Abstract

Thanks to its polyphenols and phytochemicals, green tea is believed to have a number of health benefits, including protecting from heart disease, but its mechanism of action at the molecular level is still not understood. Here we explore, by means of atomistic simulations, how the most abundant of the green tea polyphenols, (−)-Epigallocatechin 3-Gallate (EGCg), interacts with the structural C terminal domain of cardiac muscle troponin C (cCTnC), a calcium binding protein that plays an important role in heart contractions. We find that EGCg favourably binds to the hydrophobic cleft of cCTnC consistently with solution NMR experiments. It also binds to cCTnC in the presence of the anchoring region of troponin I (cTnI(34–71)) at the interface between the E and H helices. This appears to affect the strength of the interaction between cCTnC and cTnI(34–71) and also counter-acts the effects of the Gly159Asp mutation, related to dilated cardiomyopathy. Our simulations support the picture that EGCg interacting with the C terminal domain of troponin C may help in regulating the calcium signalling either through competitive binding with the anchoring domain of cTnI or by affecting the interaction between cCTnC and cTnI(34–71).

Introduction

Tea, produced largely from the plant *Camellia Sinensis*, has been cultivated since ancient times and has been associated with Oriental medicine in the treatment of a variety of ailments. Green tea makes up nearly 20% of the total tea produced worldwide and is prepared in a way which prevents the oxidation of the leaf so that its polyphenolic content is preserved. Green tea polyphenols have been investigated in relation to a number of diseases, such as HIV, influenza, cancer, Alzheimer’s, Parkinson’s and cardiovascular diseases [1–11], with the final goal to ascertain whether the biochemical effects of these polyphenols can be demonstrated and understood and, if so, by which precise mechanism they are carried out. The most abundant of green tea polyphenols by weight is (−)-Epigallocatechin 3-Gallate (EGCg), which is shown in Fig. 1. It is a powerful antioxidant, which includes a benzenediol ring (label A) joined to a tetrahydropyran moiety (C), a pyrogallol ring (B) and a galloyl ring (B’). Rings B and B’ can assume a number of orientations with respect to A and C, depending on the torsional angles indicated as γ, θ and ϕ in Fig. 1.

Experimental studies have shown that green tea polyphenols, when interacting with the protein troponin C, found in cardiac muscles, may modulate the binding of Ca^{2+} ions to the tissue. [12] The binding of calcium ions to cardiac tissue is essential for heart contraction and several diseases are known to modify such an interaction. Therefore the biochemical effects of these polyphenols are of great interest as they could potentially lead to prevention or new treatments of heart disease. [8,13,14].

Troponin is a protein which exists in both cardiac and skeletal muscles and has different functions depending on the location. Cardiac muscle troponin (cTn) plays a critical role in regulating relaxation and contraction in the cardiac thin filament. The protein is attached to tropomyosin and lies between actin filaments found in muscle tissue. It is composed of three distinct protein subunits which have crucial roles in this regulatory function. Troponin C (cTnC) binds Ca^{2+} and regulates the thin filament activation, troponin I (cTnI) inhibits contraction in the absence of Ca^{2+} and troponin T (cTnT) attaches the troponin complex to tropomyosin on the actin filament. [15,16] As shown by solution NMR [17], cTnC is a dumbbell shaped protein consisting of two domains, comprising the C and N terminals, connected by an elongated flexible linker. Each domain contains two EF-hand helix-loop-helix motifs which act to bind divalent cations, though in the N domain one of these sites is inactive. The hydrophobic core of the regulatory N terminal (cNTnC) domain is buried inside the protein, but unlike skeletal TnC, which exposes its hydropho-
bic core when bound to Ca\(^{2+}\), the regulatory domain is not thought to undergo major conformational changes upon binding Ca\(^{2+}\). [17–21] The Ca\(^{2+}\) binding sites on the C terminal of TnC (cCTnC) are of high affinity and thought to remain bound throughout the cardiac cycle; without these ions bound the domain remains unstructured. [22].

cCTnC has been investigated as a possible target for drugs which modulate the calcium sensitivity of the protein (related to its propensity to bind Ca\(^{2+}\) and pass this signal onto the rest of the protein complex). This is believed to be affected by mutations associated to dilated and hypertrophic cardiomyopathy. [12,23,24] Drugs in current use as treatment for heart diseases tend to alter the cytosolic calcium homeostasis and may provoke arrhythmia and even death. Hence there is a pressing need to find alternative therapies, such as calcium sensitisers and desensitisers. The structure of the calcium sensitiser EMD 57033, which we will also study for comparison, is shown in Fig. 1.

The C terminal domain of human cardiac troponin C in complex with EGCg has been studied by solution NMR. [13] A possible binding site was identified at the surface of cCTnC hydrophobic cleft. A three-dimensional structure for the cCTnC\(\cdot\)2Ca\(^{2+}\)\cdot\)EGCg complex has been proposed and is shown in Fig. 2; residues with significant nuclear Overhauser enhancement (NOE) contacts include Met120, Leu121, Leu136, Met157 and Val160 and are highlighted, together with Gly159 whose mutation to Asp is important for dilated cardiomyopathy. Mutations in the N terminal domain have been found to affect calcium sensitivity. [20] The binding site for EGCg in cCTnC is similar to that of the calcium sensitiser EMD 57033, which was also resolved by solution NMR spectroscopy [25] and is shown in Fig. 2. When the anchoring region of troponin I (cTnI[34–71]) was present (occupying the hydrophobic cleft), EGCg formed a ternary compound with cCTnC and cTnI [13]; no structure at atomic resolution of this ternary compound has been proposed based on spectroscopy experiments.

It is the aim of this study to provide an atomistic picture of the interactions of EGCg with the C terminal domain of cardiac muscle troponin C, which complements and most importantly expands the available experimental information. To achieve this goal, we used a series of computational techniques including ligand-protein docking, molecular dynamics (MD) and metadynamics [26,27], a method for accelerating rare events and sampling free energy landscapes, to elucidate the mechanisms of binding of EGCg to cCTnC, both in the absence and presence of the anchoring domain of cTnI. Moreover, we analysed the EGCg influence on the Ca\(^{2+}\) ions and assessed the effects of the mutation of Gly159Asp, which is related to dilated cardiomyopathy. We also compared the behaviour of EGCg with that of the calcium sensitiser EMD 57033 in complex with cCTnC. This extensive computational investigation provides a wealth of insights into the mechanisms of interaction and the dynamics of binding of EGCg with the C terminal domain of cardiac troponin C that may be useful for the development of new, complementary or preventative therapies for cardiovascular diseases.

**Methods**

EGCg partial charges were evaluated so to reproduce the electrostatic potential (ESP) [28] in a density functional theory (DFT) calculation with the BLYP gradient corrected exchange and correlation functional [29,30] and a 6–31\(^+\)G\(^{**}\) basis set, using the Gaussian03 package. [31].

Docking simulations were carried out with AUTODOCK [32] for EGCg docked into respectively cCTnC\(\cdot\)2Ca\(^{2+}\) and cCTnC\(\cdot\)2Ca\(^{2+}\) plus the anchoring domain of cTnI (cCTnC\(\cdot\)2Ca\(^{2+}\)\cdot\)cTnI[34–71]); both these systems have been investigated experimentally in complex with EGCg. [13] The

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**Figure 1. The structures of EGCg and EMD 57033.** Top: the green tea polyphenol EGCg with labels for carbon bonded hydrogen atoms (left), and rings and dihedral angles (right). Bottom: the calcium sensitiser EMD 57033.

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structure of cCTnC:2Ca$^{2+}$ used for the docking was the lowest energy structure from the solution NMR experiment (PDB ID: 2KDH). [13] Preliminary blind docking studies including the entire protein were carried out, showing a clear preference for docking at the hydrophobic core. EGCg was then docked within a 2Ca$^{2+}$ complex with EMD 57033 [25] (model M$_{exp}$) in violet, [13] while the docked poses corresponding to models M$_1$, M$_2$ and M$_3$ are respectively in grey, red and cyan. Right: cCTnC in complex with EMD 57033 (25) (model M$_{EMD}$).
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![Figure 2. The C terminal domain of troponin C in complex with EGCg and EMD 57033.](image)

Left: cCTnC in complex with EGCg. Calcium ions are shown in mauve. The residues Met120, Leu121, Leu136, Met157 and Val160 are in orange, Gly159 in yellow. The solution NMR binding pose for EGCg (model M$_{exp}$) is in violet, [13] while the docked poses corresponding to models M$_1$, M$_2$ and M$_3$ are respectively in grey, red and cyan. Right: cCTnC in complex with EMD 57033 (25) (model M$_{EMD}$).
Surface Area (PBSA) scheme. Trajectory snapshots used in the binding enthalpy calculations were taken every 10 ps of MD and a salt concentration of 0.1 mM was considered. A dielectric constant of 2.0 was used for the proteins, which contained several charged residues (e.g. 49% of cTnC residues are charged), as this has been shown to improve the accuracy of the enthalpic contributions in such cases. [42-44] Entropic contributions were calculated using normal mode analysis on a sample of 20 snapshots. [41,45,46]

There are a number of approximations in the MM/PBSA and normal mode scheme which limit its usefulness in predicting absolute free energy values; however meaningful trends can be obtained by comparing similar structures. [47,48]

Metadynamics simulations [26,27] were also performed in order to explore the free energy landscape of cTnC-2Ca$^{2+}$-EGCg as a function of selected relevant distances between the EGCg rings and cTnC residues with significant experimental NOE contacts. The well-tempered scheme was chosen due to its efficient sampling speed and reduced error in free energy landscape reconstruction. [49] These calculations were undertaken using the PLUMED plug-in for AMBER10. [50] Specifically three distances between the average position of the methyl hydrogen atoms in the side chains of Met120, Leu136 and Val160 and the average positions of respectively EGCg H12/H13 on the A/C ring, H9/H10 on the B’ ring and H16/H17 on the B ring (see Fig. 1) were used as collective variables (CVs). These distances were selected as they are related to those inferred from the NOE intensities and will be indicated as CV$_{Met}$, CV$_{Leu}$ and CV$_{Val}$. Gaussians of height equal to 0.1 kcal/mol and width 0.2 Å were deposited every ps and a bias factor of 10 was used. The sampled distances were limited by imposing a potential wall of magnitude 100 kcal/mol at 12 Å. All other computational details were as in the MD simulations. Free energies were averaged over 100 profiles. The starting configuration of cTnC-2Ca$^{2+}$ for the metadynamics simulation was chosen to be the solution NMR experimental structure equilibrated by MD.

Results and Discussion

**cTnC-2Ca$^{2+}$-EGCg**

We first investigated the interaction of EGCg with the C terminal domain of TnC in order to provide details of the interaction mechanisms that extend the experimental solution NMR information. We carried out a series of molecular dynamics simulations for the cTnC-2Ca$^{2+}$-EGCg complex starting from different initial configurations so to test whether EGCg can bind in different ways. One of the starting configurations for MD was the lowest energy experimental structure (M$_{\text{exp}}$). We also tested three docked structures as follows: (i) the lowest energy configuration from the most populated and lowest energy cluster (M$_1$), which is positioned towards the edge of the hydrophobic cleft; (ii) a configuration in the same cluster as M$_1$ but roughly rotated by 180° with respect to the symmetry axis of the galloyl (B’ ) ring so that the positions of the A and B rings are swapped (M$_2$); (iii) a configuration which does not dock directly to the hydrophobic cleft but slightly further out by the terminal strand (M$_3$) belonging to a cluster characterized by the third lowest energy and the second highest population. The initial positions of EGCg interacting with cTnC are shown in Fig. 2. Additionally a model (M$_\text{mut}$) obtained by introducing the Gly159Asp mutation in M$_{\text{exp}}$ and a model with EMD 57033 (M$_{\text{MD}}$) from solution NMR spectroscopy [25] were also studied. EMD 57033 binds more deeply inside the cleft than EGCg as shown in Fig. 2.

All models were stable as shown by the behaviour of the root mean square displacements (RMSD) of the backbone with respect to the optimized structure. In the 45 ns of simulation used for statistical analysis EGCg (and EMD 57033) remained bound in all models. The mutated cTnC had relatively larger backbone fluctuations (on average 4.11 ± 0.62 Å) than the wild type models (e.g. 3.09 ± 0.32 Å for M$_{\text{exp}}$), which were reduced to 3.92 ± 0.40 Å upon binding EGCg.

Hydrogen bonds formed by EGCg with the solvent and the protein were monitored throughout the MD simulations. EGCg alone in water showed significant hydrogen bonding to the surrounding solvent; e.g. in a 20 ns MD at ambient conditions of which the last 15 ns were used to calculate averages, EGCg formed a total of 18.0 hydrogen bonds with water molecules of which 11.1 with its oxygens as acceptors and 7.7 as donors. A small amount of intra-molecular hydrogen bonds (on average 0.0) on the EGCg B and B’ rings were detected, with very short individual lifetimes (less than 10% of the total simulation time) due to the competition with hydrogen bonds with the solvent; in fact, this amount increases to 1.7 in vacuo. The average number of hydrogen bonds EGCg formed with the C terminal domain of troponin C are listed in Table 1, together with residues involved in the bonds with large occupancies (>30%). The total average number of hydrogen bonds formed by EGCg when in complex with cTnC-2Ca$^{2+}$ in water is reduced compared to the case in water alone, as expected by the proximity with the hydrophobic cleft. Residues Gly159 and Glu161 on the H helix are significant contributors to these interactions in all cases except M$_3$ where EGCg attaches more closely to the G and F helices. The interaction with Gly159 may be important as evinced by the increased RMSD fluctuations noted for the simulations of the mutated system, which becomes heavily influenced by the binding of EGCg to the cleft. In the M$_\text{mut}$ simulation EGCg is seen to shift slightly closer to the terminal strands of cTnC enabling hydrogen bonding with both strands, effectively reducing fluctuations and stabilising the system. Interestingly it was not the mutated negatively charged residue at position 159, but its positively charged neighbour Lys158 to interact with EGCg through a hydrogen bond. With respect to EGCg, EMD 57033 has fewer atom groups capable of forming hydrogen bonds and bound deeply into the hydrophobic core of cTnC leading to a substantially reduced overall amount of hydrogen bonds with both protein and solvent.

| Model | H$_2$O | cTnC | Total | cTnC Residues |
|-------|-------|------|-------|---------------|
| expM | 12.5  | 3.9  | 16.4  | Met120, Gly159, Glu161 |
| M    | 10.7  | 5.2  | 15.9  | Phe156, Gly159, Glu161 |
| M    | 12.5  | 4.5  | 17.0  | Glu126, Gly159, Glu161 |
| M    | 11.3  | 4.2  | 15.5  | Thr124, Glu135, Asp139, Gly140 |
| mutM | 11.9  | 3.9  | 15.8  | Gly91, Met120, Lys158 |
| MD   | 4.8   | 0.1  | 4.9   | --            |

cTnC residues which formed on average more than 0.3 hydrogen bonds with EGCg are listed. [docid:10.1371/journal.pone.0070556.t001]
cCTnC contains aromatic amino acids whose rings may participate in \( \pi-\pi \) interactions with the aromatic rings of EGCg. Of the EGCg models, the only significant \( \pi-\pi \) stacking behaviour was observed in \( M_3 \) between EGCg and Phe156 which participated in 0.5 \( \pi-\pi \)-interactions on average. In \( M_3 \) the looser binding of the ligand allowed the rings to more easily re-arrange into favourable stacking positions than in the other models. In the binding of the ligand allowed the rings to more easily re-arrange with the protein with respect to EGCg.

Phe104, with EMD 57033 rings A and B. The A ring was essentially with Phe156, and to a much smaller degree (3.6%) with M

H10 on ring B

9

with the protein. The A ring was responsible for 69% of this interaction. The increased amount of \( \pi-\pi \) interactions compensated for fewer viable hydrogen bonds with the protein with respect to EGCg.

The experimental NOE-related proton-proton distances [13] were compared with those evaluated in the MD simulations as shown in Fig. 3. The labels for the EGCg hydrogen atoms are the same used in the solution NMR reference [13] and are shown in Fig. 1. Equivalent atoms [H16 and H17 on ring B and H9 and H10 on ring B'] were averaged. Of 17 experimental NOE-related distances <6 Å, 15 were also detected in the MD simulation of \( M_{\text{exp}} \). In that of \( M_1 \), 12 in that of \( M_2 \) and only 5 in that of \( M_3 \). As expected \( M_{\text{exp}} \) is in closest agreement with experiments followed by \( M_2 \), where EGCg has a similar initial ring orientation with respect to the protein cleft as \( M_{\text{exp}} \). Despite an alternative docking orientation, \( M_1 \) still shows a number of NOE-related distances consistent with experiments. \( M_3 \) is the model that deviates the most from experiments, as expected from its binding at the edge of the hydrophobic cleft.

Further exploration of the binding site was done using the metadynamics method for \( M_{\text{exp}} \) to map the free energy landscape as a function of selected collective variables which represent distances between groups of atoms of cCTnC and of EGCg related to significant NOE distances as described in the previous section.

The three CVs map the distance between Val160 and the pyrogallol (B) ring (CVVal,B); Met120 and the tetrahydrouran (C) ring (CVMet) and Leu136 and the galloyl (B') ring (CVLeu). As apparent from the free energy maps in Fig. 4, there are several minima separated by small barriers at distances below 6 Å, plus some at larger distances, revealing that EGCg can easily readjust itself in various favourable slightly different positions with respect to representative residues at the hydrophobic cleft. These minima might act as stepping points for EGCg to bind and unbind to the hydrophobic cleft, although this did not occur in the simulations due to the imposed potential walls.

Readjusting the orientation of the rings through torsions about the dihedral angles \( \gamma, \theta \) and \( \phi \) implies overcoming large energy barriers, except for \( \phi \) at specific values of \( \theta \) and \( \gamma \). In fact, during the MD simulations, in all models \( \gamma \) maintained the equivalent 0° or 180° values, \( \theta \) was mostly close to 90° with a small population at 150° for \( M_{\text{exp}} \). \( M_1 \) and \( M_2 \), while it was mostly at 150° with a small population at 90° in \( M_2 \). In \( M_{\text{exp}} \), \( M_1 \) and \( M_2 \), \( \phi \) remained around 90° confirming that the pyrogallol ring did not substantially move in spite of relatively small torsional barriers due to the constraints imposed by the protein interactions. Conversely the pyrogallol ring of \( M_3 \) was exposed to the solvent and, therefore not feeling the interaction with the protein, could easily rotate.

The behaviours of the calcium ions at the two binding sites, which we labelled I and II in Fig. 2, were monitored and compared for the various models. While these sites are of high affinity and do not instigate the contraction-relaxation cycle in the same manner as the calcium binding to the N terminal domain, it has been shown that mutations which affect calcium sensitivity on the N terminal will also affect the C terminal sites [51] and thus the reverse may also hold true. The distributions of the RMSD of each calcium ion calculated with respect to their positions at the end of the MD equilibration for all the models aligned with respect to the cCTnC backbone are shown in Fig. 5. For cCTnC with no ligand, deformation to reduce solvent exposure of the hydrophobic core tended to confine the calcium ion at site I while having the opposite effect on site II. The destabilising effect on site II was so large that the corresponding calcium ion began a transition to a new location after nearly 48 ns (the simulation was extended a further 4 ns in order to map this transition). Interestingly, the mutated systems showed larger fluctuations overall, especially for the calcium ion at site I, though no transition to a different location was observed. This is consistent with the experimental observation that calcium ions show a reduction of binding affinities in cardiac tissue affected by dilated cardiomyopathy. [52–54] The calcium sensitisier EMD 57033 appears to highly restrict the movement of both calcium ions, which may provide a comparative example of sensitisier and/or regulator behaviour. The addition of EGCg binding to the hydrophobic cleft also markedly reduced fluctuations of both site I and II. Results for the cCTnC-2Ca\(^{2+}\)-TnC case will be discussed in the following section.

The binding free energies of EGCg and EMD 57033 in the selected models, calculated within the MM/PBSA and normal mode analysis scheme, are shown in Table 2. The entropic contributions are fairly similar, as expected for related systems involving the same protein. The free energies should be considered relatively to each other rather than in absolute terms, as the enthalpic contribution depends on the chosen dielectric constant. In \( M_{\text{exp}} \), \( M_1 \) and \( M_2 \) EGCg is similarly bound, with \( M_1 \) having the lowest average binding energy, showing that EGCg can bind in different poses with similar energetic contributions. However in \( M_3 \) it is more loosely bound, as expected from the previous analysis. The average binding free energy in the presence of the Gly159Asp mutation is lower than that for wild type models.
Experimental estimations from the dissociation constant for EGCG interacting with cCTnC depend on method and concentration [13,55] and suggests a binding energy of at least 24.1 kcal/mol or lower [13]. EMD 57033 binds more strongly to cCTnC than EGCG consistently with a comparison of dissociation constants from NMR experiments. [13,25].

We then investigated the interaction of EGCG with the C terminal of troponin C in the presence of the anchoring part of troponin I. We studied the cCTnC:2Ca^2+:EGCG complex in a manner similar to cCTnC through docking with EGCG followed by MD simulations. We considered models with and without EGCG, and with and without the Gly159Asp mutation. The goals of these calculations were to locate the binding site of EGCG when the hydrophobic cleft of cTnC is occupied by the anchoring part of cTnI, as well as monitor the effects of EGCG on the interaction between cCTnC and the cTnI anchoring peptide. Previous solution NMR experiments did not completely discern the binding site [13] in this case, though measured chemical shifts suggested likely binding areas. The effect of EGCG on the interaction between cCTnC and cTnI may influence the anchoring of cCTnC to the thin filament, affecting calcium sensitivity. Hence our simulations can provide very important details to elucidate the mechanisms of EGCG interaction which are still experimentally unavailable.

As the binding site was not known, a blind docking was undertaken using a grid which covered the entire cCTnC portion of the cCTnC:2Ca^2+:TnI(34–71) complex. The docking results did not indicate a clear binding site as in the cCTnC case, where the poses were highly clustered at low energy values. However, the poses found for EGCG were predominantly close to helix E or at the interface between helix E and H. The lowest energy pose that was chosen as the initial structure for MD simulations with EGCG, was initially interacting only with helix E, but within the equilibration time it reoriented at the interface between helix E and H. The lowest energy pose that was chosen as the initial structure for MD simulations with EGCG was initially interacting only with helix E, but within the equilibration time it reoriented at the interface between helix E and H. The lowest energy pose that was chosen as the initial structure for MD simulations with EGCG was initially interacting only with helix E, but within the equilibration time it reoriented at the interface between helix E and H.

Figure 5. Mobility of the calcium ions. Distribution of the RMSD of Ca^2+ at site I versus site II for: (a) cCTnC with no ligand, (b) M_1MD, (c) M_2MD, (d) M_1, (e) M_2, (f) M_3, (g) cCTnC:2Ca^2+:TnI(34–71), (h) cCTnC:2Ca^2+:TnI(34–71):EGCG. Data related to the Gly159Asp mutation are shown in red. doi:10.1371/journal.pone.0070556.g005
interactions were formed by EGCg with cCTnC: residues also including Glu94 (2.05) and Asp151 (0.41), besides which 13.3 with water and 3.4 with cCTnC, through a number of Asp151 (1.2) on helix H as shown in Fig. 6. In the presence of the Glu94 (on average 0.9) and Glu95 (2.3) on helix E as well as cCTnC. Hydrogen bonds between EGCg and cCTnC were acted to anchor the EGCg molecule to the E and H helices of which considerably restricted the torsion of the P helix, interface, along the F helix and the E-F loop, although discriminating between direct contact with EGCg and conformational changes due to EGCg binding to the complex cCTnC:2Ca$^{2+}$:Tnl(34–71) was not possible. In ref. [13] the binding site was hypothesized to be close to the F helix; however our simulations would suggest a binding site at the E-H interface, which is still consistent with the experimental chemical shifts at this site. EGCg may also be able to bind in more than one interface, which is still consistent with the experimental chemical shifts for cTnI(34–71) through competitive binding with the hydrophobic cleft, increasing the propensity of binding the inhibitory domain cTnI(128–147) to cCTnC:2Ca$^{2+}$ [13,25]. In the presence of EGCg, the largest chemical shifts for cCTnC were recorded close to the E-H helix inter-protein binding between the anchoring part of cTnl with cCTnC in the investigated models. Replica simulations are shown on consecutive rows.

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| Model                        | $\Delta H$ (kcal/mol) | $\Delta T\Delta S$ (kcal/mol) | $\Delta G$ (kcal/mol) | $\Delta G$ (kcal/mol) |
|------------------------------|-----------------------|-------------------------------|-----------------------|-----------------------|
| [cCTnC-2Ca$^{2+}$:EGCg]-M    | -27.6±5.2             | 20.2±4.3                      | -7.4±6.7              | 0.0                   |
| [cCTnC-2Ca$^{2+}$:EGCg]-M    | -27.7±4.5             | 19.7±4.2                      | -8.1±6.2              | -0.7                  |
| [cCTnC-2Ca$^{2+}$:EGCg]-M    | -250±3.6              | 19.0±3.0                      | -5.9±4.7              | 1.5                   |
| [cCTnC-2Ca$^{2+}$:EGCg]-M    | -219±3.5              | 18.9±5.0                      | -3.1±6.1              | 4.3                   |
| [cCTnC-2Ca$^{2+}$:EGCg]-M    | -281±3.5              | 18.8±3.8                      | -9.4±5.1              | -2.0                  |
| [cCTnC-2Ca$^{2+}$:EMD 57033]-M | -311±3.7              | 20.1±2.5                      | -11.0±4.5             | -3.6                  |
| [cCTnC-2Ca$^{2+}$:Tnl:EGCg]  | -9.4±2.5              | 15.8±2.5                      | 6.5±3.5               | 0.0                   |
| [cCTnC-2Ca$^{2+}$:Tnl:EGCg]  | -11.6±3.6             | 17.9±2.0                      | 6.3±4.1               | -0.2                  |
| [cCTnC-2Ca$^{2+}$:Tnl:EGCg]  | -14.0±3.9             | 18.1±3.9                      | 4.1±5.6               | -2.4                  |
| [cCTnC-2Ca$^{2+}$:Tnl:EGCg]  | -9.1±4.4              | 16.2±3.6                      | 7.0±5.7               | 0.5                   |
| [cCTnC-2Ca$^{2+}$:Tnl:EGCg]  | -128.7±8.4            | 55.5±5.9                      | -74.2±10.3            | 0.0                   |
| [cCTnC-2Ca$^{2+}$:Tnl:EGCg]  | -130.1±8.1            | 56.2±5.5                      | -73.9±9.8             | 0.3                   |
| [cCTnC-2Ca$^{2+}$:Tnl:EGCg]  | -139.3±10.1           | 55.8±6.3                      | -83.5±11.9            | -9.3                  |
| [cCTnC-2Ca$^{2+}$:Tnl:EGCg]  | -137.0±11.7           | 53.5±5.6                      | -83.4±13.1            | -9.2                  |
| [cCTnC-2Ca$^{2+}$:Tnl:EGCg]  | -121.0±10.7           | 55.1±5.1                      | -65.8±11.9            | 8.4                   |
| [cCTnC-2Ca$^{2+}$:Tnl:EGCg]  | -123.1±9.9            | 52.8±5.8                      | -70.3±11.4            | 3.9                   |
| [cCTnC-2Ca$^{2+}$:Tnl:EGCg]  | -126.4±12.5           | 56.7±5.7                      | -69.7±13.7            | 4.5                   |
| [cCTnC-2Ca$^{2+}$:Tnl:EGCg]  | -121.6±6.9            | 52.6±5.6                      | -70.0±8.9             | 4.2                   |

The top part of the table estimates the binding of EGCg or EMD 57033 to cCTnC; the middle part estimates the binding of EGCg when the anchoring part of cTnl is present; the bottom part of the table evaluates the inter-protein binding between the anchoring part of cTnl with cCTnC in the investigated models. Replica simulations are shown on consecutive rows.

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averaged over the eight residues and the eight MD simulations (with and without EGCg and with and without the Gly159Asp mutation), the C terminal strand of the cTnl fragment was seen to have fluctuations of 2.87 ± 0.58 Å; the largest fluctuations for this terminal were observed in the cCTnC$\text{mut}$-2Ca$^{2+}$~Tnl(34–71):EGCg simulations, without which the average value reduced to 2.70 ± 0.30 Å. The N terminal strand of the cTnl peptide had on average fluctuations of 2.68 ± 0.67 Å; in this case the fluctuations were larger when EGCg was present and for the mutated systems with respect to the wild type.

When interacting with cCTnC:2Ca$^{2+}$:Tnl(34–71), EGCg formed on average 19.2 hydrogen bonds of which 4.4 were with cCTnC and 14.8 with the solvent. Protein-ligand hydrogen bonds were observed with the hydroxyl groups of the B and B' rings, which considerably restricted the torsion of the $\theta$ and $\phi$ angles and acted to anchor the EGCg molecule to the E and H helices of cCTnC. Hydrogen bonds between EGCg and cCTnC were formed with the side-chains of the negatively charged residues Gly94 (on average 0.9) and Gly95 (2.3) on the E-F helix as well as Asp151 (1.2) on helix H as shown in Fig. 6. In the presence of the Gly159Asp mutation, EGCg formed 16.7 hydrogen bonds, of which 13.3 with water and 3.4 with cCTnC, through a number of residues also including Gly94 (2.05) and Asp151 (0.41), besides Ser98 (0.24), Asp105 (0.12), Ala108 (0.11) and Tyr150 (0.44); a snapshot of this simulation is also shown in Fig. 6. 0.4 π-π interactions were formed by EGCg with cCTnC:2Ca$^{2+}$:Tnl(34–71):EGCg and 0.5 with cCTnC$\text{mut}$-2Ca$^{2+}$:Tnl(34–71):EGCg, in both cases with the Tyr150 on helix H.

It was shown by NMR spectroscopy that EGCg forms a ternary compound with cCTnC:2Ca$^{2+}$:Tnl(34–71), although decreased chemical shift perturbations and lower affinity with respect to the case when the cTnl anchoring fragment was absent were observed [13]. This is at variance from EMD 50733, which does not bind concurrently with cTnl(34–71), but does bind when the inhibitory domain cTnl(128–147) is present [25]; hence it was argued that EMD 50733 action as a calcium sensitizer may be related to the weakening of the interaction with cTnl(34–71) through competitive binding with the hydrophobic cleft, increasing the propensity of binding the inhibitory domain cTnl(128–147) to cCTnC:2Ca$^{2+}$ [13,25]. In the presence of EGCg, the largest chemical shifts for cCTnC were recorded close to the E-H helix interface, along the F helix and the E-F loop, although discriminating between direct contact with EGCg and conformational changes due to EGCg binding to the complex cCTnC:2Ca$^{2+}$:Tnl(34–71) was not possible. In ref. [13] the binding site was hypothesized to be close to the F helix; however our simulations would suggest a binding site at the E-H interface, which is still consistent with the experimental chemical shifts at this site. EGCg may also be able to bind in more than one location when the cCTnC hydrophobic cleft is occupied by the anchoring section of cTnl.

The relative binding free energy in Table 2 shows a weakening in the binding of EGCg, with respect to the models discussed in the previous section: in fact EGCg interacts more loosely with cCTnC when the hydrophobic cleft, i.e. the obvious binding cavity, is occupied by the cTnl anchoring peptide, consistently with the docking results and as observed in the experiments [13]. When entropic contributions are included, the binding free energy becomes slightly positive; however this energy should not be considered as an absolute value and EGCg remained bound to...
cCTnC:2Ca^{2+}:TnI(34–71) for the entire length of the MD simulations, both in the wild type and mutated systems.

The root mean square displacements of the cCTnC bound calcium ions were monitored throughout the MD simulations and are shown in Fig. 5. In the simulations including cTnI with and without EGCg, the ions appear less restrained than previously observed without the cTnI peptide. As the cTnI fragment occupied the cCTnC hydrophobic cleft, the variation in position of each calcium ion was similar with respect to each other: the occupation of the cleft, by a ligand or protein fragment, seemed to prevent significant structural changes of the calcium binding sites affecting the calcium binding. The presence of EGCg did not seem to strongly affect the calcium RMSD in the wild type system, while it slightly increased them in the mutated system.

We then analyzed the interaction between cCTnC and cTnI(34–71) in the absence and presence of EGCg. As shown in Table 3, cCTnC and cTnI(34–71) interacted via several hydrogen bonds most of which are salt bridges between pair of amino acids of opposite charge as evident from Fig. 7. The observed dominance of salt bridges is consistent with the experimental suggestion that cTnC-cTnI binding is electrostatically driven [56]. The total average number of hydrogen bonds was increased with the mutation of Gly159 with the negatively charged Asp, as we would intuitively expect. However the increased number of hydrogen bonds in the mutated systems, which lead to stronger binding, seems to be in contrast with NMR experiments which detected a slight weakening. [24] Overall, the presence of EGCg reduced or maintained the average number of hydrogen bonds, which may be related to a weakening of the interaction between cCTnC and cTnI(34–71).

The behaviour of the hydrogen bonds is mirrored by the trend observed in the free energies of binding between cCTnC and cTnI(34–71), shown in Table 2: the effect of the Gly159Asp mutation is to strengthen the inter-protein binding while the interaction with EGCg tends to weaken such an interaction. Differences in the average values of the cTnI fragment binding free energy in the cCTnC:2Ca^{2+}:TnI(34–71):EGCg replicas may be attributed to the cTnI N terminal strand fluctuations; the calculated values are however within the statistical error. The low values of the binding enthalpies are due to the flexible cTnI N terminal strand interacting with cCTnC and in fact can be reduced by cutting this tail and performing the calculations for a simplified model, without modifying the trend.

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**Table 3.** Average number of the inter-protein hydrogen bonds and salt bridges between cCTnC:2Ca^{2+} and cTnI(34–71) (wild type and with the Gly159Asp mutation) in the presence and absence of EGCg.

| Model                        | Hydrogen Bonds | Salt Bridges |
|------------------------------|----------------|--------------|
| [cCTnC:2Ca^{2+}] [TnI]       | 11.2           | 8.8 (78%)    |
| *                            | 10.5           | 7.8 (74%)    |
| [cCTnCmut:2Ca^{2+}] [TnI]    | 15.9           | 11.5 (72%)   |
| *                            | 15.6           | 11.3 (72%)   |
| [cCTnC:2Ca^{2+}:EGCg] [TnI]  | 10.6           | 8.0 (73%)    |
| *                            | 10.6           | 8.5 (80%)    |
| [cCTnCmut:2Ca^{2+}:EGCg] [TnI]| 12.5           | 8.6 (69%)    |
| *                            | 12.5           | 9.5 (76%)    |

The percentage of salt bridges with respect to the total number of hydrogen bonds is shown in brackets. Replica simulations are shown on consecutive rows.

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*Figure 6. EGCg in complex with cCTnC when the anchoring region of cTnI is present.* Molecular dynamics snapshots of cCTnC:2Ca^{2+}:TnI(34–71), wild type (left) and with the Gly159Asp mutation (right), interacting with EGCg. cCTnC is shown in green and cTnI(34–71) in red. The cTnC residue at position 159 is in blue. The cTnC residues which formed on average more than 0.4 hydrogen bonds with EGCg are in orange.
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Conclusions

In this work the binding site of the green tea polyphenol EGCG in the C terminal domain of cardiac muscle troponin C was computationally confirmed to be in the proximity of the hydrophobic cleft, consistently with solution NMR data [13]: we found that EGCG can bind in different orientations with fairly similar energies. The molecular interactions which occurred between EGCG and the cardiac protein were thoroughly analyzed. The two calcium ions bound to cCTnC behaved similarly to each other when either EGCG or EMD 57033 bind to cCTnC, although the latter, known as a calcium sensitiser, binds more strongly and deeply into the hydrophobic cleft. The relatively rigid A/C and B’ rings of EGCG tended to orient themselves so to minimise the hydrophobic cleft exposure to the solvent, leaving the B ring able to rotate flexibly. The hydroxyl groups attached to the EGCG aromatic rings were then able to favourably form hydrogen bonds with residues at the edge of the cleft.

Moreover, EGCG was shown to bind, although less strongly, to the cCTnC-2Ca\(^{2+}\)-TnI(34–71) complex at the interface between the E and H helices. In this position EGCG appeared to weaken the binding affinity of cCTnC for cTnI(34–71), which would then modulate the anchoring of the cTnC protein, changing the force generation response.

When the Gly159Asp mutation was present, in all systems but cCTnC-2Ca\(^{2+}\)-TnI(34–71) the high affinity calcium sites showed on average larger displacements than in the corresponding wild type systems, and this effect may be exacerbated for the N terminal calcium site as this is already of low affinity. The Gly159Asp mutation strengthened the interaction, mostly mediated by salt bridges between cCTnC and cTnI(34–71) so the presence of EGCG may compensate this effect. Further investigations on other known cardiac mutations would be important to ascertain whether a cooperative effect on calcium affinity and thin filament binding occurs.

Although the mechanisms of calcium sensitisation and desensitisation are very complex functions of subsequent events and hence difficult to unravel, these results suggest that EGCG may have a “regulating” effect on the troponin system to which it can bind both in the absence (or in competition) and presence of CnI, affecting in the latter case the strength of the protein–protein interaction and hence the calcium signalling transmission. Hence EGCG is an interesting candidate to be considered as a basis for further drug development or simply as a preventative aid against cardiovascular disease. In addition, the E-H helix interface of cCTnC should be considered for further studies as, even if binding with low affinity, EGCG was able to affect the inter-protein interaction and hence the calcium signalling transmission. Hence EGCG is an interesting candidate to be considered as a basis for further drug development or simply as a preventative aid against cardiovascular disease. In addition, the E-H helix interface of cCTnC should be considered for further studies as, even if binding with low affinity, EGCG was able to affect the inter-protein interaction and drugs which can effectively target this region may prove invaluable in treatments of heart disease; alternative binding locations could also be explored.

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Author Contributions

Conceived and designed the experiments: DB FF CM. Performed the experiments: DB. Analyzed the data: DB CM. Contributed reagents/materials/analysis tools: GF. Wrote the paper: DB CM. Revised the manuscript: GF FF. Approved final version: DB GF FF CM.

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