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## Selective monophosphorylation of cyclic diols and polyols *via* hemiboronic acid catalysis†

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The phosphorylation of organic molecules is a biologically essential chemical transformation. Consequently, there is high demand for methods that allow for the direct, selective, and catalytic monophosphorylation of diols and complex polyols. Due to their ability to form reversible covalent bonds with hydroxy (–OH) groups, hemiboronic acids present the unique capacity to catalytically activate diols in a nucleophilic manner. Herein, we disclose a hemiboronic acid-catalyzed monophosphorylation protocol, amenable to a variety of acyclic and cyclic diols, along with the site-selective functionalization of polyols including saccharides. Mechanistic analyses comprising of kinetic experiments and computational investigation were performed to probe for the origin of the observed site-selectivities. We propose that the observed site-selectivity originates from a difference in calculated nucleophilicity between the diol oxygens in the lower energy epimer of the reactive complex, which also exhibits a kinetically stabilizing hydrogen bonding effect with the approaching electrophile.

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## Introduction

Coined as one of Nature's favourite modifications, the phosphorylation of organic molecules is a fundamental transformation central to numerous cellular signalling pathways, energy transduction, protein synthesis, and maintenance of genetic material.<sup>1–3</sup> When considering that modern chemical and biological methods often aim to mimic natural processes, it is unsurprising that the synthetic manipulation of phosphorylation processes is of significant interest and remains one of the great challenges in organic chemistry.<sup>3</sup> In the pharmaceutical industry, phosphorylation is a commonly used prodrug tactic, affording improved bioavailability of therapeutics by bestowing high membrane permeability and aqueous solubility.<sup>4,5</sup> In this approach, upon administration of the phosphate ester-bearing prodrug, the active pharmaceutical agent is released by endogenous phosphatases.<sup>4,6,7</sup> This strategy has been exemplified in countless marketed polyol therapeutics such as fosamprenavir, fludarabine phosphate, and etoposide phosphate (Fig. 1a).<sup>8–10</sup> Synthetic methods for the generation of phosphate monoesters are based on the facile nucleophilic attack of alcohols onto electrophilic phosphate sources. Despite this apparent simplicity,

catalytic activation and/or prerequisite substrate derivatization are often necessary to increase hydroxy group reactivity and achieve site-selectivity in substrates bearing multiple reactive sites.<sup>11–15</sup> Using kinase enzymes, Nature phosphorylates complex biomolecules with high selectivity, circumventing the need for protecting groups. Paradoxically, the adoption of biocatalytic or pseudo-biocatalytic strategies for the preparation of therapeutic and agrochemical compounds has been limited due to active site specificity requiring substrates similar to the native ligand.<sup>16</sup> Despite this challenge, a pioneering report by Miller and coworkers exemplified this strategy using a peptide-based kinase mimic for the synthesis of *D-myo*-inositol-1-phosphate in excellent yield and enantioselectivity (Fig. 1b).<sup>17</sup> Yet, current biomimetic methods pale in comparison to kinases as they require impractical conditions, achieve suboptimal yields, and present poor tolerance in substrate variability. The synthetic phosphorylation of alcohols is typically performed using P(v) reagents such as chlorophosphates or pyrophosphates but can also be achieved using P(III) reagents such as phosphoramidites, phosphites or phosphorochloridites, followed by an oxidation step that is subject to chemoselectivity issues with complex substrates.<sup>11,18</sup> Amongst these strategies, phosphate transfer using P(v) reagents is prevalent, using either Lewis acid catalysis for hydroxy group activation, or nucleophilic catalysts for phosphosphorylating agent activation. Numerous procedures have achieved the catalytic phosphorylation of alcohols using catalysts such as pyridine-*N*-oxide derivatives,<sup>2,18</sup> tetrabutylammonium hydrogen sulfate,<sup>19</sup> Ti(IV) complexes,<sup>20–22</sup> Cu(II) salts,<sup>11</sup> Zn(II) salts,<sup>23</sup> Mo(VI) salts,<sup>24</sup> and dimethyl-

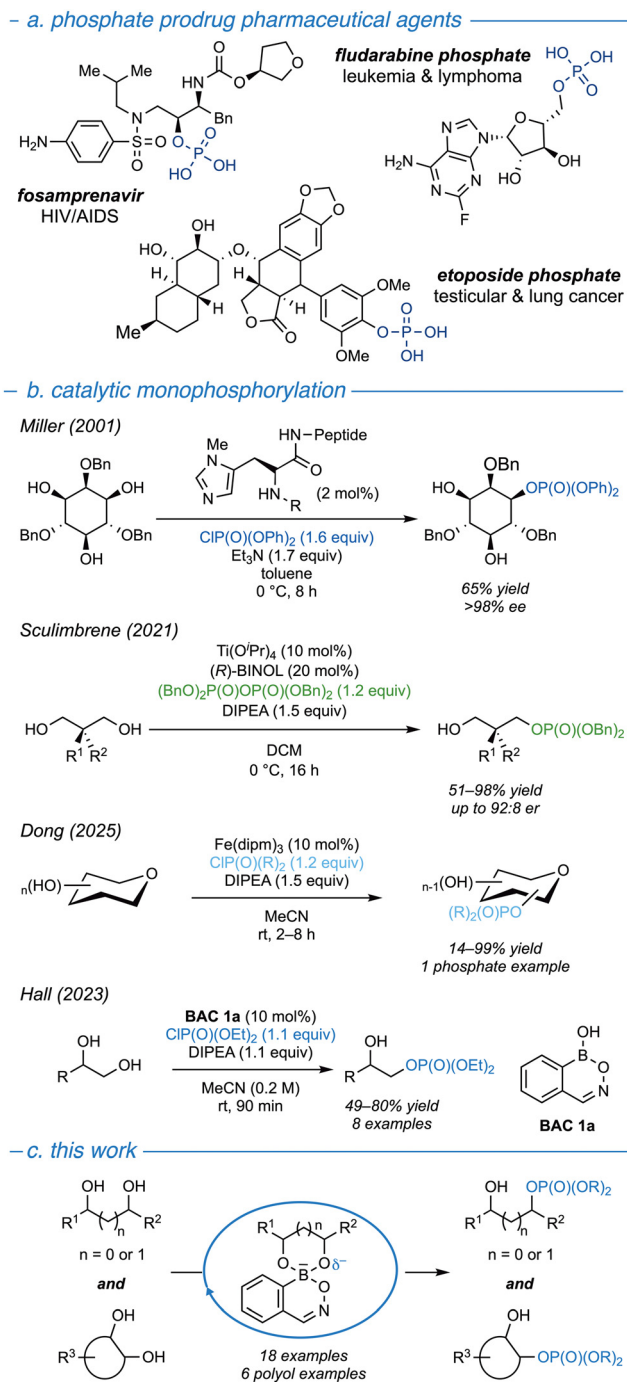
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**Fig. 1** Introduction. (a) Phosphate prodrug pharmaceutical agents. (b) Previous work in catalytic monophosphorylation. (c) This work: hemiboronic acid catalyzed monophosphorylation of acyclic and cyclic diols, and polyols.

aminopyridine (DMAP) derivatives.<sup>25</sup> Conversely, strategies achieving diol monophosphorylation have been seldomly reported, with key reports using catalytic  $\text{Ti}(\text{O}^i\text{Pr})_4$ , attaining moderate success when extended to enantioselective diol desymmetrization (Fig. 1b).<sup>26,27</sup> Recently, Dong and coworkers reported on the  $\text{Fe}(\text{II})$ -catalyzed site-selective monophosphinoylation

of carbohydrate derivatives, however monophosphorylation is not readily amenable to this system (Fig. 1b).<sup>28</sup> Finally, Dong and coworkers recently devised a method employing catalytic  $\text{SnCl}_2$  for the phosphorylation of alcohols, diols, and protected carbohydrates.<sup>29</sup> Under the reported conditions, the monophosphorylation of carbohydrate derivatives bearing three unprotected hydroxy groups was unsuccessful; monophosphinoylation was instead achieved. It remains clear that the selective monophosphorylation of unprotected diols and polyols using mild catalytic conditions is of significant interest, and there is considerable demand for innovative methods.

Boron based catalysis (boronic acid, hemiboronic acid, and borinic acid) has transformed the field of selective diol functionalization. It operates through the ability to dynamically exchange with hydroxy groups and form an anionic tetrahedral boron adduct, temporarily increasing the nucleophilicity of the diol oxygens (Fig. 1c).<sup>30</sup> Borinic acids readily form the desired tetrahedral adduct through the single exchangeable site at boron, however they suffer from poor oxidative stability, requiring special precautions such as employing the more robust ethanolamine adduct as a pre-catalyst.<sup>31</sup> In recent years, the synthetic applicability of these species in catalytic diol and polyol activation for selective carbohydrate monoacylation, sulfonylation, and alkylation has been exemplified by Taylor and coworkers.<sup>31,32</sup> Hemiboronic acids present the unique ability of having both a single exchangeable site and superior oxidative stability, with tunable Lewis acidity tailored to the desired catalytic application. Despite having first been synthesized in the 1950s by Snyder and Dewar, benzoxazaborine and boroxarophenanthrene have only recently been demonstrated as competent catalysts for the nucleophilic activation of 1,2-diols and 1,3-diols, respectively.<sup>33–36</sup> Using benzoxazaborine (**BAC 1a**), we previously reported the selective monophosphorylation of a small subset of acyclic 1,2-diols, but the wider potential of this system for the site-selective monophosphorylation of biologically relevant diols and polyols remained to be explored (Fig. 1b).<sup>33</sup>

Herein, the successful implementation of hemiboronic acid catalysis to the selective monophosphorylation of a variety of acyclic and cyclic 1,2-diols, and unprotected polyols is disclosed. In addition, the determination of preferred catalyst electronics, and computational insight into the origin of the observed site-selectivity with carbohydrates are described (Fig. 1c).

## Results and discussion

### Optimization

We began our investigation by subjecting *cis*-1,2-cyclohexanediol to our previously reported conditions, which resulted in the formation of monophosphorylated product **2a** in an 81% yield, in addition to pyrophosphate **3** as an undesired by-product in a 20% yield (Table 1, entry 2).<sup>33</sup> This unreactive species resulted in onerous purification, as it was difficult to separate from the desired monophosphorylated product. We





Table 1 Reaction optimization

Entry	Deviation	Yield 2a <sup>a</sup> (%)
1	None	82% [7%] (<1%)
2	No 3 Å MS	81% [20%]
3	No 3 Å MS, 1 equiv. H <sub>2</sub> O	35% [27%]
4	BAC 2	0%
5	BAC 3	6%
6	BAC 4	0%
7	BAC 5	22%
8	BAC 6	0%
9	BAC 7	0%
10	BAC 8	40%
11	BAC 9	42%
12	2,6-lutidine	12% (0%)
13	2,6-di- <i>tert</i> -butylpyridine	0% (0%)
14	DBU	13% (0%)
15	pyridine	20% (21%)
16	K <sub>2</sub> CO <sub>3</sub>	62% (0%)
17	Et <sub>3</sub> N	70% (0%)
18	ClP(O)(OMe) <sub>2</sub>	10%
19	ClP(O)(OPh) <sub>2</sub>	45% <sup>b</sup>
20	ClP(O)(OBn) <sub>2</sub>	70%
21	(BnO) <sub>2</sub> P(O)OP(O)(OBn) <sub>2</sub>	0%
22	( <sup>i</sup> Pr) <sub>2</sub> NP(Cl)OCH <sub>2</sub> CH <sub>2</sub> CN	0%

<sup>a</sup> All reactions performed on a 0.1 mmol scale, with yields determined by <sup>1</sup>H NMR using 1,3,5-trimethoxybenzene as the internal standard.

<sup>b</sup> Yield of cyclized phosphate product obtained (see ESI for details†). Yields in parentheses denote background product formation in the absence of boronic acid catalyst. Yields in brackets denote byproduct 3 formation.

hypothesized that formation of 3 stemmed from the presence of residual water, which was corroborated by the observed increase in by-product formation with the stoichiometric addition of water (entry 3). The incorporation of 3 Å molecular sieves suppressed the formation of 3 (entry 1). Numerous cyclic hemiboronic acid catalysts of varying Lewis acidity were screened as potential catalysts, yet benzoxazaborine (BAC 1a) was found to be the most efficient at catalyzing this transformation (entries 4–11).<sup>33,34,37</sup> A variety of non-nucleophilic bases were tested both in the presence and absence of BAC 1a to monitor background reactivity, unveiling *N,N*-diisopropylethylamine (DIPEA) as the ideal base for this system (entries 12–17). Chlorophosphates are the most commonly employed P(v) reagents as they are generally commercially available and are synthetically straightforward to use.<sup>20,22,38</sup> Within this sub-

class, diethyl-, dimethyl-, diphenyl-, and dibenzylchlorophosphate were assessed as phosphorylating agents (entries 18–20). Both dimethyl- and diphenylchlorophosphate were not competent phosphorylating agents due to poor yields with the former, and product cyclization in the latter as was also observed by Dong and coworkers with certain substrates (see ESI for details†).<sup>29</sup> As diethylchlorophosphate is commercially available and led to slightly increased yields in comparison to dibenzylchlorophosphate, it was selected as the primary phosphorylating agent for this study. Finally, tetrabenzylpyrophosphate, and 2-cyanoethyl *N,N*-diisopropylchlorophosphoramidite, popular in oligonucleotide synthesis,<sup>39</sup> were not reactive in this system (entries 21 & 22).

## Scope

With optimized conditions in hand, we turned our attention to the scope of diols and polyols. In all cases, scaling the reaction up from 0.1 mmol to 0.3 mmol resulted in a higher isolated yield in the latter than what was observed by <sup>1</sup>H NMR yield during optimization. Monophosphorylated products of cyclic *cis*-1,2-diols were obtained in near quantitative yields for 5- (2b), 6- (2a), and 8-membered rings (2c), and in moderate yield for 12-membered ring (2d) and anhydroerythritol (2f) (Fig. 2). Despite *cis*-1,2-diols being the preferred motif for boronic acid complexation,<sup>30,40</sup> monophosphorylated *trans*-1,2-cyclohexanediol (2e) was isolated in a 98% yield. Acyclic diols were also re-examined under these optimal conditions, focusing on 1,2-disubstituted-1,2-diols and 1,3-diols that were not studied in our prior work.<sup>33</sup> Both 2*R*,3*R*-2,3-butanediol (2j) and 1*R*,2*R*-1,2-diphenylethanediol (2g) lead to moderate isolated yields. Furthermore, 4-*CF*<sub>3</sub> (2h) and 2-Me (2i) derivatives of 1*R*,2*R*-1,2-diphenylethanediol, were successfully monophosphorylated in a 51% and 74% yield, respectively, demonstrating tolerance of electron poor diols, and increased steric hindrance. In addition to acyclic 1,2-diols, the products originating from 2-phenyl-1,3-propanediol (2k) and chloramphenicol (2l) were isolated in moderate to poor yields, with selective functionalization of the primary alcohol in the latter. Remarkably, the site-selective monophosphorylation of six pyranose derivatives (2m–r), including the lincosamide antibiotic clindamycin (2r), proceeded with excellent to near quantitative yields and complete site-selectivity. In all cases (2m–r), monophosphorylation occurred at the C3 position, as has previously been observed in catalytic nucleophilic functionalization of this substrate subclass.<sup>28,29,31,32</sup> Site-selectivity assignments were performed using 2D gCOSY NMR to first assign the <sup>1</sup>H signals around the sugar ring. In conjunction with this analysis, <sup>31</sup>P decoupled <sup>1</sup>H NMR data was recorded. The spectrum exhibited a simplification in the coupling pattern for the <sup>1</sup>H resonance on the carbon bound to the phosphorylated oxygen through the removal of the observed <sup>3</sup>J <sup>1</sup>H–<sup>31</sup>P (see ESI for details†). Unfortunately, this synthetic methodology was not conducive to the site-selective monophosphorylation of furanose derivatives, as it resulted in an equal mixture of constitutional isomers (see ESI for details†). Likewise, complex polyols bearing more than three unprotected hydroxy groups or pos-





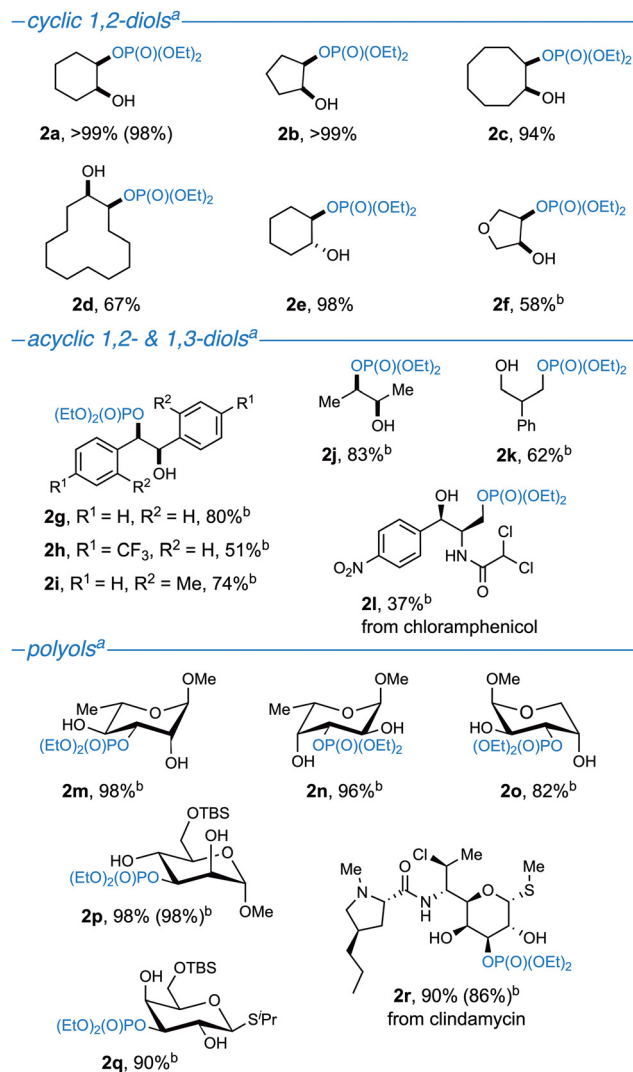


Fig. 2 Reaction scope. <sup>a</sup>Reactions performed on a 0.3 mmol scale, with diethylchlorophosphate (1.1 equiv.), DIPEA (1.1 equiv.), BAC 1a (10 mol%), in MeCN (0.1 M) with 3 Å MS, at room temperature, for 1.5 h. Isolated yields reported. Yields in parenthesis run on 1.0 mmol scale. <sup>b</sup>Reaction run for 3 h.

sessing unprotected phenol moieties afforded poor site-selectivity (see ESI for details†).

### Phosphate ester deprotection

As phosphate prodrugs are almost exclusively found in their free phosphoric acid form, it is important to demonstrate the applicability of this method to the synthesis of these medicinally valuable compounds. Since current methods to access polyol-based phosphate drugs require multiple additional steps for protecting group installation and cleavage, our catalytic system provides a notable improvement.<sup>41a–c</sup> Despite the commercial availability of diethylchlorophosphate, the harsh deprotection of phosphorylated polyols using conventional methods, such as the McKenna deprotection, was unsuccessful in our hands.<sup>42a–c</sup> We then sought to identify a different

phosphate ester that is deprotectable under milder conditions. As dibenzylchlorophosphate is both easily prepared from commercially available dibenzylphosphite, and the deprotection of benzyl esters is well established, it was selected as an alternative phosphorylating agent.<sup>43,44</sup> Thus, it was applied to the monophosphorylation of four previous substrates, in low to moderate yields (Fig. 3, examples 4a, 4m, 4p, 4r). Subsequently, the corresponding benzyl phosphate products were subjected to standard hydrogenation conditions, yielding the corresponding phosphoric acid products in near quantitative yields, circumventing column chromatography purification (Fig. 3, examples 5a, 5m, 5p, 5r).<sup>43,44</sup> In the case of 5p, the global deprotection of 4p was achieved, with concurrent removal of the silyl protecting group.<sup>45</sup>

## Mechanistic investigation

### Kinetic studies

It is well established that the pK<sub>a</sub> of a boronic acid catalyst has a significant impact on its catalytic efficacy.<sup>30,33</sup> Catalysts possessing a moderate pK<sub>a</sub> are typically sought after for nucleophilic activation, as they allow a compromise between efficient diol complexation and sufficient build-up of negative charge within the complex, while also enabling catalyst turnover. Previous work by our group alluded to a three-step mechanism for boronic acid catalyzed diol monophosphorylation (Fig. 4a): initial catalyst complexation (benefitting from a low pK<sub>a</sub> catalyst), nucleophilic attack on the phosphorylating agent (benefitting from a higher pK<sub>a</sub> catalyst), and catalyst turnover *via* product dissociation.<sup>33</sup> We hypothesized that by synthesizing electronic variants of benzoxazaborine catalyst (BAC 1a) possessing a range of pK<sub>a</sub> values, we could investigate the impact catalyst pK<sub>a</sub> has on the kinetic profile of the reaction. We successfully synthesized four new variants with pK<sub>a</sub> values ranging

### —phosphate ester deprotection—

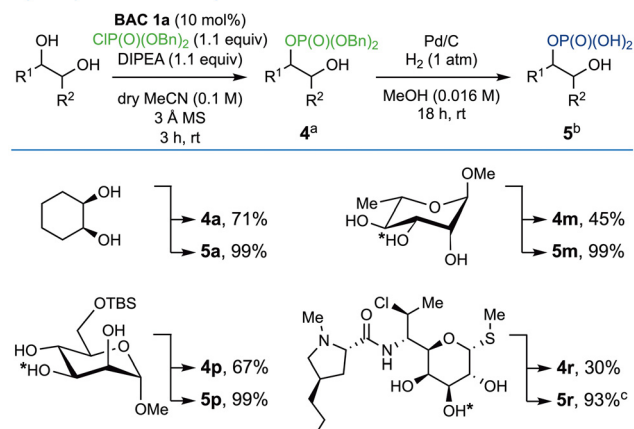
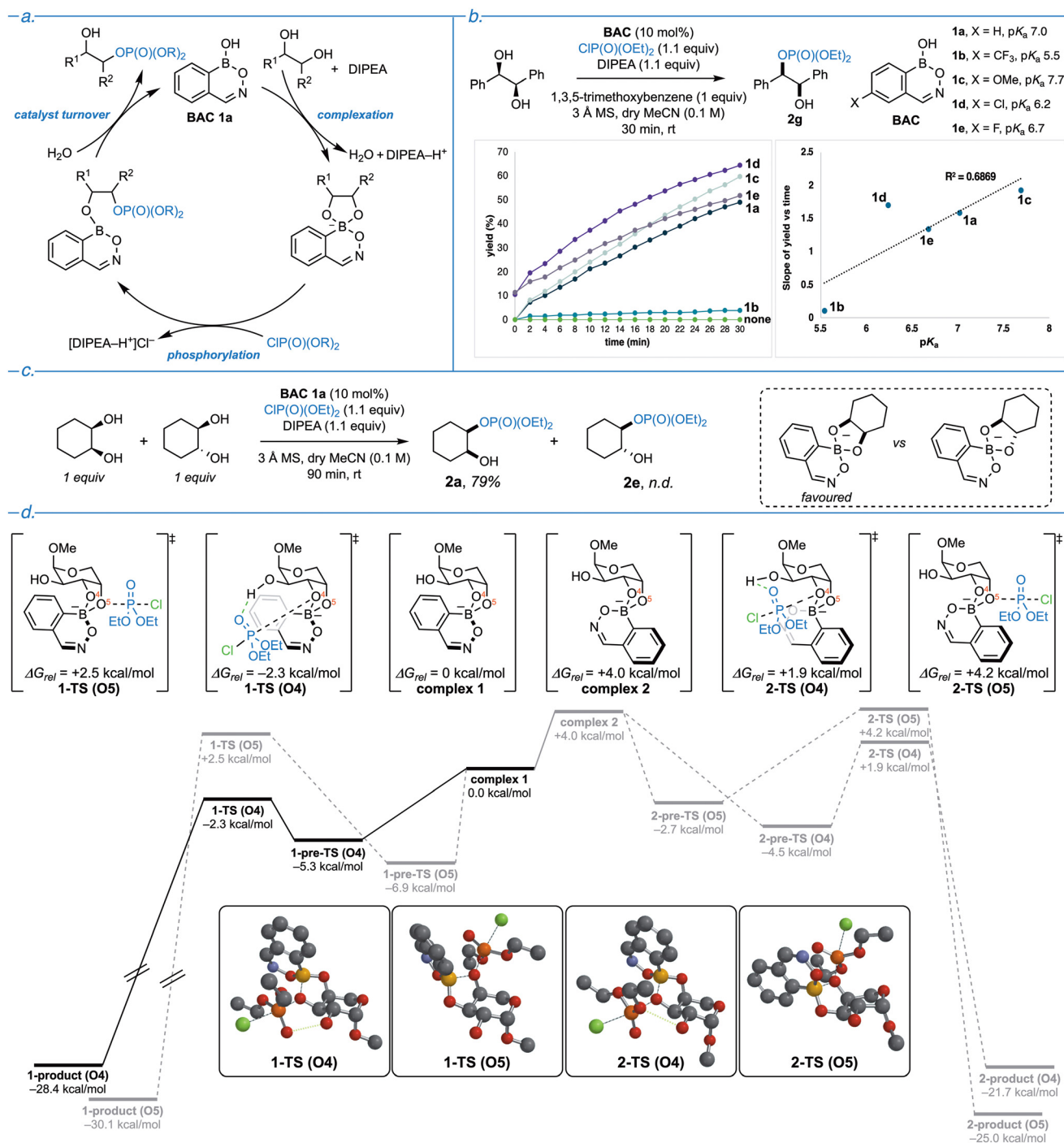


Fig. 3 Product deprotection. <sup>a</sup>Reactions performed on a 1.0 mmol scale, isolated yields reported. <sup>b</sup>Reactions performed on a 0.3 mmol scale, isolated yields reported. <sup>c</sup>Reaction run with MeOH (0.1 M). \*Denotes hydroxy group that is phosphorylated.



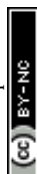




**Fig. 4** Mechanistic investigations. (a) Proposed catalytic cycle. (b) Initial rates kinetics study studying the impact of hemiboronic acid catalyst  $pK_a$ . Reactions performed on a 0.1 mmol scale, with 1*R*,2*R*-1,2-diphenylethanediol (1.0 equiv.), diethylchlorophosphate (1.1 equiv.), DIPEA (1.1 equiv.), BAC (10 mol%), 1,3,5-trimethoxybenzene (1.0 equiv.) in MeCN (0.1 M) with 3 Å MS, for 30 min, at room temperature in an NMR tube. NMR yields reported, using 1,3,5-trimethoxybenzene as the internal standard. (c) *cis*- & *trans*-1,2-cyclohexanediol competition study. Reactions performed on a 0.3 mmol scale. Isolated yield reported. (d) Transition state analysis via DFT computations (SMD(MeCN)/ $\omega$ B97X-V(6-311+G(2df,2p))// $\omega$ B97X-D(6-31G(d)).

from 5.5–7.7 (Fig. 4b, **BAC 1a–1e**), with substitution *para* to boron, maximizing the electronic effects at the boron centre, while minimizing possible steric interactions with the incoming diol or phosphorylating agent. The relative ordering of

determined  $pK_a$  values agrees with the substituents' established  $\sigma_p$  values.<sup>46</sup> The selected substrate for this experiment was 1*R*,2*R*-1,2-diphenylethanediol, as phosphorylation occurred on an appropriate time scale for monitoring by  $^1\text{H}$





NMR. In order to minimize possible manifestations of product inhibition, kinetics were only examined for the first 30 minutes of reaction time (Fig. 4b). The lowest  $pK_a$  catalyst (**BAC 1b**) had the slowest onset of reactivity, and nearly inexistent catalytic activity. Catalyst **BAC 1d** had the quickest onset of reactivity and is the highest yielding throughout the experiment. Interestingly, catalysts **BAC 1a**, **1d**, **1e**, exhibit the beginning of a plateau in yield, while the highest  $pK_a$  catalyst (**BAC 1c**) shows a slow but steady increase in yield throughout the experiment. These observations are in line with either the phosphorylation or catalyst turnover steps being rate limiting, as higher  $pK_a$  catalysts are superior. To assess the possible correlation between the initial rate of the reaction and catalyst Lewis acidity,  $pK_a$  values were plotted against the initial slopes of product formation for each catalyst, showing good correlation ( $R^2 = 0.9670$ ) between the two parameters for **BAC 1a**, **1b**, **1c**, and **1e** (Fig. 4b). However, catalyst **BAC 1d** did not fall within this trend, which is not entirely unexpected given previous studies showing that a weaker correlation between the Hammett parameters and  $pK_a$  values of 4-halo substituted boron heterocycles appears to exist.<sup>47</sup>

### Origin of selectivity

Given the well-established preference for cyclic hemiboronic acids to bind to *cis*-diols, and the observed site-selectivity with polyol substrates, the nearly quantitative yield obtained with product **2e** was surprising, as reaction conditions were identical to *cis*-diol examples. <sup>11</sup>B NMR complexation experiments confirmed that when subjected to the reaction conditions, both *cis*- and *trans*-1,2-cyclohexanediol form a tetrahedral boronate complex, with characteristic signals at 6.9 and 7.0 ppm, respectively (see ESI for details†). Thus, to better understand the site-selectivity observed with polyol examples **2m–2r**, a competition experiment with equimolar *cis*- and *trans*-1,2-cyclohexanediol was performed. Remarkably, only the *cis*-diol product **2a** was monophosphorylated and isolated, albeit in a slightly diminished yield (Fig. 4c). This outcome demonstrates that when both *cis*- and *trans*-diols are present, this system is fully selective for the former, while if only the latter is present, it will still react.

### Computational studies

To gain insight into the origin of the observed site-selectivity with monosaccharides, Fukui indices ( $f_k^-$ ) were determined for example **2n**, assessing **O4** and **O5** in both complex **1** and complex **2** (Fig. 4d). This metric is a means of evaluating the relative nucleophilicity of various atoms in a molecular structure, which is calculated by comparing the Mulliken charges of each nucleophilic atom at ground state and upon the removal of one electron from the system.<sup>48</sup> This method was successfully employed by Taylor and coworkers to rationalize their observed site-selectivity in the borinic acid-catalyzed acylation of the same class of carbohydrate derivatives.<sup>32</sup> Unfortunately, we observed that this parameter is not suitable in the current system owing to the electronic differences between hemiboronic acid catalyst **BAC 1a** with the previously

reported borinic acid catalyst (see ESI for details†). Using density functional theory calculations (SMD(MeCN)/ $\omega$ B97X-V (6-311+G(2df,2p))// $\omega$ B97X-D/6-31G(d)), transition state analysis was conducted using example **2n** to better understand the observed site-selectivity (Fig. 4d). As two possible tetracoordinate boron epimers can form upon condensation of **BAC 1a** onto **2n**, both geometries were optimized (Fig. 4d, complex **1** and complex **2**), revealing a 4.0 kcal mol<sup>−1</sup> energy difference in favour of complex **1**. Condensation of **BAC 1a** onto **2n** activates two oxygen atoms (**O4** and **O5**) that can undergo nucleophilic attack onto the incoming diethylchlorophosphate. As such, the transition states were calculated for each oxygen atom of both epimers of the **BAC 1a** condensate. Starting from the more energetically favourable complex **1**, pre-transition state and transition state energies were obtained for the phosphorylation at **O4** (observed product) and **O5**. Despite **1-pre-TS (O4)** being higher in energy than **1-pre-TS (O5)**, the activation barrier from **1-pre-TS (O4)** to **1-TS (O4)** is only 3.0 kcal mol<sup>−1</sup>, while it is 9.4 kcal mol<sup>−1</sup> from **1-pre-TS (O5)** to **1-TS (O5)**. Similarly, for complex **2**, nucleophilic attack from **O4** is more kinetically accessible over **O5**. Interestingly, a hydrogen bonding interaction was observed between the diethyl chlorophosphate and (**H**) **O3** within **1-TS (O4)** and **2-TS (O4)**. It is plausible that this interaction is key to the exquisite site-selectivity by stabilizing the transition state of the nucleophilic attack at **O4** over **O5**.

## Conclusion

We have disclosed a practical methodology leveraging the nucleophilic mode of cyclic hemiboronic acid catalysis for the site-selective monophosphorylation of a variety of cyclic diols and polyols, as well as acyclic diols. This method is amenable to the preparation of phosphate ester and free phosphate containing polyol drugs under ambient conditions. Mechanistic studies have uncovered a plausible explanation for the observed site-selectivity of the transformation, and a study of initial-rates kinetic study revealed that this system benefits from a more electron-rich boron catalyst. Future work will be driven towards developing a catalyst capable of achieving enantioselective monophosphorylation of symmetrical, prochiral substrates, in addition to extending the tolerance of this system towards polyols of increased complexity.

## Author contributions

D. G. H. obtained funding. D. G. H. and G. F. O. conceptualized the study. G. F. O. led investigation and methodology development, and supervised D. L. A. D. J. K. carried out computational studies. G. F. O. wrote the original manuscript, with review and editing in collaboration with D. G. H. and D. J. K.

## Conflicts of interest

The authors declare no conflicts of interest.





## Data availability

Data for this article, including experimental details, analytical data, and spectral reproductions for all new compounds, and details of all mechanistic and computational data have been included as part of the ESI.†

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