Discovery of selective inhibitors of tyrosyl-DNA phosphodiesterase 2 by targeting the enzyme DNA-binding cleft

Bradley R Kossmann^a, Monica Abdelmalak^b, Sophia Lopez^b, Gabrielle Tender^b, Chunli Yan^a, Yves Pommier^b, Christophe Marchand^{b*} and Ivaylo Ivanov^{a*}

^aDepartment of Chemistry, Center for Diagnostics and Therapeutics, Georgia State University, Atlanta, Georgia 30302, United States

^bDevelopmental Therapeutics Branch, Center for Cancer Research, National Cancer Institute, National Institutes of Health, Bethesda, Maryland, 20892, United States

ARTICLE INFO

Article history:

Received

Revised

Accepted

Available online

ABSTRACT

Tyrosyl-DNA phosphodiesterase 2 (TDP2) processes protein/DNA adducts resulting from abortive DNA topoisomerase II (Top2) activity. TDP2 inhibition could provide synergism with the Top2 poison class of chemotherapeutics. By virtual screening of the NCI diversity small molecule database, we identified selective TDP2 inhibitors and experimentally verified their selective inhibitory activity. Three inhibitors exhibited low-micromolar IC₅₀ values. Molecular dynamics simulations revealed a common binding mode for these inhibitors, involving association to the TDP2 DNA-binding cleft. MM-PBSA per-residue energy decomposition identified important interactions of the compounds with specific TDP2 residues. These interactions could provide new avenues for synthetic optimization of these scaffolds.

Keywords: Tyrosyl-DNA phosphodiesterase 2; TDP2; inhibitor; virtual screening; rational drug design

2009 Elsevier Ltd. All rights reserved.

Tyrosyl-DNA phosphodiesterase (TDP) activity is necessary to cleave the tyrosyl-DNA linkage between a trapped topoisomerase and its substrate DNA¹. In humans, there are two known TDPs: TDP1 cleaves 3' type-IB topoisomerase-DNA linkages², while TDP2 cleaves 5' type-II topoisomerase-DNA linkages^{3,4}.

Topoisomerase II (Top2) poisons act by trapping Top2 on its DNA substrate, causing the normally transient Top2-mediated double strand breaks to become permanent, resulting in cell death⁵. For this reason, Top2 poisons are widely used cancer therapeutics. TDP2 activity reduces the efficacy of Top2 poisons and is therefore an attractive anticancer drug target, with TDP2 deficient cells exhibiting extreme sensitivity to Top2 poisons^{3,6}. The viability of TDP2 knockout mice indicates that TDP2 inhibition is theoretically possible without unacceptable side effects⁶. A TDP2 inhibitor could have great potential for synergistic effects when used in combination with Top2 poisons and could greatly increase the efficacy of such treatments.

Small-angle X-ray scattering analysis shows that TDP2 consists of a ~110-residue, disordered N-terminal domain and a 255-residue, globular catalytic domain^{7, 8}. Only the catalytic domain is necessary for phosphodiesterase activity, while the N-terminal tail is thought to interact with cellular signaling machinery. Although no structures exist for human TDP2 (hTDP2), multiple crystal structures of TDP2 have been solved including *C. elegans* (cTDP2)⁸, *D. rerio* (zTDP2)⁸ and *M. musculus* (mTDP2)⁷. mTDP2 serves as an excellent structural

homologue to hTDP2, with the variants' catalytic domains sharing 78% sequence identity and nearly 100% sequence similarity (Supplementary Figure S1). A structure of mTDP2* bound to a substrate analog (PDB accession code 4GZ1) shows a short DNA-binding cleft, contacting 3 DNA phosphates, leading directly into the active site (Supplementary Figure S2). Kinetic studies indicate that mammalian TDP2 is highly specific for 5'-tyrosine overhangs, as opposed to 3' overhangs, blunt ends or other adducts, with the exception of p-nitrophenol, a compound frequently used as a DNA adduct in screening assays^{7,9}. Interestingly, TDP2 shares very similar active site geometry and catalytic mechanism with the base excision repair enzyme apurinic/apyrimidinic endonuclease 1 (APE1), along with 14% sequence identity and 30% sequence similarity⁸. Despite these similarities, TDP2 does not show endonuclease activity. From mutational and structural data, Schellenberg, et al. proposed that hydrophobic contacts made to W307 and F325 by the substrate DNA backbone serve as the basis for the specificity toward 5'-tyrosine overhangs by forming favorable Van der Waals interactions with the substrate deoxyribose ring⁷. Crystal structures of the *D. rerio* and *C. elegans* TDP2 homologues in complex with substrate DNA provide further evidence that hydrophobic contacts are primarily responsible for substrate binding⁸. Binding of a 3'-tyrosine substrate would juxtapose a

* Corresponding author.

+1(404) 413-5529 iivanov@gsu.edu (I. Ivanov)

+1(301) 435-2463 marchanc@mail.nih.gov (C. Marchand)

phosphate group in place of a ribose ring, unfavorably forcing a negatively charged group to contact hydrophobic sidechains. Moreover, the mutation of certain residues lining the DNA-binding cleft greatly alters catalytic activity in hTDP2. Of note are R231, R266, W297 and F315, all of which are important to the activity of TDP2 on 4-nucleotide overhang 5'-tyrosine substrate DNA, with mutations being very deleterious to catalysis⁷.

In contrast to TDP2, TDP1 cleaves 3' adducts and is capable of acting on a relatively broad range of substrates¹⁰. TDP1 also displays some activity against 5'-tyrosine overhangs, causing specificity overlap with TDP2^{11,12}. Designing a TDP2 inhibitor is complicated by the similar catalytic activity and substrate characteristics of TDP1. A potential inhibitor must be strongly selective for TDP2 to limit binding competition by TDP1. Because the TDP2 binding cleft may hold the key for TDP2 substrate specificity, it presents a promising region for targeting inhibitors that do not have activity against TDP1.

The discovery of selective TDP2 inhibitors based on toxoflavin and deazaflavin scaffolds has recently been reported, in which the authors began the inhibitor search with a high-throughput *in vitro* screening of 100,000 compounds¹³. Small molecule docking of this scaffold shows that inhibitors of this type are likely to bind in the active site of TDP2, directly blocking catalysis. While these compounds show promise, they exhibit slight inhibition of TDP1 at 100 μ M. In addition, toxoflavins and deazaflavins also have undesirable characteristics for drug scaffolds. Toxoflavins are susceptible to redox activity and deazaflavins have poor cell permeability.

The difficulty in predicting the binding selectivity of a compound necessitates a high-throughput lead screening and optimization protocol. To this end, we have carried out an inhibitor discovery protocol (Figure 1) to identify selective TDP2 inhibitors. Our protocol exploits the large scale docking of 11,000 compounds from the 250,000-compound Open National Cancer Institute (NCI) Database¹⁴ followed by virtual screening (VS) and *in vitro* assays using whole cell extract (WCE). With this protocol, we have discovered three potent and selective small molecule inhibitors of TDP2. Results from molecular docking, molecular dynamics (MD), molecular mechanics – Poisson-Boltzmann surface area (MM-PBSA) calculations, biochemical and kinetics experiments provide evidence that our inhibitors bind in the DNA-binding cleft of TDP2, effectively blocking substrate binding and catalysis. The scaffolds of these inhibitors have the potential to be further optimized by targeting specific contacts made to residues in the TDP2 binding cleft, increasing both potency and selectivity for TDP2.

Compound 21 (Supplementary Figure S3), an APE1 inhibitor provided by Dr. Neamati, was chosen to initiate our screening protocol to identify hTDP2-selective inhibitors because it inhibited hTDP2 with an IC₅₀ of 12, 20 μ M (n=2) without affecting hTDP1 up to 200 μ M (data not shown). To predict how a ligand will bind to hTDP2, a homology model of the catalytic domain was constructed from the substrate analog-bound mTDP2 structure (PDB accession code 4GZ1) using the program Modeller¹⁵. Although the homologues have extraordinary sequence similarity (Supplementary Figure S1), and therefore nearly identical secondary structure, minor alterations to the surface of the enzyme can influence preferred ligand binding positions. Compound 21 was first docked into the hTDP2 homology model (Supplementary Figure S3). Docking was performed with the program AutoDock4¹⁶, after preparing a ligand/protein atom-pair interaction grid with AutoGrid4 within the search area. The search area was comprised of both the DNA

Figure 1. Flowchart overview of our TDP2 inhibitor discovery process.

binding cleft and active site, the two regions of the enzyme known to directly impact catalytic activity. The three best-scoring docking poses that were distinct from the others with respect to ligand conformation and ligand-protein contacts were chosen for further analysis. Each pose was simulated for 30ns in MD, using the NAMD2.9 package¹⁷. The MD trajectory which resulted in the most stable binding pose was clustered with respect to the root mean square deviation (rmsd) of the atomic coordinates of the small molecule using the hierarchical clustering algorithm ($\epsilon=10.0$), as implemented in the ptraj¹⁸ MD analysis program. The representative frame from the most populated cluster was then used for VS. VS against this dominant conformation was performed as follows: sets of fifty random conformers for each compound in the NCI database were first constructed using the Openeye Software program Omega¹⁹ and screened for structural and electrostatic similarity to the dominant MD conformer using Openeye's ROCS²⁰ and EON²¹ programs, respectively. The 11,000 top hits from the VS, as determined by averaging structural and electrostatic Tanimoto overlaps with the lead compound, were then docked using the AutoDock4 program into our hTDP2 homology model.

From the top 500 docking hits, a set of 178 was selected for optimal structural diversity. From these 178 compounds, 95 were available and were tested against recombinant human TDP2 (hTDP2) at a single concentration of 1 mM (Figure 1). 48 active compounds were further tested at a single concentration of 100 μ M against hTDP2. 11 TDP2-selective inhibitors were further evaluated in dose response against both recombinant human TDP2 and TDP1 (hTDP2 and hTDP1) enzymes (Figure 1). Four TDP2-selective low micromolar inhibitors i.e. not inhibiting hTDP1 up to 111 μ M, were checked by LC/MS (Figure S5) to assess for their presence in the vial and evaluated for their ability to inhibit endogenous hTDP2 within whole cell extracts from human TDP2-complemented DT40 chicken cells (hTDP2 WCE).

Figure 2. Inhibition of human recombinant TDP2 (hTDP2) and endogenous human TDP2 from whole cell extracts (hTDP2 WCE) by NSC375976, NSC114532, and NSC3198. Concentrations of compounds are 0.5, 1.4, 4.1, 12.3, 37 and 111 μ M. Concentrations of the positive control Compound 1²² are 0.005, 0.017, 0.05, 0.15, 0.46, 1.4 μ M.

The TDP2-selective inhibitors, NSC375976, NSC114532 and NSC3198 were found to inhibit hTDP2 with IC₅₀ values of 3.5 ± 1.4 , 4.1 ± 0.9 and 9.3μ M, respectively while all being inactive against recombinant hTDP1 up to 111 μ M (Figure 2 and Table 1). All 3 compounds show robustness while being exposed to an increased complexity reaction mixture because they retain low micromolar activity against whole cell extracts (WCE) containing endogenous hTDP2 (Figure 2).

Table 1. IC₅₀ values against human (*H. sapiens*, hTDP2), mouse (*M. musculus*, mTDP2) and zebrafish (*D. rerio*, zTDP2) TDP2 enzymes and against human TDP1 (hTDP1).

Compound	Structure	IC ₅₀ (μ M)			
		hTDP2	mTDP2	zTDP2	hTDP1
NSC375976		3.5 ± 1.4 (n=4)	15 (n=1)	5.2 (n=1)	>111
NSC114532		4.1 ± 0.9 (n=4)	>111	15 (n=1)	>111
NSC3198		9.0, 9.5 (n=2)	>111	20 (n=1)	>111

The inhibition of TDP2 by these 3 compounds was then assessed across species by comparing their potencies against human (*H. sapiens*, hTDP2), mouse (*M. musculus*, mTDP2) and zebrafish (*D. rerio*, zTDP2) TDP2 enzymes (Figure 3). Only NSC375976 inhibited efficiently all three enzymes with IC₅₀ values of 15 and 5.2 μ M for mTdp2 and zTDP2, respectively (Figure 3 and Table 1). NSC114532 and NSC3198 both inhibited zTDP2 with IC₅₀ values of 15 and 20 μ M, respectively but interestingly, mTDP2 was totally resistant to both compounds up to 111 μ M (Figure 3 and Table 1). These results suggest that the TDP2 binding site for both NSC114532 and NSC3198 is not conserved in the mouse enzyme similarly to what was recently observed for the first reported selective TDP2 inhibitor Compound 1 (Figure 2)²².

A potential concern with the chemotype of NSC114532 and NSC3198 is the presence of a disulfide bond in the center of the structure (Table 1). The presence of a high concentration of DTT (1mM) in the reaction buffer could therefore potentially cleave the molecule in two. To study the potential impact of the presence of a reducing agent on the stability of NSC114532 and

Figure 3. Inhibition of human (*H. sapiens*, hTDP2), mouse (*M. musculus*, mTDP2) and zebrafish (*D. rerio*, zTDP2) TDP2 enzymes by NSC375976, NSC114532, and NSC3198. Concentrations of compounds are 0.5, 1.4, 4.1, 12.3, 37 and 111 μ M.

NSC3198, we tested their ability to inhibit hTDP2 in the presence or absence of DTT. We did not observe any difference in the inhibition of hTDP2 by these two compounds under these conditions (Figure S6), suggesting that both compounds are not affected by the composition of the reaction buffer.

Molecular mechanics Poisson-Boltzmann surface area (MM-PBSA) calculations²³, as implemented in the AmberTools14 analysis suite²⁴, were performed on NSC375976, NSC114532 and NSC3198. MM-PBSA is a thermodynamic cycle based method, whereby the solvation enthalpies of the complex ($\Delta H_{complex}$), receptor ($\Delta H_{receptor}$) and ligand (ΔH_{ligand}) are estimated independently, then subtracted to obtain the binding enthalpy ($\Delta H_{binding}$) via:

$$\Delta H_{binding} = \Delta H_{complex} - \Delta H_{receptor} - \Delta H_{ligand}$$

From MM-PBSA, NSC114532 and NSC3198 were determined to have binding enthalpies of -18 ± 12 and -30 ± 7.9 kcal/mol, respectively. NSC114532 and NSC3198 have essentially similar binding enthalpies, as expected from their structural similarity and similar IC₅₀ values. NSC375976 has a higher binding enthalpy, 0.1 ± 4.7 kcal/mol, than both NSC114532 and NSC3198, although it possesses the lowest IC₅₀ of the group. This result is less surprising given the dissimilarity of the ligands. It is also possible that the IC₅₀ values do not correlate perfectly with the binding enthalpies due to ligand interactions with other species in WCE. To compare the binding modes of the three inhibitors, we have performed MM-PBSA per-residue energy decomposition²⁵. This method calculates the enthalpy of each residue's non-bonded interactions with the ligand, highlighting residue-ligand contacts that are essential to ligand binding, as well as those that are detrimental to it.

NSC114532 and NSC3198 are symmetrical molecules around the disulfide bond. Because these two compounds are very close structural analogs, their modes of binding are strikingly similar, with the majority of the binding enthalpy coming from hydrogen bonding and electrostatic interactions between the ligands' carboxy groups and R231 and R266 (Figure 4A and 4B).

NSC114532 and NSC3198 each also form favorable hydrophobic interactions with residues W297, L313 and F315 that lie in the DNA binding cleft (Figure 4D and 4E). NSC375976 does not share the basic chemical scaffold of NSC114532 and NSC3198, yet binds in nearly the same position (Figure 4C and 4F). In contrast to NSC114532 and NSC3198, NSC375976 binds predominantly to hydrophobic residues, with W297, L313 and F315 each contributing ~ 1 kcal/mol in enthalpy. The NSC375976 sulfonate group is not able to form favorable electrostatic interactions with R231 and R266, with the adjacent phenyl group clashing slightly with R266. All three

Figure 4. Binding poses and per-residue energy decomposition for residues important to binding. Per-residue energy decomposition results for (A) NSC114532, (B) NSC3198 and (C) NSC375976. Dominant binding poses of (D) NSC114532, (E) NSC3198 and (F) NSC375976.

inhibitors form favorable van der Waals interactions with the T230 methyl group. The most unfavorable interaction between these inhibitors and hTDP2 is with D350. Each inhibitor places hydrophobic functional groups near the charged D350 side chain, incurring an enthalpic penalty relative to a solvent-exposed D350.

Residues R231, R266, W297, L313 and F315 have all been shown via mutational studies to interact favorably with the DNA substrate, with W297 and F315 having been proposed to form the basis of TDP2 selectivity towards 5'-tyrosine adducts⁷. Clearly, ligand binding in this region of the protein precludes substrate recognition by TDP2, thus inhibiting catalytic activity. Forming favorable contacts with the residues that select for 5' adduct binding is likely to be the cause of these inhibitors' selectivity towards TDP2, as 5' adducts have been shown to be very poor substrates for TDP1. It should be noted that the two distinct molecular scaffolds represented by NSC114532, NSC3198 and NSC375976 both take advantage of the same structural characteristics of the DNA-binding groove, leaving open the possibility of the discovery of other unique scaffolds that have similar binding modes.

To further probe the basis of selectivity of NSC114532, NSC3198 and NSC375976, we docked each into hTDP1, using the same methodology as with hTDP2, again including both the active site and DNA-binding groove in the search area. The top docking poses from each compound (Supplementary Figure S4) identify the hTDP1 active site as the primary binding target. These results do not indicate that NSC114532, NSC3198 or NSC375976 bind hTDP1. Rather, they serve simply as an indication that, when forced to interact with hTDP1 *in silico*, these inhibitors preferentially interact with the active site. This provides another line of evidence supporting the hypothesis that DNA-binding cleft interactions dictate substrate specificity between hTDP1 and hTDP2 and that our inhibitors select for hTDP2 over hTDP1 based on many of the same interactions. The docking poses of NSC114532 and NSC3198 in hTDP1 are again similar, with both compounds forming hydrogen bonds with S399 and K495. NSC3198 stacks one phenyl ring against H263, while leaving approximately half of the molecule solvent-exposed with no notable favorable interactions with the enzyme.

NSC114532 forms no notable favorable stacking interactions with either phenyl ring and forms an electrostatic clash between a carboxylate group and G538. NSC375976 remains largely solvent-exposed by packing against one wall of the active site while reaching slightly into the active site pocket. The only favorable hydrophobic interaction NSC375976 forms is between the phenylsulfonate ring and the aliphatic portion of the S459 sidechain, while the only hydrogen bond to the enzyme is made between the sulfonate group and H263. The combination of unfavorable interactions and inability to take advantage of nonspecific Van der Waals interactions explains these ligands' inability to bind to hTDP1 and, therefore, their selectivity for hTDP2.

NSC114532, NSC3198 and NSC375976 can each be tailored to increase binding affinity to hTDP2 by improving electrostatic complementarity to the binding cleft to enhance binding enthalpy while leaving the Van der Waals contacts in place to maintain selectivity. It is important to note that NSC3198 and NSC114532 form very favorable electrostatic interactions with R231 and R266, residues that are important to the catalytic function of hTDP2. In the future development of these scaffolds, care must be taken to preserve these interactions, meaning that the carboxy groups, as well as the 4-atom linker that separates them, appear to be absolutely essential to effective binding. NSC375976 can likely form very favorable interactions between its sulfonate group and R231 and R266, if the sulfonate group is brought more proximal to the center of the compound. NSC375976 holds the lowest IC₅₀ of the compounds studied, taking advantage of this potentially very favorable interaction may greatly improve binding characteristics.

Acknowledgments

This work was supported by a National Science Foundation CAREER award MCB-1149521 and National Institute of Health grant R01GM110387. This work was also supported in part by the Intramural Research Program of the NIH, National Cancer Institute, Center for Cancer Research (Z01 BC 006161). Computational resources were provided in part by a National Science Foundation XSEDE allocation CHE110042 and through an allocation at NERSC supported by the U.S. Department of

Energy Office of Science contract DE-AC02-05CH11231. The authors wish to thank Drs. Gary Pauly and Joel Schneider from the Chemical Biology Laboratory, CCR, NCI for their support in purity analysis.

Supplementary data

Supplementary data associated with this article can be found in the online version at

References and notes

- Pommier, Y.; Huang, S.Y.; Gao, R.; Das, B.B.; Murai, J., Marchand, C. *DNA Repair (Amst)*. **2014**, 19, 114-129.
- Liu, C.; Pouliot, J.J., Nash, H.A. *Proc. Natl. Acad. Sci. U.S.A.* **2002**, 99, 14970-14975.
- Zeng, Z.; Cortes-Ledesma, F.; El Khamisy, S.F., Caldecott, K.W. *J. Biol. Chem.* **2011**, 286, 403-409.
- Caldecott, K.W. *Nat. Struct. Mol. Biol.* **2012**, 19, 1212-1213.
- Pommier, Y.; Leo, E.; Zhang, H., Marchand, C. *Chem. Biol.* **2010**, 17, 421-433.
- Gomez-Herreros, F.; Romero-Granados, R.; Zeng, Z.; Alvarez-Quilon, A.; Quintero, C.; Ju, L.; Umans, L.; Vermeire, L.; Huylebroeck, D.; Caldecott, K.W., Cortes-Ledesma, F. *PLoS Genet.* **2013**, 9, e1003226.
- Schellenberg, M.J.; Appel, C.D.; Adhikari, S.; Robertson, P.D.; Ramsden, D.A., Williams, R.S. *Nat. Struct. Mol. Biol.* **2012**, 19, 1363-1371.
- Shi, K.; Kurahashi, K.; Gao, R.; Tsutakawa, S.E.; Tainer, J.A.; Pommier, Y., Aihara, H. *Nat. Struct. Mol. Biol.* **2012**, 19, 1372-1377.
- Adhikari, S.; Karmahapatra, S.K.; Elias, H.; Dhopeswarkar, P.; Williams, R.S.; Byers, S.; Uren, A., Roy, R. *Anal Biochem.* **2011**, 416, 112-116.
- Raymond, A.C.; Staker, B.L., Burgin, A.B., Jr. *J. Biol. Chem.* **2005**, 280, 22029-22035.
- Murai, J.; Huang, S.Y.; Das, B.B.; Dexheimer, T.S.; Takeda, S., Pommier, Y. *J. Biol. Chem.* **2012**, 287, 12848-12857.
- Bahmed, K.; Nitiss, K.C., Nitiss, J.L. *Proc. Natl. Acad. Sci. U.S.A.* **2010**, 107, 4057-4062.
- Raoof, A.; Depledge, P.; Hamilton, N.M.; Hamilton, N.S.; Hitchin, J.R.; Hopkins, G.V.; Jordan, A.M.; Maguire, L.A.; McGonagle, A.E.; Mould, D.P.; Rushbrooke, M.; Small, H.F.; Smith, K.M.; Thomson, G.J.; Turlais, F.; Waddell, I.D.; Waszkowycz, B.; Watson, A.J., Ogilvie, D.J. *J. Med. Chem.* **2013**, 56, 6352-6370.
- Ihlenfeldt, W.D.; Voigt, J.H.; Bienfait, B.; Oellien, F., Nicklaus, M.C. *J. Chem. Inf. Model.* **2002**, 42, 46-57.
- Sali, A., Blundell, T.L. *J. Mol. Biol.* **1993**, 234, 779-815.
- Morris, G.M.; Huey, R.; Lindstrom, W.; Sanner, M.F.; Belew, R.K.; Goodsell, D.S., Olson, A.J. *J. Comput. Chem.* **2009**, 30, 2785-2791.
- Phillips, J.C.; Braun, R.; Wang, W.; Gumbart, J.; Tajkhorshid, E.; Villa, E.; Chipot, C.; Skeel, R.D.; Kale, L., Schulten, K. *J. Comput. Chem.* **2005**, 26, 1781-1802.
- Roe, D.R., Cheatham, T.E. *J. Chem. Theory Comput.* **2013**, 9, 3084-3095.
- Hawkins, P.C.; Skillman, A.G.; Warren, G.L.; Ellingson, B.A., Stahl, M.T. *J. Chem. Inf. Model.* **2010**, 50, 572-584.
- Hawkins, P.C.; Skillman, A.G., Nicholls, A. *J. Med. Chem.* **2007**, 50, 74-82.
- Tresadern, G.; Bemporad, D., Howe, T. *J. Mol. Graph. Model.* **2009**, 27, 860-870.
- Marchand, C.; Abdelmalak, M.; Kankanala, J.; Huang, S.Y.; Kiselev, E.; Fesen, K.; Kurahashi, K.; Sasanuma, H.; Takeda, S.; Aihara, H.; Wang, Z., Pommier, Y. *Chem. Biol.* **2016**,
- Miller, B.R.; McGee, T.D.; Swails, J.M.; Homeyer, N.; Gohlke, H., Roitberg, A.E. *J. Chem. Theory Comput.* **2012**, 8, 3314-3321.
- Wang, J.; Wang, W.; Kollman, P.A., Case, D.A. *J. Mol. Graph. Model.* **2006**, 25, 247-260.
- Gohlke, H.; Kiel, C., Case, D.A. *J. Mol. Biol.* **2003**, 330, 891-913.

Graphical Abstract

Discovery of selective inhibitors of tyrosyl-DNA phosphodiesterase 2 by targeting the enzyme DNA-binding cleft

Bradley R Kossmann, Monica Abdelmalak, Sophia Lopez, Gabrielle Tender, Chunli Yan, Yves Pommier, Christophe Marchand and Ivaylo Ivanov