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Green tea catechins in chemoprevention of cancer: A molecular docking investigation into their interaction with glutathione S-transferase (GST P1-1)

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Abstract

The anti- and pro-oxidant effects of green tea catechins have been implicated in the alterations of cellular functions determining their chemoprotective and therapeutic potentials in toxicity and diseases. The glutathione S-transferases (GSTs; EC 2.5.1.18) family is a widely distributed phase-II detoxifying enzymes and the GST P1-1 isoenzyme has been shown to catalyze the conjugation of GSH with some alkylating anti-cancer agents, suggesting that over-expression of GST P1-1 would result in tumor cell resistance. Here we report the docking study of four green tea catechins and four alkylating anticancer drugs into the GST P1-1 model, as GSTs were found to be affected by tea catechins. The EGCG ligands exhibit higher docking potential with respect to the anticancer agents, with a ligand-receptor interaction pattern indicating a high conformational stability. Consequently, the competition mechanisms favourable for the green tea catechins could lead to enzyme(s) desensitisation with a reduction of the alkylating drugs metabolism. The results provide a useful theoretical contribution in understanding the biochemical mechanisms implicated in the chemotherapeutic use of green tea catechins in oxidative stress-related diseases.

Keywords: Green tea chemoprevention, epigallocatechingallate, molecular docking, alkylating anti-cancer drugs, glutathione S-transferases (GSTs, EC 2.5.1.18), phase-II detoxifying enzymes

Abbreviations: GSH, L-glutamyl-L-cysteinyl-glycine; HGST, human glutathione S-transferase; GTP, green tea polyphenols; EGCG, epigallocatechin gallate; ECG, epicatechin gallate; GCG, gallicocatechin gallate; CG, epicatechin gallate; RGS, regulators of G-protein signaling

Introduction

Advances in molecular medicine resulting in new drugs with specific cellular targets are not still sufficient to cover the multidisciplinary aspects related to drug resistance and individual patient variability [1]. In particular, the resistance of human tumor to multiple chemotherapeutic drugs (multidrug resistance, MDR) is known to be a major reason for the failure of cancer therapy [2]. Drug-metabolizing enzymes (DME) play a key role in the activation and deactivation of drugs, including a number of cytotoxics. They are subject

to significant inter- and intra-individual variability, displaying both genetic polymorphism and in some cases inducibility. The glutathione S-transferases (GSTs; EC 2.5.1.18), is a family of widely distributed phase-II detoxifying enzymes (DME) involved in the cells protection (against many xenobiotic substances and products of oxidative stress) and detoxification, by catalytic conjugation of toxic electrophiles (including carcinogens and cytotoxic drugs) with the tripeptide glutathione [L-glutamyl-L-cysteinyl-glycine, (GSH)], making them less toxic and more readily excretable from

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the body [3,4]. Beside detoxification, the GST system possesses an additional function in the development of cytotoxic resistance in cancer cells, playing a negative regulatory role on the mitogen-activated protein (MAP) kinase pathway (through which many cytotoxics induce apoptosis). This effect is mediated by a direct binding to c-Jun-N-terminal kinase I (JNK) that is involved in the stress response and apoptosis by activation of the MAP kinase system [5]. Glutathione is the most abundant intracellular non-protein thiol [6] and its levels are primarily regulated by the enzyme glutamate-cysteine ligase (GCL), which catalyzes the rate-limiting step in overall GSH biosynthesis (Figure 1).

Mammalian cytosolic glutathione S-transferases (GSTs) exist either as homo- or hetero-dimers with a subunit molecular mass of about 25 kDa and with one active site per monomer [7]. The active site can be divided into two adjacent functional regions, a highly specific G-site binding the tripeptidic substrate and a non-specific H-site for binding non polar electrophilic substrate. Most GSTs exist as soluble enzymes, although a small family of microsomal GSTs has been characterized [8,9] and a mitochondrial GST (GST Kappa) has also been identified [10]. GSTs have been classified into at least five distinct gene classes (alpha, mu, pi, sigma and theta) based on substrate specificity and primary structures [11–13]. Three polymorphisms have been reported to decrease or abolish the GST activity [5,14,15], one of these, the human pi-class GST (hGST P1-1), was found to be over expressed in many tumor types, where it catalyzes the conjugation of GSH with some alkylating anti-cancer agents (such as chlorambucil, cyclophosphamide and metabolites and melphalan) [16]; hGST P1-1 is also implicated in the development of tumor resistance towards various anti-cancer drugs [17–20]. The introduction of the enzymes

in colon, stomach, pancreas, uterine cervix, breast and lung tumor cell lines reduces the cytotoxic activity of these drugs, suggesting that over-expression of GSTs (in particular the hGST P1-1 isozyme [21]) would result in tumor cell resistance. Moreover, hGST P1-1 selective inhibitors increase the efficacy of anti-cancer drugs chemotherapeutic in resistant tumor cells [22]. Both results stress the importance of using highly potent hGST P1-1 selective inhibitors, since they could increase the therapeutic outcome of anti-cancer agents [23–26].

Green tea polyphenols, mainly catechins like epigallocatechin gallate (EGCG), epicatechin gallate (ECG) and epicatechin (EC), have been implicated, in protection against cancer, diabetes, cardiovascular and neurodegenerative disorders [27]. Following oral ingestion, EGCG are found in the blood and tissues in high concentrations [28]. Catechins are believed to react with biomolecules either directly or after cellular metabolism and thus alter their functions. Catechins are antioxidants and scavengers of reactive oxygen (ROS) and nitrogen species, and are used in the prevention of oxidative stress related disorders [29]. On the contrary, the most abundant green tea catechin EGCG and other catechins have been shown to spontaneously generate H_2O_2 *in vitro*, with subsequent cell death [30]. EGCG targets multiple cells signalling pathways and control the cell proliferation and apoptosis, especially in cancer cells [31]. In particular, EGCG has important effects on the expression of GSTs and these effects are cancer-specific [32].

Nowadays, the use of antioxidant help to alleviate the toxic chemotherapeutic side effects and increase the efficacy of chemotherapy [33–37], but much debate has arisen about whether and how antioxidant supplementation alters the efficacy of cancer chemotherapy.

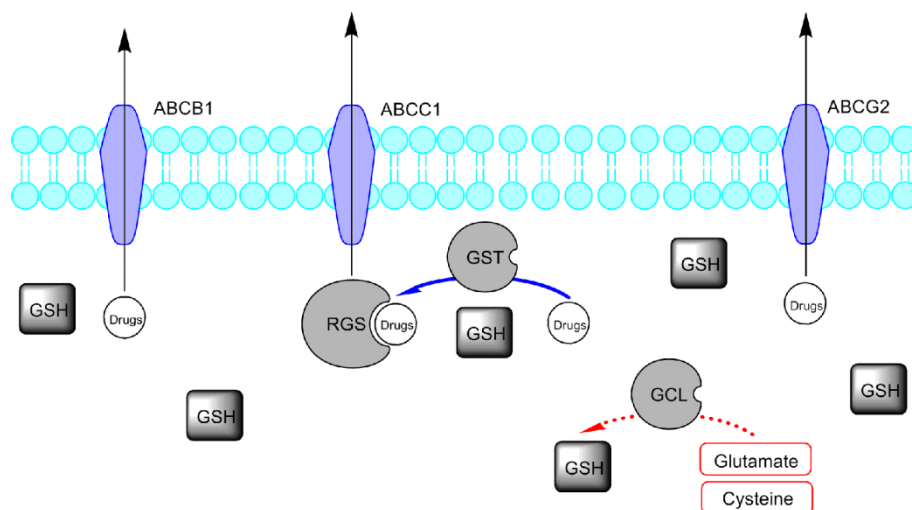


Figure 1. Representative scheme of the glutathione pathway and ABC-transporters family role in drug elimination and defence. Glutathione S-transferases (GSTs) conjugate GSH to drugs and drug metabolites facilitating the ATP-dependent elimination of drugs or drug-GS conjugates, involving ABC transporters (like ABCB1, ABCC1 and ABCG2). The drug/drug metabolites and intracellular GSH equilibrium is a major determinant of blood drug level.

Even if numerous cancer-related proteins are affected by green tea polyphenols [38,39], the molecular bases for green tea polyphenol-mediated cancer prevention remain unknown [25,26]. In particular, they significantly interfere with GSTs activity [40] and the induction of these enzymes by either naturally or synthetic agents represents a promising chemoprevention strategy [41,42]. A recent study have shown that a four weeks green catechins administration in healthy subjects leads to differential effects on lymphocytes GST activity, depending on the its baseline activity. In fact, in this study it has been observed that individuals with the highest level of GST baseline exhibited a statistically significant decrease in mean GST activity. This effect is particularly relevant in clinical protocols where the green tea catechins are combined with anti-cancer drugs, as the GST baseline level is already high for the chemotherapy. In this case the catechins principally could reduce the anti-cancer agents metabolism through GSTs inhibition, affecting the pharmacokinetics and pharmacodynamics of the anticancer drugs. For this reason the achieving of further insight into the mechanism of interaction between green tea catechins

and GST isoform(s) could give deepen confidence to their chemotherapeutic efficacy.

As an X-ray diffraction model of the complexes between the isoform hGST-P1 with the clinically used anti-cancer drugs are still absent, we have performed computational docking calculations to define the putative interactions of four alkylating anti-cancer drugs (cyclophosphamide, ifosfamide, melphalan and chlorambucil) and four natural green tea polyphenols ((-)-EGCG, (-)-GCG, (-)-ECG and (-)-CG) [30] (Figure 2). In the binding model obtained, the natural EGCG ligand exhibits high docking potential over the other studied anti-cancer drugs to the hGST-P1-1, with a ligand-receptor interaction pattern indicating an high conformational stability. This competition in the ligand-enzyme interaction favourable for the catechins with respect to the conventional alkylating drugs, could lead to a decreased tumour cell drug resistance.

Materials and methods

Molecular modelling

Molecular modelling and graphics manipulations were performed using an optimized Mac OSX version

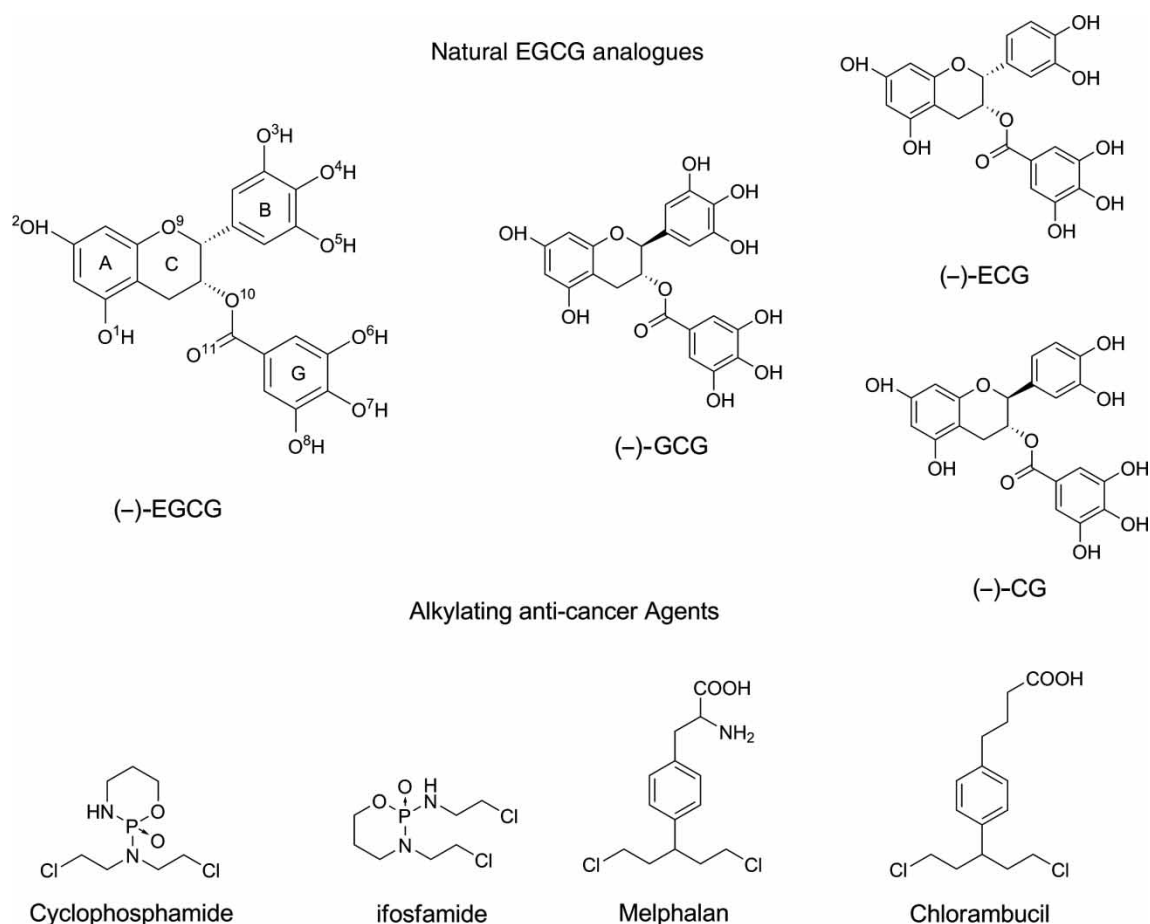


Figure 2. Chemical structures of the green tea catechins and of the alkylating anti-cancer drugs docked. The oxygen numbering scheme and the ring nomenclature [43] used throughout the text is indicated only for (-)-EGCG.

NAMD [44], AutoDock [45] and UCSF Chimera software packages [46] on an Apple® MacPro quad-Xeon workstation running Mac OSX 10.4 (Tiger). Model building and geometry optimizations of the studied compounds were accomplished with the Gaussian 03 (6-31g* base set) [47] quantum mechanical calculations package. The output from AutoDock and all modeling studies as well as images were built with PyMOL [48] and Accelrys DSVisualizer [49] and rendered with POV-Ray [50]. DSVisualizer was used to calculate the hydrogen bonds distances measured between the hydrogen and its assumed binding partner.

Ligand setup

The green tea polyphenols starting conformations were optimized with quantum mechanical calculations by means of Gaussian 03 suite (6-31g* base set) [47]. All ligands atomic charges were assigned using the Gasteiger-Marsili formation, which is the atomic charges type used in calibrating the AutoDock empirical free energy function.

Protein setup

The three-dimensional GST P1-1 starting model was obtained from the Protein Data Bank (PDB code: 1J9H [51]) and it was checked through the OSX version of the AutoDock Tools (ADT, [52]) and UCSF Chimera to guarantee the system conformity with the molecular modelling programs. All the protonation states were set to the normal ionisation forms at pH 7.0 and His residue and both molecular topology and connectivity have been created. The GST P1-1 aminoacidic chain has been terminated with $-\text{COO}^-$ and $-\text{NH}_3^+$ groups in their zwitterionic form and the polar hydrogen atoms were added in their calculated positions. The GST-substrate complex was built docking the green tea polyphenols into the equilibrated GST P1-1 structure and the resulting system was then equilibrated with a series of minimizations interspersed by short molecular dynamics simulations. The resulting structure was submitted for the 1 ns molecular dynamics simulation and then it was optimized by means of an AMBER force field as implemented in the NAMD package [44]. The simulation was performed at constant temperature and pressure (NPT ensemble) in a periodic cubic box of TIP3P water molecules [53,54]. The water bond distances and angles were constrained using the SETTLE algorithm [55], while the protein bond lengths were constrained with the LINCS algorithm [56]. The coupling time was set to 1.0 ps and the isothermal compressibility was set to $4.6 \times 10^{-5} \text{ bar}^{-1}$. Both protein and solvent were independently coupled to a temperature of 298°K, with a coupling time of 0.1 ps and the pressure was held at 1 bar, with a coupling time of 0.2 ps, using

a Berendsen thermostat maintaining temperature and pressure constant. The time step used was 1.0 fs. A total of 6000 snapshots were saved every 0.2 ps. Hydrogen bonds and close contacts were automatically identified using the “contact” module of CCP4 [57] and UCSF Chimera [46], while the other interactions were identified visually.

Docking simulations

Docking of green tea catechins and alkylating drugs to hGST P1-1 was carried out using AutoDock version 4.0 [45]. The AutoDock suite uses an automated docking approach that allows ligand flexibility, and it is able to locate docking poses in a consistent way with respect to the X-ray crystal structures [58]. Default parameters (including a distance-dependent dielectric “constant”) were used as described in the AutoDock manual and both the protein crystal structure and the ligands were prepared for docking by following the default protocols (except for the changes mentioned below). AutoDock uses an empirical scoring function able to approximate the binding free energies, as it includes a solvation free energy term. The energy-scoring grid was prepared as a $20 \times 20 \times 20 \text{ \AA}$ box centered around the S-Hexylglutathione crystallographic molecule, with a 0.375 Å grid resolution. The ligands were limited to move inside this searching space during docking. Atomic solvation parameters were assigned to the protein and the default parameters for the Lamarckian genetic algorithm were used as search protocol, except for the maximum number of energy evaluations, which were changed to 10 million (the population size was raised to 100). For the genetic algorithm, the default parameters were kept for mutation, crossover, and elitism. The docked energy also includes the ligand internal energy or the ligand intramolecular interaction energy. AutoDock also reports a binding free energy excluding the ligand internal energy but including a torsional free energy term, based on the rotatable ligand bonds.

AutoDock docking and hybrid QM/MM scoring

To find out the binding regions of the green tea polyphenols derivatives into the GST structure automated docking simulations were implemented with the AutoDock program version 4.0. As stated above, 100 independent docking runs were performed for each docking experiment. The best generated conformations of each ligand were ranked (clustered) into families of similar binding modes, with a root mean square deviation (RMSD) clustering tolerance of 2 Å. In the second step, we have docked the ligands in the binding sites previously found (“refined docking”). The poses obtained from the docking run were scored using the pseudo-bond *ab-initio* QM/MM approach

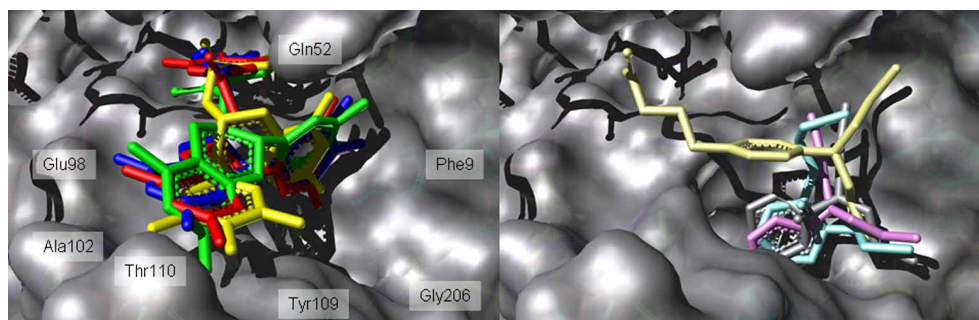


Figure 3. Left: binding into hGST P1-1 of (–)-EGCG red, (–)-GCG yellow, (–)-EGC blue and (–)-CG green. Right: binding into hGST P1-1 of cyclophosphamide gray, ifosfamide light blue, melphalan pink and chlorambucil gold. The protein is represented as a light grey Connolly surface, while the ligands as stick models.

as implemented in Gaussian03 [47]. For the QM/MM calculations, the GST-ligands resulting from the docking were partitioned into a QM and MM subsystems. The reaction system used a smaller QM subsystem formed by the ligand and aminoacids side chains within 3.5 Å from the centre of the catalytic site, while the rest of the protein (the MM subsystem) was treated using the AMBER force field, together with a low memory convergence algorithm. The boundary problem between the QM and MM subsystems was treated using the pseudo-bond approach. An iterative optimization procedure was applied to the QM/MM system, using a B3LYP/3–21G* QM/MM calculations, leading to an reactants optimized structure. The convergence criterion was set in order to obtain an energy gradient less than 10^{-4} using the twin – range cut-off method for non-bonded interactions, with a long- and short-range cut-off of 14 Å and 8 Å, respectively.

Results and discussion

Docking experiments were performed using the crystal structure of hGST P1-1 complexed with the S-Hexylglutathione (PDB entry code: 1J9H), in order to get a better insight into the interaction of the natural EGCG analogs and of the antitumoral alkylating agents with hGST P1-1. An analysis of principles and methods adopted by AutoDock for energy calculations, conformational search and clustering, and for the QM/MM interaction energy ranking is briefly presented in Materials and Methods. The conformations resulting from the automated docking runs of the green tea polyphenols were clustered and most of them (up to 89% of the docking solutions) were found

at the interface between the two GST P1-1 subunits, close to the crystallographic S-Hexylglutathione ligand. The most important residues (within 5 Å from the ligands) of this binding site are Asp⁹⁵, Glu⁹⁸, Asp⁹⁹ and Ala¹⁰² (α -subunit), Tyr⁸, Phe⁹, Arg¹⁴, Trp³⁹, Gln⁵², Pro⁵⁴, Gln⁶⁵, Ser⁶⁶, Asn⁶⁷, Glu⁹⁸, Arg¹⁰¹, Lys¹⁰³ and Tyr¹⁰⁴, Tyr¹⁰⁹, Thr¹¹⁰, Asn²⁰⁵ and Gly²⁰⁶ (β -subunit). The conformer population generated during the docking simulations into the GST P1-1 active site revealed a convergent binding mode of the EGCG analogs, in contrast to what found for the considered alkylating drugs, where the antitumoral chlorambucil is characterized by a divergent orientation with respect to the other (see Figure 3).

The (–)-EGCG ligand exhibits the most favourable energy score (see Table I), with a ligand-receptor interactions pattern indicative of a high conformational stability.

The rather hydrophobic A–C rings (see Figure 2) are oriented towards the β -subunit binding pocket, with the gallate group projected up into the α – β subunits interface, bridging the two walls of the binding cleft. The B ring points towards Tyr⁸(β), which is partially exposed to the solvent and located in a hydrophobic pocket surrounded by Phe⁸, Val¹⁰, Gly¹², Arg¹⁴ and Pro⁵³. (–)-EGCG fills the majority of the binding cleft, as evidenced by drawing the water-accessible surface after the docking into the binding site, as depicted by a ribbon structure (Figure 4).

In addition, (–)-EGCG is characterized by one carbonyl oxygen and eight polar hydrogen's available for H-bonding and among these three actively participate to the formation of an extensive hydrogen

Table I. Heavy atoms (HA), molecular weight (MW) and QM/MM interaction energies (kcal/mol) for the eight studied compounds.

Ligand	HA	MW	Int. Energy	Ligand	HA	MW	Int. Energy
(–)-EGCG	33	458	– 24.7	cyclophosphamide	14	230	– 7.1
(–)-GCG	33	458	– 20.6	ifosfamide	14	230	– 7.3
(–)-EGC	32	442	– 23.5	melphalan	19	305	– 11.3
(–)-CG	32	442	– 24.4	chlorambucil	19	304	– 10.0

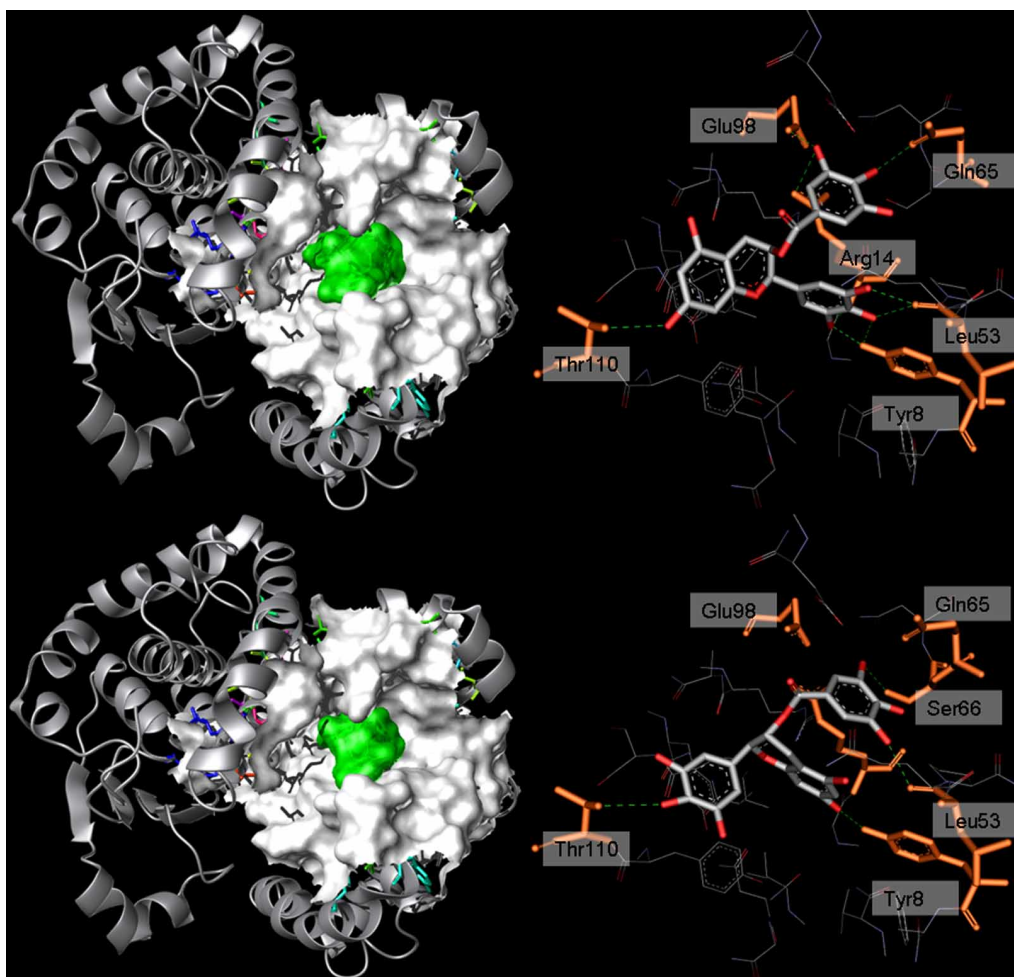


Figure 4. Left: 3D structure of hGST P1-1 (solid ribbon) in complex with (–)-EGCG (top) and (–)-GCG (bottom). Residues lining the ligand position are represented as a white Connolly surface, while the ligands are represented by a green Connolly surface. Right: binding mode of (–)-EGCG (top) and (–)-GCG (bottom) within hGST P1-1. Ligand (CPK) and the residues involved in interactions (orange, labelled) are represented in stick while the hydrogen bonds are shown as green dotted lines.

bonds network (Figure 4). The gallate ring interacts with Arg¹⁴, Gln⁶⁵ and Glu⁹⁸ through O(6) and O(7), while the only interaction of the A–C ring is between O^{γ1}Thr¹¹⁰(β) and O(2). The B ring is located near Arg¹⁴, Leu⁵³ and Tyr⁸ (H-site), and is involved in five H-bonds, O(3)···OLeu⁵³, O(4)···OLeu⁵³, O(4)···HOTyr⁸, O(5)···HOTyr⁸ and O(6)···NArg¹⁴. In particular the O(5)···HOTyr⁸ interaction has a key role for the hGST activity, as indicated by site-directed mutagenesis studies [59,60] showing that its catalytic role is due to the formation of an hydrogen bond between the HOTyr⁸ and GSH stabilizing the thiolate anion form of the substrate. In hGST the direct hydrogen bond between Tyr⁸ and GSH is replaced by an indirect one, mediated through a water molecule [61]. The same binding behaviour is common to all the EGCG analogs except (–)-GCG, the catechin presenting the worse interaction energy. In GCG the A–C ring flips to exchange its position with the B ring, leading to an interaction between the gallate ring and Leu⁵³ and

Ser⁶⁶ through O(6) and O(8). The A–C ring O(2) oxygen is now in contact with HOTyr⁸(β) and the O(4) B ring oxygen hydrogen bonds O^{γ1}Thr¹¹⁰(β), replacing the A–C ring O(2) oxygen atom, while the gallate ring points towards Tyr⁸(β).

The binding mode of the alkylating anti-cancer agents shows the conservation of the Tyr⁸ H-bond, confirming the importance of this residue for GST enzymatic activity. Cyclophosphamide and ifosfamide are completely buried within the H-Site and present the worst energy scores compared with the other studied compounds (Table I). Their phosphamidic nitrogen atoms bind to Tyr⁸, giving rise to one and two H-bonds, respectively. The H-bond network is completed by an additional contact for both compounds between chlorine and NLeu⁵³ (see Figure 5). On the other hand, chlorambucil and melphalan are located close to the catechin positions, with the carboxylic group projected up into the α–β subunits interface and the chloro-ethane groups directed towards the H-Site. Consequently, chlorambucil

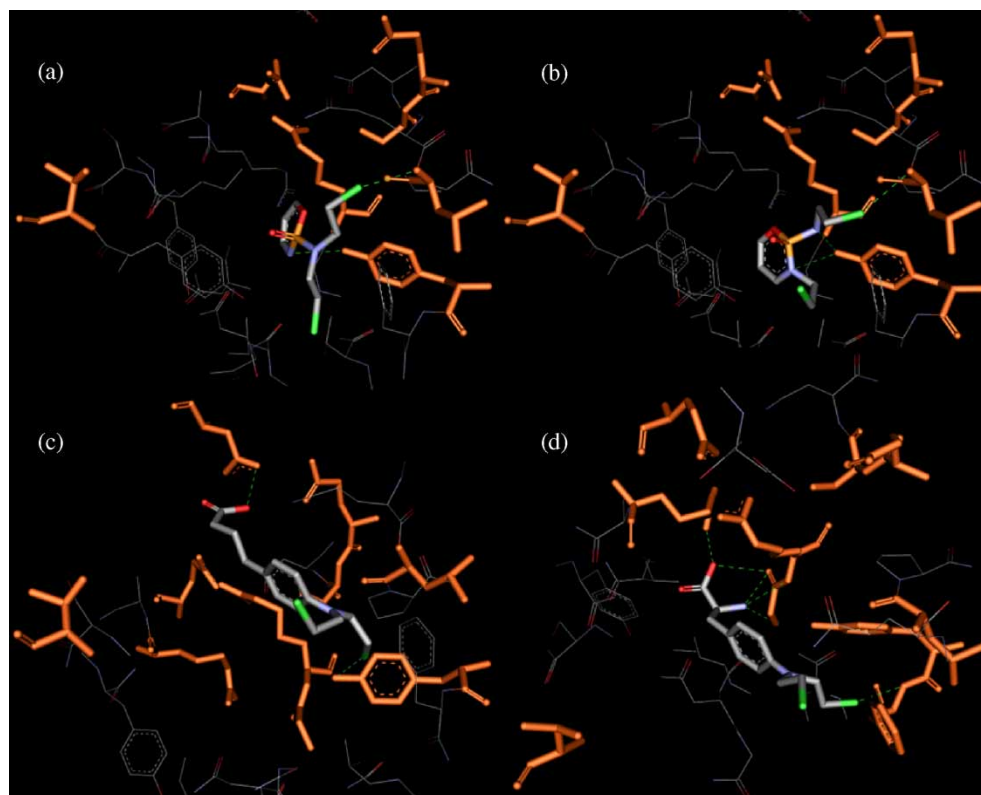


Figure 5. Binding within hGST P1-1 of the considered drugs: cyclophosphamide (a), ifosfamide (b), chlorambucil (c) and melphalan (d). Ligand (CPK) and interacting key residues (orange) are represented as stick models, hydrogen bonds are shown as green dotted lines.

makes two strong H-bonds, between $\text{Cl} \cdots \text{HOTyr}^8$ and between the carboxylic oxygen and $\text{O}^{\delta 2}\text{Asp}^{99}$. Melphalan forms a network of H-bonds: two between the carboxylic oxygen and $\text{NH}_1\text{Arg}^{101}\text{-OGly}^{13}$, two between the aliphatic nitrogen and $\text{NArg}^{14}\text{-NGly}^{13}$ and the last between the chlorine and NPhe^9 and it is the only ligand that do not contact Tyr^8 .

Conclusions

In conclusion, this molecular docking study has been carried out to deepen characterize the possible interactions between EGCGs and alkylating agents on the GST catalytic site. In particular, the interaction mechanisms of the main green tea catechins and of four alkylating drugs with GST P1-1 has been investigated, which may be responsible for the greater systemic concentration of several conventional anticancer drugs. The docking experiments have shown that catechins and the examined drugs compete for the same catalytic binding site, close to the substrate GSH, using a comparable binding mode. This latter involve hydrophilic and steric interactions and, in particular, a direct and strong contact with Tyr^8 (except melphalan), that provides an indirect evidence of the competition between alkylating drugs and catechins. This hypothesis is also supported by the values of the calculated QM/MM interaction energies,

showing that all the green tea catechins bind the hGST P1-1 with higher affinity with respect to the alkylating agents. This could lead to a enzyme desensibilization, decreasing the anti-cancer drugs metabolism resulting in a greater systemic concentration of alkylating agents.

The results presented in this study are useful in view of the potential use of the green tea polyphenols both as anticancer as well as chemopreventive agents. In particular, we have determined the potentially bioactive conformations of EGCGs and anticancer drugs which may constitute the basis for generating GSTs binding models. On the other hand, we have confirmed the importance of the interaction with Tyr^8 for the GST enzymatic activity. Finally, the observed differences in potency between the EGCGs and anticancer drugs have been explained through their different binding site occupancy.

Declaration of interest: The authors report no conflicts of interest. The authors alone are responsible for the content and writing of the paper.

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