

Cite this: *Chem. Sci.*, 2024, 15, 5950

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Investigation on the substrate specificity and *N*-substitution tolerance of PseF in catalytic transformation of pseudaminic acids to CMP-Pse derivatives†

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Pseudaminic acid (Pse) belongs to a class of bacterial non-2-ulosonic acids, and has been implicated in bacterial infection and immune evasion. Various Pse structures with diverse *N*-substitutions have been identified in pathogenic bacterial strains like *Pseudomonas aeruginosa*, *Campylobacter jejuni*, and *Acinetobacter baumannii*. In this study, we successfully synthesized three new Pse species, including Pse5Ac7Fo, Pse5Ac7(3_RHb) and Pse7Fo5(3_RHb) using chemical methods. Furthermore, we investigated the substrate specificity of cytidine 5'-monophosphate (CMP)-Pse synthetase (PseF), resulting in the production of *N*-modified CMP-Pse derivatives (CMP-Pses). It was found that PseF was promiscuous with the Pse substrate and could tolerate different modifications at the two nitrogen atoms. This study provides valuable insights into the incorporation of variable *N*-substitutions in the Pse biosynthetic pathway.

Received 1st February 2024

Accepted 20th March 2024

DOI: 10.1039/d4sc00758a

rsc.li/chemical-science

Introduction

Bacterial non-2-ulosonic acids, including pseudaminic acid (Pse), legionaminic acid (Leg), acinetaminic acid (Aci), fusaminic acid, and their epimers, have been identified from pathogenic bacteria, and play crucial roles in bacterial morphology, bacterial pathogenicity, immune escape, and other related processes.^{1–6} Pse derivatives present as key components of surface glycans in various clinical pathogens, which are widely distributed in lipopolysaccharides (LPS),⁷ capsular polysaccharides (CPS),^{8,9} exopolysaccharides (EPS),¹⁰ pili,¹¹ and flagella.¹² Unlike the case of sialic acids in mammalian cells, Pse species bear variable *N*5/*N*7-substitutions and are connected to other sugars through both α and β linkages (Fig. 1). The fascinating structures and less discovered biological functions of these compounds make them attractive

subjects for investigation in the field of synthetic glycobiology.^{13–15}

The biosynthetic pathway and associated enzymes for Pse5Ac7Ac **1** have been elucidated clearly by Logan *et al.*, including PseB, PseC, PseH, PseG and PseI (Fig. 2a).¹⁶ Based on this, Fascione *et al.* developed the enzymatic synthetic approach of Pse5Ac7Ac **1** and CMP-Pse5Ac7Ac **7**,¹⁷ and realized the enzymatic glycosylation of CMP-Pse5Ac7Ac by promiscuous sialyltransferases¹⁸ and Pse-specific glycosyltransferase KpsS1.¹⁹ The identification of novel formyltransferase PseJ²⁰ and amidine synthase PseA²¹ filled in the map of the biosynthetic pathway of Pse5Ac7Fo and Pse5Ac7Am, respectively. However, unlike sialic acid, which can be produced *via* industrial scale fermentation, the variable *N*-substitutions of Pse species and largely unknown enzymes for *N*-modifications make the biosynthetic approach more challenging, especially for large-scale *in vitro* biosynthesis.

To investigate the biological significance of Pse at a molecular level, we started a research program to chemically synthesize Pse and its derivatives,^{22,23} with which we develop probes²⁴ and explore their applications as synthetic antibacterial vaccines.²⁵ On the other forefront, we aim to use synthetic Pse derivatives to study the biosynthesis and biological function of Pse. It is noted that, after the biosynthesis of Pse, the pseudaminic acid cytidyltransferase PseF is responsible for converting Pse into corresponding cytidine 5'-monophosphate (CMP) derivatives,^{16,26} which is then transferred to other glycans to form Pse glycosides. At this stage, the Pse-specific glycosyltransferase has just been identified.¹⁹ Furthermore, natural Pse carries diverse *N*-acylations at two nitrogen atoms, which makes

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† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d4sc00758a>

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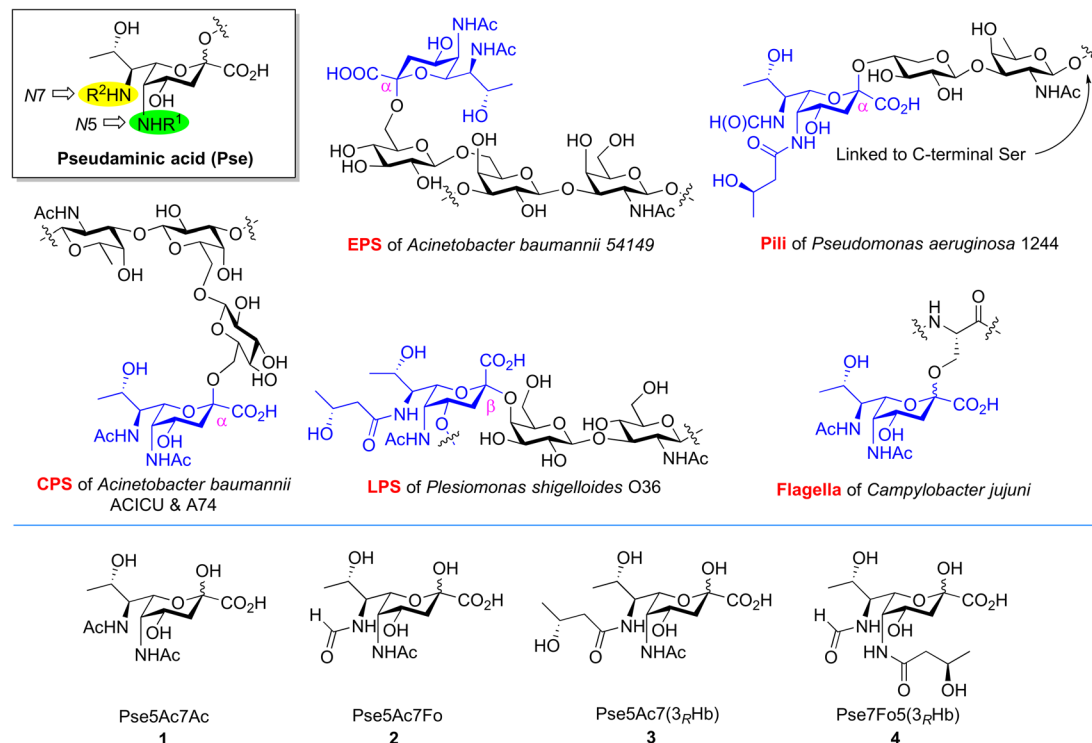


Fig. 1 Occurrence of Pse in bacterial glycans bearing different *N5/N7*-substitutions and structures of Pse species selected for synthesis and enzymatic transformation studies.

the process more complicated. The promiscuity and specificity of PseF and the glycosyltransferase, as well as the order of *N*-acylation and transglycosylation are open questions. To this end, we started to answer the question related to PseF by using *N*-modified Pse derivatives, which were only available *via* chemical synthesis at this stage. Given that the two amino groups in Pse derivatives are naturally modified by variable *N*-substitutions, in this study, we first developed chemical synthetic approaches to prepare three new naturally existing Pse derivatives 2–4 selected from bacterial glycans that were modified with acetyl, formyl and (*R*)-3-hydroxybutyryl on *N5/N7* sites, respectively, followed by the enzymatic studies of PseF with Pse derivatives 1–4 to form the corresponding Pse-CMP compounds 7–10 (Fig. 2b). We showed for the first time that PseF was well compatible with Pse derivatives *in vitro*, which indicates the variable *N*-substitution installation most likely occurs before the formation of pseudaminosides in the biosynthetic pathway. Deeper understanding of the Pse biosynthetic pathway may accelerate the development of novel inhibitors to fight pathogenic Gram-negative bacteria.

Results and discussion

Chemical synthesis of Pse5Ac7Fo, Pse5Ac7(3_RHb) and Pse7Fo5(3_RHb)

In 2017, we have reported the *de novo* total synthesis of Pse5Ac7Ac **1** from the 5*N*-Troc/7*N*-Cbz thioglycoside donor.²² The orthogonal protections installed on *N5* and *N7* sites during this *de novo* approach perfectly fit our needs in the synthesis of Pse

derivatives bearing different *N5/N7*-substitutions. Herein, starting from the 5*N*-Troc/7*N*-Cbz *S*-adamantyl thioglycoside donor **11** developed in 2019,²³ we were able to complete the synthesis of Pse5Ac7Fo **2**, Pse5Ac7(3_RHb) **3** and Pse7Fo5(3_RHb) **4**, respectively.

As illustrated in Scheme 1, the synthesis of Pse5Ac7Fo **2** started from the β -selective glycosylation of donor **11** with benzyl alcohol. The benzyl group installed at the glycosidic site of **12** served as a protecting group during the following synthetic steps. *N5*-Troc was then transformed to *N5*-Ac *via* Zn mediated orthogonal deprotection and *N*-acetylation. During the modification of *N7*, to avoid the *O8*-to-*N7* acetyl transfer when removing Cbz, the *O4/O8*-acetyl groups in **13** were first removed by treating with 5 mM MeONa in MeOH. The obtained intermediate **14** containing methyl ester generated by partial isopropyl/methyl exchange was used in the following transformations, since both ester groups would be hydrolyzed at the end stage. Next, the selective hydrogenolysis of Cbz without removing glycosidic OBn was achieved using the pyrimidine deactivated Pd/C catalyst (see ESI† for details) after extensive screening. Several nitrogen-containing compounds have been developed as deactivators to regulate the activity of the Pd/C catalyst.^{27–31} In our case, other deactivators (*e.g.*, ammonium acetate, tertiary amine, ethylenediamine, pyridine, 2,2'-bipyridine) gave inferior chemoselectivity. After *N7*-formylation, the isopropyl/methyl ester in **15** was hydrolyzed by LiOH, and Pse5Ac7Fo **2** was obtained in 79% yield after Bn removal. We also tried to avoid the methyl ester contamination by directly removing *N7*-Cbz with *O8*-Ac intact, since the *O8*-to-*N7* acyl





Fig. 2 Enzymatic and chemoenzymatic syntheses of CMP-Pse bearing variable N5/N7-substitutions. (a) Fascione's enzymatic synthesis of CMP-Pse. (b) Chemoenzymatic synthesis of N5/N7-modified CMP-Pse in this work.

transfer could be partially suppressed by shortening the reaction time. Unfortunately, product **16** was obtained in only 25% yield, which was inferior to the above 3-step process (50% for **15** from **13**). This established approach was also used to synthesize Pse5Ac7(3_RHb) **3**, where symmetric anhydride **17** was used for N7-acylation instead of formic anhydride.

For the synthesis of Pse7Fo5(3_RHb) **4**, the approach we developed during the total synthesis of the *Pseudomonas aeruginosa* 1244 pilin trisaccharide²² was adopted, where the O4-to-N5 acyl transfer was used to install the N5-(R)-3-hydroxybutyryl group. As illustrated in Scheme 2, the two acetyl groups in thioglycoside **11** were removed under acidic conditions to avoid



Scheme 1 Synthesis of the Pse5Ac7Fo and Pse5Ac7(3_RHb).



Scheme 2 Synthesis of the Pse7Fo5(3_R)Hb.

destroying the Troc group. Two (*R*)-3-hydroxybutyryl groups were installed on the *O*4/*O*8 hydroxyl groups in **19**, followed by β -selective glycosylation of **20** with benzyl alcohol. Intermediate **21** was subjected to the TBAF mediated Troc removal conditions developed by Fukase.³² The key carbamoyl fluoride intermediate proposed by Coudert *et al.*³³ was hydrolyzed *in situ* by the trace amount of H₂O to release the amino group for acyl transfer. The addition of acetic acid to partially buffer the basicity of the fluoride anion significantly suppressed side reactions, and mild heating ensured full conversion in 48 hours. To install the *N*7-formyl group, the Cbz was selectively removed by hydrogenolysis, and the amino group was formylated immediately to minimize *O*8-to-*N*7 acyl transfer. The 3-step process used in the synthesis of **2** and **3** was abandoned here due to significant side reactions during MeONa treatment. Intermediate **23**, isolated in moderate 31% yield, was then subjected to saponification and hydrolysis to give rise to Pse7Fo5(3_R)Hb **4** in 76% yield.

PseF catalyzed enzymatic synthesis of *N*5/*N*7-modified CMP-Pse

To evaluate the substrate tolerance of PseF, we designed the single step catalytic activity assay using purified PseF and synthetic *N*-substituted Pse derivatives **1**–**4**. The full-length *pseF* gene was cloned from *Acinetobacter baumannii* clinical isolates and the recombinant PseF protein containing a His tag was expressed in *E. coli* BL21 cells and purified *via* a nickel column (see ESI† for details).

With the recombinant PseF protein in hand, we validated its catalytic activity in the reported model reaction. According to the conditions suggested by Logan *et al.*¹⁶ and Fascione *et al.*,¹⁷ we observed smooth formation of CMP-Pse5Ac7Ac **7** by treating synthetic Pse5Ac7Ac **1** and CTP with PseF. In contrast, no conversion was observed when sialic acid was tested as the substrate, which showcased the Pse-specificity of the enzyme. In

order to explore the substrate tolerance of PseF towards other *N*-substituted Pse derivatives in a quantitative way, we first conducted quantitative kinetic analysis of this enzymatic transformation using a HILIC column. As depicted in Fig. 3, a rapid enzymatic reaction rate was revealed. The conversion of Pse5Ac7Ac **1** to CMP-Pse5Ac7Ac **7** reached up to 66% within a 30 minutes timeframe. Further time-dependent data are presented in the ESI (Fig. S2).†

To investigate the influence of metal ions on the activity of PseF, we conducted initial experiments (Fig. 4). PseF was subjected to assays using Pse5Ac7Ac **1** (0.9 mM) and CTP (2 mM) as substrates in the presence of various metal cations or EDTA (20 mM) as a metal ion sequestering agent. The reactions were carried out in a sodium phosphate buffer (30 mM, pH 7.4) for

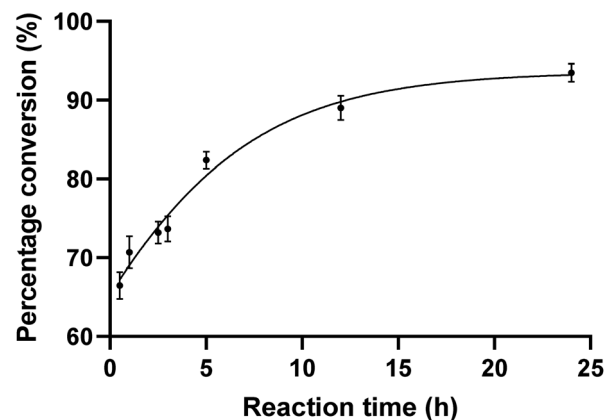


Fig. 3 CMP-Pse5Ac7Ac percentage conversion (%) at varying reaction times. PseF (3.8 μ M) was assayed using Pse5Ac7Ac (0.9 mM), CTP (2 mM) as substrates in the presence of MgCl₂ (20 mM) and sodium phosphate (30 mM, pH 7.4) at 37 °C for 0.5, 1, 2.5, 3, 5, 12, and 24 h, respectively. The error bars were derived from three parallel replicates of each experiment.



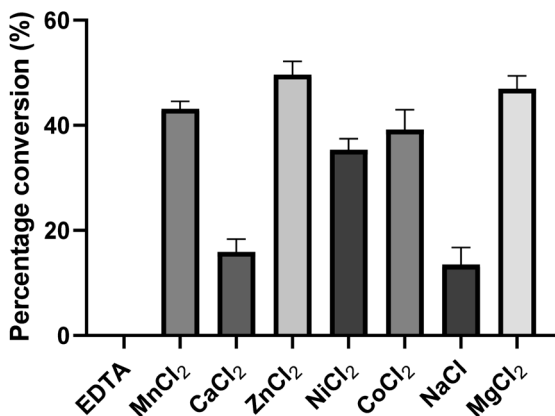


Fig. 4 Metal ion effects of PseF. PseF (2.27 μM) was assayed using Pse5Ac7Ac (0.9 mM), CTP (2 mM) as substrates in the presence of a metal cation or EDTA (20 mM), and sodium phosphate (30 mM, pH 7.4) at 37 $^{\circ}\text{C}$ for 5 minutes. Note: 5 minutes is a suitable time to compare their differences. The error bars were derived from three parallel replicates of each experiment.

a duration of 5 minutes. The obtained results clearly demonstrated the requirement of metal cations for the enzymatic activity of PseF, as no activity was observed in the presence of EDTA. Among the Mn^{2+} , Ca^{2+} , Zn^{2+} , Ni^{2+} , Co^{2+} , Na^{+} and Mg^{2+} , only Ca^{2+} and Na^{+} gave obviously lower efficiency. In contrast, Zn^{2+} and Mg^{2+} gave 49% and 47% conversion, respectively. To keep the consistency with reported conditions,^{16,17} Mg^{2+} was chosen as the metal ion in the following assays. Mg^{2+} is known to be involved in the enzymatic generation of sugar-UDP³⁴ and sugar-CMP³⁵ species under physiological conditions (intracellular $c[\text{Mg}^{2+}] \sim 1.5 \text{ mM}$ ³⁶). According to Pohl's report, the strong binding of Mg^{2+} to UTP led to the Mg^{2+} -UTP complex serving as the active substrate of the uridylyltransferase enzymes in the sugar-UDP synthesis.³⁷ In the PseF catalyzed Pse-CMP synthesis, a similar role of Mg^{2+} can be proposed. The pH and temperature profiles of PseF are described in Fig. S3 and S4.[†] Moreover, to identify a suitable concentration of CTP, we measured the

reaction rates under fixed $c(\text{Pse5Ac7Ac})$ concentration with varied $c(\text{CTP})$. As illustrated in Fig. S5,[†] $c(\text{CTP})$ higher than $K_m(\text{CTP})$ ($0.6295 \pm 0.065 \text{ mM}$) should be used to ensure rapid conversion.

With the optimized conditions (0.9 mM Pse analogue, 0.1 mg mL^{-1} PseF, 6 mM CTP, 20 mM MgCl_2 , pH 7.4, 37 $^{\circ}\text{C}$) in hand, we tested the *N*-substitution tolerance of PseF using our synthetic Pse derivatives 1–4. We extended the incubation time to 24 hours in each case to maximize the conversion, followed by isolation of the four CMP-Pse derivatives 7–10 through a Bio-Gel P-2 resin column. The isolated yields of CMP-Pse5Ac7Ac 7, CMP-Pse5Ac7Fo 8, CMP-Pse5Ac7(3_RHb) 9 and CMP-Pse7Fo5(3_RHb) 10 were 72%, 72%, 70% and 24%, respectively. For compounds 7–9, 1.1–1.2 mg products could be obtained from 2.3–2.7 μmol scale transformations. This is the first report to show that PseF is compatible with other *N*-substituted Pse derivatives *in vitro*, in addition to Pse5Ac7Ac. This result indicates that PseF is a promiscuous enzyme with the Pse substrate, showing higher tolerance to *N*7-substitutions than *N*5. Also, the installation of variable *N*-substitution most likely occurs before the formation of CMP-Pse and pseudaminosides in the biosynthetic pathway.

To gain a comprehensive understanding of PseF's characteristics and to elucidate the underlying reasons for the relatively low yield of CMP-Pse7Fo5(3_RHb), measurement of PseF catalytic kinetics was conducted. To determine the apparent kinetic parameters for PseF activity, we selected a time point of 15 seconds based on Fig. S2,[†] where the percentage conversion of CMP-Pse5Ac7Ac was 7% using 0.9 mM of Pse. This early time point was chosen to mimic the initial velocity and facilitate the calculation of PseF apparent kinetic parameters. Once the pre-determined time was reached, the reaction samples were immediately quenched by MeCN and subsequently subjected to UPLC analysis on a HILIC column. A high concentration of CTP (2 mM) giving rapid conversion at high $c(\text{Pse})$ could lead to a narrow time window (shorter than 15 s) for initial velocity measurement, which was difficult for us to achieve reliable accuracy. Thus, PseF was assayed using a fixed moderate concentration of CTP (0.8 mM) and varying Pse concentrations

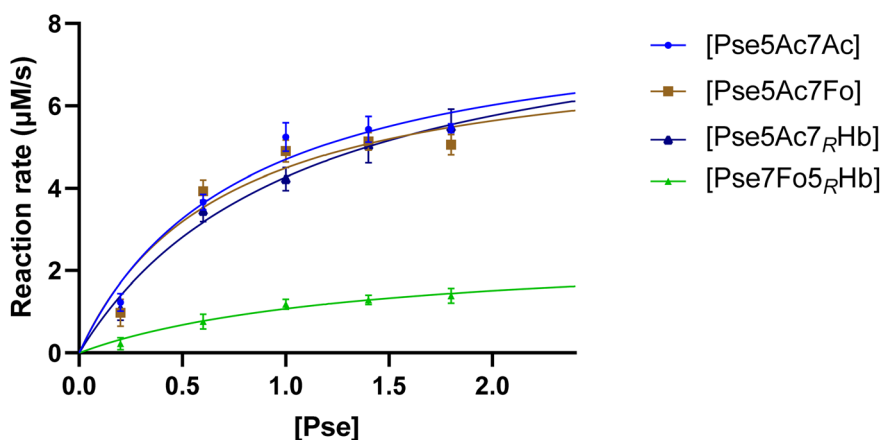


Fig. 5 PseF kinetic assay results at varying Pse concentrations. PseF (2.27 μM) was assayed using a fixed concentration of CTP (0.8 mM) and varying Pse concentrations (ranging from 0.2 to 1.8 mM) in the presence of MgCl_2 (20 mM) and sodium phosphate (30 mM, pH 7.4) at 37 $^{\circ}\text{C}$ for 15 seconds. The error bars were derived from three parallel replicates of each experiment.



Table 1 Calculated apparent kinetic parameters of PseF for the catalytic transformation of Pse derivatives 1–4^{a,b}

| Pse | V_{\max} ($\mu\text{M s}^{-1}$) | K_m (mM) | k_{cat} (s^{-1}) | k_{cat}/K_m ($\text{s}^{-1} \text{mM}^{-1}$) |
|----------------------------|-------------------------------------|-------------------|--------------------------------------|---|
| Pse5Ac7Ac | 8.377 ± 0.816 | 0.783 ± 0.185 | 3.690 ± 0.359 | 4.710 |
| Pse5Ac7Fo | 7.575 ± 0.865 | 0.688 ± 0.202 | 3.337 ± 0.381 | 4.853 |
| Pse5Ac7(3 _R Hb) | 8.863 ± 0.936 | 1.077 ± 0.239 | 3.904 ± 0.412 | 3.624 |
| Pse7Fo5(3 _R Hb) | 2.543 ± 0.524 | 1.367 ± 0.536 | 1.120 ± 0.231 | 0.819 |

^a Reactions were performed with 0.8 mM CTP and 2.27 μM PseF in 20 mM MgCl_2 , 30 mM sodium phosphate (pH 7.4) at 37 °C for 15 seconds.

^b Estimated conversion was determined by UV absorption *via* UPLC analysis on a HILIC column.

(ranging from 0.2 to 1.8 mM) in the presence of MgCl_2 (20 mM) and sodium phosphate (30 mM, pH 7.4) for 15 seconds (Fig. 5). The corresponding Michaelis–Menten best-fit values are summarized in Table 1. It is clear that Pse7Fo5(3_RHb) **4** showed around one quarter of k_{cat} value and one fifth of k_{cat}/K_m value than other Pse derivatives, which demonstrated that the lower yield of CMP-Pse7Fo5(3_RHb) **10** originated from inferior conversion. This kinetic discrimination of *N*5-substituted Pse is also consistent with our observations in metabolic labelling studies, where the *N*2-azidoacetyl modified probe that was metabolically transformed into *N*5-modified Pse gave much lower labelling efficiency compared to the *N*4-modified probe.²⁴

Conclusions

In summary, we finished the *de novo* total synthesis of three new *N*-substituted Pse derivatives, including Pse5Ac7Fo, Pse5Ac7(3_RHb) and Pse7Fo5(3_RHb), from orthogonally protected Pse thioglycoside donor **11**. The synthetic strategies we demonstrated can be useful for the chemical synthesis of other bacterial non-2-ulosonic acids, since the variation of *N*-substitutions is widely observed in this family of molecules. Furthermore, we probed the enzymatic substrate specificity of PseF by measuring the enzymatic kinetics of the transformation of our synthetic Pse derivatives to corresponding CMP-Pse products. To our delight, the results proved that PseF could specifically recognize a series of *N*-substituted Pse derivatives, but not sialic acid. Thus, PseF could be regarded as a universal Pse cytidyltransferase that is compatible with variable *N*-substitution. More critically, this is a piece of evidence that variable *N*-substitutions of Pse species in bacterial glycans are possibly decorated before the biosynthetic incorporation of Pse units into polysaccharides. This also indicates that the CMP-Pse compounds with variable *N*-substitution may be recognized by the same pseudaminyltransferase, which may reduce the difficulty of *in vitro* experiments to search for putative pseudaminyltransferases in future.

By gaining a more comprehensive understanding of the Pse biosynthetic pathway, researchers can potentially develop novel inhibitors to target pathogenic Gram-negative bacteria.^{38,39} This could lead to innovative therapeutic approaches in combating drug-resistant infections and improving the overall efficacy of antibiotic treatments. Moreover, this study may contribute to the ongoing exploration of bacterial glycosylation and its implications in human health and diseases by guiding the design of molecular probes with better labelling efficiency.

Experimental section

All experimental details are provided in the ESI† of this article.

Data availability

The data that support the findings, including the details of the synthesis, protein expression, enzymatic reactions, and copies of NMR spectra, are available in the ESI† of this article.

Author contributions

X. Li, S. Chen and P. Li conceived the project. X. Guo, Y. C. Cheung and C. Li did the experiments. X. Guo, H. Liu, S. Chen and X. Li analyzed the data. H. Liu and X. Li wrote the manuscript with the assistance of other contributors.

Conflicts of interest

The authors declare no conflict of interest.

Acknowledgements

The work was supported by Health and Medical Research Fund (20190802), Collaborative Research Fund (C7003-20G), and General Research Fund (17312822).

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