

Document downloaded from:

<http://hdl.handle.net/10251/88053>

This paper must be cited as:

Bosca Mayans, F.; Sastre Navarro, G.; Andreu, J.M.; Jornet, D.; Tormos Faus, R.E.; Miranda Alonso, M.Á. (2015). Drug-tubulin interactions interrogated by transient absorption spectroscopy. *RSC Advances*. 5(61):49451-49458. doi:10.1039/C5RA05636E.



The final publication is available at

<http://doi.org/10.1039/c5ra05636e>

Copyright Royal Society of Chemistry

Additional Information

RSC Advances



This is an *Accepted Manuscript*, which has been through the Royal Society of Chemistry peer review process and has been accepted for publication.

Accepted Manuscripts are published online shortly after acceptance, before technical editing, formatting and proof reading. Using this free service, authors can make their results available to the community, in citable form, before we publish the edited article. This *Accepted Manuscript* will be replaced by the edited, formatted and paginated article as soon as this is available.

You can find more information about *Accepted Manuscripts* in the [Information for Authors](#).

Please note that technical editing may introduce minor changes to the text and/or graphics, which may alter content. The journal's standard [Terms & Conditions](#) and the [Ethical guidelines](#) still apply. In no event shall the Royal Society of Chemistry be held responsible for any errors or omissions in this *Accepted Manuscript* or any consequences arising from the use of any information it contains.

Drug-tubulin interactions interrogated by transient absorption spectroscopy

F. Boscá^a, G. Sastre^a, J. M. Andreu^b, D. Jornet^a, R. Tormos^{a*} and M. A. Miranda^{a*}

Received 00th January 20xx,
Accepted 00th January 20xx

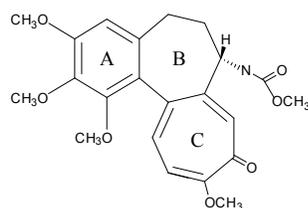
DOI: 10.1039/x0xx00000x

www.rsc.org/

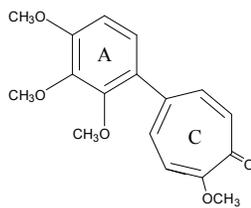
Colchicine (COL) is a bioactive molecule with antitumor properties. When COL binds to tubulin (TU), it inhibits microtubule assembly dynamics. We have investigated COL-TU interactions using laser flash photolysis (LFP) technique and performing fully flexible molecular dynamics simulations. Excitation of COL at 355 nm in aqueous medium did not lead to any transient absorption spectrum. By contrast, in the presence of TU a transient peaking at λ_{max} ca. 420 nm was registered and assigned as triplet excited COL complexed with TU ($^3\text{COL}^*\text{@TU}$). In aerated medium, the lifetime was τ ca 160 μs and the quantum yield was 0.138. Likewise, when the bicyclic COL analog MTC was submitted to LFP in the presence of TU, $^3\text{MTC@TU}^*$ was detected with a lifetime of ca. 62 μs and a quantum yield of 0.296. Aqueous solutions of MTC did not produce any signal in the microsecond timescale. The triplet energy of MTC was obtained by means of emission measurements and found to be ca. 200 kJ mol^{-1} , a value that matches with that previously reported for COL (188 kJ mol^{-1}). Molecular dynamic simulations, both with the ground and triplet excited state, reveal a strong interaction between COL and TU to give stabilized complexes with restricted mobility inside the protein binding site. These results demonstrate that LFP is a useful methodology to study the binding of COL derivatives to TU and open a new way to evaluate the interactions of non-fluorescent anticancer drugs with this protein.

Introduction

Tubulin (TU), the major component of microtubules, is a heterodimeric protein formed by α and β subunits that plays a crucial role in biological processes such as mitosis, intracellular transport or cell growth [1-4]. This protein is the target for anticancer drugs, which inhibit or promote its assembly into microtubules [5-10].



COL



MTC

Chart 1 Colchicine (COL) and 2-methoxy-5-(2',3',4'-trimethoxyphenyl)-2,4,6-cycloheptatrien-1-one (MTC) molecular structures

Colchicine (COL, Chart 1) is a natural alkaloid obtained from meadow saffron (*Colchicum autumnale*) [11]. It is an important bioactive drug as well as a neurotoxin in animal models of Alzheimer's disease and epilepsy [12-14]. Although COL has been shown to possess antitumor properties [15,16], its therapeutic use is limited by toxicity problems; this has led to the consideration of analogs [17-19]. When COL binds to TU, it inhibits the assembly into

microtubules and microtubule dynamics [20]. The binding process is slow and strongly temperature-dependent [21], while dissociation is kinetically unfavorable due to its high activation energy [22]. The binding site is mostly buried in the intermediate domain of the β subunit [6].

The structure of COL consists of three rings, the trimethoxybenzene ring (A), the methoxytropone (C) and the seven-membered ring with an acetamido substitution at C-7 (B) [23-24]. Interaction of COL with TU has been attributed to the simultaneous binding of its A and C rings. A properly positioned oxygen atom in ring C is essential feature for powerful inhibition of microtubule assembly [25-27]. The middle connecting B ring, which interacts with the α subunit, is involved in the peculiar binding kinetics [23], but it is not an essential requirement for TU binding. Thus, the MTC analog (Chart 1) lacking the B ring binds rapidly and reversibly to the high affinity COL binding site of TU, thereby inhibiting microtubule assembly [28-30].

The toxicity of COL has promoted computational studies in an attempt to find new TU binding agents with improved properties [6,31-33]. Interestingly, using a combined computational and cytotoxicity study, the binding energies of several COL derivatives have been ranked according to systematic modifications of rings A-C and correlated with cytotoxicity [32].

In general, protein binding parameters can be determined by a variety of techniques [34-41]. Fluorescence has been found to be one of the most convenient methods, due to the dramatic enhancement of the emission quantum yield of COL upon TU binding [42]. More recently, laser flash photolysis (LFP) has been introduced as a new tool to investigate drug-protein interactions

and to determine binding parameters such as affinity constants or population of the binding sites [43, 44]. Interestingly, the absence of a triplet excited state detectable in LFP has revealed that COL does not bind to serum albumins [45]. This technique is very sensitive and does not require separation of free and complexed drug. However, it has not yet been applied to TU binding studies.

With this background, the aim of the present work is to investigate COL-TU and MTC-TU interactions by means of a combined experimental and theoretical approach, using in parallel LFP and fully flexible molecular dynamics simulations. The obtained results confirm that the LFP methodology is indeed a powerful tool for interrogating the TU binding behavior of COL and its derivatives.

Experimental

General. COL was purchased from Sigma-Aldrich Chemical Company. MTC was a kind gift from T.J. Fitzgerald, Florida A&M University [46]. TU was purified from bovine brain and stored as described [47]. Before use, concentrated TU was diluted more than 50-fold into PG buffer to give the desired final concentration and employed within the next 4 h. PG buffer: 10 mM sodium phosphate and 0.1 mM GTP at pH 7.0.

Absorption and emission spectra. Optical spectra in different media were measured on a Perkin-Elmer Lambda 35 UV/vis spectrophotometer. Phosphorescence measurements were recorded with a time-resolved spectrometer (TimeMaster fluorescence lifetime spectrometer TM-2/2003) from Photon Technology International. The spectra of the samples were recorded at 77 K in acetonitrile and in 1mM PB aqueous solution using a single-cell Peltier cooler. The emission spectral band widths were set to 10 nm, and the excitations were performed at 350 nm. The phosphorescence emission spectra were recorded with a delay time of 0.5 ms and a total gate time of 10 ms.

Laser flash photolysis experiments. A pulsed Nd:YAG laser was used for the excitation at 355 nm. The single pulses were ~10 ns duration, and the energy was from 10 to 1 mJ pulse⁻¹. A pulsed xenon lamp was employed as detecting light source. The LFP apparatus consisted of the pulsed laser, the Xe lamp, a monochromator and a photomultiplier made up of a tube, housing and power supply. The output signal from the oscilloscope was transferred to a personal computer.

The laser pulse was probed by a system containing a fiber that synchronizes the LFP with the digitizer operating in the pretrigger mode. All transient spectra were recorded using 10 x 10 mm² quartz cells with 4 mL capacity, and they were bubbled during 20 min with N₂. All the experiments were carried out at room temperature. The solutions of COL and MTC were prepared at 5 x 10⁻⁵ M concentration.

Experiments with COL and MTC in buffered aqueous media in absence and presence of tubulin. Aqueous solutions of 5 x 10⁻⁵ M COL and MTC were prepared in PG buffer at 25 C, with and without the presence of TU under aerobic and anaerobic conditions. The samples containing protein needed special manipulation, avoiding bubbling the protein solutions to remove oxygen. Thus, N₂O and N₂ were introduced inside the sample quartz cells flowing the gas during 20 min without generating foam and stirring the solution. Transient absorption spectra at different times after the laser pulse

were obtained for each sample in the absence and the presence of TU (0.5 x 10⁻⁵ up to 1 x 10⁻⁵ M in the case of COL and only 1 x 10⁻⁵ M using MTC). The absorptions were registered twice (with two different samples) and the results include the average.

MTC triplet excited state measurements. The molar absorption coefficient of MTC triplet state (³MTC*) in acetonitrile was estimated by monitoring the energy transfer reaction between ³MTC* and β-carotene ground state (β-Car). As the intersystem crossing quantum yield (Φ_{ISC}) of β-Car is exceedingly small, ³β-Car* can only be populated via energy transfer (equation 1). Indeed, ³MTC* can act as energy donor because of the energy of ³β-Car* (ca. 79 kJ mol⁻¹) [48].



Measurements were performed in deaerated acetonitrile solutions of MTC with and without the presence of β-Car (1 x 10⁻⁵ - 5 x 10⁻⁵ M). Then, the molar absorption coefficient (ε) of ³MTC* was calculated using equation 2:

$$k_2/(k_2 - k_1) \times \Delta A({}^3\beta\text{-Car}^* (520 \text{ nm})) \times \epsilon({}^3\text{MTC}^* (400 \text{ nm})) = \Delta A({}^3\text{MTC}^* (400 \text{ nm})) \times \epsilon({}^3\beta\text{-Car}^* (520 \text{ nm})) \quad (2)$$

where the ΔA values refer to the absorbance at 400 nm of ³MTC* at the beginning of the reaction and ³β-Car* (at 520 nm) at the end of the reaction, k₁ is the ³MTC* decay rate constant without β-Car, and k₂ is the ³MTC* decay rate constants obtained at different concentrations of β-Car. The molar absorption coefficient of ³β-Car* in acetonitrile at 520 nm was taken as 100000 M⁻¹ cm⁻¹ (as described in toluene) [49].

The intersystem crossing quantum yield (Φ_{ISC}) of MTC was obtained by the comparative method [50] assuming that ε(³MTC*) is similar in all solvents. Hence, excitation of benzophenone (BP) and MTC was carried out separately using solutions with identical absorbance at the excitation wavelength (0.3 at 355 nm). Then equation 3 was applied:

$$\Phi_{\text{ISC}}(\text{MTC}) = \Phi_{\text{ISC}}(\text{BP}) \times \Delta A({}^3\text{MTC}^* (400 \text{ nm})) \times \epsilon({}^3\text{BP}^* (525 \text{ nm})) / \Delta A({}^3\text{BP}^* (525 \text{ nm})) \times \epsilon({}^3\text{MTC}^* (400 \text{ nm})) \quad (3)$$

where the ΔA values refer to the absorbance of ³MTC* at 400 nm and ³BP* at 525 nm. The benzophenone triplet molar absorption coefficient (ε(³BP*(525 nm))) and its triplet state quantum yield (Φ_{ISC}(BP)) in acetonitrile were taken to be 6500 M⁻¹ cm⁻¹ and 1, respectively [51].

The ³MTC* quenching rate constants by oxygen and ³β-Car were determined using the Stern-Volmer equation 4:

$$1/\tau = 1/\tau_0 + k [\text{Quencher}] \quad (4)$$

Computational methods. With the aim of unveiling the dynamic behavior of the system COL-TU, molecular dynamics runs were performed using LAMMPS [52] (Large-scale Atomic/Molecular Massively Parallel Simulator) software, which integrates Newton's equations of motion for all the atoms of the system, that interact via short- and long-range forces with an input consisting on the

initial location of all the atoms. The TU geometry was taken from Ravelli et al. [6], where the TU structure is solved at 3.5 Å resolution in complex with COL. Starting from this initial geometry for the complex, 1000 water molecules were added and treated explicitly in the simulation using a known method [53], where the advantages of explicit versus continuum solvent models have been optimized with parallel techniques for computational efficiency. The molecular dynamics simulations have been performed within the NVT ensemble at 298 K and 1 atm for 1 ns with the explicit relaxation of all the atoms of the system using a timestep of 1 fs. The system was initially energy minimized at 10 K using the following algorithms: steepest-descent (50000 steps) and conjugated-gradients (100000 steps). Then, three subsequent dynamic cycles of 10000 steps (at 98, 198, and 298 K) were performed for equilibration. Finally, 1 million steps of dynamics at 298 K allowed obtaining 1 ns of molecular dynamics behavior.

Periodic boundary conditions were implemented within a rectangular box of [230,200,120] Å in [x,y,z]. The force field employed was UFF [54], including an electrostatic part obtained from a MOPAC2012 [55,56] calculation using the minimized system as single point. For the short-range and long-range parts, a method with Lennard-Jones and Coulombics computed via damped shifted forces with a damping parameter 0.05 Å⁻¹ and cutoffs of 10.0 Å (Lennard-Jones) and 12.0 Å (Coulombic) was employed.

Diffusion of the triplet excited state within TU has been considered by means of a two-steps strategy. Thus, the triplet geometry of COL and MTC has been calculated using first principles based on density functional theory (DFT). The geometries, as well as the corresponding charge distributions, have been compared with those of the ground states. Then, a reparameterization of UFF has been made so that the specific triplet geometries can be reproduced within this force field and the modified UFF has been employed to perform a new molecular dynamics run.

The first-principles geometry optimizations have also been performed using the PBE and ωB97X-D [57] functionals, with TZVP [58] basis set, obtaining very similar results. For calculation of excitation energies by means of DFT methods, the performance has been analyzed and rationalized according to five different categories of functional such as: LDA (local density approach), GGA (Generalized Gradient Approximation), meta-GGA (which includes not only the density and its first derivative in the exchange-correlation potential, as GGA, but also the second derivative); GH (global hybrids, which include a predetermined amount of exact exchange), and LCH (long-range-corrected hybrids, whose fraction of exchange depends on the interelectronic distance) [59-61]. The results of a GGA functional (PBE) and a LCH (ωB97X-D) have been compared giving similar results.

Results and discussion

Colchicine-tubulin interactions in the triplet excited state

In organic solvents, LFP of COL gives rise to the triplet excited state (³COL*) as a well characterized transient with absorption maximum at λ_{max} ca. 420 nm [45]. In PG buffer, 355 nm laser excitation of 5 × 10⁻⁵ M COL, either under aerobic or anaerobic conditions, did not lead to any transient absorption spectrum, in agreement with previous results obtained in aqueous medium [45,62]. As a control experiment, LFP of a TU solution gave no signal either. By contrast,

when COL (5 × 10⁻⁵ M in GTP) was submitted to LFP in the presence of TU (0.5 and 1 × 10⁻⁵ M) a transient absorption spectrum peaking at λ_{max} ca. 420 nm was indeed registered (see Fig. 1). This reveals generation of an intermediate detectable in the microsecond timescale. Its quenching by oxygen and its similarity to the transient absorption spectrum obtained for ³COL* in acetonitrile [45] support the assignment as ³COL* complexed with TU (³COL*@TU). Moreover, in aerated medium, the determined lifetime (τ ca 160 μs) was more than 500 times longer than that obtained for ³COL* in acetonitrile (τ = 0.29 μs), which agrees well with the markedly longer triplet lifetimes usually measured in the intraprotein microenvironment [43,44,63].

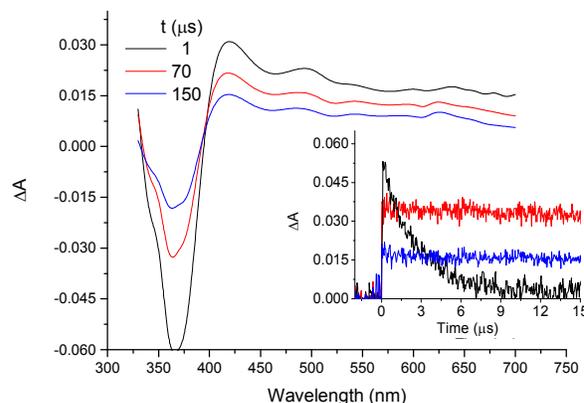


Fig.1 Transient absorption spectra of COL in aerated aqueous PG buffer, in the presence of 10⁻⁵ M TU at different times after 355 nm laser excitation. Inset: Comparison between the decay traces at 420 nm after LFP of 5 × 10⁻⁵ M COL in deaerated ACN (black) and in aerated aqueous 10⁻³ M GTP buffer with TU at 5 × 10⁻⁵ M (blue) or 1 × 10⁻⁵ M (red) concentration.

Comparison of the decay traces of ³COL* in acetonitrile and ³COL*@TU in aqueous solutions revealed that the initial intensity of the signal immediately after the laser pulse was higher in the organic solvent (Fig 1, Inset). A proper analysis of this observation should consider the following factors in aqueous medium: a) the TU concentrations are five or ten times lower than those of COL, b) the affinity constant between COL and TU is very high [22] and c) in the absence of TU, COL does not give rise to any observable signal in the microsecond timescale. Whereas a) and b) favor full complex formation, c) implies that all the observed signal can be safely attributed to ³COL*@TU.

At this point, if it is assumed that the molar absorption coefficient of the complex is nearly the same as that reported for ³COL* in acetonitrile, the absorbance of the transient arising from the LFP of 5 × 10⁻⁵ COL should be five or ten times lower in the presence of 0.5 and 1 × 10⁻⁵ M TU than in acetonitrile. Since the real absorbance is close to four times higher than the expected value, the ³COL* quantum yield must be markedly higher inside the protein (0.138 vs 0.037 in acetonitrile [45]). This enhancement suggests a restricted mobility in the protein microenvironment and consequently, limited degrees of freedom, leading to less efficient non-radiative decay pathways.

Interactions of MTC with tubulin

When MTC was submitted to LFP in the presence of TU, a transient absorption spectrum very similar to that of ³COL* with a lifetime of ca 62 μs was obtained (Fig. 2). In view of the parallel behavior of both substrates, it was inferred that the transient

absorption species was the MTC triplet excited state ($^3\text{MTC}^*$). However, before tackling a detailed analysis of the interactions between MTC and TU, it was considered of interest to undertake a more detailed characterization of the transient. With this purpose, LFP experiments were performed with MTC (5×10^{-5} M) in acetonitrile, methanol, ethanol and H_2O under aerated and deaerated conditions.

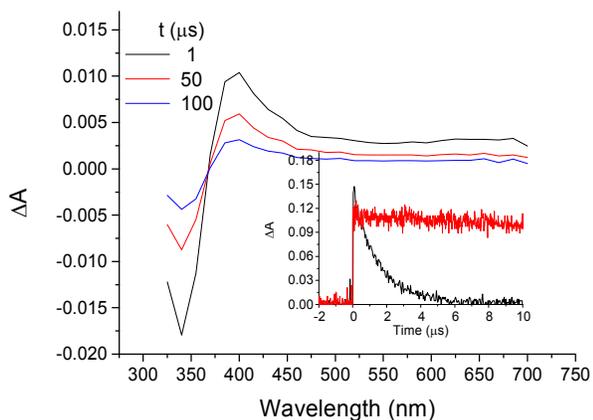


Fig.2 Transient absorption spectra of aerated 5×10^{-5} M MTC in aqueous PG buffer in the presence of 1×10^{-5} M TU at different times after laser excitation. Inset: Decay traces at 430 nm of MTC (5×10^{-5} M) in a deaerated acetonitrile solution (black) and in an aerated aqueous 1×10^{-3} M GTP in the presence of 1×10^{-5} M TU (red).

As previously observed with COL, aqueous solutions of MTC did not produce any signal in the microsecond timescale, while a COL-like transient species with λ_{max} ca. 400 nm was observed in the other solvents (Fig. 3). The initial absorption of this species decreased with the increasing protic character of the solvent. As expected for a triplet excited state, it was quenched by molecular oxygen in acetonitrile with rate constant of $3 \times 10^9 \text{ M}^{-1} \text{ s}^{-1}$ (Fig. 3, inset).

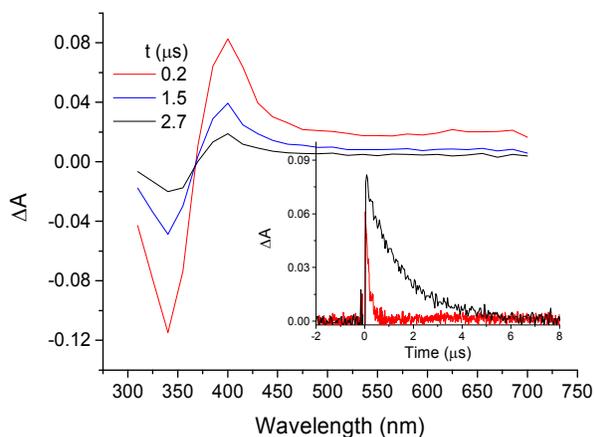


Fig.3 Transient absorption spectra of 5×10^{-5} M MTC in deaerated acetonitrile at different times after laser excitation. Inset: Comparison between the decay traces at 400 nm after LFP of 5×10^{-5} M MTC solutions in ACN under N_2 (black) and air (red) atmospheres.

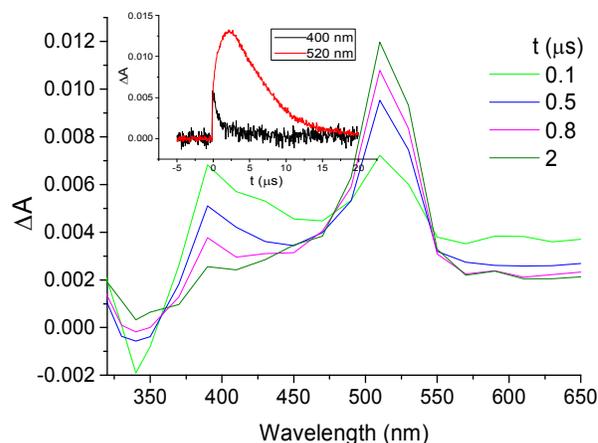


Fig.4 Laser flash photolysis spectra of 5×10^{-5} M MTC in acetonitrile, in the presence of 5×10^{-5} M $\beta\text{-Car}$, 0.1 μs , 0.5 μs , 0.8 μs and 2 μs after the laser pulse. Inset: Decay and growth traces at 400 nm (black) and 520 nm (red), corresponding to $^3\text{MTC}^*$ and $^3\beta\text{-Car}^*$, respectively.

This assignment was confirmed by energy transfer to β -carotene ($\beta\text{-Car}$) in acetonitrile (equation 1 in the experimental section). Thus, the decay at λ_{max} ca. 400 nm was concomitant with the growth of $^3\beta\text{-Car}^*$ at 520 nm (Fig. 4). This reaction was diffusion-controlled (k_q ca. $2 \times 10^{10} \text{ M}^{-1} \text{ s}^{-1}$), in agreement with the expectations based on the low $^3\beta\text{-Car}^*$ energy (79 kJ mol^{-1}) [48].

Table 1. Photophysical properties of MTC in different solvents.

Solvent	Absorption (λ_{max} , nm)	τ_T (μs)	Φ_{ISC}	Solvent dielectric constant	HBD ability
H_2O	343	---	0	78.3	1.19
CH_3OH	341	2	0.020	32.6	0.98
$\text{CH}_3\text{CH}_2\text{OH}$	340	1.9	0.029	24.5	0.86
CH_3CN	335	1.8	0.074	37.5	0.19

The triplet molar absorption coefficient (ϵ) at 400 nm was found to be $31200 \text{ M}^{-1} \text{ cm}^{-1}$ in acetonitrile, as indicated in the experimental section (equation 2). When this parameter was used to determine the intersystem crossing quantum yield (Φ_{ISC}) of MTC (see equation 3 in experimental section), important solvent dependence was observed (Table 1). The hydrogen bond donating (HBD) ability of the solvents [64] seems to be better correlated with the Φ_{ISC} changes than the polarity or the dielectric constant. Besides, the triplet lifetimes in all solvents were very similar (ca. 2 μs , Table 1).

The triplet energy (E_T) was obtained by means of emission measurements. Thus, the phosphorescence spectra of MTC in MeCN as well as in 1 mM PB aqueous medium at 77 K consisted of a structured band with two maxima at 615 and 670 nm. From these spectra, the E_T was found to be ca 200 kJ mol^{-1} . This value matches with that obtained for COL in the same way and is close to that estimated for COL (188 kJ mol^{-1}) in a previous study [65].

After characterization of the transient, the interaction of MTC with TU was investigated. Assuming that the molar absorption coefficient for the $^3\text{MTC}^*\text{@TU}$ complex at 430 nm is similar to that determined for $^3\text{MTC}^*$ in acetonitrile, it is possible to estimate the percentage of MTC bound to TU. As in the case of COL, comparison

of the maximum intensity of the decay signals in organic solution and in the presence of TU showed again that, at the employed concentrations, the value in acetonitrile was more than seven times higher than in the presence of the protein. Using the reported equilibrium constant (K_a ca $5 \times 10^5 \text{ M}^{-1}$ at 25 °C) [28,29] it was possible to establish that the degree of occupancy of the colchicine sites of TU by MTC at 25 °C is between 98 and 95 %. Based on these results, and following the same reasoning as in the case of ${}^3\text{COL}^*\text{@TU}$, the quantum yield of the ${}^3\text{MTC@TU}^*$ complex was found to be 0.296, ca. 4 times higher than that found for ${}^3\text{MTC}^*$ in acetonitrile (0.074).

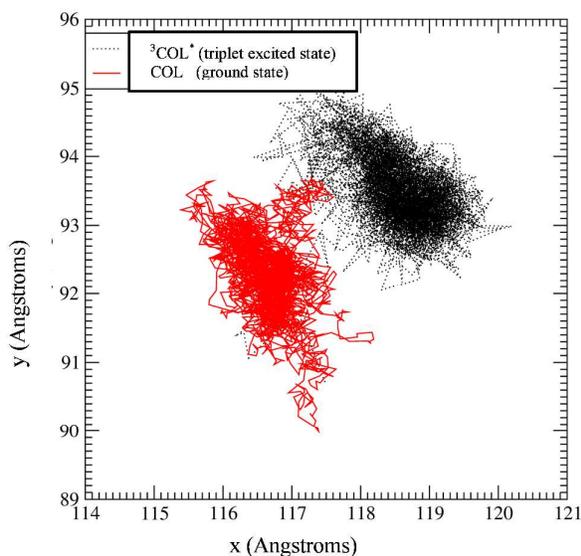


Fig.5 Trajectories across xy of the centre-of-mass coordinates of COL triplet excited state in TU obtained from the molecular dynamics run (500 ps) in which the force field has been adapted to reproduce the triplet excited state geometry as obtained from the DFT calculation (Fig. 8). In this graph, the corresponding trajectory (500 ps) of the COL ground state has been superimposed in order to show their similar diffusivity.

Computational analysis

Molecular dynamics simulations at 1 atm and 298 K, allowed us to obtain the positions of all the atoms of the system during 1 ns. The possibility of extending the time window was considered, but this would only make sense to simulate the long trajectory from the bulk solution to the intraprotein microenvironment. This does not seem necessary when the ligand is initially placed in the known active site, as in our case. Once there, only minimal motion is anticipated, which requires a much shorter time window. As a matter of fact, extension of the simulation period up to 10 ns (see supplementary information) did not lead to significant changes. Fig. 5 shows the positions of COL (red) as it moves around within TU, whose atoms are not displayed because the interest was focused on the relative diffusional paths. Thus, COL spends all the simulation time near the same area within a rectangular box of $[\Delta x, \Delta y, \Delta z] = [3, 5, 4] \text{ \AA}$; hence it is strongly adsorbed at its known binding site.⁶ A general view of the COL@TU complex, at the end of the simulation, is shown in Fig. 6.

The total energy of COL-TU and MTC-TU interactions (Fig. 7) can be split into its short-range (van der Waals) and long-range (Coulombic) contributions. These results explain the reduced mobility of COL (Fig. 5), and also its irreversible binding to TU, due to the strong adsorption energy (-330 kJ mol^{-1} on average).

Diffusion of the COL triplet excited state within TU is also shown in Fig. 5. This has been considered because lifetimes of this species inside TU are much longer than the 1 ns simulation time (in the order of 160 μs microseconds). Moreover, triplet excited states may show significant differences in geometry and/or polarity, which would exert a direct influence on the diffusivity inside TU.

While the previous molecular dynamics study based on the Universal force field (UFF) is capable to reproduce the geometries of the drugs in their ground state, this force field would be totally useless to predict and simulate excited state geometries. Hence, to study the diffusivity of excited triplet states a two-steps strategy have been employed. First, the triplet geometry of COL has been calculated using first principles based on density functional theory (DFT). This geometry, as well as the corresponding charge distribution, has been compared with those of their ground states to find the main differences that may result in specific triplet diffusivity. Secondly, a reparameterization of UFF has been made so that the specific triplet geometries can be reproduced within this force field. With the modified UFF, a molecular dynamics run has been performed in order to check whether there are significant differences between the COL@TU diffusivity in the ground and triplet states.

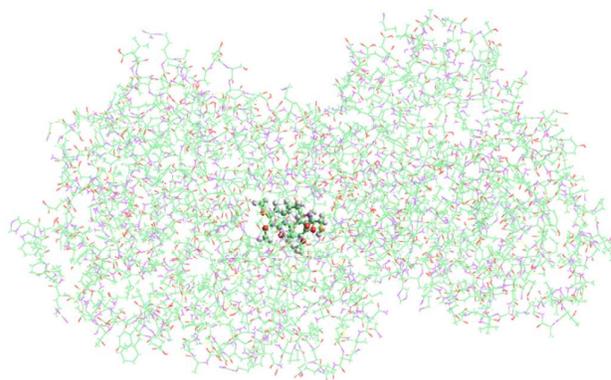


Fig.6 General view of the COL@TU complex near the equilibrium, at the end of the simulation time.

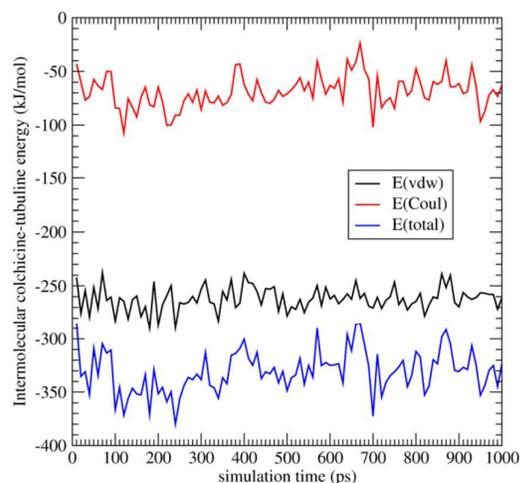


Fig.7 Intermolecular interaction energy and its components (short-range vdw and long-range Coulomb) corresponding to the molecular dynamics runs, COL@TU.

The first-principles geometry optimizations have also been performed using the PBE [66,67] and ω B97X-D [57] functionals, with TZVP [58] basis set, obtaining very similar results. For calculation of excitation energies by means of DFT methods, the performance has been analyzed and rationalized according to five different categories of functional such as: LDA (local density approach), GGA (Generalized Gradient Approximation), meta-GGA (which includes not only the density and its first derivative in the exchange-correlation potential, as GGA, but also the second derivative); GH (global hybrids, which include a predetermined amount of exact exchange), and LCH (long-range-corrected hybrids, whose fraction of exchange depends on the interelectronic distance) [59-61].⁵⁹⁻⁶¹ The results of a GGA functional (PBE) and a LCH (ω B97X-D) have been compared giving similar results.

For the sake of brevity we only show the results obtained with ω B97X-D, which are more confident because of the improvements in the excitation energies when including exact exchange in the functional. In addition, ω B97X-D includes a dispersion term, which contributes to improve the optimized geometries of large molecules when there is a significant non-covalent short-range interaction. Fig. 8 shows the COL ground (top) and triplet (bottom) state geometries, whereas Fig. 9 shows the corresponding MTC geometries. In both cases the diffusional profiles are very similar and hence the small differences observed in the ground and triplet geometries are not expected to affect the diffusivity significantly. In other words, it can be expected that the mobility of COL and MTC within TU will not be strongly dependent on the electronic state of the molecule.

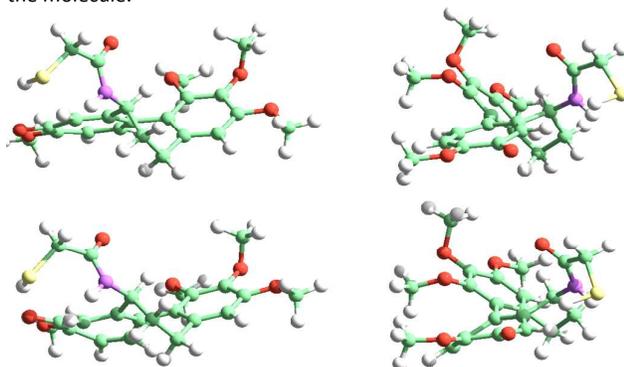


Fig.8 Two views (left and right) of the ω B97X-D optimized geometry of COL in ground (top) and triplet (bottom) electronic states. Top vs. bottom views in each case (left and right) indicated a very similar diffusional profile.

In order to check this hypothesis, a new molecular dynamics run of 500 ps was performed with COL@TU under the abovementioned conditions. With the COL triplet geometry, including the new values of the relevant distances, angles, etc, the UFF force field was modified and loaded as input of the LAMMPS simulation. Fig. 5 shows the new diffusivity of $^3\text{COL}^*$ superimposed to that of COL ground state. It can be seen that the COL-ground and triplet states spend most of their time in a very close location (ca. 2 Å), exhibiting roughly the same mobility. A comparable behavior of their corresponding states is expected for MTC, given their similar geometries (Fig. 9).

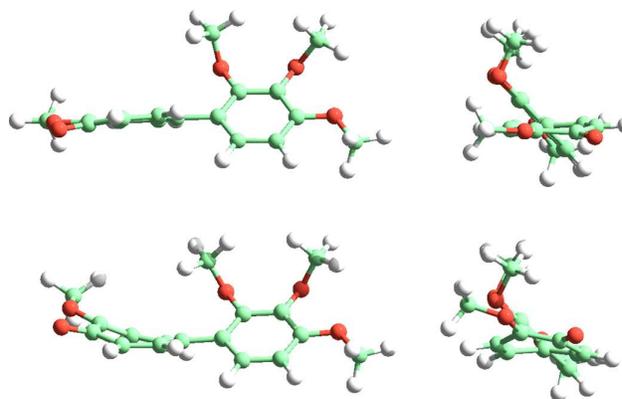


Fig.9 Two views (left and right) of the ω B97X-D optimized geometry of MTC in ground (top) and triplet (bottom) electronic states. Top vs. bottom views in each case (left and right) indicated a very similar diffusional profile.

Conclusions

For the first time, laser flash photolysis has been applied as a new tool to study the interaction between tubulin and drugs such as COL or MTC. Laser excitation of COL in buffered aqueous solutions does not show formation of any short-lived species. However, in the presence of tubulin, COL gives rise to a triplet-triplet transient absorption spectrum very similar to that registered in organic solvents, revealing location of the drug in a hydrophobic region of the biomolecule. Complex formation is accompanied by a remarkable enhancement of the intersystem crossing quantum yields and triplet lifetimes compared to acetonitrile. Likewise, the triplet excited state of MTC has been characterized in several solvents; its quantum yield decreases in protic media (down to nearly zero in water), whereas its lifetime remains practically constant. Again, complexation of MTC with tubulin in aqueous solution leads to a triplet-triplet transient absorption spectrum similar to that obtained in organic solution. The lifetime of this species in the complex is much longer than that measured in organic solvents and the formation quantum yield is also markedly higher. Molecular dynamic simulations, both with the ground and triplet excited state, reveal a strong interaction between COL and tubulin to give stabilized complexes with reduced mobility inside the protein binding site. This effect justifies the enhanced quantum yields and lifetimes of the triplet excited states observed in the drug@protein complexes. These results demonstrate that laser flash photolysis is a useful methodology to study the binding of COL derivatives to tubulin and open a new way to evaluate the interactions of other drugs with this protein. The obtained results can be highly relevant to understand the primary event in the anticancer activity of drugs binding to the COL site of tubulin.

Acknowledgements

Financial support from the Spanish Government (grants CTQ2010-19909; BFU2011-23416 and SEV 2012-0267), the Generalitat Valenciana (Prometeo II/2013/005) and Comunidad de Madrid (S2010/BMD-2353) is gratefully acknowledged.

Notes and References

^a Departamento de Química/Instituto Universitario Mixto de Tecnología Química UPV-CSIC, Avenida de los Naranjos s/n, E-46022 Valencia, Spain.

^b Centro de Investigaciones Biológicas, CSIC, Ramiro de Maeztu 9 E-28040 Madrid, Spain.

*Corresponding author, RT: rtormos@gim.upv.es, MAM: mmiranda@gim.upv.es.

- [1] T. Mitchison and M. Kirschner, *Nature*, 1984, **312**, 237-242.
 [2] R. L. Margolis and L. Wilson, *Cell*, 1978, **13**, 1-8.
 [3] A. Desai and T. J. Mitchison, *Annu. Rev. Cell Dev. Biol.*, 1997, **13**, 83-117.
 [4] J. Howard and A.A. Hyman, *Nature*, 2003, **422**, 753-758.
 [5] M. A. Jordan and L. Wilson, *Nat. Rev. Cancer*, 2004, **4**, 253-265.
 [6] R. B. Ravelli, B. Gigant, P.A. Curmi, I. Jourdain, S. Lachkar, A. Sobel and M. Knossow, *Nature*, 2004, **428**, 198-202.
 [7] A. Cormier, M. Marchand, R. B. G. Ravelli, M. Knossow and B. Gigant, *EMBO Rep.*, 2008, **9**, 1101-1106.
 [8] A. E. Prota, K. Bargsten, J. F. Diaz, M. Marsh, C. Cuevas, M. Liniger, M. Neuhaus, J. M. Andreu, K. H. Altmann and M. O. Steinmetz, *Proc. Nat. Acad. Sci. USA*, 2014, **111**, 13817-13821.
 [9] A. E. Prota, K. Bargsten, D. Zurwerra, J. J. Field, J. F. Diaz, K.H. Altmann and M. O. Steinmetz, *Science*, 2013, **339**, 587-590.
 [10] A. E. Prota, K. Bargsten, P. T. Northcote, M. Marsh, K. Altmann, J. H. Miller, J. F. Diaz and M. Steinmetz, *Angew. Chem. Int. Ed.*, 2014, **53**, 1621-1625.
 [11] F. Santavý, *Collect. Czech. Chem. Commun*, 1951, **16**, 655-675.
 [12] A. Brossi, H. J. C. Yeh, M. Chrzanowska, J. Wolff, E. Hamel, C. M. Lin, F. Quin, M. Suffness, and J. Silverton, *Med. Res. Rev*, 1988, **8**, 77-94.
 [13] J. L. Weiner, A.V. Buhler, V. J. Whatley, R. A. Harris and T. V. Dunwiddie, *J. Pharmacol. Exp. Ther*, 1998, **284**, 95-102.
 [14] M. Imazio, R. Trincherio and Y. Adler, *Future Cardiol*, 2008, **4**, 599-607.
 [15] R. de Vincenzo, G. Scambia, C. Ferlini, M. Distefano, P. Filippini, A. Riva, E. Bombardelli, D. Pocar, M.L. Gelmi, P.B. Panici and S. Mancuso, *Anti-cancer Drug Des.*, 1998, **13**, 19-33.
 [16] M. Fakh, A. Yagoda, T. Reploge, J. E. Lehr and K. J. Pienta, *Prostate*, 1995, **26**, 310-315.
 [17] R.M. Lee and D. A. Gerwitz, *Drug Dev. Res.*, 2008, **69**, 352-358.
 [18] A. Abad, J. L. Lopez-Perez, E. del Olmo, L. F. Garcia-Fernandez, A. Francesch, Ch. Trigili, I. Barasoain, J. M. Andreu, J. F. Diaz and A. San Feliciano, *J. Med. Chem.*, 2012, **55**, 6724-6737.
 [19] R. Alvarez, P. Puebla, J. F. Diaz, A. C. Bento, R. Garcia-Navas, J. de la Iglesia-Vicente, F. Mollinedo, J. M. Andreu, M. Medarde and R. Pelaez, *J. Med. Chem.*, 2013, **56**, 2813-2827.
 [20] D. Panda, J. E. Daijo, M. A. Jordan and L. Wilson, *Biochemistry*, 1995, **34**, 9921-9929.
 [21] A. Lambeir and Y. Engelborghs, *J. Biol. Chem.*, 1981, **256**, 3279-3282.
 [22] J. F. Diaz and J. M. Andreu, *J. Biol. Chem.*, 1991, **266**, 2890-2896.
 [23] J. M. Andreu and J. M. Timasheff, *Biochemistry*, 1982, **21**, 534-543.
 [24] L. Atwell, A. Brossi, M. A. Iorio, T. H. Williams, R. H. Sik, C. F. Chignell, *J. Med. Chem.*, 1981, **24**, 257-261.
 [25] B. Pérez-Ramirez, M. J. Gorbunoff and S. N. Timasheff, *Biochemistry*, 1998, **37**, 1646-1661.
 [26] S. B. Hastie, R. C. Jr. Williams, D. Puett and T. L. Macdonald, *J. Biol. Chem.*, 1989, **264**, 6682-6688.
 [27] Ch. Dumortier, Q. Yan, S. Bane and Y. Engelborghs, *Biochem. J.*, 1997, **327**, 685-688.
 [28] S. Bane, D. Puett, T.L. Macdonald and R. C., Jr. Williams, *J. Biol. Chem.*, 1984, **259**, 7391-7398.
 [29] J. M. Andreu, M. J. Gorbunoff, J. C. Lee and Timasheff, S. N., *Biochemistry*, 1984, **23**, 1742-1752.
 [30] Y. Engelborghs and T. J. Fitzgerald, *J. Biol. Chem.*, 1987, **262**, 5204-5209.
 [31] T. L. Nguyen, C. McGrath, A. R. Hermone, J. C. Burnett, D. W. Zaharevitz, B. W. Day, P. Wipf, E. Hamel and R. Gussio, *J. Med. Chem.*, 2005, **48**, 6107-6116.
 [32] J. T. Huzil, P. Winter, L. Johnson, A. L. Weis, T. Bakos, A. Banerjee, R. F. Luduena, S. Damaraju and J. A. Tuszynski, *Chem. Biol. Drug Design*, 2010, **75**, 541-550.
 [33] R. Cao, M. Liu, M. Yin, Q. Liu, Y. Wang and N. Huang, *J. Chem. Inf. Model.*, 2012, **52**, 2730-2740.
 [34] N. Laing, B. Dahlloef, B. Hartley-Asp, S. Ranganathan and K. D. Tew, *Biochemistry*, 1997, **36**, 871-878.
 [35] K. K. Gireesh, A. Rashid, S. Chakraborti, D. Panda and T. Manna, *Biochem. Pharmacol.* 2012, **84**, 633-645.
 [36] N. Gunasekera, G. Xiong, K. Musier-Forsyth and E. Arriaga, *Anal. Biochem.*, 2004, **330**, 1-9.
 [37] F. J. Medrano, J. M. Andreu, M. J. Gorbunoff and S. N. Timasheff, *Biochemistry*, 1991, **30**, 3770-3777.
 [38] K. C. Morrison and P. J. Hergenrother, *Anal. Biochem.*, 2012, **420**, 26-32.
 [39] S. B. Hastie and R.P. Rava, *J. Am. Chem. Soc.*, 1989, **111**, 6993-7001.
 [40] B. Bhattacharyya, S. Kapoor and D. Panda, *Meth. Cell Biol.*, 2010, **95**, 301-329.
 [41] P. S. Sardar, S. S. Maity, L. Das and S. Ghosh, *Biochemistry*, 2007, **46**, 14544-14556.
 [42] B. Bhattacharyya and J. Wolff, *Proc. Nat. Acad. Sci. USA*, 1974, **71**, 2627-2631.
 [43] V. Lhiaubet-Vallet, Z. Sarabia, F. Bosca and M. A. Miranda, *J. Am. Chem. Soc.*, 2004, **126**, 9538-9539.
 [44] I. Vaya, V. Lhiaubet-Vallet, M. C. Jimenez and M.A. Miranda, *Chem. Soc. Rev.* 2014, **43**, 4102-4122.
 [45] F. Bosca and R. Tormos, *J. Phys. Chem. B*, 2013, **117**, 7528-7534.
 [46] T. J. Fitzgerald, *Biochem Pharmacol.*, 1976, **25**, 1383-1387.
 [47] J. M. Andreu, *Meth. Mol. Medicine*, 2007, **137**, 17-28.
 [48] S. L. Murov, I. Carmichael and G.L. Hug, *Handbook of Photochemistry*, Marcel Dekker, Inc., New York, 2nd edn. (1993).
 [49] J. N. Silva, F. Bosca, J. P. C. Tome, E. M. P. Silva, M. G. P. M. S. Neves, J. A. S. Cavaleiro, L. K. Patterson, P. Filipe, J.-C. Maziere, R. Santus and P. Morliere, *J. Phys. Chem. B*, 2009, **113**, 16695-16704.
 [50] E. J. Land, *Biochimie*, 1980, **62**, 207-221.
 [51] R. V. Bensasson and J. C. Gramain, *J. Chem. Soc.-Faraday Trans.*, 1980, **76**, 1800-1810.
 [52] S. Plimpton, *J. Comp. Phys.*, 1995, **117**, 1-19.
 [53] P. J. in't Veld, S. J. Plimpton and G. S. Grest, *Comp. Phys. Commun.*, 2008, **179**, 320-329.
 [54] A. K. Rappe, C. J. Casewit, K. S. Colwell and W. A. III Goddard and W. M. Skiff, *J. Am. Chem. Soc.*, 1992, **114**, 10024-10035.
 [55] MOPAC2009, James J. P. Stewart, Stewart computational chemistry, version 13.207L; web: [HTTP://OpenMOPAC.net](http://OpenMOPAC.net)
 [56] J. D. Carvalho Maia, G. A. Urquiza Carvalho, C. Peixoto Manguera, S. Ramos Santana, L. A. Formiga Cabral and G. B. Rocha, *J. Chem. Theory Comput.*, 2012, **8**, 3072-3081.
 [57] J.-D. Chai and M. Head-Gordon, *Phys. Chem. Chem. Phys.*, 2008, **10**, 6615-6620.
 [58] A. Schaefer, C. Huber and R. Ahlrichs, *J. Chem. Phys.*, 1994, **100**, 5829-5835.
 [59] D. Jacquemin, W. Wathélet, E. A. Perpète and C. Adamo, *J. Chem. Theory Comput.*, 2009, **5**, 2420-2435.
 [60] D. Jacquemin, E. A. Perpète, I. Ciofini and C. Adamo, *J. Chem. Theory Comput.*, 2010, **6**, 1532-1537.
 [61] M. J. G. Peach, P. Benfield, T. Helgaker and D. J. Tozer, *J. Chem. Phys.*, 2008, **128**, 044118.

COMMUNICATION

Journal Name

- [62]P. Bartovsky, R. Tormos and M. A. Miranda, *Chem. Phys. Lett.*, 2009, **480**, 305-308.
- [63]I. Vaya, C. J. Bueno, M.C. Jimenez and M. A. Miranda, *Chem. Eur. J.*, 2008, **14**, 11284-11287.
- [64]Y. Marcus, *Chem. Soc. Rev.*, 1993, **22**, 409-416.
- [65]A. L. P. Nery, F. H. Quina, P. F. Jr. Moreira, C. E. R. Medeiros, W.J. Baader, K. Shimizu, L. H. Catalani and E. J. H. Bechara, *Photochem. Photobiol.*, 2001, **73**, 213-218.
- [66]J. P. Perdew, K. Burke and M. Ernzerhof, *Phys. Rev. Lett.* 1996, **77**, 3865-3868.
- [67]J. P. Perdew, K. Burke and M. Ernzerhof, *Phys. Rev. Lett.*, 199, **78**, 1396.

Table of Contents

The triplet excited state of complexed COL and MTC gives well defined transient spectra undetectable in the absence of TU.

