

Structural and functional characterization of the interaction of the photosensitizing probe methylene blue with *Torpedo californica* acetylcholinesterase

Aviv Paz,^{1,2} Esther Roth,¹ Yacov Ashani,¹ Yechun Xu,^{1,2} Valery L. Shnyrov,³ Joel L. Sussman,² Israel Silman,¹ and Lev Weiner^{4*}

¹Department of Neurobiology, Weizmann Institute of Science, Rehovot 76100, Israel

²Department of Structural Biology, Weizmann Institute of Science, Rehovot 76100, Israel

³Department of Biochemistry and Molecular Biology, Universidad de Salamanca, Salamanca 37007, Spain

⁴Department of Chemical Research Support, Weizmann Institute of Science, Rehovot 76100, Israel

Received 18 March 2012; Revised 16 May 2012; Accepted 16 May 2012

DOI: 10.1002/pro.2101

Published online 1 June 2012 proteinscience.org

Abstract: The photosensitizer, methylene blue (MB), generates singlet oxygen that irreversibly inhibits *Torpedo californica* acetylcholinesterase (TcAChE). In the dark, it inhibits reversibly. Binding is accompanied by a bathochromic absorption shift, used to demonstrate displacement by other acetylcholinesterase inhibitors interacting with the catalytic “anionic” subsite (CAS), the peripheral “anionic” subsite (PAS), or bridging them. MB is a noncompetitive inhibitor of TcAChE, competing with reversible inhibitors directed at both “anionic” subsites, but a single site is involved in inhibition. MB also quenches TcAChE’s intrinsic fluorescence. It binds to TcAChE covalently inhibited by a small organophosphate (OP), but not an OP containing a bulky pyrene. Differential scanning calorimetry shows an ~8° increase in the denaturation temperature of the MB/TcAChE complex relative to native TcAChE, and a less than twofold increase in cooperativity of the transition. The crystal structure reveals a single MB stacked against Trp279 in the PAS, oriented down the gorge toward the CAS; it is plausible that irreversible inhibition is associated with photooxidation of this residue and others within the active-site gorge. The kinetic and spectroscopic data showing that inhibitors binding at the CAS can impede binding of MB are reconciled by docking studies showing that the conformation adopted by Phe330, midway down the gorge, in the MB/TcAChE crystal structure, precludes simultaneous binding of a second MB at the CAS. Conversely, binding of ligands at the CAS dislodges MB from its preferred locus at the PAS. The data presented demonstrate that TcAChE is a valuable model for understanding the molecular basis of local photooxidative damage.

Keywords: acetylcholinesterase; singlet oxygen; photosensitizer; methylene blue; tryptophan; radiation damage

Additional Supporting Information may be found in the online version of this article.

Aviv Paz’s current address is Department of Physiology, David Geffen School of Medicine at UCLA, Los Angeles, CA 90095, USA.

Yechun Xu’s present address is Drug Discovery and Design Center, Shanghai Institute of Materia Medica, Chinese Academy of Sciences, Shanghai 201203, China.

Grant sponsors: Benoziyo Center for Neuroscience and the Nalvyco Foundation; V.S. acknowledges travel support from the Kimmelman Center for the Study of Biomolecular Structure and Assembly

*Correspondence to: Lev Weiner, Department of Chemical Research Support, Weizmann Institute of Science, Rehovot 76100, Israel. E-mail: lev.weiner@weizmann.ac.il

Introduction

An important category of photosensitized compounds are those that generate reactive oxygen species (ROS).^{1,2} Light excitation of such compounds raises them to an excited state (singlet or triplet), which stimulates generation of the ROS by one or both of two mechanisms. Superoxide radical ($O_2^{\cdot-}$) and hydroxyl radical ($HO^{\cdot}$) are generated by electron transfer and singlet oxygen (1O_2) by energy transfer.³ ROS can produce damage to cells by various mechanisms, including lipid peroxidation,^{4,5} DNA degradation,^{6,7} and chemical modification of proteins.^{8–10} Photosensitized compounds are utilized in a clinical context for photodynamic therapy (PDT) of malignancies,¹¹ and are often targeted to specific “addresses” by conjugation with antibodies or peptide hormones.^{3,12,13} Photoactivated inhibitors can similarly serve as valuable tools for selective modification of enzymes and other proteins,¹⁴ both *in vitro* and *in situ*, by binding specifically to their active sites before photoactivation. For optimal utilization of photosensitizers in PDT, it is necessary to understand how they interact with their targets, whether nonspecifically or specifically, the pathway(s) taken by the reactive species generated by photoexcitation, the molecular targets, and the specific chemical damage produced. These issues are both under active investigation and discussion.¹⁵

We earlier showed that *Torpedo californica* acetylcholinesterase (*TcAChE*) provides a useful model for studying the effects of ROS. Thus, $HO^{\cdot}$ radicals generated by the Fenton reaction¹⁶ mimic the effects produced by oxidative stress, namely, peptide bond cleavage and conversion of the native *TcAChE* dimer to an inactive partially unfolded molten globule state¹⁷ with structural features typical of a molten globule.^{18,19} Furthermore, the photosensitive reagent, hypericin,²⁰ was shown to specifically target the molten globule state of *TcAChE*, and to crosslink it by generation of 1O_2 .^{21,22} In addition, we were able to identify specific disulfide bond cleavage by X-irradiation.²³ Thus, *TcAChE* has served as a valuable model protein for studying the molecular processes underlying radiation damage.

The photosensitizer, rose Bengal, as well as other xanthene dyes, are reversible inhibitors of acetylcholinesterase (AChE) that act as irreversible inhibitors under illumination.^{24,25} Another photosensitive dye, methylene blue (MB)²⁶ (see Scheme 1 for the structures of MB and of other compounds used in this study), was shown to act as a reversible inhibitor of bovine erythrocyte AChE²⁷ and *Electrophorus electricus* AChE,²⁸ as well as of human and horse serum butyrylcholinesterase.^{27–29} Ozer and Küçükiling reported that their kinetic studies of the inhibition of BChE by MB and related compounds revealed more than one binding site and cooperative binding, whereas the inhibition of AChE

was consistent with a single binding site, with no evidence for cooperativity.²⁸

Recently, we showed that MB acts as an irreversible inhibitor of *TcAChE* under illumination, due to the action of 1O_2 generated by its photosensitizing action.¹⁰ Plausible targets for this irreversible action effect are the tryptophan (Trp) residues within the active-site gorge, associated with either the anionic subsite of the catalytic anionic site (CAS), near the bottom of the active-site gorge, Trp84, or with the peripheral anionic site (PAS), near the entrance to the gorge, Trp279.³⁰ To further explore the mode of interaction of MB with *TcAChE* and the site(s) with which it interacts, we studied its reversible inhibition of the enzyme in the dark. In addition, we used a repertoire of steady state spectroscopic methods, differential scanning calorimetry (DSC), computer docking and X-ray crystallography to characterize the reversible MB/*TcAChE* complex. These data provide a valuable molecular platform for investigating the pathways and possible targets of 1O_2 generated *in situ*.

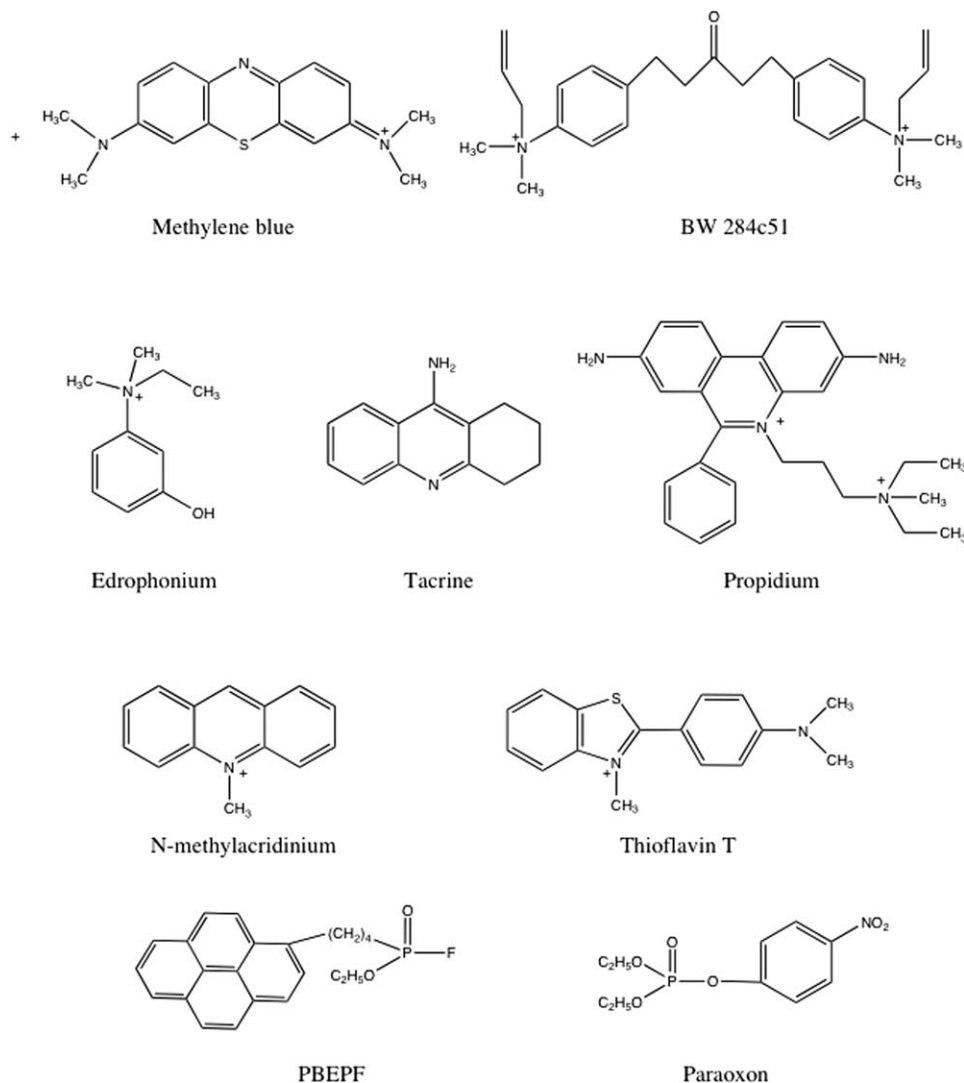
Results

Photodynamic inactivation of *TcAChE*

Figure 1(A) shows that *TcAChE* undergoes time- and concentration-dependent irreversible inactivation by MB under irradiation. Similar rates of photo-inactivation were obtained if white light was used or if illumination at wavelengths below 600 nm was cut off by a red-light filter (not shown). Detailed evidence demonstrating that this irreversible inactivation involves generation of singlet oxygen (1O_2) was presented earlier.¹⁰

As mentioned in the Introduction, MB has been shown to act as a reversible inhibitor of *Electrophorus* AChE.²⁸ It is, thus, plausible that it binds reversibly within the active site-gorge of *TcAChE* at the “anionic” subsite of the CAS, near the bottom of the active-site gorge, and/or at the PAS near the entrance to the gorge. This assumption was supported by the fact that various reversible inhibitors of *TcAChE*, known to bind at the CAS (edrophonium, EDR), at the PAS (propidium, PROP), or to span the active-site gorge, BW284c51 (BW), were shown to retard MB-stimulated photoinactivation [Fig. 1(B)]. Furthermore, kinetic studies recently showed that MB displays essentially complete competition with thioflavin (ThT) T,³¹ which has been shown to be bound at the PAS in its crystalline complex with *TcAChE*.³²

Accordingly, we decided to carefully study the reversible binding of MB to *TcAChE* by both steady-state spectroscopy and kinetics, under conditions in which irreversible photoinactivation was practically excluded (see Materials and Methods section).



Scheme 1.

Spectroscopic monitoring of the interaction of MB with *TcAChE*

Figure 2(A) shows the effect of increasing concentrations of *TcAChE* on the absorption spectrum of MB. The absorption maximum is shifted from 662 to 682 nm, with an isosbestic point at ~671 nm. Figure 2(B) displays a plot showing the dependence of the concentration of bound MB on the *TcAChE* concentration.

As mentioned above, binding of MB to *TcAChE* could be occurring at the CAS, at the PAS, or at both. We used our finding of the large optical shift (20 nm) between free and bound MB as a tool to monitor the displacement of MB by ligands whose binding-site within the active-site gorge of *TcAChE* had already been identified. Accordingly, we examined the influence on the spectrum of the MB/*TcAChE* complex of a CAS inhibitor, EDR,³⁰ of a PAS inhibitor, PROP,³³ and of a bifunctional inhibitor, BW, which spans the active-site gorge, binding at both the CAS and PAS simultaneously.³⁴

Figure 3(A,B) shows that neither a large excess of the CAS inhibitor alone, nor of the PAS inhibitor alone, completely reverses the change in the spectrum of MB produced by *TcAChE*, but that both together, or the gorge-spanning ligand, are capable of doing so [Fig. 3(C)].

Figure 4 shows an experiment in which, initially, *TcAChE* was titrated to saturation with MB. Upon subsequent titration with EDR, the ratio of the absorbance at 682 nm to that at 661 nm decreased from a maximum of ~1.45 to a plateau level of ~0.8, due to displacement of the MB from the active-site gorge of the enzyme by EDR. But upon addition of PROP, the ratio further decreased to ~0.5, approaching the value of 0.42 characteristic of free MB. In a control experiment, it was shown that if EDR was added first, MB that was added subsequently displaced the bound EDR.

The spectroscopic data thus suggest that MB can bind both to the PAS of *TcAChE* and, with much lower affinity, to the CAS. It is possible, however,

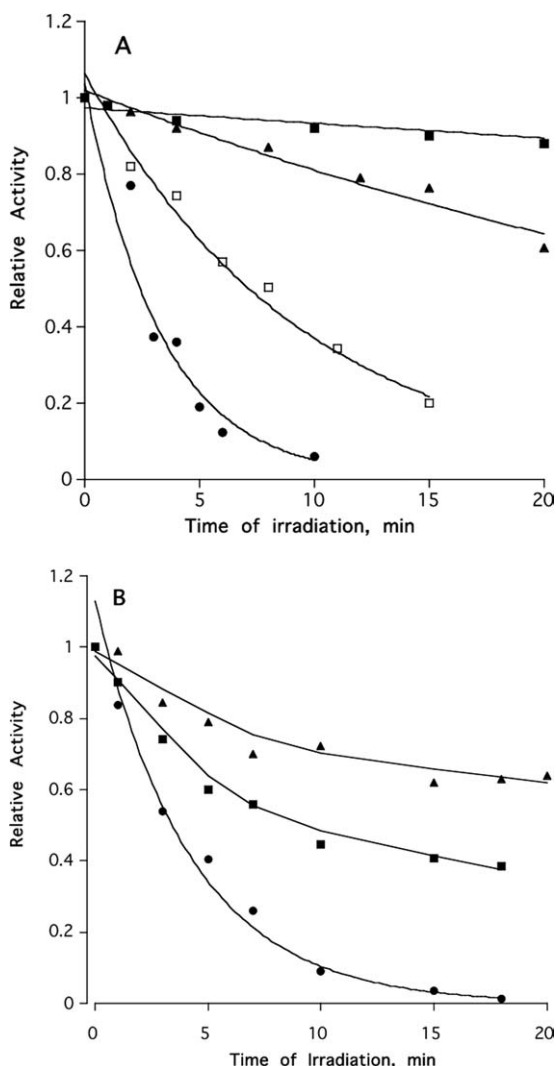


Figure 1. (A) Irreversible photoinactivation of TcAChE stimulated by MB. TcAChE samples were incubated in the presence of various concentrations of MB under illumination, as described under Materials and Methods, with aliquots being withdrawn for assay at the appropriate times. The TcAChE concentration was 2 μM in buffer 1, at 23°C. ▲, 0.2 μM MB; □, 5 μM MB; ●, 10 μM MB; ■, 5 μM MB in the absence of illumination. (B) Effects of reversible AChE inhibitors on the photoinactivation of TcAChE stimulated by MB. TcAChE, 1.5 μM ; MB, 1 μM ; EDR, 20 μM ; BW, 20 μM . ●, MB; ■, MB + EDR; ▲, MB + BW.

that the apparent weak binding in fact reflects an allosteric effect of its binding to the PAS.

Kinetics of reversible inhibition of TcAChE by MB

Kinetic data have been used extensively to investigate the interaction of inhibitors with the CAS and PAS of AChE.³⁵ Accordingly, we decided to complement the spectroscopic data presented above for the interaction of MB with TcAChE by kinetic measurements.

The reciprocal plots of $1/V$ versus $1/S$, in the presence of increasing fixed concentrations of MB per plot, show that the inhibitor does not change the K_m of acetylthiocholine iodide (ATC), but decreases $V_{\max}$ (Supporting Information Fig. S1). MB seems, therefore, to act as a classical noncompetitive inhibitor of TcAChE; thus, ATC maintains the same substrate affinity for the $[E][MB]$ complex as for the free enzyme, and MB binds to the $[E][S]$ complex with the same affinity as it binds to the free enzyme. $V_{\max}$ decreases with increasing MB concentration, most likely due to increasing formation of a nonproductive ternary $[E][MB][S]$ complex of the enzyme with ATC and MB. Similar values were obtained for K_I , 35.6 nM, and for αK_I , 33.1 nM, derived, respectively, from secondary plots of $1/V_{\max}$ versus $[MB]$, and from the slopes of the reciprocal plots of $1/V$

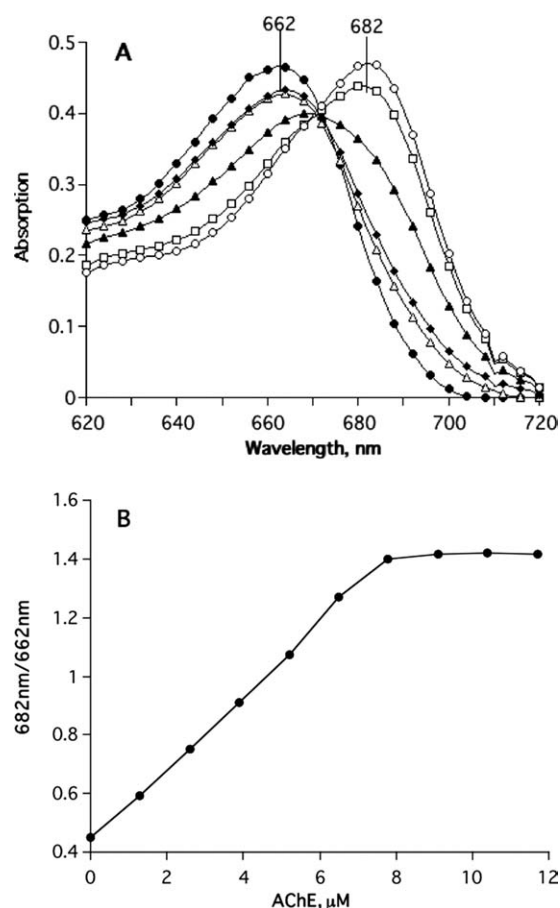


Figure 2. Binding of MB to TcAChE monitored by absorption spectroscopy. (A) The spectra shown monitor titration of 5- μM MB in buffer 1, with a concentrated solution of TcAChE in the same buffer. The values given represent the final concentrations of TcAChE; under the conditions used, the dilution of MB is negligible, as is evident from the sharp isosbestic point. ●, no TcAChE; △, 0.65 μM TcAChE; ◆, 1.3 μM TcAChE; ▲, 3.9 μM TcAChE; □, 6.5 μM TcAChE; ○, 9.2 μM TcAChE. (B) Saturation curve calculated using spectroscopic data similar to those presented in (A).

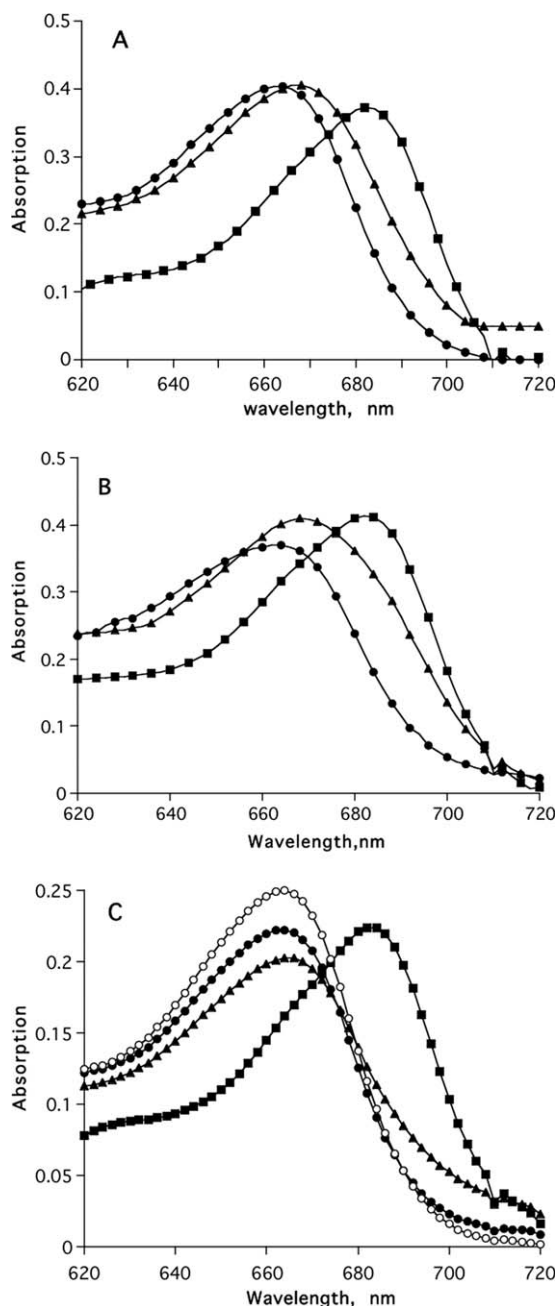


Figure 3. Effect of reversible AChE inhibitors on the binding of MB to *TcAChE* as monitored by absorption spectroscopy. (A) Displacement of MB by EDR. Concentrations were *TcAChE*, 12 μM ; MB, 5 μM ; EDR, 60 μM . $\bullet$, MB; $\blacksquare$, MB + *TcAChE*; $\blacktriangle$, MB + *TcAChE* + EDR. (B) Displacement of MB by PROP. Concentrations were *TcAChE*, 12 μM ; MB, 5 μM ; PROP, 60 μM . $\bullet$, MB; $\blacksquare$, MB + *TcAChE*; $\blacktriangle$, MB + *TcAChE* + PROP. (C) Displacement of MB by EDR+PROP or by BW. Concentrations were *TcAChE*, 5.2 μM ; MB, 2.5 μM ; EDR, 60 μM ; PROP, 60 μM ; BW, 40 μM . $\bullet$, MB; $\blacksquare$, MB + *TcAChE*; $\blacktriangle$, MB + *TcAChE* + EDR + PROP; $\circ$, MB + *TcAChE* + BW.

versus $1/S$ against $[\text{MB}]$. These observations are consistent with the equilibrium scheme shown for non-competitive inhibition in Scheme 2, where $\alpha \approx 1$.

The Lineweaver–Burk plots displayed in Figure 5(A,B)³⁶ show that EDR and PROP, respectively,

both compete with MB for binding to *TcAChE*, even though the former binds at the CAS, and latter at the PAS.

The number of MB binding sites involved in the inhibition of *TcAChE* was determined by plotting $\log [(V_0/V) - 1]$ versus $\log [\text{MB}]$ at a fixed concentration of the substrate, ATC, in accordance with the following equation,³⁶ that is analogous to the Hill equation for determining the number of substrate-binding sites³⁷:

$$\log[(V_0/V) - 1] = n \log [\text{MB}] - \log K \quad (1)$$

where V_0 and V are the velocities in absence and presence of MB, respectively, and the slope of the line, n , is the number of binding sites. Figure 6 shows that the plots are linear for a wide range of fixed ATC concentrations per plot, and for the increasing MB concentrations used. The average number of MB binding sites obtained on its basis, for an ATC concentration range of 0.02–1.5 mM, is 1.06 ± 0.07 (SD; $n = 6$). As the K_m of *TcAChE* for ATC, as determined by us, is ~ 0.08 mM, the level of occupancy of the enzyme by the substrate clearly does not influence the number of binding sites for MB that inhibits enzyme activity. The value obtained for the number of binding sites for MB in the *Torpedo* enzyme is in agreement with that reported earlier for electric eel AChE.²⁸

Fluorescence spectroscopy

MAC is a competitive inhibitor of AChE that binds at the catalytic site of the enzyme with concomitant concentration-dependent quenching of the fluorescence of both ligand and enzyme.^{38,39} Figure 7(A) shows the quenching of MAC fluorescence by increasing concentrations of *TcAChE*. Figure 7(B) shows that quenching of the fluorescence of MAC is reduced significantly by both MB and BW, proving that, as might be predicted, they can both displace MAC from the CAS. Figure 8(A) shows that MB itself can fully quench the intrinsic fluorescence of *TcAChE*. Figure 8(B) shows that BW also quenches the intrinsic fluorescence, but less efficiently than MB, because the quenching produced by 10 μM MB alone is reduced in the presence of BW.

Figure 9 displays the effect of two organophosphate (OP) inhibitors of AChE on the interaction of MB with *TcAChE*. The two OP inhibitors shown in Scheme 1, paraoxon and pyrenebutyl ethyl phosphorofluoridate (PBEPF),⁴⁰ both inhibit AChE irreversibly, by covalent reaction with its active-site serine, S200; but the pyrenebutyl ethyl phosphoryl moiety of the conjugate produced by PBEPF is much bulkier than the diethylphosphoryl moiety in the conjugate obtained by reaction with paraoxon. Addition of the diethylphosphoryl-*TcAChE* conjugate to MB produces a large “red” shift in its absorption

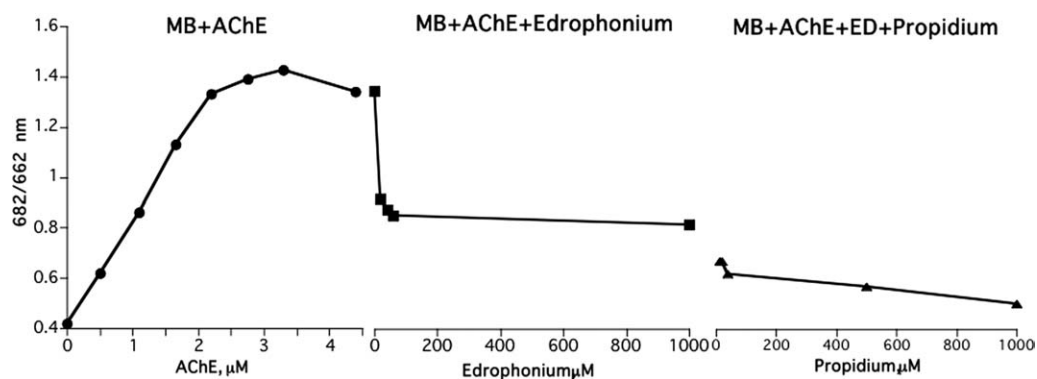


Figure 4. Sequential displacement of MB from *TcAChE* by EDR and PROP. A 5- μ M solution of MB in buffer 1 was titrated with a concentrated solution of *TcAChE* until a plateau value of the 682 nm/662 nm ratio was reached. Subsequent titration with concentrated EDR reduced the ratio to a lower plateau, and further titration with PROP reduced the ratio further.

maximum, to a value (~ 678 nm) approaching that produced by the free enzyme; in contrast, addition of the PBEP-*TcAChE* conjugate only slightly shifts the absorption maximum (~ 667 nm), suggesting a substantial decrease in the affinity of MB for the PBEPF/*TcAChE* conjugate.

The mamba venom polypeptide toxin, fasciculin II (FasII) is a very potent inhibitor of vertebrate AChEs.⁴¹ It exerts its action by binding tightly to the PAS, thus blocking entrance of substrate to the active-site gorge.^{42,43} Figure 10 shows that FasII completely prevents interaction of MB with *TcAChE*.

Thermal inactivation and differential scanning calorimetry of the MB/*TcAChE* complex

We earlier showed that thermal denaturation irreversibly transforms *TcAChE* into a molten globule.^{44,45} DSC permitted quantitative characterization of this transition,^{44,45} as well as of the thermal stabilization conferred by inhibitors, osmolytes, and divalent cations.^{22,46}

Figure 11 shows that MB, like other reversible AChE inhibitors,⁴⁶ provides effective protection of *TcAChE* against thermal inactivation.

Thermal denaturation of the MB/*TcAChE* complex, as of native *TcAChE*, gives rise to well-defined DSC transitions (Fig. 12). As we showed earlier for the native enzyme,⁴⁴ the value of the temperature of the excess heat capacity maximum (T_m) is dependent on the temperature scan rate (not shown). Thermal denaturation of *TcAChE*, under the experimental conditions used, is always calorimetrically irreversible, since on reheating of the same protein sample no thermal effect is observed. Such behavior is characteristic of an irreversible, kinetically controlled process.^{47,48} Accordingly, analysis of the DSC transitions under these conditions utilized a simple two-state irreversible model (see Materials and Methods). The results of the fitting of the experimental data to this model are displayed in Figure 12

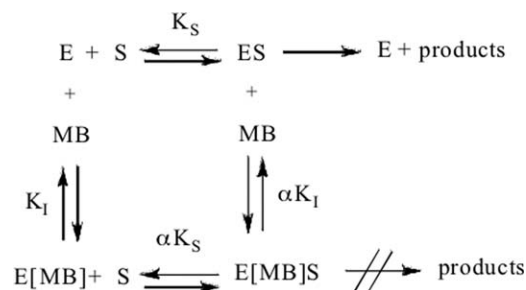
(solid lines) and in Table I. As can be seen, a good approximation was achieved.

It is clear that binding of MB to *TcAChE* substantially sharpens the cooperativity of the transition (more than twofold) as measured by the ratio $T_m/\Delta T$, where ΔT is the temperature range at the half-maximum of the DSC transition: namely, 6.1 for native *TcAChE* and 13.5 for the MB/*TcAChE* complex.

Crystal structure of the MB/*TcAChE* complex

The kinetic and spectroscopic data presented above showed that both ligands that bind at the catalytic site and those that bind at the PAS, as well as gorge-spanning ligands, can interfere with the binding of MB. We decided to use X-ray crystallography to identify the site of interaction of MB with the enzyme at the atomic level. Accordingly, MB was soaked into trigonal crystals of *TcAChE*, and data collection for the complex obtained, to 2.4 Å resolution, followed by solution of its structure by molecular replacement, and subsequent refinement, as described under Materials and Methods.

The crystal structure revealed a single molecule of MB, bound at the PAS, near the entrance of the active-site gorge (Fig. 13, upper panels). The MB molecule is surrounded by six aromatic residues at a distance of <4 Å: Tyr70, Tyr121, Tyr334, Trp279, Phe330, and Phe331, all six of which are among the



Scheme 2.

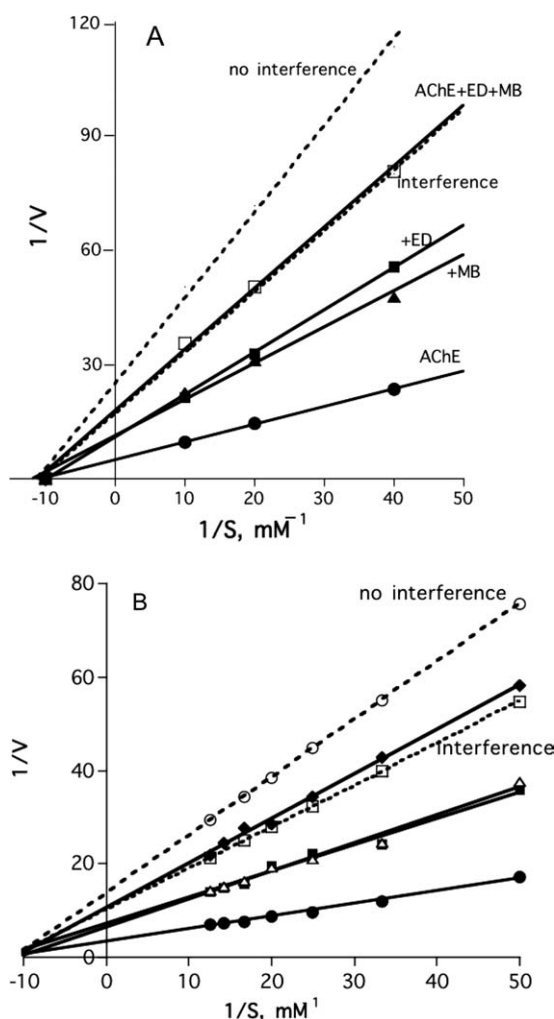


Figure 5. Lineweaver-Burk plots for hydrolysis of ATC by *TcAChE* in the presence and absence of reversible inhibitors. (A) Lineweaver-Burk plots demonstrating the interference of EDR with binding of MB. The dotted lines represent the theoretical plots for no interference of EDR with binding of MB and for interference, namely, complete competition for the same binding site. ●, *TcAChE* alone; ▲, *TcAChE* + 0.5 nM MB; ■, *TcAChE* + 5 nM EDR; □, *TcAChE* + 0.5 nM MB + 5 nM EDR. (B) Lineweaver-Burk plots demonstrating the interference of PROP with the binding of MB. The dotted lines represent the theoretical plots for no interference of PROP with binding of MB and for interference, namely, complete competition for the same binding site. ●, *TcAChE* alone; ■, *TcAChE* + 0.5 nM MB; △, *TcAChE* + 5 nM PROP; ◆, *TcAChE* + 0.5 nM MB + 5 nM PROP.

14 conserved aromatic residues which line the active-site gorge⁴⁹ (Fig. 13, lower left). In particular, MB forms a π - π stacking interaction with the indole ring of Trp 279, with a distance of 3.7 Å between the rings. Superposition of the MB/*TcAChE* structure on that of native *TcAChE* (PDB code 1EA5) shows that all the gorge main chains and side chains maintain their native conformation in the complex (data not shown).

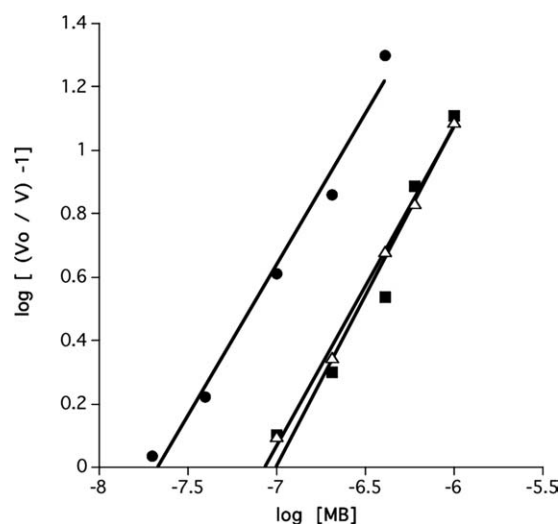


Figure 6. Inhibition of hydrolysis of ATC by *TcAChE* as a function of the MB concentration. Double logarithmic plots are displayed at three ATC concentrations. The slopes of these plots yield the number of binding sites for MB on the catalytic subunit involved in inhibition of enzymic activity.³⁶ ●, 0.1 mM ATC; △, 0.5 mM ATC; ■, 1.0 mM ATC.

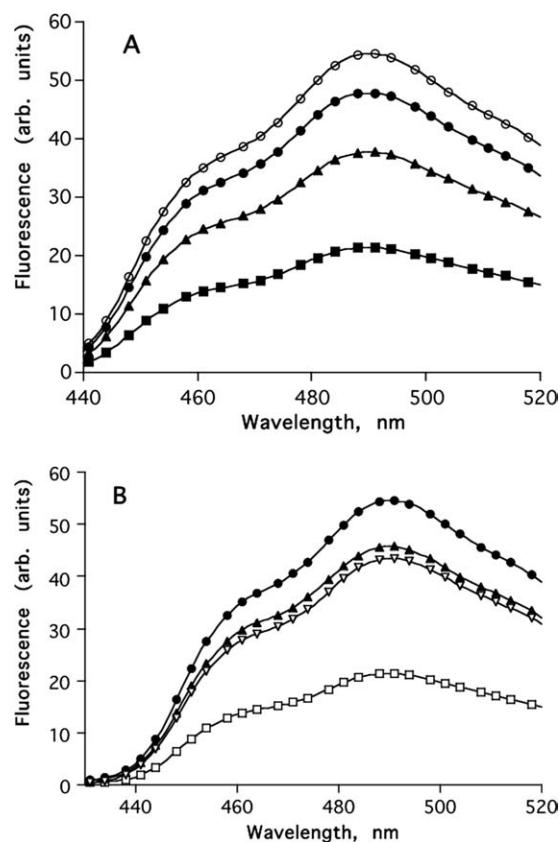


Figure 7. (A) Quenching of *N*-MAC fluorescence by *TcAChE*. ○, 20 μ M *N*-MAC; ●, 20 μ M *N*-MAC + 1.5 μ M *TcAChE*; ▲, + 3 μ M *TcAChE*; ■, + 6 μ M *TcAChE*; (B) Displacement of *N*-MAC from *TcAChE* by MB and by BW. ●, 20 μ M *N*-MAC; □, 20 μ M *N*-MAC + 6 μ M *TcAChE*; ▽, 20 μ M *N*-MAC + 6 μ M *TcAChE* + 0.200 μ M BW; ▲, 20 μ M *N*-MAC + 6 μ M *TcAChE* + 20 μ M MB.

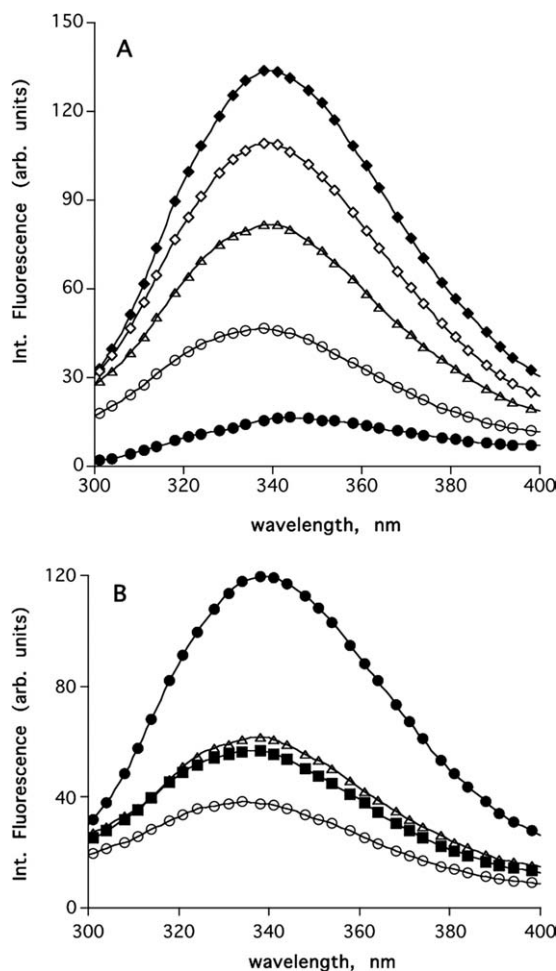


Figure 8. (A) Quenching of intrinsic fluorescence of *TcAChE* by MB. $\blacklozenge$, 1.5 μ M *TcAChE*; $\square$, 1.5 μ M *TcAChE* + 0.5 mM MB; $\triangle$, + 1 mM MB; $\circ$, + 10 mM MB; $\bullet$, + 10 mM MB. (B) Competition of MB and BW. $\bullet$, 1.5 μ M *TcAChE*; $\circ$, 1.5 μ M *TcAChE* + 20 μ M MB; $\triangle$, 1.5 μ M *TcAChE* + 100 μ M BW; $\blacksquare$, 1.5 μ M *TcAChE* + 20 μ M MB + 100 μ M BW.

The structure of the MB/*TcAChE* complex was compared to that of the ThT/*TcAChE* complex,³² in which a single molecule of ThT is similarly bound at the PAS. The lower right panel in Figure 13 shows that the ThT and MB molecules in their respective complexes are almost superimposed, as had earlier been predicted.³¹ These structures differ from that obtained for the complex with *TcAChE* of tacrine, whose chemical structure is quite similar to that of MB (see Scheme 1). In the tacrine/*TcAChE* complex, the single tacrine molecule bound is stacked against both Trp 84 and Phe330 at the CAS.^{30,46}

Thus, while the kinetic and spectroscopic data obtained in solution showed that both PAS and CAS inhibitors could prevent binding of MB to *TcAChE*, suggesting that it might bind at both the CAS and the PAS, the X-ray data revealed only a single MB molecule, bound at the PAS.

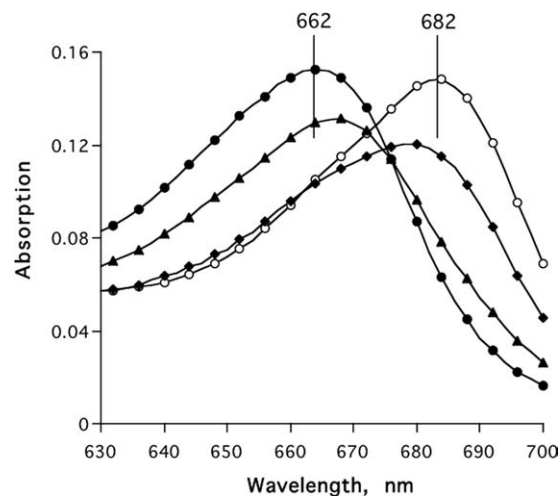


Figure 9. Effects of paraoxon and PBEPF on binding of MB to *TcAChE*. $\bullet$, 5 μ M MB; $\circ$, 5 μ M MB + 6 μ M *TcAChE*; $\blacklozenge$, 5 μ M MB + 6 μ M paraoxon/*TcAChE* conjugate; $\blacktriangle$, 5 μ M MB + 6 μ M PBEPF/*TcAChE* conjugate. The conjugates were prepared as described under Materials and Methods section.

Discussion

The use of photosensitizers for selective inactivation of targets in a clinical context is a rapidly developing area.¹⁴ In particular, PDT is providing novel therapeutic approaches to the treatment of malignancies.¹¹ We earlier showed that *TcAChE* serves as a valuable model system for studying the effects of irradiative damage by nonspecific generation of ROS, namely, OH radical¹⁶ and $^1\text{O}_2$.²¹ Recently, we reported that MB produces targeted inactivation of *TcAChE* under irradiation.¹⁰

In this study, we have shown that MB is a strong reversible inhibitor in the dark, and utilized an extensive arsenal of techniques to characterize its binding properties and to clarify its site(s) of interaction.

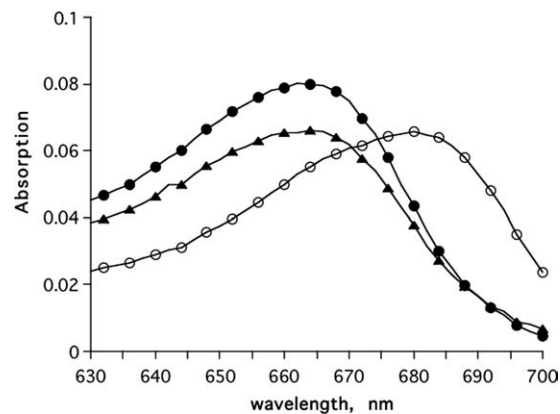


Figure 10. Effect of FasII on the binding of MB to *TcAChE*. $\bullet$, 1 μ M MB; $\circ$, 1 μ M MB + 2 μ M *TcAChE*; $\blacktriangle$, 1 μ M MB + 2 μ M *TcAChE* + 2 μ M FasII.

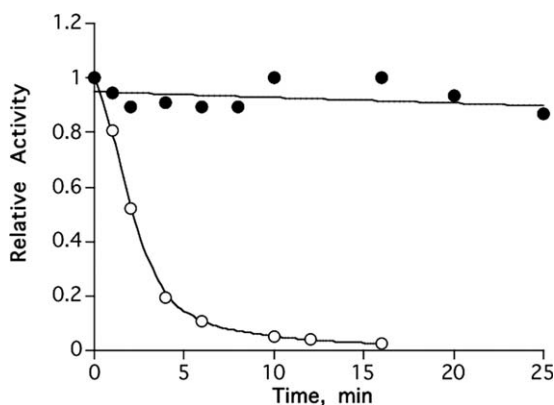


Figure 11. Protection of *TcAChE* from thermal inactivation by MB. ○, Incubation of 1.5 μM *TcAChE* at 39°C; ●, Incubation of 1.5 μM *TcAChE* + 50 μM MB at 39°C.

As expected, binding of MB to *TcAChE* substantially raised its thermal stability (Fig. 11), and DSC measurements showed an increase in the temperature of the thermal transition, accompanied by a substantial increase in cooperativity (Fig. 12). This effect is similar to that observed with other reversible inhibitors binding at either the CAS or the PAS or spanning the two.⁴⁶

Our main concern was to reconcile the data generated by the variety of experimental tools used, and to put forward a unified interpretation. As described under Results, binding of MB to *TcAChE* enzyme could be readily monitored spectroscopically, as it is accompanied by a 20-nm red shift in the absorption maximum of the probe [Fig. 2(A)]. This red shift could be reversed by displacement of the bound MB by a reversible inhibitor specific for the CAS, EDR, and by one specific for the PAS, PROP. However, neither inhibitor alone completely reversed the red shift. This could, however, be achieved by addition of the CAS and PAS inhibitors together, or by the potent gorge-spanning inhibitor, BW.³⁴ The spectroscopic experiments in which MB was titrated with *TcAChE* irreversibly inhibited by either paraoxon or PBEPF also support binding of MB at the PAS, as the paraoxon conjugate produced a large red shift in MB, whereas the pyrenebutyl conjugate had a much more limited effect (Fig. 9). It is plausible that the bulky pyrene moiety stacks against W279 and impedes association of the conjugate with MB.

Data showing that MB could displace ligands binding at the *TcAChE* active site were also obtained using the fluorescent probe, MAC [Fig. 7(A,B)]. Moreover, MB itself strongly quenched the intrinsic Trp fluorescence that is believed to be associated with the Trp residue at the active site,³⁹ which is now known to be Trp84 (*Torpedo* numbering).

The data obtained in the kinetic interference experiments in which MB competes with either EDR or with PROP on binding to *TcAChE* indicate that

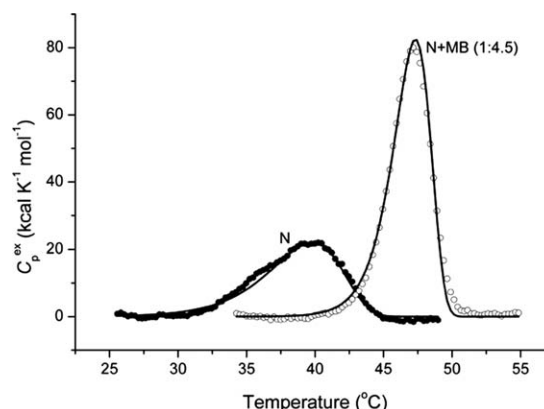


Figure 12. Temperature dependence of the excess molar heat capacity of *TcAChE* (N) and of the MB/*TcAChE* complex (N+MB) at a scan rate of 60 K h⁻¹ in 0.1M NaCl/10 mM HEPES, pH 7.5. Solid lines represent the best fit for each experimental curve to a simple two-state model of protein denaturation (see Materials and Methods section). The enzyme concentration was 6.9 μM , and the concentration of MB in the solution of the complex was 31 μM .

both CAS and PAS ligands affect the binding of MB to *TcAChE*. Thus, a mixture of either of these ligands with MB generated a Lineweaver–Burk plot that resembled the theoretical plot³⁶ calculated on the assumption of mutual interference between the ligand and MB, whether at the CAS or PAS.

In view of the 1:1 stoichiometric ratio established for the inhibition of *TcAChE* by MB (Fig. 7), simultaneous binding of the ligand at the CAS and PAS is unlikely to occur. Strong support is provided by the crystal structure of the MB/*TcAChE* complex (Fig. 13, upper and lower left panels), which clearly shows a single MB molecule bound at the PAS, stacked against W279, in a position very similar to that assumed by ThT in its corresponding *TcAChE* complex³² (Fig. 13, lower right panel). This complex is further stabilized by a cation- π interaction with F330 (Fig. 13, lower left panel). In this conformation, the F330 should also hinder the binding of a

Table I. Arrhenius Equation Parameter Estimates for *TcAChE* and for Its Complex with MB

Function	<i>TcAChE</i>	MB/ <i>TcAChE</i> (1:4.5)
ΔH (kcal mol ⁻¹)	544.4 $\pm$ 4.2	765.8 $\pm$ 5.2
E_A (kcal mol ⁻¹)	74.4 $\pm$ 1.6	146.8 $\pm$ 1.8
T^* (°C)	42.6 $\pm$ 0.2	47.9 $\pm$ 0.1
R	0.9992	0.9985

The correlation coefficient (r), used as a criterion for fitting accuracy, was calculated according to the equation,

$$r = \sqrt{\sum_{i=1}^n (y_i - y_i^{\text{calc}})^2 / \sum_{i=1}^n (y_i - y_i^{\text{m}})^2}$$
, where y_i and y_i^{calc} are, respectively, the experimental and calculated values of the measurable parameter; y_i^{m} is the mean of the experimental values of the measurable parameter and n the number of points.

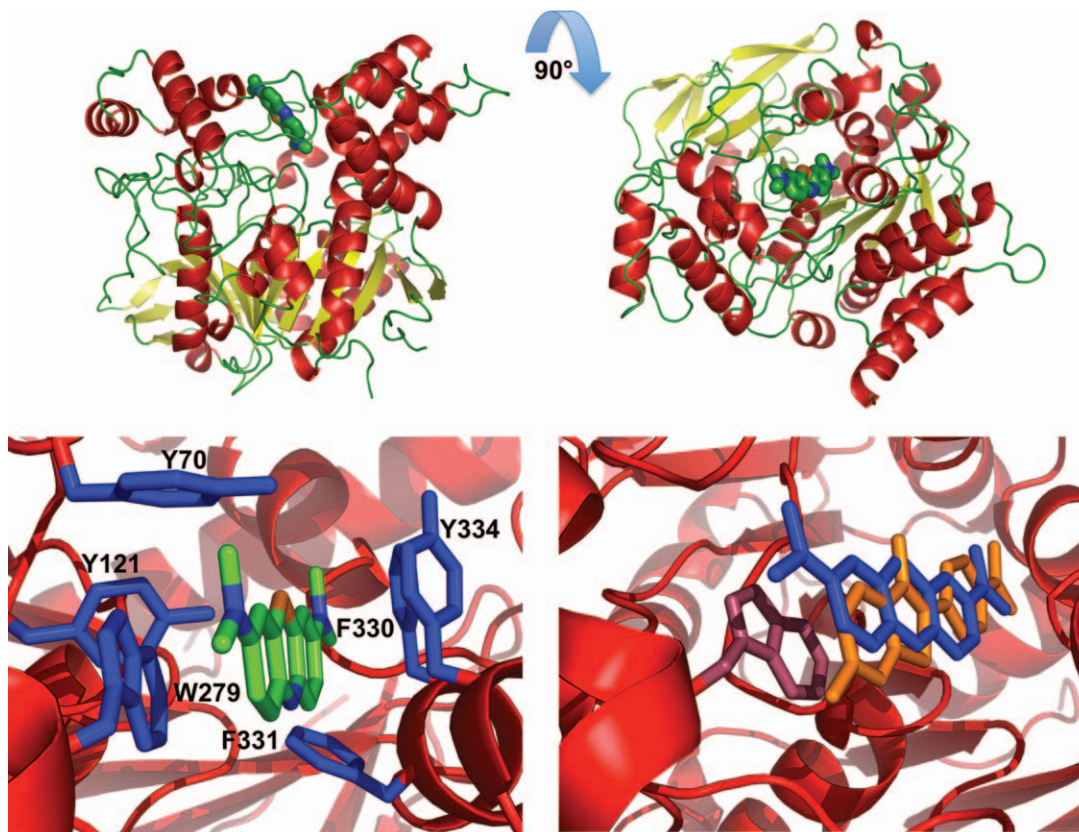


Figure 13. Crystal structure of the MB/*TcAChE* complex. Upper left: overall view of the complex. The *TcAChE* molecule is oriented with the entrance to the active-site gorge at the top. MB is seen in the PAS, where it is stacked against W279. MB carbon atoms are displayed in green, α -helices in red, β -sheets in yellow, and turns in green; upper right, same representation as upper left, but rotated by 90° around the x-axis so that the viewer is looking down the active-site gorge; lower left: view of the PAS in the MB/*TcAChE* complex. The enzyme is shown as a red ribbon, MB as a stick model with carbon atoms in green, nitrogen in blue and oxygen in red, and aromatic side chains as blue sticks. MB is within 4 Å of the six conserved aromatic residues displayed; lower right: superposition of the PAS regions of the MB/*TcAChE* and ThT/*TcAChE* complexes. MB is shown as blue sticks, ThT as orange sticks, and *TcAChE* in a red ribbon representation. The side chain of Trp279 is displayed in raspberry color. As the conformation of *TcAChE* in the crystal structures of the two complexes is practically identical, only that of the ThT/*TcAChE* structure (PDB code 2J3Q) is shown.

second MB at the CAS. Conversely, if, using AutoDock, MB is docked at the CAS, the F330 phenyl ring stacks against it in the model structure very similarly to F330 in the tacrine/*TcAChE* crystal structure (Fig. 14). Overall, the crystal structure of the MB/*TcAChE*, together with the docking data, do not seem to favor two MB molecules residing simultaneously within the active-site gorge of *TcAChE*, a conclusion consistent with the 1:1 stoichiometry of MB inhibition of enzymic activity.

In the EDR-*TcAChE* complex (2ACK.pdb), the phenyl group of F330 is not in close contact with either the quaternary nitrogen or the aromatic ring of EDR.³⁰ Nevertheless, it is rotated 90° relative to its orientation in the native crystal structure (1EA5.pdb). Furthermore, replacement of the homologous aromatic residue by Ala in both human and mouse AChE, that is, generation of the Y337A mutant, decreases the affinity for EDR 10–15-fold.^{50,51} Thus, in aqueous solution, F330 seems to influence

the binding of EDR. As MB stacks against F330, and this interaction most likely contributes to the stabilization of the MB-*TcAChE* complex, it is plausible that at a sufficiently high concentration, EDR will interact preferentially with the phenyl ring of F330 in the MB-*TcAChE* complex, thus displacing MB from its location as seen in the crystal structure. Yet EDR does not completely displace MB from the gorge, and it may be conjectured that MB remains stacked against the indole ring of W279 in an altered orientation, from which it can only be completely displaced by PROP. Conversely, PROP cannot completely remove the bound MB due to the latter's interaction with F330. Thus, under the experimental conditions utilized, and the ligand concentrations used, only a mixture of EDR and PROP, or a gorge-spanning ligand, such as BW, can achieve the complete removal of MB from the active-site gorge. The fluorescence data displayed in Figures 7 and 8, which show that *N*-MAC is displaced by MB, can

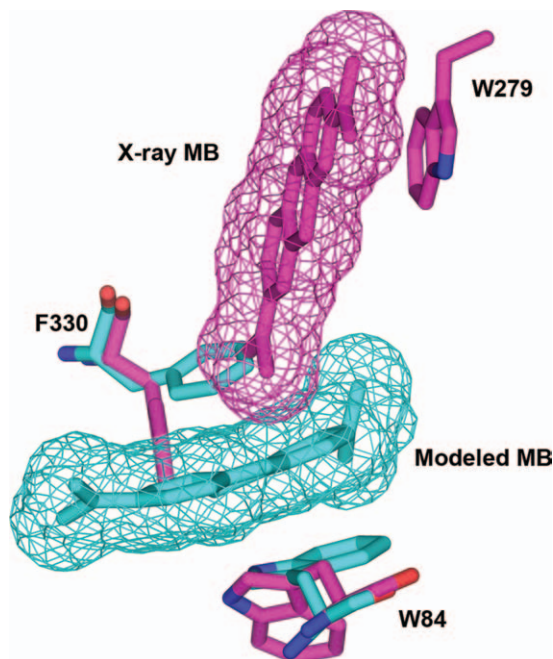


Figure 14. Overlay of the crystal structure of the MB/*TcAChE* complex and of a model structure in which MB is docked within the CAS. The overlay shows that the MB bound at the PAS (magenta) precludes simultaneous binding of a second MB at the CAS (cyan) due to its effect on the orientation of the phenyl group of F330.

also be explained in structural terms. *N*-MAC may be assumed to bind at the CAS sandwiched between W84 and F330,³⁹ analogously to tacrine.³⁰ The conformation adopted by the phenyl ring of F330 in the MB complex should thus impede binding of *N*-MAC at the CAS just as it inhibits binding of a second MB molecule.

In a cellular environment, it is well documented that $^1\text{O}_2$ reacts close to its site of generation.¹⁵ The data that we have presented here show that MB is acting within the active-site gorge of *TcAChE*, and suggest W279 as a likely target. It is, of course, important to know which other residues are photooxidized, and for this purpose, mass spectrometry provides a powerful tool.⁵²

In conclusion, we have presented kinetic, physicochemical, structural, and computational evidence that the photosensitizing agent, MB, which is a potent reversible and irreversible inhibitor of *TcAChE*, as well as of other ChEs, exerts its effect(s) by binding to the PAS of the enzyme.

Materials and Methods

Materials

TcAChE was the dimeric (G_2) glycosylphosphatidylinositol-anchored form from *T. californica* electric organ tissue, purified by affinity chromatography subsequent to solubilization with phosphatidylinositol-specific phospholipase C,⁵³ as described earlier.⁴⁶

Concentrations of stock solutions were determined by titration with paraoxon (*O,O*-diethyl-*O*-[4-nitrophenyl]phosphate).⁵⁴

MB was obtained from Merck KGaA (Darmstadt, Germany), and recrystallized from ethyl acetate. ATC, 5,5'-dithiobis-(2-nitrobenzoic acid) (DTNB), sodium azide, PEG 200, 2-(*N*-morpholino)ethanesulfonic acid (MES), and bovine serum albumin were purchased from Sigma (St. Louis, MO), as were the reversible AChE inhibitors, EDR, decamethonium bromide, tacrine hydrochloride, *d*-tubocurarine chloride, and BW284c51 (BW), and the irreversible OP inhibitor, paraoxon. Propidium ethiodide was obtained from Calbiochem (San Diego, CA). FasII was purchased from Alomone Laboratories (Jerusalem, Israel). 1-PBEPF was prepared according to Amitai and coworkers.⁴⁰ The chemical formulae of the principal probes used in this study are displayed in Scheme 1.

Determination of inhibition constants and number of binding sites

All inhibition studies of *TcAChE* by MB and other reversible ligands were performed in a 1-mL cuvette containing ~ 0.025 nM *TcAChE* in 0.1M NaCl/10 mM sodium phosphate, pH 7.0 (buffer 1), containing 0.01% gelatin, at 25°C. The inhibition constant, K_i , for inhibition of *TcAChE* by MB was determined by preincubating the enzyme with increasing concentrations of the ligand (10–90 nM) for 2 min in a dark compartment before measurement of residual enzymic activity. Measurement of activity was initiated by adding DTNB and ATC to final concentrations of 0.6 and 0.02–0.1 mM, respectively, in accordance with the protocol of Ellman and coworkers,⁵⁵ followed by monitoring the increase in absorbance at 412 nm for 1.2 min.

Preparation of conjugates of PBEPF and paraoxon with *TcAChE*

An aliquot of 6 μL of 5 mM PBEPF in dry dioxane was added to 0.5 mL of 30 μM *TcAChE* in buffer 1 at room temperature. After 5 min, an Ellman assay revealed >99% inhibition. The solution was passed over a 10-mL Biogel P-6 to separate the enzyme conjugate from free PBEPF and its hydrolysis products. The peak protein fractions were pooled, and concentrated in a Cetricon apparatus to a final concentration of 23 μM PBEP-*TcAChE*.

A diethylphosphoryl-*TcAChE* conjugate was prepared similarly by the addition of 10 μL of 2×10^{-5} M paraoxon in ethanol to 0.5 mL of 23 μM *TcAChE* in buffer 1. As for the PBEP-*TcAChE* conjugate, >99% inhibition was obtained within 5 min. Gel filtration, followed by concentration of the pooled protein fractions, were also performed similarly, to yield a stock solution of 15- μM diethylphosphoryl-*TcAChE*.

Examination of competition between MB and other reversible inhibitors for binding to TcAChE

The competition between MB and two classical cationic ligands, EDR, which binds specifically at the CAS, located deep in the catalytic gorge, and PROP, a ligand specific for the peripheral anionic-site (PAS), was examined by comparing the Lineweaver–Burk plots of AChE activity versus ATC concentration under the following experimental conditions: (a) in the absence of inhibitor; (b) in the presence of a fixed concentration of MB; (c) in presence of a fixed concentration of the CAS or PAS ligand; (d) in presence of a mixture of the same concentrations of MB and of the same CAS and PAS ligands as used in (b) and (c). The Lineweaver–Burk plots obtained were compared with two theoretically distinguishable lines suggested by Krupka³⁶ for two cases: (a) MB competes with the second ligand for the same binding site (interference); (b) Simultaneous binding of MB and the second ligand does not involve any steric overlap (no interference).

Photooxidation

Irreversible inactivation of TcAChE by photooxidation in the presence of MB was performed as described.¹⁰

Steady state spectroscopy

Absorption spectroscopy utilized a Uvikon 940 spectrophotometer (Kontron AG, Munich, Germany), and fluorescence measurements were performed on a Jasco FP-750 spectrofluorometer (Jasco, Easton, MD). All spectroscopic measurements were performed in buffer 1 at room temperature.

Differential scanning calorimetry

Calorimetric experiments utilized a MicroCal MC-2D differential scanning microcalorimeter (MicroCal, Northampton, MA) as described previously.²² All solutions were degassed by stirring under vacuum before scanning, and an overpressure of 2 atm of dry nitrogen was maintained over the liquid within the cell throughout the scan. The reversibility of the thermal transitions was checked by performing a second scan on the same sample, immediately after it had been cooled to room temperature. The molar excess heat capacity curves were smoothed and plotted using the Windows-based ORIGIN software package supplied by MicroCal (Northampton, MA), as described previously.^{46,56}

In our earlier study,⁴⁴ we showed that thermal denaturation of TcAChE could only be adequately described by a two-state model, $N \xrightarrow{k} D$, which considers only two significantly populated macroscopic states, the initial or native state (*N*) and the final or denatured (*D*) state, where *k* is a first-order kinetic

Table II. Data Collection and Refinement Statistics for the MB/TcAChE Complex

Data collection	
Space group	P3 ₁ 21
Cell parameters (Å)	<i>a</i> = 111.1 <i>b</i> = 111.4 <i>c</i> = 137.1
Molecules (a.u.)	1
Resolution (Å)	30–2.43
Total reflections	848,294
Unique reflections	35,368
Completeness (%) ^a	99.9 (97.2)
Avg. <i>I</i> /σ(<i>I</i>)	34.8 (5.55)
<i>R</i> _{merge} (%) ^b	5.6
Refinement statistics	
<i>R</i> (%) ^c	19.96
<i>R</i> _{free} (%)	24.54 (5.2% of data)
RMSD, bond lengths (Å)	0.019
RMSD, bond angles (°)	1.98
	PDB code 2w9i

^a Values in parentheses relate to the highest resolution shell, 2.52–2.43 Å.

^b $R_{\text{merge}} = \sum |I - \langle I \rangle| / \sum I$, where *I* is the observed intensity, and $\langle I \rangle$ is the average intensity obtained from multiple observations of symmetry-related reflections after rejections.

^c $R = \sum ||F_o| - |F_c|| / \sum |F_o|$, where *F*_o and *F*_c are the observed and calculated structure factors, respectively.

constant that changes with temperature, as given by the Arrhenius equation: $k = \exp[E_A(1/T^* - 1/T)/R]$, where *E*_A is the energy of activation, and *T*^{*} is the temperature at which the rate constant equals 1 min^{−1} (for details see Ref. 47).

X-Ray crystallography

Trigonal crystals of TcAChE⁴⁹ were soaked for 20 h at 4°C in 2 μL of 2.5 mM MB dissolved in the crystallization solution [46% PEG 200 (v/v)/100 mM MES, pH 6.5], using the hanging drop procedure.⁵⁷ The crystals were then transferred to cryoprotectant oil and flash cooled in a stream of liquid nitrogen, using a Oxford Cryosystems apparatus.

Data collection was performed “in-house,” using a RIGAKU RU-H3R running at 50 kV/100 mA with Osmic blue confocal optics at 120 K. Diffraction images were recorded on a RAXIS-IV++. DENZO and SCALEPACK were used to integrate and scale the data.⁵⁸ Data were truncated with the CCP4 program TRUNCATE,⁵⁹ and 5.2% of the reflections were randomly used as test reflections.

A model of MB was constructed using the GaussView 3.09 program (Gaussian, Carnegie, PA), and converted to a pdb file format using Babel.⁶⁰

Rigid body refinement in CCP4⁶¹ was based on a previously solved trigonal crystal form of native TcAChE (PDB code 1EA5), excluding water molecules and carbohydrates. Initial 2*F*_o−*F*_c and *F*_o−*F*_c electron density maps were calculated using 30–2.4 Å data, and the initial *F*_o−*F*_c map was used, with the aid of the program Coot,⁶² to fit MB into positive density at the entrance to the active-site gorge of

the enzyme, and to add 101 water molecules. Subsequent rounds of restrained refinement with overall *B*-factor refinement were performed, and 56 carbohydrate atoms were added, until convergence to values of $R_{\text{work}} = 19.96\%$ and $R_{\text{free}} = 24.54\%$ (R_{free} was as defined by Brunger⁶³). As analyzed by Molprobit, ⁶⁴ 96.0% of all residues are in favored regions, and 99.6% in allowed regions of the Ramachandran plot, the only outliers being Asp380 and Asn457, which are both glycosylated surface residues. Table II displays the data collection and refinement statistics. The crystal structure of the MB/*TcAChE* complex has been deposited in the PDB with access code 2W9I.

Computer docking and MD simulations

MB was docked into the protein conformation extracted from the X-ray structure of the tacrine/*TcAChE* complex (pdb code 1ACJ)³⁰ by use of the AutoDock program.^{65,66} The docked MB overlaid well at the location of tacrine in the complex. As the conformation of the protein during the docking process is fixed, and MB is larger than tacrine, an MD simulation was performed based on the docked MB/*TcAChE* complex. The complex, together with the crystal water molecules included in the 1ACJ structure were inserted into a box of dimensions $10.6 \times 10.6 \times 10.6 \text{ nm}^3$, with the minimal distance of the protein from its walls being 2.0 nm. This box was solvated by use of a simple point-charge water model,⁶⁷ and the resulting solvated box then submitted to energy minimization. Counterions were subsequently added, to provide a neutral simulation system. Energy minimization was then repeated on the whole system. After convergence had been reached, the solvent, the counterions and the protein, as well as MB, were coupled separately to a temperature bath at 300 K. The whole system was then equilibrated for 10 ns. MD simulations were carried out with the GROMACS package^{68,69} as described earlier.⁷⁰

Acknowledgments

JLS is the Morton and Gladys Pickman Professor of structural biology.

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