

# Discovery of peptide ligands through docking and virtual screening at nicotinic acetylcholine receptor homology models

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Venom peptide toxins such as conotoxins play a critical role in the characterization of nicotinic acetylcholine receptor (nAChR) structure and function and have potential as nervous system therapeutics as well. However, the lack of solved structures of conotoxins bound to nAChRs and the large size of these peptides are barriers to their computational docking and design. We addressed these challenges in the context of the  $\alpha 4\beta 2$  nAChR, a widespread ligand-gated ion channel in the brain and a target for nicotine addiction therapy, and the 19-residue conotoxin  $\alpha$ -GID that antagonizes it. We developed a docking algorithm, ToxDock, which used ensemble-docking and extensive conformational sampling to dock  $\alpha$ -GID and its analogs to an  $\alpha 4\beta 2$  nAChR homology model. Experimental testing demonstrated that a virtual screen with ToxDock correctly identified three bioactive  $\alpha$ -GID mutants ( $\alpha$ -GID[A10V],  $\alpha$ -GID[V13I], and  $\alpha$ -GID[V13Y]) and one inactive variant ( $\alpha$ -GID[A10Q]). Two mutants,  $\alpha$ -GID[A10V] and  $\alpha$ -GID[V13Y], had substantially reduced potency at the human  $\alpha 7$  nAChR relative to  $\alpha$ -GID, a desirable feature for  $\alpha$ -GID analogs. The general usefulness of the docking algorithm was highlighted by redocking of peptide toxins to two ion channels and a binding protein in which the peptide toxins successfully reverted back to near-native crystallographic poses after being perturbed. Our results demonstrate that ToxDock can overcome two fundamental challenges of docking large toxin peptides to ion channel homology models, as exemplified by the  $\alpha$ -GID: $\alpha 4\beta 2$  nAChR complex, and is extendable to other toxin peptides and ion channels. ToxDock is freely available at [rosie.rosettacommons.org/toxDock](http://rosie.rosettacommons.org/toxDock).

homology model | nicotinic receptor | conotoxin | docking | virtual screening

Venom peptide toxins isolated from predatory marine cone snails (conotoxins) are prized research tools due to their native potency and selectivity for ion channels (1–4). Conotoxins have played an especially important role in elucidating how the many different subtypes of the nicotinic acetylcholine receptor (nAChR), a prototypical ligand-gated ion channel, contribute to diseases like nicotine addiction, Parkinson's disease, and neuropathic pain (5–8). Peptide engineering approaches such as alanine scanning and structure-guided mutagenesis have also been remarkably successful at expanding the limits of conotoxin pharmacology to include those nAChRs not naturally targeted by conotoxins (7). However, some promising conotoxins remain challenging to optimize, and some important nAChRs are difficult to target.

The peptide  $\alpha$ -GID was the first conotoxin found to have substantial antagonism at the rat  $\alpha 4\beta 2$  nAChR, with its  $IC_{50}$  reported

as 128–152 nM and 4.8  $\mu$ M (9–11). The  $\alpha 4\beta 2$  nAChR is one of the most widely expressed nAChRs in the brain and is a validated target for smoking cessation therapeutics (11–13). The receptor is formed from five homologous subunits, consisting of either two  $\alpha 4$  subunits and three  $\beta 2$  subunits or three  $\alpha 4$  subunits and two  $\beta 2$  subunits, organized like barrel staves to form a central channel (14). Cations flow through the channel when acetylcholine (ACh) is bound at specific interfaces between subunits in the extracellular domain (14), where  $\alpha$ -GID presumably binds competitively (9, 15). In addition to the  $\alpha 4\beta 2$  nAChR,  $\alpha$ -GID potently antagonizes the rat  $\alpha 3\beta 2$  nAChR ( $IC_{50}$  reported as 3.1–3.4 nM and 36 nM) as well as the rat  $\alpha 7$  nAChR ( $IC_{50}$  = 4.5–5.1 nM) (9–11). Consequently, there has been significant interest in engineering an analog of  $\alpha$ -GID that antagonizes the  $\alpha 4\beta 2$  nAChR but not the  $\alpha 3\beta 2$  and  $\alpha 7$  nAChRs, especially for the human nAChRs (9–11, 15). A recent study by Banerjee et al. (9) used structure-guided mutagenesis to discover

## Significance

Predicting how conotoxins bind to nicotinic acetylcholine receptors (nAChRs) is hard. Not only are these venom-derived peptides large, but the structures of many nAChRs are unknown. In response, we developed an ensemble-docking algorithm named ToxDock. We used ToxDock to reliably dock the conotoxin  $\alpha$ -GID to a homology model of the  $\alpha 4\beta 2$  nAChR, a main nAChR in the brain and target for nicotine addiction therapeutics. A virtual screen with ToxDock identified four  $\alpha$ -GID analogs and, based on experimental evidence, correctly predicted their activity at the  $\alpha 4\beta 2$  nAChR in all cases. More screening showed that two of these analogs have substantially reduced antagonism at the human  $\alpha 7$  nAChR, a key step in optimizing  $\alpha$ -GID into a tool for studying brain nAChRs.

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The authors declare no conflict of interest.

Data deposition: Atomic coordinates and NMR parameters have been deposited in the Protein Data Bank, [www.pdb.org](http://www.pdb.org) (PDB ID codes 5UG3 and 5UG5).

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that  $\alpha$ -GID[V18N] is selective for the rat  $\alpha 4\beta 2$  nAChR vs. the rat  $\alpha 3\beta 2$  nAChR. Developing an  $\alpha$ -GID analog that retains its activity at the  $\alpha 4\beta 2$  nAChR while having reduced potency at the  $\alpha 7$  nAChR has proven difficult. Millard et al. (10) made significant progress in this respect by performing alanine scanning on  $\alpha$ -GID to elucidate the structural determinants of its activity at the  $\alpha 4\beta 2$  nAChR. They found that five alanine mutants of  $\alpha$ -GID retained bioactivity at the  $\alpha 4\beta 2$  nAChR, but all also remained potent antagonists of the  $\alpha 7$  nAChR (10). Given that the  $\alpha 4\beta 2$  and  $\alpha 7$  nAChRs are the most abundant nAChRs in the human brain (12, 16), developing an  $\alpha$ -GID analog that distinguishes between these two receptors has been, and remains, an important goal.

Docking has emerged as a very useful approach for rationalizing mutagenesis data on conotoxin and nAChR interactions (17–21). Less well understood is whether docking and computational methods can be used to virtually screen for conotoxins with optimized properties, such as potency and selectivity, although recent efforts suggest this may be possible (22, 23). There are two main challenges when docking and virtually screening conotoxins against nAChRs. First, the presence of highly mobile elements in peptide toxins, like the four-residue N-terminal “tail” of  $\alpha$ -GID (11), can impact the results of docking simulations (24). Second, although cocrystal structures of conotoxins and a homopentameric soluble surrogate of the nAChR ligand-binding domain [Acetylcholine Binding Protein (AChBP)] have been solved (25), no crystal structure of a conotoxin and nAChR has been reported to date. This is problematic for a virtual screening campaign that uses a homology model of a heteromeric nAChR built on a homomeric AChBP template structure with less than 30% sequence identity to the nAChR, because effective virtual screening requires that the shape of the ligand-bound complex be known to high precision (26, 27). Hypothetically, broad sampling of many different conformational states of both the conotoxin and nAChR homology model could overcome these two issues.

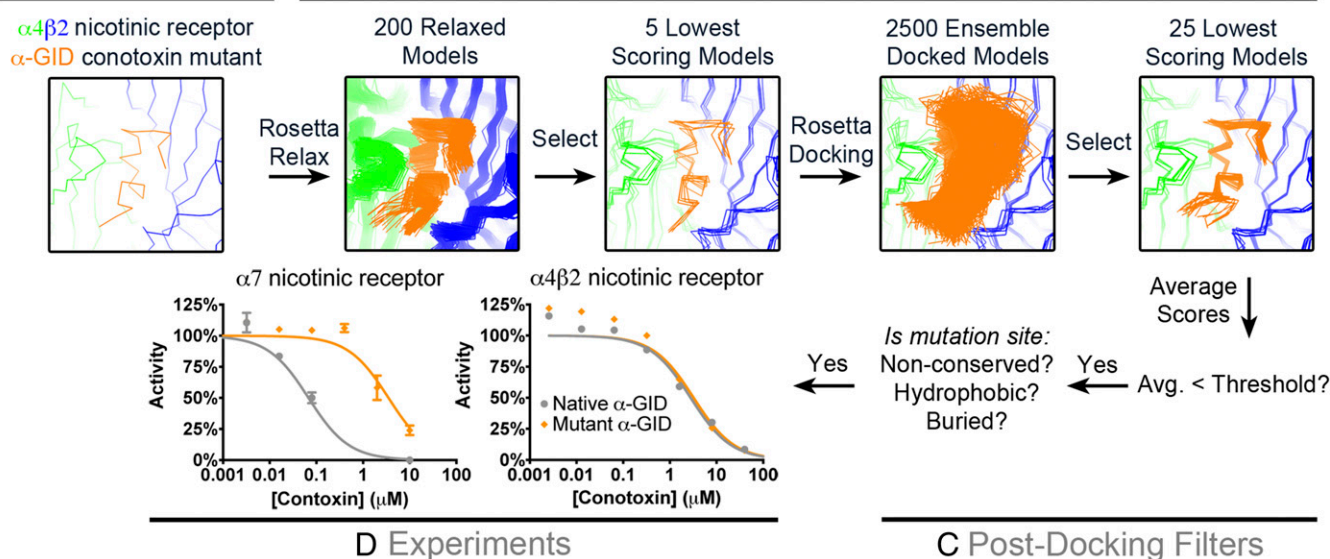
To address these dual challenges, we have developed a docking algorithm, ToxDock, in the context of the conotoxin  $\alpha$ -GID and the  $\alpha 4\beta 2$  nAChR. ToxDock employs extensive conformational

sampling of both the nAChR homology model and the conotoxin peptide to refine models of these complexes by combining two existing protocols in the Rosetta macromolecular modeling suite (28), Rosetta FastRelax and Rosetta FlexPepDock (29, 30) (Fig. 1). In the first stage of ToxDock, the entire complex is relaxed into lower energy states, followed by flexible docking of the conotoxin with ensemble docking in the second step (Fig. 1). In this manner, ToxDock is a refinement algorithm that can simulate a peptide toxin bound to an ion channel, given sufficient data for the bound complex. Using ToxDock, we investigate two principal questions. Can an algorithm that accounts for the conformational flexibility of a toxin peptide:ion channel complex be used to dock  $\alpha$ -GID and its analogs to the  $\alpha 4\beta 2$  nAChR with confidence? If so, can it also be used to discover new  $\alpha$ -GID derivatives bioactive at the  $\alpha 4\beta 2$  nAChR, and do any of these new derivatives have novel properties such as reduced potency at the human  $\alpha 7$  nAChR?

We undertook both benchmark and prospective docking screens of  $\alpha$ -GID analogs against a homology model of the  $\alpha 4\beta 2$  nAChR using ToxDock, followed by experimental assays at nAChR subtypes and structural characterization of  $\alpha$ -GID analogs. We also sought to obtain additional corroboration of our binding pose for  $\alpha$ -GID by docking it into the recently crystallized structure of the  $\alpha 4\beta 2$  nAChR in complex with nicotine (31). Docking  $\alpha$ -GID to this crystal structure could help confirm the binding mode of  $\alpha$ -GID, but studies have also shown that the binding pocket adopts distinctive interfaces depending on whether an agonist or antagonist is present (32–35). Finally, we performed redocking experiments on crystal structures of peptide toxins with two different ion channels and a binding protein to probe the generality of ToxDock. Our extensively benchmarked docking algorithm can overcome some of the fundamental limitations in docking the conotoxin  $\alpha$ -GID to the  $\alpha 4\beta 2$  nAChR, and may prove useful for other toxin peptide:ion channel systems.

## A Homology Model

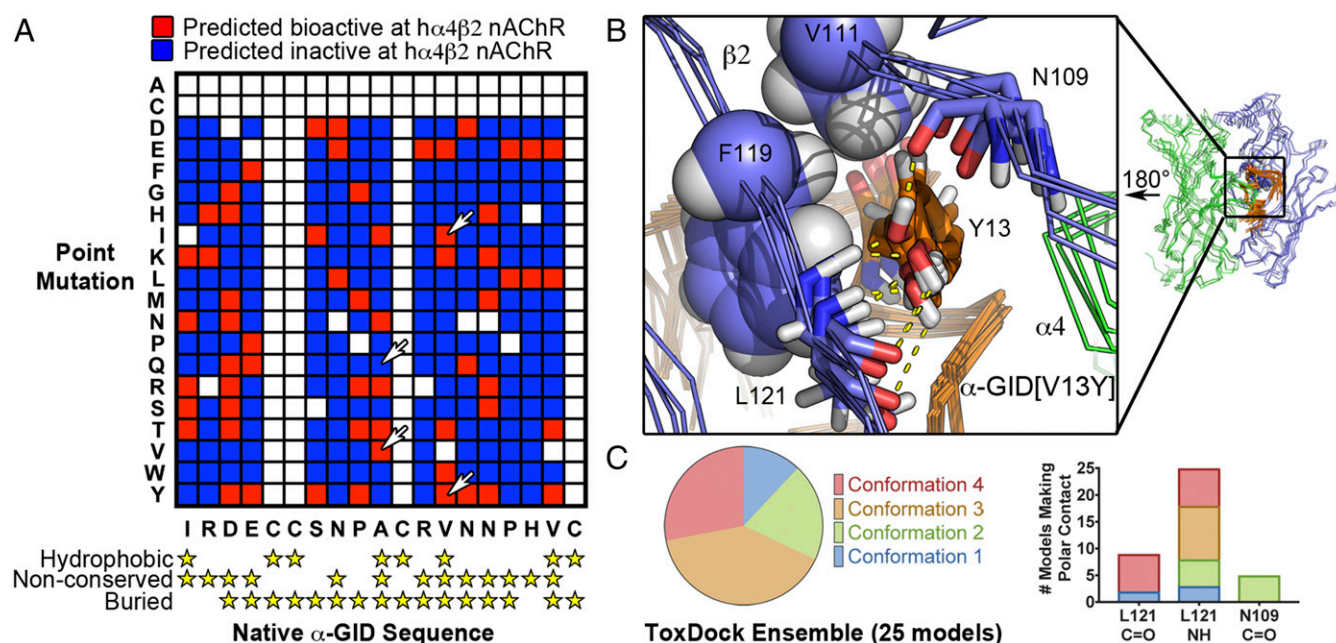
## B ToxDock



**Fig. 1.** Overview. (A) Initial homology model of an  $\alpha$ -GID conotoxin point mutant (orange) in complex with the extracellular domain of the  $\alpha 4\beta 2$  nAChR ( $\alpha 4$  subunit in green and  $\beta 2$  subunit in blue) is shown. (B) Each box shows the models of the complex considered at each step in the ToxDock protocol. Individual models are shown in ribbon representation and aligned. (C) Suitability of the conotoxin mutant for synthesis is further assessed by its average (ensemble) docking score to the receptor and the position of its point mutation. (D) Abilities of both the native (gray) and orange (mutant) conotoxins to block the responses of nicotinic receptor subtypes to ACh are assessed using fluorescent membrane potential assays. Avg., average.







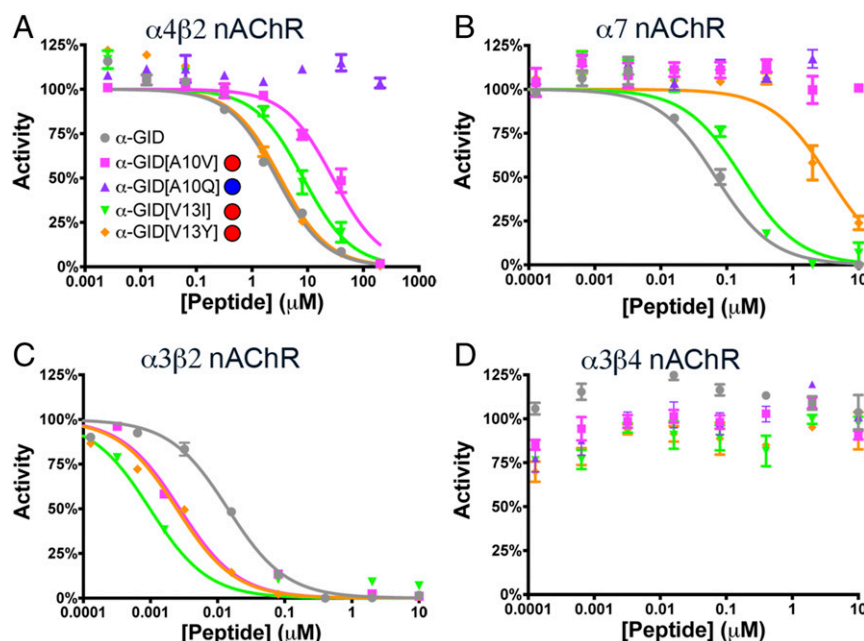
**Fig. 3.** Virtual screen and predicted binding modes. (A) Virtual screen with ToxDock of 256  $\alpha$ -GID point mutants against the  $\alpha 4\beta 2$  nAChR homology model. Each square represents an individual ToxDock simulation of a different  $\alpha$ -GID point mutant. Red squares are mutants predicted to be bioactive at the human  $\alpha 4\beta 2$  nAChR, blue squares are mutants predicted to be inactive, and white squares are mutants that were not simulated. The white arrows point to the four mutants ( $\alpha$ -GID[A10V],  $\alpha$ -GID[A10Q],  $\alpha$ -GID[V13I], and  $\alpha$ -GID[V13Y]) chosen for experimental characterization. Gold stars denote positions on  $\alpha$ -GID that are nonconserved, hydrophobic, or buried in the  $\alpha 4\beta 2$  nAChR homology model. (B) Twenty-five lowest scoring ToxDock models ("ToxDock ensemble") of the  $\alpha 4\beta 2$  nAChR homology model in complex with  $\alpha$ -GID[V13Y] are shown. Side chains are depicted as sticks, with the exception of the V111 and F119 side chains, which are shown as spheres for a single model. The  $\alpha 4$  subunit is green, the  $\beta 2$  subunit blue, and the peptide orange. Putative polar contacts are shown as dashed lines. All images were made in PyMOL. (C) Statistics of the ToxDock ensemble are shown, such as how many models in each conformation are included in the ToxDock ensemble (pie chart) and the polar contacts formed by each model (stacked bar chart). Numbering of residues is consistent with PDB ID code 5KXI.

$\alpha 4\beta 2$  nAChR (Fig. 3A), with scores equal to or below the threshold score derived from the benchmarking analysis ( $-637$  Rosetta energy units) that separates bioactive from inactive  $\alpha$ -GID mutants (SI Appendix, Fig. S3). As it was only feasible to experimentally test a subset of the  $\alpha$ -GID bioactive predictions, we applied a series of evolutionary and sequence-based filters to identify positions on  $\alpha$ -GID that were nonconserved, hydrophobic, and buried (Fig. 3A and SI Appendix, Fig. S4). These filters identified which of the 19 residues of  $\alpha$ -GID might be open to point mutations that would not negatively affect function. Previous studies have demonstrated that making point mutations at positions that are nonconserved, hydrophobic, and buried can increase the odds of successful protein design (42–44).

First, to identify the nonconserved  $\alpha$ -GID residues, a multiple sequence alignment of 20  $\alpha$ -conotoxins with the same spacing of cysteine residues (4/7) as  $\alpha$ -GID was performed. Due to their role in conferring structural stability and function, the cysteine residues at positions 5, 6, 11, and 19 were completely conserved. Serine 7 and Proline 9, which comprise the well-known S-X-P motif (3), also had 95% and 90% sequence identity, respectively (Fig. 3A and SI Appendix, Fig. S4A). The remaining 13 positions, which all had sequence identity of less than or equal to 75%, were considered nonconserved (SI Appendix, Fig. S4A; marked with yellow stars in Fig. 3A). Second, using Chimera (45), the Kyte–Doolittle hydrophobicity was calculated for each position on  $\alpha$ -GID, resulting in eight positions being identified as hydrophobic (hydropathy score  $> 0$ ) (SI Appendix, Fig. S4B; marked with yellow stars in Fig. 3A). Third, Chimera was also used to calculate the solvent-accessible surface area (SASA) for each residue on  $\alpha$ -GID in the ToxDock refined model. Three  $\alpha$ -GID positions were clearly solvent-exposed, while the remaining 16 had a per-residue SASA of less than  $150 \text{ \AA}^2$  and

were classified as buried (SI Appendix, Fig. S4C; marked with yellow stars in Fig. 3A). Two positions that met all three criteria of being nonconserved, hydrophobic, and buried were Ala-10 and Val-13 (Fig. 3A and SI Appendix, Fig. S4). Finally, phylogenetic analysis by maximum likelihood was performed on a set of 46  $\alpha$ -conotoxins from 24 different cone snail species. We found that position 13 is undergoing positive selection (SI Appendix, Table S2). Based on the three postdocking filters, we chose positions 10 (A10) and 13 (V13) of  $\alpha$ -GID for making point mutations.

Specifically, two  $\alpha$ -GID analogs with point mutations at position 10 (A10V and A10Q) and two analogs with point mutations at position 13 (V13I and V13Y) were chosen for synthesis and experimental assays. One of the peptides,  $\alpha$ -GID[A10Q], had a score above threshold and was predicted to be inactive (Fig. 3A). Three of these peptides,  $\alpha$ -GID[A10V],  $\alpha$ -GID[V13I], and  $\alpha$ -GID[V13Y], had scores below threshold and were predicted to be bioactive at the human  $\alpha 4\beta 2$  nAChR (Fig. 3A). These three predicted bioactives were chosen to select one relatively modest mutation ( $\alpha$ -GID[V13I]) and two "riskier" mutations ( $\alpha$ -GID[A10V] and  $\alpha$ -GID[V13I]) that might also have novel function. For example, the prediction that the V13Y mutation would be bioactive (Fig. 3A) was surprising, as previous reports had speculated that aromatic residues could not be accommodated at that position due to the tight fit between the conotoxin and receptor (9, 18). However, inspection of the 25 lowest energy models that comprise the ToxDock ensemble showed that the hydroxyl group on the tyrosine's aromatic ring is capable of participating in buried polar interactions with backbone atoms on the receptor that could make this interaction favorable (Fig. 3B). Notably, all 25 models that comprise the ToxDock ensemble made polar contacts with the backbone of  $\beta 2$ [L121]. In addition, depending on



**Fig. 4.** Fluorescent membrane potential experiments. Inhibition of the human  $\alpha 4 \beta 2$  (A),  $\alpha 7$  (B),  $\alpha 3 \beta 2$  (C), and  $\alpha 3 \beta 4$  (D) nAChRs as a function of peptide concentration for native  $\alpha$ -GID and the four peptides designed by ToxDock is shown. The peptides predicted to be bioactive at the  $\alpha 4 \beta 2$  nAChR have red circles by their names, and the peptide predicted to be inactive has a blue circle. All dose-response experiments were performed using fluorescent membrane potential assays on HEK cell lines stably transfected with the respective combination of nAChR subunits. ACh concentrations used to evoke the currents were 30  $\mu$ M for the  $\alpha 7$  nAChR, 0.5  $\mu$ M for the  $\alpha 4 \beta 2$  nAChR, 1  $\mu$ M for the  $\alpha 3 \beta 2$  nAChR, and 5  $\mu$ M for the  $\alpha 3 \beta 4$  nAChR.

which conformer was considered, additional polar contacts were possible (Fig. 3C). Finally, the tyrosine aromatic ring appeared to pack tightly against  $\beta 2$ [V111] and  $\beta 2$ [F119] (Fig. 3B).

**Experimental Characterization of  $\alpha$ -GID Derivatives Identified with ToxDock at nAChRs.** The four peptides identified by ToxDock ( $\alpha$ -GID[A10V],  $\alpha$ -GID[A10Q],  $\alpha$ -GID[V13I], and  $\alpha$ -GID[V13Y]) were synthesized, purified, and oxidatively folded as described previously (46) (SI Appendix, Fig. S5 A and B and Table S3). NMR structures of  $\alpha$ -GID[A10V] (SI Appendix, Fig. S5C) and  $\alpha$ -GID[V13Y] (SI Appendix, Fig. S5D) were solved (SI Appendix, Table S4) and indicated that both peptides adopted the expected conformations, with correct bonding of Cys-5 to Cys-11 and Cys-6 to Cys-19. The disulfide bond connectivities were determined based on the NMR data alone and were consistent with those found in wild-type  $\alpha$ -GID.

Each  $\alpha$ -GID derivative peptide was characterized experimentally with a fluorescent membrane potential assay to assess its ability to block the response to ACh of human  $\alpha 4 \beta 2$ ,  $\alpha 7$ ,  $\alpha 3 \beta 2$ , and  $\alpha 3 \beta 4$  nAChRs in stably transfected HEK cells (14) (Fig. 4 and Table 1). All three peptides predicted to be bioactive by ToxDock ( $\alpha$ -GID[A10V],  $\alpha$ -GID[V13I], and  $\alpha$ -GID[V13Y]) at the human  $\alpha 4 \beta 2$  nAChR exhibited significant inhibition of this receptor (Fig. 4A).  $\alpha$ -GID[V13Y] was the most potent analog, with an  $IC_{50}$  indistinguishable from the native peptide ( $IC_{50} = 3 \mu$ M), followed by  $\alpha$ -GID[V13I] and  $\alpha$ -GID[A10V] ( $IC_{50} = 8 \mu$ M and  $IC_{50} = 30 \mu$ M, respectively) (Fig. 4A and Table 1). Although  $\alpha$ -GID[A10V] had the weakest antagonism overall, it was still bioactive at the human  $\alpha 4 \beta 2$  nAChR. The peptide predicted by ToxDock to be inactive at the human  $\alpha 4 \beta 2$  nAChR,  $\alpha$ -GID[A10Q], displayed no antagonism of the receptor when tested at up to 200  $\mu$ M (Fig. 4A). The ToxDock predictions were also confirmed when the peptides were tested with two-electrode voltage clamp electrophysiology (TEVC) at the rat  $\alpha 4 \beta 2$  nAChR expressed in *Xenopus* oocytes (SI Appendix, Fig. S6A). Specifically,  $\alpha$ -GID[A10V],  $\alpha$ -GID[V13I], and  $\alpha$ -GID[V13Y] were all bioactive at the rat  $\alpha 4 \beta 2$  nAChR, while  $\alpha$ -GID[A10Q] was inactive (SI Appendix, Fig. S6A).

The TEVC results were reassuring, given that the rat and human  $\alpha 4 \beta 2$  nAChR receptor binding pockets are nearly identical (SI Appendix, Fig. S6B).

The inhibitory activity of the  $\alpha$ -GID derivative peptides was also examined at the human  $\alpha 3 \beta 2$ ,  $\alpha 3 \beta 4$ , and  $\alpha 7$  nAChRs (Fig. 4B–D and Table 1). Perhaps due to the high conservation between the binding pockets of the  $\alpha 3 \beta 2$  and  $\alpha 4 \beta 2$  nAChRs, which differ by only a single methyl between the  $\alpha 3$  and  $\alpha 4$  subunits in the vicinity of these mutations (9) (SI Appendix, Fig. S7), all of the peptides bioactive at the  $\alpha 4 \beta 2$  nAChR were also potent antagonists of the  $\alpha 3 \beta 2$  nAChR ( $IC_{50} = 0.003 \mu$ M,  $IC_{50} = 0.001 \mu$ M, and  $IC_{50} = 0.002 \mu$ M for  $\alpha$ -GID[A10V],  $\alpha$ -GID[V13I], and  $\alpha$ -GID[V13Y], respectively) (Fig. 4C). In addition, just as is the case with the native  $\alpha$ -GID, none of the peptides displayed any antagonism at the  $\alpha 3 \beta 4$  nAChR subtype (Fig. 4D and Table 1). Conotoxin  $\alpha$ -AUIB was used as a positive control for the  $\alpha 3 \beta 4$  nAChR and displayed the expected antagonism ( $IC_{50} = 0.1 \mu$ M; SI Appendix, Fig. S8).

While native  $\alpha$ -GID is a strong antagonist of the human  $\alpha 7$  nAChR ( $IC_{50} = 0.1 \mu$ M; Table 1), two of the peptides designed with ToxDock,  $\alpha$ -GID[A10V] and  $\alpha$ -GID[V13Y], had strongly diminished activity at the human  $\alpha 7$  nAChR up to 10  $\mu$ M (Fig. 4B).  $\alpha$ -GID[V13I] had no discernable inhibition at up to 10  $\mu$ M, although additional testing at 50  $\mu$ M revealed

**Table 1.** Summary of experimental findings at human nAChRs

Peptide	$\alpha 4 \beta 2$ nAChR		$IC_{50}$ at additional nAChRs, $\mu$ M		
	ToxDock prediction	$IC_{50}$ , $\mu$ M	$\alpha 7$	$\alpha 3 \beta 2$	$\alpha 3 \beta 4$
$\alpha$ -GID		3	0.1	0.01	>10
$\alpha$ -GID[A10V]	Bioactive	30	>10	0.003	>10
$\alpha$ -GID[A10Q]	Inactive	>200	>10	>10	>10
$\alpha$ -GID[V13I]	Bioactive	8	0.2	0.001	>10
$\alpha$ -GID[V13Y]	Bioactive	3	4	0.002	>10



significant (73%), but not complete, inhibition of the human  $\alpha 7$  nAChR at that concentration (*SI Appendix*, Fig. S9). It was also found that  $\alpha$ -GID[V13Y] had reduced antagonism at the  $\alpha 7$  nAChR, relative to the native, with an  $IC_{50}$  of 4  $\mu$ M (Fig. 4B and Table 1). Taken together, these data indicate that  $\alpha$ -GID[V13Y] and especially  $\alpha$ -GID[A10V] discriminate against the human  $\alpha 7$  nAChR.

**Docking  $\alpha$ -GID to the  $\alpha 4\beta 2$  nAChR Crystal Structure.** As this paper was being prepared, the crystal structure of the  $\alpha 4\beta 2$  nAChR in complex with the agonist nicotine became available (31). We examined if our pose for  $\alpha$ -GID in the homology model matched when docking it to this crystal structure (Fig. 5). Initially, there was a clash between loop C of the  $\alpha 4$  subunit and  $\alpha$ -GID after the conotoxin was superimposed onto the receptor. This clash was cleared by the Rosetta Relax phase of ToxDock (*SI Appendix*, Fig. S10). In the final pose after docking, the peptides in the crystal structure and homology model had a 1-Å C $\alpha$  rmsd when aligned to each other. The largest differences in the poses were in the N- and C-terminal regions, which are the most flexible parts of the conotoxin (*SI Appendix*, Fig. S11). Key aromatic residues in the  $\alpha 4\beta 2$  nAChR binding pocket (47) had the same positions and rotamers in the two models (Fig. 5). On the  $\alpha 4$  subunit, TyrC1, TyrC2, and TrpB all occupied the same positions and rotamers in the two models, and TrpD on the  $\beta 2$  subunit was in nearly the same position (Fig. 5). There were some differences, such as in the side-chain orientation of R12 on  $\alpha$ -GID. Nonetheless, these data suggested that  $\alpha$ -GID adopted largely the same docked poses in the crystal structure and homology model.

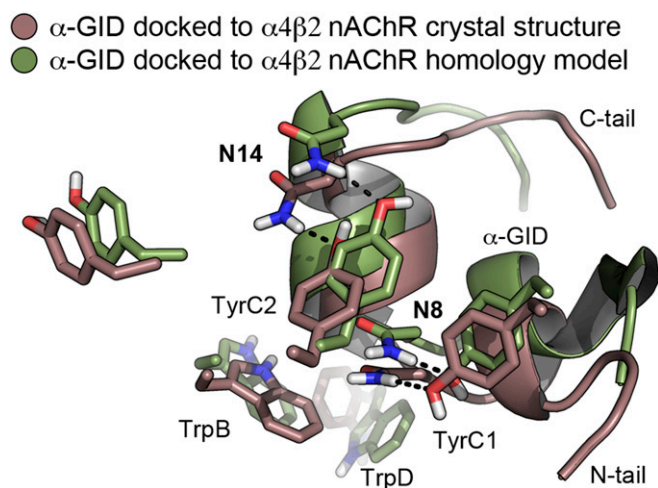
**ToxDock Applied to Other Toxin Peptide:Ion Channel Systems.** The generality of ToxDock was examined by measuring its redocking performance on three proteins of varying size and structure that bind peptide toxins and for which crystal structures have been determined. These were  $\alpha$ -PnIA[A10L,D14K] and AChBP ( $\alpha$ -PnIA:AChBP), Psalmotoxin and human acid-sensing ion channel (PcTx1:ASIC1a), and Charybdotoxin and a potassium channel (CTX:K<sup>+</sup>) (36, 48, 49) (Fig. 6 A–C and *SI Appendix*,

Figs. S12 and S13 A–C). In the redocking experiments, the peptide toxin was moved from its crystallographic pose using rotations and translations (*SI Appendix*, Fig. S12), and ToxDock was then applied to the perturbed pose to see if it could recover the native crystallographic orientation. In all three cases, at least one model in the ToxDock ensemble had a peptide C $\alpha$  rmsd of less than 2.6 Å and within a factor of 2 of the lowest overall peptide C $\alpha$  rmsd, indicating that the ensemble could capture near-native poses (*SI Appendix*, Fig. S13 D and E). Specifically, for the 16-residue  $\alpha$ -PnIA[A10L,D14K], the lowest peptide C $\alpha$  rmsd was 0.6 Å (Fig. 6A) and the mean over the ToxDock ensemble was  $1.6 \pm 0.5$  Å (*SI Appendix*, Fig. S13 A and E). For Psalmotoxin, which is more than twice the size, with 40 amino acids, the lowest peptide C $\alpha$  rmsd was 1.2 Å (Fig. 6B) and the mean peptide C $\alpha$  rmsd over the ToxDock ensemble was  $2.3 \pm 0.4$  Å (*SI Appendix*, Fig. S13 B and E). Charybdotoxin was the most challenging case overall, with a mean peptide C $\alpha$  rmsd over the ToxDock ensemble of  $5.7 \pm 2.0$  Å (*SI Appendix*, Fig. S13 C and E), but near-native poses were also captured as well; the lowest overall peptide C $\alpha$  rmsd was 1.3 Å (Fig. 6C) and the lowest peptide C $\alpha$  rmsd in the ensemble was 2.6 Å. Taken together, these data indicate that ToxDock is capable of modeling how medium to large peptide toxins dock to diverse ion channels.

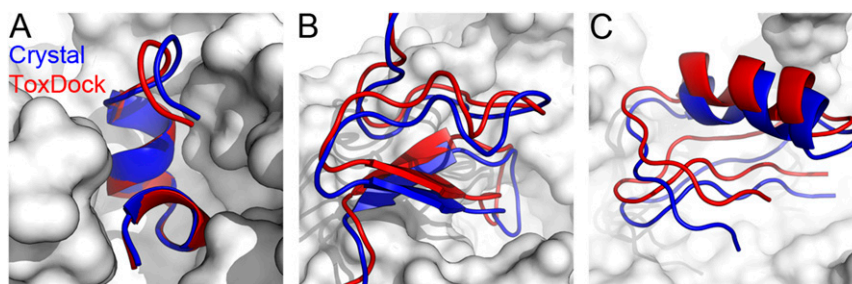
## Discussion

Peptide toxins isolated from venomous creatures have played a major role in elucidating the structure and function of ion channels (1). In addition to nAChRs (50), venom peptides are used to study the structure and function of sodium (51), potassium (49), calcium (52), ASIC (48), and transient receptor potential (TRP) channels (53). Given the utility of animal venom peptides as probe compounds, their estimated abundance in the millions, and the challenge of synthesizing and screening them (46), new computational methods are needed to accelerate their discovery and development. Next-generation sequencing and proteomic discovery strategies would also benefit from computational approaches that help to narrow down which of the large number of putative toxins they identify are potentially active or selective for a given target (54). Structure-based virtual screening could be particularly useful in these contexts, but its potential has remained unclear due to the size of peptide toxins and the lack of crystal structures of the ion channels that they target. We addressed this challenge in the context of a homology model of the  $\alpha 4\beta 2$  nAChR, a prototypical ligand-gated ion channel, and the conotoxin  $\alpha$ -GID that antagonizes it. Our two broad goals were to see if an algorithm for correctly docking  $\alpha$ -GID and its analogs to the  $\alpha 4\beta 2$  nAChR could be developed by accounting for the conformational flexibility of these complexes, and if so, whether it could be used to discover new  $\alpha$ -GID derivatives and those with functionally interesting properties.

One of our main findings is that, with extensive treatment of its conformational flexibility, a homology model of nAChR with less than 30% sequence identity to its template can be refined so that it is capable of discriminating between inactive and bioactive mutants of a conotoxin (Figs. 1 and 2). Using functional assays, we found that all four predictions from a virtual screen based on the homology model (Fig. 3) were correct (Fig. 4 and Table 1). The ability to identify bioactive  $\alpha$ -GID mutants is particularly useful because the majority of mutants of  $\alpha$ -GID that have been synthesized and tested are inactive. Two of the  $\alpha$ -GID point mutants identified in the virtual screen,  $\alpha$ -GID[A10V] and  $\alpha$ -GID[V13Y], have substantially reduced antagonism at the human  $\alpha 7$  nAChR. Using ToxDock, we showed that docking  $\alpha$ -GID into two ostensibly dissimilar structures, the  $\alpha 4\beta 2$  nAChR crystallized with an agonist and an  $\alpha 4\beta 2$  nAChR homology model based on the AChBP/conotoxin template structure, resulted in similar poses for the peptide (Fig. 5). Finally, we confirmed the broader applicability of our docking program on



**Fig. 5.** Docking to  $\alpha 4\beta 2$  nAChR crystal structure in the desensitized state. The lowest energy poses of  $\alpha$ -GID docked into the  $\alpha 4\beta 2$  nAChR homology model based on AChBP (green) and agonist-bound crystal structure (brown) with ToxDock are shown. Key aromatic residues on the nAChR are shown as sticks and labeled (TrpB, TyrC1, TyrC2, and TrpD). Residues on  $\alpha$ -GID that participate in putative polar interactions (black dashed lines) are shown in sticks and labeled with bold text (N8 and N14). The receptor's backbone is omitted for clarity.



**Fig. 6.** Redocking experiments on diverse crystal structures. The crystallographic (blue) and lowest scoring redocked (red) pose of the toxin peptide in the ToxDock ensemble are shown for  $\alpha$ -PnIA[A10L,D14K] bound to AChBP (A), Psalmotoxin bound to ASIC1a (B), and Charybdotoxin bound to a potassium channel (C). The surfaces of the binding protein and ion channels are shown in white. Renderings were made in PyMOL.

crystal structures of peptide toxins in complex with ion channels and a binding protein (Fig. 6). These findings are discussed in more detail below, along with possible future directions and improvements to the ToxDock method.

**Importance of Conformational Sampling in Modeling the  $\alpha$ -GID: $\alpha$ 4 $\beta$ 2 nAChR Complex.** Our benchmarking experiments demonstrated that as more conformational sampling was included in the docking procedure, the ability of the refined homology model to discern between bioactive and inactive  $\alpha$ -GID analogs improved (Fig. 24). The identification of  $\alpha$ -GID[V13Y] in the virtual screen also highlights how this extensive sampling is critical for making new predictions borne out by experiment. In the case of  $\alpha$ -GID[V13Y], if the score for its interaction with the  $\alpha$ 4 $\beta$ 2 nAChR had only used models based on a single conformation of the complex, then the favorable nature of its binding would have been underestimated. This is because different conformations of the channel and peptide captured different interactions, but no single conformation comprised all three interactions that were possible between the tyrosine hydroxyl group and the backbone amide and carbonyl groups of the receptor (Fig. 3C). Specifically, models based on conformation 2 captured that the hydroxyl group can form polar contacts with the amide group of  $\beta$ 2[L121] and the carbonyl group of  $\beta$ 2[N109], but did not predict any interactions with the carbonyl group of  $\beta$ 2[L121]. Conversely, models based on conformation 4 had polar contacts with the amide and carbonyl of  $\beta$ 2[L121], but not with  $\beta$ 2[N109]. Thus, modeling  $\alpha$ -GID[V13Y] as a superposition of these different conformers allowed the favorable nature of its interactions with the  $\alpha$ 4 $\beta$ 2 nAChR to be captured more fully than if only a single conformation had been used and led to the correct prediction that it would be bioactive. Experimental validation of this prediction was particularly gratifying, given that the nearly identical  $\alpha$ -GID[V13F] has been shown to be inactive at the rat  $\alpha$ 4 $\beta$ 2 nAChR (9).

Extensive computational sampling also played a role when docking  $\alpha$ -GID to the agonist-bound crystal structure of the  $\alpha$ 4 $\beta$ 2 nAChR (Fig. 5). It was unclear if docking  $\alpha$ -GID to the  $\alpha$ 4 $\beta$ 2 nAChR agonist-bound structure could lead to agreement with the homology model because the nAChR binding pocket undergoes significant rearrangement depending on whether an agonist or antagonist is present (25, 33, 34). We found that the poses of  $\alpha$ -GID in the  $\alpha$ 4 $\beta$ 2 nAChR crystal structure or AChBP-derived homology model were similar after refinement with ToxDock, and that initial steric collisions between the conotoxin and loop C were alleviated through global relaxation of the receptor backbone. The concordance between the poses of  $\alpha$ -GID in the two models suggests they have captured the binding pose of  $\alpha$ -GID reasonably well. More broadly, our finding raises the possibility that the extracellular domains of agonist-bound nAChR structures could potentially be used as an alternative to, or in conjunction with, AChBP-derived homology models

for docking conotoxins, provided they are subject to appropriate refinement.

**$\alpha$ -GID Analogs with Useful Properties.** A key finding from this study is that two of the three  $\alpha$ -GID analogs identified by the virtual screen,  $\alpha$ -GID[A10V] and  $\alpha$ -GID[V13Y], had the long-sought-after property of reduced activity at the human  $\alpha$ 7 nAChR without sacrificing their activity at the  $\alpha$ 4 $\beta$ 2 nAChR. These findings suggest that even a relatively modest virtual screen of 256  $\alpha$ -GID point mutants, further focused by the use of post-docking filters, can uncover novel peptide ligands with favorable properties. This result agrees with the recent  $\alpha$ -GID modeling study of Suresh and Hung (15). Using sophisticated umbrella sampling and free-energy calculations, they came to the conclusion that a point mutation on the N-terminal tail, a different region of  $\alpha$ -GID than the one we focused on, could potentially also impart specificity for the  $\alpha$ 4 $\beta$ 2 nAChR (15). More generally, our findings are in accord with published results of Hogg et al. (55) and Luo et al. (56), who showed that a single-residue alteration in a similar conotoxin,  $\alpha$ -PnIA, is sufficient for switching its nAChR subtype selectivity.

A mechanistic hypothesis for why  $\alpha$ -GID[V13Y] has reduced activity at the human  $\alpha$ 7 nAChR can be developed by comparing the binding pockets of the  $\alpha$ 4 $\beta$ 2 and  $\alpha$ 7 nAChRs. In the case of  $\alpha$ -GID[V13Y], the lack of a T-shaped pi-stack with  $\beta$ 2[F119] (which is a glutamine in the  $\alpha$ 7 nAChR) and the additional steric hindrance from a leucine in the  $\alpha$ 7 nAChR (which is Val-111 in the  $\alpha$ 4 $\beta$ 2 nAChR) positioned close to the tyrosine side chain may have sufficed to cause the loss of activity at the  $\alpha$ 7 nAChR. In a broad sense, chemical novelty conferred new biological properties, which is often the case for docking and virtual screening studies of small molecules (57). Mutations at other positions may also result in  $\alpha$ -GID derivatives that discriminate against the  $\alpha$ 7 nAChR. For example, the hydrophobic, conservation, and buried residue filters used to pinpoint positions 10 and 13 for making mutations were also in agreement that position 18 would be a viable choice for making mutations. Since this is the position at which the  $\alpha$ -GID mutant selective for the rat  $\alpha$ 4 $\beta$ 2 nAChR vs. the  $\alpha$ 3 $\beta$ 2 nAChR was discovered, it implies that integrating additional sources of information such as evolutionary data can help to accelerate the discovery of optimized mutations by pinpointing the most promising sites to search for mutations (54, 58).

Combining the mutations identified in this study with those previously found (9–11) may facilitate the development of an  $\alpha$ -GID derivative that is truly selective for the  $\alpha$ 4 $\beta$ 2 nAChR. However, to be effective as a tool compound, the potency of such an  $\alpha$ -GID analog at the  $\alpha$ 4 $\beta$ 2 nAChR would also need to be increased into the low nanomolar range, commensurate with other classical conotoxin probes such as MII (4). One particularly promising route for achieving this aim could be developing  $\alpha$ -GID analogs with noncanonical amino acids (NCAAs), which



Hopping et al. (18) used to increase the potency of the conotoxin  $\alpha$ -PnIA at the  $\alpha 7$  nAChR. The recent incorporation of NCAs into Rosetta could also facilitate such an effort (59). Finally, the design of more potent  $\alpha$ -GID analogs could exploit our finding that  $\alpha$ -GID[V13Y] is  $\sim 1,000$ -fold weaker at the  $\alpha 4\beta 2$  nAChR than at the  $\alpha 3\beta 2$  nAChR despite only one residue differing between the  $\alpha 3$  and  $\alpha 4$  subunits in the vicinity of  $\alpha$ -GID[Y13]. Since this residue is a threonine on the  $\alpha 4$  subunit and a serine on the  $\alpha 3$  subunit, it appears that the presence of a single additional methyl group in  $\alpha 4$  may account for the much weaker potency of  $\alpha$ -GID[V13Y] at the  $\alpha 4\beta 2$  nAChR. Thus, a design consideration when choosing canonical or noncanonical mutations to  $\alpha$ -GID appears to be avoiding steric clashes with  $\alpha 4$ [Thr157].

As specialized methods for peptide toxin synthesis and nAChR screening continue to be developed, such as parallelized oxidative folding (60) and high-throughput electrophysiology (61), it may become possible to test the majority of bioactive predictions made by a virtual screen with ToxDock to discover additional analogs with novel activity.

**Improvements to ToxDock.** Two improvements to ToxDock could be made that would enhance its generality and utility. First, both computational and experimental efforts have demonstrated that the interaction of peptide toxins with the cellular membrane can play a key role in their docking to ion channels (62–64). Thus, the addition of an explicit or implicit representation of the membrane to ToxDock might help it to model peptide toxin:ion channel docking for systems like TRP and sodium channels where membrane interactions are important. Second, combining ToxDock with a protocol for estimating relative binding affinities would enable ToxDock to go beyond prediction of bioactivity to guide improvements in potency of peptide toxin analogs. Since one of the most important components of an affinity calculation is a realistic binding pose, which ToxDock appears to achieve reasonably well for both homology models and crystal structures, it is possible that using ToxDock-derived poses as the basis for free-energy perturbation (FEP) calculations could yield estimates of relative binding affinities for peptide toxin mutants. Recent efforts suggest that computing relative binding affinities for peptide toxin and channel systems is within the domain of applicability of FEP (65). However, given that FEP calculations are sensitive to the amount of conformational reorganization that a protein undergoes (66, 67), it is possible that specialized enhanced sampling protocols may need to be developed to fully accommodate the flexibility of conotoxins like  $\alpha$ -GID.

In summary, this study demonstrates the feasibility of a combined computational and experimental approach for an  $\alpha$ -conotoxin and nAChR and opens up the possibility of it being applied to the growing number of identified peptide toxins and their ion channel targets. Moreover, it highlights the value in using extensive conformational sampling to model peptide toxin:ion channel complexes and what kind of functional and structural data are necessary for validating those efforts. To facilitate such studies, we have made ToxDock publicly available as a server that can be freely and easily used by any member of the academic research community ([rosie.rosettacommons.org/tox\\_dock](http://rosie.rosettacommons.org/tox_dock)).

## Materials and Methods

**Homology Modeling.** The  $\alpha 4\beta 2$  nAChR extracellular domain dimer was modeled using Modeller (68) version 9.11 based on the X-ray structures of AChBP from *A. californica* in complex with the peptide toxins  $\alpha$ -PnIA [A10L, D14K],  $\alpha$ -Iml,  $\alpha$ -BuIA, and  $\alpha$ -TxIA[A10L] [Protein Data Bank (PDB) ID codes 2BR8, 2C9T, 2BYF, 4EZ1, and 2UZ6, respectively] (33, 36–38). The alignment between AChBP and the nAChR sequences was based on a multiple sequence alignment presented previously (47). The sequence identity between AChBP and the extracellular domains was 27.8% and 23.4% for the  $\alpha 4$  and  $\beta 2$  subunits, respectively.

**Docking and Virtual Screening.** Docking and virtual screening of  $\alpha$ -GID and  $\alpha$ -GID analogs against the  $\alpha 4\beta 2$  nAChR homology model was performed using our ensemble-docking algorithm, ToxDock, using Rosetta revision 57232 and the talaris2013 weight set. ToxDock is available at [rosie.rosettacommons.org/tox\\_dock](http://rosie.rosettacommons.org/tox_dock). Models of  $\alpha$ -GID mutants were based on its NMR structure (PDB ID code 1MTQ) (11). Apo rotamers for AChBP were obtained from PDB ID code 2Y7Y (69).

**Molecular Dynamics Simulation.** Molecular dynamics simulations were completed using DESMOND version 4.4 (70) with the OPLS3 forcefield after preparation with the System Builder workflow (Schrödinger).

**Peptide Synthesis.** All peptides were synthesized by The Rockefeller University Proteomics Center using Fmoc solid-phase peptide synthesis as previously described (46).

**NMR Solution Structure Derivation.** Standard homonuclear NMR techniques were used to solve the solution structures of  $\alpha$ -GID[A10V] and  $\alpha$ -GID[V13Y]. Data were collected at 7 °C, and structures were obtained via simulated annealing and explicit water refinement, using nuclear Overhauser effect (NOE) distance restraints and chemical shift-derived angular restraints as input for the calculations. The temperature of 7 °C was chosen to maximize the number and intensity of observable NOE distance restraints and also for consistency with previous NMR studies of  $\alpha$ -GID (11). For final calculations, hydrogen and disulfide bonds consistent with the NMR data and preliminary structures were also included in the calculations.

**Fluorescent Membrane Potential Assay.** For functional tests of nAChRs expressed in HEK cells, we used a FLEXstation (Molecular Devices) bench-top scanning fluorimeter as described by Kuryatov et al. (14). To increase the expression level of  $\alpha 3\beta 2$  nAChRs, the plates were incubated at 29 °C for 20 h before being tested. The  $\alpha 7$  cell line was tested as described by Kuryatov et al. (71). A membrane potential fluorescent indicator kit (Molecular Devices) was used according to the manufacturer's protocols. Peptides were added 1 h before recordings. ACh concentrations used to evoke currents were 30  $\mu$ M for the  $\alpha 7$  nAChR, 0.5  $\mu$ M for the  $\alpha 4\beta 2$  nAChR, 1  $\mu$ M for the  $\alpha 3\beta 2$  nAChR, and 5  $\mu$ M for the  $\alpha 3\beta 4$  nAChR. All recordings were performed at 29 °C. Each data point was averaged from three responses from separate wells. IC<sub>50</sub> values were calculated from a nonlinear fit of the Hill equation to the data (GraphPad Prism version 7.0). Native  $\alpha$ -GID was purchased from Smartox Biotechnology. Conotoxins AulB and MII were purchased from Tocris.

**Electrophysiological Measurements.** The nAChR cDNAs were provided by J. Patrick, Baylor College of Medicine, Houston, TX, and subcloned into the oocyte expression vector pNK52. The cRNA was synthesized with the SP6 mMessage mMachine kit (Ambion), and *Xenopus laevis* (Nasco International) oocytes were injected with 50-nL aliquots of cRNA (0.05 mg/mL). The nAChR subunits were mixed at the ratio of 5:1 ( $\alpha 4$ : $\beta 2$ ).

Antagonist dose–response curves were measured as described (72) in ND96 (96 mM NaCl, 2 mM KCl, 1 mM CaCl<sub>2</sub>, 1 mM MgCl<sub>2</sub>, and 5 mM Hepes at pH 7.4). Briefly, current responses to 100  $\mu$ M ACh were recorded at –70 mV using a Turbo Tec 05X Amplifier (NPI Electronic) and Cell Works software. A fast and reproducible solution exchange (<300 ms) was achieved with a 50- $\mu$ L funnel-shaped oocyte chamber combined with a fast solution flow ( $\sim 150$   $\mu$ L s<sup>–1</sup>) fed through a custom-made manifold mounted immediately above the oocyte. Agonist pulses were applied for 2 s at 4-min intervals. Peptides were applied for 3 min in a static bath. IC<sub>50</sub> values were calculated from a nonlinear fit of the Hill equation to the data (GraphPad Prism version 6.0). Data are presented as mean  $\pm$  SE from at least three experiments.

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