Macroscopic and Microscopic Activation of α7 Nicotinic Acetylcholine Receptors by the Structurally Unrelated Allosteric Agonist-Positive Allosteric Modulators (ago-PAMs) B-973B and GAT107

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ABSTRACT

B-973 is an efficacious type II positive allosteric modulator (PAM) of α7 nicotinic acetylcholine receptors that, like 4BP-TQS and its active isomer GAT107, can produce direct allosteric activation in addition to potentiation of orthosteric agonist activation, which identifies it as an allosteric activating (ago)-PAM. We compared the properties of B-973B, the active enantiomer of B-973, with those of GAT107 regarding the separation of allosteric potentiation and activation. Both ago-PAMs can strongly activate mutants of α7 that are insensitive to standard orthosteric agonists like acetylcholine. Likewise, the activity of both ago-PAMs is largely eliminated by the M254L mutation in the putative transmembrane PAM-binding site. Allosteric activation by B-973B appeared more protracted than that produced by GAT107, and B-973B responses were relatively insensitive to the noncompetitive antagonist mecamylamine compared with GAT107 responses. Similar differences are also seen in the single-channel currents. The two agents generate unique profiles of full-conductance and subconductance states, with B-973B producing protracted bursts, even in the presence of mecamylamine. Modeling and docking studies suggest that the molecular basis for these effects depends on specific interactions in both the extracellular and transmembrane domains of the receptor.

Initial attempts to record α-BTX-sensitive nicotinic currents from peripheral neurons were largely unsuccessful until efforts were made to achieve especially fast agonist applications (Zhang et al., 1994), something that was later understood to be due to the particularly rapid desensitization of α7 receptors exposed to high concentrations of agonist (Papke and Thinschmidt, 1998; Uteshev et al., 2002). Unlike other nAChRs, which produce their maximal peak current responses upon rapid saturation of all their agonist binding sites, α7 receptors progressively enter nonconducting states at high levels of agonist occupancy so that peak currents occur in a narrow range of agonist concentration and often occur before fast drug application is complete (Papke and Porter Papke, 2002; Williams et al., 2012). The unique desensitization properties of α7 receptors fundamentally limit the ion-channel currents that can be produced by typical agonists; however, the existence of the receptor’s unusual desensitized conformations has also made α7 receptors uniquely sensitive

Introduction

Nicotine and α-bungarotoxin (α-BTX) bind with high affinity to two different populations of receptors in the brain (Clarke et al., 1985; Jones et al., 1999). The molecular cloning of nicotinic acetylcholine receptor (nAChR) subunits led to identification of the brain receptors with high affinity for nicotine as protein complexes composed of α and β subunits (Wada et al., 1988) containing primarily α4 and β2 subunits (Wada et al., 1989). The α-BTX binding sites in the brain were later shown to be associated with α7 subunits, which can form functional homomeric receptors (Couturier et al., 1990).

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ABBREVIATIONS: ABNR, Adopted Basis Newton-Raphson; ACh, acetylcholine; ago-PAM, allosteric activating positive allosteric modulator; 4BP-TQS, 4-(4-bromophenyl)-3a,4,5,9b-tetrahydro-3H-cyclopenta[c]quinoline-8-sulfonamide; B-973B, (3-(3,4-difluorophenyl)-N-(1-(6-(4-(pyridin-2-yl)piperazin-1-yl)pyrazin-2-yl)ethyl)propenamide); α-BTX, α-bungarotoxin; CAP, cholinergic anti-inflammatory pathway; CTPE, constant pressure constant temperature ensemble; DAA, direct allosteric activation; Dc, PAM-insensitive desensitization; Ds, PAM-sensitive desensitization; DMEM, Dulbecco’s modified Eagle’s medium; ECD, extracellular domain; GAT107, (3aR,4S,9bS)-4-(4-bromophenyl)-3a,4,5,9b-tetrahydro-3H-cyclopenta[c]quinoline-8-sulfonamide; IFD, induced-fit docking; M2, second transmembrane domain; M3, third transmembrane domain; M4, fourth transmembrane domain; nAChR, nicotinic acetylcholine receptor; PAMs, positive allosteric modulators; PNU-120596 (1-[5-chloro-2-(4-dimethoxyphenyl)-3-[5-(methylsioxazol-3-yl)-urea]; QX-222, 2-[(2,6-dimethylphenyl)aminio]-N,N,N-trimethyl-2-oxoethaniminium chloride; TMD, transmembrane domain; TQS, 3a,4,5,9b-tetrahydro-4-(1-naphthalenyl)-3H-cyclopenta[c]quinoline-8-sulfonamide.
to selective positive allosteric modulators (PAMs) (Williams et al., 2011c), some of which eliminate the limitations imposed by desensitization by converting desensitized receptors into receptors that can show particularly long bursts of openings (Williams et al., 2011b, 2012). Efficacious type II PAMs (Grønlien et al., 2007), like PNU-120596, perturb the stability of agonist-induced desensitization by binding to a site within the receptor’s transmembrane domain (TMD) (Young et al., 2008). In the absence of agonist, application of a PAM to wild-type α7 receptors will not produce activation but can prime receptors to generate a large potentiated response to a subsequent agonist application. The duration of such priming effects depends on the specific PAM and the concentration applied (Williams et al., 2011b; Gill et al., 2012; Papke et al., 2014b).

Whereas typical PAMs, like PNU-120596 and TQS (Grønlien et al., 2007), produce enhanced activation through coupling the transmembrane PAM site with binding of orthosteric agonists, other agents structurally related to TQS produce activation without the requirement of an orthosteric agonist (Gill et al., 2011, 2012; Gill-Thind et al., 2015), identifying them as ago-PAMs. One of the most studied ago-PAMs is GAT107, the active isomer of 4BP-TQS (Papke et al., 2014b, 2018; Bagdas et al., 2016; Horenstein et al., 2016). Several lines of evidence indicate that the direct allosteric activation (DAA) produced by GAT107, when applied without agonist, relies on binding to both the transmembrane PAM site and a novel DAA binding site in the extracellular domain (ECD) (Horenstein et al., 2016). Whereas dissociation of GAT107 from the DAA site is relatively rapid, receptors remained primed for potentiated activation by orthosteric agonists for a prolonged period (Papke et al., 2018).

Numerous ago-PAMs and allosteric antagonists have been reported that are structurally related to TQS (Gill-Thind et al., 2015; Horenstein et al., 2016). The recent report on B-973 (Post-Munson et al., 2017), an ago-PAM structurally unrelated to TQS, offers the opportunity to explore both the common features and the differences in the allosteric activation of α7 produced by these agents. In this study, we expanded on the macroscopic characterization of B-973 using the active isomer B-973B (Garai et al., in press) for comparison with published studies of GAT107. We also studied the activity of these two agents on the level of single-channel currents. Efficacious PAMs, like PNU-120596, in combination with agonists, induced protracted bursts of channel activation (Williams et al., 2011b, 2012; Andersen et al., 2016) with both full-conductance and subconductance states, and initial characterizations of 4BP-TQS and GAT107 single-channel activity showed similar results (Paleczynska et al., 2012; Horenstein et al., 2016). We additionally investigated the effects of noncompetitive nAChR antagonists on the macroscopic and microscopic currents stimulated by the ago-PAMs.

By evaluating our results in the context of high-resolution molecular models, the present studies provided the opportunity for new insights into the molecular mechanisms of receptor function and new avenues for future drug development.

Materials and Methods

Chemicals. Solvents and reagents were purchased from Sigma-Aldrich (St. Louis, MO). Cell-culture supplies were from Life Technologies (Carlsbad, CA). Dulbecco’s modified Eagle’s medium (DMEM) was from Corning (Tewksbury, MA). Fetal bovine serum was purchased from Omega Scientific (Tarzana, CA). Antibodies 2-aminoo-2,7-dideoxy-D-glycero-a-D-gluco-heptopyranosyl-(1→4)-O-[3-deoxy-4-C-methyl-3-(methylamino)-β-L-arabinopyranosyl-(1→6)]-2-deoxy-D-streptamine disulfate (GAT18) and hygromycin B were from Gibco and Invitrogen (Thermo Fisher, Waltham, MA), respectively. The Hank’s balanced salt solution (HBSS) (Life Technologies) contained the following components expressed in millimolars: 1.26 CaCl2, 0.493 MgCl2, 0.407 MgSO4, 5.53 KCl, 0.441 KH2PO4, 4.17 NaHCO3, 137.93 NaCl, 0.338 Na2HPO4, and 5.56-glucose. EDTA was purchased from Sigma-Aldrich. GAT107 ((3aR,4S,9bS)-4-(4-bromophenyl)-3a,4,5,9b-tetrabhydro-3H-cyclopenta[c]quinoline-8-sulfonamide) was synthesized as described previously (Kulkarni and Thakur, 2013; Thakur et al., 2013). B-973B (3-(3,4-difluorophenyl)-N-(1-((6-(4-(pyridin-2-yl)piperazin-2-yl)pyrazin-2-yl)ethyl)propenamide) was synthesized by Dr. Ganesh Thakur as previously described (Garai et al., 2018). Mecamylamine ([1S,2R,4R]-N,2,3,3-tetramethylbicyclo[2.2.1]heptan-2-amine) and tetracaine were purchased from Sigma. PNU-120596 (1-(5-chloro-2,4-dimethoxyphenyl)-3-(5-methylisoxazol-3-yl)-urea) was synthesized by Dr. Jingyi Wang and Dr. Kinga Chojnacka as described previously (Williams et al., 2011b). Fresh ACh stock solutions were made each day of experimentation. PNU-120596, GAT107, and B-973B stock solutions were prepared in dimethylsulfoxide (DMSO), stored at −20°C, and used for up to a year. PNU-120596, GAT107, B-973B, and mecamylamine solutions were prepared freshly at the desired concentration from the stored stock solutions and dissolved in the internal solution containing (in millimolars) 185 NaCl, 5 KCl, 2 CaCl2, 5 HEPES (4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid; Sigma Aldrich), 0.001 atropine and 10-glucose, pH 7.3. Tested solutions contained either 10 μM of GAT107 or 10 μM B-973B in internal solution, alone or with 30 μM mecamylamine.

Heterologous Expression of α7 nAChRs in Xenopus Oocytes. The cDNA clones of human α7 nAChR and human resistance-to-cholinesterase 3 (Ric-3) were provided by Dr. Jon Lindstrom (University of Pennsylvania, Philadelphia, PA) and Dr. Millet Treinin (Hebrew University, Jerusalem, Israel), respectively. After linearization and purification of the plasmid cDNAs, RNAs were prepared using the mMessage mMACHINE in vitro RNA transcription kit (Ambion, Austin, TX). Alpha7 mutants were made using the QuikChange Site-Directed Mutagenesis Kit (Agilent Technologies, Santa Clara, CA) as described (Papke et al., 2011). The Y99C and C190S mutants were made in α7C116S to ensure no spurious disulfide bonding, and the C116S mutation had no effect on α7 expression or responses compared with wild-type α7 (Papke et al., 2011).

Oocytes were surgically removed from mature female Xenopus laevis frogs (Nasco, Ft. Atkinson, WI) and injected with RNAs of α7 nAChR and Ric-3 as described previously (Papke and Stokes, 2010). The Ric-3 chaperone protein can improve and accelerate α7 expression with no effects on the pharmacologic properties of the receptors (Halevi et al., 2003). Frogs were maintained in the animal care service facility of the University of Florida, and all procedures were approved by the University of Florida Institutional Animal Care and Use Committee. All studies were carried out in accordance with the Guide for the Care and Use of Laboratory Animals as adopted and promulgated by the U.S. National Institutes of Health. In brief, the frog was first anesthetized for 15–20 minutes in 1.5-liter frog tank water containing 1 g of 3-aminoenzoate methanesulfonate (MS-222) buffered with sodium bicarbonate. The harvested oocytes were treated with 1.4 mg/ml collagenase (Worthington Biochemicals, Freehold, NJ) for 3 hours at room temperature in a calcium-free Barth’s solution (88 mM NaCl, 1 mM KCl, 2.38 mM NaHCO3, 0.82 mM MgSO4, 15 mM HEPES, and 12 mg/l tetracycline, pH 7.6) to remove the follicular layer. Stage 5 oocytes were subsequently isolated and injected with 50 nl of 6-ng α7 nAChR subunit cRNA and 3 ng Ric-3 cRNA. Recordings were carried out 1–7 days after injection.

Two-Electrode Voltage-Clamp Electrophysiology of Oocytes. Experiments were conducted using OupaXpress 6000A (Molecular Devices, Union City, CA) (Papke and Stokes, 2010). Both the voltage and current electrodes were filled with 3 M KCl. Oocytes were voltage-clamped
at ~60 mV. The oocytes were bath-perfused with Ringer’s solution (115 mM NaCl, 2.5 mM KCl, 1.8 mM CaCl$_2$, 10 mM HEPES, and 1 μM ATP, pH 7.2) with a flow rate of 2 ml/min. To evaluate the effects of experimental compounds on ACh-evoked responses of α7 nAChRs expressed in oocytes, two initial control responses to applications of ACh were recorded before test applications of experimental drugs alone or coapplied with the ACh. The agonist solutions were applied from a 96-well plate via disposable tips, and the drugs were either coapplied with ACh by the OpusXpress pipette delivery system or bath-applied using the OpusXpress system to switch the running buffer. Drug applications were 12 seconds long, followed by a 181-second washout period, and usually alternated between controls and test solutions. The control concentration of ACh was 60 μM for wild-type α7. After experimental drug applications, follow-up control applications of ACh were made to determine primed potentiation, desensitization, or rundown of the receptors.

Data were collected at 50 Hz, filtered at 20 Hz, and analyzed by Clampfit 9.2 or 10.0 (Molecular Devices) and Excel (Microsoft, Redmond, WA). Data were expressed as means ± S.E.M. from at least five oocytes for each experiment, and plotted with Kaleidagraph 4.5.2 (Abelbeck Software, Reading, PA). Concentration response data were fit to the Hill equation using the Levenberg-Marquardt algorithm. Multicell averages were calculated for comparisons of complex responses. Averages of the normalized data were calculated for each of the 10,322 points in each of the 206.44-second traces (acquired at 50 Hz), as well as the S.E. for those averages.

Single-Channel Patch-Clamp Electrophysiology of Cultured Cells. The A7R3HC10 cells stably expressing human α7 and human RIC-3 were generated from low-passage-number human embryonic kidney (HEK) 293 cells obtained from American Type Culture Collection (Manassas, VA) as previously described (Williams et al., 2012). The A7R3HC10 cells stably expressing human α7 and human RIC-3 were routinely cultured in DMEM supplemented with 10% fetal bovine serum in the presence of 0.45 mg/ml G418 and 0.015 mg/ml hygromycin B at 37°C with 5% CO2, 5% or 10% fetal bovine serum in the presence of 0.45 mg/ml O-2-amino-2,7-dideoxy-β-D-glucopyranosyl-(1→4)-O-[3-deoxy-4-C-methyl-3-(methylamino)-β-L-arabinopyranosyl-(1→6)-2-deoxy-D-streptamine disulfate (G418) and 0.015 mg/ml hygromycin B at 37°C with 5% CO2. For normal passaging, cells were dissociated with 1 mM EDTA in calcium-free and magnesium-free HBSS to avoid nonsedative damage to the α7 nAChRs expressed on the cell surface. Cells with fewer than 50 passages after stable transfection were used for patch-clamp recordings. For cell-attached patch-clamp experiments, cells were plated on poly-L-lysine-coated coverslips, cultured in DMEM supplemented with 5% or 10% fetal bovine serum in the presence of 0.45 mg/ml G418 and 0.015 mg/ml hygromycin B at 37°C with 5% CO2, and used 1–10 days after plating.

Cell-Attached Patch-Clamp Electrophysiology. Single-channel currents were recorded in the cell-attached configuration at room temperature using an MultiClamp 700B amplifier (Molecular Devices) as described previously (Horenstein et al., 2016). A solution containing (millimolar) 165 NaCl, 5 KCl, 2 CaCl$_2$, 10 g-HEPES, and 5 HEPES, and 1 μM atropine, with pH adjusted to 7.3, with NaOH was used for the bath (intracellular solution and had a resistance of 1.5 MΩ). The membrane was giga-ohm sealed using a P-97 micropipette puller to a tip diameter of 1 to 2 μm, fire-polished, and coated with SigmaCote (Sigma-Aldrich). Patch pipettes were then back-filled with the appropriate drug dissolved in the above-mentioned internal solution and had a resistance of 1.5–10 MΩ. Recordings were low-pass-filtered to 6 kHz with the built-in amplifier filter (four-pole Bessel) and digitized at 50 kHz with a DigiData 1440A (Molecular Devices) with Clampfit 10.3.0.2 (Molecular Devices). Additional analysis was conducted with QuB 1.5.0.39 (University at Buffalo, Buffalo, NY). Multiple recordings for each experimental condition were obtained from recording dates. For single-channel experiments, cell-attached recordings were analyzed if on-cell seal access resistances before a recording were greater than 1 GΩ. In cell-attached patches, single-channel events were recorded at 0, 10, and 20 mV pipette potentials from a pipette holding potential of ~60 mV.

Data from at least four individual patches for each condition were pooled to obtain enough number of events for analysis. Portions of data traces containing only single-channel activity were selected for analysis, and any section of recording with two or more channels opened simultaneously was excluded from the analysis. To determine intra-burst closure, subconductance, and open durations, 1631 protracted bursts (584 in GAT107 and 1047 in B-973B) were selected at random for careful idealization. Burst analysis was conducted to define groups of one or more apparent channel openings and closures that arose from an individual channel. Single bursts were defined by closing time greater than t$_{crit}$, a critical parameter that defines the interval that separates closures within a burst (interburst closures) from closures between different bursts (interburst closures). In our analysis, we set the t$_{crit}$ to 50 milliseconds, 5-fold longer than the longest time constant in the intra-burst closed time distribution so that less than 1% of these interburst closures would have been misidentified. All traces were filtered at 6 kHz, corrected for baseline drift, and analyzed with Clampfit 10.3.0.2 (Molecular Devices). Data traces were idealized within QuB using the segmental 6-means method (Qi, 2004) and were idealized at a bandwidth of 6 kHz. After the automated idealization, the fit was manually inspected event by event, and corrections were made as necessary to the idealization. A temporal resolution limit of 1.3× filter rise time was set at 40 microseconds and imposed in all analyses (sampling rate, 50 kHz; interval, 20 microseconds). The probability of channel opening was determined according to the equation reported by Williams et al. (2011b). The mean current amplitudes measured at potentials of 0, 10, and 20 mV relative to the cells’ intrinsic resting potentials were used to generate I–V plots and estimate single-channel conductances. I–V curves were plotted by Kalidagraph 4.1.1, and data were represented as the mean ± S.E.M. of five cells.

Amplitude dwell-time distribution histograms were generated for each event (i.e., closings, openings, and subconductances) and fitted with an appropriate number of Gaussian components. To exclude brief events that could not be properly resolved by our analysis owing to the limit of detection of the filter applied, we discarded dwell-time values shorter than 0.3 millisecond. We calculated the burst-length distributions and the average burst length and identified three major classes of channel openings, namely, full openings and two levels of subconductances. To discriminate between different classes of subconductances, we applied the following criteria and grouped similar subconductances accordingly: for each potential (0, 10, or 20 mV), relative full-opening average amplitude was set to 100% and named “large open.” The average of the smallest subconductances detected was 25% of the full-channel-opening amplitude, and the corresponding openings were identified as small open. A second group of subconductances was defined by opening amplitudes 50% of the large open or intermediate amplitude and named “intermediate open.” Three distinct groups of openings were therefore detected and analyzed separately. Statistical analysis was done using the unpaired t test with equal variances. All differences were considered significant at P < 0.05.

Homology Modeling of the α7 ECD. The α7 nAChR homology model was generated using the Prime module of the Schrödinger 2014-2 Suite (Jacobson et al., 2002, 2004). The epibatidine-bound α7-ACHBP chimera protein structure (PDB ID: 3SQ6) (Li et al., 2011) was used as the homology model template, and the ECD of the mature human α7 subunit sequence was used as the model template (Uniprot P36544). Nonconserved residues were optimized based on energy minimization, and loops smaller than five residues were refined by Prime. Subunit symmetry was not enforced while generating the model. The mature protein α7 numbering was used to number the residues, and the five subunits were named A, B, C, D, and E in a clockwise fashion when viewing the receptor from above on the extracellular side.

Generation of Ligand Parameters. All quantum mechanics calculations for epibatidine, GAT107, TQS, and B-973B were run using Gaussian 09 (Frisch et al., 2016). The ligand molecules we investigated were drawn using Molview or Molden 5.9 (Schaftenaar and Noordik, 2000). Stereochmistry was considered for GAT107, but it was omitted for TQS and B-973B. These structures were then subjected to a two-step geometry optimization procedure, an HF/6-31G* optimization, followed by a B3LYP/6-31G** optimization. Electrostatic potentials of the optimized structures were calculated.
by single-point calculations at the HF/6-31G* level. The atomic charges were generated via restricted electrostatic potential method using the antechamber module of AmberTools16 (Jakalian et al., 2002). Ligand parameters were generated using the leap module with GAFF (Wang et al., 2004) as the force field. Charges and atom types assigned by leap were manually inspected for any discrepancies. Atom names in PDB files were adapted to the nomenclature used in .mol2 file of the corresponding ligand. GAT107, TQS, and B-973B molecules were neutral, and epibatidine had an overall charge of +1.

**Molecular Dynamics of the α7 ECD Model.** The leap routine was used for protein parameterization. Protein residues were parametrized using ff14SB (Maier et al., 2015), and epibatidine was parametrized using GAFF force field for bonded, nonbonded, and charge terms. Hydrogens were added to protein residues under neutral conditions along with missing terminal atoms. Disulfide bond-forming Cys residues were defined by changing the names of residue CYX into CYX in the PDB file. Each subunit had two disulfide bonds between residues C128-C142 and C190-C191 based on numbering of the mature α7 receptor. Ten Na+ ions were added to neutralize the epibatidine-bound protein system. The system was solvated in an octahedral box extending 8 Å from protein residues in all directions containing ~12,000 TIP3P water molecules. Hydrogen mass repartitioning was applied to nonwater molecules of the solvated parameter file.

Two-step minimization was applied where a restrained minimization was followed by an unrestrained minimization. Restrainted minimization (min1) put 10 kcal/mol Å2 force on nonwater atom types CA, C, O, and N; 5000 steps of steepest descent minimization were followed by 45,000 steps of Adopted Basis Newton-Raphson (ABNR) minimization or until a local minimum was reached. The resulting restart file was submitted to the unrestrained minimization step (min2), where 5000 steps of steepest descent minimization was followed by 45,000 steps of ABNR minimization or until a local minimum was reached without any restraints. The restart file from the second minimization step was used for the heating step, in which the system was gradually heated from 0 to 300 K over 3 nanoseconds with a step size of 4 femtoseconds with a restraint of 5 kcal/mol Å2 force on nonwater atom types CA, C, O, and N. Langevin dynamics with a collision frequency of 5 picoseconds−1 and without pressure control was used. The SHAKE algorithm was used to constrain bonds involving hydrogens. The heated structure was equilibrated through two 20-nanosecond CTPE calculations at 1 atm pressure and 300 K temperature first with a restraint of 2 kcal/mol Å2 and then 0.5 kcal/mol Å2. A CTPE MD simulation at 1 atm pressure and 300 K temperature was run for 200 nanoseconds, and the final pose of this structure was minimized for 5000 steps of steepest descent minimization, followed by 15,000 steps of ABNR minimization or until a local minimum was reached without any restraints. Only the TMD portion of this structure was used for the docking calculations.

**Docking.** Schrödinger 2014-2 was used for all docking calculations. The final minimized structure of the ECD or the TMD model was used as the docking model for the docking calculations with GAT107 and B-973B. For the ECD docking calculations, the GlideXP module of the Schrödinger Suite (Friesner et al., 2006) was used such that the ligand conformers were allowed, but the protein structure was rigid. Docking was done for all five interfaces of the ECD model (IAB, IBC, ICD, IBD, and IEC). The top 10 structures were kept for each ligand at each interface based on docking scores where the top pose had the lowest Glide score. For the TMD model docking, an initial Glide docking calculation was run, but it failed to yield an overlapping binding site for both GAT107 and B-973 or produced no poses at all, depending on the interface. As a result, an induced-fit docking (IFD) approach was used to allow optimization of the binding environment within the TM region. The IFD method involved a flexible ligand approach and allowed the protein refinement of the side chain residues within 5 Å of the ligand. An implicit membrane was placed manually to cover the transmembrane helix residues, based on the prediction of the membrane residues by the PPM server. Up to 20 structures were recorded for each ligand and the corresponding protein geometry. The IFD method was used for interfaces that produced some glide poses during the first trial. The poses resulting from IFD calculations were similar for the remaining interfaces, so we used the poses at IAB for our analyses.

**Results**

**Macroscopic Responses of Human α7 nAChR to B-973B.** As previously reported for the racemic compound B-973, the active enantiomer B-973B (Garai et al., 2018) is a very effective modulator of α7 ion channel activation. DAA was observed at concentrations of 1 μM and greater, with persistent primed potentiation of subsequent ACh responses (Fig. 1A). It is also a potent and efficacious PAM of ACh-evoked responses (Fig. 1B). Note that once coapplied at high concentrations with ACh, currents persisted beyond the washout of the drug from the bath (Fig. 1B, lower trace). Whereas DAA (Fig. 2A) and primed potentiation (PP, Fig. 2B) were observed only at concentrations greater than 1 μM, significant (P < 0.001) PAM activity was seen with coapplication of 300 nM B-973B compared with ACh alone (Fig. 2C).
The concentration-response data for allosteric activation and primed potentiation were not suitable to be fit to the Hill equation since the responses continued to increase over the range tested (1–100 μM), and higher concentrations produced responses that were too large to be voltage-clamped. The PAM activity, however, could be fit to the Hill equation, provided the net charge was calculated with our standard protocol (Papke and Porter Papke, 2002) for just the 120-second period immediately after the start of drug delivery because, as noted already, at high concentrations, responses did not return to baseline and tended to become unstable. The \( I_{\text{max}} \) values were 51 ± 9 and 476 ± 40 (relative to 60 μM ACh controls) for peak current and net charge, respectively. The EC\(_{50}\) values were 2.2 ± 1.8 and 3.6 ± 1.1 μM for peak current and net charge, respectively.

Shown in Fig. 3 is a comparison of the allosteric activation produced by 10 μM B-973B and GAT107, along with the structures of the two active isomers. Although comparable in net charge, the GAT107 responses were more reversible than the B-973B responses, and comparison of the peak currents associated with the application of the ago-PAMs alone compared with ACh controls from the same cells indicated that peak currents during the allosteric activation phase were greater (\( P < 0.05 \)) for GAT107 compared with B-973B. The kinetics of B-973B DAA appear slower than those of GAT107. Specifically, GAT107 activation reverses immediately upon washout, whereas B-973B activation still increases after washout begins. This may be due to slower unbinding in the ECD, different coupling to the PAM site, or both these factors.
Additionally, the two ago-PAMs appear to differ in the rates and manner they equilibrate activation and the PAM-sensitive ($D_s$) and PAM-insensitive ($D_i$) desensitized states (Williams et al., 2011b). A prolonged application of 10 µM GAT107 produced a large transient phase of activation before an equilibration toward a steady-state level of activation, as seen with ACh co-applied with PNU-120596 (Williams et al., 2011b). In contrast, activation by 10 µM B-973B was more uniform over the same period (Supplemental Fig. 1).

One interesting aspect of the allosteric activation of α7 by GAT107 is that it can overcome the limitations to orthosteric activation imposed by various point mutations, including mutation of the key aromatic residue Y93 and elimination of the vicinal disulfide between cysteines 190 and 191 at the tip of the C-loop (Papke et al., 2014b; Horenstein et al., 2016). B-973B also effectively activated these mutants (Fig. 4A).

Consistent with their allosteric activity on wild-type receptors (Fig. 3), GAT107 produced larger transient peak currents than B-973B, but the net-charge responses were not different.

The effects of the M254L mutation, previously shown to limit the allosteric activity of PNU-120596 (Young et al., 2008), 4BP-TQS (Gill et al., 2011), and GAT107 (Papke et al., 2014b), were evaluated with B-973B and compared with similar experiments with GAT107 (Fig. 4B). It is interesting to note that whereas this mutation was more than 90% effective at reducing the allosteric and potentiating effects of GAT107, it was essentially 100% effective on eliminating allosteric activation and potentiating effects of B-973B.

As shown in Figs. 1 and 3, cells did not recover to baseline after coapplication of ACh with B-973B at concentrations of ≥10 µM. We have previously shown that mecamylamine is an effective inhibitor of PNU-120596- (Williams et al., 2011b) and GAT107-potentiated currents (Papke et al., 2018) with an IC$_{50}$ for the inhibition of PNU-120596-potentiated currents, similarly to inhibition of responses evoked by ACh alone (Peng et al., 2013). Curiously, we observed that mecamylamine was relatively ineffective at bringing the protracted currents activated by B-973B and ACh back to baseline (Fig. 5).

The kinetics of B-973B DAA appear slower than those of GAT107. That is, GAT107 activation reversed immediately upon washout, whereas B-973B activation still increased after washout begins, possibly due to slower unbinding in the ECD, different coupling to the PAM site, or both these factors. Additionally, the two ago-PAMs appear to differ in the rates and manner that they equilibrate activation and the PAM-sensitive ($D_s$) and PAM-insensitive ($D_i$) desensitized states (Williams et al., 2011b). A prolonged application of 10 µM GAT107 produced a large transient phase of activation during the coapplication of ACh and B-973B at the concentrations indicated. Data represent average (±S.E.M.) responses, normalized to the average of two initial ACh controls of at least five cells under each concentration. Plotted are both the peak currents (left axis) and net charge (right axis) of the responses. The peak current data were not appropriate to fit to the Hill equation. Peak currents, in general, have limited value as measurements of α7 nAChR function since they are confounded by the kinetics of solution exchange and the dynamics of receptor desensitization (Papke and Porter Papke, 2002). The net-charge data were fit to the Hill equation using the Levenberg-Marquardt algorithm with a Hill coefficient of 0.9 ± 0.2, $I_{\text{Max}} = 476 ± 40$, $n = 0.89 ± 0.18$, and EC$_{50} = 3.66 ± 1.15$ µM.
Macroscopic and Microscopic α7 nAChR Allosteric Activation

before an equilibration toward a steady-state level of activation, as seen with ACh coapplied with PNU-120596 (Williams et al., 2011b). In contrast, activation by 10 μM B-973B was more uniform over the same period (Supplemental Fig. 1).

Microscopic (Single-Channel Currents)

General Observations. In control experiments with untransfected HEK cells, neither GAT107 (n = 3, 15-minute recordings) nor B-973B (n = 3, 15-minute recordings) stimulated single-channel currents, whereas allosterically activated channel openings were observed in cell-attached patches on A7R3HC10 cells (Williams et al., 2012), and HEK cells stably transfected with α7 and RIC3. Openings occurred primarily in protracted bursts (Patzynska et al., 2012; Horenstein et al., 2016) (Fig. 6A), like those produced by PNU-120596 when coapplied with an orthosteric agonist (Williams et al., 2012; Andersen et al., 2016). The α7 ago-PAM activity of GAT107 and B-973B resulted primarily in prolonged channel openings consisting mainly of isolated protracted bursts of several hundred milliseconds of duration, consistent with large sporadic increases in single-channel P_open, so that each burst is suitable for analysis as the behavior of one channel. We define burst as a group of openings and closures arising from an individual channel; single bursts were defined by closing time greater than the t_crit. The t_crit is a critical parameter to define the interval separating closures within a burst (intraburst closures) from closures between different bursts (interburst closures). In our analysis, we set the t_crit to 50 milliseconds so that every closure greater than 50 milliseconds defined a single and isolated burst in our analysis (see Materials and Methods). The t_crit of 50 milliseconds was not applied to subconductances within big channel openings.

In general, bursts recorded in the presence of 10 μM B-973B were longer than those activated by 10 μM GAT107, averaging 2.14 ± 0.08 seconds (n = 1047) compared with 1.64 ± 0.11 seconds (n = 584). For both ago-PAMs, the burst-duration distributions (Fig. 6B) were best fit with three components with similar time constants (Table 1); however, the relative area of the longest component was greater for B-973B than for GAT107. Bursts were further characterized by the intraburst components (Fig. 7; Table 2). On average, approximately 130 intraburst state conversions occurred with B-973B and 84 with GAT107. The major differences were apparent in α7-state stabilization and intraburst opening-time durations. For example, 80% of the large openings in B-973B were fit to a time constant of 28.2 ± 0.08 milliseconds and 16% to a longer component of 51.3 ± 0.4 milliseconds, whereas for GAT107, 23% were fit to 11.7 ± 3.2 milliseconds and 30% to the longest time constant of 18.9 ± 1.7 milliseconds. Additionally, 50% of all intraburst state conversions for B-973B were to the large open state and only 32% for GAT107 to the large open state.

B-973B also induced and stabilized prolonged subconductance α7 states, both within bursts of large opening and occasionally as isolated bursts without large openings. Although still able to induce subconductances, GAT107 subconductances were much less frequent and prolonged. Indeed, GAT107 subconductances appeared as brief, isolated intraburst openings and only very briefly and rarely as isolated bursts, consistent with our previous observations (Horenstein et al., 2016). Therefore, resolution and analysis of GAT107 subconductances were more challenging compared with B-973B.

Within most recordings, B-973B presented all three different openings (large open, intermediate and small subconductance openings). To discriminate between the three different openings, we subdivided them according to the amplitude (picoampere) of the currents. The intermediate and small subconductance openings were analyzed separately and as to whether they were isolated from large openings. Current-voltage analysis of the three current levels indicated conductances of 100.9 ± 8.0, 59.5 ± 5.8, and 23.5 ± 1.8 pS for the large, intermediate, and small channel openings, respectively, with no significant differences observed between the currents stimulated by GAT107 and B-973B.

Starting from the total number of patches, for each drug, we analyzed the type of bursts present and calculated the relative percentages (Table 1). For GAT107, approximately 18% of patches analyzed contained exclusively subconductances, and 78% of the patches with large channel openings also displayed...
detectable intraburst subconductances (0.25 and 0.5 openings). In the case of B-973B, 30% of patches analyzed contained exclusively subconductances, and 100% of the patches with large openings also displayed intraburst subconductances (0.25 and 0.5 openings). Although the data suggest that for both drugs subconductances were frequent, there were notable differences. For example, only for B-973B did patches with large openings always also display subconductances.

The apparent intraburst closures were mostly comparable in duration for the two ago-PAMs (\(\sim 0.5\) millisecond) with an additional component around 2 milliseconds (Table 2), even though their frequencies were quite different. Note that for this analysis, closed-time values shorter than 0.3 millisecond were discarded to exclude brief events that could not be resolved properly. Both GAT107 and B-973B occasionally (5\%--10\% of the times) induced intraburst closures longer than 10 milliseconds, suggesting an infrequent intraburst closed state of greater stability. With our burst criteria selection of 50 milliseconds (Materials and Methods), it is likely that some of what we identified as separate bursts actually were part of the same burst; most of the interburst closed times were much longer than 50 milliseconds, but we did not investigate this in detail.

For the dwell-time analyses, we separately analyzed large openings, intermediate openings, and small openings. Additionally, we evaluated whether the intermediate openings and subconductances states occurred in isolation or within bursts containing large openings.

As noted already, on average, the long bursts of openings evoked and potentiated by 10 \(\mu\)M GAT107 and 10 \(\mu\)M B-973B were quite different (Fig. 7; Table 2). For GAT107, two different intraburst open times were observed that averaged 0.64 \(\pm\) 0.23 millisecond and 18.9 \(\pm\) 1.67 milliseconds. In contrast, the intraburst large conductance openings with B-973B were well fit by a single time constant with an average opening time duration of 28.2 \(\pm\) 0.08 milliseconds. These data highlight significant differences in ago-PAM behavior on \(\alpha 7\) nACHR, indicating that GAT107 evoked more brief and fast openings compared with B-973B, which usually induced longer large conductance openings.

We also detected significant differences in the duration and frequency of the subconductance openings evoked by the two tested compounds. Average isolated subconductance openings were fit to several time constants for GAT107 and B-973B, with larger percentages being of longer duration for B-973B (Table 2). These differences may parallel the previously
Mecamylamine Effects on B-973B- and GAT107-Evoked α7 nAChR Currents. In the presence of the neuronal nAChR antagonist mecamylamine, GAT107 and B-973B α7 receptor activation was affected in very different ways (Fig. 8). GAT107-stimulated currents antagonized by mecamylamine showed both large conductances and subconductances in 29% of the patches and had profoundly changed opening durations, resulting in very brief channel openings with long intraburst closing times, as evidenced by the representative traces displayed in Fig. 8A. These effects are similar to those reported by Peng et al. (2013) for mecamylamine on PNU-120596-potentiated currents, where the drug both decreased the duration of large openings and caused the appearance of a subconductance. In contrast, B-973B coapplication with mecamylamine resulted in complete abolition of large conductances of the receptor and the appearance of a novel subconductance (Fig. 8B) in bursts of increased duration (Fig. 9) compared with those recorded with B-973B alone (Fig. 6).

The short durations of the open events observed in the presence of mecamylamine made the estimation of single-channel amplitudes difficult, especially in the case of the shortened GAT107 currents. The subconductance preserved in the presence of B-973B and mecamylamine was estimated to be 60 ± 25 pS, consistent with the intermediate state observed in the absence of mecamylamine. The conductance levels in the presence of mecamylamine and GAT107 corresponded most closely to the large conductance and small subconductance observed in the absence of the antagonist.

The kinetics of the currents stimulated by the ago-PAMs GAT107 or B-973B in co-application with 30 μM mecamylamine showed numerous differences, especially compared with currents observed when no antagonist was present. As noted, in the absence of mecamylamine, burst duration distributions were best described by three exponentials with an overall average length of 1.64 seconds for a GAT107 burst. Relatively few bursts were observed in the presence of GAT107 and mecamylamine (n = 53 bursts, 14 patches), and the distribution was fit with two exponentials. The overall average burst duration was reduced to 0.42 second ± 79 milliseconds. In remarkable contrast, the B-973B bursts, which had an overall average of 2.14 seconds in the absence of mecamylamine, were increased in frequency by mecamylamine (n = 151 bursts, three patches) and increased in average duration to 3.08 second (Fig. 8C). They were uniform in amplitude (Fig. 8A), and the distribution was fit by a single exponential (τ = 2.8 second, Fig. 9). Analysis of intraburst activity also showed reduced richness of channel behavior in the presence of mecamylamine for both ago-PAMs and a special tendency to constrain the channel’s behavior to long sequences of small openings in the case of B-973B-evoked activity. These dissimilarities indicate a qualitatively different mechanism for channel block, dependent not on the antagonist but rather on the specific channel activator, suggesting that the two allosteric activators provide distinctly different environments for drug binding and for the coupling of that binding to further conformational changes. The protracted bursts of small conductance openings in the presence of B-973B and mecamylamine are consistent with the sequential open channel block first described by Neher and Steinbach (1978) of 2-[(2,6-dimethylphenyl)amino]-N,N,N-trimethyl-2-oxethaniminium chloride (QX-222) and the muscle nAChR channel and are also consistent with the relative resistance of the B-973B macroscopic currents to mecamylamine (Fig. 5). The sequential open-channel block model predicts that, with shorter open times per burst and longer bursts, the total open time within a burst should be the same with and without the blocker since it represents the average time required for the blocker to dissociate. We calculated the average open and closed times across all bursts for the four conditions and calculated the Poopen within the bursts as the average open time divided by the sum of the average open and closed time (Table 3). Multiplying that value by the average burst duration gives the average open time within a burst. Averaging across all bursts, in the absence of mecamylamine, open time in B-973B bursts was 1854.3 milliseconds; in the presence of mecamylamine, the average open time per burst was
unchanged, 1,849.4 milliseconds. Using this to estimate an off-rate of 0.54 second\(^{-1}\) for mecamylamine from B-973B-activated channels and the difference in open times to estimate an on-rate of 2,300,000 M\(^{-1}\)s\(^{-1}\), we can further estimate the \(K_d\) of mecamylamine as 230 nM for B-973B-activated channels.

In contrast, the effects of mecamylamine in the presence of GAT107 seem more consistent with a parallel block of both open and closed channels (Papke and Oswald, 1989). Averaging across all bursts, in the absence of mecamylamine, open time in GAT107 bursts was 1340.7 milliseconds, and in the presence of mecamylamine, average open time per GAT107 burst was reduced to 277.3 milliseconds.

**In Silico Analysis of Ago-PAM Binding and Activity.** As discussed previously, the allosteric activation of \(\alpha_7\) receptors by ago-PAMs has been proposed to arise from binding these agents to both of two different classes of sites, a PAM site in the TMD and a DAA site in the ECD that is, at least in part, distinct from the binding site for ACh or other orthosteric agonists (Horenstein et al., 2016). Given the large differences in the structure of GAT107 and B-973B (Fig. 3), it is reasonable that they might show very different activities at one or both classes of sites. Although there are no perfect models of \(\alpha_7\) structure, significant advancements have been made in recent years toward developing homology models thanks to the resolution of protein structures with higher homology to the \(\alpha_7\) receptor (see Materials and Methods) that give some possible insights into the basis for the difference in GAT107 and B-973B activity we observed.

To explore ligand binding poses at the orthosteric and the DAA sites of the ECD of the receptor, we ran 20 nanoseconds molecular dynamics (MD) simulations with a wild-type \(\alpha_7\) ECD homology model to refine the amino acid sidechain geometries. The last snapshot of this MD trajectory was minimized and used as the docking model.

We have previously identified a putative DAA binding site in a deep pocket in the vestibule of the receptor (Horenstein et al., 2016). GAT107 and B-973B showed poses at this DAA site, as well as some poses that overlapped the orthosteric binding site. The minimum pharmacophore of an \(\alpha_7\) agonist is

<table>
<thead>
<tr>
<th>Drug</th>
<th>(A_1)%</th>
<th>(t_1) ms</th>
<th>(A_2)%</th>
<th>(t_2) ms</th>
<th>(A_3)%</th>
<th>(t_3) ms</th>
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<td>GAT107</td>
<td>19</td>
<td>74</td>
<td>60</td>
<td>1058</td>
<td>21</td>
<td>3560</td>
</tr>
<tr>
<td>B-973B</td>
<td>8</td>
<td>86</td>
<td>57</td>
<td>1224</td>
<td>35</td>
<td>3330</td>
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</tbody>
</table>

**Fig. 6.** (A) Representative \(\alpha_7\) single-channel traces of cell-attached patch recordings in 10 \(\mu\)M GAT107 or 10 \(\mu\)M B-973B, shown at different time scales as indicated. Currents were sampled at 50 kHz and low-pass filtered at 6 kHz. In the lower right trace, the various conductance levels are indicated: closed channels (C), small subconductance (S), intermediate subconductance (I), and full (large) open (F). (B) Normalized burst duration distributions for 584 bursts in GAT104 and 1047 in B-973B. Bursts were defined, and sequences of single-channel events were separated from other channel activity by a closed time of \(>50\) milliseconds. The distributions were best fit by three exponential components. See Table 1 for the fit values.
a charged nitrogen (Horenstein et al., 2008), which forms π-cation interactions with the aromatic residues of the orthosteric site (Zhong et al., 1998) and an electrostatic interaction with the backbone oxygen of α7W149. Other agonists, such as ACh and epibatidine, have their hydrophobic groups buried inside a hydrophobic pocket among residues W149, L109, and L119 and may form additional hydrogen bonds through their heteroatoms; however, because neither GAT107 nor B-973B has a permanently charged nitrogen group, they cannot form the landmark electrostatic interactions with the aromatic residues within the orthosteric site. As a result, the orthosteric poses of these compounds are different from those expected for orthosteric full and partial agonists and so would not be consistent with conventional mechanism for channel activation.

Interestingly, we found that the docking of the ago-PAMs was different at each of the subunit interfaces in our model (Supplemental Fig. 1; Supplemental Table 1). Similar non-equivalence of ACh binding sites has been reported previously in other in silico studies (Henchman et al., 2003; Law et al., 2005) and is consistent with studies of large partial agonists crystallized in the AChBP (Hibbs et al., 2009) that showed ligands bound in different orientations at various interfaces. Preferred docking poses for GAT107 and B-973B at one sample interface (I_{BC}) are shown in Fig. 10A. At this interface, the GAT107 poses were closest to the orthosteric site, whereas
B-973B showed a pose in contact with both the orthosteric and putative DAA sites.

The tendency of α7 receptors to convert to nonconducting (desensitized) states at high levels of agonist occupancy (Papke et al., 2000; Uteshev et al., 2002; Williams et al., 2011) may have to do with a bias toward the induction of specific conformational states associated with binding to a particular nonequivalent interface. Although this is an interesting hypothesis, it is beyond the scope of either existing experimental data or in silico modeling to test. Since we cannot directly interpret the significance of the difference in binding among the subunit interfaces, these observations are described in the Supplemental Data.

To delineate the poses important to the ago-PAM activity, we docked TQS, a type II PAM that does not exhibit DAA, into the same binding sites as the ago-PAMs (Fig. 10B). Poses shared by TQS and GAT107 or B-973B would seem unlikely to be responsible for ago-PAM activity. The TQS poses at $I_{AB}$, $I_{BC}$, and $I_{DE}$ were similar to those of GAT107. At $I_{CD}$, although TQS and GAT107 both occupied the DAA site, the binding configurations were different. The sulfonamide group of GAT107 was buried inside the hydrophobic pocket of the DAA site, whereas for TQS, this was the naphthalene group. Furthermore, the GAT107 sulfonamide nitrogen formed a hydrogen bond with the backbone of the L92 residue at the receptor subunit-subunit interface, and a similar interaction was observed for B-973B (data not shown), suggesting a possible significance of this interaction for allosteric activation. This residue is, of course, adjacent to Y93, an essential residue for ordinary orthosteric activation, but not allosteric, as indicated by the ago-PAM activity on the Y93C mutant (Fig. 4).

We also used a homology model of the α7 TMD for our docking studies to analyze the differences at the putative PAM binding site (see Materials and Methods). Other studies (Gill et al., 2011; Newcombe et al., 2018), using Torpedo nAChR structures (Unwin, 2005; Unwin and Fujiyoshi, 2012) as the starting point, have suggested binding sites in the lower regions of the TMD within or between TM helices. For our work, we used the TMDs of the recently resolved α4β2 structure (Hibbs et al., 2009) as a starting point. We applied an induced-fit approach to allow for interactions between the ligand molecules and TMD residues and generated a maximum of 10 structures for each ligand. Our data suggest that a binding site that puts the ligands in direct contact with the region below the M2-M3 linker is more likely, which seems reasonable, considering the importance of the M2-M3 linker and Cys-loop interactions for gating (Purohit and Auerbach, 2007; Lee et al., 2009). The TMD helices of the subunit rotated and tilted during the MD simulations, consistent with the proposed channel opening motion (Law et al., 2005). Both ago-PAMs docked to two different regions close to each other. One was a region situated below the M2-M3 linker of the TMD defined by the residues F253, M254, V257, P269, I271, F275, L215, and C219 of the positive-face subunit and the residues L209, Y210, L213, and N214 of the negative face subunit (Fig. 11A), and the other was between the neighboring M2 helices of the receptor, where ligand binding could block further rotation of the subunit A (Fig. 11B). The best GAT107 poses were under the M2–M3 linker, but some lower-ranking poses appeared between the M2 helices as well. The B-973B poses were more dispersed overall, but two of the best three poses were between the M2 helices, whereas only one was under the M2–M3 linker. Although most of the GAT107 molecule was buried inside the TMD helices, the larger size of B-973B caused a portion of the molecule to stick out and interact with other TMD residues of the neighboring subunits and helices such as Y211, N214, L215, P218, and A258 (Fig. 11B). These additional contacts of B-973B may be the reason behind the differences in the ago-PAM profiles of

### TABLE 2

<table>
<thead>
<tr>
<th>Intraburst Statistics</th>
<th>GAT107</th>
<th>B-973B</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Type of Intraburst Event</strong></td>
<td><strong>A%</strong></td>
<td><strong>[t] ms</strong></td>
</tr>
<tr>
<td>Closures</td>
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<td>0.44 ± 0.33</td>
</tr>
<tr>
<td>Isolated subconductances</td>
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<tr>
<td></td>
<td>4 ± 0.9</td>
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<tr>
<td>Large openings</td>
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<tr>
<td></td>
<td>35 ± 24</td>
<td>4.83 ± 0.68</td>
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<tr>
<td></td>
<td>18 ± 34</td>
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<tr>
<td></td>
<td>39 ± 11</td>
<td>0.64 ± 0.23</td>
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<tr>
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<td>8 ± 10</td>
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<td>Intermediate openings</td>
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<td>13 ± 11</td>
<td>7.95 ± 1.45</td>
</tr>
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<td>Small openings</td>
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<td>6 ± 4</td>
<td>2.50 ± 10.5</td>
</tr>
<tr>
<td></td>
<td>17 ± 16</td>
<td>12.5 ± 1.51</td>
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Fig. 8. (A) Representative α7 single-channel traces of cell-attached patches for GAT107 and B-973B coapplied with 30 μM mecamylamine shown at different time scales as indicated. Currents were sampled at 50 kHz and low-pass filtered at 6 kHz. (B) Representative normalized all points (amplitude) histograms for 1-second intervals of channel-bursting activity in the presence of the ago-PAMs alone (red) or the ago-PAMs plus 30 μM mecamylamine (blue). (C) Since burst-duration distributions are exponential functions and unsuitable for parametric comparisons, burst durations were converted to their corresponding log10 values, generating Gaussian-like distributions. Shown are the average log10 burst durations for the two ago-PAMs ± S.E.M. of those values; t test comparisons indicated significant differences between the values for GAT107 alone compared with B-973B alone, that GAT107 bursts were shorter in mecamylamine, and that B-973B bursts were longer in the presence of mecamylamine. A Bonferroni correction was applied so that P values for a given level of significance are reduce by a factor of 3. The P value
B-973B and GAT107, particularly regarding the availability of binding sites within the channel for mecamylamine and possibly other noncompetitive antagonists.

The M254L mutation can affect a variety of type I and type II PAM molecules to different extents (Bertrand et al., 2008; Young et al., 2008; Gill et al., 2011). We made the in silico

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**Fig. 9.** Kinetic analysis of single-channel α7 activity in bursts evoked by 10 μM GAT107 or 10 μM B-973B. Fit log_{10} probability histograms displaying burst durations and apparent open and close durations. Note that in the presence of B-973B and mecamylamine, only a single type of opening was resolved.

For comparison of GAT107 log_{10} burst durations to the B973B burst durations was $3.6 \times 10^{-5}$. The $P$ value for the comparison of GAT107 log_{10} burst durations to the GAT107 burst durations in the presence of mecamylamine was $3.8 \times 10^{-6}$. The $P$ value for the comparison of B-973B log_{10} burst durations to the B-973B burst durations in the presence of mecamylamine was $1.7 \times 10^{-5}$, so that all were significant at an adjusted value $<0.00033$. We also calculated the false discovery rate correction for multiple comparisons, and those corrected values were $1.07 \times 10^{-5}$, $5.7 \times 10^{-6}$, and $1.7 \times 10^{-5}$, respectively, for the values reported above. ***$P < 0.0001$.**
a7M254L mutation using UCSF chimera (Pettersen et al., 2004) to see how it affected the interactions among the TMD residues (Fig. 11C). All possible rotamers of L254 had clashes with the nearby F275, which not only restricted the PAM binding pocket but could also cause global disruption of the TMD geometry owing to the compact network of hydrophobic residues at the TMD. GAT107 directly enters the binding pocket without passing through other TMD helices. When B-973B binds to this site, however, it does so like a hook where the difluorinated phenyl ring sits in the hydrophobic pocket, whereas the other part of the molecule passes over M254 and interacts with other TMD residues. Because GAT107 is smaller than B-973B and has fewer interactions in the direction of M254, it is less likely to suffer from a clash with L254, which would explain why the M254L mutation affected B-973B more significantly than GAT107 (Fig. 11D).

### Discussion

Although structurally very different, both GAT107 and B-973B are ago-PAMs, capable of producing allosteric activation when the orthosteric site is disabled by mutations. Both will synergize with orthosteric agonists and are more potent as PAMs than as allosteric agonists, evoking greater channel activation when coapplied with ACh than when applied alone. Both the potentiating effects and allosteric agonism rely on the TMD PAM binding site. Both apparently destabilize desensitized states induced by agonist binding; however, GAT107-potentiating effects appear to be limited by the induction of D1 (PAM-insensitive desensitization) at high concentrations, accounting for an inverted-U concentration-response relationship (Papke et al., 2014b), whereas B-973B-potentiating effects are limited at high concentrations, and persistent activation can be evoked, even with limited exposure to ACh (Figs. 2 and 3).

On the microscopic level, both ago-PAMs caused individual a7 receptors to show huge increases in single-channel P_{open} by manifesting long-lasting bursts, as previously reported for currents promoted by PNU-120596 (Williams et al., 2011b) and other type II PAMs (Andersen et al., 2016). Although ideally it would be best to compare macroscopic and microscopic response in the same system, comparisons of the potentiation of single-channel currents in HEK cells to whole-cell responses in oocytes suggests that, at the concentration studied, more channels were quiescent than activated at any time, permitting the resolution of protracted single-channel bursts with little or no overlapping activation of other channels. Whereas on the macroscopic level the allosteric activation produced by 10 μM B973B had a net charge only twice that stimulated by 60 μM ACh (Fig. 2), the total open time within B-973B single-channel bursts (Table 3) was 30,000-fold greater than the average 60 μM ACh-activated currents (Williams et al., 2011b).

Under our experimental conditions, when only the ago-PAM is in the pipette, we can be confident that we are observing only DAA. Likewise, when we contrast that activity to what is observed when mecamylamine is added, we know that we observed the effects of mecamylamine only on allosterically activated channels.

Both ago-PAMs promote a large conductance state that is similar in amplitude, but far more stable, than that promoted by orthosteric agonists (Williams et al., 2011b). The PAM-associated open state(s) are coupled to unstable “flip states” (Lape et al., 2008), promoting bursting. Burst-duration distributions suggest at least three different bursting states. At the concentration tested, bursts of longer duration were promoted more effectively by B-973B than by GAT107. In addition, both drugs also promoted subconductances connected to the same or to different flip states.

One interesting difference was that B-973B-evoked currents, unlike those evoked by GAT107, were resistant to mecamylamine, a neuronal nAChR-selective channel blocker (Webster et al., 1999). In mecamylamine, GAT107 bursts were fewer, shorter, and contained proportionately more small-conductance openings, suggesting that large or full openings are more easily blocked and that channels do not have to reopen to release the blocker. In contrast, B-973B bursts in the presence of mecamylamine were longer and consisted exclusively of intermediate and small conductance openings. Channels appeared to be prevented from entering the large open state, which may have to do with the size and positioning of B-973B in the TMD site. It is known that sensitivity to mecamylamine and other noncompetitive antagonists is regulated by sequence at sites in the pore-forming M2 domain (see Table 4). Specifically, the residues at the 10' and 6' positions (Miller, 1989) in the muscle β1 subunit protect muscle receptors (Webster et al., 1999). Perhaps the binding of B-973B in the PAM site partially occludes or otherwise distorts the channel, preventing complete block by mecamylamine. Partial penetration of the channel by mecamylamine may then exclusively permit the subconductance state that is then blocked in a sequential open-state-dependent manner, leading to protracted bursts, as described for QX-222 inhibition of muscle-type receptors (Neher and Steinbach, 1978). The same M2 residues regulate sensitivity to other antagonists such as tetracaine (Papke et al., 2001), and we determined that B-973B currents are also relatively insensitive to 100 μM tetracaine (Supplemental Fig. 3).

Although PAMs have been characterized that differ greatly in structure (Williams et al., 2011c), this is the first case for such diversity in ago-PAMs, which are likely to bind at both the PAM site in the TMD and the allosteric site(s) in the ECD. We used in silico methods to test the hypothesis that they shared the same ECD site; however, the docking studies suggested multiple binding modes or sites in the ECD, including the putative DAA site and atypical binding to the

<table>
<thead>
<tr>
<th>Condition</th>
<th>Average Open, ms</th>
<th>Average Closed, ms</th>
<th>P_{open}</th>
<th>Open Time (ms) per Burst</th>
</tr>
</thead>
<tbody>
<tr>
<td>10 μM GAT107</td>
<td>8.13</td>
<td>1.79</td>
<td>0.82</td>
<td>1340.7</td>
</tr>
<tr>
<td>10 μM B973B</td>
<td>19.87</td>
<td>2.88</td>
<td>0.87</td>
<td>1854.3</td>
</tr>
<tr>
<td>GAT107 + Mec.</td>
<td>1.70</td>
<td>0.89</td>
<td>0.66</td>
<td>277.3</td>
</tr>
<tr>
<td>B973B + Mec.</td>
<td>2.53</td>
<td>1.68</td>
<td>0.60</td>
<td>1849.4</td>
</tr>
</tbody>
</table>
canonical orthosteric site (Supplemental Data). We know that high levels of binding to the orthosteric sites promote transitions to desensitized states, some of which are perturbed by PAMs, but no data have related this to different binding modes or activity at different interfaces; however, the differences detected in the in silico study may suggest that binding sites at the multiple subunit interfaces may play different roles in coupling ligand binding to specific conformational transitions of the receptor.

Both ago-PAMs produced complex patterns of subconductances and different burst states, unusual for nAChR in the absence of allosteric modulation but more common for glutamate-gated ion channels (Traynelis et al., 2010). The different open states we observed may be due to different numbers of sites occupied at either of the allosteric sites, or by different poses at nonequivalent sites at the subunit interfaces or elsewhere in the ECD. These factors would be expected to change dynamically throughout the course of a recording since the ligands were used at a subsaturating concentration. It should be noted that the extreme gain-of-function $\alpha 7 L9 T$ mutation promotes currents very much like a type II PAM and couples the binding of competitive antagonists to channel activation (Bertrand et al., 1997). The presence of this mutation in $\alpha 7$ diminishes the relative effects of PAMs (Williams et al., 2011b; Newcombe et al., 2018), suggesting that the mutation itself partially destabilizes desensitization in a manner similar to a PAM. The distinction between PAMs and ago-PAMs—that PAMs require agonist coapplication whereas ago-PAMs do not, can also be reduced or eliminated by mutations in the W55 residue of $\alpha 7$ (Horenstein et al., 2016; Newcombe et al., 2018), a mutation that also perturbs desensitization produced by conventional agonists (Gay et al., 2008; Williams et al., 2009). It may also be the case that nonconventional binding of the ago-PAMs to the orthosteric site could also couple to activation with the binding of the agents to the TMD PAM site. We observe similar behavior with weak partial agonists in the presence of the PAM PNU-120596 (Papke et al., 2014a).

The environment of the channel itself differs in the open states promoted by the two drugs so that current-dependent signaling may also differ, especially regarding the limitations imposed by the D state. We know from earlier work (Peng et al., 2013) that ions and channel blockers affect the PNU-potentiated channel differently from how they affect the “normal” open state. We have shown that mecamylamine affects GAT107- and B-973B-potentiated channels and the B-973B conductance states differently.

Although superficially similar in some respects, the two agents have different balances in their activities at the two allosteric sites and subsequent coupling to the orthosteric site. These differences may have further implications about how the intracellular domains couple to the interactome and subsequent signal transduction. The subtleties of the pharmacologic manipulation of $\alpha 7$ conformational states may have significance beyond the integrated effects of ion channel currents since $\alpha 7$ nAChRs have been shown to play a key role in the regulation of inflammatory disease and neuropathic pain through the cholinergic anti-inflammatory pathway (CAP) (Rosas-Ballina and Tracey, 2009). Although this study has focused on the modulation of ion channel activity produced by these ago-PAMs, it remains the case that most channels remain closed for most of the time, and modulation of CAP appears to rely on such nonconducting states (Papke et al., 2015; Horenstein and Papke, 2017). GAT107 has already been shown to be effective in models of neuropathic and inflammatory pain (Bagdas et al., 2016), and preliminary data (Garai et al., 2018) indicate that B-973B has similar effects.

**Fig. 10.** (A) The top two docking poses of GAT107 (right) and the top four docking poses of B-973 (left) from the outside (top) and the vestibular side (bottom) at $\alpha 7$ ECD I$_D$. The GAT107 poses shown are at the orthosteric site, whereas the B-973B poses are in contact with both the orthosteric site and the DAA site. (B) The best docking pose of GAT107 (left) and TQS (right) at the I$_D$ DAA site viewed from the vestibular side. The sulfonamide group of each molecule is circled in white. (C) A magnified view of (B).
The limitations on α7 channel activation, associated with its homomeric composition, have made it a fascinating and challenging target for drug development (Papke, 2014). Highly selective agonists and partial agonists (Horenstein et al., 2008) were developed in the hopes that they might produce treatments for Alzheimer disease. With the discovery of α7-selective PAMs (Williams et al., 2011c) and more recently, ago-PAMs, the limitations of channel activation have been removed, and the discovery of CAP demonstrated significance for α7 targeting drugs beyond channel-dependent processes (de Jonge and Ulloa, 2007). The TQS scaffold on which GAT107 is based has already been shown to be a useful starting point for the development of related agents with diverse properties (Gill-Thind et al., 2015; Horenstein et al., 2016; Newcombe et al., 2018). Characterization of B-973B as a structurally unrelated ago-PAM with unique properties provides an exciting new starting point for further development of potential therapeutic agents.

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Authorship Contributions

Participated in research design: Quadri, Horenstein, Papke.

Conducted experiments: Quadri, Stokes, Gulsevin.

Contributed new reagents or analytic tools: Garai, Thakur, Gulsevin.

Performed data analysis: Quadri, Gulsevin, Papke.

Wrote or contributed to the writing of the manuscript: Quadri, Thakur, Stokes, Gulsevin, Horenstein, Papke.

References


Barkas T, and Ballivet M (1990) A neuronal nicotinic acetylcholine receptor subunit (alpha 7) is developmentally regulated and forms a homo-oligomeric channel blocked by alpha-BTX. J Neurosci 10: 1306–1313.


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References


TABLE 4

Sequence in the second transmembrane domain of select nAChR subunits with 6’ and 10’ residues in bold

<table>
<thead>
<tr>
<th>Domain</th>
<th>Sequence</th>
</tr>
</thead>
<tbody>
<tr>
<td>α1</td>
<td>MTLISLVESSVTVLTVLLVibilePST</td>
</tr>
<tr>
<td>γ</td>
<td>CTVAIOGLLAQVTVLPLVAXKVPST</td>
</tr>
<tr>
<td>Delta</td>
<td>TSVVIASVLLASQVLLISSKRLPST</td>
</tr>
<tr>
<td>α3</td>
<td>VTCI5LEVLLSVTVTTLITIPST</td>
</tr>
<tr>
<td>β1</td>
<td>MGLSI1PAAIITTVLTLALLADKVPST</td>
</tr>
<tr>
<td>β4</td>
<td>MTLCI5AVLALTFVLLISSKIVPST</td>
</tr>
<tr>
<td>β2</td>
<td>MTLGC6AVLALTFVLLISSKIVPST</td>
</tr>
<tr>
<td>α7</td>
<td>6’10’ ISLGI1VTVLTVFNVLVAEINPST</td>
</tr>
</tbody>
</table>

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