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Identification of aspartic acid-203 in human thymidine phosphorylase as an important residue for both catalysis and non-competitive inhibition by the small molecule “crystallization chaperone” 5’-tritylinosine (KIN59)

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3 **Identification of aspartic acid-203 in human thymidine phosphorylase as an important**
4 **residue for both catalysis and non-competitive inhibition by the small molecule**
5 **“crystallization chaperone” 5’-*O*-tritylinosine (KIN59)**
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Abbreviations

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3 6A5BU: 6-amino-5-bromouracil; 6AT: 6-aminothymine; 7-DX: 7-deazaxanthine; BVDU:
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5 (E)-5-(2-bromovinyl)-2'-deoxyuridine; CAM: chicken chorio-allantoic membrane; dThd:
6
7 thymidine; EDTA: ethylenediaminetetraacetic acid; KIN59: 5'-O-tritylinosine; GST:
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9 glutathione S-transferase; HPLC: high-pressure liquid chromatography; MD: molecular
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11 dynamics; MM-GBSA: molecular mechanics-generalized Born surface area; NMA: normal
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13 mode analysis; PMSF: phenylmethylsulfonylfluoride; RESP: restrained electrostatic potential;
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15 rmsd: root mean square deviation; SASA: solvent accessible surface area; TFT: 5-
16
17 trifluorothymidine; TP: Thymidine phosphorylase ; TP65: 9-[8-phosphonoctyl]-7-
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19 deazaxanthine; TPI: 5-chloro-6-[1-(2-iminopyrrolidinyl)methyl]uracil hydrochloride; TS,
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21 transition state.
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Keywords:

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31 Thymidine phosphorylase, 5'-O-tritylinosine (KIN59), allosteric inhibitor, modeling,
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33 mutagenesis
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ABSTRACT

1 Thymidine phosphorylase (TP) is a catabolic enzyme in thymidine metabolism that is
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3 frequently upregulated in many solid tumors. Elevated TP levels are associated with tumor
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5 angiogenesis, metastasis and poor prognosis. Therefore, the use of TP inhibitors might offer a
6
7 promising strategy for cancer treatment. The tritylated inosine derivative 5'-*O*-tritylinosine
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9 (previously designated KIN59) is a non-competitive inhibitor of TP which was previously
10
11 found to be instrumental for the crystallization of human TP. A combination of computational
12
13 studies including normal mode analysis, automated ligand docking and molecular dynamics
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15 simulations were performed to define a plausible binding site for 5'-*O*-tritylinosine on human
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17 TP. A cavity in which 5'-*O*-tritylinosine could fit was identified in the vicinity of the Gly405-
18
19 Val419 loop at a distance of about 11 Å from the substrate-binding site. In the X-ray crystal
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21 structure, this pocket is characterized by an intricate hydrogen-bonding network in which
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23 Asp203 was found to play an important role to afford the loop stabilization that is required for
24
25 efficient enzyme catalysis. Site-directed mutagenesis of this amino acid residue afforded a
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27 mutant enzyme with a severely compromised catalytic efficiency (V_{\max}/K_m of mutant enzyme
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29 ~ 50 -fold lower than for wild-type TP) and pronounced resistance to the inhibitory effect of
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31 5'-*O*-tritylinosine. In contrast, the D203A mutant enzyme kept full sensitivity to the
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33 competitive inhibitors 6-aminothymine and 6-amino-5-bromouracil, which is in line with the
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35 kinetic properties of these inhibitors. Our findings reveal the existence of a previously
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37 unrecognized site of TP that can be targeted by small molecules to inhibit the catalytic activity
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1. Introduction

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3 Thymidine phosphorylase (TP) is a key enzyme of the nucleoside salvage pathway [1] that
4 catalyzes the reversible conversion of thymidine and 2'-deoxyuridine to their respective bases
5 and α -D-2-deoxyribose-1-phosphate. TP not only recognizes natural 2'-deoxynucleosides but
6 is also able to catalyze the phosphorolysis of several nucleoside analogues endowed with
7 antiviral or antitumor activity, such as 5-trifluorothymidine (TFT), 5-(E)-(2-bromovinyl)-2'-
8 deoxyuridine (BVDU) and 5-fluoro-2'-deoxyuridine [2-4]. In addition to its catabolic role in
9 the nucleoside salvage pathway, TP is also implicated in tumor progression as it has been
10 shown to stimulate angiogenesis [5-8] and protect tumor cells from apoptosis [9-11]. A wide
11 range of tumors, including breast, colon and oesophageal carcinomas overexpress TP, and
12 high levels of TP have been correlated with microvessel density, metastasis and poor patient
13 survival [10].

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15 Thus, TP represents a promising chemotherapeutic target and TP inhibitors may (i)
16 improve the antiviral and antitumor activity of certain nucleoside analogues, and (ii) inhibit
17 the angiogenic and anti-apoptotic activities of TP. During the last decade, several classes of
18 novel TP inhibitors have been developed [12]. For more than 30 years, the reference TP
19 inhibitors were 6-aminouracil derivatives including 6-aminothymine (6AT) and 6-amino-5-
20 bromouracil (6A5BU)[13]. In 2000, Fukushima et al. reported on TPI (5-chloro-6-[1-(2-
21 iminopyrrolidinyl)methyl]uracil hydrochloride) as one of the most effective TP inhibitors to
22 date, with an IC_{50} value of 35 nM [14]. In fact, a combination of TPI and TFT is designated
23 TAS-102 which is currently being evaluated in clinical trials for the treatment of various solid
24 tumors, including metastatic colon and breast carcinomas [15, 16].

25
26 Our research groups have described 7-deazaxanthine (7-DX), the first purine
27 derivative with inhibitory activity against TP and angiogenesis in the 'chicken chorioallantoic
28 membrane' (CAM) assay [17], and also TP65 (9-[8-phosphonoctyl]-7-deazaxanthine), the

1 first multisubstrate inhibitor of TP that was shown to interact with both the thymidine and the
2 phosphate binding site of TP [18, 19]. More recently, we identified the purine nucleoside 5'-
3 *O*-tritylinosine as a novel TP inhibitor [20, 21], in which the presence of the trityl moiety was
4 proven to be crucial for both its inhibitory activity against TP and its anti-angiogenic effect in
5 the CAM assay. 5'-*O*-Tritylinosine is also a rather unique TP inhibitor because (i) it promotes
6 the degradation of pre-existing immature vessels in the CAM assay, and (ii) in contrast to all
7 previously described TP inhibitors, it inhibits TP in a non-competitive fashion with respect to
8 both thymidine and phosphate. This latter finding implies that 5'-*O*-tritylinosine does not bind
9 to the substrate-binding sites of TP, but rather interacts with a novel, yet unknown, allosteric
10 site of TP that could also play a role in the biological activities of this protein.
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23 It may be mentioned that diffraction-grade crystals of human TP in complex with TPI
24 [22] were only obtained after limited treatment with trypsin, yielding two protein fragments
25 consisting of residues 12–408 and 411–482. The structure was solved at 2.7 Å resolution,
26 revealing a single protomer in the asymmetric unit, but the polypeptide stretch encompassing
27 amino acids 407–414 could not be determined. In contrast, no proteolysis was required and
28 well-ordered crystals with diffraction at 2.3 Å resolution were obtained when both thymidine
29 (2 mM) and 5'-*O*-tritylinosine (1 mM) were present under the crystallization conditions [23].
30 In this latter structure, two dimers appeared in the asymmetric unit, and residues 406–415
31 were visible as an arginine-rich surface loop that extends across the active site cleft and
32 appears to stabilize the closed conformation of the enzyme and the dimer interface through
33 hydrogen bonds. Taking these two crystal structures together, it is clear that the overall fold of
34 human TP is similar to that previously reported for *E. coli* TP [24, 25] and comprises a large
35 α/β domain connected by three loops to a smaller α domain. These three loops act as a hinge
36 that allows the motion of one domain relative to the other so that the enzyme can interconvert
37 from an open, inactive form to a closed conformation that is catalytically active. Intriguingly,
38 although the presence of 5'-*O*-tritylinosine was instrumental for yielding good-quality
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1 diffracting crystals of TP, no clear electron density was found for this non-competitive
2 inhibitor, which in light of these findings may be regarded as a synthetic small-molecule, so
3 called “crystallization chaperone” [26]. For this reason, we have attempted to find a plausible
4 binding site for 5'-*O*-tritylinosine on human TP and to provide a rationale for its behavior as a
5 non-competitive inhibitor. Our proposal has been indirectly validated by site-directed
6 mutagenesis of Asp-203, an amino acid that has turned out to be important not only for
7 selective inhibition of the enzyme by the 5'-*O*-tritylinosine ligand but also for full catalytic
8 competence of the enzyme.
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20 **2. Methods and materials**

23 **2.1 Reagents**

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27 5'-*O*-Tritylinosine (earlier designated KIN59) was synthesized as described previously [20,
28 27]. The synthesis of 6AT and 6A5BU was performed as described [13, 28]. Thymidine
29 (dThd) and thymine (Thy) were purchased from Sigma (St-Louis, MO). The structural
30 formulae of these compounds, as well as the reference inhibitor TPI, are depicted in Figure 1.
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40 **2.2 Molecular modeling.**

41 **2.2.1 Normal mode analysis.**

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44 The X-ray crystal structure of human TP (PDB code 2J0F)[22] at 2.31 Å resolution, was
45 retrieved from the Protein Data Bank (<http://www.rcsb.org/pdb/>). To probe the flexibility of
46 this enzyme, an elastic network model was used in which all non-hydrogen protein atoms
47 (within a cutoff of 10Å) were modeled as point masses and C α atoms were connected by
48 springs representing the interatomic force fields. Monomer A was analyzed as a large set of
49 coupled harmonic oscillators using the normal mode analysis, deformation and refinement
50 (NOMAD-Ref) server [29] and default parameters. The conformational changes were
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deduced by calculating the 10 lowest-frequency normal modes, which are those with the highest amplitudes and those most often related to large-scale structural rearrangements in proteins. Each mode was explored in its two opposite directions, thus resulting in a series of conformations different from the crystallographic structure within a maximum root-mean-square deviation (rmsd) value of 2 Å.

2.2.2 Quantum mechanical methods, molecular mechanics force field and atom point charges.

The geometry of 5'-*O*-tritylinosine was first refined by means of the semi-empirical quantum mechanical program molecular orbital package (MOPAC) [30], using the AM1 Hamiltonian and PRECISE stopping criteria, and further optimized using a restricted Hartree-Fock (RHF) method and a 6-31G(d) basis set, as implemented in the *ab initio* program Gaussian 03[31]. The resulting wavefunctions were used to calculate electrostatic potential-derived (ESP) charges employing the restrained electrostatic potential (RESP) methodology, as implemented in the assisted model building with energy refinement (AMBER) suite of programs (<http://amber.scripps.edu/>). The missing bonded and nonbonded parameters were assigned, by analogy or through interpolation from those already present in the AMBER database, in a way consistent with the AMBER force field [32]. The geometry and atom point charges for the transition state (TS) of the phosphorolysis reaction were retrieved from previous work [33].

2.2.3 Docking of 5'-*O*-tritylinosine into human TP.

A box encompassing both the active site and the loop regions was defined for the exploration of possible binding modes. The Lamarckian genetic algorithm implemented in AutoDock 3.0.5 [<http://www.scripps.edu/pub/olson-web/doc/autodock>] was then used to generate docked conformations of this inhibitor in either the original protein structure or those derived from the NMA by randomly changing the overall orientation of the molecule as well as the torsion angles of all rotatable bonds. Default settings were used except for number of runs, population

size, and maximum number of energy evaluations, which were fixed at 250, 100, and 250,000, respectively. Rapid intra- and intermolecular energy evaluation of each configuration was achieved by having the receptor's atomic affinity potentials for aliphatic and aromatic carbon, oxygen, nitrogen, and hydrogen atoms precalculated in a three-dimensional grid with a spacing of 0.375 Å. A distance-dependent dielectric function was used in the computation of electrostatic interactions.

2.2.4 Molecular dynamics (MD) simulations.

The MD simulations were carried out using the AMBER 8.0 suite of programs [34]. A model of D203A TP was constructed using the mutagenesis module available in the molecular graphics program PyMOL (<http://www.pymol.org/>). The complexes of wild-type and D203A TP with the transition state (TS) of the reaction [33] and of wild-type TP with KIN59 were neutralized by addition of sodium ions [35] and immersed in truncated octahedrons of TIP3P water molecules [36]. Periodic boundary conditions were applied and electrostatic interactions were treated using the smooth particle mesh Ewald method [37] with a grid spacing of 1 Å. The cutoff distance for the non-bonded interactions was 9 Å. The SHAKE algorithm [38] was applied to all bonds and an integration step of 2.0 fs was used throughout. Solvent molecules and counterions were relaxed by energy minimization and allowed to redistribute around the positionally restrained solute ($25 \text{ kcal}\cdot\text{mol}^{-1}\cdot\text{Å}^{-2}$) during 50 ps of MD at constant temperature (300 K) and pressure (1 atm). These initial harmonic restraints were gradually removed and the resulting systems were allowed to equilibrate further in the absence of any restraints for 10.0 ns during which system coordinates were collected every 2 ps for further analysis.

2.2.5 MM-GBSA binding calculations.

The binding free energy describing protein-ligand associations is the difference between the free energy, G , of the complex and that of the respective binding partners, in our case the

wild-type protein (TP) and the transition state of the reaction (TS), calculated over a series of snapshots (*i*) taken from the MD trajectory:

$$G_{\text{bind}} = G_{(\text{TP-TS})} (i) - (G_{\text{TP}} (i) + G_{\text{TS}} (i)) \quad (1)$$

The value of *G* for each species was calculated using the following energy decomposition scheme, which is implemented in the Molecular Mechanics/Generalized Born/Surface Area (MM-GBSA) method within AMBER 8 (script mm_gbsa.pl) [39, 40]:

$$G = E_{\text{gas}} + G_{\text{sol}} - TS$$

$$E_{\text{gas}} = E_{\text{int}} + E_{\text{ele}} + E_{\text{vdw}}$$

$$E_{\text{int}} = E_{\text{bond}} + E_{\text{angle}} + E_{\text{torsion}} \quad (3)$$

$$G_{\text{sol}} = G_{\text{GB}} + G_{\text{nonpolar}}$$

$$G_{\text{nonpolar}} = \gamma \text{SASA}$$

E_{gas} is the gas-phase energy, which is calculated using the AMBER force field [41, 42] as the sum of internal energies (E_{int}), encompassing bond lengths (E_{bond}), bond angles (E_{angle}) and torsional angles (E_{torsion}), and coulombic (E_{ele}) and van der Waals (E_{vdw}) non-bonded energies. The solvation free energy, G_{sol} , is decomposed into polar and nonpolar contributions: the polar contribution (G_{GB}) is calculated by solving the generalized Born equation [43] using dielectric constants of 1 and 78.5 for solute and solvent, respectively, whereas the nonpolar contribution, G_{nonpolar} , is estimated from the solvent accessible surface area (SASA) [44], which is determined using a water probe with a radius of 1.4 Å. The surface tension constant γ was set to 0.005 kcal/mol/Å². *T* and *S* are the temperature and the total solute entropy, respectively. *S* is typically estimated on the basis of classical statistical formulae and NMA of representative snapshots of energy minimized structures from a MD trajectory. In the present investigation this cumbersome procedure was obviated so that, in a first approximation, the solute entropy estimate were not included in the calculation of the free energy changes by

assuming, on the basis of previous work [45], that its contribution to ΔG_{bind} would be negligible.

On the other hand, the use of a single trajectory to calculate G_{bind} is conducive to a much better agreement with the experimental data than when multiple trajectories are used [46]. It is then expected that, by using a single trajectory, error cancellation will overcome the insufficient sampling of the conformational space [45, 47].

2.3 Site-directed mutagenesis of human TP

The pMOAL-10T vector containing the human TP gene (fused to glutathione S-transferase [GST]) was described and kindly provided by Prof. R. Bicknell (Oxford, UK)[48]. The mutant TP enzyme containing the D203A mutation was derived from this vector. Site-directed mutagenesis was performed using the Quick Change site-directed mutagenesis kit (Stratagene, Westburg, Leusden, The Netherlands) according to manufacturer's instructions. The oligonucleotide primers that were used are 5' cgg aat cct ata tgc agc cag agc tgt gac agc cac cgt gg-3' and its reverse primer (Invitrogen, Merelbeke, Belgium). After linear amplification of the plasmid using *Pfu* DNA polymerase (Stratagene) in a temperature cycler program (30 min at 95 °C followed by 20 cycles of 30 min at 95 °C, 1 min at 55 °C and 12 min at 68 °C), wild-type (methylated) plasmid was digested with *DpnI* restriction enzyme (Stratagene) and the mutant (unmethylated) DNA was transformed into *E. coli* XL1-Blue super competent cells (Stratagene). The presence of the desired mutation in the TP gene was confirmed by sequencing on an ABI Prism 3100 sequencer (Applied Biosystems, Foster City, CA), using the ABI Prism Big Dye terminator cycle sequencing ready reaction kit (Applied Biosystems).

2.4 Purification of human recombinant TP

1 The pMOAL-10T vector containing the wild-type TP or the mutant D203A TP was
2 transfected into *E. coli* BL21(DE3)pLysS. Bacteria were grown overnight at 37 °C in 2YT
3 medium containing ampicillin (Bristol Myers Squibb, Brussels, Belgium) (100 µg/mL) and
4 chloramphenicol (Sigma) (100 µg/mL), diluted 1:10 in fresh medium and further incubated
5 for 5 h at 27 °C. Then isopropyl-β-D-thiogalactopyranoside (IPTG) (Sigma) was added to a
6 final concentration of 0.1 mM to induce the expression of the GST-TP fusion protein. After a
7 further incubation for 15 h at 27 °C, the cells were pelleted (10,000×g for 10 min at 4 °C) and
8 resuspended in lysis buffer (50 mM Tris, pH 7.5, 1 mM dithiothreitol, 5 mM EDTA, 10%
9 glycerol, 1% Triton X-100, 0.1 mM phenylmethylsulfonylfluoride [PMSF] and 0.15 mg/ml
10 lysozyme). The bacterial suspensions were passed through the SLM Aminco French Pressure
11 Cell Press (Beun de Ronde, La Abcoude, The Netherlands) and centrifuged at 20,000×g for
12 15 min at 4 °C. The GST-TP fusion protein was further purified from the supernatant using
13 Glutathione Sepharose 4B (GE Healthcare, Little Chalfont, Buckinghamshire, UK), following
14 the instructions of the manufacturer. Briefly, a 50% glutathione-agarose slurry was added to
15 the bacterial supernatant, incubated for 30 min at 4 °C, and then washed three times with 10
16 bed volumes (7.5 mL) of lysis buffer without lysozyme and PMSF. Bound proteins were
17 eluted in 50 mM Tris (pH 8.0) containing 0.1% Triton X-100 and 10 mM glutathione. Protein
18 content of the purified fractions was determined using Bradford reagent (Sigma, St-Louis,
19 MO).

2.5 TP enzyme assays

20 The conversion of dThd to thymine was measured by high-pressure liquid chromatography
21 (HPLC) analysis. To determine the conversion activity, different amounts (ranging from 20 to
22 0.10 µg/500 µl reaction mixture) of wild-type or mutant D203A TP were initially incubated
23 with 100 µM of dThd in TP-buffer (10 mM Tris-HCl, pH 7.6, 1 mM EDTA, 2 mM
24 KH₂PO₄/K₂HPO₄ and 150 mM NaCl). At different time points (i.e. 0, 15, 30, 60 minutes), 100

1 μL aliquots of the reaction mixtures were withdrawn and heated at 95 °C for 3 min to
2 inactivate the enzyme. DThd was separated from thymine on a reverse-phase RP-8 column
3 (Merck, Darmstadt, Germany) and quantified by high pressure liquid chromatography
4 (HPLC) analysis (Alliance 2690, Waters, Milford, MA). The separation was performed by a
5 linear gradient from 98% buffer B (50 mM NaH_2PO_4 and 5 mM heptane sulfonic acid, pH
6 3.2), to 20% buffer B + 80% acetonitrile (8 min 98% buffer B + 2% acetonitrile; 5 min linear
7 gradient of 98% buffer B + 2% acetonitrile to 20% buffer B + 80% acetonitrile; 10 min 20%
8 buffer B + 80% acetonitrile, followed by equilibration at 98% buffer B + 2% acetonitrile).
9 Retention times of thymine and dThd were respectively 5.1 and 10.8 minutes. UV-based
10 detection was performed at 267 nm.
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23 In the kinetic assays, the K_m and V_{max} of wild-type TP and the mutant D203A TP were
24 determined at varying concentrations of dThd, while the concentration of inorganic phosphate
25 was kept constant at 25 mM. In these experiments, wild-type TP was used at a concentration
26 of 0.5 $\mu\text{g}/500 \mu\text{l}$ reaction mixture in the presence of 5, 3, 2, 1, 0.5 and 0.2 mM dThd, while
27 the mutant D203A TP was tested at a concentration of 10 $\mu\text{g}/500 \mu\text{L}$ reaction mixture in the
28 presence of 40, 30, 20, 15 and 10 mM dThd. Incubation was performed for 15 min. At this
29 time point, the reaction still proceeded linearly.
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40 To evaluate the inhibitory activity of 5'-*O*-tritylinosine, 6AT and 6A5BU several
41 concentrations of these TP inhibitors were added to the reaction mixtures containing 100 μM
42 (for wild-type TP) or 2.5 mM (for mutant D203A TP) of dThd. Wild-type TP and mutant
43 D203A TP were used at a concentration of 0.5 and 10 $\mu\text{g}/500 \mu\text{L}$ reaction mixture,
44 respectively. The mixtures were incubated for 15 minutes. The amount of thymine formed
45 was determined by HPLC analysis as described above.
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3. Results and discussion

3.1 Proposal of a binding mode for KIN59 on human TP

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6 Inspection of the Gly-405–Val-419 loop (which was clipped in the structure of
7
8 Norman et al. [22]), in the crystallographic structure of El Omari et al. [23], revealed an
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10 intricate hydrogen-bonding network, the most relevant part of which is schematically
11
12 represented in Figure 2: (i) the backbone NH of Arg-408 is hydrogen-bonded to a well-
13
14 defined water molecule (W98 in subunit A, designated WAT1) that is fixed in place by two
15
16 additional hydrogen bonds: one with the backbone carbonyl of Gly-149 and another with one
17
18 carboxylate oxygen from Asp-203, a residue that is situated at the terminus of α -helix 9, (ii)
19
20 the same carboxylate oxygen of Asp-203 forms a good hydrogen bond with the backbone NH
21
22 of Gly-149, which is next to Leu-148, a residue whose side-chain provides a hydrophobic
23
24 surface for interaction with the pyrimidine base of both the dThd substrate and competitive
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26 inhibitors such as TPI, (iii) the guanidinium group of Arg-408 establishes two strong
27
28 hydrogen bonds with the carboxylate oxygens of Asp-156 (not shown) and also a water-
29
30 mediated (W80 in subunit B, designated WAT2) hydrogen bond with the other Asp-203
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32 carboxylate oxygen, (iv) the carbonyl groups of the backbones of Asp-203 and Arg-408 are
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34 both accepting a hydrogen bond from a common water molecule (designated WAT3); (v) the
35
36 guanidinium group of Arg-410 is partly exposed to the solvent but also engaged in an
37
38 intersubunit hydrogen bond with the carbonyl group of Ser-65 (not shown); and (vi) the
39
40 backbone CO and NH groups of Ala-411 in one protomer establish two good hydrogen bonds
41
42 with the side-chain carboxamide of Gln-67 in the neighboring subunit (not shown). Thus, it
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44 appears that the Gly-405–Val-419 loop consists of closely interacting amino acids that
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46 contribute to the stabilization of the closed conformation of the enzyme (the only state that
47
48 presents the integral active site) by helping to hold the α and α/β domains together.
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1 When the surroundings of this loop were probed for plausible binding sites for 5'-*O*-
2 tritylinosine using the reported X-ray crystal structure and the automated docking program
3 AutoDock, no convincing solutions were found. 5'-*O*-Tritylinosine is a rather lipophilic
4 compound possessing a bulky trityl moiety for which no electron density emerged in the
5 diffraction maps. However, the molecule clearly helped stabilize the conformation of this loop
6 in the crystal lattice [23]. We reasoned this might be due to ligand (5'-*O*-tritylinosine)
7 expulsion following protein packing and local hydrophobic collapse. We therefore decided to
8 further focus on some prior conformational state(s) of the enzyme for our docking studies.
9 Since open and closed TP conformations are connected by hinge-dependent interdomain
10 motions [33], as is also the case in the related pyrimidine nucleoside phosphorylase (PyNP)
11 [49], we used normal mode analysis (NMA) to reproduce this motion and generate a series of
12 intermediate protein conformers that could help in the identification of a region capable of
13 lodging the hydrophobic and polar substituents of 5'-*O*-tritylinosine.
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30 By following the lowest frequency mode, a cavity was generated in the vicinity of the
31 Gly-405–Val-419 loop that was mainly lined by the side chains of apolar residues Leu-155,
32 Val-166 and Ile-167, and positively charged residues Arg-146 and Arg-408, all of which can
33 provide good anchoring points for the crucial trityl group of 5'-*O*-tritylinosine. Indeed, the
34 automated docking program could find satisfactory docking solutions for 5'-*O*-tritylinosine
35 making use of this hydrophobic cavity. Thus, in the optimal solution, and following complex
36 relaxation using molecular dynamics (MD), the guanidinium of Arg-146 is found stacking on
37 top of one of the phenyl rings, which in turn makes van der Waals contacts with the side
38 chains of Leu-155 and Val-166, whereas another phenyl of the trityl group interacts with the
39 side chains of Val-166 and Ile-167. As regards the remaining part of the inosine nucleoside,
40 the 3' hydroxyl of the ribose is engaged in a hydrogen bond with the carboxamide group of
41 Gln-187 (not shown), whereas the hypoxanthine moiety gets sandwiched between the tip of
42 the Gly-405–Val-419 loop and the C-terminus of helix 9. In this location, the heteroaromatic
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ring partially sits on top of the side-chain methyl of Ala-200 (not shown), N3 is close to Nε of Arg-146 and N1 is oriented towards the carboxylate of Asp-203 and the isopropyl of Val-204 (not shown) (Figure 3).

This docking proposal accounts for the mechanism of non-competitive inhibition, insofar as 5'-*O*-tritylinosine does not make use of the dThd or phosphate binding sites, and is qualitatively consistent with the known structure-activity relationship for this class of 5'-*O*-tritylinosines [50]. Indeed, it is known that one of the phenyl rings in the essential trityl substituent can accept a chlorine atom at the *para* position leading to enhanced potency, and also that incorporation of a methylcyclopropyl or methylcyclohexyl substituent at the N1 position of hypoxanthine results in increased inhibitory activity. The former finding can be explained because the chlorine atom can be held in one of the hydrophobic pockets made up by Leu-155, Val-166, and Ile-167 whereas the latter may originate from improved hydrophobic interactions between the cycloalkyl substituent and the side chain of Val-204.

Given the central role ascribed to Asp-203 in the stabilization of the Gly-405–Val-419 loop over the active-site cleft in the closed conformation, we decided to mutate this residue to alanine in order to assess the impact of this amino acid replacement on both enzyme activity and modulation of catalysis by TP inhibitors.

3.2 Enzymatic activity and kinetic parameters of wild-type and mutant D203A TP

The enzymatic activities of purified recombinant wild-type and mutant D203A TP were compared by using different amounts of enzyme (0.5 and 0.1 μg/500 μl reaction mixture for wild-type TP and 20, 5 and 0.5 μg/500 μl reaction mixture for mutant D203A TP). The enzymes were incubated for 0, 15, 30 and 60 minutes in the presence of 100 μM of dThd (Fig. 4). The mutant D203A TP converted markedly less substrate to thymine, as compared to the corresponding wild-type enzyme. The reaction proceeded linearly for up to at least 60 min at the lower TP concentrations and up to 15 min at the highest enzyme concentrations. Based on

1 the enzyme concentrations used, it was estimated that the mutant D203A TP converts dThd to
2 thymine at least 20-fold less efficiently than wild-type enzyme in the presence of 100 μM
3 dThd (Fig. 4). Thus, the mutation of aspartic acid to alanine at amino acid position 203 of TP
4 resulted in a significant loss of efficiency to perform the phosphorolysis of dThd. Therefore,
5 higher concentrations were used for the mutant D203A enzyme than wild-type TP to obtain
6 reliable enzyme kinetic data that could be compared.
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13 To determine the kinetic parameters for wild-type and mutant D203A TP the enzyme
14 concentrations used were 0.5 $\mu\text{g}/500 \mu\text{l}$ reaction mixture and 10 $\mu\text{g}/500 \mu\text{l}$ reaction mixture,
15 respectively. Under these experimental conditions, the enzyme-catalytic activity proceeded
16 linearly during the incubation (15 min) time. Both enzymes were incubated in the presence of
17 varying concentrations of dThd at a saturating concentration of inorganic phosphate. The K_m
18 value of the mutant D203A TP (9.8 mM) proved to be 23 times higher than that of the wild-
19 type enzyme (K_m : 414 μM), which suggests a markedly diminished affinity of the mutant
20 enzyme for its natural substrate dThd. Moreover, a 2.5-fold reduction in the V_{max} value (1,026
21 nmol/min/ μg for wild-type TP *versus* 400 nmol/min/ μg for mutant D203A TP) indicated that
22 the mutation also affected (albeit to a lesser extent) the rate of dThd phosphorolysis (Table 1).
23 Overall, the phosphorolytic capacity of the mutant D203A TP proved to be ~ 60 -fold lower
24 than that of wild-type TP using dThd as the natural substrate.
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45 ***3.3 Simulation of the transition state of the reaction in the wild-type and mutant D203A TP*** 46 ***enzymes*** 47

48 In light of the preceding results we sought to determine the structural basis for the decrease in
49 catalytic performance of the mutant D203A TP relative to the wild-type enzyme using
50 computational methods. Since the geometry of the phosphorolysis reaction transition state
51 (TS) has been previously characterized in detail [33], unrestrained MD simulations of each
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enzyme in the presence of an active site-bound TS were undertaken to analyze the relative binding affinities in the two complexes.

In the wild-type enzyme the TS appears to be stabilized by a number of interactions involving both polar and apolar active-site residues (Figure 5): (i) the pyrimidine ring gets sandwiched between the side chains of Tyr-199 and Leu-148, on one side, and that of Ile-218 on the other side, and places the methyl group in the vicinity of Val-241 (not shown); (ii) the O2 and O4 of thymine accept hydrogen bonds from the amino group of Lys-221 and the guanidinium group of Arg-202, respectively, whereas the hydroxyl of Ser-217 accepts a hydrogen bond from N3 and the imidazole ring of His-116 makes a bifurcated hydrogen bond with N1 and O2; (iii) the O5' of the deoxyribose accepts a H-bond from the hydroxyl of Thr-151 and donates one to the carbonyl of the backbone of Gly-119 while the backbone NH of Thr-154 hydrogen bonds to O3' of the deoxyribose, which in turn donates another H-bond to the phosphate, precisely to the same oxygen (OP1) that accepts a second H-bond from the hydroxyl of Thr-154; and (iv) the phosphate OP3 accepts two hydrogen bonds, one from the amino group of Lys-115 and another from the hydroxyl of Ser-117 (not shown). This elaborate hydrogen-bonding pattern, which is maintained during the whole length of the MD simulation, stabilizes the TS geometry and facilitates the nucleophilic attack of the phosphate O2P onto the sugar ring atom C1'.

By hydrogen-bonding to the NH of Gly-149, the carboxylate of Asp-203 was thus seen to play a crucial role in facilitating the correct placement of the side-chain of Leu-148. This amino acid residue has been previously shown to be essential for TP activity [7], most likely because it provides a hydrophobic platform for the stacking of both the phenol ring of Tyr-199 (which is fixed in place with the aid of a hydrogen bond from the hydroxyl group to the backbone carbonyl of Arg-146) and the pyrimidine base of the substrate and competitive inhibitors. In fact, the major difference observed within the slightly more open active site in the simulated TS-TP complex for the mutant D203A enzyme involved the positioning of Leu-

148 and when the MM-GBSA method was used to calculate the binding energies of the TS within both enzymes along the simulation period, the binding energy corresponding to the mutant D203A TP was indeed found to be consistently decreased relative to the wild-type protein (Figure 6). Moreover, the loss of binding energy originated in the van der Waals term because the electrostatic and desolvation components were virtually identical (data not shown). Therefore, even though these binding energy values have to be regarded as qualitative rather than quantitative (since entropic effects, which are cumbersome to compute and subject to large numerical errors, were not included in the calculation), the trend is very clear and is in agreement with the experimental findings: the loss of the hydrogen bond between the carboxylate of Asp-203 and the backbone NH of Gly-149 results in a repositioning of the side chain of Leu-148 and in a more open active site, both of which are detrimental for dThd binding, TS stabilization and efficient catalysis.

3.4 Inhibition of wild-type and mutant D203A TP by competitive and non-competitive inhibitors

It has been suggested that mouse liver TP contains more than one allosteric (i.e. thymine binding) site, and that when thymine is bound to one of these sites, the enzyme is inhibited [51]. We found thymine (100 μM) inhibitory to the human TP-catalysed hydrolysis of 100 μM dThd, but thymine had no effect on the inhibitory activity of 5'-*O*-tritylinosine (200 μM and 100 μM) (data not shown), making it unlikely that 5'-*O*-tritylinosine may bind to one of such potential thymine-binding sites.

However, we previously showed that 5'-*O*-tritylinosine inhibits TP in a non-competitive fashion [20]. If Asp-203 is part of a putative allosteric site to which this inhibitor binds or contributes to its formation, one would expect that mutation of this amino acid residue to alanine would affect the inhibitory activity of this compound against TP. To test this hypothesis, wild-type and mutant TP enzymes were incubated for 15 minutes with 100

1 μM dThd in the presence of various concentrations of either the non-competitive inhibitor 5'-
2 *O*-tritylinosine or the known competitive inhibitors 6AT and 6A5BU. As shown in Figure 7,
3 5'-*O*-tritylinosine was completely unable to inhibit the enzymatic activity of the D203A
4 mutant TP regardless of the concentrations used (100, 50 and 25 μM) whereas a
5 concentration-dependent inhibition of the wild-type enzyme was observed (Fig. 7, panel A).
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10 In sharp contrast, the competitive inhibitors 6AT and 6A5BU, which have been shown to
11 interact with the dThd-binding site of TP [20], were able to dose-dependently decrease the
12 enzymatic activity of wild-type and mutant D203A TP to a similar extent (Fig. 7, panels B
13 and C) (6AT and 6A5BU inhibitor concentrations used: 50, 25 and 5 μM for wild-type TP and
14 1,250, 625 and 125 μM for mutant D203A TP in the presence of 100 μM dThd for wild-type
15 TP and 2,500 μM dThd for mutant D203A TP). These findings are compatible with the view
16 that binding of dThd, 6AT and 6A5BU to the active site is similarly compromised in the
17 mutant enzyme relative to the wild-type TP whereas the putative allosteric site for 5'-*O*-
18 tritylinosine binding is no longer present on the mutant D203A TP due to loop disruption.
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35 **3.5 Concluding remarks**

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37 Earlier site-directed mutagenesis studies on TP have confirmed the importance for catalysis of
38 Lys-115, Leu-148, Arg-202, Ser-217, and Lys-221 [7, 8]. All of these are active-site residues
39 directly involved in the stabilization of the substrates (dThd and phosphate) and the TS, as
40 explained above. Asp-203, on the other hand, is positioned at the end of an α -helix and
41 extends its side chain towards a solvent region comprised between the glycine-rich 144–153
42 loop and the more mobile arginine-rich 406–415 loop. In this location, the Asp-203
43 carboxylate can accept two hydrogen bonds: one from the backbone NH of Gly-149 in the
44 former loop (Fig. 2 and 5) and a second one from a water molecule that is, in turn, hydrogen
45 bonded to the guanidinium of Arg-408 in the latter loop (Fig. 2). In addition, the Asp-203
46 carbonyl can also make a water-mediated hydrogen bond with the carbonyl of Arg-408 (Fig.
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2). It thus appears that Asp-203 plays a crucial role in loop stabilization, and this action is important for both catalysis and inhibition by the non-competitive inhibitor 5'-*O*-tritylinosine.

5'-*O*-Tritylinosine has been described previously as an inhibitor of both human and bacterial TP enzymes [20, 21]. Given the non-competitive nature of this inhibition, this purine riboside derivative was not expected to interact directly with either the pyrimidine nucleoside-binding or phosphate-binding site of the enzyme. On the contrary, this novel inhibitory mechanism was indicative of the possible existence of a yet unidentified allosteric site in the enzyme. Nonetheless, a crystal structure of human TP produced in the presence of 5'-*O*-tritylinosine revealed no traces of this inhibitor neither in the active site, which was occupied by the reaction product, thymine, nor anywhere else.

By using a combination of NMA, docking and MD simulations methods, a location for this binding site is now proposed that relies on the loop stabilization brought about by Asp-203. The site-directed mutant D203A TP, which was constructed to assess the importance of this residue, displayed a ~60-fold reduction in phosphorolytic capacity (V_{\max}/K_m) relative to the wild-type enzyme. Importantly, both wild-type and mutant TP were inhibited to a similar extent by the competitive inhibitors 6AT and 6A5BU whereas the non-competitive inhibitor KIN59 lost its inhibitory activity against the mutant D203A enzyme. These data thus support a role for Asp-203 in loop stabilization, which is important for catalytic activity, as well as for creating a rather unique binding site for 5'-*O*-tritylinosine. A multiple sequence alignment of human, mouse, rat and *E. coli* TP (Figure 8) shows that Asp-203 is highly conserved during evolution, which reinforces the idea that this amino acid may indeed be crucial for the biological activities of TP. We previously hypothesized that this putative allosteric site might be involved in the angiogenic activity of TP [20, 21]. Therefore, further investigations on the angiogenic and anti-apoptotic properties of the mutant D203A enzyme would be warranted.

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FIGURE LEGENDS

1
2 Figure 1. Chemical structures of relevant TP inhibitors.
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10 Figure 2. Scheme depicting the most relevant hydrogen bonds that contribute to loop
11 stabilization in the vicinity of Asp-203. For simplicity, only the atoms of one water molecule
12 are shown.
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22 Figure 3. Proposed binding site for 5'-*O*-tritylinosine (previously designated KIN59 [20]) in
23 human TP. (a) Ribbon representation of the enzyme with protein residues enveloped by a
24 semitransparent solvent-accessible surface. Carbon atoms of 5'-*O*-tritylinosine are colored in
25 magenta. (b) Enlarged view of the framed area shown in (a). Some of the protein residues
26 relevant to the discussion have been labeled, and their side chains are shown as sticks.
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39 Figure 4. Conversion of thymidine to thymine by wild-type and mutant D203A TP as a
40 function of time and at different enzyme concentrations. The dThd concentration was 100
41 μM .
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51 Figure 5. Detail of the active site of wild-type TP containing the transition state of the reaction
52 (black dashes represent bonds that are being broken/created). The most relevant residues have
53 been labeled and hydrogen bonds are shown as yellow dashed lines.
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Figure 6. Time evolution of the overall interaction energy between the transition state of the reaction and either wild-type (blue) or D203A (magenta) human TP during the molecular dynamics simulations.

Figure 7. Effect of the inhibitors 5'-O-tritylinosine (panel A), 6AT (panel B) and 6A5BU (panel C) on the enzymatic activity of TP. Wild-type and mutant D203A TP were incubated at 37 °C for 15 minutes at an enzyme concentration of 0.5 and 10 µg/500 µL reaction mixture, respectively. The dThd concentration in the assays were 100 µM (WT TP) and 2,500 µM (mutant D203A TP) to compensate for the different Km values of both enzymes. The conversion of dThd to thymine in the presence of different amounts of inhibitors was determined by HPLC analysis. Values are presented as the mean of at least three different independent experiments ± standard error.

Figure 8. ClustalW multiple sequence alignment of the amino acid sequences of mouse, rat, human and *E. coli* TP. (*) identical residues, (.) and (:) conserved and semi-conserved substitutions, respectively. Asp-203 is shown in red, amino acids participating in substrate binding are shown in blue, and residues proposed to be involved in 5'-O-tritylinosine (previously designated KIN59 [20]) are shown in green.

TABLES

Table 1. Kinetic parameters of human wild-type and mutant D203A TP

	K_m^a (μM)	V_{\max}^b ($\mu\text{mol/mg}$ protein/min)	V_{\max}/K_m^c
Wild-type TP	414 ± 96	1026 ± 460	2.48
D203A mutant TP	9828 ± 3200	400 ± 136	0.041

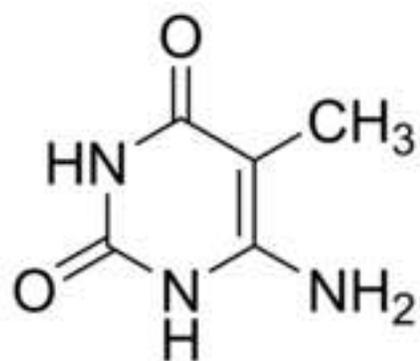
Enzyme activity is expressed as the amount of dThd converted to thymine [$\mu\text{mol/mg}$ protein/min].

^a K_m is the Michaelis-Menten constant of the enzyme for dThd.

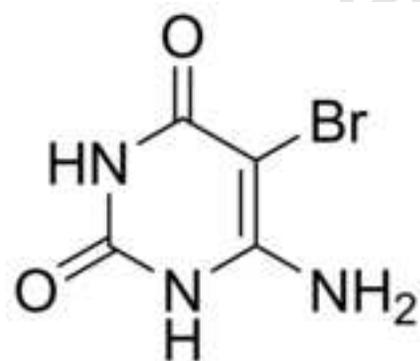
^b V_{\max} is the maximum velocity of the enzyme reaction.

^c V_{\max}/K_m ratio or catalytic phosphorylytic capacity of the enzyme.

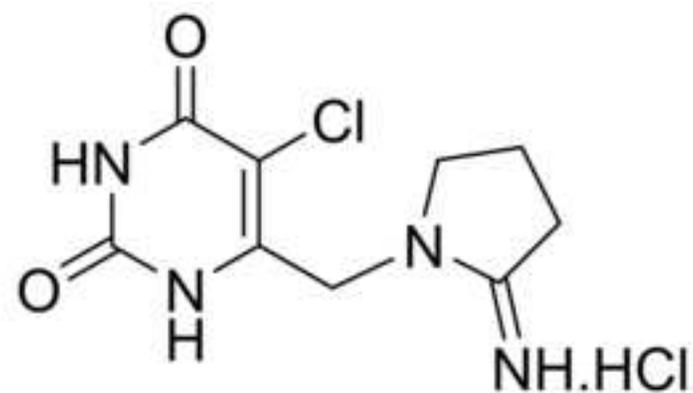
Values are presented as means \pm standard deviation of at least three independent experiments.



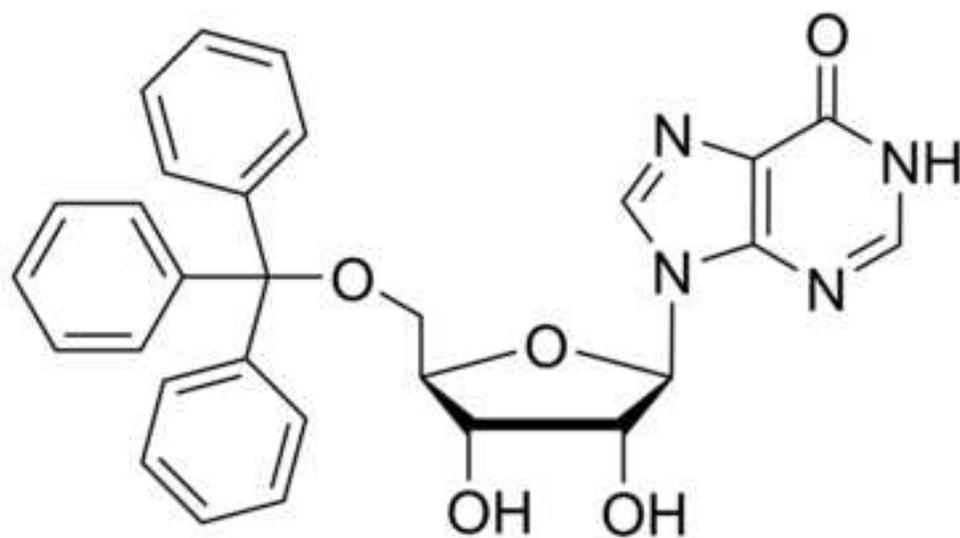
6AT



6A5BU

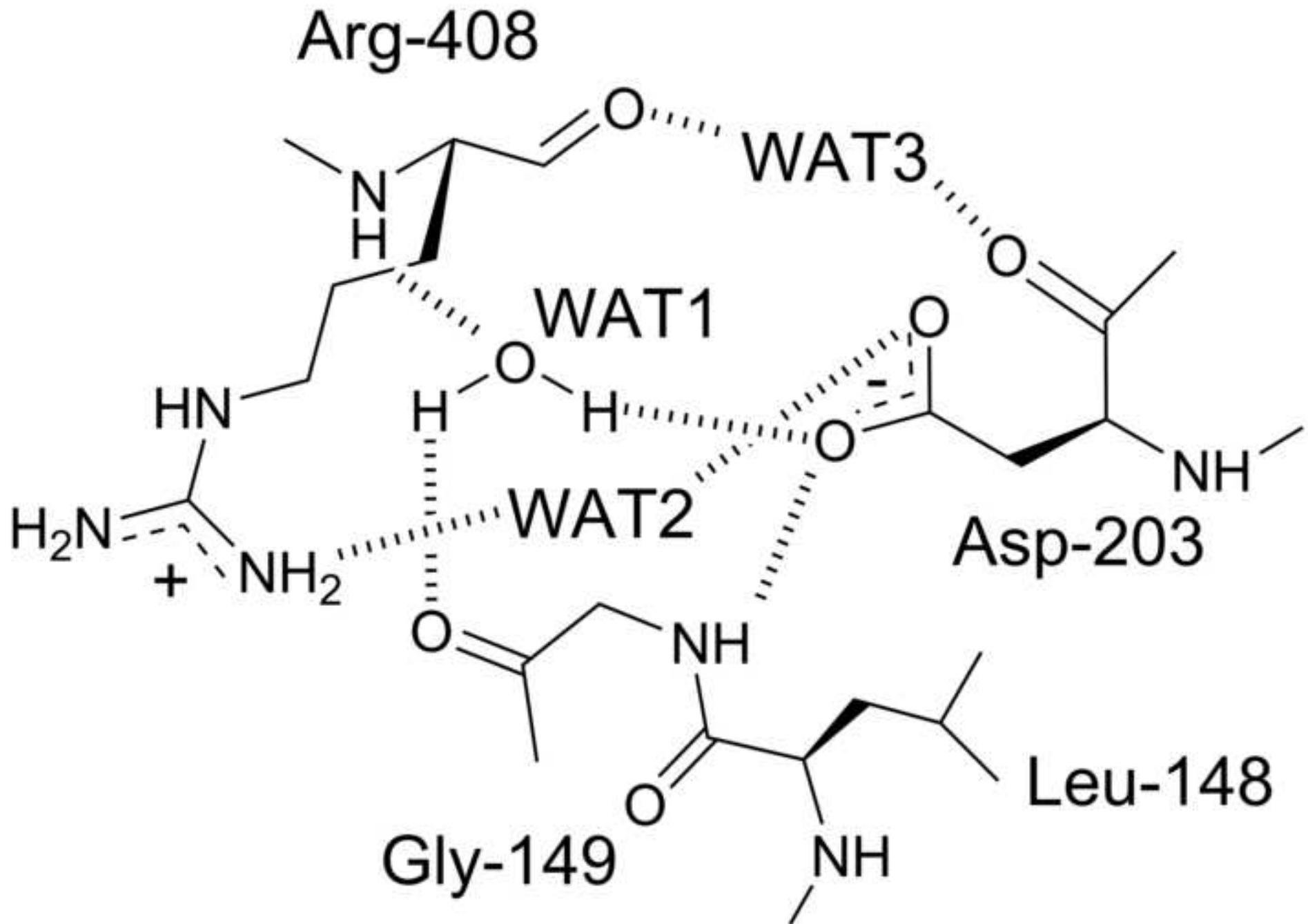


TPI



5'-O-tritylinosine (KIN59)

Figure 2



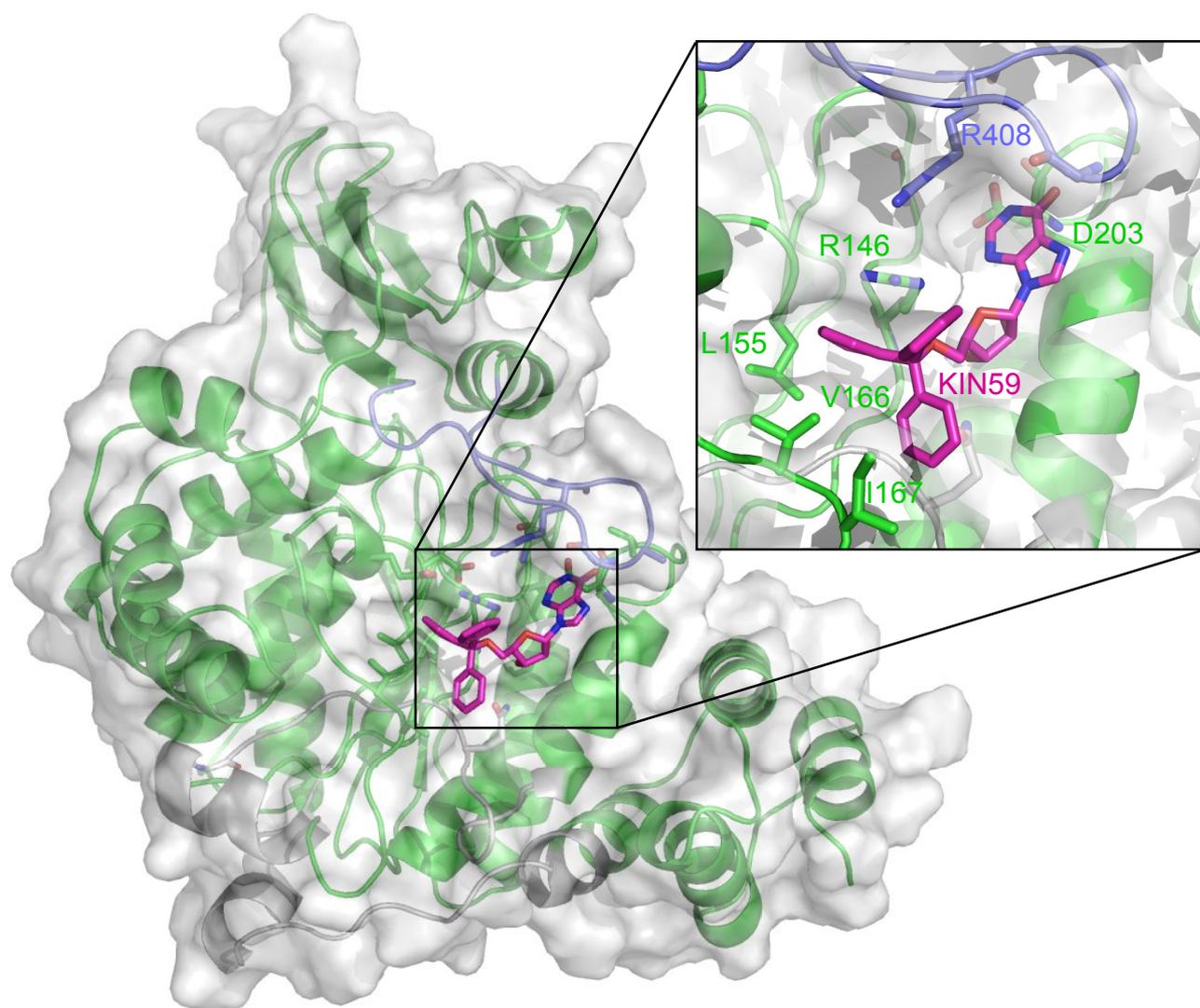


Fig. 3

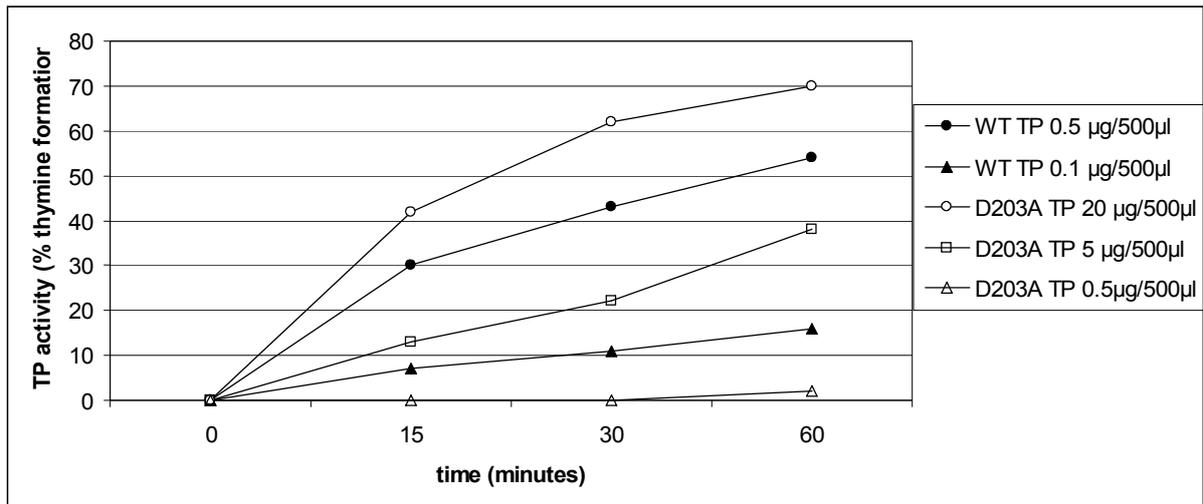


Fig.4

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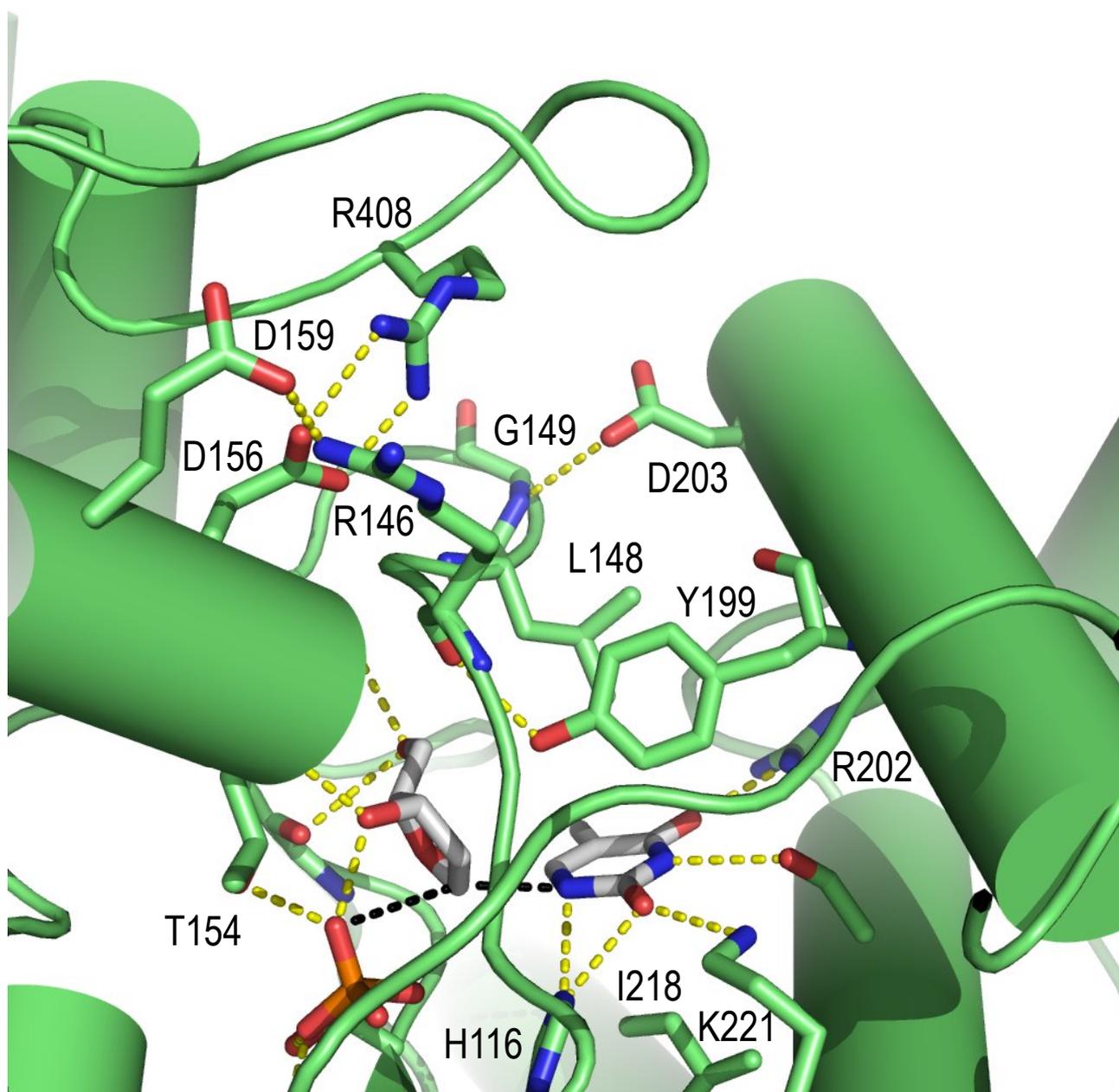


Fig. 5

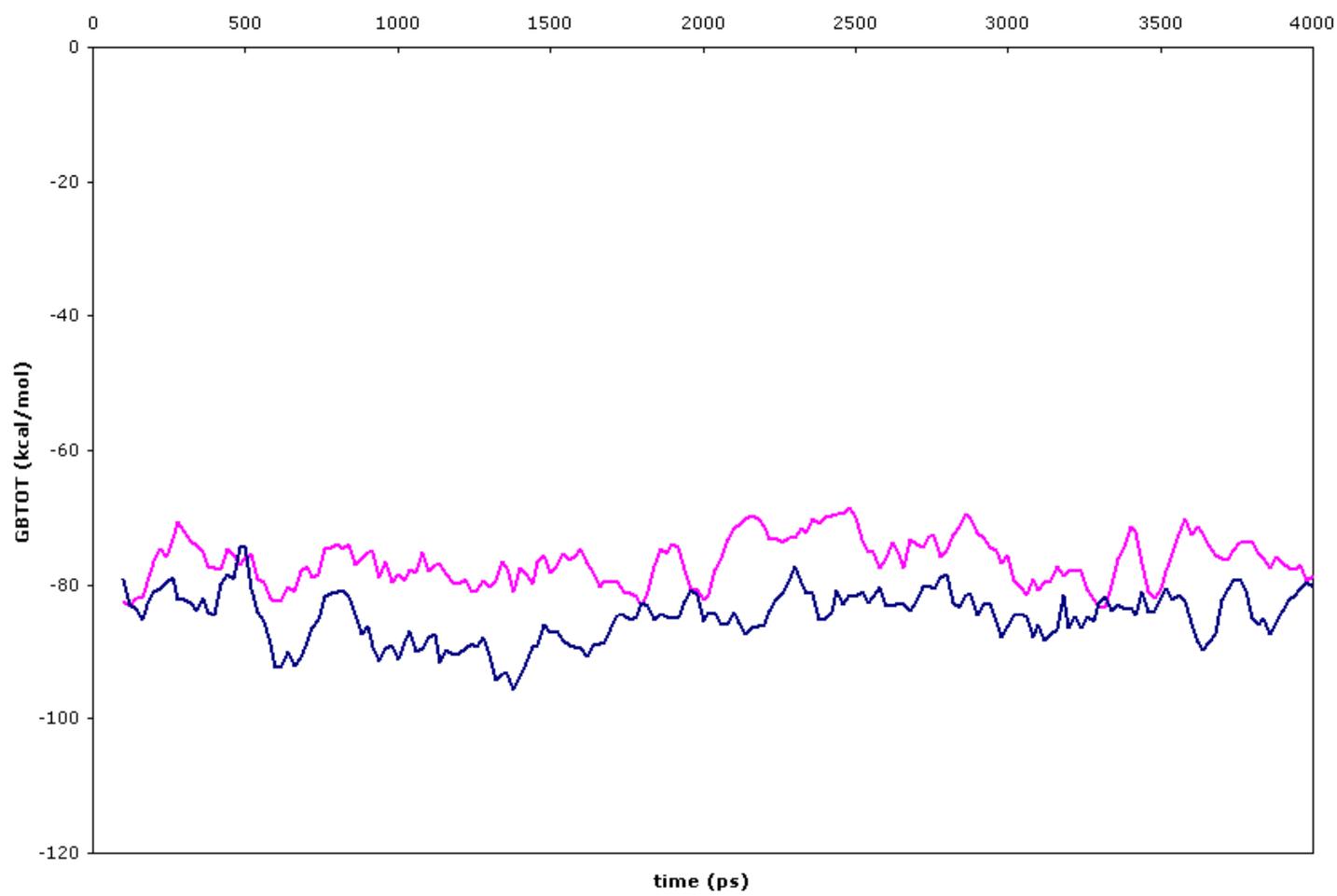


Fig.6

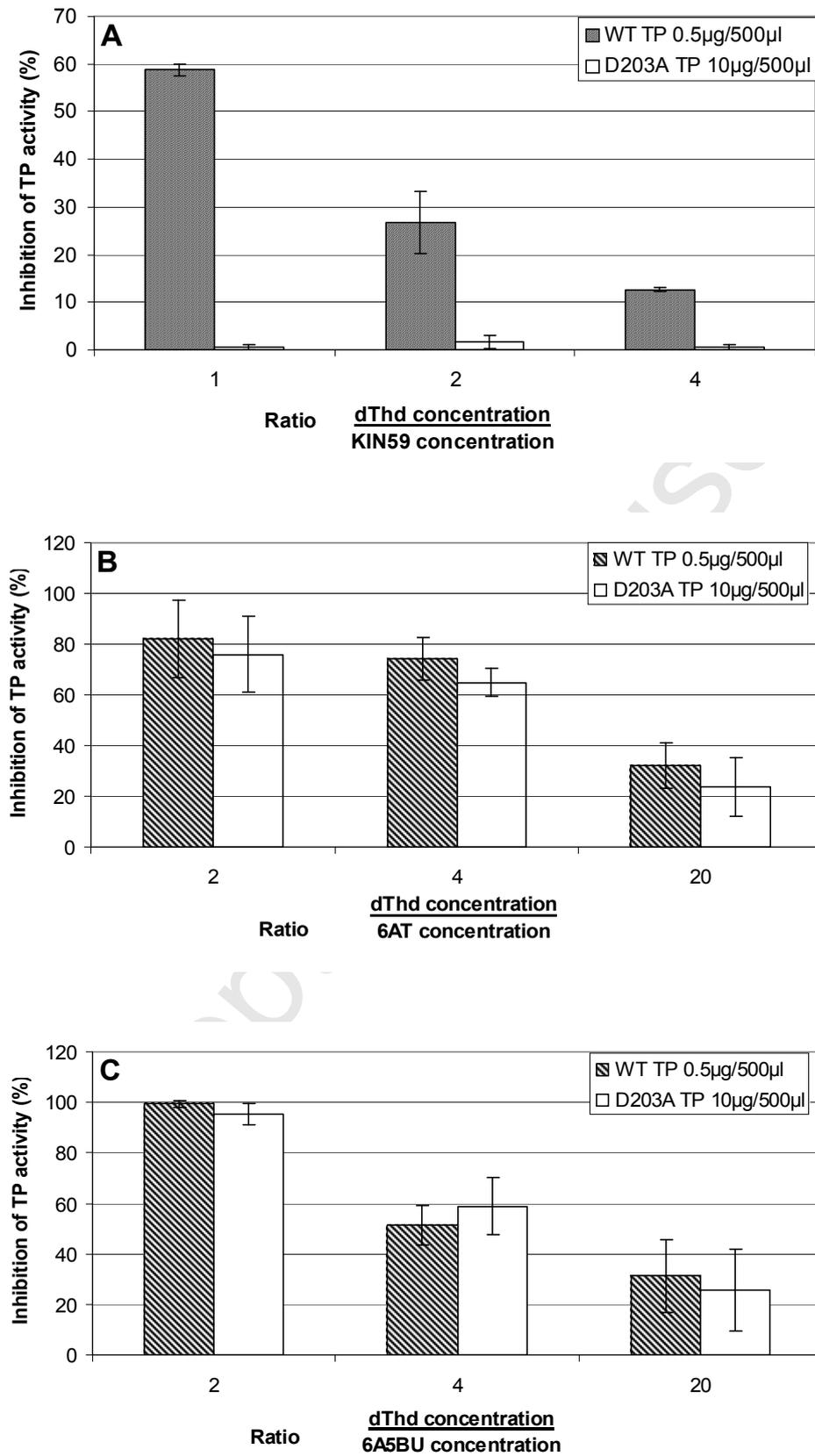
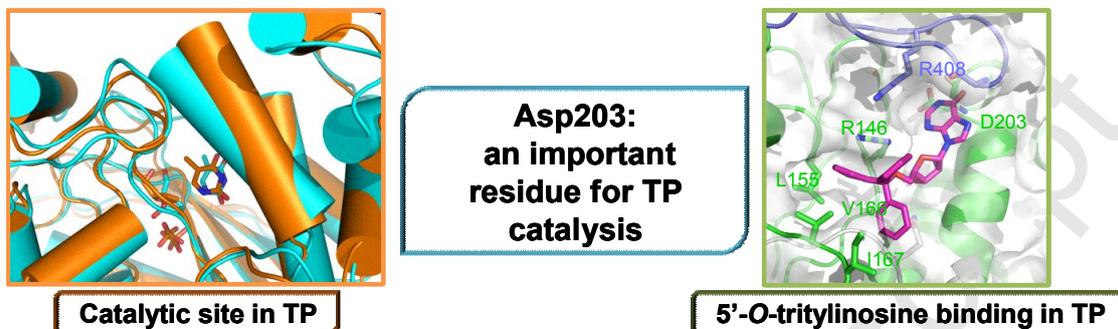


Fig. 7

Graphical abstract



The search for the binding site of the allosteric inhibitor 5'-*O*-tritylinosine has allowed the identification of Asp203 as an important residue for catalysis in human thymidine phosphorylase.