are convenient, non-hazardous, and eco-friendly. We believe that this greener approach will be useful for the preparation of several common acylated carbohydrate building blocks in organic synthesis.

## Author contributions

SD: conceptualisation, methodology, validation, formal analysis, data curation and writing (original draft); DG: methodology, validation and formal analysis; AN: methodology, validation and formal analysis. AS: conceptualisation, funding acquisition, supervision, project administration, reviewing, and editing.

## Data availability

The data that support the findings of this study are available in the ESI.†

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

The authors appreciate the financial support from SERB and MoE-STARS, India (SRG/2023/000034 & MoE-STARS/STARS-2/2023-0126) and the Indian Institute of Technology Hyderabad (IITH). Instrumental support was provided by the IIT Hyderabad.

## References

- 1 A. Orita, C. Tanahashi, A. Kakuda and J. Otera, *Angew. Chem., Int. Ed.*, 2000, **39**, 2877–2879.
- 2 F. Piazzolla and A. Temperini, *Tetrahedron Lett.*, 2018, **59**, 2615–2621.
- 3 T. G. Bonner and P. McNamara, *J. Chem. Soc. B*, 1968, 795–797.





- 4 A. Temperini, D. Annesi, L. Testaferri and M. Tiecco, *Tetrahedron Lett.*, 2010, **51**, 5368–5371.
- 5 N. Anbu, N. Nagarjun, M. Jacob, J. M. V. K. Kalaiarasi and A. Dhakshinamoorthy, *Chemistry*, 2019, **1**, 69–79.
- 6 Z. Zhang, F. Cheng, X. Ma, K. Sun, X. Huang, J. An, M. Peng, X. Chen and B. Yu, *Green Chem.*, 2024, **26**, 7331–7336.
- 7 Z. Zhang, Z. Zhao, M. Liu, H. Liu, Q. Li, J. Xiang, T. Wu and B. Han, *Green Chem.*, 2022, **24**, 9763–9771.
- 8 Z. Zhong, H. Wu, X. Chen, Y. Luo, L. Yang, X. Feng and X. Liu, *J. Am. Chem. Soc.*, 2024, **146**, 20401–20413.
- 9 C. Li, J. Cheng, X. Wan, J. Li, W. Zu, Y. Xu, Y. Huang and H. Huo, *J. Am. Chem. Soc.*, 2024, **146**, 19909–19918.
- 10 A. Santra, G. Guchhait and A. K. Misra, *Green Chem.*, 2011, **13**, 1345–1351.
- 11 S. Dey, S. Dwivedi, D. Giri and A. Sau, *Org. Lett.*, 2024, **26**, 10351–10355.
- 12 S. Dey, S. Dwivedi and A. Sau, *Carbohydr. Res.*, 2025, **552**, 109447.
- 13 P. Tiwari, R. Kumar, P. R. Maulik and A. K. Misra, *Eur. J. Org. Chem.*, 2005, **2005**, 4265–4270.
- 14 P. M. Bhaskar and D. Loganathan, *Tetrahedron Lett.*, 1998, **39**, 2215–2218.
- 15 P. Tiwari and A. K. Misra, *Carbohydr. Res.*, 2006, **341**, 339–350.
- 16 B. Yu, J. Xie, S. Deng and Y. Hui, *J. Am. Chem. Soc.*, 1999, **121**, 12196–12197.
- 17 E. F. Scriven, *Chem. Soc. Rev.*, 1983, **12**, 129–161.
- 18 G. Höfle, W. Steglich and H. Vorbrüggen, *Angew. Chem., Int. Ed. Engl.*, 1978, **17**, 569–583.
- 19 R. Wang, Y. Chen, B. Fei, J. Hu, J. Chen, Y. Luo and Y. Xia, *Org. Lett.*, 2023, **25**, 2970–2974.
- 20 C. D. Hubbard and J. F. Kirsch, *Biochemistry*, 1972, **11**, 2483–2493.
- 21 W. P. Jencks and J. Carriolo, *J. Biol. Chem.*, 1959, **234**, 1272–1279.
- 22 J. Gelas, in *Advances in Carbohydrate Chemistry and Biochemistry*, ed. R. S. Tipson and D. Horton, Academic Press, 1981, vol. 39, pp. 71–156.
- 23 A. I. Vogel, *A text-book of practical organic chemistry including qualitative organic analysis*, Longmans, 1956, vol. 2, pp. 676–681.
- 24 N. P. Bizier, S. R. Atkins, L. C. Helland, S. F. Colvin, J. R. Twitchell and M. J. Cloninger, *Carbohydr. Res.*, 2008, **343**, 1814–1818.
- 25 S. K. Das, K. A. Reddy, V. L. N. R. Krovvidi and K. Mukkanti, *Carbohydr. Res.*, 2005, **340**, 1387–1392.
- 26 C. Limousin, J. Cléophas, A. Petit, A. Loupy and G. Lukacs, *J. Carbohydr. Chem.*, 1997, **16**, 327–342.
- 27 F. Dasgupta, P. P. Singh and H. C. Srivastava, *Carbohydr. Res.*, 1980, **80**, 346–349.
- 28 L. Shi, G. Zhang and F. Pan, *Tetrahedron*, 2008, **64**, 2572–2575.
- 29 K. P. R. Kartha and R. A. Field, *Tetrahedron*, 1997, **53**, 11753–11766.
- 30 S. Toorchi Roudsari and S. Sadjadi, *ChemistrySelect*, 2023, **8**, e202204067.
- 31 C.-A. Tai, S. S. Kulkarni and S.-C. Hung, *J. Org. Chem.*, 2003, **68**, 8719–8722.
- 32 J.-C. Lee, C.-A. Tai and S.-C. Hung, *Tetrahedron Lett.*, 2002, **43**, 851–855.
- 33 R. Ghosh, A. Chakraborty and S. Maiti, *Tetrahedron Lett.*, 2004, **45**, 9631–9634.
- 34 S. Ahmad and J. Iqbal, *J. Chem. Soc., Chem. Commun.*, 1987, **2**, 114–115.
- 35 J.-L. Montero, J.-Y. Winum, A. Leydet, M. Kamal, A. A. Pavia and J.-P. Roque, *Carbohydr. Res.*, 1997, **297**, 175–180.
- 36 K.-C. Lu, S.-Y. Hsieh, L. N. Patkar, C.-T. Chen and C.-C. Lin, *Tetrahedron*, 2004, **60**, 8967–8973.
- 37 H. Binch, K. Stangier and J. Thiem, *Carbohydr. Res.*, 1998, **306**, 409–419.
- 38 B. Mukhopadhyay, *Tetrahedron Lett.*, 2006, **47**, 4337–4341.
- 39 A. K. Misra, P. Tiwari and S. K. Madhusudan, *Carbohydr. Res.*, 2005, **340**, 325–329.
- 40 J. Song and W.-H. Zheng, *Org. Lett.*, 2022, **24**, 2349–2353.
- 41 P. M. Bhaskar and M. Pallooru, *Synlett*, 1999, 129–131.
- 42 R. Kumareswaran, K. Pachamuthu and Y. D. Vankar, *Synlett*, 2000, 1652–1654.
- 43 T. Hirose, B. G. Kopek, Z.-H. Wang, R. Yusa and B. W. Baldwin, *Tetrahedron Lett.*, 2003, **44**, 1831–1833.
- 44 M. H. S. Troftgruben, I. Oldenburg, A. M. Pama, B. Igawa, S. Deguara, M. Shields, N. K. Viswanathan, J. X. Silverio, C. Gleason, J. S.-M. Wang, J. S. Ford, H. Lazaro and S. Eagon, *J. Org. Chem.*, 2024, **89**, 17850–17852.
- 45 S. A. Forsyth, D. R. MacFarlane, R. J. Thomson and M. von Itzstein, *Chem. Commun.*, 2002, 714–715.
- 46 W.-X. Lv, H. Chen, X. Zhang, C. C. Ho, Y. Liu, S. Wu, H. Wang, Z. Jin and Y. R. Chi, *Chem*, 2022, **8**, 1518–1534.
- 47 Y.-G. Liu, Z. Zhong, Y. Tang, H. Wang, S. V. C. Vummaleti, X. Peng, P. Peng, X. Zhang and Y. R. Chi, *Nat. Commun.*, 2025, **16**, 54.
- 48 G. Nie, T. Tu, T. Liao, D. Liu, W. Ye and S.-C. Ren, *Green Chem.*, 2024, **26**, 5397–5408.
- 49 J. Jin, J. Guo, F. Xu, Y. Lv, G. Wang, J. Song, W.-X. Lv, T. Li and Y. R. Chi, *Green Chem.*, 2024, **26**, 5997–6004.
- 50 G. Xiao, G. A. Cintron-Rosado, D. A. Glazier, B. Xi, C. Liu, P. Liu and W. Tang, *J. Am. Chem. Soc.*, 2017, **139**, 4346–4349.
- 51 C. J. Pedersen, *Angew. Chem., Int. Ed. Engl.*, 1988, **27**, 1021–1027.
- 52 V. I. Lakshmanan and S. Vijayan, in *Extraction 2018*, ed. B. R. Davis, M. S. Moats, S. Wang, D. Gregurek, J. Kapusta, T. P. Battle, M. E. Schlesinger, G. R. Alvear Flores, E. Jak, G. Goodall, M. L. Free, E. Asselin, A. Chagnes, D. Dreisinger, M. Jeffrey, J. Lee, G. Miller, J. Petersen, V. S. T. Ciminelli, Q. Xu, R. Molnar, J. Adams, W. Liu, N. Verbaan, J. Goode, I. M. London, G. Azimi, A. Forstner, R. Kappes and T. Bhambhani, Springer International Publishing, Cham, 2018, pp. 1913–1930.



- 53 S. Blair, E. Kempen and J. Brodbelt, *J. Am. Soc. Mass Spectrom.*, 1998, **9**, 1049–1059.
- 54 Y. Song, H. Jing, B. Li and D. Bai, *Chem. – Eur. J.*, 2011, **17**, 8731–8738.
- 55 S. L. Silva, M. S. Valle and J. R. Pliego, *J. Org. Chem.*, 2020, **85**, 15457–15465.
- 56 R. A. Sheldon, *Chem. Commun.*, 2008, **29**, 3352–3365.
- 57 R. A. Sheldon, *Green Chem.*, 2016, **18**, 3180–3183.

