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# Transketolase catalysed upgrading of L-arabinose: the one-step stereoselective synthesis of L-glucuheptulose†

Fabiana Subrizi,<sup>a</sup> Max Cárdenas-Fernández,<sup>b</sup> Gary J. Lye,<sup>b</sup> John M. Ward,<sup>b</sup> Paul A. Dalby,<sup>b</sup> Tom D. Sheppard<sup>a</sup> and Helen C. Hailes<sup>\*a</sup>

Conversion of biomass using biocatalysis is likely to become a technology that contributes significantly to the future production of chemical building blocks, materials and transport fuels. Here the synthesis of a value-added chemical from L-arabinose, a major component of the carbohydrates in sugar beet pulp (SBP), in a concise and sustainable manner has been investigated. Biocatalytic conversions using transketolase variants have been developed for the efficient, scalable synthesis of a rare naturally occurring ketoheptose, L-glucuheptulose, from L-arabinose. New active *E. coli* TK mutants that readily accept L-arabinose were identified using a versatile colorimetric screening assay and the reaction was performed on a preparative scale.

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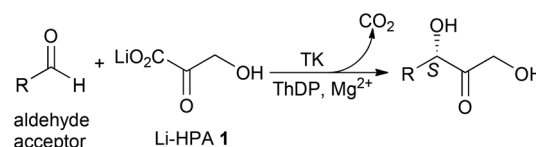
## Introduction

Sustainability issues related to the use of crude oil are progressively directing the chemical and pharmaceutical industries towards renewable sources for the production of organic synthons. Indeed, biomass represents an attractive alternative as a renewable source for chemical building blocks, materials and biofuels.<sup>1</sup> In a bio-based economy, biocatalysis has been recognised as a green, environmentally friendly technology with significant potential due to the mild reaction conditions used and high chemo- and enantioselectivities that can be achieved.<sup>2</sup>

Transketolase (TK), a key enzyme in metabolic regulation providing a link between the pentose phosphate pathway and glycolysis, is a thiamine diphosphate (ThDP) dependent enzyme that has been used in stereospecific carbon-carbon bond formation.<sup>3</sup> *In vivo*, TK catalyses the reversible transfer of a two-carbon fragment from a five-carbon ketose (D-xylulose-5-phosphate) to a five-carbon aldose (D-ribose-5-phosphate) to form a seven-carbon ketose (D-sedoheptulose-7-phosphate). The ability of TK to perform the reaction irreversibly with the loss of carbon dioxide, when using β-hydroxypyruvate (HPA) as the ketol donor, makes it an attractive biocatalyst in industrial

synthesis.<sup>4,5</sup> To achieve commercial viability in large-scale processes with non-phosphorylated substrates, variants of TK with greater specific activity are required. The TK reaction with lithium hydroxypyruvate (Li-HPA **1**) as a donor has been studied with a range of aldehyde acceptors (Scheme 1) and TK variants exhibiting enhanced activity towards non-phosphorylated aliphatic and aromatic substrates have been developed.<sup>6</sup>

Ketoheptoses are rare seven-carbon sugars found in nature, that have significant pharmacological potential as inhibitors of sugar metabolism. For example, in the 1960s D-mannoheptulose was isolated from avocados and reported to inhibit glucose metabolism and insulin secretion *in vivo* and *in vitro*.<sup>7</sup> Following the identification of D-mannoheptulose as an inhibitor of glucokinase, interest in ketoheptoses for biomedical applications has increased and concise synthetic routes to these compounds are required.<sup>8</sup> Multistep syntheses of D-mannoheptulose and other ketoheptoses have been described, including the use of rearrangements and chain extensions resulting in the formation of diastereomeric mixtures and poor overall yields.<sup>9</sup> For example, the most recent approach reported the stereoselective preparation of D-manno-



**Scheme 1** TK reaction with lithium β-hydroxypyruvate (Li-HPA) **1**, highlighting the WT-TK *S*-stereochemistry in the α,α'-dihydroxyketone product.

<sup>a</sup>Department of Chemistry, University College London, 20 Gordon Street, London, WC1H 0AJ, UK. E-mail: h.c.hailes@ucl.ac.uk

<sup>b</sup>The Advanced Centre for Biochemical Engineering, Department of Biochemical Engineering, University College London, Bernard Katz Building, London, WC1H 0AH, UK

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heptulose and D-*gluco*-heptulose in eight synthetic steps and 56% overall yield, using toxic reagents.<sup>10</sup>

In a current multidisciplinary project, we are investigating the use of the waste biomass sugar beet pulp (SBP) for the synthesis of value-added chemicals. L-Arabinose 2, is one of the major sugars in this feedstock, and can be obtained from the hydrolysis of pectin.<sup>11</sup> The acceptance of L-arabinose as a substrate with TK would generate L-*gluco*-heptulose 3, a rare naturally-occurring ketoheptose with potential therapeutic applications in hypoglycaemia and cancer (Scheme 2).<sup>8</sup> Here we report for the first time the one-step biocatalytic preparation and characterisation of L-*gluco*-heptulose 3. TK libraries were screened through a medium-throughput assay and a colorimetric screening assay was also developed for high-throughput applications and successfully applied to identify productive enzymes. The use of a high-performing TK variant with L-arabinose was then used for the efficient stereoselective, one-step synthesis of L-*gluco*-heptulose 3 on a preparative laboratory scale.

## Results and discussion

### Identifying an improved transketolase for L-arabinose acceptance

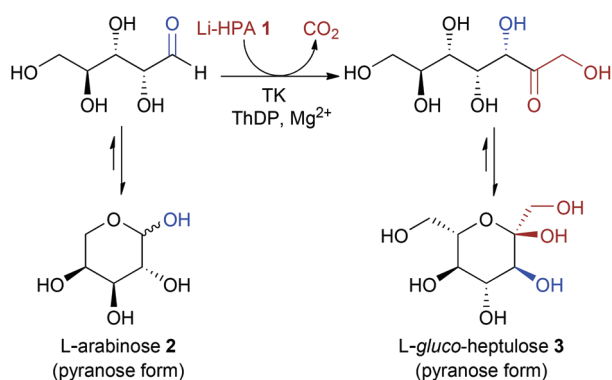
Compared to the natural aldehyde acceptors D-ribose-5-phosphate and erythrose-4-phosphate, TK in general has low affinity towards non-phosphorylated sugars.<sup>12</sup> In 1971, Axelrod *et al.* reported that L-arabinose 2 was accepted by wild-type (WT) *Saccharomyces cerevisiae* TK, with a  $K_M$  > 200-fold higher than the natural phosphorylated substrate D-ribose-5-phosphate ( $K_M$  120 mM and 0.4 mM for L-arabinose and D-ribose-5-phosphate respectively).<sup>13</sup> More recently, WT *S. cerevisiae* and *E. coli* TK specific activities of approximately 0.045 U mg<sup>-1</sup> were noted, more than 400-fold less than that for glycolaldehyde, using a pH based assay.<sup>14</sup> Studies to evolve *S. cerevisiae* TK for improved activities against D-ribose and D-glucose led to a double mutant R526Q/S252T that gave some of the highest improvements in specific activities (2.6-fold for D-ribose and 2.1-fold for D-glucose).<sup>15</sup> The pentose sugars D-ribose and

L-arabinose 2 share the same stereochemistry at C-2, but have the opposite stereochemistries at C-3 and C-4, and to the best of our knowledge no attempt to evolve TK towards the acceptor L-arabinose has been reported to date.

In our preliminary investigation of over 20 *E. coli* TK variants from our UCL-TK libraries, selected on the basis of previous studies using polyhydroxylated substrates, several single and double active-site mutants were identified that were able to accept L-arabinose 2, including WT (as previously reported), F434A and A29E/H461S (Table 1). For this smaller library a GC-MS protocol for identification of the heptulose product 3 was used (see Experimental).<sup>16</sup> GC-MS is particularly valuable in terms of sensitivity and product identification, as well as giving access to relative stereoselectivity data. The gas chromatogram for all the active mutants identified revealed a single product peak, suggesting a fully stereoselective reaction. The new stereocentre in the ketose product was assigned as the *S*-configuration (D-threo product), on the basis of the established WT-stereoselectivity for hydroxylated aldehyde substrates and a comparison of the characterization data for 3 to the enantiomer reported in the literature.<sup>3,10,12</sup> (see below and ESI†).

Although the variants showing activity towards 2 did not lead to a significant increase in activity compared to WT, it was possible to rationalise the substrate specificity observed. Notably, none of the D469X variants possessing either polar ionic (E) or uncharged (T and Y) residues led to active mutants, suggesting a crucial role for D469 in binding of the sugar 2. Computational docking analysis of L-arabinose 2 in the most abundant cyclic  $\alpha$ -pyranose (63% at equilibrium in water)<sup>17</sup> form into the active site of the *E. coli* wild-type TK (PDB ID: 1QGD) containing the ThDP-enamine intermediate (Fig. 1) highlighted a key interaction between the D469 residue and C-2 hydroxyl group of the acceptor.<sup>18</sup>

Docking analysis also provided further insights: the inner part of the substrate channel contains several His residues, and the middle section a Ser residue. These residues (H26, H261 and S385) interact with the C-1 and C-2 hydroxyl groups of the L-arabinose 2 *via* polar interactions, similar to those in the crystal structure with D-ribose-5-phosphate (R5P) in its cyclic five-membered ring form (cR5P) in the WT-TK active



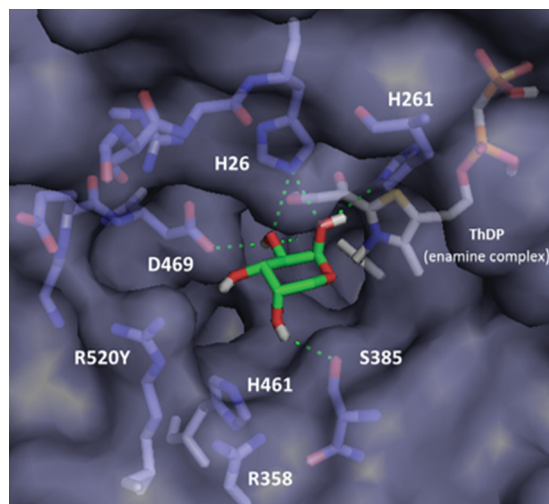
**Scheme 2** TK mediated synthesis of L-*gluco*-heptulose 3 from Li-HPA 1 and L-arabinose 2.

**Table 1** Preliminary screening of selected in house TK mutants<sup>a,b</sup>

TK <sub>E.coli</sub> variant	Heptulose 3	TK <sub>E.coli</sub> variant	Heptulose 3
WT	✓	F434A/R520Q	Traces
D469X (X = T, Y, E)	✗	A29E/H461S	✓
F434A	✓	A29E/R520V	Traces
H461S	✗	A29E/D469Y	✗
A29E	✗	H461S/R520Q	Traces
R520V	Traces	R358P/D469T	✗
D469X/R520Q (X = T, Y, E)	✗	S385X/D469 T/R520Q	✗

<sup>a</sup> Reaction conditions: Li-HPA 1 (33 mM), 2 (33 mM), ThDP (1.7 mM) and MgCl<sub>2</sub> (6 mM) in Tris-HCl (50 mM) at pH 7.0, with 33% v/v TK lysate for 24 h. <sup>b</sup> Traces corresponds to <5% conversion; ✓ corresponds to ~6–20% conversion; ✗ corresponds to no conversion.





**Fig. 1** Computational docking of *L*-arabinose **2** in the cyclic pyranose form into the active site of *E. coli* WT-TK (PDB ID: 1QGD) containing the ThDP-enamine intermediate.

site.<sup>14,18,19</sup> They can also contribute to maintaining the close proximity of the acceptor substrate to the nucleophilic centre on the ThDP-enamine intermediate in the correct orientation. In addition, the two Arg residues and the His residues in the outer part of the active site (R358, H461 and R520), interact with the phosphate moiety of cR5P *via* a mixture of hydrogen bonds and electrostatic interactions. These residues represent interesting sites for mutation to enhance the activity of TK toward non-phosphorylated long chain polyhydroxylated compounds, and have previously been highlighted in other mutational studies.<sup>6,14</sup> The same set of interactions (except for residue S385) was also identified in the docking analysis of *L*-arabinose in its acyclic form (see ESI†), which represent less than 0.1% at equilibrium in water.

On the basis of the docking calculation and preliminary data, high-throughput screening with three libraries of mutants previously prepared by saturation mutagenesis of the conserved residues (R358, H461, and R520) was performed.<sup>6</sup> Each library contained 90 randomly picked variants and three WT controls. Screening of the reaction plates, containing cell free lysate in each well, co-factors, Li-HPA **1** and *L*-arabinose **2** was performed using a new colorimetric assay.

### Development of a high-throughput colorimetric assay

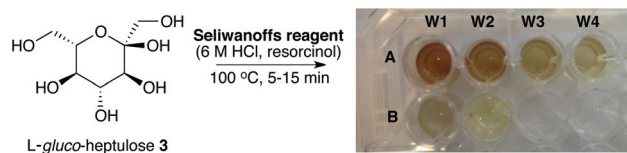
A crucial component in enzyme library screening is a rapid and reliable high-throughput method for identifying variants that catalyse the reaction. To date, colorimetric, fluorogenic and a colorimetric pH-based assay have been used to screen TK mutant libraries.<sup>14,20,21</sup> However these methods either cannot be used with  $\alpha$ -hydroxy aldehyde acceptors, are limited to a restricted substrate, or are based on the consumption of Li-HPA. Our aim was to develop an assay that could monitor the formation of ketose products in the TK reaction from *L*-arabinose **2** or other aldose sugars. Seliwanoff's test distinguishes

between ketoses and aldoses in a reaction using 6 M HCl and resorcinol.<sup>22</sup> Ketose dehydration under acidic conditions is followed by condensation with resorcinol to give a red xanthenone product. Aldoses, such as *L*-arabinose, however react more slowly than ketoses giving a yellow to green coloration. The reaction was therefore developed as a colorimetric assay to detect *L*-gluco-heptulose **3** formation in the presence of *L*-arabinose **2** *via* production of a red xanthenone. An assay for *L*-gluco-heptulose formation was performed (Scheme 3) using the Seliwanoff's reagent and solutions containing different concentrations of **3** (Scheme 3, line A). After heating, a red coloration was evident in wells containing **3**, and in addition a correlation could be observed between the amount of **3** present and intensity of the coloration generated. No coloration was observed in control reactions (also see experimental) containing *L*-arabinose alone or in combination with the co-factors (Scheme 3, line B). This qualitative assay was then used to rapidly identify mutants from the TKs libraries that accepted *L*-arabinose **2** to give **3**. After 24 h the TK reactions in 96-well plates were treated with Seliwanoff's reagent, and heated at 100 °C. Colour formation due to the presence of the ketosugar **3** (and conversion to a xanthenone) was observed within 15 min (see ESI†).

Visual analysis of the colorimetric assay plates (performed in duplicate) indicated that the *E. coli* TK libraries for R358, H461 and R520 contained active variants. In total, 31 active variants (17 from the R520 library, 11 from R358 library and 3 from the H461 library) were selected and sequenced. Among the R520 variants, two were revealed to be WT at a higher expression level while two mutations were identified for the other variants: R520Y and R520P. Similarly, three different mutants were identified from the 11 R358 variants sequenced: R358S, R358P and R358I. Finally, from the H461 library the single point mutant H461Y only was identified. In parallel a reaction plate for each library was also analysed by HPLC to validate the colorimetric screening method and confirmed the formation of **3**. The positive hits from the colorimetric assay and HPLC screening were furthermore repeated and analysed by GC-MS, which corroborated the formation of **3**.<sup>16</sup> The new TK assay is versatile and potentially applicable to a variety of sugar-like acceptor substrates, including hexoses.

### Determination of TK specific activities towards *L*-arabinose

All variants identified with the colorimetric assay were expressed and activities compared to *E. coli* WT-TK (Fig. 2).



**Scheme 3** Assay for *L*-gluco-heptulose **3**. Wells (W1–W4) line A correspond to the following concentrations of **3**: W1, 50 mM; W2, 30 mM; W3, 15 mM; W4, 5 mM; line B: W1, *L*-arabinose **2** (33 mM), ThDP (1.6 mM) and MgCl<sub>2</sub> (6 mM) and W2 *L*-arabinose **2** (33 mM).





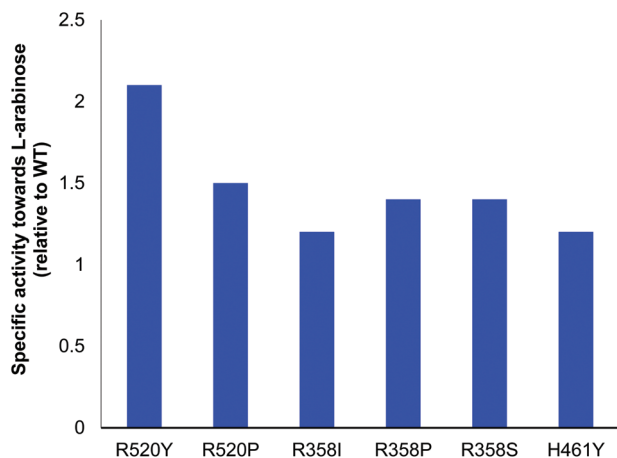


Fig. 2 *E. coli* TK specific activities ( $\mu\text{mol min}^{-1} \text{mg}^{-1}$ ) towards *L*-arabinose 1, relative to WT (clarified lysates) using Li-HPA 1 (33 mM), 2 (33 mM), ThDP (1.7 mM) and  $\text{MgCl}_2$  (6 mM) in Tris-HCl (50 mM) at pH 7.0. The specific activity of the WT-TK was  $0.021 \mu\text{mol min}^{-1} \text{mg}^{-1}$ .

Reactions were monitored using high-performance anion-exchange chromatography with pulsed amperometric detection (HPAC-PAD). The specific activities of the selected mutants R358P, R358S, H461Y, R520P and R520Y were greater than or comparable to WT-TK. The greatest improvements relative to WT were a 2.1-fold increase for R520Y, and 1.5-fold for R520P. In studies to evolve yeast TK for improved activities against *D*-ribose and *D*-glucose, R526Y (residue R526 in yeast TK is equivalent to R520 in *E. coli* TK) gave specific activities compared to WT of 1.5-fold and 0.9-fold, respectively. This highlights the variation in activities with different aldose stereochemistries and chain lengths.<sup>15</sup> The conversion of 2 to 3 was also performed for all 6 variants, and expression levels noted (Table 2).

The higher conversion yields for R520Y and H461Y reflected the generally higher reaction rates compared to the other mutants, although a higher amount of H461Y is needed with respect to R520Y. Replacement of the positively charged residues (R and H) with a hydrophobic and uncharged tyrosine

residue has previously been reported to enhance activity towards non-phosphorylated polyhydroxylated substrates.<sup>6</sup> Moreover, docking of *L*-arabinose in the linear form showed a possible interaction between the residue R520 and one of the OH groups (see ESI†), which may explain the higher specific activity for some R520 mutants.

### Preparative scale

*L*-gluco-Heptulose 3 is a rare naturally occurring ketoheptose. While synthesis of the corresponding enantiomer *D*-gluco-heptulose has been reported in eight-steps,<sup>10</sup> the synthesis and characterization of *L*-gluco-heptulose has not previously been reported. Variant H461Y was selected initially for the preparative-scale reaction due to its high conversion yields, and good enzyme stability. The reaction was stopped after 24 h and analysis of the crude product by <sup>1</sup>H NMR spectroscopy showed approximately a 50:50 ratio between the remaining *L*-arabinose 2 and the *L*-gluco-heptulose 3 formed. The crude mixture was then filtered through Amberlite® IR120 (hydrogen form) to remove both Tris buffer and some remaining 2 to give *L*-gluco-heptulose in 45% isolated yield after column chromatography, which was comparable to the conversion yield noted (Table 2). In a similar way the reaction was then carried out using R520Y mutant with a final 40% isolated yield.

### Stereoselectivity and product isolation

*L*-gluco-Heptulose 3 can exist as an equilibrium mixture of  $\alpha$ - and  $\beta$ -anomers. In the TK reaction one new stereogenic centre is formed, and spontaneous cyclisation can result in the generation of four isomers. However, GC-MS and NMR spectroscopy indicated that only one isomer was formed. Notably, NMR spectroscopic data of 3 recorded in  $\text{D}_2\text{O}$  showed the presence of a single pyranose isomer for *L*-gluco-heptulose: the <sup>1</sup>H and <sup>13</sup>C NMR data were the same as that previously reported for the enantiomer of 3, *D*-gluco-heptulose.<sup>10</sup> The large *trans*-coupling constant between the axial protons at 3-H and 4-H also confirmed that the hydroxyl group at C-3 was in an equatorial orientation giving an *S*-configured centre at this position.<sup>10</sup> The characterization data confirmed that the TK catalysed reaction showed exquisite diastereoselectivity leading to a single ketose with the *D*-threo configuration, and was consistent with the stereoselectivity observed in TK reactions with other aldose acceptors.<sup>12</sup> Only the more stable  $\alpha$ -anomer was generated and additionally, as previously reported, *L*-gluco-heptulose did not exhibit mutarotation behaviour: the optical rotation in aqueous solution remained constant over several days.<sup>10</sup> Furthermore, the absolute configuration of 3 was confirmed by correlation of the optical rotation to its enantiomer<sup>10</sup> and previous reports on the optical rotation of 3 in the literature.<sup>23</sup>

Table 2 Performance of *E. coli* TK mutants towards *L*-arabinose 2

TK	Conversion <sup>a</sup> /%	mg mL <sup>-1</sup> TK	Initial rate <sup>b</sup> (mM h <sup>-1</sup> )
WT	48	0.67	0.84
R520Y	54	0.48	1.27
R520P	31	0.28	0.54
R358I	22	0.23	0.36
R358P	35	0.48	0.84
R358S	48	0.75	1.31
H461Y	54	0.73	1.10

<sup>a</sup>Conversions were determined based on *L*-arabinose consumption after 24 h by high-performance anion-exchange chromatography with pulsed amperometric detection (HPAC-PAD). <sup>b</sup>Initial rates were determined by monitoring the production of 3 using HPAC-PAD. Reactions were performed using Li-HPA 1 (33 mM), 2 (33 mM), ThDP (1.67 mM) and  $\text{MgCl}_2$  (6 mM) in Tris-buffer (50 mM) at pH 7.0 and using clarified lysate (v/v 33%) in a total reaction volume of 900  $\mu\text{L}$ .

## Conclusions

An efficient TK-mediated asymmetric C–C bond forming reaction has been successfully applied to convert *L*-arabinose 1 into a ketoheptose with potential therapeutic applications in



hypoglycaemia and cancer. The ready availability of L-arabinose, a major component of pectin in SBP and renewable starting material, and one step synthesis of L-gluco-heptulose avoiding the use of toxic or hazardous solvents and reagents, could serve as a stimulus for the development of other green chemistry applications using SBP waste. In this study, several *E. coli* TK mutants with enhanced activity toward the non-phosphorylated sugar **2** have been identified for the first time, leading to the target product **3** in good yields with total stereo-control. A rapid high-throughput colorimetric screening assay to determine TK activity towards sugar-like substrates was also developed, which allowed the rapid identification of the higher performing mutants. The TK protocol developed could be performed on a preparative scale without further optimization, highlighting the potential for further synthetic applications in an industrial setting.

## Experimental

### General experimental and analytical methods

All chemicals were obtained from commercial source and used as supplied, except Li-HPA **1** which was prepared by modification of a previously described procedure.<sup>24</sup> TLC analysis was performed on Kieselgel 60 F254 precoated plates and compounds visualised by exposure to UV light, potassium permanganate or *p*-anisaldehyde stain. Flash column chromatography was carried out using silica gel (particle size 40–63  $\mu\text{m}$ ). NMR:  $^1\text{H}$  and  $^{13}\text{C}$  NMR spectra were recorded at 298 K at the field indicated using Bruker Avance 500 and Avance 600 machines. Coupling constants were measured in Hertz (Hz) and peaks referenced to the deuterated solvent used. Infrared spectra were recorded on a Perkin Elmer Spectrum 100 FTIR spectrometer.

**Analytical HPLC (TK reaction plates analysis).** Analysis of the plates were performed using a Dionex X500 HPLC (Camberley, UK) fitted with a Famos 120 autosampler, an Ultimate 3000 generation II isocratic pump, a LC30 Chromatography column oven, AD20 Absorbance detector and refractive index (RI) detector (Shodex RI-101 or ERC Refractomax 520). Analysis was performed using an Biorad Aminex HPX-87H column  $7.8 \times 300$  mm. Isocratic elution was carried out at  $0.6 \text{ mL min}^{-1}$  isocratic flow of 0.1% TFA (v/v) as mobile phase for 30 min, a column temperature of  $60^\circ\text{C}$ , injection volume of  $20 \mu\text{L}$  and monitoring by UV at 210 nm, and RI, for the identification of L-gluco-heptulose (retention time (RT) 8.7 min).

**Gas chromatography-mass spectrometry (GC-MS).** GC was performed on a ThermoScientific TRACE 1310 by using a capillary column ( $30 \text{ m} \times 0.25 \text{ mm}$ ,  $0.25 \text{ mm}$ , TRACE TR-5MS, ThermoScientific). The samples were injected with a TriPlus RSH autosampler and applied by splitless injection at an injection temperature of  $220^\circ\text{C}$ , with a split flow of  $50 \text{ mL min}^{-1}$  and an oven temperature of  $50^\circ\text{C}$ . After 0.5 min, the temperature of the oven was raised at a rate of  $35^\circ\text{C min}^{-1}$  to  $280^\circ\text{C}$  and held for 5 min. The flow of the carrier gas, helium, was  $1.2 \text{ mL min}^{-1}$ . MS analysis was performed with an ISQ single quadru-

pole (ThermoScientific) in tandem with the GC. MS data were acquired in  $m/z$  70–800 range with scan time 0.2 s,  $70 \text{ eV}$  and EI mode.

**High performance anion-exchange chromatography with pulsed amperometric detection (HPAC-PAD).** Quantitative analysis of L-arabinose **2**, L-gluco-heptulose **3** and Li-HPA **1** was performed using a Reagent-Free Ion Chromatography System (ICS 5000+, Dionex, Sunnyvale, CA, USA) with a Dionex Aminopac™ PA1 anion exchange column  $4 \times 250$  mm fitted with a Dionex Aminopac™ PA1 guard column  $4 \times 50$  mm, an injection volume of  $10 \mu\text{L}$ , an electrochemical detector system, and an eluent generator fitted with a KOH 500 cartridge. Analysis was carried out at isocratic flow using 15 mM KOH as the mobile phase with a flow rate of  $1.5 \text{ mL min}^{-1}$  for 18 min at  $30^\circ\text{C}$ . Standard calibration curves with L-arabinose, HPA and L-glucoheptulose (isolated from the preparative scale reaction) were used for quantification purposes. Retention times for **2**, **3** and **1** were 6.5 min, 8.3 min and 11.8 min respectively.

### Transketolase preparation

**Transketolase expression.** Selected TK glycerol stocks (pQR791, XL-10 Gold) from the UCL TK library were used to inoculate 10 mL of Luria-Bertani (LB) broth containing ampicillin ( $150 \mu\text{g mL}^{-1}$ ) and incubated for 16–18 h ( $37^\circ\text{C}$ , 250 rpm). The cells were then harvested or transferred to a larger volume of fermentation. For larger scale growths, 10% inoculum volume was used and the fermentation was performed at the same conditions for 6 h before harvesting. To harvest cells, the culture was centrifuged (4000 rpm, 10 min) and the cell pellet was used fresh or stored at  $-20^\circ\text{C}$ .

**Transketolase lysate preparation.** TK cell pellets were re-suspended in Tris-HCl buffer (50 mM, pH 7.0) and lysed by sonication on ice (10 s on and 15 s off for 10 cycles). The insoluble portion of the lysate was pelleted by centrifugation (4000 rpm, 30 min) and the supernatant used fresh or stored at  $-20^\circ\text{C}$  in aliquots for up to one month.

### Transketolase preliminary library investigations

Transketolases mutants (Table 1) were investigated for activity with L-arabinose **2** and Li-HPA **1** via the formation of L-gluco-heptulose **3**. Reaction conditions: Li-HPA **1** (33 mM), **2** (33 mM),  $\text{MgCl}_2$  (6 mM) and ThDP (1.7 mM) in Tris-HCl buffer (50 mM, pH 7.0) and using 33% v/v TK lysate. The reactions ( $300 \mu\text{L}$ ) were incubated at  $25^\circ\text{C}$  with 450 rpm orbital rotation.

**GC-MS derivatization protocol.**<sup>15</sup> After 24 h reaction,  $15 \mu\text{L}$  of D-glucose (50 mM solution in water) was added and the sample freeze dried. The residue was dissolved in dry pyridine ( $150 \mu\text{L}$ ) and a large excess of *N,O*-bis(trimethylsilyl)trifluoroacetamide (BSTFA) with 1% trimethylsilyl chloride (TMSCl) added ( $150 \mu\text{L}$ ). The reaction mixture was shaken at room temperature for 40–60 min before adding  $0.7 \text{ mL}$  HPLC grade EtOAc. After centrifugation to remove the precipitated salts, the sample was analyzed by GC-MS. Per(trimethylsilylated) L-gluco-2-heptulose in the  $\alpha$ -pyranose form, exhibited a single GC peak (RT 7.26 min);  $m/z$  204 and 539  $[\text{M} - \text{CH}_2\text{OSiMe}_3]^+$ .<sup>25</sup>



An  $m/z$  204 has been reported to be characteristic of trimethylsilylated pyrano-ring sugars.<sup>26</sup>

### Docking experiments

ThDP-enamine and L-arabinose 2 (cyclic  $\alpha$ -pyranose and linear form) structures were drawn in Chem3D Ultra v.13.0 and energy minimised by MM2 calculations. The energy-minimised ThDP-enamine was docked into the active site.<sup>6d,27</sup> A rigid docking for the TK structure (PDB: 1QGD, *E. coli*) against 2 was performed using ligand docking and binding site analysis with PyMOL and Autodock Vina<sup>28,29</sup> with the grid centered at  $-11.25$ ,  $25.858$ ,  $40.198$  and a grid size of  $28 \text{ \AA} \times 28 \text{ \AA} \times 28 \text{ \AA}$ . Conformations were selected on the basis of the lowest energy (standard deviation for the 9 conformations of the cyclic and linear forms was  $0.2$ ) and for the cyclic form from visual analysis and proximity to the ThDP-enamine. The binding affinity for cyclic L-arabinose (in Fig. 1) was  $-4.7 \text{ kcal mol}^{-1}$  and for the linear form (in ESI Fig. 3†)  $-4.8 \text{ kcal mol}^{-1}$ .

### Colorimetric assay: assaying for TK-mediated synthesis of L-glucio-heptulose 3 (preliminary assay)

A 20 mL reaction mixture containing ThDP ( $1.7 \text{ mM}$ ),  $\text{MgCl}_2$  ( $6 \text{ mM}$ ) and cell-free lysate (33% total reaction volume) in Tris-HCl ( $50 \text{ mM}$ , pH 7.0) was incubated at  $25^\circ\text{C}$  under slow magnetic stirring. After 20 min, 10 mL of the substrate solution (1 and 2,  $100 \text{ mM}$  at pH 7.0) was added and the reaction mixture incubated at  $25^\circ\text{C}$  for 24 h. At time 0 and 24 h,  $100 \mu\text{L}$  of the reaction mixture were quenched with 1 mL of Seliwanoff's reagent in a closed test tube and heated at  $100^\circ\text{C}$ . After 5 minutes a cherry red coloration appeared in the solution quenched after 24 h, which became more intense after 10 min at  $100^\circ\text{C}$ . No coloration was evident for the reaction quenched at 0 h, indicating there was no formation of 3 and no coloration interference from any of the reaction components. Moreover, three control experiments without either TK, 1 or 2, were performed in parallel and no coloration after treatment with Seliwanoff's reagent and heating was observed.

### Library selection for activity

**Preparation of the reaction plates.** From the master plates, 96 deep-square well (DSW) plates containing LB broth ( $0.8 \text{ mL}$  with  $100 \mu\text{g mL}^{-1}$  ampicillin) were inoculated, using a colony replicator, and incubated for 24 h at  $37^\circ\text{C}$  with 85% humidity with gas-permeable sealing. The plates were centrifuged for 10 min at  $4000 \text{ rpm}$  and  $4^\circ\text{C}$  (polypropylene sealing). The supernatant was removed and the pellets stored at  $-80^\circ\text{C}$  or submitted directly to cell lysis. A commercial cell lysis reagent containing lysozyme was used to obtain the lysates. The pellets were re-suspended in  $150 \mu\text{L}$  of BugBuster Protein Extraction Reagent containing lysozyme and incubated for 20 min at room temperature (plate shaker,  $400 \text{ rpm}$ ) then centrifuged 35 min at  $4000 \text{ rpm}$  and  $4^\circ\text{C}$ . The supernatants were transferred into 96-well reaction plates and then used. The reaction plates, containing  $50 \mu\text{L}$  of lysate in each well, were directly submitted to the screening reactions by adding cofactors ( $50 \mu\text{L}$  stock solution  $\text{MgCl}_2$  ( $18 \text{ mM}$ ) and ThDP ( $5 \text{ mM}$ )). After

incubation for 20 min at room temperature, the reaction was initiated by addition of the substrates ( $50 \mu\text{L}$  stock solution 1 ( $100 \text{ mM}$ ) and 2 ( $100 \text{ mM}$ ) in Tris-HCl ( $50 \text{ mM}$ ), pH 7.0). After 24 h the plates were analyzed either by HPLC or GC-MS or the colorimetric assay.

**HPLC assay.** After 24 h the plates were diluted and directly analysed by HPLC following the protocol described above. The positive hits from primary screening were re-evaluated by reaction at  $25^\circ\text{C}$  as before but in a volume of  $300 \mu\text{L}$  and analysed by GC-MS following the protocols described above.

**Colorimetric screening.** A Seliwanoff's reagent solution ( $1 \text{ mg mL}^{-1}$  resorcinol in  $6 \text{ M HCl}$ ) was prepared. After 24 h,  $30 \mu\text{L}$  of the TK reaction was quenched by addition of  $120 \mu\text{L}$  of Seliwanoff's reagent to each well, and the plate was incubated at  $100^\circ\text{C}$ . Colour formation due to the presence of a ketosugar was observed after 10 min. Colorimetric screening of the TK mutants was carried out in duplicate for each library.

### Specific activities of selected mutants

Glycerol stocks of selected mutants were streaked from the library master plate onto LB agar containing  $150 \mu\text{g mL}^{-1}$  ampicillin. Individual colonies were selected and grown in  $10 \text{ mL}$  LB with  $150 \mu\text{g mL}^{-1}$  ampicillin for 18 h at  $200 \text{ rpm}$ ,  $37^\circ\text{C}$ . The cells were then transferred to a larger volume of fermentation and the cell lysate was prepared as previously described. The specific activities of selected TK mutants were obtained for reactions of 1 ( $33 \text{ mM}$ ) with 2 ( $33 \text{ mM}$ ), ThDP ( $1.7 \text{ mM}$ ),  $\text{MgCl}_2$  ( $6 \text{ mM}$ ), in Tris-HCl ( $50 \text{ mM}$ ) at pH 7.0. Lysate (33% (v/v)) was used in duplicate or triplicate reactions with a final volume of  $900 \mu\text{L}$ . At a time point during the first 24 h,  $50 \mu\text{L}$  of the solution was sampled and quenched by adding  $50 \mu\text{L}$  of  $\text{H}_2\text{SO}_4$  ( $0.5\%$  v/v aqueous solution). These samples were then analysed by HPAC-PAD as described above. TK concentrations were determined by SDS-PAGE densitometry as previously described for soluble TK after clarification.<sup>6d</sup> Densitometry of samples electrophoresed on a 12% SDS-Page gel (Bio-Rad) was used. The Coomassie stained gel was imaged using Imagesoft software to calculate the TK band density. Total protein in the samples was determined using a standard Bradford assay. A range of concentrations of commercial BSA was run in each SDS-Page gel, and used to calculate a standard curve based on integrated optical density for calibration of the enzyme concentration.

### Preparative scale biocatalytic reaction

A 20 mL reaction mixture containing ThDP ( $1.7 \text{ mM}$ ),  $\text{MgCl}_2$  ( $6 \text{ mM}$ ) and cell-free lysate (33% total reaction volume,  $0.39$  and  $1 \text{ mg mL}^{-1}$  TK final concentration for R520Y and H461Y respectively) in Tris-HCl ( $50 \text{ mM}$ , pH 7.0) was incubated at  $25^\circ\text{C}$  under slow magnetic stirring. After 20 min,  $10 \text{ mL}$  of the substrate solution (1 and 2,  $100 \text{ mM}$  at pH 7.0 in Tris-HCl buffer ( $50 \text{ mM}$ )) was added and the reaction mixture incubated at  $25^\circ\text{C}$ . After 24 h,  $60 \text{ mL}$  of MeOH was added and the white precipitate formed removed by centrifugation ( $10 \text{ min}$ ,  $4000 \text{ rpm}$ ). The sample was then filtered twice through a strong cationic resin (Amberlite® IR-120 hydrogen form,  $40 \text{ mL}$ ) to





remove both Tris buffer and some 2. The filtrate, containing *L*-gluco-heptulose **3**<sup>23</sup> was dry-loaded on a silica gel column chromatography (EtOAc/2-PrOH/H<sub>2</sub>O) to give *L*-gluco-heptulose **3** as a colourless oil (95 mg, 45% for H461Y; 80 mg, 40% for R520Y). *R*<sub>f</sub> 0.33 (EtOAc/2-PrOH/H<sub>2</sub>O, 7 : 3 : 1); [ $\alpha$ ]<sub>D</sub><sup>20</sup> −67.0 (c 1, H<sub>2</sub>O), lit., [ $\alpha$ ]<sub>D</sub><sup>20</sup> −67.8 (c 2.5, H<sub>2</sub>O);<sup>23</sup> IR (film/cm<sup>−1</sup>) 3314, 2925, 2469, 1419; <sup>1</sup>H NMR (D<sub>2</sub>O; 600 MHz) 3.37 (1H, t, *J* 9.6 Hz, 5-H), 3.46 (1H, d, *J* 9.6 Hz, 3-H), 3.49 (1H, d, *J* 12.0 Hz, 1-HH), 3.65 (1H, d, *J* 12.0 Hz, 1-HH), 3.68 (1H, t, *J* 9.6 Hz, 4-H), 3.70–3.75 (2H, m, 6-H, 7-HH), 3.79 (1H, m, 7-HH); <sup>13</sup>C NMR (D<sub>2</sub>O; 151 MHz) 61.3 (C-7), 64.3 (C-1), 70.2 (C-5), 70.9 (C-3), 72.9 (C-6), 74.3 (C-4), 98.2 (C-2); *m/z* HRMS (ES<sup>+</sup>) found [MNa]<sup>+</sup> 233.0645, C<sub>7</sub>H<sub>14</sub>O<sub>7</sub>Na requires 233.0637.

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