

## Interactions between Human Glutamate Carboxypeptidase II and Urea-Based Inhibitors: Structural Characterization<sup>†</sup>

Cyril Barinka,<sup>‡</sup> Youngjoo Byun,<sup>§</sup> Crystal L. Dusich,<sup>§,#</sup> Sangeeta R. Banerjee,<sup>§,#</sup> Ying Chen,<sup>§,#</sup> Mark Castanares,<sup>||</sup> Alan P. Kozikowski,<sup>⊥</sup> Ronnie C. Mease,<sup>§</sup> Martin G. Pomper,<sup>\*,§,||</sup> and Jacek Lubkowski<sup>\*,‡</sup>

Center for Cancer Research, National Cancer Institute at Frederick, Frederick, Maryland 21702, Russell H. Morgan Department of Radiology and Radiological Sciences, Department of Pharmacology & Molecular Sciences, Johns Hopkins Medical Institutions, Baltimore, Maryland 21231, Department of Medicinal Chemistry and Pharmacognosy, University of Illinois at Chicago, Chicago, Illinois 60612

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Urea-based, low molecular weight ligands of glutamate carboxypeptidase II (GCPII) have demonstrated efficacy in various models of neurological disorders and can serve as imaging agents for prostate cancer. To enhance further development of such compounds, we determined X-ray structures of four complexes between human GCPII and urea-based inhibitors at high resolution. All ligands demonstrate an invariant glutarate moiety within the S1' pocket of the enzyme. The ureido linkage between P1 and P1' inhibitor sites interacts with the active-site Zn<sub>1</sub><sup>2+</sup> ion and the side chains of Tyr552 and His553. Interactions within the S1 pocket are defined primarily by a network of hydrogen bonds between the P1 carboxylate group of the inhibitors and the side chains of Arg534, Arg536, and Asn519. Importantly, we have identified a hydrophobic pocket accessory to the S1 site that can be exploited for structure-based design of novel GCPII inhibitors with increased lipophilicity.

### Introduction

Glutamate carboxypeptidase II (GCPII, E.C. 3.4.17.21) is a type II transmembrane exopeptidase expressed in the nervous system<sup>1</sup> where it hydrolyzes endogenous *N*-acetylaspartyl glutamate (NAAG<sup>a</sup>) to *N*-acetylaspartate and glutamate, the latter being the most abundant excitatory neurotransmitter.<sup>2</sup> Under normal conditions, glutamate is indispensable for physiological processes such as learning, memory, and developmental plasticity.<sup>3</sup> However, excessive glutamate production and release can result in neuronal cell death and is implicated in a variety of neurological conditions including neuropathic/diabetic pain, stroke, trauma, amyotrophic lateral sclerosis, and schizophrenia.<sup>4,5</sup> The upregulation of GCPII expression is also observed in prostate carcinoma and within the neovasculature of solid tumors. The expression of GCPII is confined predominantly to

the cell surface, and the enzyme is not extensively shed into the circulation. Therefore, GCPII represents an attractive target for the imaging and therapy of a variety of cancers.<sup>6–8</sup>

Given the GCPII involvement in a variety of pathologies, discovery and development of GCPII inhibitors as diagnostic or therapeutic agents have been extensively pursued for the last decade (refs 9 and 10 and references therein). GCPII inhibitors reported to date occupy either solely the S1' (pharmacophore) pocket, or the enzyme-inhibitor interactions can extend over both S1 and S1' sites. The former type of inhibitor is usually a derivative or mimetic of glutamic acid linked to a zinc-binding group such as phospho(i)onate or thiol, and such inhibitors preferentially bind to the S1' site of GCPII.<sup>11–13</sup> The latter group encompasses analogues of dipeptides (such as NAAG, the natural GCPII substrate) with the peptide bond substituted by a hydrolysis-resistant surrogate.<sup>14–17</sup> Among others, this category includes dipeptide analogues connected by a urea linkage, previously developed by us.<sup>18,19</sup> The relative ease of synthesis of the urea-based inhibitors of GCPII facilitates SAR studies. Furthermore, compared to the similar phosphinates, the urea derivatives are less polar and thus better suited for applications requiring increased lipophilicity of inhibitors such as stroke therapy. The efficacy of the urea-based compounds has been demonstrated in various animal models of neurological disorders.<sup>19–23</sup> Additionally, radiolabeled derivatives were successfully applied for in vivo imaging in experimental models of prostate cancer as well as for in vitro identification of GCPII in rodent and human brains.<sup>24–26</sup>

The extracellular catalytic portion of GCPII (amino acids 44–750) consists of three structurally distinct domains, each of them contributing residues implicated in substrate binding.<sup>27</sup> The S1' (pharmacophore) pocket accepts the C-terminal part of a substrate/inhibitor and preferentially binds glutamate or glutamate-like moieties. The most prominent feature of the S1 pocket in GCPII is the "arginine patch", comprising arginines 463, 534, and 536, which is mainly associated with the enzyme's preferences for negatively charged P1 residues of the ligand.<sup>11,27–30</sup> Compared to the S1' site, the S1 pocket is more

<sup>†</sup> PDB accession numbers. Atomic coordinates of the present structures together with the experimental diffraction amplitudes have been deposited at the RCSB Protein Data Bank with accession numbers 3D7G (the complex with **1**, (DCMC)), 3D7F (the complex with **2**, (DCIT)), 3D7D (the complex with **3**, (DCFBC)), and 3D7H (the complex with **4**, (DCIBzL)).

\* To whom correspondence should be addressed. for J.L.: phone, 301-846-5494; fax, 301-846-7517; E-mail, jacek@ncifcrf.gov; address: Macromolecular Crystallography Laboratory, 539 Boyles Street, National Cancer Institute at Frederick, Frederick, MD 21702. For M.G.P.: phone, 410-955-2789; fax, 443-956-5055; E-mail, mpomper@jhmi.edu; address: 1550 Orleans Street, 492 CRB II, Johns Hopkins Medical Institutions Baltimore, MD 21213.

<sup>‡</sup> Center for Cancer Research, National Cancer Institute at Frederick.

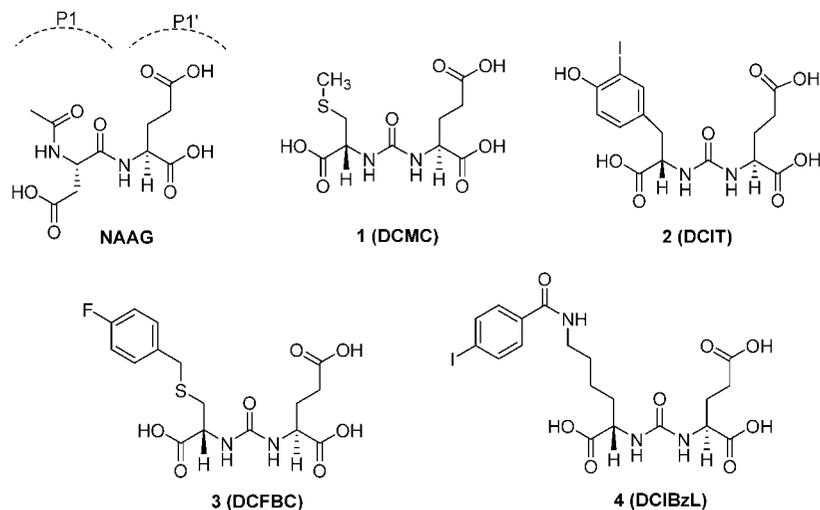
<sup>§</sup> Russell H. Morgan Department of Radiology and Radiological Sciences, Johns Hopkins Medical Institutions.

<sup>||</sup> Department of Pharmacology & Molecular Sciences, Johns Hopkins Medical Institutions.

<sup>⊥</sup> Department of Medicinal Chemistry and Pharmacognosy, University of Illinois at Chicago.

<sup>#</sup> These authors contributed equally.

<sup>a</sup> Abbreviations: rhGCPII, recombinant human glutamate carboxypeptidase II; DCIT, (S)-2-(3-((S)-1-carboxy-2-(4-hydroxy-3-iodophenyl)ethyl)ureido)pentanedioic acid; DCMC, (S)-2-(3-((R)-1-carboxy-2-methylthio)ethyl)ureido)pentanedioic acid; DCFBC, (S)-2-(3-((R)-1-carboxy-2-(4-fluorobenzylthio)ethyl)ureido)pentanedioic acid; DCIBzL, (S)-2-(3-((S)-1-carboxy-(4-iodobenzamido)pentyl)ureido)pentanedioic acid; NAAG, *N*-acetylaspartylglutamate; SAR, structure–activity relationship.



**Figure 1.** Chemical structures of NAAG and urea-based GCPII inhibitors.

flexible in terms of structural modifications of the GCPII inhibitors.<sup>14,16,19</sup>

Recently, we have reported several high-resolution structures of the GCPII ectodomain in complex with small-molecule ligands such as mimetics and derivatives of glutamic acid as well as phosphinate-based analogues of acidic dipeptides.<sup>11,29</sup> These structures advanced our understanding of the architecture of the GCPII active site but did not reveal the complete picture of GCPII–inhibitor interactions. First, out of a variety of zinc-binding groups utilized for the design of GCPII inhibitors, only phospho(i)nates have so far been successfully co-crystallized with GCPII.<sup>11,27,29</sup> Furthermore, lack or limited variability of the P1 side chains in inhibitors studied previously hampers detailed analysis of the S1 site. Also, this lack of variability limits a structure-based design of new GCPII inhibitors by overlooking the advantage of modifications in the P1 position that might provide higher affinity interaction with the enzyme.

This manuscript aims to extend our understanding of interactions between human GCPII and low molecular weight ligands. We describe the first detailed structures of complexes between human GCPII and urea-based inhibitors. Our data provide an explanation for earlier SAR studies, offer a deeper insight into both the architecture of the S1 pocket and enzyme–inhibitor interactions, and form the basis for the structure-aided design and development of the next generation of dipeptide-based GCPII inhibitors and substrates.

## Results and Discussion

**Synthesis of Inhibitors.** Urea-based inhibitors with which GCPII crystal complexes were determined in this study include (*S*)-2-(3-((*R*)-1-carboxy-2-methylthio)ethyl)ureido)pentanedioic acid (**1**, DCMC in Figure 1), (*S*)-2-(3-((*S*)-1-carboxy-2-(4-hydroxy-3-iodophenyl)ethyl)ureido)pentanedioic acid (**2**, DCIT in Figure 1), (*S*)-2-(3-((*R*)-1-carboxy-2-(4-fluorobenzylthio)ethyl)ureido)pentanedioic acid (**3**, DCFBC in Figure 1), and (*S*)-2-(3-((*S*)-1-carboxy-(4-iodobenzamido)pentyl)ureido)pentanedioic acid (**4**, DCIBzL in Figure 1). All four inhibitors were stereochemically pure and can be viewed as nonhydrolyzable analogues of NAAG, with the *N*-acetylaspartate moiety replaced by four different side chains. Common features of these compounds are (a) the ureido linkage substituting for the peptide bond of a parental dipeptide, and (b) the C-terminal glutarate moiety (Figure 1). The synthesis of **1** has been reported previously.<sup>18</sup> Our synthesis of **1** differs from the previous protocol by the use of *p*-methoxybenzyl esters as protecting

groups instead of benzyl esters. Preparation of **2** is different from that used to prepare [<sup>125</sup>I]**2**,<sup>24</sup> where the radioiodine is added in the last step of the radiosynthesis. Experimental details and analytical data of **1** and **2** are described in the Supporting Information. Compounds **3** and **4** were prepared as described previously.<sup>31,32</sup> Compounds **1**, **2**, and **3** are potent inhibitors of GCPII, with IC<sub>50</sub> values of 17, 0.5, and 14 nM, respectively.<sup>24,32,33</sup> Compound **4** is even more potent with an IC<sub>50</sub> value of 0.06 nM.

**Overall Structural Comparison.** Structures of rhGCPII/**2**, rhGCPII/**1**, rhGCPII/**3**, and rhGCPII/**4** complexes were determined by difference Fourier methods and refined at the resolutions of 1.54, 1.75, 1.69, and 1.55 Å, respectively (Table 1). The orientation of the individual inhibitors in the rhGCPII active site was unequivocally identified from the positive density peaks in the  $F_o - F_c$  omit maps (Supporting Information, Figure S1) with the invariant C-terminal glutamate facing the S1' pocket of the enzyme and variable positioning of the P1 moieties. All four complexes of rhGCPII share common fold, with the root-mean-square deviations between 0.32 and 0.52 Å for ~695 equivalent C $\alpha$ -atoms in any two complexes compared.

**The S1' Pocket.** The S1' site of human GCPII has strong preference for glutamate or glutamate-like residues.<sup>28,34</sup> Consistent with this observation, the S1' pocket is occupied by the invariant glutamate of the urea inhibitors. The  $\alpha$ -carboxylate group of glutamate of the inhibitor forms strong H-bonds with the guanidinium group of Arg210 (2.8 Å; distances present here are from rhGCPII/**2**, the complex refined at the highest resolution) and the hydroxyl group of Tyr552 (3.1 Å), and the  $\gamma$ -carboxylate is engaged by the side chains of Asn257 (N $\delta$ 2, 2.9 Å) and Lys699 (N $\zeta$ , 2.7 Å). The shape of the S1' pocket is defined by Gly518 and the side chains of Phe209 and Leu428, which also contribute nonpolar interactions to the inhibitor binding. Additional stabilization of inhibitor is contributed by the water interactions of  $\alpha$ - and  $\gamma$ -carboxylate groups. The overall architecture of the S1' pocket, position of the C-terminal glutamate, and the enzyme–inhibitor interactions are virtually identical to those observed for complexes of rhGCPII with free glutamate and phospho(i)nate analogues of glutamate/NAAG<sup>11,27,29</sup> (Figure 2). Evidently, the S1' pocket is “optimized” for glutamate binding and glutamate-like residues are the best choice as to be positioned at the C-terminus of GCPII inhibitors. Our structural observations are fully supported by available SAR studies showing low tolerance for substitutions at the P1' position of the inhibitors.<sup>14,16,19</sup>

**Table 1.** Data Collection and Refinement Statistics

	rhGCPII/1	rhGCPII/2	rhGCPII/3	rhGCPII/4
Data Collection Statistics				
wavelength (Å)			1.000	
temperature (K)			100	
space group			I222	
unit cell parameters <i>a</i> , <i>b</i> , <i>c</i> (Å)	101.8, 129.7, 159.3	101.4, 130.0, 158.5	101.9, 130.5, 159.2	101.3, 130.5, 158.7
resolution limits (Å)	30–1.75 (1.81–1.75) <sup>a</sup>	30–1.54 (1.60–1.54)	30–1.69 (1.75–1.69)	30–1.55 (1.61–1.55)
number of unique reflections	105078 (10129)	144557 (9835)	116666 (11057)	143433 (9756)
redundancy	6.8 (5.2)	6.1 (4.0)	6.5 (4.5)	7.1 (4.5)
completeness (%)	99.6 (96.9)	95.0 (65.3)	99.5 (95.3)	94.6 (64.9)
<i>I</i> / <i>σ</i> <i>I</i>	21.5 (2.4)	17.6 (2.2)	14.8 (2.3)	21.4 (2.2)
<i>R</i> <sub>merge</sub>	0.070 (0.49)	0.071 (0.43)	0.094 (0.51)	0.068 (0.43)
Refinement Statistics				
resolution limits (Å)	15.0–1.75 (1.79–1.75)	15.0–1.54 (1.58–1.54)	15.0–1.69 (1.74–1.69)	15.0–1.55 (1.59–1.55)
no. of reflections	103375 (6801)	142946 (6620)	114765 (7473)	141846 (6804)
no. of reflections in test set	1550 (111)	1446 (66)	1730 (97)	1445 (76)
<i>R</i>	0.170 (0.252)	0.177 (0.279)	0.179 (0.255)	0.183 (0.270)
<i>R</i> <sub>free</sub>	0.204 (0.272)	0.195 (0.276) [0.155 (0.226)] <sup>b</sup> [0.179 (0.254)]	0.199 (0.293)	0.208 (0.290) [0.161 (0.213)] [0.190 (0.257)]
total number of non-H atoms	6552	6603	6580	6529
number of ligand atoms	21	26	28	31
number of ions	4	4	4	4
number of water molecules	617	657	626	592
average B-factor (Å <sup>2</sup> )				
protein atoms	26.4	25.7	26.8	27.9
waters	38.7	38.6	38.4	39.6
inhibitor	24.5	25.7	26.8	25.4
rms deviations				
bond lengths (Å)	0.021	0.018 [0.015]	0.020	0.018 [0.015]
bond angles (deg)	1.92	1.78 [1.60]	1.79	1.74 [1.57]
Ramachandran plot (%)				
most favored	90.3	89.2	89.7	89.3
additionally allowed	9.2	10.2	9.8	10.0
generously allowed	0.3	0.3	0.3	0.5
disallowed	0.2 (Lys207)	0.2 (Lys207)	0.2 (Lys207)	0.2 (Lys207)
missing residues	44–54, 541–543, 654–655	44–54, 545–547, 654–655	44–54, 654–655	44–54, 654–655

<sup>a</sup> Values in parentheses correspond to the highest resolution shells. <sup>b</sup> Values in brackets correspond to refinement utilizing the anisotropic model for B-factors.

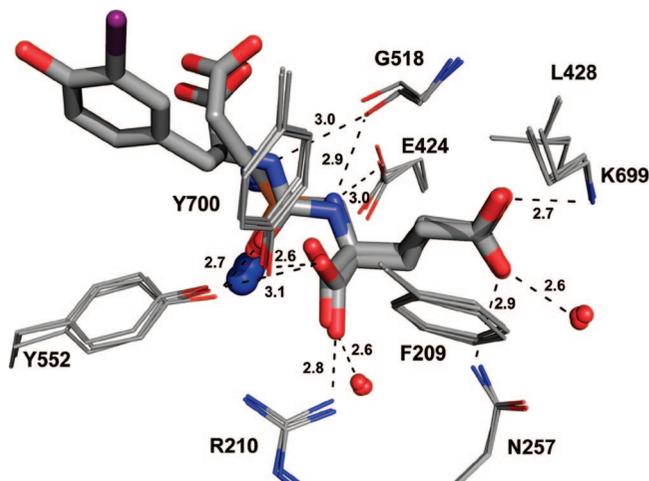
**The Ureido Linkage.** The ureido group of the inhibitors mimics a planar peptide bond of a GCPII substrate, such as NAAG, but acts as an amide-bioisostere due to its resistance to hydrolysis by the enzyme.<sup>35</sup> In the structures reported here, the ureido carbonyl oxygen is engaged by the side chains of Tyr552 (OH, 2.7 Å) and His553 (Nε2, 3.2 Å), and further interacts with the activated water molecule (2.8 Å) and the Zn<sub>1</sub><sup>2+</sup> ion (2.6 Å). The N2 is H-bonded to the Glu424 γ-carboxylate (3.0 Å), Gly518 carbonyl oxygen (2.9 Å), and the activated water molecule (3.1 Å), while the second ureido nitrogen donates only one H-bond to the Gly518 main-chain carbonyl (3.0 Å).

It is interesting to note that the active-site arrangement of the rhGCPII–ureido complexes mirrors the situation in the rhGCPII(E424A)/NAAG complex (Barinka, unpublished) with the distance between the two active-site zinc ions ~3.3 Å and the activated water molecule positioned symmetrically in between them (2.0 Å). This finding validates the prediction of the urea bioisostere as a true surrogate of the peptide bond.

**The S1 Site and an “Accessory Hydrophobic Pocket.”** The major structural signature of the S1 pocket of the enzyme is the “arginine patch” comprising arginines 463, 534, and 536.<sup>29,30</sup> The Arg534 is kept in an invariant position via its interaction with the S1-bound chloride anion, but both Arg536 and, to a

lesser extent Arg463, are quite flexible. The Arg536 side chain can adopt two distinct conformations referred to as “stacking” and “binding,” and transition between these two states is associated with shift of the Arg463 side chain between “up” and “down” positions, respectively (Supporting Information, Figure S2). Such flexibility likely contributes toward less stringent substrate specificity within the S1 site of the enzyme (in comparison to the S1′ site) and also modulates affinity of P1 diversified GCPII inhibitors.<sup>28,29,36</sup>

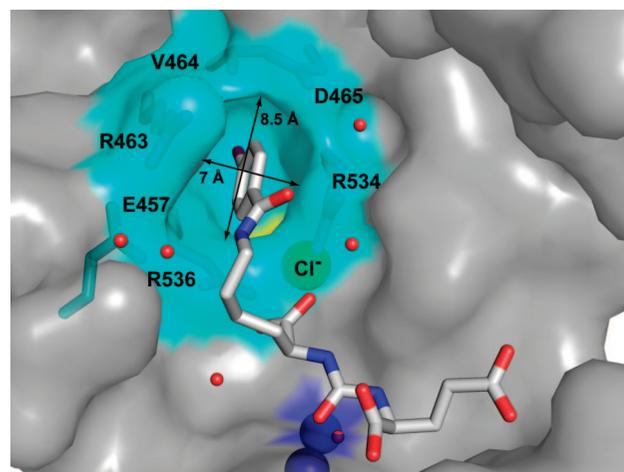
While the C-terminal glutamate and the ureido group of all four inhibitors structurally overlap, there are considerable positional differences within the S1 pocket of the enzyme (Figure 3A). The only common denominator of the S1 sites in four complexes is the engagement of the P1 carboxylate in four direct contacts with the side chains of Asn519 (Nδ, 2.9 Å), Arg534 (Nη1, 2.9 Å) and Arg536 (Nη1, 2.9 Å; Nη2, 3.0 Å) and two water mediated polar interactions (2.8 and 2.7 Å; Figure 3B). Hydrogen bonds between the P1 carboxylate and the guanidinium group of Arg536 stabilize the latter in its “binding” conformation in a manner similar to the binding of NAAG. Because the P1 carboxylate of the studied urea-based compounds contributes prominently to the GCPII–inhibitor interactions in the S1 pocket, it is reasonable to suggest that this



**Figure 2.** Structural similarity of the glutamate binding in the S1' pocket. The X-ray structures of rhGCPII/EPE (PDB code 3bi0), rhGCPII/glutamate complex (PDB code 2c6g), and the rhGCPII/2 complexes are superimposed using corresponding C $\alpha$ -atoms. The active site ligands are in stick representation, residues shaping the S1' pocket are shown as lines, Zn<sup>2+</sup> ions as blue spheres, and conserved water molecules as red spheres. The H-bonds are indicated by dashed lines with distances shown in Ångstroms (from the rhGCPII/2 complex).

structural feature should be preserved in future generations of such inhibitors to retain high-affinity binding for GCPII.

The P1 methylcysteine side chain of **1** is placed into a "hydrophobic section" of the S1 pocket, defined by the side-chains of Tyr549, Tyr552, and Tyr700, but its fit is rather loose with no signature interactions observed. A similar conclusion can be drawn for the P1 side chain of **2**, with the exception of H-bonds between the P1 hydroxyl group and side-chains of Glu457 (O $\epsilon$ 1, 2.6 Å) and Arg463 (N $\eta$ 2, 3.2 Å). Each of **2**, **3**, and **4** feature a phenyl ring as a terminal part of their P1 side chain but they differ in the length of a linker connecting the ring to the ureido group (Figure 1). The linker in **2** consists of only one methylene group, while those in **3** and **4** feature equivalents of three and six methylene groups, respectively (Figure 1). Consequently, the phenyl groups of **3**, and especially

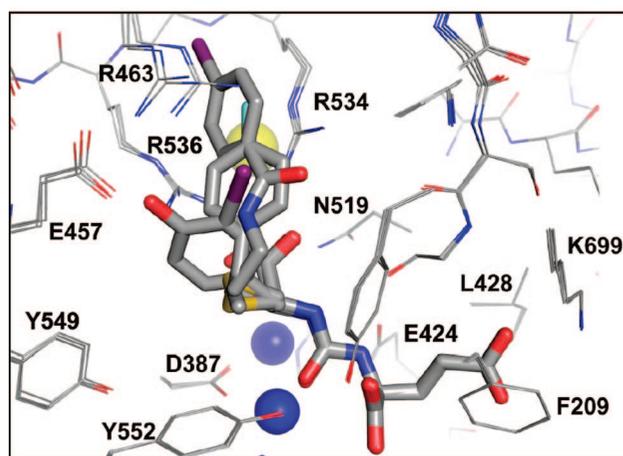


**Figure 4.** The hydrophobic pocket accessory to the S1 site. The dissected substrate-binding cavity of GCPII is shown in semitransparent surface representation (gray). The side chains of amino acids delineating the "accessory hydrophobic pocket" are shown in stick representation and colored cyan. The active-site Zn<sup>2+</sup> and S1-bound Cl<sup>-</sup> are colored blue and yellow, respectively, and water molecules are represented by red spheres. Compound **4** bound to the active site is in stick representation.

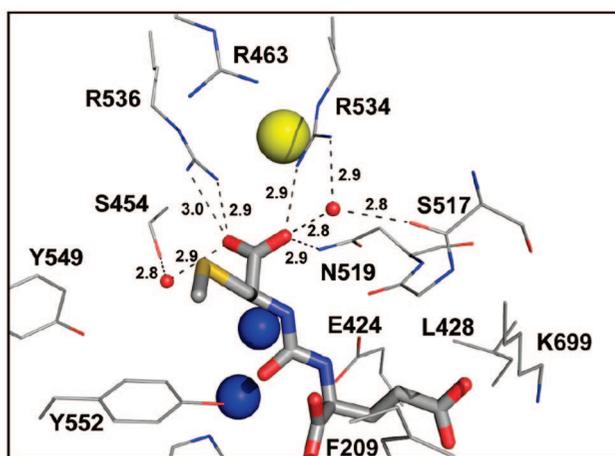
of **4**, have more positional freedom and could extend further into the S1 pocket.

Given the positional freedom of the terminal phenyl ring of **3**, it is interesting to find it partially inserted into a pocket accessory to the S1 site with approximate dimensions of 8.5 Å  $\times$  7 Å  $\times$  9 Å. The bottom of this hydrophobic "accessory pocket" is defined by portions of  $\beta$ -sheets  $\beta$ 13 (Arg463–Asp465) and  $\beta$ 14 (Arg534–Arg536), while the walls are shaped by the side chains of Glu457, Arg463, Asp465, Arg534, and Arg536 (Figure 4). The formation of such a pocket is only possible by simultaneous "atypical" positioning of Arg536 into the "binding" configuration and Arg463 into the "up" position. On the contrary, when NAAG, the natural GCPII substrate, is bound in the active site of the enzyme, the Arg536 side chain is stabilized in the "binding" conformation, but Arg463 relocates

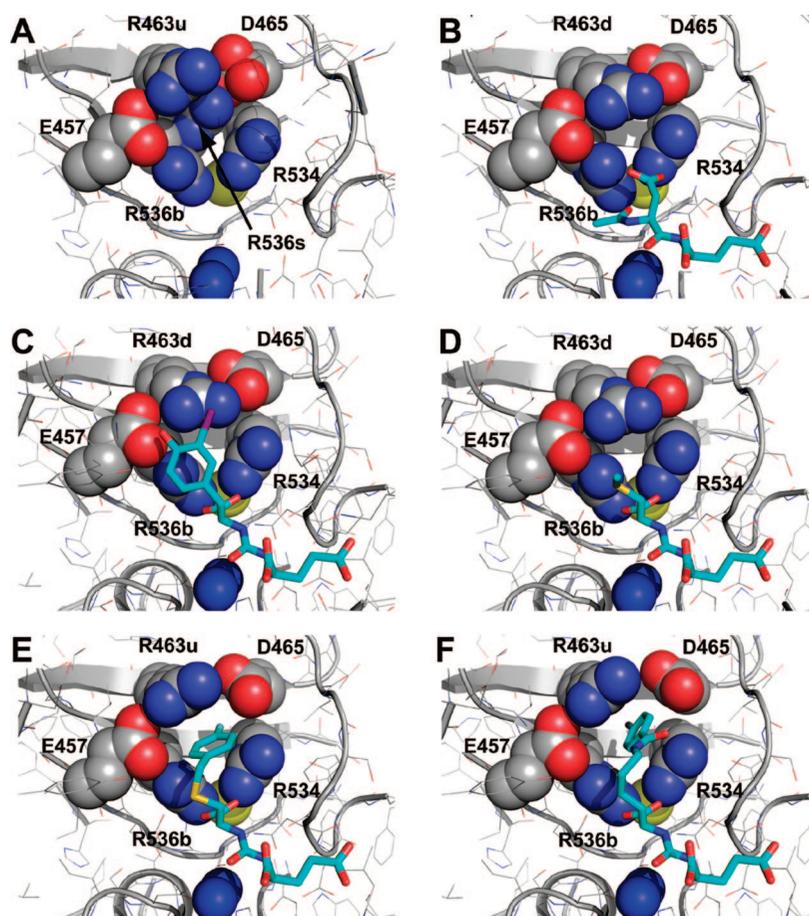
**A**



**B**



**Figure 3.** (A) Superposition of urea-based inhibitors in the active-site of rhGCPII. The rhGCPII–inhibitor complexes were superimposed on corresponding C $\alpha$ -atoms. The inhibitors are shown in stick representation and protein residues are shown as lines. Note invariant positioning of the P1' glutamate contrasting with inhibitor conformational variability in the S1 pocket. (B) Hydrogen-bonding network in the S1 site of GCPII. Hydrogen bonding interactions (indicated with dashed lines) and distances (in Ångstroms) between **1** bound in the active site and S1 residues of GCPII. The zinc ions, chloride anion, and water molecules located in the active site are shown as blue, yellow and red spheres, respectively. The protein and inhibitor atoms are colored red (oxygen), blue (nitrogen), yellow (sulfur), violet (iodine), cyan (fluorine), gray (carbons).



**Figure 5.** The flexibility of S1 arginines 463 and 536 defines size of the “accessory pocket”. Dissected view into the active-site of GCPII. The protein moiety is shown in combination of cartoon and line representation. The residues shaping walls of the accessory pocket are shown as spheres and inhibitor (substrate) residues are in stick representation. The active site  $\text{Zn}^{2+}$  and S1 bound  $\text{Cl}^-$  are colored blue and yellow, respectively. The R463u and R463d denotes the side chain of Arg463 in the “up” and “down” position, respectively, while R536b and R536s denotes the side chain of Arg536 in the “binding” and “stacking” configuration, respectively. Notice the closure of the accessory pocket in the unliganded GCPII structure (A). Shown are complexes of rhGCPII with **2** (C), **1** (D), **3** (E), and **4** (F). The complex between the E424A active-site mutant of GCPII and NAAAG, the GCPII natural substrate, is included for comparison (PDB code 3bxm, (B)).

by 2 Å into the “down” position at the same time, effectively closing the “accessory pocket.” Similarly, under conditions when the S1 site is unoccupied or is occupied by a moiety that fails to enforce the Arg536 “binding” conformation, the Arg536 side chain is found in both “binding” and “stacking” alternate positions (accompanied by the “down” and “up” position of Arg463, respectively) resulting in closure (disappearance) of the “accessory pocket” (Figure 5). In the case of **4**, the phenyl ring is fully inserted into the pocket (due to presence of longer P1 spacer), and this fact likely contributes to the tighter inhibitor binding that is translated into an  $\text{IC}_{50}$  value of  $\sim 10$ -fold lower as compared to **2** and **3**.

The existence of such “remote hydrophobic binding register” has been predicted in the study by Berkman’s group, where the authors showed that lengthening of the methylene linker between the phosphoramidate surrogate and the terminal phenyl group results in stronger inhibitor binding to GCPII.<sup>15</sup> Clearly, the incorporation of a longer spacer into the P1 side chain allows for the full insertion of the phenyl ring into the accessory hydrophobic pocket with concomitant decrease in inhibition constants. Our observations thus provide a structural rationale for Berkman’s inhibition data and further underscore the importance of Arg463 and Arg536 flexibility in influencing the affinity of various inhibitors to GCPII.<sup>29</sup>

Thermal displacement parameters for the P1 moieties of the inhibitors are considerably higher ( $6.3\text{--}15.0 \text{ \AA}^2$ ) when compared

to the corresponding C-termini (Supporting Information, Figure S3). This observation is consistent with site-directed mutagenesis studies, SAR data, and findings reported for the X-ray structures of GCPII and phosphinate-based inhibitors.<sup>14–17,19,29,37</sup> Clearly, the C-terminal glutamate or glutamate-like residue contributes prominently to the inhibitor affinity towards GCPII, while the P1 moiety is less important in this respect. Concurrently, however, the hydrophobic pocket accessory to the S1 site identified here could be exploited for further optimization of GCPII inhibitors by modification of the P1 moiety. The appendage of a hydrophobic functionality (with the linker of the appropriate length) at the P1 position can both enhance inhibitor affinity towards GCPII and increase lipophilicity of such compounds as a first step toward facilitating blood–brain barrier penetration, if desired.

## Conclusions

We determined high-resolution structures of human GCPII and four urea-based inhibitors and identified a hydrophobic pocket accessory to the S1 specificity site. Our data provide a mechanistic explanation for prior SAR studies on GCPII inhibitors and can serve as a useful platform for the design of novel GCPII inhibitors with improved pharmacokinetic properties.

## Experimental Section

**Expression and Purification of Recombinant Human GCPII (rhGCPII).** Human GCPII (the extracellular part, amino acids 44–750) was heterologously overexpressed in *Drosophila*

Schneider's S2 cells and purified to homogeneity as described previously.<sup>28</sup> This construct is designated rhGCPII (recombinant human GCPII). The final protein preparation in 20 mM Tris-HCl, 100 mM NaCl, pH 8.0 was concentrated to 8 mg/mL and stored at -80°C until further use.

**Crystallization and X-ray Data Collection.** The inhibitors were dissolved in distilled water to a final concentration of 40 mM, and the pH of the solution was adjusted to 8.0 with 1 M NaOH. The rhGCPII stock solution (8 mg/mL) was mixed with 1/10 v/v of the individual inhibitor, and the complexes were crystallized using the hanging drop vapor diffusion setup at 293 K. Crystallization droplets were made by combining 1  $\mu$ L of the rhGCPII-inhibitor mixture and 1  $\mu$ L of the reservoir solution containing 33% pentaerythritol propoxylate (PO/OH 5/4; Hampton Research), 1% PEG3350, and 100 mM Tris-HCl, pH 8.0. Crystals belonging to the space group *I*222 typically appeared within two days and reached their final size with approximate dimensions of 0.3 mm  $\times$  0.4 mm  $\times$  0.1 mm within a week. For data collection crystals were flash-frozen in liquid nitrogen directly from the reservoir solution, and diffraction intensities were collected at 100 K using synchrotron radiation at the SER-CAT sector 22 beamlines of the Advanced Photon Source (Argonne, IL) at the X-ray wavelength of 1.0 Å. In all cases, the diffraction data were collected from a single crystal, recorded on a CCD detector, and processed using the HKL2000 software package.<sup>38</sup>

**Structure Determination and Refinement.** Structures of rhGCPII/inhibitor complexes were determined by difference Fourier methods using the coordinates of unliganded rhGCPII (PDB code 2oot) as a starting model.<sup>30</sup> Refinement calculations were performed with Refmac 5.1<sup>39</sup> and manual rebuilding of the models was carried out using the program Xfit.<sup>40</sup> Approximately 1% (corresponding to 1445–1730 reflections) of the data were selected to monitor the progress of the refinement by calculating  $R_{\text{free}}$ . The inhibitor moieties were easily modeled in strong positive  $F_o - F_c$  electron density observed at the expected place in the substrate binding cavity of rhGCPII. At the later stages of refinement, the mixed anisotropic/isotropic refinement protocol was employed with the anisotropic model of the displacement parameters (B-factors) applied to "heavy atoms" (i.e., I, S, Zn<sup>2+</sup>, Ca<sup>2+</sup>, and Cl<sup>-</sup>) of all complexes. Additionally, we used the fully anisotropic refinement model for rhGCPII/2 and rhGCPII/4, the two complexes with the highest resolution data. The anisotropic refinements resulted in better refinement statistics, including lower  $R$  and  $R_{\text{free}}$  and more favorable model geometry (Table 1). At the same time, root mean square deviations between corresponding models refined with the mixed anisotropic/isotropic versus anisotropic protocols were 0.16 Å for each of rhGCPII/2 and rhGCPII/4 complexes, suggesting virtual identity of the final models. Despite of improving the stereochemistry of the structures and their agreement with experimental data, an implementation of the anisotropic refinement of B-factors did not result in additional information. Because with  $\sim$ 2.4 experimental data per parameter refined, there is a possibility of the model being "over-refined" using the fully anisotropic protocol. Accordingly, we based our subsequent structure analysis on models refined in the mixed anisotropic/isotropic mode. The quality of the final models was assessed with the program PROCHECK.<sup>41</sup> The data collection and refinement statistics are listed in Table 1.

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**Supporting Information Available:** Density maps of the substrate binding cavity of human GCPII in complex with **3**, **2**, **1**, and **4**; flexibility of S1 Arg463 and Arg536; thermal displacement parameters of urea-based inhibitors; experimental procedures and analytical data for all intermediates of **1** and **2**. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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