

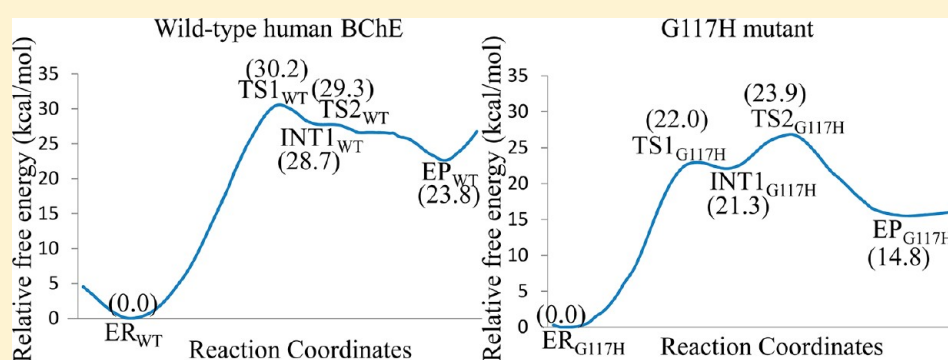
# Why Does the G117H Mutation Considerably Improve the Activity of Human Butyrylcholinesterase against Sarin? Insights from Quantum Mechanical/Molecular Mechanical Free Energy Calculations

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**ABSTRACT:** Human butyrylcholinesterase (BChE) is recognized as the most promising bioscavenger for organophosphorus (OP) warfare nerve agents. The G117H mutant of human BChE has been identified as a potential catalytic bioscavenger with a remarkably improved activity against OP nerve agents such as sarin, but it still does not satisfy the clinical use. For further design of the higher-activity mutants against OP nerve agents, it is essential to understand how the G117H mutation improves the activity. The reaction mechanisms and the free energy profiles for spontaneous reactivation of wild-type BChE and its G117H mutant phosphorylated by sarin have been explored, in this study, by performing first-principles quantum mechanical/molecular mechanical free energy calculations, and the remarkable role of the G117H mutation on the activity has been elucidated. For both the wild-type and G117H mutant enzymes, H438 acts as a general base to initiate the spontaneous reactivation that consists of two reaction steps: the nucleophilic attack at the phosphorus by a water molecule and decomposition of the pentacoordinated phosphorus intermediate. The calculated overall free energy barriers, i.e., 30.2 and 23.9 kcal/mol for the wild type and G117H mutant, respectively, are in good agreement with available experimental kinetic data. On the basis of the calculated results, the mutated residue (H117 in the G117H mutant) cannot initiate the spontaneous reactivation as a general base. Instead, it skews the oxyanion hole and makes the phosphorus more open to the nucleophilic water molecule, resulting in a remarkable change in the rate-determining step and significantly improved catalytic activity of human BChE.

Organophosphorus (OP) chemical warfare nerve agents, e.g., sarin, soman, tabun, VX, and VR, are highly neurotoxic compounds. They inhibit cholinesterases, including both acetylcholinesterase (AChE, EC 3.1.1.7) and butyrylcholinesterase (BChE, EC 3.1.1.8), by phosphorylation of the catalytic serine residue. Phosphorylation of AChE at neuronal and neuromuscular synapses is the main cause of symptoms of OP nerve agent poisoning.<sup>1–3</sup> The production, stockpiling, and use of OP chemical warfare nerve agents are now prohibited under the terms of the Chemical Weapons Convention (CWC) of the United Nations, and thus, the threat of military use has been diminished. However, there is still perceived to be a serious threat of use by terrorists. In the 1980s, sarin was used against Kurdish communities in Iraq.<sup>4</sup> In the 1990s, sarin was

released by Japanese terrorists in Matsumoto, Nagano, and the Tokyo subway.<sup>5–8</sup> Moreover, VX was also used in an assassination in Japan by the same group in 1994.<sup>9</sup>

Cholinesterases phosphorylated by OP chemical warfare nerve agents can undergo three possible postinhibitory reactions: (1) spontaneous reactivation by a water molecule to restore the enzyme activity, (2) nucleophile-induced reactivation to restore the enzyme activity, and (3) aging reaction by dealkylation of the phosphorylated cholinesterase, resulting in the truly irreversible inhibition. On the basis of

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these reactions, many strategies, e.g., protective inhibitor, reactivator, and scavenger approaches, have been developed for the pretreatment and post-treatment against the effects of the chemical warfare nerve agents in the past 50 years.<sup>10</sup> The protective inhibitor, such as carbamate, may be able to block AChE against the reactions with the chemical warfare nerve agents. However, after an exposure, there will always be a time during which acetylcholine builds up pending reactivation of the enzyme, causing a period of incapacitation. The reactivator, such as oxime, must reach the phosphorylated enzyme and restore the enzyme activity in time to protect the life of the organism if it is to be effective. In the mean time, the developed stronger anticholinergics and superior anticonvulsants must be delivered in a timely fashion; otherwise, the reactivator strategy is useless. Therefore, the best way to completely and efficiently protect humans from the OP nerve agents is to design and discover an effective scavenger to degrade the OP nerve agents in plasma. A desirable scavenger for this purpose is capable of intercepting and destroying an OP nerve agent before it reaches a critical target without causing any incapacitation or behavioral deficits. There are several approaches to achieving this goal. One approach is to isolate and characterize an organophosphorus acid anhydride hydrolase (OPAAH), for example, carboxylesterase, paraoxonase, and a bacterial OPAAH from *Pseudomonas diminuta*. The bacterial OPAAH is also known as phosphotriesterase (PTE) in the literature.<sup>11–15</sup> Unfortunately, none of them can defend the effects of the chemical warfare nerve agents without measurable side effects. A more efficient approach is to construct a highly active OPAAH against the nerve agents by using protein engineering techniques. The progress was achieved by using human BChE as a scavenger for all of the OP chemical warfare nerve agents.<sup>16,17</sup> Human BChE was chosen as the scavenger for several reasons. (1) The OP-inhibited human BChE can restore the enzyme activity and generate a nontoxic product via a spontaneous reactivation reaction. (2) Human BChE is present in human serum and could be used as a therapeutic agent injected into blood. (3) Human BChE has a history of use in the clinic, and no adverse effects have been reported.<sup>18</sup> (4) So far, ~65 different naturally occurring mutants of human BChE have been identified, and none of them is antigenic.

The rationale for the initial attempt involving site-directed mutagenesis of human BChE was based on a hypothesis proposed by Jarv<sup>19</sup> that organophosphates are hemisubstrates for the cholinesterases. In the hemisubstrate hypothesis, the first stage, i.e., phosphorylation, is similar to the acylation stage of acetylcholine hydrolysis, but the second stage is significantly different from the deacylation stage of acetylcholine hydrolysis. In the deacylation stage, a histidine-bound water molecule displaces the acyl group from the catalytic serine, whereas this histidine-bound water molecule is prevented sterically from attacking the appropriate face of the tetrahedral phosphorus atom in the corresponding stage of the OP hydrolysis and thus prevents the reactivation of the phosphorylated enzyme by water. Therefore, introducing a new histidine residue into the active site of the enzyme might generate another nucleophilic center that activates a water molecule to the opposite side of the phosphorus–serine bond and thereby initiates the reactivation reaction. Given this scheme, G115, G117, E119, G121, L286, and V288 in human BChE were mutated to histidine, and only the G117H mutant was reported to have the ability to catalyze the hydrolysis of OP compounds.<sup>20,21</sup> The experimental data indicated that the catalytic rate constant of

the G117H mutant of human BChE against sarin has been increased by at least ~174-fold versus that of wild-type human BChE.<sup>21,22</sup> However, the significantly improved catalytic activity of the G117H mutant against sarin, where the catalytic rate constant is  $8.7 \times 10^{-5} \text{ s}^{-1}$ ,<sup>21,22</sup> is still too low for clinical use. The search for new mutants of human BChE with a further improved catalytic activity continues.

The rationale for the G117H mutation was to introduce a new nucleophilic center where H117 may act as a general base in the hydrolysis of OP. An empirical valence bond (EVB) simulation on a cluster model of the G117H mutant predicted the conformation of the introduced H117 residue, and on the basis of the EVB results, the authors suggested that H117 may act as a general base to hydrolyze OP.<sup>23</sup> If this suggestion were correct, then the spontaneous reactivation reaction would show a bell-shaped pH dependence curve with the rates,<sup>21</sup> similar to those in the deacylation and dephosphorylation reactions catalyzed by wild-type cholinesterases.<sup>19</sup> However, neither the recent X-ray crystal structure of the G117H mutant of human BChE phosphorylated by VX<sup>24</sup> nor the relationship between the reactivation rate and pH value<sup>21</sup> supports this suggestion. Understanding why and how the G117H mutation increases the activity against OP is therefore becoming very important. There have been several efforts to understand the mechanism for the reaction of the G117H mutant. On the basis of molecular mechanics calculations, a hypothesis proposed by Fortier et al.<sup>25</sup> involves a phosphoryl–histidine intermediate that is subsequently hydrolyzed by a water or hydroxide ion. However, the formation of the phosphoryl–histidine intermediate is prevented by the steric effect of OP, and hence, this process should be very slow, as mentioned by Broomfield et al.<sup>10,22</sup> This suggests that the spontaneous reactivation reaction of the G117H mutant of human BChE is not really initiated by the introduced H117 residue. Hence, it is possible that the mechanism for the spontaneous reactivation of the G117H mutant might be similar to that proposed for the spontaneous reactivation of wild-type human BChE,<sup>26,27</sup> implying that the H438 residue still acts as a general base to activate a water molecule that initiates the nucleophilic attack on the phosphorus atom. If this is true, the introduced H117 residue might play an indirect role in accelerating the reaction process. One hypothesis attributes the improved rate constant to the presence of a positive charge at the H117 residue (as it was assumed that H117 is protonated).<sup>22</sup> However, the G117K mutation, which also increases the positive charge, makes the dephosphorylation much slower than that of the G117H mutant, meaning that the hypothesis is not correct.<sup>22</sup> Another hypothesis<sup>20</sup> is that the G117H mutation causes a couple of structural changes. The first one is to distort the oxyanion hole, resulting in the P=O bond sitting askew and H438 no longer blocking attack by water. The other is to destabilize the covalent bond between sarin and the enzyme.<sup>20</sup> The nature of the G117H mutation improving the catalytic activity for the OP hydrolysis, which is very important in the design of high-activity mutants, remains uncovered. Understanding the role of the G117H mutation is therefore crucial.

In this study, we have performed first-principles quantum mechanical/molecular mechanical free energy perturbation (QM/MM-FE) calculations<sup>28–31</sup> to uncover the role of the G117H mutation by exploring the detailed reaction mechanisms for spontaneous reactivation of both wild-type human BChE and the G117H mutant that are phosphorylated by chemical warfare nerve agent sarin. Sarin was chosen because it

is a typical representative of the G-type nerve agents and its experimental kinetic data are well-established. In the QM/MM-FE method, the first-principles QM/MM reaction coordinate calculations were followed by free energy perturbation (FEP) calculations to more reasonably account for the dynamic effects of the protein environment on the free energy barriers for the enzymatic reactions. Our QM/MM simulations are based on the pseudobond first-principles QM/MM approach,<sup>29–32</sup> which has been demonstrated to be a powerful tool in simulating a variety of enzymes,<sup>28,33–50</sup> and some theoretical predictions<sup>39,49,51</sup> were subsequently confirmed by the experimental studies.<sup>35,49,52,53</sup> The computational results clearly reveal the reaction mechanisms and free energy profiles of the reactivation reactions of sarin-phosphorylated wild-type human BChE and the G117H mutant. The rate-determining steps are identified, and the roles of the essential residues, including those forming the catalytic triad and oxyanion hole, are discussed on the basis of the determined free energy profiles and the QM/MM-optimized geometries. By comparing the detailed reactivation reaction mechanisms of the sarin-phosphorylated wild-type enzyme and G117H mutant of human BChE, we are able to understand the role and the remarkable effects of the G117H mutation.

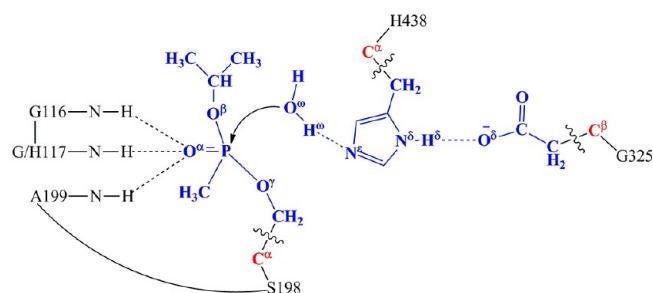
## COMPUTATIONAL METHODS

**Structure Preparation.** The initial model of BChE was constructed on the basis of the X-ray crystal structure of human BChE [Protein Data Bank (PDB) entry 1P0M].<sup>54</sup> All small molecules and the carbohydrate in the crystal structure were removed except the crystal water molecules. The missing residues (D2, D3, D378, D379, Q455, and Q486) in the X-ray crystal structure were rebuilt by using the automated homology modeling tool Modeler/InsightII (Accelrys, Inc.).<sup>55,56</sup> It should be noted that residues 17, 455, 481, and 486 are all asparagines (glycosylation sites) in wild-type BChE. One had to mutate these asparagines to glutamines through site-directed mutagenesis to obtain a single crystal for structural analysis by X-ray crystallography. Therefore, the artificially engineered residues Q17, Q455, Q481, and Q486 were modified back to N17, N455, N481, and N486, respectively, to obtain the wild-type BChE structure. In addition, G117 was mutated to histidine, arginine, or valine for the G117H, G117K, or G117V mutant, respectively, examined in this study, by using the LEaP module in Amber 9.<sup>57,58</sup> The missing hydrogen atoms were also added by using the LEaP module in Amber 9.<sup>57,58</sup> Then, sarin-phosphorylated wild-type human BChE and G117H mutant conjugates were built by referring to the orientation of the X-ray crystal structure of sarin-inhibited mouse AChE (non-aged enzyme, PDB entry 2JGG).<sup>59</sup> Each system was solvated in an orthorhombic box of TIP3P water molecules<sup>60</sup> with a minimal solute–wall distance of 10 Å.

The molecular dynamics (MD) simulations in this study were conducted using the Sander module of Amber 9.<sup>57,58</sup> The prepared systems were fully energy-minimized, followed by equilibration via the gradual increase in the temperature from 10 to 298.15 K. Then, the production MD simulations at 298.15 K were kept running for ~2 ns. In the MD simulation, the time step of the simulation was 2 fs with a cutoff of 10 Å for nonbonded interactions. The Shake procedure<sup>61,62</sup> was employed to constrain all bonds involving hydrogen atoms. The partial atomic charges for the atoms in the nonstandard residue (sarin-phosphorylated S198) were calculated by using

the restrained electrostatic potential (RESP) protocol implemented in the Antechamber module of Amber 9.<sup>57,58</sup>

**QM/MM-FE Simulation.** The initial structures used for the QM/MM calculations were chosen from snapshots of the MD simulations of sarin-phosphorylated wild-type human BChE and the G117H mutant, and the chosen structures were close to the corresponding average simulated structures. All of the QM/MM calculations were performed by using a pseudobond QM/MM method<sup>29,30</sup> implemented recently in a revised version<sup>49</sup> of Gaussian03<sup>63</sup> and AMBER8.<sup>64</sup> The QM–MM interface was treated by a pseudobond approach, where a seven-valence electron atom with an effective core potential is constructed to replace the boundary atom of the environment part and to form a pseudobond with the boundary atom of the active part. In the QM/MM calculation, the attacking water molecule and the side chains of the sarin-phosphorylated S198, H438, and E325 (that form the catalytic triad) were considered as the QM atoms, whereas the other atoms were regarded as MM atoms (Figure 1). The QM/MM calculations were



**Figure 1.** Division of the QM/MM systems for simulating spontaneous reactivation reactions of sarin-phosphorylated wild-type human BChE and the G117H mutant. Atoms colored blue were treated by the QM method. Three boundary carbon atoms colored red were treated with the improved pseudobond parameters.<sup>29</sup> All other atoms belong to the MM subsystem.

performed using an iterative energy minimization procedure<sup>31</sup> at the B3LYP/6-31G\*:AMBER level; i.e., the QM calculations were conducted at the B3LYP/6-31G\* level, whereas the MM calculations were conducted by using the AMBER force field implemented in AMBER8. For the QM subsystem, the convergence criterion for the geometry optimizations follows the original Gaussian03 defaults; for the MM subsystem, the geometry optimization convergence criterion is a root-mean-square deviation (rmsd) of the energy gradient of <0.1 kcal mol<sup>-1</sup> Å<sup>-1</sup>. An iterative restrained optimization procedure was then repeatedly applied to different points along the reaction coordinate, resulting in a minimal-energy path. Full QM/MM geometry optimizations were conducted at the B3LYP/6-31G\*:AMBER level, followed by vibrational frequency analyses at the same level to characterize the reactants, intermediates, and transition states. The contribution of the QM subsystem fluctuation to the free energy change was then calculated with the obtained vibrational frequencies using the harmonic approximation. In addition, single-point energy calculations were conducted at the QM/MM (MP2/6-31+G\*:AMBER) level for each geometry along the minimal-energy path.

The free energy changes associated with the QM–MM interaction were then determined by using the free energy perturbation (FEP) method<sup>31</sup> using a revised version<sup>40</sup> of AMBER8. The FEP calculations allowed us to more reasonably determine the relative free energy changes due to the QM–

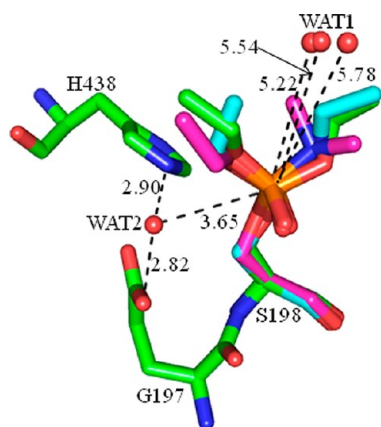


MM interaction. In the FEP calculations, sampling of the MM subsystem was conducted with the QM subsystem frozen at different states along the reaction path.<sup>31</sup> Technically, the final (relative) free energy determined by the QM/MM-FE calculations is the QM part of the QM/MM energy (excluding the Coulombic interaction energy between the point charges of the MM atoms and the ESP charges of the QM atoms) plus the relative free energy change determined by the FEP calculations. In FEP calculations, a time step of 2 fs was used, and bond lengths involving hydrogen atoms were constrained. When the MM subsystem was sampled by MD simulations, the temperature was maintained at 298.15 K. Each FEP calculation (for each geometry of the QM subsystem along the minimal-energy path) consisted of equilibration for 50 ps and sampling for 300 ps.

The MD simulations and QM/MM-FE calculations were performed on a supercomputer (e.g., IBM X-series Cluster with 340 nodes or 1360 processors) at the University of Kentucky Computer Center. The other less-time-consuming modeling and computations were conducted on SGI Fuel workstations and a 34-processor IBM x335 Linux cluster in our own lab.

## RESULTS AND DISCUSSION

**Determination of the Nucleophilic Attacking Water Molecule.** A water molecule is involved and acts as a nucleophile in the spontaneous reactivation of sarin-phosphorylated wild-type BChE. Such a water molecule needs to be close to the phosphorus atom of sarin to perform the nucleophilic attack. From the stereochemical perspective, this water molecule was proposed to be located on the opposite side of the enzyme–phosphorus bond because the enzyme molecule (specifically the oxygen atom of the S198 side chain) is the leaving group of the reactivation reaction.<sup>19</sup> As shown in Figure 2, a water molecule (denoted WAT1) was found at such a



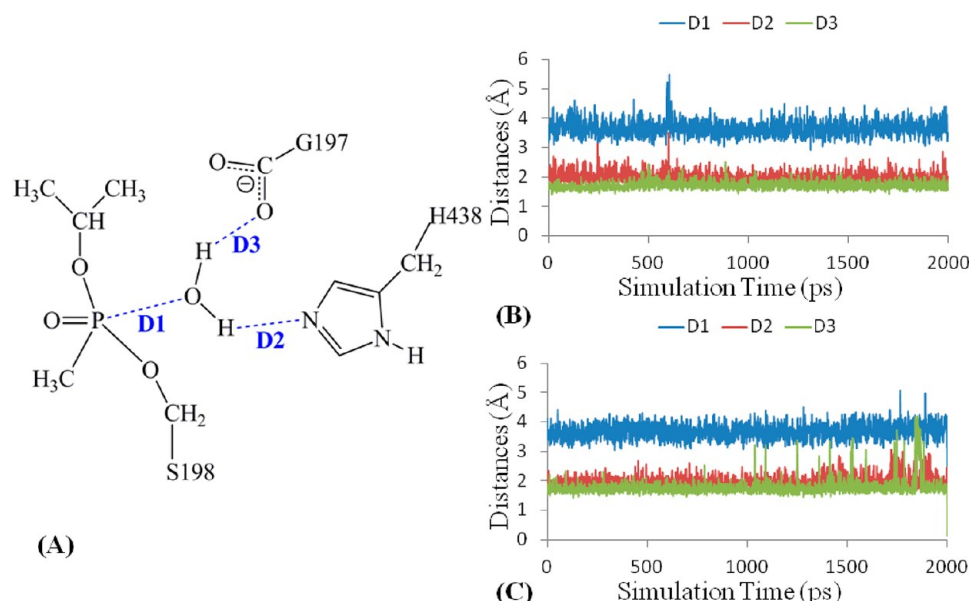
**Figure 2.** Superimposed conformations of two different locations of water in the active sites of the three crystal structures of human BChE phosphorylated by different organophosphorus compounds (PDB entries 1XLW, 3DJY, and 2WIJ) in which the carbon atoms of organophosphorus compounds are colored green, magenta, and cyan, respectively. The distances are given in angstroms.

location in three crystal structures, i.e., the diethylphosphorylated BChE (PDB entry 1XLW)<sup>65</sup> and the non-aged forms of human BChE inhibited by tabun (PDB entry 3DJY)<sup>66</sup> and by tabun analogue TA5 (PDB entry 2WIJ).<sup>67</sup> However, WAT1 was too far from the phosphorus atom to initiate the nucleophilic attack. The respective distances between the

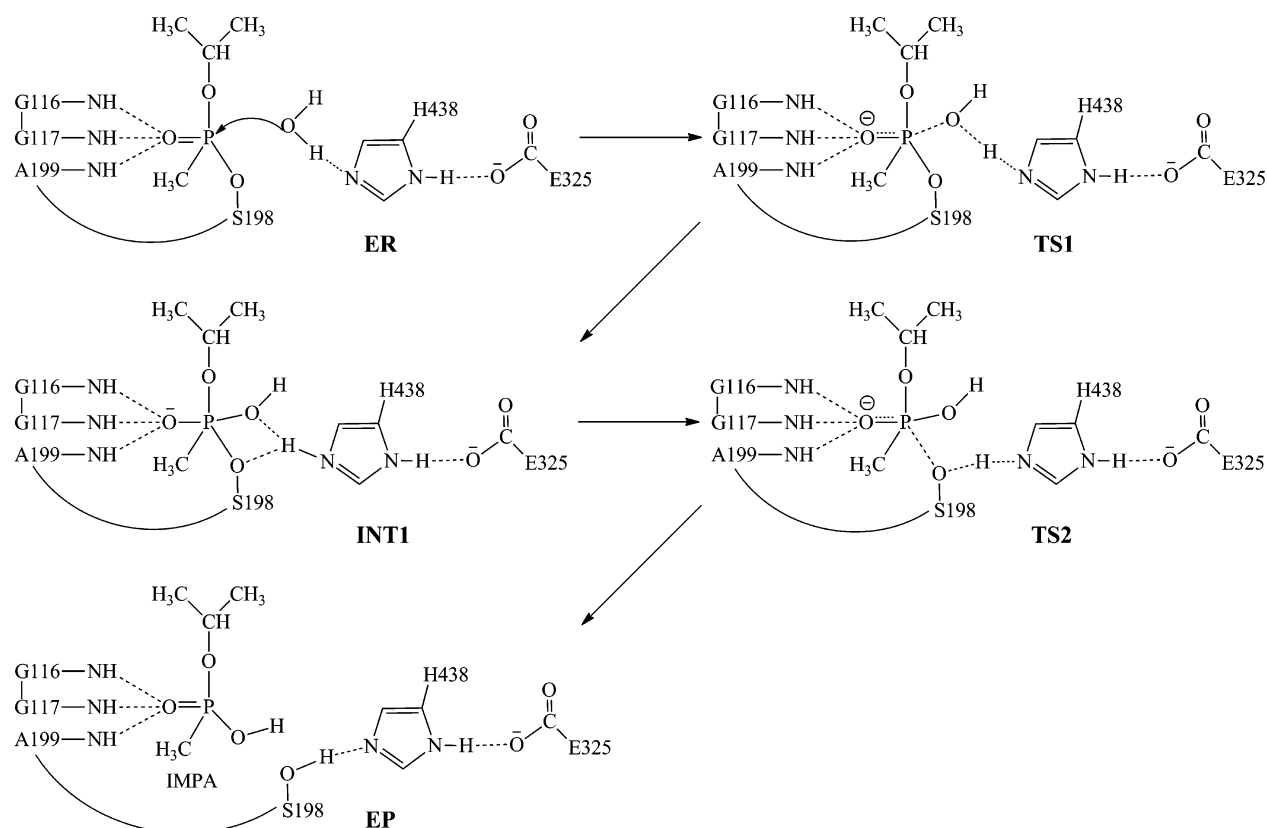
oxygen atom of WAT1 and the phosphorus atom of the OP were 5.78, 5.22, and 5.54 Å, respectively, in these crystal structures. Considering the significant effects of the protein environment, the true nucleophilic water molecule may not necessarily sit well on the opposite side of the enzyme–phosphorus bond to initiate a linear nucleophilic attack on the phosphorus atom. Another water molecule close to the phosphorus atom might be more appropriate for performing a nonlinear nucleophilic attack on the phosphorus atom. By carefully checking the surrounding environment of the phosphorus atom, we noted that another water molecule (denoted WAT2) was located on the opposite side of the ethoxy group toward the acyl pocket in the crystal structures of the diethylphosphorylated BChE mentioned above. The distance between the phosphorus atom and the oxygen of WAT2 was 3.65 Å, which was 2.13 Å shorter than the corresponding distance for the oxygen of WAT1. In addition, there were two hydrogen bonds between WAT2 and the side chains of two essential residues, E197 and H438. The distances between the oxygen atom of WAT2 and the oxygen atom of the E197 side chain and the nitrogen (N<sup>e</sup>) atom of the H438 side chain were 2.82 and 2.90 Å, respectively, indicating that WAT2 was strongly stabilized by these two hydrogen bonds. Clearly, a water molecule at the location of WAT2 is favored to serve as the nucleophile to initiate the nucleophilic attack on the phosphorus atom.

To further identify which water molecule, WAT1 or WAT2, should be the nucleophile, an ~2 ns MD simulation was performed on sarin-phosphorylated wild-type human BChE. At the end of the MD simulation, because of the dynamic distribution of water molecules, another water molecule (denoted WAT532) was found to be located on the original position of WAT2 and WAT532 formed hydrogen bonds with the carboxylate group of the E197 side chain and the nitrogen (N<sup>e</sup>) atom of the H438 side chain. The traces of simulated key distances associated with WAT532 are given in Figure 3B, including the distance ( $D_1$ ) between the oxygen atom of WAT532 and the phosphorus atom as well as distances  $D_2$  and  $D_3$  reflecting the hydrogen bonds from WAT532 to H438 and E197, respectively. The average value of  $D_1$  is  $3.67 \pm 0.28$  Å, suggesting that WAT532 is appropriate for the attack by the phosphorus atom. The average values of  $D_2$  and  $D_3$  are  $2.00 \pm 0.19$  and  $1.72 \pm 0.12$  Å, respectively, indicating that WAT532 has strong hydrogen bonding interactions with E197 and H438. In addition, through the hydrogen bond between WAT532 and H438, it is possible that H438 accepts a proton from WAT532 and serves as a general base to facilitate the nucleophilic attack on the phosphorus atom by WAT532. On the basis of the results and discussion above, WAT532 is determined to be the nucleophile to initiate the spontaneous reactivation reaction of sarin-phosphorylated wild-type human BChE.

**Spontaneous Reactivation Reactions of the Sarin-Phosphorylated Wild-Type Enzyme and G117H Mutant of Human BChE.** *Reaction Pathway for Wild-Type Human BChE.* The QM/MM reaction coordinate calculations at the B3LYP/6-31G\*:AMBER level revealed that the spontaneous reactivation reaction consists of two reaction steps. The first step is the nucleophilic attack on the phosphorus atom by the oxygen atom of WAT532, and the second step is the decomposition of the pentacoordinate intermediate, resulting in the departure of the oxygen of the S198 side chain and the generation of a nontoxic product, i.e., *O*-isopropyl methylphosphonic acid (IMPA). The schematic representation of the



**Figure 3.** Key internuclear distances (A) ( $D_1$ – $D_3$ ) vs the simulation time in the molecular dynamics simulation of the sarin-phosphorylated (B) wild-type enzyme and (C) G117H mutant of human BChE.

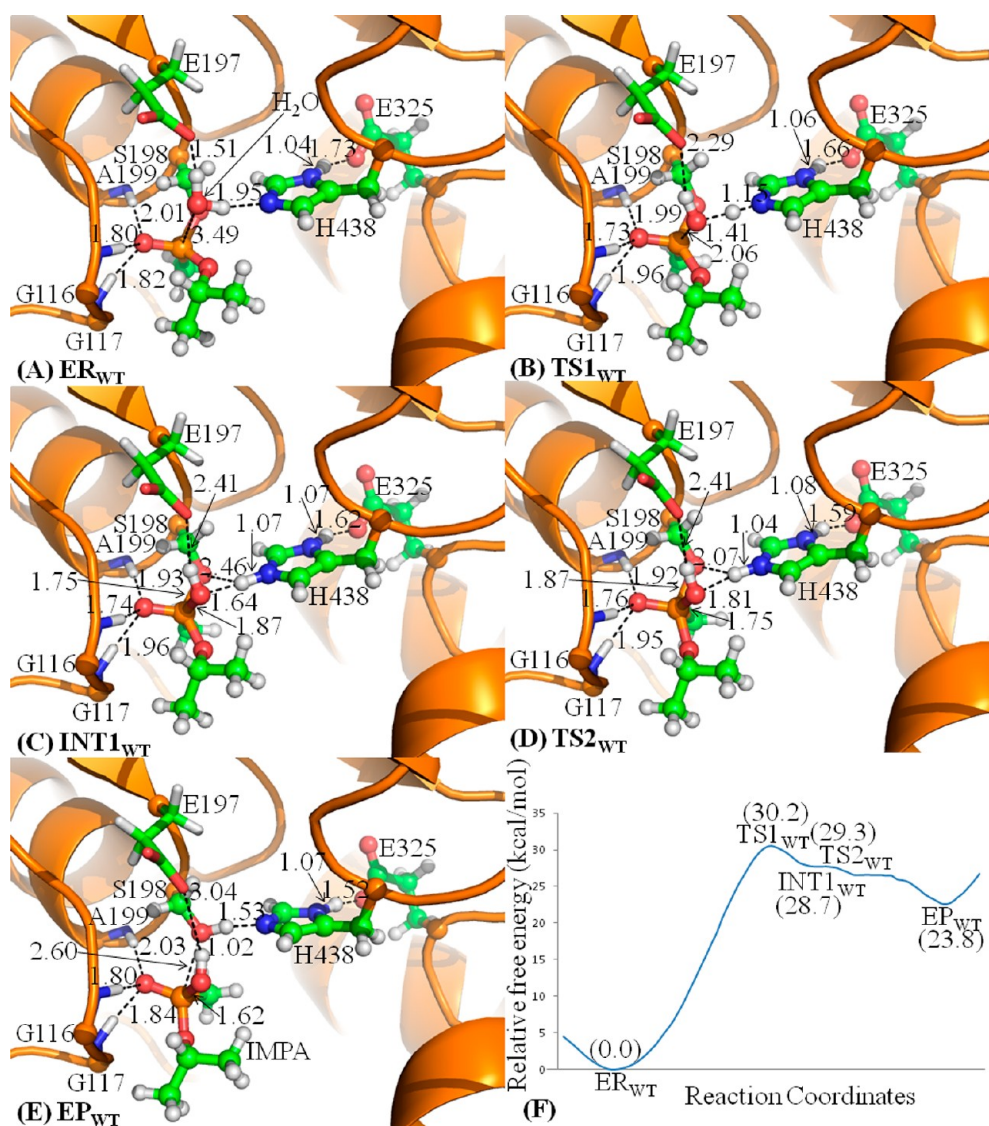


**Figure 4.** Schematic representation of the pathway for the spontaneous reactivation reaction of sarin-phosphorylated wild-type human BChE. The dashed lines in the transition states represent the covalent bonds that form or break during the reaction steps.

pathway is shown in Figure 4. The optimized geometries of the reactant, transition states, intermediate, and product are depicted in Figure 5. Below, we discuss each step in detail.

In the nucleophilic attack process, the oxygen atom ( $O^w$ ) of WAT532 gradually approaches and attacks the phosphorus atom (P), while a hydrogen atom ( $H^w$ ) of WAT532 is gradually transferred to the nitrogen atom ( $N^e$ ) of the H438 side chain.

Because this reaction step involves the formation of both the  $P-O^w$  and  $N^e-H^w$  bonds and the breaking of the  $O^w-H^w$  bond, the distances between the P and  $O^w$  atoms ( $R_{P-O^w}$ ), between the  $N^e$  and  $H^w$  atoms ( $R_{N^e-H^w}$ ), and between the  $O^w$  and  $H^w$  atoms ( $R_{O^w-H^w}$ ) reflect the nature of the current chemical reaction step, and thus, the reaction coordinate was set as  $R_{O^w-H^w} - R_{P-O^w} - R_{N^e-H^w}$ .



**Figure 5.** Optimized geometries (A–E) and free energy profile (F) of the spontaneous reactivation reaction of sarin-phosphorylated wild-type human BChE. The QM atoms are represented as balls and sticks. Carbon, oxygen, nitrogen, phosphorus, and hydrogen atoms are colored green, red, blue, orange, and white, respectively. The key distances are given in angstroms. Figures 6–8 below are represented using the same method.

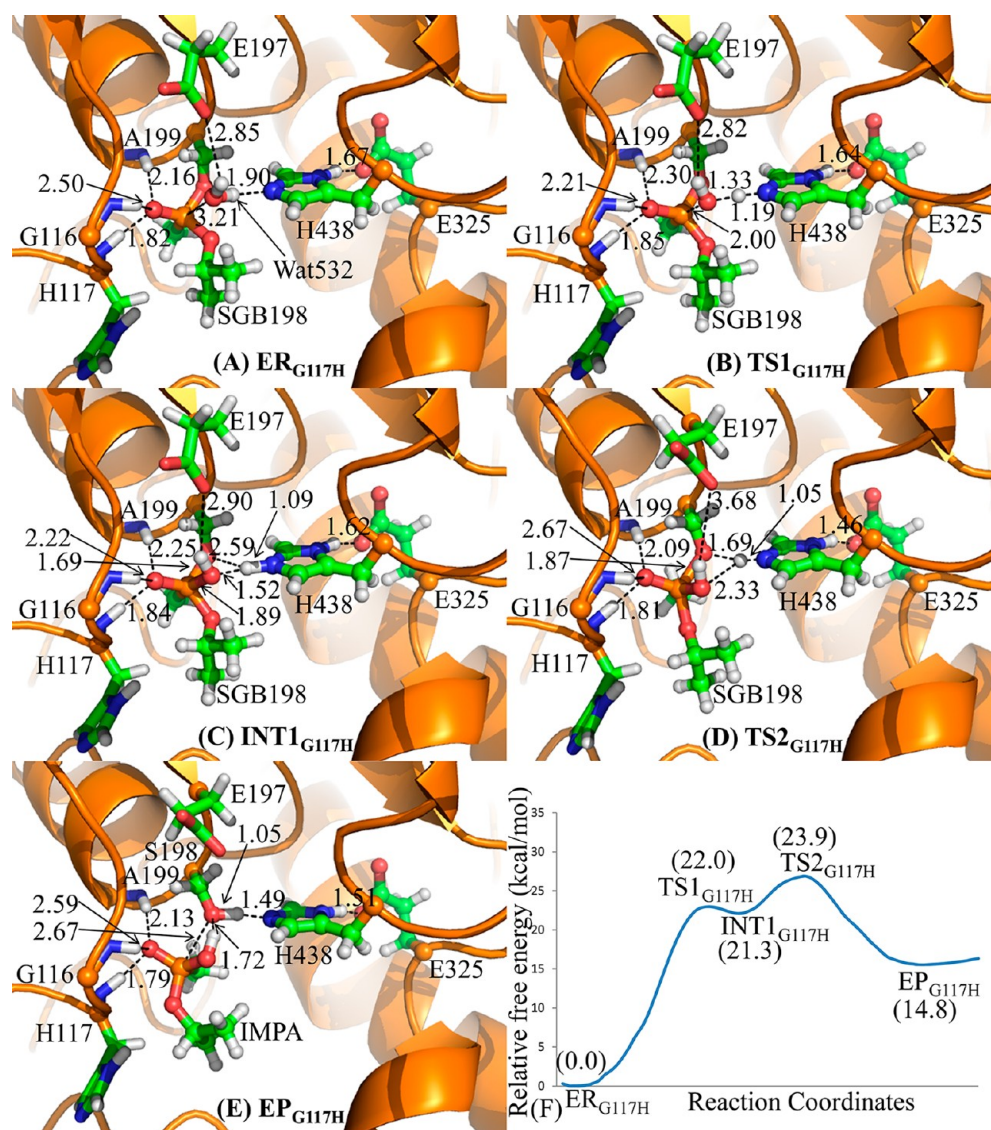
As shown in Figure 5,  $ER_{WT}$  has a tetrahedrally coordinated phosphorus conformation. The distances between the P and  $O^w$  atoms is 3.49 Å in  $ER_{WT}$  and 1.87 Å in  $INT1_{WT}$ , indicating the formation of a P– $O^w$  bond during the reaction process via transition state  $TS1_{WT}$ . The distance between the  $N^e$  and  $H^w$  atoms is 1.95 Å in  $ER_{WT}$  and 1.07 Å in  $INT1_{WT}$ , along with the distance between the  $O^w$  and  $H^w$  atoms being elongated to 1.64 Å in  $INT1_{WT}$ , illustrating the completion of the transfer of the proton ( $H^w$ ) from WAT532 to H438.  $INT1_{WT}$  is a pentacoordinated phosphorus intermediate and has a trigonal bipyramidal conformation with the  $O^w$  atom and methyl group of sarin in the axial orientation. As a general base, H438 accepts a proton ( $H^w$ ) from WAT532 to increase the nucleophilicity of WAT532 and facilitates the nucleophilic attack on the phosphorus atom, resulting in the protonated H438 in  $INT1_{WT}$ . With regard to the role of E325, which is one of the three members of the catalytic triad in human BChE, the optimized distances between the  $H^d$  and  $O^d$  atoms in all states indicate that no proton transfer between H438 and E325 takes place. However, a strong  $N^d$ – $H^d$ ... $O^d$  hydrogen bond is formed

between the side chains of these two residues, and this hydrogen bond distance is shortened to 1.62 Å in  $INT1_{WT}$  from 1.73 Å in  $ER_{WT}$ , suggesting that the hydrogen bonding interaction becomes stronger during the reaction process. The stronger hydrogen bonding interaction is useful for stabilizing the developed positive charge of H438 in  $TS1_{WT}$  and  $INT1_{WT}$ .

During the decomposition process of the pentacoordinate intermediate, the P– $O^v$  bond length gradually increases while the proton ( $H^w$ ) in the protonated H438 is gradually transferred to the  $O^v$  atom of S198. Because this reaction step consists of the cleavages of the P– $O^v$  and  $N^e$ – $H^w$  bonds and the formation of the bond between the  $O^v$  and  $H^w$  atoms, the distances between the P and  $O^v$  atoms ( $R_{P-O^v}$ ), between the  $N^e$  and  $H^w$  atoms ( $R_{N^e-H^w}$ ), and between the  $O^v$  and  $H^w$  atoms ( $R_{O^v-H^w}$ ) reflect the nature of the current chemical reaction step, and thus, the reaction coordinate was set as  $R_{P-O^v} + R_{N^e-H^w} - R_{O^v-H^w}$ .

As shown in Figure 5, the P– $O^v$  bond length is 1.75 Å in  $INT1_{WT}$  and 2.60 Å in  $EP_{WT}$ , and the P atom is tetrahedrally coordinated in  $EP_{WT}$ , indicating the cleavage of the P– $O^v$  bond





**Figure 6.** Optimized geometries (A–E) and free energy profile (F) of the spontaneous reactivation reaction of the sarin-phosphorylated G117H mutant of human BChE.

and the generation of the nontoxic IMPA. The  $N^{\epsilon}$ – $H^{\omega}$  bond length is 1.07 Å in INT1<sub>WT</sub> and 1.53 Å in EP<sub>WT</sub>, and the distance between the  $O^{\gamma}$  and  $H^{\omega}$  atoms is 2.46 Å in INT1<sub>WT</sub> and 1.02 Å in EP<sub>WT</sub>, showing that the proton ( $H^{\omega}$ ) has been transferred from the nitrogen atom of the H438 side chain to the  $O^{\gamma}$  atom of the S198 side chain. Therefore, the enzyme activity is restored. As in the first reaction step, E325 is also not involved in the formation or cleavage of any covalent bond during the decomposition reaction (second reaction step) as the  $N^{\delta}$ – $H^{\delta}$  bond length is almost unchanged, but the  $O^{\delta}$  atom of the E325 side chain forms a strong  $N^{\delta}$ – $H^{\delta}$ ... $O^{\delta}$  hydrogen bond with H438. The length of this hydrogen bond in EP<sub>WT</sub> is shorter than that in INT1<sub>WT</sub>, and this is probably caused by the rotation of H438 such that it facilitates the transfer of the proton from H438 to S198.

It is well-known that the oxyanion hole consists of the backbone NH groups of G116, G117, and A199. It is interesting to know the catalytic role of the oxyanion hole in the reaction process. On the basis of the geometries shown in Figure 5, the phosphoryl oxygen ( $O^{\alpha}$ ) atom forms strong N– $H^{\alpha}$ ... $O^{\alpha}$  hydrogen bonds with the backbone NH groups of G116,

G117, and A199 during the whole reaction process, and these three hydrogen bonds help to stabilize the developed negative charge of the  $O^{\alpha}$  atom of the sarin-phosphorylated S198 in TS1<sub>WT</sub> and INT1<sub>WT</sub>.

On the basis of the QM/MM-optimized geometries at the B3LYP/6-31G\*:AMBER level, the QM/MM single-point energy calculations were performed at the MP2/6-31+G\*:AMBER level for each geometry along the minimum-energy path for the spontaneous reactivation reaction of sarin-phosphorylated human BChE. For each geometry along the minimum-energy path, the ESP charges determined in the QM subsystem of the QM/MM single-point energy calculation were used in further FEP calculations for estimating the free energy changes along the reaction path. The zero-point and thermal corrections for the QM subsystem are included in the free energy profile determined by the QM/MM-FE calculations and shown in Figure 5F. As seen in Figure 5F, the first reaction step, i.e., the nucleophilic attack on the phosphorus atom by the oxygen of WAT532, is the rate-determining step with the free energy barrier being 30.2 kcal/mol. The calculated free energy barrier of 30.2 kcal/mol is consistent with the available

experimental kinetic data<sup>22</sup> that revealed no significant reaction or a  $k_{\text{cat}}$  of  $<5 \times 10^{-5} \text{ min}^{-1}$  (within experimental error). Therefore, it has been known that  $k_{\text{cat}} < 5 \times 10^{-5} \text{ min}^{-1}$ , although the exact  $k_{\text{cat}}$  value remains to be determined. According to the conventional transition-state theory,<sup>68</sup> when  $k_{\text{cat}} < 5 \times 10^{-5} \text{ min}^{-1}$ , the activation free energy should be at least higher than 25.7 kcal/mol.

**Reaction Pathway for the G117H Mutant of Human BChE.** After the G117H mutant had been constructed by using the LEaP module in Amber 9, an  $\sim 2 \text{ ns}$  MD simulation was performed to identify the conformation of the sarin-phosphorylated G117H mutant of human BChE. With regard to the possibility that H117 acts as a general base, there is no water molecule observed close to H117 or sarin-phosphorylated S198. The H117 side chain is very close to the O-isopropyl group of sarin, and there is clearly not enough space between the sarin-phosphorylated S198 and H117 to accommodate a water molecule, showing that H117 cannot serve as a general base for the reactivation reaction. Similar to the reaction pathway for wild-type human BChE, a water molecule (denoted WAT532) is located on the position of WAT2 and forms two hydrogen bonds with the nitrogen ( $\text{N}^{\text{e}}$ ) atom of the H438 side chain and the carboxyl group of E197 in the sarin-phosphorylated G117H mutant. The traces of simulated key distances associated with WAT532, including the distance ( $D_1$ ) between the oxygen atom of WAT532 and the phosphorus atom of sarin as well as the hydrogen bonding distances ( $D_2$  and  $D_3$ ) between WAT532 and H438 and E197, are shown in Figure 3C. The average value of  $D_1$  is  $3.68 \pm 0.20 \text{ \AA}$ , suggesting that WAT532 is in a position that is appropriate for the attack on the phosphorus atom. The average values of  $D_2$  and  $D_3$  are  $2.00 \pm 0.16$  and  $1.78 \pm 0.14 \text{ \AA}$ , respectively, indicating strong hydrogen bonding interactions with both H438 and E197. Through the hydrogen bond between WAT532 and H438, it is very possible that H438 serves as a general base to accept a proton from WAT532 and facilitate the nucleophilic attack on the phosphorus atom by the oxygen of WAT532.

The reaction pathway for the spontaneous reactivation of the sarin-phosphorylated G117H mutant of human BChE has been investigated by the QM/MM reaction coordinate calculations at the B3LYP/6-31G\*:AMBER level. The results indicate that it follows a reaction pathway similar to that for the spontaneous reactivation of sarin-phosphorylated wild-type human BChE shown in Figure 4. The whole reaction process consists of two steps, i.e., the nucleophilic attack and the decomposition of the pentacoordinated phosphorus intermediate. For each reaction step, the same reaction coordinate used to represent the reaction for wild-type BChE was used to represent the reaction for the G117H mutant. The optimized geometries of the reactant, transition states, intermediate, and product are depicted in Figure 6.

In the first reaction step, the oxygen atom ( $\text{O}^{\text{w}}$ ) of WAT532 gradually approaches and attacks the phosphorus atom (P) of sarin, while the hydrogen atom ( $\text{H}^{\text{w}}$ ) of WAT532 is gradually transferred to the nitrogen atom ( $\text{N}^{\text{e}}$ ) of the H438 side chain. As seen in Figure 6, the distance between the P and  $\text{O}^{\text{w}}$  atoms is  $3.21 \text{ \AA}$  in  $\text{ER}_{\text{G117H}}$  and  $1.89 \text{ \AA}$  in  $\text{INT1}_{\text{G117H}}$ , indicating the formation of the P– $\text{O}^{\text{w}}$  bond during the reaction process via transition state  $\text{TS1}_{\text{G117H}}$ . The distance between the  $\text{N}^{\text{e}}$  and  $\text{H}^{\text{w}}$  atoms is  $1.90 \text{ \AA}$  in  $\text{ER}_{\text{G117H}}$  and  $1.09 \text{ \AA}$  in  $\text{INT1}_{\text{G117H}}$ , along with the distance between the  $\text{O}^{\text{w}}$  and  $\text{H}^{\text{w}}$  atoms being elongated to  $1.52 \text{ \AA}$  in  $\text{INT1}_{\text{G117H}}$ , revealing the transfer of the

proton ( $\text{H}^{\text{w}}$ ) from WAT532 to H438. During the reaction process, the distance between the  $\text{H}^{\text{e}}$  and  $\text{O}^{\text{e}}$  atoms is almost unchanged, indicating that there is no transfer of a proton between H438 and E325, but the strong  $\text{N}^{\text{e}}\text{--}\text{H}^{\text{e}}\cdots\text{O}^{\text{e}}$  hydrogen bond between H438 and E325 helps to stabilize the positively charged H438 in  $\text{TS1}_{\text{G117H}}$  and  $\text{INT1}_{\text{G117H}}$ .

In the second reaction step, the P– $\text{O}^{\text{y}}$  bond length gradually increases while the proton ( $\text{H}^{\text{w}}$ ) in the protonated H438 is gradually transferred to the  $\text{O}^{\text{y}}$  atom of S198. As seen in Figure 6, in  $\text{INT1}_{\text{G117H}}$ , the distance between the  $\text{O}^{\text{y}}$  and  $\text{H}^{\text{w}}$  atoms is  $2.59 \text{ \AA}$ , indicating a very weak  $\text{N}^{\text{e}}\text{--}\text{H}^{\text{w}}\cdots\text{O}^{\text{y}}$  hydrogen bond. This distance is  $0.13 \text{ \AA}$  longer than that in  $\text{INT1}_{\text{WT}}$ , suggesting that it is not as favorable as that in the wild-type human BChE for the transfer of the proton ( $\text{H}^{\text{w}}$ ) from H438 to S198. During the reaction process via  $\text{TS2}_{\text{G117H}}$ , the P– $\text{O}^{\text{y}}$  bond is elongated and the imidazole group of the protonated H438 residue is rotated to place the  $\text{H}^{\text{w}}$  atom closer to the  $\text{O}^{\text{y}}$  atom. The distance between the  $\text{O}^{\text{y}}$  and  $\text{H}^{\text{w}}$  atoms is shortened to  $1.69 \text{ \AA}$  in  $\text{TS2}_{\text{G117H}}$ . This change may facilitate the transfer of the proton ( $\text{H}^{\text{w}}$ ) from the protonated H438 to S198. At the same time, an important angle among the  $\text{O}^{\text{y}}$ , P, and  $\text{O}^{\text{w}}$  atoms ( $\angle\text{O}^{\text{y}}\text{PO}^{\text{w}}$ ) increases to  $132.2^\circ$  in  $\text{TS2}_{\text{G117H}}$  from  $110.7^\circ$  in  $\text{INT1}_{\text{G117H}}$ . This spatial reorganization results in the  $\text{O}^{\text{y}}$  atom being in the axial orientation that is favored for the cleavage of the P– $\text{O}^{\text{y}}$  bond. Finally, in  $\text{EP}_{\text{G117H}}$ , the P– $\text{O}^{\text{y}}$  bond is broken, with the distance between the P and  $\text{O}^{\text{y}}$  atoms being  $2.67 \text{ \AA}$ . The phosphorus atom is tetrahedrally coordinated, indicating that the nontoxic product, i.e., IMPA, is released. The angle  $\angle\text{O}^{\text{y}}\text{PO}^{\text{w}}$  equals  $160.1^\circ$  in  $\text{EP}_{\text{G117H}}$ , meaning that  $\text{O}^{\text{y}}$  and  $\text{O}^{\text{w}}$  are located in the axial orientation before the complete cleavage of the P– $\text{O}^{\text{y}}$  bond. The distances between the  $\text{N}^{\text{e}}$  and  $\text{H}^{\text{w}}$  atoms and between the  $\text{O}^{\text{y}}$  and  $\text{H}^{\text{w}}$  atoms are  $1.49$  and  $1.05 \text{ \AA}$ , respectively, showing that the transfer of the proton ( $\text{H}^{\text{w}}$ ) from the H438 side chain to the S198 side chain has been completed. During the reaction process, the strong  $\text{N}^{\text{e}}\text{--}\text{H}^{\text{e}}\cdots\text{O}^{\text{e}}$  hydrogen bond remains between H438 and E325 and there is no proton transfer between them.

It should be pointed out that the spatial reorganization observed for the G117H mutant has not been observed in the corresponding step of the reaction for wild-type human BChE, which reveals a remarkable difference between the reactivation reactions of the sarin-phosphorylated wild-type enzyme and the G117H mutant of human BChE.

Another significant difference can be noted in the hydrogen bonding interaction between the phosphoryl oxygen ( $\text{O}^{\text{a}}$ ) and the oxyanion hole during the reaction process. There are three  $\text{N}\cdots\text{H}\cdots\text{O}$  hydrogen bonds between the  $\text{O}^{\text{a}}$  atom and the NH groups of the three residues (G116, H117, and A199) forming the oxyanion hole during the reaction process for the G117H mutant, which is similar to the reaction process for wild-type human BChE. However, there are relatively larger changes in the  $\text{H}\cdots\text{O}^{\text{a}}$  distances of these hydrogen bonds, particularly those with G116 and A199, in the reaction of the G117H mutant compared to that of wild-type human BChE. For example, the  $\text{H}\cdots\text{O}^{\text{a}}$  distance with the NH group of the G116 backbone increases from  $2.22 \text{ \AA}$  in  $\text{INT1}_{\text{G117H}}$  to  $2.67 \text{ \AA}$  in  $\text{TS2}_{\text{G117H}}$ , while the  $\text{H}\cdots\text{O}^{\text{a}}$  distance with the NH group of the A199 backbone decreases from  $2.25 \text{ \AA}$  in  $\text{INT1}_{\text{G117H}}$  to  $2.09 \text{ \AA}$  in  $\text{TS2}_{\text{G117H}}$ . In comparison, there are small changes (within  $0.02 \text{ \AA}$ ) in the corresponding distances in the reaction of wild-type human BChE from  $\text{INT1}_{\text{WT}}$  to  $\text{TS2}_{\text{WT}}$ .

Further, the free energy profile depicted in Figure 6F reveals that the second reaction step is the rate-determining step for

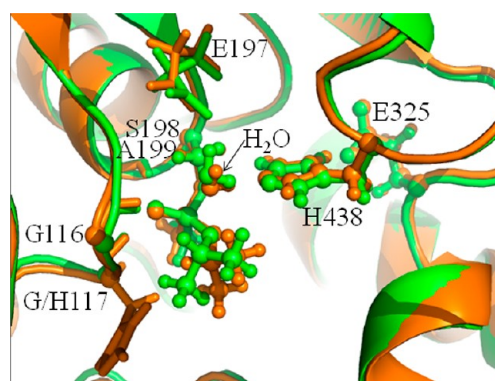


the G117H mutant, and the overall free energy barrier is the free energy change from  $ER_{G117H}$  to  $TS2_{G117H}$ . The calculated overall free energy barrier is 23.9 kcal/mol. The calculated free energy barrier of 23.9 kcal/mol is in good agreement with the available experimental kinetic data ( $k_{cat} = 8.7 \times 10^{-5} s^{-1}$ ).<sup>21,22</sup> According to the conventional transition-state theory,<sup>68</sup> when  $k_{cat} = 8.7 \times 10^{-5} s^{-1}$ , the activation free energy should be 23.0 kcal/mol, which is close to our computational value of 23.9 kcal/mol.

**Effects of the G117H Mutation.** On the basis of the results discussed above, it is evident that the spontaneous reactivation reactions of the sarin-phosphorylated wild-type enzyme and G117H mutant of human BChE are all catalyzed by H438, which acts as a general base. The G117H mutation cannot introduce a second nucleophilic center into the active site of human BChE, which was supposed to activate a water molecule to initiate the spontaneous reactivation reaction. Our results do not support the aforementioned hypothesis<sup>19</sup> that the H117 residue in the G117H mutant acts as a general base to catalyze the OP hydrolysis. A careful comparison between the uncovered spontaneous reactivation mechanisms of the sarin-phosphorylated wild-type enzyme and G117H mutant of human BChE reveals some remarkable differences caused by the G117H mutation, as discussed below.

**G117H Mutation-Caused Change in the Oxyanion Hole Interacting with the Phosphoryl Oxygen.** As discussed above, the oxyanion hole consists of the backbone NH groups of G116, G117 (or H117 in the G117H mutant), and A199. As seen in Figure 5, the oxyanion hole in wild-type human BChE forms three strong hydrogen bonds with the phosphoryl oxygen and the average values of the  $H \cdots O^{\alpha}$  distances of these three hydrogen bonds with G116, G117, and A199 in all of the optimized geometries (including  $ER_{WT}$ ,  $TS1_{WT}$ ,  $INT1_{WT}$ ,  $TS2_{WT}$ , and  $EP_{WT}$ ) in Figure 5 are  $1.77 \pm 0.03$ ,  $1.90 \pm 0.07$ , and  $1.98 \pm 0.05$  Å, respectively, indicating that these hydrogen bonds are rather strong and stable during the reaction process. Similarly, for the G117H mutant of human BChE shown in Figure 6, the oxyanion hole also forms three hydrogen bonds with the phosphoryl oxygen. However, the average values of the  $H \cdots O^{\alpha}$  distances of these three hydrogen bonds with G116, H117, and A199 in all of the optimized geometries in Figure 6 are  $2.44 \pm 0.21$ ,  $1.82 \pm 0.02$ , and  $2.19 \pm 0.09$  Å, respectively. Clearly, the G117H mutation makes the hydrogen bond of the phosphoryl oxygen with H117 in the G117H mutant stronger and more stable than that with G117 in wild-type human BChE, while making the other two hydrogen bonds (with G117 and A199) in the G117H mutant weaker and less stable than those in wild-type human BChE. When G117 is replaced with H117, the local environment of the oxyanion hole is changed by the larger side chain of H117 and the three NH groups of the oxyanion hole are redistributed, resulting in the changes in the hydrogen bonds between the oxyanion hole and the phosphoryl oxygen. Thus, the oxyanion hole is skewed by the G117H mutation.

**G117H Mutation-Caused Change in the Conformation of Sarin-Phosphorylated S198.** The skewed oxyanion hole makes some change in the conformation of sarin-phosphorylated S198. Depicted in Figure 7 are the superimposed conformations of  $ER_{WT}$  and  $ER_{G117H}$ , showing that the O-isopropyl group of sarin in the G117H mutant of human BChE rotates away from the side chain of H117. This crucial structural difference caused by the G117H mutation makes the phosphorus atom in the G117H mutant more open to the

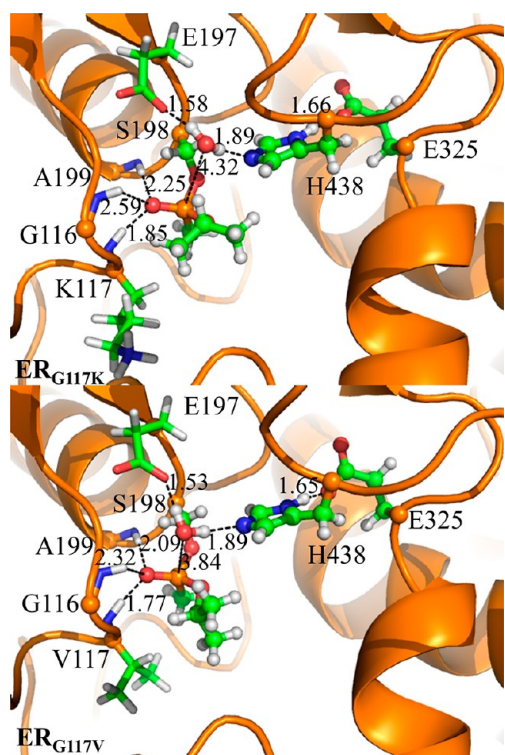


**Figure 7.** Superimposed conformations of  $ER_{WT}$  and  $ER_{G117H}$ . The wild type and G117H mutant of human BChE are colored green and orange, respectively.

nucleophilic water molecule and, thus, decreases the steric hindrance for the nucleophilic attack. According to the structures of  $ER_{WT}$  and  $ER_{G117H}$  depicted in Figures 5 and 6, the distances between the P and  $O^{\alpha}$  atoms are 3.49 and 3.21 Å, respectively, illustrating that the water molecule in the G117H mutant is closer to the phosphorus atom than in the wild type. Hence, this conformational change caused by the G117H mutation facilitates the nucleophilic attack on the phosphorus atom of sarin by the water molecule.

**G117H Mutation-Caused Change in the Free Energy Profile.** On the basis of the free energy profile for the spontaneous reactivation reaction of sarin-phosphorylated wild-type human BChE given in Figure 5F, the relative free energy of  $TS1_{WT}$  is 30.2 kcal/mol, which is 0.9 kcal/mol higher than that of  $TS2_{WT}$ , meaning that the first reaction step is the rate-determining step and the overall free energy barrier is 30.2 kcal/mol for the reaction of the wild-type enzyme. However, after the G117H mutation, the relative free energy of  $TS1_{G117H}$ , shown in Figure 6F, becomes 22.0 kcal/mol, which is 1.9 kcal/mol lower than that of  $TS2_{G117H}$ , meaning that the second reaction step is the rate-determining step and that the overall free energy barrier is 23.9 kcal/mol for the reaction of the G117H mutant. Apparently, the G117H mutation remarkably changes the rate-determining step from the first reaction step to the second reaction step and effectively decreases the overall free energy barrier of the spontaneous reactivation reaction of sarin-phosphorylated human BChE.

All of the geometric and energetic results discussed above indicate that the role of the G117H mutation is to skew the oxyanion hole and to make the phosphorus atom more open to the nucleophilic attack by a water molecule so that the overall free energy barrier for the spontaneous reactivation reaction of sarin-phosphorylated human BChE can be decreased effectively. For comparison, we also examined the  $ER$  structures of the G117V ( $ER_{G117V}$ ) and G117K ( $ER_{G117K}$ ) mutants by using the same computational protocol, i.e., the MD simulation for 2 ns followed by the QM/MM geometry optimization, for each of the mutants. It turned out that in the QM/MM-optimized  $ER_{G117V}$  and  $ER_{G117K}$  structures (see Figures 8), the distances between the P and  $O^{\alpha}$  atoms are 3.84 and 4.32 Å, respectively. The  $P \cdots O^{\alpha}$  distances in both the  $ER_{G117V}$  (3.84 Å) and  $ER_{G117K}$  (4.32 Å) structures are significantly longer than that in the  $ER_{WT}$  structure (3.49 Å). Therefore, whereas the G117H mutation made the  $P \cdots O^{\alpha}$  distance shorter, the G117V and G117K mutations made the  $P \cdots O^{\alpha}$  distance longer compared to that in  $ER_{WT}$  (3.21 Å). On the basis of these computational



**Figure 8.** QM/MM-optimized geometries of ER<sub>G117V</sub> and ER<sub>G117K</sub>.

data, one may reasonably expect that the G117V and G117K mutations would increase the free energy barrier, whereas the G117H mutation decreased the free energy barrier for the spontaneous reactivation reaction of sarin-phosphorylated human BChE. In fact, the G117K mutant was examined in a previously reported experimental study,<sup>22</sup> and no spontaneous reactivation reaction was observed for the G117K mutant. The experimental observation for the G117K mutant was qualitatively consistent with our computational insights.

## CONCLUSION

Spontaneous reactivation reactions of the sarin-phosphorylated wild-type enzyme and G117H mutant of human BChE have been studied by using the first-principles QM/MM-FE calculations. For wild-type human BChE, the attacking water molecule is located on the opposite side of the methyl group toward the acyl pocket of human BChE. The water molecule is stabilized in this location by E197 and H438 through strong hydrogen bonds. The reaction pathway consists of two steps. The first step is the nucleophilic attack on the phosphorus atom of sarin by the oxygen atom of the water molecule, along with the transfer of a proton from the water molecule to the H438 side chain. The second reaction step is the decomposition of the pentacoordinate intermediate, resulting in the active form of human BChE. According to the calculated free energy profile, the first reaction step is the rate-determining step of the reaction with the calculated free energy barrier being 30.2 kcal/mol, which is consistent with the reported experimental observation that the rate constant ( $k_{\text{cat}}$ ) should be at least smaller than  $5 \times 10^{-5} \text{ min}^{-1}$  (or the activation free energy should be at least higher than 25.7 kcal/mol).

For the G117H mutant of human BChE, the attacking water molecule occupies the same position as in wild-type human BChE, and no water molecule can be close to H117, demonstrating that H117 cannot act as a general base in the

spontaneous reactivation. The reaction of the G117H mutant also consists of two reaction steps. The first step is similar to that in the spontaneous reactivation reaction of sarin-phosphorylated wild-type human BChE. However, the second step is remarkably different. A spatial reorganization, which is not involved in the second step of the reaction of wild-type human BChE, is involved in the second step of the reaction of the G117H mutant. The spatial reorganization results in the oxygen atom of S198 side chain sitting in the axial orientation and being close to the transferring proton in the protonated H438, thus facilitating the departure of the oxygen of the S198 side chain from the phosphorus atom. The second reaction step is the rate-determining step of the entire reaction of the G117H mutant. The overall free energy barrier is the free energy change (23.9 kcal/mol) from the initial reactant ER<sub>G117H</sub> to transition state TS<sub>G117H</sub>. The calculated overall free energy barrier of 23.9 kcal/mol is in good agreement with the experimentally derived activation free energy of 23.0 kcal/mol (based on the reported experimental rate constant of  $8.7 \times 10^{-5} \text{ s}^{-1}$ ).

For both wild-type human BChE and the G117H mutant, E325 serves as one of the three members of the catalytic triad to stabilize the developed positive charge on H438, but there is no proton transfer between H438 and E325. The oxyanion hole forms three hydrogen bonds with the phosphoryl oxygen to stabilize the developed negative charge on the phosphoryl oxygen during the reactivation reaction.

Further, with regard to how the G117H mutation accelerates the reactivation reaction process, the obtained structural and energetic data reveal the specific role of the G117H mutation. It has been demonstrated that the G117H mutation does not introduce a second nucleophilic center into the active site of human BChE, which was supposed to activate the water molecule to initiate the spontaneous reactivation reaction. Instead, the G117H mutation skews the oxyanion hole and makes the phosphorus atom more open to the nucleophilic water molecule, resulting in the dramatically decreased overall free energy barrier of the reactivation reaction. In comparison, the G117V and G117K mutations place the phosphorus atom farther from the nucleophilic water molecule.

The obtained mechanistic insights provide a valuable guide to the rational design of new mutants of human BChE with an improved catalytic activity against sarin. In particular, future work should be focused on the computational identification of the new mutants that can more effectively stabilize the transition state associated with the second reaction step.

In general, this computational study demonstrates how a single amino acid mutation on an enzyme can dramatically change the rate-determining step of the enzymatic reaction from one reaction step to another and considerably improve the activity against a substrate of interest. The obtained novel mechanistic insights should be interesting to researchers in a variety of fields of chemistry and related disciplines, such as biochemistry, neurochemistry, phosphorus chemistry, agricultural chemistry, computational chemistry, chemical defense, human health sciences, catalyst design, drug design and discovery, and protein engineering design.

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# Author Contributions

Y.Y. and J.L. contributed equally to this work.

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# Notes

The authors declare no competing financial interest.

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# REFERENCES

- (1) MacPhee-Quigley, K., Taylor, P., and Taylor, S. (1985) Primary structures of the catalytic subunits from two molecular forms of acetylcholinesterase. A comparison of NH<sub>2</sub>-terminal and active center sequences. *J. Biol. Chem.* 260, 12185–12189.
- (2) Marrs, T. C. (1993) Organophosphate poisoning. *Pharmacol. Ther.* 58, 51–66.
- (3) Taylor, P., Radic, Z., Hosea, N. A., Camp, S., Marchot, P., and Berman, H. A. (1995) Structural bases for the specificity of cholinesterase catalysis and inhibition. *Toxicol. Lett.* 82–83, 453–458.
- (4) Black, R. M., Clarke, R. J., Read, R. W., and Reid, M. T. (1994) Application of gas chromatography-mass spectrometry and gas chromatography-tandem mass spectrometry to the analysis of chemical warfare samples, found to contain residues of the nerve agent sarin, sulphur mustard and their degradation products. *J. Chromatogr. A* 662, 301–321.
- (5) Fidder, A., Hulst, A. G., Noort, D., de Ruiter, R., van der Schans, M. J., Benschop, H. P., and Langenberg, J. P. (2002) Retrospective Detection of Exposure to Organophosphorus Anti-Cholinesterases: Mass Spectrometric Analysis of Phosphorylated Human Butyrylcholinesterase. *Chem. Res. Toxicol.* 15, 582–590.
- (6) Nagao, M., Takatori, T., Matsuda, Y., Nakajima, M., Iwase, H., and Iwade, K. (1997) Definitive evidence for the acute sarin poisoning diagnosis in the Tokyo subway. *Toxicol. Appl. Pharmacol.* 144, 198–203.
- (7) Noort, D., Hulst, A. G., Platenburg, D. H., Polhuijs, M., and Benschop, H. P. (1998) Quantitative analysis of O-isopropyl methylphosphonic acid in serum samples of Japanese citizens allegedly exposed to sarin: Estimation of internal dosage. *Arch. Toxicol.* 72, 671–675.
- (8) Polhuijs, M., Langenberg, J. P., and Benschop, H. P. (1997) New method for retrospective detection of exposure to organophosphorus anticholinesterases: Application to alleged sarin victims of Japanese terrorists. *Toxicol. Appl. Pharmacol.* 146, 156–161.
- (9) Tsuchihashi, H., Katagi, M., Nishikawa, M., and Tatsuno, M. (1998) Identification of metabolites of nerve agent VX in serum collected from a victim. *J. Anal. Toxicol.* 22, 383–388.
- (10) Broomfield, C. A., and Kirby, S. D. (2001) Progress on the road to new nerve agent treatments. *J. Appl. Toxicol.* 21 (Suppl. 1), S43–S46.
- (11) Koca, J., Zhan, C. G., Rittenhouse, R. C., and Ornstein, R. L. (2003) Coordination number of zinc ions in the phosphotriesterase active site by molecular dynamics and quantum mechanics. *J. Comput. Chem.* 24, 368–378.

- (12) Zheng, F., Zhan, C. G., and Ornstein, R. L. (2001) Theoretical studies of reaction pathways and energy barriers for alkaline hydrolysis of phosphotriesterase substrates paraoxon and related toxic phosphofluoridate nerve agents. *J. Chem. Soc., Perkin Trans. 2*, 2355–2363.
- (13) Koca, J., Zhan, C. G., Rittenhouse, R. C., and Ornstein, R. L. (2001) Mobility of the active site bound paraoxon and sarin in zinc-phosphotriesterase by molecular dynamics simulation and quantum chemical calculation. *J. Am. Chem. Soc.* 123, 817–826.
- (14) Zhan, C. G., de Souza, O. N., Rittenhouse, R., and Ornstein, R. L. (1999) Determination of two structural forms of catalytic bridging ligand in zinc-phosphotriesterase by molecular dynamics simulation and quantum chemical calculation. *J. Am. Chem. Soc.* 121, 7279–7282.
- (15) Zheng, F., Zhan, C.-G., and Ornstein, R. L. (2002) Theoretical Determination of Two Structural Forms of the Active Site in Cadmium-Containing Phosphotriesterases. *J. Phys. Chem. B* 106, 717–722.
- (16) Broomfield, C. A., Maxwell, D. M., Solana, R. P., Castro, C. A., Finger, A. V., and Lenz, D. E. (1991) Protection by butyrylcholinesterase against organophosphorus poisoning in nonhuman primates. *J. Pharmacol. Exp. Ther.* 259, 633–638.
- (17) Raveh, L., Grauer, E., Grunwald, J., Cohen, E., and Ashani, Y. (1997) The stoichiometry of protection against soman and VX toxicity in monkeys pretreated with human butyrylcholinesterase. *Toxicol. Appl. Pharmacol.* 145, 43–53.
- (18) Gorelick, D. A. (1997) Enhancing cocaine metabolism with butyrylcholinesterase as a treatment strategy. *Drug Alcohol Depend.* 48, 159–165.
- (19) Jarv, J. (1984) Stereochemical Aspects of Cholinesterase Catalysis. *Bioorg. Chem.* 12, 259–278.
- (20) Lockridge, O., Blong, R. M., Masson, P., Froment, M. T., Millard, C. B., and Broomfield, C. A. (1997) A single amino acid substitution, Gly117His, confers phosphotriesterase (organophosphorus acid anhydride hydrolase) activity on human butyrylcholinesterase. *Biochemistry* 36, 786–795.
- (21) Millard, C. B., Lockridge, O., and Broomfield, C. A. (1995) Design and expression of organophosphorus acid anhydride hydrolase activity in human butyrylcholinesterase. *Biochemistry* 34, 15925–15933.
- (22) Broomfield, C. A., Lockridge, O., and Millard, C. B. (1999) Protein engineering of a human enzyme that hydrolyzes V and G nerve agents: Design, construction and characterization. *Chem.-Biol. Interact.* 119–120, 413–418.
- (23) Amitay, M., and Shurki, A. (2009) The structure of G117H mutant of butyrylcholinesterase: Nerve agents scavenger. *Proteins* 77, 370–377.
- (24) Nachon, F., Carletti, E., Wandhammer, M., Nicolet, Y., Schopfer, L. M., Masson, P., and Lockridge, O. (2011) X-ray crystallographic snapshots of reaction intermediates in the G117H mutant of human butyrylcholinesterase, a nerve agent target engineered into a catalytic bioscavenger. *Biochem. J.* 434, 73–82.
- (25) Albaret, C., Masson, P., Broomfield, C. A., El Kaim, L., and Fortier, P. L. (1998) Mechanical aspects of the phosphotriesterase activity of human butyrylcholinesterase G117H mutant. In *Structure and Function of Cholinesterases and Related Proteins* (Doctor, B. P., Ed.) pp 399–405, Plenum Press, New York.
- (26) Kovach, I. M. (1988) Structure and dynamics of serine hydrolase-organophosphate adducts. *J. Enzyme Inhib.* 2, 199–208.
- (27) Qian, N., and Kovach, I. M. (1993) Key active site residues in the inhibition of acetylcholinesterases by soman. *FEBS Lett.* 336, 263–266.
- (28) Hu, P., and Zhang, Y. K. (2006) Catalytic mechanism and product specificity of the histone lysine methyltransferase SET7/9: An ab initio QM/MM-FE study with multiple initial structures. *J. Am. Chem. Soc.* 128, 1272–1278.
- (29) Zhang, Y. K. (2005) Improved pseudobonds for combined ab initio quantum mechanical/molecular mechanical methods. *J. Chem. Phys.* 122, 024114.



- (30) Zhang, Y. K., Lee, T. S., and Yang, W. T. (1999) A pseudobond approach to combining quantum mechanical and molecular mechanical methods. *J. Chem. Phys.* 110, 46–54.
- (31) Zhang, Y. K., Liu, H. Y., and Yang, W. T. (2000) Free energy calculation on enzyme reactions with an efficient iterative procedure to determine minimum energy paths on a combined ab initio QM/MM potential energy surface. *J. Chem. Phys.* 112, 3483–3492.
- (32) Zhang, Y. K. (2006) Pseudobond ab initio QM/MM approach and its applications to enzyme reactions. *Theor. Chem. Acc.* 116, 43–50.
- (33) Cheng, Y., Zhang, Y., and McCammon, J. A. (2005) How Does the cAMP-Dependent Protein Kinase Catalyze the Phosphorylation Reaction: An ab Initio QM/MM Study. *J. Am. Chem. Soc.* 127, 1553–1562.
- (34) Cheng, Y. H., Zhang, Y. K., and McCammon, J. A. (2006) How does activation loop phosphorylation modulate catalytic activity in the cAMP-dependent protein kinase: A theoretical study. *Protein Sci.* 15, 672–683.
- (35) Cisneros, G. A., Wang, M., Silinski, P., Fitzgerald, M. C., and Yang, W. (2004) The protein backbone makes important contributions to 4-oxalocrotonate tautomerase enzyme catalysis: Understanding from theory and experiment. *Biochemistry* 43, 6885–6892.
- (36) Corminboeuf, C., Hu, P., Tuckerman, M. E., and Zhang, Y. (2006) Unexpected Deacetylation Mechanism Suggested by a Density Functional Theory QM/MM Study of Histone-Deacetylase-Like Protein. *J. Am. Chem. Soc.* 128, 4530–4531.
- (37) Hu, P., Wang, S., and Zhang, Y. (2008) How Do SET-Domain Protein Lysine Methyltransferases Achieve the Methylation State Specificity? Revisited by Ab Initio QM/MM Molecular Dynamics Simulations. *J. Am. Chem. Soc.* 130, 3806–3813.
- (38) Ke, Z., Wang, S., Xie, D., and Zhang, Y. (2009) Born–Oppenheimer ab Initio QM/MM Molecular Dynamics Simulations of the Hydrolysis Reaction Catalyzed by Protein Arginine Deiminase 4. *J. Phys. Chem. B* 113, 16705–16710.
- (39) Liu, H., Zhang, Y., and Yang, W. (2000) How Is the Active Site of Enolase Organized To Catalyze Two Different Reaction Steps? *J. Am. Chem. Soc.* 122, 6560–6570.
- (40) Liu, J., Hamza, A., and Zhan, C.-G. (2009) Fundamental Reaction Mechanism and Free Energy Profile for (–)-Cocaine Hydrolysis Catalyzed by Cocaine Esterase. *J. Am. Chem. Soc.* 131, 11964–11975.
- (41) Liu, J., Zhang, Y., and Zhan, C.-G. (2009) Reaction Pathway and Free-Energy Barrier for Reactivation of Dimethylphosphoryl-Inhibited Human Acetylcholinesterase. *J. Phys. Chem. B* 113, 16226–16236.
- (42) Wang, L., Yu, X., Hu, P., Broyde, S., and Zhang, Y. (2007) A Water-Mediated and Substrate-Assisted Catalytic Mechanism for *Sulfolobus solfataricus* DNA Polymerase IV. *J. Am. Chem. Soc.* 129, 4731–4737.
- (43) Wang, S., Hu, P., and Zhang, Y. (2007) Ab Initio Quantum Mechanical/Molecular Mechanical Molecular Dynamics Simulation of Enzyme Catalysis: The Case of Histone Lysine Methyltransferase SET7/9. *J. Phys. Chem. B* 111, 3758–3764.
- (44) Wu, R., Hu, P., Wang, S., Cao, Z., and Zhang, Y. (2010) Flexibility of Catalytic Zinc Coordination in Thermolysin and HDAC8: A Born–Oppenheimer ab Initio QM/MM Molecular Dynamics Study. *J. Chem. Theory Comput.* 6, 337–343.
- (45) Wu, R., Wang, S., Zhou, N., Cao, Z., and Zhang, Y. (2010) A Proton-Shuttle Reaction Mechanism for Histone Deacetylase 8 and the Catalytic Role of Metal Ions. *J. Am. Chem. Soc.* 132, 9471–9479.
- (46) Xiao, C., and Zhang, Y. (2007) Catalytic Mechanism and Metal Specificity of Bacterial Peptide Deformylase: A Density Functional Theory QM/MM Study. *J. Phys. Chem. B* 111, 6229–6235.
- (47) Zhang, Y., Kua, J., and McCammon, J. A. (2002) Role of the Catalytic Triad and Oxanion Hole in Acetylcholinesterase Catalysis: An ab initio QM/MM Study. *J. Am. Chem. Soc.* 124, 10572–10577.
- (48) Zhang, Y., Kua, J., and McCammon, J. A. (2003) Influence of Structural Fluctuation on Enzyme Reaction Energy Barriers in Combined Quantum Mechanical/Molecular Mechanical Studies. *J. Phys. Chem. B* 107, 4459–4463.
- (49) Zheng, F., Yang, W., Ko, M. C., Liu, J., Cho, H., Gao, D., Tong, M., Tai, H. H., Woods, J. H., and Zhan, C. G. (2008) Most efficient cocaine hydrolase designed by virtual screening of transition states. *J. Am. Chem. Soc.* 130, 12148–12155.
- (50) Zhou, Y., Wang, S., and Zhang, Y. (2010) Catalytic Reaction Mechanism of Acetylcholinesterase Determined by Born–Oppenheimer Ab Initio QM/MM Molecular Dynamics Simulations. *J. Phys. Chem. B* 114, 8817–8825.
- (51) Cisneros, G. A., Liu, H., Zhang, Y., and Yang, W. (2003) Ab Initio QM/MM Study Shows There Is No General Acid in the Reaction Catalyzed by 4-Oxalocrotonate Tautomerase. *J. Am. Chem. Soc.* 125, 10384–10393.
- (52) Metanis, N., Brik, A., Dawson, P. E., and Keinan, E. (2004) Electrostatic Interactions Dominate the Catalytic Contribution of Arg39 in 4-Oxalocrotonate Tautomerase. *J. Am. Chem. Soc.* 126, 12726–12727.
- (53) Poyner, R. R., Larsen, T. M., Wong, S. W., and Reed, G. H. (2002) Functional and structural changes due to a serine to alanine mutation in the active-site flap of enolase. *Arch. Biochem. Biophys.* 401, 155–163.
- (54) Nicolet, Y., Lockridge, O., Masson, P., Fontecilla-Camps, J. C., and Nachon, F. (2003) Crystal structure of human butyrylcholinesterase and of its complexes with substrate and products. *J. Biol. Chem.* 278, 41141–41147.
- (55) Sali, A., and Blundell, T. L. (1990) Definition of general topological equivalence in protein structures. A procedure involving comparison of properties and relationships through simulated annealing and dynamic programming. *J. Mol. Biol.* 212, 403–428.
- (56) Sali, A., and Blundell, T. L. (1993) Comparative protein modelling by satisfaction of spatial restraints. *J. Mol. Biol.* 234, 779–815.
- (57) Case, D. A., Cheatham, T. E., Darden, T., Gohlke, H., Luo, R., Merz, K. M., Onufriev, A., Simmerling, C., Wang, B., and Woods, R. J. (2005) The Amber biomolecular simulation programs. *J. Comput. Chem.* 26, 1668–1688.
- (58) Case, D. A., Darden, T. A., Cheatham, T. E., III, Simmerling, C. L., Wang, J., Duke, R. E., Luo, R., Merz, K. M., Pearlman, D. A., Crowley, M., Walker, R. C., Zhang, W., Wang, B., Hayik, S., Roitberg, A., Seabra, G., Wong, K. F., Paesani, F., Wu, X., Brozell, S., Tsui, V., Gohlke, H., Yang, L., Tan, C., Mongan, J., Hornak, V., Cui, G., Beroza, P., Mathews, D. H., Schafmeister, C., Ross, W. S., and Kollman, P. A. (2006) *Amber 9*, University of California, San Francisco.
- (59) Hörnberg, A., Tunemalm, A.-K., and Ekström, F. (2007) Crystal Structures of Acetylcholinesterase in Complex with Organophosphorus Compounds Suggest that the Acyl Pocket Modulates the Aging Reaction by Precluding the Formation of the Trigonal Bipyramidal Transition State. *Biochemistry* 46, 4815–4825.
- (60) Jorgensen, W. L., Chandrasekhar, J., Madura, J. D., Impey, R. W., and Klein, M. L. (1983) Comparison of Simple Potential Functions for Simulating Liquid Water. *J. Chem. Phys.* 79, 926–935.
- (61) Miyamoto, S., and Kollman, P. A. (1992) Settle: An Analytical Version of the Shake and Rattle Algorithm for Rigid Water Models. *J. Comput. Chem.* 13, 952–962.
- (62) Ryckaert, J. P., Ciccotti, G., and Berendsen, H. J. C. (1977) Numerical-Integration of Cartesian Equations of Motion of a System with Constraints: Molecular-Dynamics of N-Alkanes. *J. Chem. Phys.* 23, 327–341.
- (63) Frisch, M. J., Trucks, G. W., Schlegel, H. B., Scuseria, G. E., Robb, M. A., Cheeseman, J. R., Montgomery, J. A., Jr., Vreven, T., Kudin, K. N., Burant, J. C., Millam, J. M., Iyengar, S. S., Tomasi, J., Barone, V., Mennucci, B., Cossi, M., Scalmani, G., Rega, N., Petersson, G. A., Nakatsuji, H., Hada, M., Ehara, M., Toyota, K., Fukuda, R., Hasegawa, J., Ishida, M., Nakajima, T., Honda, Y., Kitao, O., Nakai, H., Klene, M., Li, X., Knox, J. E., Hratchian, H. P., Cross, J. B., Bakken, V., Adamo, C., Jaramillo, J., Gomperts, R., Stratmann, R. E., Yazyev, O., Austin, A. J., Cammi, R., Pomelli, C., Ochterski, J. W., Ayala, P. Y., Morokuma, K., Voth, G. A., Salvador, P., Dannenberg, J. J., Zakrzewski, V. G., Dapprich, S., Daniels, A. D., Strain, M. C., Farkas, O., Malick, D. K., Rabuck, A. D., Raghavachari, K., Foresman, J. B., Ortiz, J. V., Cui,

Q., Baboul, A. G., Clifford, S., Cioslowski, J., Stefanov, B. B., Liu, G., Liashenko, A., Piskorz, P., Komaromi, I., Martin, R. L., Fox, D. J., Keith, T., Al-Laham, M. A., Peng, C. Y., Nanayakkara, A., Challacombe, M., Gill, P. M. W., Johnson, B., Chen, W., Wong, M. W., Gonzalez, C., and Pople, J. A. (2004) *Gaussian 03*, revision C.02, Gaussian, Inc., Wallingford, CT.

(64) Case, D. A., Darden, T. A., Cheatham, T. E., III, Simmerling, C. L., Wang, J., Duke, R. E., Luo, R., Merz, K. M., Wang, B., Pearlman, D. A., Crowley, M., Brozell, S., Tsui, V., Gohlke, H., Mongan, J., Hornak, V., Cui, G., Beroza, P., Schafmeister, C., Caldwell, J. W., Ross, W. S., and Kollman, P. A. (2004) *AMBER 8*, University of California, San Francisco.

(65) Nachon, F., Asojo, O. A., Borgstahl, G. E., Masson, P., and Lockridge, O. (2005) Role of water in aging of human butyrylcholinesterase inhibited by echothiophate: The crystal structure suggests two alternative mechanisms of aging. *Biochemistry* 44, 1154–1162.

(66) Carletti, E., Li, H., Li, B., Ekstrom, F., Nicolet, Y., Loiodice, M., Gillon, E., Froment, M. T., Lockridge, O., Schopfer, L. M., Masson, P., and Nachon, F. (2008) Aging of cholinesterases phosphorylated by tabun proceeds through O-dealkylation. *J. Am. Chem. Soc.* 130, 16011–16020.

(67) Carletti, E., Aurbek, N., Gillon, E., Loiodice, M., Nicolet, Y., Fontecilla-Camps, J. C., Masson, P., Thiermann, H., Nachon, F., and Worek, F. (2009) Structure-activity analysis of aging and reactivation of human butyrylcholinesterase inhibited by analogues of tabun. *Biochem. J.* 421, 97–106.

(68) Truhlar, D. G., and Garrett, B. C. (1984) Variational Transition State Theory. *Annu. Rev. Phys. Chem.* 35, 159–189.