

# Higher-order interactions bridge the nitric oxide receptor and catalytic domains of soluble guanylate cyclase

Eric S. Underbakke<sup>a</sup>, Anthony T. Iavarone<sup>b</sup>, and Michael A. Marletta<sup>a,1</sup>

<sup>a</sup>Department of Chemistry, The Scripps Research Institute, La Jolla, CA 92037; and <sup>b</sup>Department of Chemistry and California Institute for Quantitative Biosciences, University of California, Berkeley, CA 94720

Edited by Gregory A. Petsko, Brandeis University, Waltham, MA, and approved March 20, 2013 (received for review January 30, 2013)

**Nitric oxide (NO) signaling pathways mediate diverse physiological functions, including vasodilation and neurotransmission. Soluble guanylate cyclase (sGC), the primary NO receptor, triggers downstream signaling cascades by producing the second messenger cGMP. NO binds the sGC heme cofactor to stimulate cyclase activity, yet the molecular mechanisms of cyclase activation remain obscure. Although structural models of the individual sGC domains are available, the structure of the full sGC heterodimer is unknown. Understanding the higher-order domain architecture of sGC is a prerequisite to elucidating the mechanisms of NO activation. We used protein footprinting to map interdomain interaction surfaces of the sGC signaling domains. Hydrogen/deuterium exchange mass spectrometry revealed direct interactions between the Per/Arnt/Sim domain and the heme-associated signaling helix of the heme-NO/O<sub>2</sub> binding (H-NOX) domain. Furthermore, interfaces between the H-NOX and catalytic domains were mapped using domain truncations and full-length sGC. The H-NOX domain buries surfaces of the  $\alpha$ 1 catalytic domain proximal to the cyclase active site, suggesting a signaling mechanism involving NO-induced derepression of catalytic activity. Together, our data reveal interdomain interactions responsible for communicating NO occupancy from H-NOX heme to the catalytic domain active site.**

HDX-MS | hemoprotein | gas receptor | nucleotide cyclase

The diatomic gas nitric oxide (NO) serves as a unique signaling molecule in mammalian physiology. NO signaling controls diverse circulatory and neural processes, including vasodilation, neurotransmission, myocardial function, and platelet aggregation (1, 2). Accordingly, disruptions in NO signaling have been linked to heart disease, erectile dysfunction, stroke, hypertension, and neurodegeneration (3, 4).

Soluble guanylate cyclase (sGC) is the primary cellular NO receptor (5, 6). sGC responds to NO with exquisite sensitivity, amplifying the signal to downstream effectors by catalyzing the formation of the second messenger cGMP from GTP. As a central arbiter of cGMP signaling, sGC represents an attractive target for therapeutic development. Indeed, several sGC stimulators are in clinical development for treatment of cardiovascular disease (7, 8).

NO binds to sGC through a ferrous heme cofactor. Formation of the NO-heme complex breaks the bond between the heme iron and the axial histidine ligand to initiate a largely uncharacterized conformational change that ultimately stimulates cyclase activity several hundred fold.

The predominant and best-characterized sGC isoform is a heterodimeric hemoprotein composed of two homologous subunits:  $\alpha$ 1 and  $\beta$ 1. Each subunit is composed of four domains: an N-terminal receptor domain, a Per/Arnt/Sim (PAS)-like domain, an amphipathic helix, and a C-terminal catalytic domain (Fig. 1). The N-terminal domain of the  $\beta$ 1 subunit shares sequence homology with a broader family of gas sensors dubbed heme-NO/O<sub>2</sub> binding (H-NOX) domains (9, 10). The corresponding domain of the  $\alpha$ 1 subunit is a degenerate H-NOX that does not bind heme, and its function remains unclear. The sGC PAS and helical domains

are involved in heterodimer formation and play ambiguous roles in cyclase activation (11–13). The C-terminal catalytic domains encompass the cyclase active site (14). The active site is formed between the  $\alpha$ 1 and  $\beta$ 1 catalytic domains, with residues from both subunits contributing to catalysis.

The molecular mechanism by which NO occupancy of the sGC heme controls cyclase activity remains a central question in sGC/NO signaling. NO-induced activation presumably occurs via conformational changes communicated from the sGC heme to the catalytic domain. However, this regulatory mechanism remains poorly understood largely because the domain architecture of the full enzyme heterodimer is unknown. Although the structure of full-length sGC remains undetermined, high-resolution structures have been reported for sGC domain truncations and homologs (Fig. 1). We sought to gain insight into the structural organization of the sGC holoenzyme by studying the interactions of its isolated domains.

To explore the higher-order architecture of sGC, we used protein footprinting strategies amenable to mapping interdomain interactions. Protein footprinting reports on the solvent accessibility and local chemical environment of protein residues by monitoring their susceptibility to chemical modification. Changes in the propensity for modification at particular protein regions can be mapped to define conformational changes, domain interactions, and protein–protein interactions. Hydrogen/deuterium exchange MS (HDX-MS) is a widely used footprinting technique that relies on measuring the rate of exchange of peptide bond amide protons with the deuterons of a D<sub>2</sub>O solvent (15–17). Residues buried by protein interactions or conformational changes are detected as decreases in exchange rates. Here, we report results of HDX-MS studies to map interactions between functional sGC domain truncations. Interaction surfaces link heme-associated H-NOX surfaces with the PAS domain. Furthermore, the H-NOX–PAS domains jointly impact regulatory lobes of the catalytic domain. Taken together, the interdomain interaction footprints reveal a potential allosteric pathway by which NO occupancy of the H-NOX heme is communicated to the catalytic domain active site.

## Results

**H-NOX–PAS Domain Interactions.** NO interacts with sGC via the heme cofactor of the N-terminal  $\beta$ 1 H-NOX domain, initiating conformational changes that ultimately activate the C-terminal catalytic domain. The two intervening domains, PAS and helical, likely participate in communicating the heme occupancy state through the enzyme. Accordingly, we reasoned that direct inter-

Author contributions: E.S.U. and M.A.M. designed research; E.S.U. and A.T.I. performed research; E.S.U., A.T.I., and M.A.M. analyzed data; and E.S.U. and M.A.M. wrote the paper.

The authors declare no conflict of interest.

This article is a PNAS Direct Submission.

<sup>1</sup>To whom correspondence should be addressed. E-mail: marletta@scripps.edu.

This article contains supporting information online at [www.pnas.org/lookup/suppl/doi:10.1073/pnas.1301934110/-DCSupplemental](http://www.pnas.org/lookup/suppl/doi:10.1073/pnas.1301934110/-DCSupplemental).





small Sf9 cell cultures (Fig. 3A). Catalytic activity of the basal (i.e., unstimulated) and NO-stimulated enzyme were assessed with cGMP-specific enzyme immunoassays (EIAs; Fig. 3B).

Variants of sGC exhibiting little change in either basal or NO-stimulated activity largely localize to the N-terminal region, outside of the PAS domain footprint. The exceptions, T110R and R116E, reside on the periphery of the interface; the side chains of these residues may extend away from the H-NOX–PAS interaction. Interestingly, alanine substitution at T110 has been reported to elevate basal activity (22). The D106K variant exhibited the greatest impairment of activity (>500-fold inhibition). Accordingly, D106 resides in the middle of the H-NOX–PAS domain interface. Mutation of several clustered residues (I41E, R40E, and D45A) in the N-terminal lobe also substantially reduces NO stimulation of cyclase activity. The I41E, R40E, and D45A variants colocalize to the  $\alpha$ B– $\alpha$ C loop overlaying the heme pocket (Fig. 3A, *Inset*). These results agree well with previous alanine scanning studies that revealed the functional importance of the  $\alpha$ B– $\alpha$ C loop (20, 22). HDX-MS results showed little significant perturbations to the  $\alpha$ B– $\alpha$ C loop due to the PAS domain (Fig. 2A). Taken together, these results suggest that the  $\alpha$ B– $\alpha$ C loop participates in regulatory interactions with the catalytic domain.

**Catalytic Domain Interactions.** The sGC catalytic domain is composed of the C-terminal domains of the  $\alpha$ 1 and  $\beta$ 1 subunits, with both subunits contributing essential active-site residues. Catalytic domain truncations,  $\alpha$ 1[467–690] and  $\beta$ 1[414–619], can be expressed independently and recombined to reconstitute cyclase activity (14). Direct interactions between the sGC H-NOX/PAS and catalytic domain are thought to be responsible for regulating cyclase activity. In fact, coinubation of the catalytic domain truncations ( $\alpha\beta_{\text{cat}}$ ) with H-NOX truncations represses cyclase activity, suggesting an autoinhibitory regulatory role for the sGC

H-NOX (14). Furthermore, the  $\beta$ 1[1–385] construct inhibits  $\alpha\beta_{\text{cat}}$  more potently than the shorter  $\beta$ 1[1–194] truncation, indicating that the PAS domain and/or dimerization contribute to repression of catalytic domain activity. To explore the mechanisms underlying autoinhibition of the catalytic domain, HDX-MS was used to map catalytic domain surfaces buried by regulatory domain interactions.

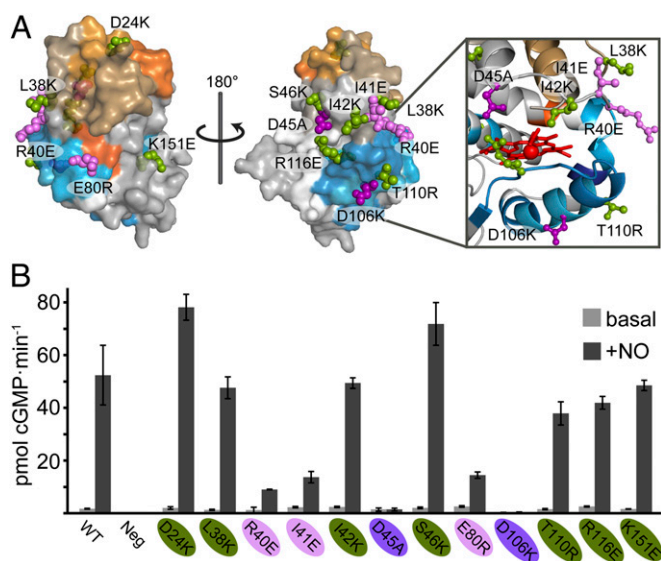
**Mapping  $\alpha\beta_{\text{cat}}$  interactions with  $\beta$ 1[1–385].** When added *in trans*, several molar equivalents of H-NOX domain are required to inhibit  $\alpha\beta_{\text{cat}}$  activity. To ensure detectable interactions via HDX-MS, the concentration dependence of  $\beta$ 1[1–385] inhibition of  $\alpha\beta_{\text{cat}}$  activity was determined (Fig. S2). A 10-fold molar excess of  $\beta$ 1[1–385] over  $\alpha\beta_{\text{cat}}$  inhibited cyclase activity by more than 70%. H/D exchange time courses (10 time points in triplicate, ranging from 5 s to 60 min) were performed comparing  $\alpha\beta_{\text{cat}}$  alone vs.  $\alpha\beta_{\text{cat}}$  incubated with 10 molar equivalents of  $\beta$ 1[1–385] (see Fig. S3 for full exchange time course plots).

Peptide coverage was plotted to primary amino acid sequence for both  $\alpha_{\text{cat}}$  and  $\beta_{\text{cat}}$  (Fig. 4A).  $\beta$ 1[1–385]–induced changes in the extent of exchange ( $\Delta\%D$ ) were color-coded and mapped to both primary sequence and structure of  $\alpha\beta_{\text{cat}}$  (Fig. 4A and B). The two catalytic domain subunits exhibit asymmetries in  $\beta$ 1[1–385]–dependent exchange rate effects. The C-terminal third of the  $\alpha_{\text{cat}}$  subunit exhibits extensively reduced exchange rates. On the other hand, the  $\beta_{\text{cat}}$  subunit contains far fewer peptide fragments displaying lower exchange rates, and conversely, undergoes increased exchange in two discrete regions.

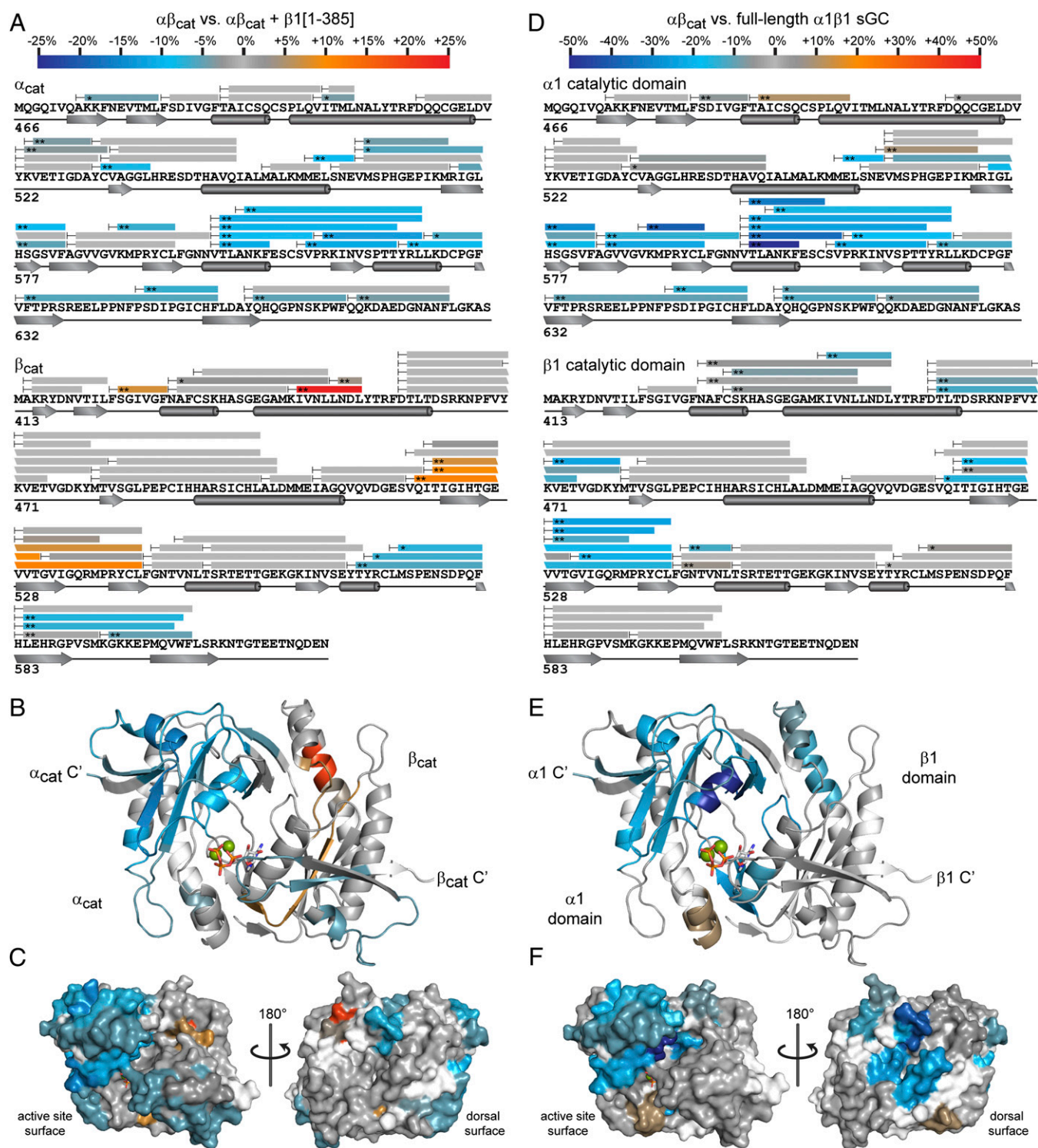
Mapped to the structure of the human  $\alpha\beta_{\text{cat}}$ , the regions protected by  $\beta$ 1[1–385] cluster to one prominent lobe bordering the active site pocket (Fig. 4B). The smaller region of protected surface on the  $\beta_{\text{cat}}$  subunit also envelops the active site pocket. Interestingly, most of the  $\alpha\beta_{\text{cat}}$  surfaces affected by the  $\beta$ 1[1–385] interaction surround the active site, leaving the opposite surface largely unaffected (Fig. 4C, *Right*).

**Mapping catalytic domain interactions in full-length sGC.** Comparing domain truncations interacting *in trans* is invaluable for isolating domain-specific contributions to the architecture of the sGC holoenzyme. Nevertheless, isolated truncations are unrestrained by the linkers and intervening domains that constitute the native sGC heterodimer. We sought to examine regulatory interactions with the catalytic domain in the context of full-length  $\alpha$ 1 $\beta$ 1 sGC. The interaction surfaces of the catalytic domain in the native enzyme can be revealed by comparing the intrinsic solvent accessibility of the isolated  $\alpha\beta_{\text{cat}}$  vs. the catalytic domain ensconced within the sGC holoenzyme.

Full-length sGC presents additional challenges to HDX-MS analysis. The large size (~150 kDa) and limited protein expression are potential complications for HDX-MS analysis. Nevertheless, high-resolution Orbitrap LC-MS afforded excellent sequence coverage, and large magnitude changes in exchange rates between free  $\alpha\beta_{\text{cat}}$  and full-length sGC were strikingly evident (Fig. 4C; Fig. S4). Unsurprisingly, the catalytic domain tethered within the full sGC holoenzyme exhibited larger exchange rate perturbations compared with  $\alpha\beta_{\text{cat}}$  incubated with free  $\beta$ 1[1–385]. Magnitude differences aside, comparing  $\alpha\beta_{\text{cat}}$  vs.  $\beta$ 1[1–385] with  $\alpha\beta_{\text{cat}}$  vs. full-length sGC reveals interesting parallels. Much like incubation with  $\beta$ 1[1–385], full-length sGC buries catalytic domain surfaces surrounding the active site (Fig. 4D). Again, the regions exhibiting the most conspicuously slowed exchange rates cluster to the C-terminal lobe of the  $\alpha$ 1 catalytic subunit. The majority of the  $\beta$ 1 catalytic subunit remains unaffected by inclusion of the full sGC. However, full-length sGC does appear to bury small regions of both  $\alpha$ 1 and  $\beta$ 1 on the “dorsal” catalytic domain surface, opposite the active site (Fig. 4F). This region corresponds to the attachment point of the helical domains. Burial at the dorsal surface may be attributable to catalytic–helical domain interactions. Accordingly, this interface is not evident in the interactions with  $\beta$ 1[1–385], which lacks the C-terminal helical domain (Fig. 4C).

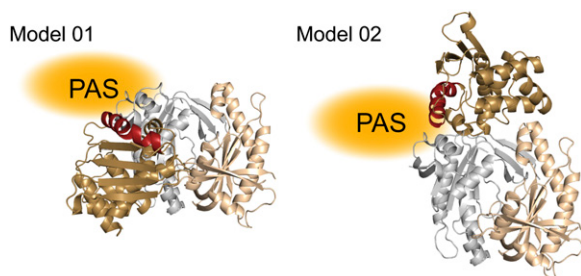


**Fig. 3.** Survey of basal and NO-stimulated cyclase activity of H-NOX variants. (A) H-NOX variants were designed to probe regions implicated in interactions and/or conformational changes in the presence of the PAS domain. Mutated residues are shown on a surface model of sGC  $\beta$ 1 H-NOX derived from the *Nostoc* sp. PCC 7120 H-NOX. Residues are color-coded based on results of activity assays, as in B. (B) WT, negative control ( $\beta$ 1 sGC subunit alone), or H-NOX variants of full-length  $\alpha$ 1 $\beta$ 1 sGC were enriched from Sf9 cell lysates. Cyclase activity of both basal (i.e., minus NO) or NO-stimulated enriched sGC was assayed with cGMP-specific EIA and normalized to WT expression levels. Error bars represent the SD of three experimental replicates. Variants exhibiting similar NO-stimulated activity to WT, moderately reduced activity (<30% of WT), or highly reduced activity (<5% of WT) are color-coded green, pink, and purple, respectively.



**Fig. 4.** Mapping interdomain interactions of the sGC catalytic domains with HDX-MS. (A) Comparison of HDX-MS exchange kinetics of  $\alpha\beta_{cat}$  in the presence or absence of a 10-fold molar excess of  $\beta 1[1-385]$ . Peptide coverage is represented by bars superimposed over catalytic domain primary sequence annotated with secondary structural features. Regions of  $\alpha\beta_{cat}$  exhibiting exchange rate differences ( $\Delta\%D$ ) induced by  $\beta 1[1-385]$  are color-coded according to the scale bar ( $*P < 0.01$ ,  $**P < 0.005$ ; two-tailed, unpaired  $t$  test). Peptides exchanging more slowly in the presence of  $\beta 1[1-385]$  are color-coded in the blue range, whereas peptides exchanging more rapidly are in the red range. Peptides exhibiting nonsignificant changes ( $P > 0.01$ ) are colored gray. (B) Exchange rate differences of  $\alpha\beta_{cat}$  induced by addition of  $\beta 1[1-385]$  are mapped to the structure of human  $\alpha\beta_{cat}$ . Regions lacking peptide coverage are colored white. (C) Surface representations of  $\alpha\beta_{cat}$  were color-coded as in B to highlight potential interdomain interfaces with H-NOX and PAS domains. (D) Comparison of HDX-MS exchange kinetics of  $\alpha\beta_{cat}$  vs. the  $\alpha 1\beta 1$  catalytic domain in the context of full-length sGC. Exchange rate differences are denoted as in A with color bar rescaled to accommodate the larger magnitude changes in full-length sGC. Catalytic domain regions exchanging more slowly in full-length sGC than in the free  $\alpha\beta_{cat}$  truncation are scaled in shades of blue. (E) Likewise, rate differences between free  $\alpha\beta_{cat}$  and the catalytic domain of full-length sGC are color-coded and mapped to the  $\alpha\beta_{cat}$  structure. (F) Results in D were mapped to surface representations of the catalytic domains.





**Fig. 5.** Modeling the H-NOX–catalytic domain interface. The model of sGC  $\beta 1$  H-NOX (based on *Nostoc* sp. PCC 7120 H-NOX) and structure of truncated human catalytic domain were submitted to the ClusPro 2.0 server (24, 25). Collected HDX-MS results served as attraction and repulsion restraints (SI Materials and Methods). ClusPro models representative of the two most highly populated classes of interdomain interactions are pictured with  $\beta 1$  H-NOX (brown with red “signaling helix”),  $\alpha 1$  (gray), and  $\beta 1$  (tan) catalytic domain subunits. Hypothetical PAS domain locations are noted in orange.

Previous reports on the higher-order architecture of sGC position the NO sensor H-NOX domain in contact with the catalytic domains (10, 14, 21, 22). Nevertheless, the structural details of this crucial regulatory interaction remain ambiguous. To explore possible modes of interaction between H-NOX and catalytic domains, a homology-based model of  $\beta 1$  H-NOX and the structure of human  $\alpha\beta_{\text{cat}}$  were docked using the ClusPro server (24, 25). HDX-MS and mutagenesis results were factored into the modeling as attraction and repulsion constraints. The top 10 balanced models were evaluated for conformations that satisfy experimental constraints. The two best models are shown in Fig. 5. Both models allow for mutual contacts between H-NOX–PAS and H-NOX–catalytic domains. Notably, the heme-associated signaling helices of both models are poised to allow simultaneous PAS domain interaction and proximity to the proposed regulatory  $\alpha 1$  lobe.

Taken together, the parallel HDX-MS experiments comparing catalytic domain interfaces both point to a regulatory role for the C-terminal subdomain of the  $\alpha 1$  catalytic subunit. Importantly, this region folds over to adjoin the loops containing the two catalytic  $\text{Mg}^{2+}$ -binding Asp residues and has implications for catalytic regulation (discussed below).

## Discussion

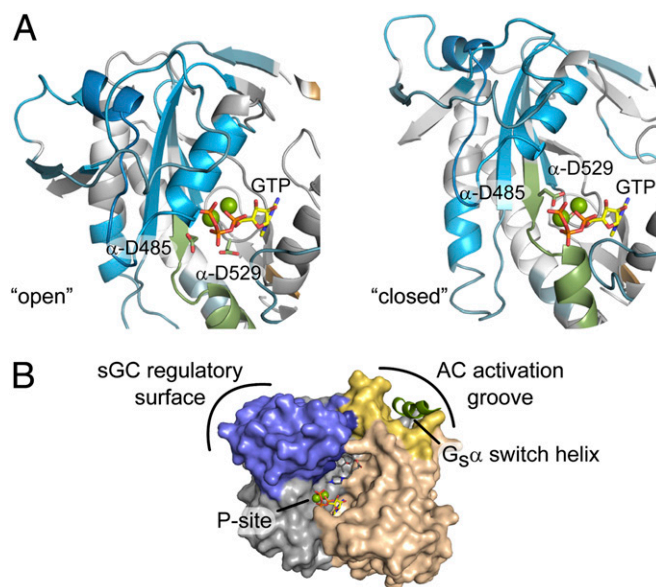
**Interdomain Interactions of sGC H-NOX and PAS Domains.** HDX-MS revealed H-NOX–PAS domain interactions encompassing the signaling helix ( $\alpha\text{F}$ ) with likely implications for interdomain conformational changes. On NO binding to the sGC heme, the bond to the axial His-105 ligand breaks. Consequently, heme distortion is relieved, and the newly freed signaling helix rearranges conformation (10, 19, 20). Stabilization of local structure may also contribute to the observed protection of the signaling helix, potentially expanding the extent of the affected surface. In addition, due to the intrinsic affinity of the  $\beta 1$  PAS domain, homodimerization of  $\beta 1$ [1–385] may also influence local structure. Nevertheless, inclusion of the PAS domain strongly influences the heme-associated helix, likely exerted, in part, by direct interaction. With the PAS domain abutting the signaling helix, the NO-induced conformational change is likely propagated through the PAS domain on to the catalytic domain. Indeed, mutational screens have implicated several PAS domain residues in NO activation of sGC (12).

The surfaces of sGC H-NOX responsible for interdomain communication also extend beyond the signaling helix. Mutagenesis studies demonstrate a functional role for the  $\alpha\text{B}$ – $\alpha\text{C}$  loop and contiguous surfaces (20, 22). The observation that  $\beta 1$ [1–194] represses  $\alpha\beta_{\text{cat}}$  activity *in trans* also indicates that direct H-NOX–catalytic domain interactions contribute to cyclase regulation. Nevertheless, the isolated H-NOX represses  $\alpha\beta_{\text{cat}}$  activity far less po-

tently than the H-NOX–PAS construct (14). Taken together, these studies suggest that cooperative interdomain interactions involving both H-NOX and PAS domains regulate sGC catalytic activity.

**Interdomain Interactions of the sGC Catalytic Domain.** The results of our HDX-MS investigations implicate defined, contiguous catalytic domain surfaces engaged in interdomain interactions in both  $\alpha\beta_{\text{cat}}$  and full-length sGC. Specifically, the  $\beta 1$  H-NOX and/or PAS domain bury the C-terminal lobe of the  $\alpha 1$  catalytic domain. The proximity of the affected surfaces to the sGC active site suggests a role in controlling cyclase activity. Although  $\alpha\beta_{\text{cat}}$  surfaces buried by  $\beta 1$ [1–385] largely localize to the  $\alpha_{\text{cat}}$  domain, subtle decreases in exchange rate were also observed in the  $\beta_{\text{cat}}$  subunit.  $\beta 1$ [1–385] interactions with  $\beta_{\text{cat}}$  are confined to a C-terminal feature, pseudosymmetrical to the buried lobe of  $\alpha_{\text{cat}}$ . However, buried  $\beta 1$  catalytic subunit surfaces were not evident in full-length sGC. The homodimerized  $\beta 1$  H-NOX domains of  $\beta 1$ [1–385] may bury more surface area than the full-length sGC, which substitutes a pseudo-H-NOX in the  $\alpha 1$  subunit.

Comparing our HDX-MS results with the proposed activation mechanisms of adenylate cyclase (AC) yields insights into the regulation of sGC activity. The sGC catalytic domain shares the overall dimeric, wreath-like protein fold as AC and may undergo similar conformational changes (26). AC is activated on interaction with the  $\alpha$  subunit of a heterotrimeric G protein ( $\text{G}_s\alpha$ ). Stimulation of cyclase activity involves closing the active site (e.g., the P-site lid) to position two catalytic aspartate residues into coordination with the  $\text{Mg}^{2+}$ -triphosphate of the ATP substrate (27–29). The active AC conformation is represented by the costructure of the AC catalytic domain and  $\text{G}_s\alpha$ , a stimulatory G protein (30). The inactive conformation is thought to resemble the structure of a low-activity homodimeric catalytic domain. Activation of AC is



**Fig. 6.** Conformational changes involved in catalytic domain activation. (A) HDX-MS results ( $\alpha\beta_{\text{cat}}$  vs.  $\alpha\beta_{\text{cat}}+\beta 1$ [1–385]) mapped to two proposed activity-related conformations of nucleotide cyclases: the putatively inactive conformation (Left, PDB: 3UVJ) and a model of sGC catalytic domains based on an active adenylate cyclase structure (Right, PDB: 1AZS). The two conformations were aligned via the  $\beta 1$  subunit alone using PyMol. The P-site lid is colored drab green.  $\text{Mg}^{2+}$ -coordinating, active-site aspartate residues ( $\alpha\text{-D485}$  and  $\alpha\text{-D529}$ ) are highlighted as sticks. (B) Regulatory surfaces of sGC and AC mapped to activated AC bound to  $\text{G}_s\alpha$  (PDB: 1AZS). Buried sGC surfaces identified by HDX-MS are mapped to the homologous regions of AC in blue. The activation groove of AC is colored yellow with the bound  $\text{G}_s\alpha$  helix in green. The rest of  $\text{G}_s\alpha$  is omitted for clarity.

proposed to involve an  $\sim 7^\circ$  rotation between the two subunits to enclose the substrate within the P site (29, 30).

To explore activation mechanisms of sGC catalytic domains, the interdomain interactions determined by HDX-MS were mapped to models of active and inactive sGC (Fig. 6A). Inactive sGC, represented by an “open” structure of the human sGC catalytic domain, exhibits a relatively compacted  $\alpha 1$  C-terminal lobe (Fig. 6A, *Left*). Consequently, the P-site lid is pushed open, displacing the catalytic aspartate residues (D485 and D529) away from the modeled substrate triphosphate. In contrast, a model of activated sGC catalytic domain based on AC demonstrates a realignment of the  $\alpha 1$  lobe, drawing the catalytic aspartate residues into coordination with  $Mg^{2+}$ -triphosphate (Fig. 6A, *Right*).

Interestingly, the sGC regulatory interfaces identified herein differ from those of AC. Although HDX-MS implicates the C-terminal  $\alpha 1$  lobe in sGC regulation, the costructure of AC catalytic domain with  $G_s\alpha$  demonstrates a stimulatory role for an N-terminal groove in the subunit corresponding to  $\beta 1$  sGC subunit (Fig. 6B). Sprang and coworkers (30) proposed that binding of  $G_s\alpha$  to the AC groove favors the closed catalytic domain conformation, thereby orienting the P site lid into the catalytically competent position. Likewise, activation of sGC may proceed via an analogous intersubunit “closing” mechanism, albeit by relieving repressive interactions through an opposing surface (Movie S1).

The results of these mapping studies suggest mechanisms by which NO binding to the sGC N-terminal H-NOX controls cyclase activity at the C terminus. We anticipate that future structural investigations will address remaining questions. For instance, the interdomain interactions of the  $\alpha 1$  pseudo-H-NOX and helical domains remain to be elucidated. Nevertheless, the interdomain interactions defined herein highlight an allosteric pathway through which the sGC heme can communicate with the cyclase active site. These HDX-MS results, in conjunction with high-resolution structural investigations (10, 12–14, 19), contribute toward building

a model of the sGC holoenzyme (8). Ultimately, a refined model of the higher-order architecture of sGC will empower the development of next-generation therapeutics targeting sGC.

## Materials and Methods

**HDX-MS Analysis.**  $\beta 1[1-194]$  (18),  $\alpha_{cat}/\beta_{cat}$  (14), and full-length sGC (31) were purified as previously described with modifications detailed in *SI Materials and Methods*. H/D exchange was performed as three independent time courses for each treatment. Protein stocks were diluted into buffered  $D_2O$  (pD 8.0, 25 °C) to initiate exchange. Aliquots were withdrawn at various time points, quenched to pH 2.5 with trifluoroacetic acid, and frozen. Rapidly thawed samples were digested (3 min, pH 2.5, 4 °C) with immobilized pepsin (Pierce), frozen in LC-MS vials, and stored at  $-80^\circ C$  until MS analysis. Mass spectra for  $\beta 1[1-194]$  and  $\beta 1[1-385]$  samples were collected on an Agilent 1100/Waters LCT Premier XE LC-MS. Full-length sGC and  $\alpha_{cat}$  were analyzed on an Agilent 1200/Thermo Fisher Scientific LTQ Orbitrap XL LC-MS. C8 columns were maintained at 4 °C throughout LC-MS analysis. HDX-MS data were analyzed using HX Express (32) and HDX Workbench (33) software for  $\beta 1[1-194]/\beta 1[1-385]$  and  $\alpha_{cat}$  experiments, respectively (Fig. S5).

**Cyclase Activity Assays.** Recombinant, His<sub>6</sub>-tagged H-NOX variants of sGC were expressed in 25 mL Sf9 cell culture using the Bac-to-Bac baculovirus expression system (Invitrogen). sGC variants were enriched from Sf9 cell lysates using magnetic Ni-NTA agarose beads (Qiagen). Cyclase activity was assayed in triplicate in the presence or absence of 130  $\mu M$  DEA-NONOate at 25 °C. Endpoint activity assays were initiated with 2 mM GTP and quenched after 6 min. The cGMP product was quantified using the cGMP EIA kit (Enzo Life Sciences), per the manufacturer's instructions.

Full experimental details are described in *SI Materials and Methods*.

**ACKNOWLEDGMENTS.** We thank Bruce Pascal, Michael Chalmers, and Patrick Griffin (Scripps Florida) for data analysis advice and HDX Workbench software assistance. LC-MS/MS instrumentation was acquired with National Institutes of Health (NIH) support (Grant 1510RR022393-01). E.S.U. received financial support from NIH Postdoctoral Fellowship 5F32GM093564. This is The Scripps Research Institute Manuscript 22075.

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