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# A unified strategy for the synthesis of aldohexoses by boronate assisted assembly of $\text{CH}_2\text{X}_2$ derived $\text{C}_1$ -building blocks†‡

Sujenth Kirupakaran, Glib Arago and Christoph Hirschhäuser \*

A synthetic strategy for all aldohexoses with individually addressable protecting groups from dihalomethane  $\text{C}_1$ -units is reported. The underlying synthesis of  $\text{C}_6$ -sugar alcohols relies on three consecutive Matteson sequences, vinylation and bishydroxylation. *Erythro* and *threo* isomers have been realized for every glycol motif by strategic variation of the sequence.

Carbohydrates are of immense biological importance as a source of energy and as complex chiral scaffolds that participate in numerous recognition processes.<sup>1</sup> Deciphering this “glycocode” is a task, which requires modern analytics as well as organic synthesis.<sup>2</sup> Nowadays oligosaccharides can even be prepared in an automated manner from orthogonally protected monosaccharides.<sup>3</sup> Syntheses of the latter often still rely on ex-chiral pool strategies, each of which has to face the challenge of differentiating five similar hydroxyl groups.<sup>4</sup> *De novo* syntheses, such as the ones developed by Sharpless,<sup>5</sup> Danishefsky<sup>6</sup> or MacMillan<sup>7</sup> approach this problem from the bottom up. Strategies based on  $\text{C}_1$ -building blocks, like those by Fischer,<sup>8</sup> Dondoni<sup>9</sup> and Matteson<sup>10</sup> could allow for maximal protecting group variability and enable isotopic labelling of individual atoms.<sup>11</sup> However, each of these methods has its individual limitations<sup>12</sup> and no unified  $\text{C}_1$ -based strategy to aldohexoses had been reported until now. One key to our route is the Matteson homologation (MH) shown in Scheme 1A.<sup>13</sup> This sequence employs chiral boronic esters (**1**), which react with a lithiated dihalomethane and  $\text{ZnCl}_2$  at low temperatures. In the resulting ate complex (**2**) electrostatic interactions<sup>14</sup> between the zinc bound chloride atoms and the carbenoid C–H favour an anti-periplanar arrangement of one C–X-bond relative to the boronate R-group. Upon warming 1,2-rearrangement results in the diastereoselective formation of  $\alpha$ -halo boronates (**3**). Reaction with various nucleophiles yields  $\alpha$ -chiral boronates (**4**) under stereochemical inversion. Thus MHs are highly useful for preparing heteroatom rich motifs.<sup>16</sup>

As shown in Scheme 1B, iterative MH and substitution with alkoxides can lead to carbohydrate like structures. This was applied by Matteson to the synthesis of *L*-ribose.<sup>10</sup> While MH

and substitution with  $\text{LiOBn}$  worked well for the  $\text{C}_1$ -building block **5**, and two more homologs, further homologation proved to be problematic. Attempts to react **6** with  $\text{LiCHBr}_2$  led to intractable mixtures and the use of  $\text{LiCHCl}_2$  only allowed for the indirect detection of product traces. The route was thus concluded by homologation with  $\text{LiCH}_2\text{Cl}$ , which does not allow for installation of another stereocenter. For detailed discussion of this surprising limitation, an explanatory hypothesis and supporting evidence see the ESI.† Importantly this restricted Matteson’s synthesis to ribose (an aldopentose),<sup>15</sup> while most biologically relevant monosaccharides are hexoses. Thus, a  $\text{C}_1$ -based synthesis of aldohexoses that allows for (i) installing individual protecting groups, (ii) choosing the configuration at each stereocenter and (iii) potentially introducing isotopic labels at every individual atom, remained an open challenge.<sup>11</sup>



Scheme 1 Matteson reactions in the synthesis of monosaccharides. YouTube tutorial: <https://youtu.be/vXy5oVavJUU>.

University of Duisburg-Essen, Universitätsstr. 5-7, 45117 Essen, Germany. E-mail: Christoph.hirschhaeuser@uni-due.de

† Dedicated to Prof. Hans-Günther Schmalz on the occasion of his 66th birthday.  
‡ Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d3sc03778a>



We achieved this by preparing orthogonally protected versions of prototypical sugar alcohols from  $\text{CH}_2\text{X}_2$  building blocks through three MHs, vinylation and bishydroxylation (Scheme 1C). By strategically combining different homologation and vinylation strategies, both *erythro* and *threo* isomers were realized. Conversion of the sugar alcohols into aldohexoses can be achieved by oxidation of either terminal hydroxyl functionality. By combining this with a short synthesis of the vinylation agent from  $\text{CH}_2\text{X}_2$  building blocks we paved the way for the late-stage introduction of isotopic labels.

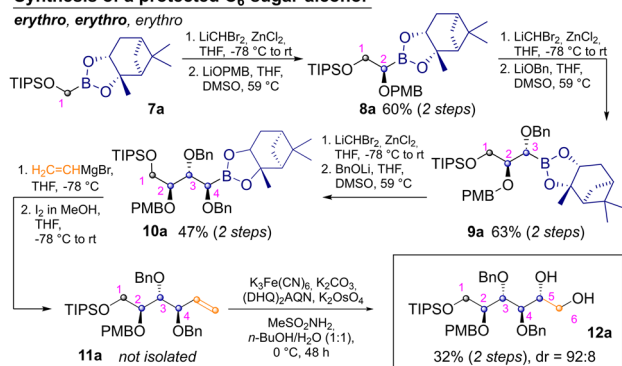
To start the discussion with the stereochemically most basic example, the synthesis of allitol **12a** is depicted in Scheme 2. It begins with the  $\text{CH}_2\text{Br}_2$  derived  $\text{C}_1$ -building block **7a**. MH with  $\text{LiCHBr}_2$  and substitution with  $\text{LiOPMB}$  delivered the  $\text{C}_2$ -building block **8a**. Two consecutive MHs, which are followed by substitution with an alkoxide, produce a masked *erythro* glycol motif. Thus, a second MH and substitution with  $\text{LiOBn}$  delivered **9a**, with an *erythro* relationship between  $\text{C}^2$  and  $\text{C}^3$  (as IUPAC priorities change during the route, carbon atoms are numbered according to their introduction in this article). Another homologation and substitution with  $\text{LiOBn}$  yielded **10a** in 47% yield after two steps.<sup>18</sup> While other alkoxide based protecting groups could have been used here, a second benzyl group was chosen, to allow for confirmation of the relative configuration by direct comparison (see ESI†). Vinylation of **10a** was achieved by Zweifel-olefination.<sup>17</sup> Although this reaction had not been described for the sterically hindered and thermodynamically stable pinanediol boronic esters, it proceeded reasonably well after some optimization (see ESI†) yielding **11a**. The product contained some unidentified contaminations, which were removed after the next step, in which Sharpless bishydroxylation delivered the desired allitol-derivative **12a** (32% yield, calculated over both steps). Several silylethers at  $\text{C}^1$

were tested but neither the use of TBS- (*tert*-butyldimethylsilyl) nor TBDPS-groups (*tert*-butyldiphenyl-silyl) on the first hydroxyl group allowed for introduction of a fourth carbon atom (**13** → **14**). This was quite surprising as a third Matteson-sequence had worked well for the benzyl-derivative **6**. Indeed, homologation of **13** to bromides **15** proceeded with reasonable efficiency, but subsequent substitution with  $\text{LiOBn}$  led to decomposition. This was attributed to competing nucleophilic attack of benzoxide on the silyl ether. We suspected that the latter was activated by an intramolecular O–B interaction, which simultaneously deactivates the boron-atom as an electrophile. By using the TIPS-group (triisopropylsilyl) this side reaction was avoided, through better shielding of the silicon atom (further discussion in ESI†).<sup>19</sup>

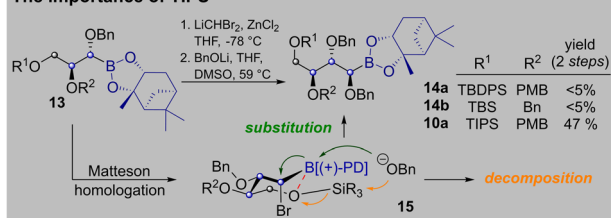
In order to extend the route to other diastereomers it was necessary to modify the synthesis, so that *threo*-glycols could be obtained. This was a particular challenge for the three stereocenters generated by Matteson homologation as the exchange of the pinanediol director is quite cumbersome.<sup>20</sup> Thus the route was modified as shown in Scheme 3.

In order to establish a *threo*-relationship between  $\text{C}^2$  and  $\text{C}^3$  the synthesis starts with *S,S*-dicyclohexylethylenediol (*S,S*-DICHED)<sup>21</sup> boronic ester **7b**. Homologation and substitution analogously to Scheme 2 delivered **8b-1**. The greater thermodynamic stability of pinanediol boronic esters<sup>22</sup> allowed for transesterification to **8b-2** with (+)-pinanediol in  $\text{Et}_2\text{O}$ ,<sup>15</sup> as well as recovery of the precious *S,S*-DICHED auxiliary. However, once this card had been played and the more stable pinanediol boronate was formed, a different strategy had to be applied. In order to establish a *threo* relationship between  $\text{C}^3$  and  $\text{C}^4$ , the third Matteson homologation was followed by substitution with vinylmagnesium bromide<sup>23</sup> and oxidation to **16b**. In contrast to Scheme 2 the C–B bond is now converted into the new C–O bond, while the vinyl group is introduced under inversion.

### Synthesis of a protected $\text{C}_6$ -sugar-alcohol

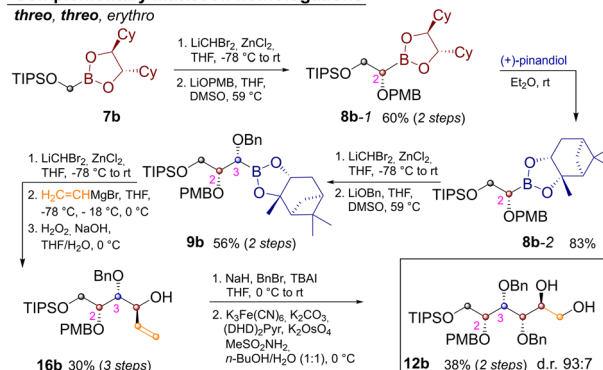


### The importance of TIPS

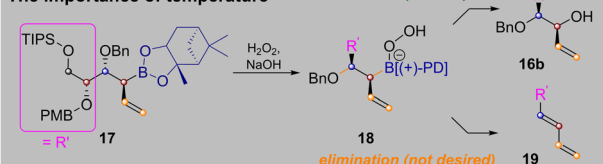


Scheme 2 Synthesis of protected allitol and associated challenges.

### Complementary Matteson homologations



### The importance of temperature



Scheme 3 Synthesis of protected L-glucitol and associated challenges.





## Conclusions

All in all we have developed a highly modular route that opens up a vast field of opportunities for the synthesis of differentially protected sugar alcohols and carbohydrates. By combining the different approaches depicted in Schemes 2–4 a wide variety of C<sub>6</sub>-sugar alcohols and by extension all natural and unnatural aldohexoses become available. Two consecutive MHs, followed by substitution with an alkoxide lead to *erythro*-C<sub>3</sub> building block **9a**. To introduce a C<sup>2</sup>–C<sup>3</sup> *threo*-relationship the chiral director can be changed from DICHD to pinanediol. The transesterification proceeds readily and allows for recovery of the valuable DICHD auxiliary. Both enantiomers of DICHD and pinanediol are available, so that all stereoisomers of C<sub>3</sub>-building blocks of type **9** are accessible. To gain control over the relationship between C<sup>3</sup> and C<sup>4</sup> a strategic crossroad was incorporated in the next homologation. A vinyl group was introduced either by Zweifel olefination (*erythro*) or Matteson substitution (*threo*). Thereby C<sub>3</sub>-building blocks of type **9** can be converted into vinyl tetrol of type **11**, again with the potential for making all stereoisomers. For installing the final glycol moiety Sharpless bishydroxylation was employed. Unfortunately overwhelming substrate control only allowed for the direct synthesis of sugar alcohols of type **12** with a C<sup>4</sup>–C<sup>5</sup>-*erythro* configuration. In order to obtain a C<sup>4</sup>–C<sup>5</sup>-*threo* configuration at this position conversion into epoxide **20** was necessary. Some first attempts at epoxide opening to a diol of type **12** (with a C<sup>4</sup>–C<sup>5</sup>-*threo* configuration) were plagued by side reactions (see ESI†). Fortunately this only affects the synthesis of monosaccharides with both C<sup>2</sup>–C<sup>3</sup>-*threo* and C<sup>4</sup>–C<sup>5</sup>-*threo* configurations (*i.e.*, galactose and idose). For these cases the corresponding epoxides might be better converted into hexoses along the lines of Sharpless's carbohydrate synthesis.<sup>31</sup> In all other cases conversion into the desired aldohexoses can be achieved by appropriate cyclisation *via* C<sup>1</sup> or C<sup>6</sup> due to the orthogonal silyloxy group at C<sup>1</sup> (Scheme 5). Another advantage of these two cyclisation options arises as half of the carbon scaffold is introduced in the last homologation/vinylation sequence. The required vinyl metal species can be prepared from two (CH<sub>2</sub>X<sub>2</sub> derived) C<sub>1</sub> building blocks (Scheme 5). By choosing the appropriate cyclisation route an isotopic label could thus be placed at every position in the aldohexose scaffold. Thus this C<sub>1</sub> based *de novo* approach to aldohexoses is uniquely suited for the synthesis of labelled aldohexoses, which we plan to pursue in the near future.

## Data availability

All experimental and characterization data, as well as pictures of NMR spectra are available in the ESI.†

## Author contributions

S. K. developed the synthesis of aldohexoses; G. A. developed the synthesis of vinyl metal species from C<sub>1</sub>-units; C. H. conceived the project and wrote the manuscript.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

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