An Alternative Mechanism for the Methylation of Phosphoethanolamine Catalyzed by *Plasmodium falciparum* Phosphoethanolamine Methyltransferase

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**Background:** In *Plasmodium*, phosphoethanolamine methyltransferase (PfPMT) is essential for normal growth and development.

**Results:** Computational, biochemical, and structural studies suggest catalytic and structural roles for Asp-128 in PfPMT.

**Conclusion:** Asp-128 is critical for specific methylation of phosphoethanolamine and not other phosphobases.

**Significance:** This work provides new insight on the evolution of multiple substrate activity in PMT from different organisms.

The phosphobase methylation pathway catalyzed by the phosphoethanolamine methyltransferase in *Plasmodium falciparum* (PfPMT), the malaria parasite, offers an attractive target for antiparasitic drug development. PfPMT methylates phosphoethanolamine (pEA) to phosphocholine for use in membrane biogenesis. Quantum mechanics and molecular mechanics (QM/MM) calculations tested the proposed reaction mechanism for methylation of pEA involving the previously identified Tyr-19–His-132 dyad, which indicated an energetically unfavorable mechanism. Instead, the QM/MM calculations suggested an alternative mechanism involving Asp-128. The reaction coordinate involves the stepwise transfer of a proton to Asp-128 via a bridging water molecule followed by a typical S$_2$-type methyl transfer from S-adenosylmethionine to pEA. Functional analysis of the D128A, D128E, D128Q, and D128N PfPMT mutants shows a loss of activity with pEA but not with the final substrate of the methylation pathway. X-ray crystal structures of the PfPMT-D128A mutant in complex with S-adenosylhomocysteine and either pEA or phosphocholine reveal how mutation of Asp-128 disrupts a hydrogen bond network in the active site. The combined QM/MM, biochemical, and structural studies identify a key role for Asp-128 in the initial step of the phosphobase methylation pathway in *Plasmodium* and provide molecular insight on the evolution of multiple activities in the active site of the PMT.

Methyltransferases are ubiquitous in biochemical and cellular processes (1). Using S-adenosylmethionine (AdoMet)$^4$ as the methyl donor, this diverse group of enzymes catalyzes the modification of macromolecules and metabolites using a range of chemical strategies, including proximity, desolvation effects, acid-base catalysis, and metal-dependent deprotonation (1–3). The action of these enzymes is crucial for metabolic function, controlling gene regulation, and is linked to assorted diseases (3). Recent studies also suggest an important role for methyltransferases in plants, nematodes, and the apicomplexan parasite *Plasmodium falciparum* in the synthesis of phospholipid precursors (4–6).

In plants, nematodes, and *Plasmodium*, the phosphobase methylation pathway is used for the synthesis of phosphocholine (pCho) for phosphatidylcholine production (Fig. 1, top). The methylations that convert phosphoethanolamine (pEA) to phosphonomonomethylethanolamine (pMME), phosphodimethylethanolamine (pDME), and pCho are performed by phosphoethanolamine methyltransferases (PMT). Three types of PMT with varied domain architectures are found in different organisms (6). Plant PMT are di-domain enzymes (~450 amino acids) with one active site (PMT1) catalyzing the methylation of pEA to pMME, and a second active site (PMT2) that catalyzes the conversion of pMME to pDME and pDME to pCho (7). In nematodes, two PMT retain the di-domain organization (~430–480 amino acids) of the plant enzyme, except one of the domains in each nematode PMT is vestigial (8–11). This leads to distinct PMT1 and PMT2 enzymes in *Caenorhabditis elegans* and other nematodes. Unlike plants and nematodes, *P. falciparum* contains a single domain PMT (PfPMT; 266 amino acids) that catalyzes all three methylation steps in the phosphobase pathway.

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* The abbreviations used are: AdoMet, to S-adenosylmethionine; AdoHcy, adenosylhomocysteine; PfPMT, *P. falciparum* phosphoethanolamine methyltransferase; pEA, phosphoethanolamine; pCho, phosphocholine; QM/MM, quantum mechanics and molecular mechanics; pMME, phosphonomonomethylethanolamine; pDME, phosphodimethylethanolamine; PMT, phosphoethanolamine methyltransferase; PDB, Protein Data Bank; ITC, isothermal titration calorimetry; MD, molecular dynamics; r.m.s.d., root mean square deviation.

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PfPMT plays an important role in the metabolism of the malaria parasite by supporting the synthesis of phospholipids during reproduction and growth. Disruption of the PfPMT gene leads to a reduced phosphatidylcholine synthesis from serine, growth and survival defects, and the loss of parasite transmission to mosquitoes (12–14). Because of its critical role in Plasmodium and the lack of PMT homologs in mammals, PfPMT, along with other enzymes in phospholipid metabolism, is a target for antiparasitic drug development (5, 6, 15–17).

X-ray crystal structures of PfPMT in complex with various substrates and inhibitors and extensive biochemical characterization provide insight on the organization of the AdoMet and phosphobase-binding sites in the enzyme (Fig. 1, bottom) (18, 19). In particular, electrostatic interactions provided by Arg-179 and Lys-247, together with hydrogen bond interactions contributed by hydroxyl groups of several tyrosine residues, are critical for phosphobase binding. In addition, Tyr-19 and His-132, which are positioned between AdoMet and phosphobase-binding sites, were suggested as catalytic residues (18). In the proposed reaction mechanism (Fig. 1, bottom), His-132 functions as a general base to abstract a proton from the hydroxyl group of Tyr-19. Next, the negatively charged tyrosine oxygen interacts with a hydrogen atom on the substrate amine, orienting the lone pair of electrons toward the methyl group of AdoMet to facilitate the bimolecular nucleophilic substitution ($S_n2$) methyl transfer to pEA.

In this study, we combine computational modeling with protein crystallography and site-directed mutagenesis to re-examine the methylation of pEA catalyzed by PfPMT. Initially, we tested the ability of the proposed catalytic Tyr-19–His-132 dyad in deprotonating pEA, which suggests an energetically unfavorable mechanism. Instead, the quantum mechanical/molecular mechanical (QM/MM) calculations suggest an alternative mechanism involving Asp-128 to deprotonate pEA via a bridging water molecule (Fig. 1, bottom). Biochemical analysis of site-directed mutants targeting Asp-128 shows a loss of activity with pEA as a substrate but retains activity with pDME. The x-ray crystal structures of the PfPMT-D128A mutant in complex with $S$-adenosylhomocysteine (AdoHcy) and either pEA or pCho reveal structural changes in the active site. Over-
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EXPERIMENTAL PROCEDURES

Computational Model Setup—The initial structure of PfPM'T-pEA-AdoMet ternary complex was modeled from the crystal structure of PfPM'T in complex with pEA and AdoHcy (18; PDB code 3UJ7). AdoHcy was amended to AdoMet by adding one methyl group on S by based on the geometric parameters of AdoMet bound to PfPM'T (PDB code 3UJ7 (18)). The system was prepared using Protein Wizard (20) within the Schrodinger suite to optimize hydrogen-bonded networks and assign proper protonation states for ionizable residues at pH 7. In particular, all histidines were singly protonated (neutral) with His-82 and His-218 having the proton at their Nε with all other histidines protonated at the Nδ. The system was then equilibrated with 1 ns of molecular dynamics (MD) in a water box using weak harmonic constraints (5 kcal mol⁻¹Å²) for all heavy atoms to maintain their active site configuration.

The protonation state of the amine group in pEA was estimated using PROPKA (21). Five different PfPM'T-pEA-AdoMet conformations, including the crystal structure and four MD snapshots along the equilibration, were used and resulted in an estimated pKₐ of 10.72 ± 1.51 (Table 1). Addition of AdoMet to the active site model decreases the pKₐ to 8.68 ± 0.52 because of electrostatic interaction with the positive charge of the AdoMet methyl group. A similar pKₐ shift (from ~11 to ~8) in the presence of the AdoMet cofactor was reported for lysine methyltransferase (22). Thus, we assigned the pEA substrate in its protonated form as NH₃⁺-pEA.

QM/MM Reaction Profiles—All QM/MM calculations were carried out using the QSite program of the Schrodinger Suite (QSite, version 5.7; Schrödinger, LLC: New York). The QM part was treated at the M06/6-31G(d,p) level of theory (23), and the MM part was described with the OPLS force field (24). The hydrogen link atom approach was used for the QM/MM boundary treatment. During the Q/MM geometry optimization, the QM region and all MM atoms within a distance of 10 Å from any atom in NH₃⁺-pEA and AdoMet were allowed to relax. To calculate the QM/MM reaction profiles, we performed a linear scanning approach by imposing distance constraints. For each reaction profile, we carried out three iterative scans driving the reaction coordinate forward, backward, and repeatedly forward between reactant and product states. Analytical harmonic second derivatives were calculated to confirm the nature of the stationary point, with no imaginary eigenvalues for minima and only one imaginary eigenvalue for transition states.

Site-directed Mutagenesis, Protein Expression, and Protein Purification—Generation of PfPM'T D128A, D128E, D128Q, and D128N mutants used the QuikChange PCR mutagenesis method with the pET28a-PfPM'T vector (18) as a template. Wild-type and mutant PfPM'T were expressed in Escherichia coli BL21(DE3) and purified by nickel-affinity and size-exclusion chromatographies, as described previously (18).

Enzyme Assays and Isothermal Titration Calorimetry—Standard assay conditions for PM'T activity were 0.1 M Hepes-KOH (pH 8), 2 mM Na₂EDTA, 10% glycerol, 2.5 mM AdoMet (100 nCi of [methyl-¹³C]AdoMet), and 5 mM phosphoethanolamine substrate (either pEA or pDM) in 100 µl. Protein (10 µg) was incubated for 10 min at room temperature with reactions terminated and product quantified as described previously (8, 9). Steady-state kinetic analysis of wild-type and mutant PfPM'T was performed as described previously (8, 9, 18). Isothermal titration calorimetry (ITC) was used to monitor AdoHcy and pCho binding to wild-type and mutant PfPM'T, as described previously (10, 18).

Protein Crystallography—Crystals of the PfPM'T-D128A-AdoHcy-pEA and PfPM'T-D128A-AdoHcy-pCho complexes were grown by the vapor diffusion method in hanging drops of a 1:1 mixture of protein (13.5 mg/ml) and crystallization buffer.
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**TABLE 2**

| Crystallographic statistics | PfPMT-D128A-AdoHcy-pCho | PfPMT-D128A-AdoHcy-pEA |
|----------------------------|--------------------------|------------------------|
| Crystal space group        | P2                       | P2                     |
| Cell dimensions            | a = 77.17 Å, b = 44.06 Å, c = 89.15 Å; β = 108.1° | a = 76.83 Å, b = 43.65 Å, c = 88.05 Å; β = 107.8° |
| Data collection            |                          |                        |
| Wavelength (Å)             | 0.979                    | 0.979                  |
| Resolution range (Å) (highest shell resolution) | 30.4–1.59 (1.62–1.59) | 38.7–2.60 (2.64–2.60) |
| Reflections (total/unique) | 270,106/76,333           | 57,984/17,579          |
| Completeness (highest shell) | 99.6% (99.6%)           | 99.4% (99.3%)          |
| R(merge) (highest shell)   | 22.5 (2.2)               | 9.1 (1.9)              |
| Rsym (highest shell)       | 7.9% (52.1%)             | 16.5% (62.7%)          |
| Model and refinement       |                          |                        |
| Rcryst/Rfree               | 15.9%/18.8%              | 20.3%/25.8%            |
| No. of protein atoms       | 4,310                    | 4,242                  |
| No. of water molecules     | 715                      | 68                     |
| No. of ligand atoms        |                          |                        |
| r.m.s.d., bond lengths (Å) | 0.007                    | 0.002                  |
| r.m.s.d., bond angles (°)  | 1.026                    | 0.585                  |
| Average B-factor (Å²)      | 22.2, 36.8, 18.4         | 27.4, 28.7, 25.7       |
| Stereochemistry: most favored, allowed, outliers | 99.0, 1.0, 0% | 97.9, 2.1, 0% |

| Structure factor          |            |            |
|---------------------------|------------|------------|
| r.m.s.d., bond angles (°)  | 1.026      | 0.585      |
| Stereochemistry: most favored, allowed, outliers | 99.0, 1.0, 0% | 97.9, 2.1, 0% |

**RESULTS**

Modeling of Phosphoethanolamine Methylation Catalyzed by PfPMT—Starting from the crystal structure of PfPMT-pEA-AdoMet complex after a 1-ns equilibration, we examined the previously proposed mechanism in which Tyr-19 and His-132 deprotonate pEA (18). The QM region included pEA, Tyr-19, His-132, and part of AdoMet (excluding the adenine base) for a total of 76 atoms. We performed QM/MM scanning where the reaction coordinate was defined as the distance between the Nε of His-132 and the hydroxyl hydrogen of Tyr-19. In doing so, a proton was pulled from Tyr-19 to His-132 starting from a distance of 1.8 Å (at the reactant) to 1.0 Å (at the product). The proton transfer reaction profile displayed a 21 kcal/mol endothermic profile, with energy increasing monotonically and no energy minimum corresponding to products/intermediates (Fig. 2A). Upon the approach of a proton from Tyr-19 to His-132, simultaneous proton transfer from NH3+-pEA to a negatively charged (deprotonated) Tyr-19 did not take place. Moreover, the forced product of this proton transfer path (Tyr-19-His-1322-) reverts to the tyrosine when removing the constraint on the reaction coordinate.

(20% PEG-8000, 0.1 M sodium cacodylate (pH 6.5), 0.2 M sodium acetate, 20 mM tris(2-carboxyethyl)phosphine, 5 mM AdoHcy, and 5 mM pEA/pCho). Diffraction data were collected at beamline 19-ID of the Argonne National Lab Advanced Photon Source. HKL3000 (32) was used for indexing, integration, and scaling of diffraction data. The crystal structure of each complex was determined by a molecular replacement using PHASER (33) and the PfPMT complex was determined by a molecular replacement using PHASER (33) and the PfPMT-D128A complex after a1-ns equilibration, we examined the previously proposed mechanism in which Tyr-19 and His-132.

Based on this lack of deprotonation from the Tyr-His dyad, we manually took a proton from pEA and performed a methyl transfer linear scanning, aiming to observe an enhanced structure for proton abstraction. Thus, we perturbed the distance...
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Potential Role of Asp-128 in the Methylation of Phosphoethanolamine—From the QM/MM results described above, an alternative mechanism where Asp-128 acts as a general base to deprotonate pEA through a water molecule, Wat (Fig. 2B). This conformational rearrangement allows for hydrogen bond interactions between pEA and Wat1 (2.5 Å N–O distance) and between Wat1 and Asp-128 (2.6 Å O–O distance). These distances were obtained after a QM/MM minimization, and inclusion of missing vibrational (temperature and zero point) energies would increase these distances to ~2.7 Å. In this orientation, although Asp-128 is located almost 6 Å away from the amine group of pEA, a hydrogen bond network via a water bridge may contribute to catalysis. The same conformational rearrangement results were obtained when repeating the QM/MM calculations by extending the QM region to include Wat and Asp-128, for a total of 96 atoms.

Potential Role of Asp-128 in the Methylation of Phosphoethanolamine—From the QM/MM results described above, an alternative mechanism where Asp-128 acts as a general base to deprotonate pEA through a water molecule is possible. In this orientation, although Asp-128 is located almost 6 Å away from the amine group of pEA, a hydrogen bond network via a water bridge may contribute to catalysis. The same conformational rearrangement results were obtained when repeating the QM/MM calculations by extending the QM region to include Wat and Asp-128, for a total of 96 atoms.

Methyl transfer step undergoes a typical $S_n^2$ process with an energy barrier of 10.0 kcal/mol to produce a highly stable product ($\text{NH}_2\text{CH}_3\text{H}^+_2\text{pEA}$-$\text{Ash}$-$128$-$\text{AdoHcy}$). The overall reaction is exothermic by 35.3 kcal/mol. The structures of the reaction sequence (Fig. 3, B–E) suggest that the intermediate presents $S_n^2$-ready spatial ordering, where the amine lone pair electrons point directly to the methyl carbon of AdoMet. In this orientation, the distance between them is about 3.0 Å, and the angle is ~171°. The methyl transfer transition state has an imaginary frequency of ~424 cm$^{-1}$, exhibits a linear AdoMet:$\text{S}_n^2$-$\text{pEA}$:$\text{Ash}$:$128$-$\text{AdoHcy}$ configuration with an angle of 173°, and is almost equidistant between bond breaking (2.18 Å) and forming (2.20 Å). All attempts to produce a concerted proton transfer/methyl transfer pathway dramatically increased the energy barriers to ~70 kcal/mol.

Water Network Stability and the Methylation of Phosphoethanolamine—To probe the presence of the water network connecting pEA and Asp-128, three 10-ns MD runs were performed. In each simulation, the stability of the water network was observed (Fig. 4), although several water exchanges with bulk solvent were observed. Moreover, in 70% of the simulations, the network consists of two water molecules (inset of Fig. 4).
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![Graphical representation of mechanism](image)

### TABLE 3

|                  | Crystal | M1   | M2   | M3   | M4   | Average |
|------------------|---------|------|------|------|------|---------|
| No. of waters in bridge | 1.0     | 2.0  | 2.0  | 2.0  | 2.0  | 2.0     |
| ∆E (kcal/mol)    | 10.3    | -3.4 | -2.6 | -1.8 | 1.70 |         |

### Reactant parameters

- AdoMet:S*−AdoMet:Ce (Å) 1.91
- AdoMet:Ce=pEA:N (Å) 3.57
- S*−Ce:N (°) 165
- pEA:N=Wat1:Ow (Å) 2.64
- Wat1:Ow=Wat2:Ow (Å) 2.63
- Wat2:Ow=Asp-128:Oδ (Å) 2.60

### Summary of activation energy (EA#) and reaction energy (∆E) for proton transfer step from QM/MM calculations using different starting configurations

|                  | Crystal | M1 | M2 | M3 | M4 | Average |
|------------------|---------|----|----|----|----|---------|
| EA# (kcal/mol)   | 10.0    | 7.1 | 9.4 | 14.8| 7.3 | 9.7     |
| ∆E (kcal/mol)    | -25.0   | -31.8| -34.4| -27.0| -31.7| -30.0   |

### Reactant

- d1 (Å) 3.00
- d2 (Å) 1.83
- θ (°) 171

### Transition state

- d1 (Å) 2.20
- d2 (Å) 1.81
- θ (°) 173

### Product

- d1 (Å) 1.49
- d2 (Å) 3.28

4). Despite the presence of two waters, the QM/MM results indicate that the proton transfer to Asp-128 is associated with low energy barriers. Four additional QM/MM energy profiles were obtained from different MD snapshots (Fig. 5), with three simulations (M1, M3, and M4) presenting a two water network between substrate and Asp-128. The average barrier for proton transfer and methyl transfer steps were 4.1 ± 1.9 and 9.7 ± 3.2 kcal/mol, respectively, and are in good agreement with the simulation based on the x-ray crystal structure. Importantly, even with a two water bridge, all of the proton transfer reactions from NH₃⁺=pEA to Asp-128 undergo a concerted asynchronous mechanism with low energy barriers (4–6 kcal/mol; Table 3). The subsequent methyl transfers proceed through a mechanism consistent with an S₂,2 reaction (Table 4). All of the transition state structures exhibit similar AdoMet:S*−AdoMet:Ce=pEA:N linear geometry (~170 ± 4°) with a reactive methyl species located in the middle between bond breaking and forming (average distance of 2.22 ± 0.02 and 2.23 ± 0.05 Å, respectively). The observed fluctuation of the methyl transfer energy barriers (Table 4) is related to the closeness of the reactant conformations to the in-line transition state features. For instance, the higher barrier of the M3 model (14.8 kcal/mol) is due to the longer bond length of AdoMet:Ce=pEA:N and larger deviation of the AdoMet:S*−AdoMet:Ce=pEA:N angle from the linear attacking angle. We should underline that, for each frame, we also attempted QM/MM calculations driving deprotonation of NH₃⁺=pEA through the Tyr-19–His-132 dyad. These simulations, however, resulted again in highly endothermic and monotonically increasing energy profiles analogous to the one shown in Fig. 2A.

**Biochemical Analysis of PfPMT Asp-128 Site-directed Mutants—**

To probe the potential role of Asp-128 in the reactions catalyzed by PfPMT, four site-directed mutants (D128A, D128E, D128Q, and D128N) were generated for biochemical analysis. Each mutant was expressed in *E. coli* and was purified by nickel-affinity and size-exclusion chromatographies. All the mutants were isolated as soluble monomeric proteins with expression yields comparable with wild-type PfPMT. The methylation activity of wild-type PfPMT and the four Asp-128 mutants were compared using AdoMet and either pEA (first methylation reaction) or pDME (third methylation reaction) as substrates (Fig. 6). Using pEA and AdoMet as substrates, the D128A, D128E, D128Q, and D128N mutants displayed 1.5, 9.3, 0.6, and 0.2% of wild-type methylation activity, respectively. In contrast, using pDME and AdoMet as substrates, three of the Asp-128 mutants were either comparable (D128A and D128Q) to or had slightly higher (1.8-fold; D128E) specific activities than PfPMT. The D128N mutant showed a 4-fold reduction in specific activity with pDME and AdoMet as substrates.

Determination of steady-state kinetic parameters for the D128A and D128N mutants using either pEA or pDME showed a similar difference between these substrates (Table 5). Both mutations increased the *K_m* values and decreased the turnover rates with pEA compared with wild-type enzyme. The combination of effects reduced the catalytic efficiencies (*k_{cat}/K_m*) of the D128A and D128N mutants with pEA by 12,400- and 135,000-fold, respectively. In contrast, the *k_{cat}/K_m* values for pDME with the D128A and D128N mutants were reduced by 18- and 36-fold, respectively. These results demonstrate that Asp-128 plays a critical role in the methylation of pEA and has a greatly diminished but still significant role in the methylation of pDME.
Ligand Binding and X-ray Crystal Structure of the D128A PfPMT Mutant—To further examine the effect of substitutions of Asp-128, the PfPMT D128A mutant was chosen for additional study by ITC and x-ray crystallography. Given the loss of specific activity observed with this mutant, ITC was used to quantify ligand binding to the enzyme (Fig. 7; Table 6). This analysis shows that mutation of Asp-128 to an alanine does not significantly change the binding affinity for AdoHcy, although the enthalpic and entropic contributions to AdoHcy binding differ compared with wild-type PfPMT (Table 6). For binding of pCho to the D128A mutant, there was insufficient heat signal to accurately determine a $K_d$ value by ITC.

The structural effects of the Asp-128 to alanine mutation were determined by solving the three-dimensional structure of the PfPMT D128A mutant by molecular replacement. The x-ray crystal structures of the mutant protein in complex with AdoHcy and either pCho or pEA were solved to 1.59 and 2.60 Å resolution, respectively (Table 2). The overall structures of the PfPMT-D128A/AdoHcy and PfPMT-D128A/pCho complexes were superimposable on the wild-type structure with 0.17 and 0.18 Å r.m.s.d., respectively, for 257 Cα atoms (Fig. 8A). As described previously (18, 19), the AdoHcy-binding site of PfPMT is situated along the N-terminal
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FIGURE 8. Structural comparison of the wild-type and D128A PfPMT active sites. A, overall structure of the PfPMT-D128A-AdoHcy-pCho complex. AdoHcy and pCho are shown as space-filling models. The position of the α2-α3 loop, which includes Ser-37–Gly-40, is indicated. B, active site view of the PfPMT-AdoHcy-pEA complex (18). AdoHcy is not shown for clarity. Key water molecules are shown as red spheres. Hydrogen bonds are indicated by dashed lines. C, active site view of the PfPMT-D128A-AdoHcy-pCho complex. AdoHcy is not shown for clarity. Key water molecules are shown as red spheres. Hydrogen bonds are indicated by dashed lines. D, active site view of the PfPMT-D128A-AdoHcy-pEA complex. AdoHcy is not shown for clarity. Key water molecules are shown as red spheres. Hydrogen bonds are indicated by dashed lines.

Three different 10-ns trajectories for wild-type and D128A PfPMT were generated. The average binding free energy (ΔG) of NH$_3^+$-pEA estimated by the MM/GBSA approach (31) suggests weaker affinity to the mutant, with values of −16.8 and −7.3 kcal/mol for wild-type and D128A mutant, respectively. In addition to altering the active site architecture, removal of the carboxylate side chain in the D128A mutant allows for greater mobility of active site residues and diffusion of water molecules from the bulk, as indicated by MD simulations (Fig. 8, A and B). Analysis of water molecules within 5 Å of the AdoMet.Ce atom indicates the presence of 1–2 water molecules in the PfPMT, but 1–4 water molecules for the D128A mutant (Fig. 9C). Overall, the increased active site mobility translates into a longer distances between AdoMet and the NH$_3$ group of pEA (Fig. 9D), in agreement with the crystallographic observations, as described above. For both wild-type and D128A, the phosphate group of pEA is comparatively restrained due to multiple interactions with Tyr-27, Tyr-160, Tyr-175, Arg-179, Tyr-181, and Lys-247.

DISCUSSION

Although AdoMet-dependent methylations of macromolecules and metabolites are ubiquitous in biological systems, the enzymes that perform these transformations employ a variety of chemical strategies (1). Most methyltransferases follow a common S$_2$2 methyl transfer mechanism following proton transfer from the substrate to a catalytic base in either stepwise or concerted mechanisms (2, 36–41). The identification of a family of PMT from plants, nematodes, and the malaria parasite P. falciparum provides an opportunity to explore the structural and chemical evolution of diverse methyltransferase active site architectures that catalyze the steps of the phosphobase methylation pathways in these organisms (Fig. 1, top) (4–12).

Using PfPMT, we examined the methylation of pEA using a combined computation, biochemical, and structural approach. In pEA methylation catalyzed by PfPMT, the substrate must be deprotonated for catalysis to occur. Based on a series of x-ray crystal structures, ligand binding studies, and the analysis of site-directed mutants of PfPMT, a reaction mechanism involving the Tyr-19–His-132 dyad was proposed (Fig. 1, bottom) (18). To examine the energetics of this mechanism, QM/MM calculations were used to obtain the energy profile for deprotonation and methylation of pEA. Overall, the simulations reveal features in common with other methyltransferase reactions.
and provide new molecular insight on the first step of the phosphobase methylation pathway.

The simulations clearly indicate an unfavorable energetic reaction coordinate for the proposed catalytic role of His-132 as a general base to activate Tyr-19 by deprotonating a charged (i.e., NH$_3^+$/pEA) substrate (Fig. 2A). Interestingly, during the QM/MM simulation, rotation of the amine group of pEA toward a water molecule bound to Asp-128 was observed and suggested an alternative reaction mechanism (Fig. 2B). This rotation aids in formation of a hydrogen bond network and correlates with the relatively high value of B-factor of this moiety in x-ray crystal structure (18), which supports the mobility of the substrate amine observed in the simulation. The QM/MM reaction coordinate profiles suggest that the methylation of pEA catalyzed by PfPMT proceeds through the stepwise mechanism starting with the proton transfer step from NH$_3^+$/pEA to Asp-128 through a water bridge. This is followed by the methyl transfer from AdoMet to NH$_2$/pEA to form the product. The overall stepwise mechanism is an exothermic process with methyl transfer as the rate-limiting step; the presence of high barriers excludes a concerted methyl transfer/proton transfer pathway.

For this alternative mechanism in PfPMT, the core features of the methyltransferase reaction are consistent with other computational studies. The QM/MM calculations on multiple starting configurations of the PfPMT active site address the potential dynamical nature of the enzyme and indicate that they do not alter the mechanism (42). Our results indeed support the hypothesis from Hu and Zhang (39) that an indicator for the reactivity for such a methyl transfer reaction is a combination of the nucleophilic attack bond length ($d$) and angle ($\theta$) (Table 4). In addition, the calculated average energy barrier of 9.7 ± 3.2 kcal/mol obtained from the M06/6−31G(d,p)/OPLS calculation for the methylation of pEA is in agreement with the activation barrier of 9.6 kcal/mol estimated from the experimental value of $k_{cat} = 109$ min$^{-1}$ (18) and the simple transition state theory: $k(T) = (k_p/k_h)\exp(-\Delta G/RT)$. To further validate this mechanism, a combination of site-directed mutagenesis, biochemical assays, and protein crystallography were performed.

Biochemical analysis of the PfPMT D128A, D128N, D128Q, and D128E site-directed mutants confirms an important functional role for Asp-128 in the methylation of pEA; however, the specific activities of the mutants remain comparable with wild-type enzyme for the conversion of pDME to pCho (Fig. 6). Steady-state kinetic analysis of the D128A and D128N mutants indicates that either removal of the carboxylic acid group or substitution with an amide group reduces $k_{cat}$ with pEA by 100–300-fold (Table 5). In addition, these changes significantly increase the $K_m$ values for pEA (Table 5). Consistent with the specific activity measurements, the D128A and D128N mutants displayed more modest changes in kinetic parameters for pDME. This suggests a specific role for Asp-128 during the first step of the phosphobase methylation pathway catalyzed by PfPMT.

Ligand binding analysis of the PfPMT D128A mutant using ITC indicates that this mutation does not alter AdoHcy binding (Fig. 7A; Table 6), as expected from this residue’s position in the three-dimensional structure. In comparison, the D128A substitution alters pEA and pCho binding, even though the aspartate makes no direct contacts with phosphobase substrates. Although the bind-
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ing constant for pCho could not be accurately determined because of a lack of heat signature (Fig. 7B; Table 6), use of saturating concentrations of pEA and pCho during protein crystallization allowed for determination of the x-ray crystal structure of the PfPMT D128A mutant in the presence of each ligand (Fig. 8; Table 6). The initial inspection of the crystals confirms that mutating Asp-128 does not change the overall positioning of AdoHcy, pEA, and pCho in the active site. The network of interactions centered on Asp-128 in the wild-type enzyme, however, is disrupted in the mutant enzyme, inducing a larger mobility in pEA’s amine group. Those changes, reducing the probability of reactive conformations with reasonable distances between the methyl donor and acceptor (43), are nicely reproduced by MD simulations of the D128A mutant (Fig. 9).

Although the residue corresponding to Asp-128 of PfPMT is invariant across other PMT from plants and nematodes (6), the structural context of this aspartate differs in these enzymes. The x-ray crystal structures of PMT1 and PMT2 from the nematodes *C. elegans* and *Haemonchus contortus* reveal two distinct active site architectures (11) as follows: one for the methylation of pEA (PMT1) and another for the methylation of pMME and pDME (PMT2). Intriguingly, PMT2 from *C. elegans* and *H. contortus* do not use pEA as a substrate, even though these enzymes share an active site architecture nearly identical to PfPMT (11, 18). To understand how the PfPMT catalyzes the methylation of pEA, sequence and structural comparisons identified a difference between Ser-38 of PfPMT and a proline at the same position in the nematode PMT2 enzymes (11). These comparisons show that the interface of the AdoMet/AdoHcy and phosphobase-binding site of the nematode PMT2 is more constrained compared with PfPMT. This constriction in the nematode enzymes prevents water molecules from binding in the active site near the phosphobase amine. Because the extra space in PfPMT allows water molecules to enter the active site, it was previously suggested that the water network helps to orient the amine group of pEA for catalysis in PfPMT (11). A point mutation in PfPMT (i.e. S38P) eliminated activity of the enzyme with pEA but retained wild-type activity with pDME (11). That result mimics the effects of the D128A, D128Q, D128N, and D128E mutants in PfPMT (Fig. 6). Moreover, mutation of Asp-128 to an alanine in PfPMT shifts Ser-37–Gly-40 of the a2–a3 loop to a position similar to that of the same loop in the active site of the PMT2 from *C. elegans* and *H. contortus* (11). Additional MD simulations for an S38P mutant clearly indicate the disruption of the hydrogen bond network (and the dehydration) between pEA and Asp-128 (Fig. 10). These results highlight the subtle changes in the PfPMT active site required for use of pEA as a substrate.

In conclusion, we suggest a new catalytic role for Asp-128 in the active site of PfPMT. The first contribution of Asp-128 is that of a general base to activate a water molecule that aids in deprotonation of pEA, as suggested by the QM/MM calculations described here. In contrast to the active sites of the nematode PMT2 (11), the PfPMT active site readily accommodates water molecules that form a hydrogen bond network involving Asp-128. Importantly, this structural difference is linked to the interaction between Asp-128 and Ser-38, which shifts the a2–a3 loop to allow for the water network in PfPMT. These structural changes are important for the evolution of an alternate reaction path for pEA methylation by PfPMT.

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