Identification of Global and Ligand-Specific Calcium Sensing Receptor Activation Mechanisms[§]

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ABSTRACT

Calcium sensing receptor (CaSR) positive allosteric modulators (PAMs) are therapeutically important. However, few are approved for clinical use, in part due to complexities in assessing allostery at a receptor where the endogenous agonist (extracellular calcium) is present in all biologic fluids. Such complexity impedes efforts to quantify and optimize allosteric drug parameters (affinity, cooperativity, and efficacy) that dictate PAM structure-activity relationships (SARs). Furthermore, an underappreciation of the structural mechanisms underlying CaSR activation hinders predictions of how PAM SAR relates to in vitro and in vivo activity. Herein, we combined site-directed mutagenesis and calcium mobilization assays with analytical pharmacology to compare modes of PAM binding, positive modulation, and agonism. We demonstrate that 3-(2-chlorophenyl)-*N*-((1*R*)-1-(3-methoxyphenyl)ethyl)-1-propanamine (NPS R568) binds to a 7 transmembrane domain (7TM) cavity common to class C G

protein-coupled receptors and used by (αR)-(-)- α -methyl-*N*-[3-[3-(trifluoromethylphenyl)propyl]-1-naphthalenemethanamine (cinacalcet) and 1-benzothiazol-2-yl-1-(2,4-dimethylphenyl)-ethanol (AC265347); however, there are subtle distinctions in the contribution of select residues to the binding and transmission of cooperativity by PAMs. Furthermore, we reveal some common activation mechanisms used by different CaSR activators, but also demonstrate some differential contributions of residues within the 7TM bundle and extracellular loops to the efficacy of the PAM-agonist, AC265347, versus cooperativity. Finally, we show that PAMS potentiate the affinity of divalent cations. Our results support the existence of both global and ligand-specific CaSR activation mechanisms and reveal that allosteric agonism is mediated in part via distinct mechanisms to positive modulation.

Introduction

The human calcium sensing receptor (CaSR) is a class C G protein-coupled receptor (GPCR) that negatively regulates parathyroid hormone secretion in response to serum extracellular calcium (Ca_o^{2+}) concentrations. As such, the small-molecule positive allosteric modulator (PAM), (αR)-(-)- α -methyl-*N*-[3-[3-(trifluoromethylphenyl)propyl]-1-naphthalenemethanamine (cinacalcet), is approved for the treatment of secondary hyperparathyroidism, and some instances for the treatment of primary hyperparathyroidism (Nemeth and Goodman, 2016). Cinacalcet

has also been used off-label to correct disorders caused by naturally occurring loss-of-function CaSR mutations (Mayr et al., 2016). Although various CaSR small-molecule PAM chemotypes have been discovered, cinacalcet remains the only one on the market. Failure to translate newer CaSR PAMs to the clinic may reflect complexities associated with detecting and quantifying in vitro allosteric modulator effects at class C GPCRs (Leach and Gregory, 2017), thus hampering structure-activity relationship (SAR) optimization efforts and predictions of in vivo drug efficacy, and impeding dissection of allosteric mechanisms of action. For instance, despite cinacalcet being the first clinically approved GPCR allosteric modulator to reach the market, surprisingly little is known about how cinacalcet and other PAMs mediate positive allosteric modulation and/or agonism at the CaSR. Understanding GPCR activation mechanisms can help guide SAR optimization of ligands with desired activity.

Structural studies of class A and B GPCRs indicate that during receptor activation a rotational and vertical movement of transmembrane (TM) 6 and TM7 relative to TM3 occurs,

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ABBREVIATIONS: AC265347, 1-benzothiazol-2-yl-1-(2,4-dimethylphenyl)-ethanol; ANOVA, analysis of variance; Ca_i^{2+} , intracellular calcium; Ca_o^{2+} , extracellular calcium; CaSR, calcium sensing receptor; cinacalcet, (αR)-(-)- α -methyl-*N*-[3-[3-(trifluoromethylphenyl)propyl]-1-naphthalenemethanamine; ECL, extracellular loop; GPCR, G protein-coupled receptor; NPS R568, 3-(2-chlorophenyl)-*N*-((1*R*)-1-(3-methoxyphenyl)ethyl)-1-propanamine; PAM, positive allosteric modulator; SAR, structure-activity relationship; TM, transmembrane; 7TM, 7 transmembrane domain; VFT, Venus flytrap; WT, wild type.

Fig. 1. Studying positive allosteric modulation at the CaSR. (A) Structures of the arylalkylamine PAM, NPS R568, and the PAM-agonist, AC265347, investigated in this study. (B) According to the operational model of allosterism, the CaSR PAM effects on orthosteric agonist signaling are governed by PAM affinity (K_B), binding and/or efficacy cooperativity between the PAM and the orthosteric agonist (α and β , respectively), and the efficacy of the PAM (τ_B); $\alpha\beta$ governs the magnitude of agonist potentiation, reflected by an increase in agonist $E_{\max}$ and/or potency, and enhanced vehicle-mediated signaling plus a reduction in the slope of the transducer function, n , if ambient orthosteric agonists are present in the buffer. If the PAM is also an agonist, τ_B will dictate PAM-mediated receptor signaling in the presence of vehicle. (C) Snake diagram of the primary CaSR 7TM domain and ECL sequence, indicating residues that were mutated in the current study. Mutations at residues previously shown to be important for the binding of the PAMs, cinacalcet, and/or AC265347, their cooperativity with Ca_o^{2+} , or both (Leach et al., 2016), are shown in blue, red, and orange, respectively. All engineered mutations were substituted with alanine, with the exception of A615^{1.42}, A772^{5.39}, A840^{7.35}, and A844^{7.3}, which were substituted with valine, and E837^{7.32}, which was mutated to aspartic acid and isoleucine. The residue highlighted in green indicates where a naturally occurring mutation alters the activity of some PAMs (Cook et al., 2015). Arrows point to the $\times .50$ conserved class C amino acid residues based on a modified Ballesteros-Weinstein numbering system (Doré et al., 2014).

resulting in conformational changes at the intracellular face of the receptor that promote transducer coupling (reviewed in de Graaf et al., 2017; Manglik and Kruse, 2017). For class C GPCRs, most endogenous agonists predominantly bind in the large extracellular Venus flytrap (VFT) domain, stabilizing activating conformational rearrangements that transmit through the 7 transmembrane domain (7TM) spanning regions and/or extracellular loops (ECLs) to the intracellular side of the receptor. Many synthetic molecules that target class C GPCRs bind to allosteric sites in the 7TM domain and/or ECL regions. Furthermore, for the CaSR, the 7TM and/or ECL domains also contain additional binding sites for endogenous agonists including polyamines and Ca_o^{2+} (Ray and Northup, 2002; Leach et al., 2016). However, the 7TM molecular mechanisms that underlie class C GPCR allosteric (and indeed orthosteric) activation are largely unknown. This is due to a complete lack of active-state class C GPCR 7TM structures, but also, in part, due to the difficulties in interpreting mutation-based structure-function studies to delineate residue contributions to the three key molecular properties that govern allosteric activity; ligand affinity, cooperativity, and efficacy.

In light of the difficulties associated with studying GPCR allosteric modulator effects, we developed pharmacological and analytical methods to robustly quantify allosteric modulation of GPCRs (Leach et al., 2007), and recently employed these methods to probe class C GPCR allostery (Gregory et al., 2012, 2013, 2014; Leach et al., 2016). We demonstrated

unprecedented insights into class C GPCR drug actions by combining pharmacological, analytical, and detailed structure-function analyses. For instance, we identified a CaSR 7TM domain cavity that accommodates small-molecule CaSR PAMs and negative allosteric modulators, and showed that structurally diverse modulators bind to distinct regions in this cavity; the arylalkylamine PAM, cinacalcet, binds toward the top of the 7TM domain cavity, whereas the structurally distinct CaSR PAM-agonist, 1-benzothiazol-2-yl-1-(2,4-dimethylphenyl)-ethanol (AC265347), binds deeper in the pocket (Leach et al., 2016). Furthermore, we mapped residues that contribute to PAM potentiation of the CaSR's endogenous agonist, Ca_o^{2+} , revealing both shared and distinct activation mechanisms used by cinacalcet and AC265347 (Leach et al., 2016).

Given the importance of understanding the structural basis underlying ligand binding and activation mechanisms at class C GPCRs, the current study sought to build on our prior analysis of CaSR allostery by examining the structural basis of positive allosteric modulation mediated by the small-molecule PAM, 3-(2-chlorophenyl)-*N*-((1*R*)-1-(3-methoxyphenyl)ethyl)-1-propanamine (NPS R568). The binding of NPS R568 and other arylalkylamine PAMs is predicted to overlap with the cinacalcet binding site. Given that arylalkylamine PAMs display comparable in vitro pharmacology, we hypothesized that NPS R568 would demonstrate similar structure-function relationships to cinacalcet. We also hypothesized that the PAM-agonist, AC265347, mediates agonism via engagement of residues that transmit positive

modulation. Thus, we used mutagenesis to compare mechanisms of positive allosteric modulation and direct allosteric agonism mediated by NPS R568 and AC265347 (Fig. 1A). We identify both global and ligand-specific activation mechanisms at the CaSR, which could inform future drug discovery efforts seeking to rationally optimize the activity of CaSR activators and/or potentiators.

Materials and Methods

Materials. Dulbecco's modified Eagle's medium, blasticidin S HCl, and fetal bovine serum were obtained from Invitrogen (Carlsbad, CA), while hygromycin B was obtained from Roche (Mannheim, Germany). The Flp-In TREx human embryonic kidney 293 cells and Fluo-4 AM acetoxymethyl ester were purchased from Invitrogen. Quikchange mutagenesis kits were purchased from Agilent Technologies (Santa Clara, CA). NPS R568 was prepared as described previously (Davey et al., 2012). AC265347 and all other chemicals were from Sigma Aldrich (St Louis, MO).

Generation and Maintenance of Cell Lines Expressing Wild-Type (WT) and Mutant CaSRs. The generation of DNA and cells stably expressing c-myc-tagged WT and mutant human CaSRs in pcDNA5/ftt/TO have been described previously (Davey et al., 2012; Leach et al., 2016). Cells were maintained in Dulbecco's modified Eagle's medium containing 5%–10% fetal bovine serum, 200 $\mu\text{g}/\text{ml}$ hygromycin B, and 5 $\mu\text{g}/\text{ml}$ blasticidin S HCl.

Intracellular Calcium (Ca^{2+}) Mobilization Assays. Cells were seeded in clear 96-well plates coated with poly-D-lysine (50 $\mu\text{g}/\text{ml}^{-1}$) at 80,000 cells/well and incubated overnight in the presence of 100 ng/ml^{-1} tetracycline. The following day, cells were washed with assay buffer (150 mM NaCl, 2.6 mM KCl, 1.18 mM MgCl_2 , 10 mM D-Glucose, 10 mM HEPES, 0.1 mM CaCl_2 , 0.5% bovine serum albumin, and 4 mM probenecid at pH 7.4) and loaded with Fluo-4 AM acetoxymethyl ester (1 μM in assay buffer) for 1 hour at 37°C. Cells were washed with assay buffer prior to the addition of fresh assay buffer. For interaction studies between Ca_0^{2+} and PAMs, modulators were coadded with Ca_0^{2+} . For assays in the absence of ambient divalent cations, cells were treated as described previously, except that the final assay buffer and the buffer used for washing and loading the cells did not contain CaCl_2 or MgCl_2 .

For all studies, each well was treated with a single agonist and/or modulator concentration. The release of Ca_i^{2+} was measured at 37°C using Flexstation 1 or 3 (Molecular Devices, Sunnyvale, CA). Fluorescence was detected for 60 seconds at 485 nm excitation and 525 nm emission, and the peak Ca_i^{2+} mobilization response (approximately 12 seconds after agonist addition) was used for the subsequent determination of the agonist response. Relative peak fluorescence units were normalized to the fluorescence stimulated by 1 μM ionomycin to account for differences in cell number and loading efficiency.

Data Analysis. All nonlinear regression analyses were performed using GraphPad Prism 7 (GraphPad Software, San Diego, CA). Parametric measures of potency, affinity, and cooperativity were estimated as logarithms (Christopoulos, 1998). Interaction experiments between Ca_0^{2+} and allosteric modulators were fitted to an operational model of allosterism (Leach et al., 2007; Aurelio et al., 2009; Leach et al., 2010), which was modified to accommodate the presence of ambient agonist (Davey et al., 2012):

$$E = \frac{(E_{\max} - \text{basal}) [(A) + [A_e] (K_B + \alpha\beta[B]) + \tau_B [B] [\text{EC}_{50}]]^n}{[\text{EC}_{50}] (K_B + [B]) + [(A) + [A_e] (K_B + \alpha\beta[B]) + \tau_B [B] [\text{EC}_{50}]]^n} \quad (1)$$

where $[A]$ denotes the concentration of exogenously added orthosteric agonist to stimulate the response; $[A_e]$ is the ambient orthosteric agonist concentration in the buffer; $[B]$ is the molar allosteric ligand concentration; K_B is the functional affinity of the allosteric ligand; τ_B

denotes allosteric ligand intrinsic efficacy; α and β are the allosteric effects on orthosteric ligand binding affinity and efficacy, respectively (estimated as a composite $\alpha\beta$ value); $E_{\max}$ is the maximal possible system response; and n is the slope factor of the transducer function.

Concentration-response curves for Ca_0^{2+} , NPS R568, and AC265347 in the absence of ambient divalent cations were fitted to the following form of an operational model of agonism to obtain estimates of PAM affinity and efficacy:

$$E = \left(\frac{(E_{\max} - \text{basal}) \tau_B [B]}{(K_B + [B]) + \tau_B [B]} \right)^n \quad (2)$$

where all parameters are as described for eq. 1.

Statistics. An F test ($P < 0.001$) was used to determine whether eq. 1 best fit interaction data when NPS R568 and AC265347 were assumed to have no agonist efficacy at the WT and mutant CaSRs (i.e., whether the best-fit value for τ_B was equal to or differed from zero). Statistical differences between the Ca_0^{2+} pEC₅₀, or PAM pK_B and log $\alpha\beta$ at the WT versus the mutant receptors were determined by one-way analysis of variance (ANOVA) with Dunnett's multiple comparisons post-test, where significance was defined as $P < 0.05$. Statistical differences between PAM pK_B at the WT and mutant receptors in the absence versus the presence of ambient divalent cations was determined by two-way ANOVA with Sidak's comparisons post-test, where significance was considered as $P < 0.05$.

Molecular Modeling and Docking Studies. A three-dimensional model of the CaSR 7TM domain was constructed by homology with mGlu₅ (PDB: 5CGD) (Christopher et al., 2015) using the ICM software package (Molsoft, San Diego, CA) (Abagyan and Totrov, 1994) as previously described (Leach et al., 2016). The 7TM helices in the model were additionally refined to accommodate irregularities such as nonconserved proline residues (e.g., Pro682^{3,34}) and a one-residue insertion in the extracellular part of TM5 (Ile777^{5,44}). Compound docking relied on the pharmacophore properties of the TM cavity as well as the results of earlier and present mutagenesis studies indicating that mutation of Glu837^{7,32} abrogates activity of all arylalkylamines. A salt bridge is predicted to occur between the protonated arylalkylamine secondary amine and the Glu837^{7,32} side chain (Petrel et al., 2003, 2004; Miedlich et al., 2004; Bu et al., 2008; Leach et al., 2016); therefore, the formation of this salt bridge was enforced in our docking studies by imposing a harmonic distance restraint between the corresponding atoms. The arylalkylamine docking proceeded by extensive conformational sampling of the ligands and the residue side chains lining the pocket in the internal coordinates in the presence of this distance restraint. All complexes were further refined by local minimization in the presence of distance restraints maintaining receptor secondary and tertiary structures, and inspected manually.

Results

Rationale for the Study. We have previously evaluated the effects of CaSR 7TM and ECL mutations on allosteric modulator activity using an interaction paradigm (Davey et al., 2012; Cook et al., 2015; Leach et al., 2016). Here, the effect of a CaSR PAM can manifest as: 1) modulation of the exogenously added agonist to stimulate the response, reflected by a change in the added agonist potency and/or $E_{\max}$; 2) modulation of ambient agonist present in the buffer, reflected by an increase in the baseline response and/or a shallowing of the agonist concentration-response curve; and 3) direct PAM-mediated agonism, also reflected by an increase in the baseline response (Davey et al., 2012; Cook et al., 2015; Leach et al., 2016) (Fig. 1B). In this work, we have extended our analysis to account for ambient Ca_0^{2+} ions present in the assay buffer. Including the ambient Ca_0^{2+} concentration in the

Fig. 2. 7TM mutations alter the activity of NPS R568 at the CaSR. Concentration-response curves to Ca_o^{2+} in the absence and presence of NPS R568 were determined in Ca_o^{2+} mobilization assays to identify mutations that altered NPS R568 activity (A–E). Representative mutants are shown that reduced NPS R568 affinity (F684^{3.36}A), increased NPS R568 affinity (A844^{7.39}V), reduced NPS R568 cooperativity (V817^{6.49}A), or increased NPS R568 affinity while decreasing cooperativity (F821^{6.53}A). Data are mean + S.E.M. pooled from at least four separate experiments performed in duplicate. Curves are the best fit of the data to eq. 1. Vehicle (buffer) is plotted as the smallest $[\text{Ca}_o^{2+}]$ in accordance with the logarithmic scale.

operational model of allosterism allows delineation of PAM-mediated increases in the vehicle (buffer) response due to agonism versus potentiation of ambient Ca_o^{2+} (eq. 1). Accordingly, we are now able to quantify and delineate the effect of CaSR mutations on PAM affinity (pK_B), allosteric cooperativity with the orthosteric agonist ($\alpha\beta$), and allosteric agonism (τ_B).

Using this analytical paradigm, we examined two tool compounds; NPS R568 and AC265347. NPS R568 (Fig. 1A) was among the first identified CaSR PAMs (Nemeth et al., 1998) and a precursor to cinacalcet, which is an approved clinical treatment of human hyperparathyroidism. Both PAMs display similar in vitro pharmacology (Cook et al., 2015), despite subtle differences in their chemical structure; although NPS R568 and cinacalcet share a three carbon aliphatic chain linked to a secondary amine, they differ in their terminal aromatics: chlorophenyl and methoxyphenyl groups in NPS R568 were replaced with tri-fluoromethylphenyl and naphthyl groups in cinacalcet. With these differences in mind, we aimed to examine the SAR of NPS R568 in comparison with cinacalcet in more detail. Furthermore, whereas cinacalcet and NPS R568 exhibit little agonism, the chemically distinct CaSR PAM, AC265347, is both a PAM and an agonist (a PAM-agonist) (Cook et al., 2015). We were, therefore, interested in elucidating the different activation

mechanisms used by these chemically and pharmacologically distinct CaSR activators. Of further interest, AC265347 exhibits biased CaSR modulation when compared with cinacalcet, presumably via stabilizing different active states (Davey et al., 2012; Leach et al., 2013, 2016; Cook et al., 2015).

Therefore, in this work, we used mutagenesis in combination with an operational model of allosterism that explicitly accommodates ambient agonist levels (eq. 1), to characterize important features of NPS R568 binding and positive cooperativity, and to directly probe AC265347 agonism. 7TM and ECL mutations were selected based on those known to alter the affinity and/or cooperativity of cinacalcet and/or AC265347 (Fig. 1C).

NPS R568 Binds to a Common 7TM Allosteric Binding Site. We first confirmed that NPS R568 bound within the same 7TM cavity used by other arylalkylamine modulators (Leach et al., 2016). We fitted NPS R568 interaction data at the WT and 7TM mutant receptors to an operational model of allosterism with ambient agonist to assess the impact of 7TM amino acid substitutions on NPS R568 affinity (pK_B). WT and representative mutants are shown in Fig. 2 that exemplify the different profiles observed. Similar to our previous observations with cinacalcet (Leach et al., 2016), F668^{2.56}A, F684^{3.36}A, F688^{3.40}A, E837^{7.32}I, and I841^{7.36}A mutations abolished NPS R568 activity or reduced

Fig. 3. 7TM and ECL mutations alter the affinity and cooperativity of NPS R568 at the CaSR. (A) Concentration-response curves to Ca_o^{2+} in the absence and presence of NPS R568 determined in Ca_i^{2+} mobilization assays were fitted to an operational model of allosterism (eq. 1) to determine the change in affinity (ΔpK_B) (A) and cooperativity ($\Delta \log \alpha \beta$) (C) of NPS R568 at the mutant CaSRs in comparison with the WT receptor. White bars represent no significant change in pK_B or $\log \alpha \beta$. Colored bars that sit above and below zero represent a significant increase or decrease in pK_B or $\log \alpha \beta$, respectively. A 1.5-fold or lower reduction in pK_B and/or $\log \alpha \beta$ is represented by yellow bars, a 1.6-fold to 2.5-fold reduction is represented by orange bars, and a greater than 2.5-fold reduction or increase is represented by red and blue bars, respectively. NA, negligible PAM activity. Data are the ΔpK_B or $\Delta \log \alpha \beta$ + S.E.M. calculated from WT and mutant pK_B or $\log \alpha \beta$ values and experiment numbers shown in Table 1; *denotes significant difference in comparison with WT ($P < 0.05$, one-way ANOVA with Dunnett's multiple comparisons post-test). Affinity (B) and cooperativity (D) residues are shown on a CaSR 7TM domain homology model (gray cartoon), along with the predicted NPS R568 pose from docking studies. Residue positions are shown as ball and stick, and colors correspond to the fold change in affinity or cooperativity shown in (A) and (C).

NPS R568 pK_B , whereas F821^{6.53}A, E837^{7.32}D, and A844^{7.39}V increased NPS R568 pK_B (Fig. 2; Fig. 3A; Supplemental Fig. 1; Table 1). Furthermore, although the magnitude of NPS R568 potentiation of Ca_o^{2+} potency at the Y825^{6.57}A mutant was reduced (Supplemental Fig. 1), a significant and maximal ~2-fold increase in Ca_o^{2+} potency was reached in the presence of 0.1 μM NPS R568, suggesting the PAM pK_B was increased. To confirm this finding, we performed interaction studies between AC265347 and NPS R568. We found that 0.1–10 μM NPS R568 caused a dextral translocation of the AC265347 concentration-response curve at the Y825^{6.42}A mutant, with no apparent decrease in AC265347 E_{max} (Supplemental Fig. 2); although limitations in AC265347 solubility prevented determination of complete AC265347 concentration-response curves in the presence of the higher NPS R568 concentrations. Nonetheless, these findings suggest that the interaction between NPS R568 and AC265347 is competitive, or an allosteric interaction governed by extremely high negative cooperativity. In support of a competitive or highly negatively

cooperative interaction, when AC265347 and NPS R568 interactions at the Y825^{6.42}A mutant were fitted to a modified Hill/Gaddum/Schild equation (Motulsky and Christopoulos, 2004; Langmead et al., 2006), the Schild slope was not significantly different from unity. Under these circumstances, the concentration of NPS R568 that shifts the AC265347 EC_{50} 2-fold is equal to the NPS R568 pK_B ; this analysis confirmed that the NPS R568 pK_B was increased by the Y825^{6.57}A mutation (Table 1). A similar increase in cinacalcet pK_B has been observed at this mutant (Leach et al., 2016). Our findings are consistent with NPS R568 binding in the 7TM cavity in a similar pose to cinacalcet, concordant with previous predictions that E837^{7.32} forms a hydrogen bond with the core secondary amine in arylalkylamine modulators (Petrel et al., 2003, 2004; Miedlich et al., 2004; Bu et al., 2008; Leach et al., 2016). With this interaction, the terminal naphthyl (cinacalcet) or methoxyphenyl (NPS R568) moiety extends downward, π -stacking with F684^{3.36} and forming hydrophobic interactions with F668^{2.56}, F688^{3.40}, and I841^{7.36}.

TABLE 1

Functional affinity (pK_B) and cooperativity ($\log\alpha\beta$) parameters for NPS R568 determined in Ca^{2+} mobilization assays using an interaction paradigm

Data are mean $\pm$ S.E.M. and 95% confidence interval from the indicated number of independent experiments (n).

	NPS R568		
	pK_B (95% CI)	$\log\alpha\beta$ (95% CI) ($\alpha\beta$)	n
WT	5.99 $\pm$ 0.05 (5.89–6.09)	0.55 $\pm$ 0.01 (0.52–0.58) (3.5)	17
F612 ^{1.39} A	6.39 $\pm$ 0.15 (6.03–6.75)	0.35 $\pm$ 0.04 ^a (0.28–0.43) (2.2)	4
A615 ^{1.42} V	6.22 $\pm$ 0.13 (5.93–6.49)	0.45 $\pm$ 0.04 (0.38–0.53) (2.8)	6
F668 ^{2.56} A	<5	ND	4
G670 ^{ECL1} A	6.39 $\pm$ 0.08 (6.23–6.55)	0.09 $\pm$ 0.04 ^a (–0.07–0.25) (1.2)	4
F684 ^{3.36} A	5.28 $\pm$ 0.18 ^a (4.79–5.63)	0.35 $\pm$ 0.05 ^a (0.28–0.51) (2.2)	4
F688 ^{3.40} A	5.40 $\pm$ 0.16 ^a (5.00–5.73)	0.52 $\pm$ 0.05 (0.44–0.65) (3.3)	5
E767 ^{ECL2} A	6.36 $\pm$ 0.22 (5.93–6.77)	0.25 $\pm$ 0.03 ^a (0.18–0.33) (1.8)	3
A772 ^{5.39} V	6.33 $\pm$ 0.08 (6.17–6.48)	0.38 $\pm$ 0.02 ^a (0.35–0.42) (2.4)	4
F775 ^{5.42} A	6.11 $\pm$ 0.07 (5.95–6.27)	0.39 $\pm$ 0.02 ^a (0.36–0.42) (2.5)	4
L776 ^{5.43} A	6.40 $\pm$ 0.09 (6.20–6.58)	0.46 $\pm$ 0.03 (0.40–0.51) (2.9)	4
V817 ^{6.49} A	5.89 $\pm$ 0.26 (5.29–6.38)	0.37 $\pm$ 0.05 ^a (0.27–0.50) (2.3)	4
W818 ^{6.50} A	5.88 $\pm$ 0.05 (5.77–5.98)	0.55 $\pm$ 0.01 (0.52–0.58) (3.5)	5
F821 ^{6.53} A	7.14 $\pm$ 0.18 ^a (6.80–7.58)	0.18 $\pm$ 0.02 ^a (0.14–0.22) (1.5)	4
Y825 ^{6.57} A	6.88 $\pm$ 0.11 ^{a,b} (6.65–7.11)	0.07 $\pm$ 0.02 ^{a,c} (0.04–0.10) (1.2)	4
K831 ^{ECL3} A	5.60 $\pm$ 0.15 (5.26–5.91)	0.60 $\pm$ 0.05 (0.49–0.71) (4.0)	4
V833 ^{ECL3} A	6.07 $\pm$ 0.11 (5.77–6.33)	0.35 $\pm$ 0.02 ^a (0.31–0.40) (2.2)	4
E837 ^{7.32} D	7.13 $\pm$ 0.16 ^a (6.85–7.52)	0.26 $\pm$ 0.03 ^a (0.22–0.32) (1.9)	4
E837 ^{7.32} I	<5	ND	7
A840 ^{7.35} V	6.31 $\pm$ 0.13 (6.05–6.56)	0.33 $\pm$ 0.03 ^a (0.28–0.39) (2.1)	4
I841 ^{7.36} A	NA	NA	4
A844 ^{7.39} V	6.69 $\pm$ 0.09 ^a (6.49–6.90)	0.05 $\pm$ 0.09 ^a (–0.17–0.26) (1.1)	5
L848 ^{7.43} A	6.04 $\pm$ 0.13 (5.79–6.28)	0.29 $\pm$ 0.02 ^a (0.25–0.32) (1.9)	4

CI, confidence interval; NA, negligible PAM activity; ND, not determined due to weak cooperativity or low affinity.

^asignificantly different to WT ($P < 0.05$, one-way ANOVA with Dunnett's multiple comparisons post-test).

^b pK_B determined in interaction studies between AC265347 and NPS R568 ($n = 3$).

^c pK_B was fixed to 6.88 determined in interaction studies between AC265347 and NPS R568.

There were, however, subtle differences in the effects of some mutations on NPS R568 pK_B when compared with cinacalcet. For instance, W818^{6.50}A decreased, whereas E767^{ECL2}A, A772^{5.39}V, or V833^{ECL3}A increased, cinacalcet pK_B (Leach et al., 2016); in contrast, these mutations had no significant effect on NPS R568 pK_B (Fig. 3A; Table 1). To better understand the differences in the pK_B effects of these mutations, we constructed a homology model of the CaSR 7TM domain based on the mGlu₇ 7TM structure (Christopher et al., 2015) and interrogated this model bound to NPS R568 in silico. Since the NPS R568 core consists of an arylalkylamine, whose secondary amine is predicted to interact directly with E837^{7.32}, the distance required for hydrogen bonding between this residue and the secondary amine of NPS R568 was maintained during the docking process. Unsurprisingly, this produced docking poses for NPS R568 that were consistent with previously published predictions for cinacalcet and other arylalkylamine-based modulators (Leach et al., 2016) (Fig. 3B).

Close inspection of docking poses for NPS R568 in the CaSR homology model explained the observed differences in mutational effects on cinacalcet versus NPS R568 pK_B . First, our predicted docking poses oriented the methoxyphenyl moiety of NPS R568 toward the hydrophobic-aromatic network that surrounds the naphthyl group of cinacalcet (F684^{3.36}, F775^{5.42}, L776^{5.43}, W818^{6.50}, and I841^{7.36}), but crucially, the smaller and more hydrophilic NPS R568 moiety does not interact with W818^{6.50} (Fig. 3B; Supplemental Fig. 3). This change in binding mechanism likely underpins why W818^{6.50}A does not influence NPS R568 pK_B . Second, the predicted NPS R568 pose places the chloro-phenyl group of NPS R568 in a position identical to the tri-fluoromethyl-phenyl group of

cinacalcet, but the NPS R568 moiety occupies appreciably less chemical space. Indeed, the increase in cinacalcet pK_B observed at the E767^{ECL2}A and V833^{ECL3}A mutants is likely due to an increase in the volume of the top of the binding pocket, which is more favorable for cinacalcet but makes no difference to the binding of the smaller NPS R568 (Supplemental Fig. 3). Collectively, these data suggest that NPS R568 and cinacalcet bind to an overlapping binding site; however, due to key differences in their chemical topology, there are subtle distinctions in the contribution of select residues to the binding of the two PAMs.

Identification of Residues that Mediate Cooperativity between Ca^{2+} and NPS R568. By applying an operational model of allosterism to our data, we also revealed nine residues within the CaSR 7TM bundle that are involved in NPS R568 cooperativity ($\log\alpha\beta$) but not affinity (pK_B), with major contributions from ECL1 and 2 as well as TMs 6 and 7 (WT and representative mutants are shown in Fig. 2, quantification is shown in Fig. 3C). In contrast, five mutations influenced both affinity and cooperativity; F688^{2.56}A decreased pK_B and $\log\alpha\beta$, whereas F821^{6.53}A, E837^{7.32}D, Y825^{6.42}A, or A844^{7.39}V had differential effects on the two parameters (Fig. 3, A and C; Table 1). Thus, residues within the binding pocket differentially contribute to NPS R568 affinity versus cooperativity with Ca^{2+} .

Our previous structure-function analysis at the CaSR identified residues that contributed to the transmission of positive cooperativity between Ca^{2+} and cinacalcet (Leach et al., 2016). Comparisons between cooperativity determinants for cinacalcet and NPS R568 revealed key similarities and differences. Five mutations (E767^{ECL2}A, V817^{6.49}A, Y825^{6.57}A, A844^{7.39}V, and L848^{7.43}A) significantly reduced

TABLE 2

Efficacy (τ_B) of NPS R568 and AC265347 determined in Ca_i^{2+} mobilization assays using an interaction paradigm

Data are mean $\pm$ S.E.M. and 95% confidence intervals from the indicated number of independent experiments (n).

	NPS R568		AC265347	
	Log τ_B (95% CI) (τ_B)	n	Log τ_B (95% CI) (τ_B)	n
WT	ND	17	-0.22 ± 0.09 (-0.45 to -0.08) (0.6)	8
F612 ^{1.39} A	ND	4	0.12 ± 0.09^a (-0.10 to 0.44) (1.3)	4
A615 ^{1.42} V	ND	6	0.04 ± 0.03 (-0.01 to 0.09) (1.1)	4
F668 ^{2.56} A	ND	4	ND	4
G670 ^{ECL1} A	-0.25 ± 0.06 (-0.41 to -0.15) (0.6)	4	-0.15 ± 0.09 (-0.39 to 0.02) (0.7)	4
F684 ^{3.36} A	ND	4	ND	5
F688 ^{3.40} A	ND	5	ND	4
E767 ^{ECL2} A	ND	3	0.05 ± 0.03 (-0.02 to 0.12) (1.1)	4
A772 ^{5.39} V	ND	4	ND	4
F775 ^{5.42} A	ND	4	-0.26 ± 0.07 (-0.48 to 0.14) (0.5)	7
L776 ^{5.43} A	ND	4	0.19 ± 0.12^a (-0.11 to 0.32) (1.5)	6
V817 ^{6.49} A	ND	4	$0.33 \pm 0.04^{a,b}$ (0.26 – 0.45) (2.1)	5
W818 ^{6.50} A	ND	5	ND	4
F821 ^{6.53} A	ND	4	ND	4
Y825 ^{6.57} A	ND	4	ND	4
K831 ^{ECL3} A	ND	4	-0.05 ± 0.03 (-0.10 to 0.00) (0.9)	5
V833 ^{ECL3} A	ND	4	ND	5
E837 ^{7.32} D	ND	4	0.09 ± 0.12 (-0.18 to 0.28) (1.2)	4
E837 ^{7.32} I	NA	7	NA	5
A840 ^{7.35} V	ND	4	ND	7
I841 ^{7.36} A	NA	4	ND	4
A844 ^{7.39} V	-0.22 ± 0.07 (-0.45 to -0.11) (0.6)	5	-0.12 ± 0.03 (-0.14 to -0.09) (0.8)	6
L848 ^{7.43} A	ND	4	-0.17 ± 0.07 (-0.35 to -0.07) (0.7)	4

CI, confidence interval; NA, negligible PAM activity; ND, not determined due to negligible efficacy.

^aSignificantly different from WT ($P < 0.05$, one-way ANOVA with Dunnett's multiple comparisons post-test).

^bTo account for a reduction in cell surface expression caused by V817^{6.49}A compared with the WT CaSR (Leach et al., 2016), log τ_B was normalized to determine what it would be if this mutant was expressed at the same level as the WT receptor (Gregory et al., 2010).

the cooperativity of both NPS R568 and cinacalcet, suggesting contributions to a common network that transmits positive cooperativity by structurally similar PAMs (Fig. 3, C and D). In contrast, six of the mutations that specifically reduce NPS R568 cooperativity with Ca_o^{2+} without impacting affinity (F612^{1.39}A, G670^{ECL1}A, A772^{5.39}V, F775^{5.42}A, V833^{ECL3}A, and A840^{7.35}V) did not influence cinacalcet cooperativity. Furthermore, A615^{1.42}V, W818^{6.50}A, and K831^{ECL3}A, which we previously reported to influence cinacalcet cooperativity, had no significant effect on the cooperativity between NPS R568 and Ca_o^{2+} , demonstrating that the contribution of these residues to the transmission of positive cooperativity is ligand specific. Collectively, our data reveal differences in the network of residues that transmit cooperativity mediated by structurally similar PAMs, indicating that distinct conformational states are favored by cinacalcet and NPS R568.

Identification of Global and Ligand-Specific Activation Mechanisms at the CaSR. Having identified the NPS R568 binding site and differentiated ligand-receptor interactions governing affinity and cooperativity, we next sought to compare activation mechanisms used by different CaSR activators. To do so, we first validated the ability of the operational model of allosterism with ambient agonist (eq. 1) to distinguish between PAMs (e.g., NPS R568) and PAM-agonists (e.g., AC265347). When applied to Ca_o^{2+} versus NPS R568 interaction data, we found that the operational model of allosterism with ambient agonist fit the data best when the PAM was presumed to have no efficacy (i.e., the best fit value for τ_B did not differ from zero, F test; $P < 0.001$) (Fig. 2A; Tables 1 and 2). This agreed with our previous study showing that cinacalcet, NPS R568, and other structurally related

PAMs demonstrate negligible intrinsic agonism (Cook et al., 2015). In contrast, and also consistent with our previous study (Cook et al., 2015), the interaction between Ca_o^{2+} and the PAM-agonist AC265347 were fitted best when the model assumed AC265347 was an agonist (i.e., the best fit value for τ_B differed from zero, F test; $P < 0.001$) (Fig. 4A; Table 2). Importantly, reanalysis of these previously reported data with eq. 1 yielded similar estimates of AC265347 pK_B and $\log\alpha\beta$ (Supplemental Table 1). Taken together, these findings demonstrate that our operational model of allosterism with ambient agonist can differentiate between two PAMs with different agonist activity. Therefore, we next assessed the impact of mutations on allosteric modulator efficacy, τ_B , independently of positive cooperativity.

Alanine substitution of F684^{3.36} or F688^{3.40} was previously found to reduce Ca_o^{2+} potency (Leach et al., 2016), and in the current study also abolished AC265347-mediated Ca_i^{2+} mobilization in the presence of vehicle (while positive cooperativity with Ca_o^{2+} was retained) (Fig. 4B; Supplemental Fig. 1; Tables 1 and 2). Using a tetracycline-inducible cell line, we titrated F684^{3.36} and F688^{3.40} receptor expression levels to establish the Ca_o^{2+} pEC_{50} under conditions where it should more closely reflect its affinity (i.e., in the absence of receptor reserve) (Supplemental Fig. 4). With reduced expression levels, Ca_o^{2+} pEC_{50} values at the F684^{3.36}A (2.90 ± 0.07) or F688^{3.40}A (2.95 ± 0.03) mutants were not significantly different from WT (3.06 ± 0.04), indicating that the reduction in Ca_o^{2+} potency at the mutants versus WT under high expression conditions was due to reduced Ca_o^{2+} efficacy, not affinity. In contrast, F612^{1.39}A, L776^{5.43}A, or V817^{6.49}A increased Ca_o^{2+} potency (Leach et al., 2016) and AC265347 efficacy (Table 2). The changes in Ca_o^{2+}

Fig. 4. 7TM domain mutations alter orthosteric and allosteric agonist activity at the CaSR. Concentration-response curves to Ca_o^{2+} determined in Ca_i^{2+} mobilization assays in the absence and presence of AC265347 (A–D) have been published previously (Leach et al., 2016), and were reanalyzed with an operational model of allosterism that considered ambient divalent cations to determine the effect of mutations on AC265347 efficacy (τ_B). WT and representative mutants are shown that caused a simultaneous reduction in Ca_o^{2+} potency and loss of AC265347 efficacy (F688^{3.40}A), a simultaneous increase in Ca_o^{2+} potency and gain in AC265347 efficacy (L776^{5.43}A), and a selective loss of AC265347 efficacy (A840^{7.35}V). Data are mean $\pm$ S.E.M. pooled from at least four separate experiments performed in duplicate. Curves through the data points are the best fit of the data to eq. 1. Vehicle (buffer) is plotted as the smallest $[\text{Ca}_o^{2+}]$ in accordance with the logarithmic scale.

potency and AC265347 efficacy at the F684^{3.36}A, F688^{3.40}A, F612^{1.39}A, L776^{5.43}A, or V817^{6.49}A mutants under conditions where WT and mutant cell surface expression levels are comparable (Leach et al., 2016) suggests these residues are important for the transmission of agonism imparted by both Ca_o^{2+} and AC265347.

In contrast, W818^{6.50}A, F668^{2.56}A, A772^{5.39}V, F821^{6.53}A, Y825^{6.57}A, V833^{ECL3}A, A840^{7.35}V, or I841^{7.36}A did not reduce Ca_o^{2+} potency (several increased it), or with the exception of F668^{2.56}A did not reduce cell surface expression (Leach et al., 2016), but they all abolished AC265347 efficacy (Fig. 4D; Table 2). Qualitatively, AC265347 enhanced Ca_i^{2+} mobilization in the presence of vehicle alone at certain mutations (see, for example, Fig. 4D); however, our analysis is consistent with these baseline increases being attributable to positive modulation of ambient cations in the buffer rather than intrinsic AC265347 efficacy (Supplemental Table 1). Therefore, these eight residues are critical for mediating agonism of AC265347, but not Ca_o^{2+} .

Intriguingly, analysis of NPS R568 interactions with Ca_o^{2+} revealed two mutations (G670A^{ECL1}A and A844^{7.39}V) where there was a gain in allosteric agonism, as demonstrated by a significant increase in τ_B , but neither mutation enhanced AC265347 efficacy (Table 2). Combined with a comparison of mutations that influence Ca_o^{2+} potency (Supplemental Fig. 5) versus AC265347 efficacy (Fig. 5; Table 2), our findings indicate that some 7TM and ECL residues play a global role in agonism imparted by all classes of agonist (i.e., orthosteric and allosteric), whereas others contribute to ligand-specific receptor activation.

The finding that the G670A and A844V mutations significantly reduced NPS R568 cooperativity yet converted NPS R568 to a PAM-agonist was intriguing. Therefore, we next sought to compare the global and AC265347-specific

activation residues with the AC265347 cooperativity networks previously established (Leach et al., 2016) to determine whether there were also distinctions between cooperativity versus agonism networks used by other PAMs. Two mutations (F612^{1.39}A and L776^{5.43}A) that increased AC265347 efficacy (Fig. 5, A and B) did not change positive cooperativity, whereas V817^{6.49}A increased AC265347 efficacy while decreasing cooperativity. In contrast, the loss in AC265347 efficacy at F668^{2.56}A was opposed by increases in cooperativity. Four mutations that decreased AC265347 cooperativity (A615^{1.42}V, E767^{ECL2}A, K831^{ECL3}A, and A844^{7.39}V) had no effect on AC265347 efficacy. Collectively, these data reveal differential contributions of residues within the 7TM bundle and ECLs to efficacy versus cooperativity.

CaSR PAMs Enhance the Affinity of Orthosteric Agonists. To confirm mutational effects on PAM efficacy predicted from our analysis of interaction data, we assessed the ability of NPS R568 or AC265347 to stimulate Ca_i^{2+} mobilization in the absence of ambient Ca_o^{2+} or Mg_o^{2+} at WT and select mutant receptors. We evaluated G670^{ECL1}A and A844^{7.39}V mutations, where NPS R568 was predicted to gain agonist activity, and A840^{7.35}V as a representative mutation that decreased AC265347 efficacy (Table 2). Furthermore, we evaluated E837^{7.32}D as a control, because this mutation had no effect on the efficacy of NPS R568 or AC265347. Where possible, we fit agonist concentration response data to an operational model of agonism (eq. 2) to determine agonist affinity and efficacy at WT and/or mutant CaSRs (Table 3).

In agreement with our previous evaluation of PAM agonism (Cook et al., 2015), up to 30 μM NPS R568 did not stimulate Ca_i^{2+} mobilization in the absence of ambient cations (Fig. 6A); due to limited solubility, it was not possible to test higher concentrations. Nonetheless, these findings confirm that NPS

Fig. 5. 7TM and ECL mutations alter AC265347 efficacy at the CaSR. (A) Previously published concentration-response curves to Ca_0^{2+} in the absence and presence of AC265347 determined in Ca_i^{2+} mobilization assays (Leach et al., 2016) were reanalyzed with an operational model of allosterism that accounted for ambient divalent cations (eq. 1) to determine the change in AC265347 efficacy ($\Delta\tau_B$) at the mutant CaSRs in comparison with the WT receptor. White bars represent no significant change in τ_B . Colored bars that sit above zero represent an increase in τ_B . A significant 1.6–2.5-fold increase in τ_B is represented by the purple bar and a greater than 2.5-fold increase is represented by blue bars. ND, not determined due to negligible efficacy; NA, no PAM activity. Data are the $\Delta\tau_B$ + S.E. M. calculated from WT and mutant τ_B values and experiment numbers shown in Table 2; * demotes significant difference in comparison with WT ($P < 0.05$, one-way ANOVA with Dunnett's multiple comparisons post-test). Efficacy residues are shown on a CaSR 7TM domain homology model (B), along with the predicted AC265347 pose from docking studies. Blue and purple colors correspond to the fold change in efficacy shown in (A), whereas red corresponds to a complete loss in AC265347 efficacy.

R568 exhibits negligible agonism at the CaSR. In contrast to WT CaSR, NPS R568 stimulated concentration-dependent Ca_i^{2+} mobilization in the absence of ambient cations at G670^{ECL1}A and A844^{7.39}V (Fig. 6, B and C), suggestive of increased NPS R568 efficacy, as predicted by our operational model analysis that incorporated the presence of ambient cations (Table 2). However, the estimated NPS R568 pK_B at the G670^{ECL2}A mutant was significantly lower in the absence of ambient cations (Table 3) than in their presence (Table 1) ($P < 0.05$, two-way ANOVA with Sidak's multiple comparisons). A similar trend toward a lower NPS R568 pK_B in the absence of ambient cations was also observed at the A844^{7.39}V mutant, but this did not reach significance. At E837^{7.32}D and A840^{7.35}V, NPS R568 exhibited negligible agonist activity, similar to WT CaSR and as predicted from our operational model of allosterism analysis.

In contrast to NPS R568, AC265347 stimulated relatively robust Ca_i^{2+} mobilization at the WT CaSR in the absence of

ambient cations (Fig. 6A; Table 3), indicative of intrinsic agonism. AC265347 was also an agonist at the G670^{ECL2}A, E837^{7.32}D, and A844^{7.39}V mutants in the absence of ambient cations, although there was a small but significant decrease in its efficacy at A844^{7.39}V (Fig. 6; Table 3). Consistent with our operational model of allosterism analysis, AC265347 efficacy was reduced at the A840^{7.35}V mutant.

Similar to NPS R568 at WT CaSR and G670^{ECL2}A, A844^{7.39}V, E837^{7.32}D, or A840^{7.35}V mutants, AC265347 bound with lower affinity in the absence of ambient cations than in their presence ($P < 0.05$, two-way ANOVA with Sidak's multiple comparisons). The difference in affinities suggests that CaSR PAMs bind preferentially when an orthosteric agonist simultaneously occupies the receptor. Since cooperative allosteric effects on affinity are reciprocal (and, therefore, affect both agonist and modulator) (May et al., 2007), the elevated PAM affinity in the presence of ambient cations is expected if the PAM also enhances the

TABLE 3
Functional affinity (pK_B) and efficacy ($\log\tau_B$) parameters for NPS R568 and AC265347 determined in Ca^{2+}_i mobilization assays in the absence of ambient divalent cations
Data are mean $\pm$ S.E.M. and 95% confidence intervals from the indicated number of independent experiments (n).

NPS R568			AC265347		
pK_B (95% CI)	$\log\tau_B$ (95% CI) (τ_B)	n	pK_B (95% CI)	$\log\tau_B$ (95% CI) (τ_B)	n
WT	ND	8	5.43 ± 0.10^a (5.23–5.63)	0.05 ± 0.10 (0.01–0.11) (1.1)	8
G670 ^{ECL1A}	5.71 ± 0.18^a (5.28–6.13)	7	6.13 ± 0.08^b (5.96–6.31)	0.07 ± 0.03 (0.04–0.11) (1.2)	7
E837 ^{7.32D}	ND	8	$5.90 \pm 0.14^{a,b}$ (5.60–6.19)	-0.03 ± 0.03 (–0.10 to 0.03) (0.9)	8
A840 ^{7.35V}	ND	7	ND	ND	7
A844 ^{7.39V}	6.48 ± 0.32 (5.78–7.21)	6	6.89 ± 0.20^b (6.51–7.31)	-0.16 ± 0.04^b (–0.25 to –0.09) (0.7)	6

CI, confidence interval; ND, not determined due to negligible efficacy and/or low affinity.
^aSignificantly lower than in the presence of ambient divalent cations ($P < 0.05$, two-way ANOVA with Sidak's comparisons post-test).
^bSignificantly different from WT ($P < 0.05$, one-way ANOVA with Dunnett's multiple comparisons post-test).

affinity of the ambient cations. Thus, NPS R568 and AC265347 likely act, at least in part, by potentiating the affinity of divalent cations. This could occur if the PAMs stabilized the closed form of the VFT via allosteric interactions between the 7TM and VFT domains.

Discussion

The current study has used a structure-function approach combined with analytical pharmacology to probe CaSR 7TM domain and ECL residues that contribute to receptor

activation and positive allosteric modulation. Importantly, by incorporating the presence of ambient agonist into an operational model of allosterism, we were able to delineate the contribution of residues to PAM intrinsic efficacy (agonism) independently of positive cooperativity (potentiation of the orthosteric agonist) and affinity. We found both ligand-specific and global activation networks within the CaSR ECLs and 7TM bundle. The fact that mutations in these regions of the receptor have substantial effects on both orthosteric and allosteric ligands highlights the pivotal role that the 7TM domain and ECLs play in transmitting activating stimuli,

Fig. 6. CaSR PAMs bind with lower affinity in the absence vs. presence of ambient divalent cations, and exhibit differential efficacy at the 7TM domain and ECL mutants. Concentration-response curves to Ca^{2+}_o , NPS R568, and AC265347 were determined in Ca^{2+}_i mobilization assays in the absence of ambient divalent cations to evaluate PAM efficacy (A–E). Representative mutants were chosen based on their ability to provoke NPS R568 efficacy (G670^{ECL2A} and A844^{7.39V}), reduce AC265347 efficacy (A840^{7.35V}), or have no effect on PAM efficacy (E837^{7.32D}). Data are mean $\pm$ S.E.M. pooled from at least four separate experiments performed in duplicate. Curves are the best fit of the data to eq. 2. Vehicle (buffer) is plotted as the smallest $[Ca^{2+}_o]$ in accordance with the logarithmic scale.

irrespective of the location of the ligand binding site. Importantly, our ability to delineate mutation effects on cooperativity and efficacy reveals that positive cooperativity appears to be mediated in part by residues that are distinct to those that transmit allosteric agonism. For instance, A615^{1,42}, E767^{ECL2}, and K831^{ECL3} all contributed to the transmission of AC265347 cooperativity yet played no role in AC265347 efficacy. Conversely, F612^{1,39} and L776^{5,43} constrained AC265347 efficacy but not positive cooperativity. In other instances, there were opposing contributions of residues to cooperativity versus efficacy, as was observed for G670^{ECL2} and A844^{7,39}; mutation of these residues decreased NPS R568 cooperativity yet converted NPS R568 to an agonist. Similarly, V817^{6,49}A increased AC265347 efficacy while decreasing cooperativity, where the opposite was true for F668^{2,56}A.

Using mutagenesis to dissect the contribution of 7TM and ECL residues to NPS R568 affinity, we confirm that NPS R568 binds to an extended cavity in the CaSR's 7TM domain, much like cinacalcet, with key interactions between F668^{2,56}, F684^{3,36}, F688^{3,40}, E837^{7,32}, and I841^{7,36}. We did, however, identify some subtle differences in the binding of NPS R568 and cinacalcet, likely attributed to their different terminal aromatic groups, permitting cinacalcet to occupy more space at the top and bottom of the binding pocket. Despite this, and the identification of a number of residues that differentially contribute to NPS R568 and cinacalcet cooperativity, these two PAMs from the arylalkylamine family are pharmacologically very similar (Cook et al., 2015). The differences in NPS R568 and cinacalcet SAR appear to be overcome by shared contributions of E767^{ECL2}, V817^{6,49}, Y825^{6,57}, A844^{7,39}, and L848^{7,43} to their transmission of positive cooperativity. Importantly, E767^{ECL2}, V817^{6,49}, and A844^{7,39} are also critical for positive cooperativity mediated by the structurally distinct PAM-agonist AC265347, suggesting that these residues may contribute to a global allosteric activation network. Furthermore, mutation of these residues also increases Ca_o²⁺ potency, suggesting a reduction in the energy barrier to transition to an active CaSR 7TM domain that mirrors the effects of PAMs. In fact, greater than 70% of the 7TM/ECL mutations studied herein increase Ca_o²⁺ potency (Leach et al., 2016), supporting the idea that substitution of residues in the allosteric binding cavity lowers the activation barrier for Ca_o²⁺. Mutations that reduce Ca_o²⁺ potency (e.g., F684^{3,36}A, F688^{3,40}A, and E837^{7,32}I) were at residues critical to the binding of allosteric drugs. Collectively, the influence of these 7TM/ECL residues on both orthosteric agonist activity and small-molecule binding and agonism indicates that the 7TM allosteric binding site has evolutionary importance and raises the possibility that an endogenous allosteric modulator may act via this site.

With regards to mutational effects on agonism, it was of particular note that aside from L776^{5,43} and V817^{6,49}, which were important for the efficacy of both Ca_o²⁺ and AC265347, the majority of residues differentially contributed to orthosteric versus allosteric efficacy. This was demonstrated by the opposing effect of many mutations, which increased Ca_o²⁺ potency yet decreased AC265347 efficacy. We cannot exclude the possibility that changes in Ca_o²⁺ potency were due to altered Ca_o²⁺ affinity, but we did show this was not the case for F684^{3,36}A and F688^{3,40}A mutations (Supplemental Fig. 4). With the exception of E837^{7,32}, which we postulate may bind Ca_o²⁺, it is unlikely that mutation of other residues in this pocket significantly alter Ca_o²⁺ affinity, given that four key

Ca_o²⁺ binding sites are located in the receptor's VFT domain (Geng et al., 2016). Thus, our data support the existence of global and ligand-specific activation mechanisms in the CaSR's 7TM and ECL regions.

Another key finding of this study is the demonstration that CaSR PAMs act by potentiating the affinity of Ca_o²⁺. This is in contrast to other class C GPCR PAMs, which enhance orthosteric agonist-mediated efficacy but do not appear to potentiate their binding (Gregory et al., 2012). The significant loss in PAM affinity in the absence of ambient cations versus in their presence is reminiscent of the reciprocal interaction between Ca_o²⁺ and L-amino acids (Conigrave et al., 2000, 2004); structural studies suggest both ligands are bound to the CaSR's VFT domain to achieve a fully active receptor conformation (Geng et al., 2016; Zhang et al., 2016). However, unlike L-amino acids, our current and previous studies (Leach et al., 2016) demonstrate that small-molecule PAMs bind in an extended CaSR 7TM domain cavity.

Our findings have important implications for future drug discovery efforts at the CaSR. Cinacalcet use is restricted due to an adverse risk of hypocalcemia (Chonchol et al., 2009), in part caused by oversuppression of parathyroid hormone secretion along with stimulation of thyroid CaSRs and consequent calcitonin secretion, and/or suppression of Ca_o²⁺ reabsorption via activation of renal CaSRs (reviewed in Leach et al., 2014, 2015). There is consequently continued interest in developing CaSR activators with greater tissue selectivity. Indeed, weak (partial) agonists or PAMs would be expected to stimulate robust pharmacological responses only in tissues where strong stimulus-response coupling exists, such as high CaSR-expressing parathyroid cells. To date, no small-molecule CaSR agonists devoid of PAM activity have been identified, and there have been no reported efforts to examine the physiologic consequences of PAMs with different degrees of cooperativity. Furthermore, as demonstrated by AC265347, some PAMs also exhibit agonism, which may heighten adverse effects by further amplifying receptor signaling. A greater appreciation of the drivers of CaSR agonism versus allosteric potentiation may afford an opportunity to rationally tune in or tune out PAM and agonist activity.

In conclusion, the current study provides new insight into 7TM and ECL CaSR activation mechanisms. In the future, this information could facilitate rational structure-based design of novel CaSR activators whose pharmacological and consequent therapeutic properties could be optimized by fine-tuning their PAM and/or agonist activity.

Authorship Contributions

Participated in research design: Gregory, Leach.

Conducted experiments: Keller, Kufareva, Josephs, Diao, Mai, Gregory, Leach.

Performed data analysis: Gregory, Leach.

Wrote or contributed to the writing of the manuscript: Keller, Kufareva, Conigrave, Christopoulos, Gregory, Leach.

References

- Abagyan R and Totrov M (1994) Biased probability Monte Carlo conformational searches and electrostatic calculations for peptides and proteins. *J Mol Biol* **235**: 983–1002.
- Aurelio L, Valant C, Flynn BL, Sexton PM, Christopoulos A, and Scammells PJ (2009) Allosteric modulators of the adenosine A1 receptor: synthesis and pharmacological evaluation of 4-substituted 2-amino-3-benzoylthiophenes. *J Med Chem* **52**:4543–4547.
- Bu L, Michino M, Wolf RM, and Brooks CL, III (2008) Improved model building and assessment of the calcium-sensing receptor transmembrane domain. *Proteins* **71**: 215–226.

- Chonchol M, Locatelli F, Abboud HE, Charytan C, de Francisco AL, Jolly S, Kaplan M, Roger SD, Sarkar S, Albizem MB, et al. (2009) A randomized, double-blind, placebo-controlled study to assess the efficacy and safety of cinacalcet HCl in participants with CKD not receiving dialysis. *Am J Kidney Dis* **53**:197–207.
- Christopher JA, Aves SJ, Bennett KA, Doré AS, Errey JC, Jazayeri A, Marshall FH, Okrasa K, Serrano-Vega MJ, Tehan BG, et al. (2015) Fragment and structure-based drug discovery for a class C GPCR: discovery of the mGlu₅ negative allosteric modulator HTL14242 (3-chloro-5-[6-(5-fluoropyridin-2-yl)pyrimidin-4-yl]-benzonitrile). *J Med Chem* **58**:6653–6664.
- Christopoulos A (1998) Assessing the distribution of parameters in models of ligand-receptor interaction: to log or not to log. *Trends Pharmacol Sci* **19**:351–357.
- Conigrave AD, Mun HC, Delbridge L, Quinn SJ, Wilkinson M, and Brown EM (2004) L-amino acids regulate parathyroid hormone secretion. *J Biol Chem* **279**:38151–38159.
- Conigrave AD, Quinn SJ, and Brown EM (2000) L-amino acid sensing by the extracellular Ca²⁺-sensing receptor. *Proc Natl Acad Sci USA* **97**:4814–4819.
- Cook AE, Mistry SN, Gregory KJ, Furness SG, Sexton PM, Scammells PJ, Conigrave AD, Christopoulos A, and Leach K (2015) Biased allosteric modulation at the CaS receptor engendered by structurally diverse calcimimetics. *Br J Pharmacol* **172**:185–200.
- Davey AE, Leach K, Valant C, Conigrave AD, Sexton PM, and Christopoulos A (2012) Positive and negative allosteric modulators promote biased signaling at the calcium-sensing receptor. *Endocrinology* **153**:1232–1241.
- de Graaf C, Song G, Cao C, Zhao Q, Wang MW, Wu B, and Stevens RC (2017) Extending the structural view of class B GPCRs. *Trends Biochem Sci* **42**:946–960.
- Doré AS, Okrasa K, Patel JC, Serrano-Vega M, Bennett K, Cooke RM, Errey JC, Jazayeri A, Khan S, Tehan B, et al. (2014) Structure of class C GPCR metabotropic glutamate receptor 5 transmembrane domain. *Nature* **511**:557–562.
- Geng Y, Mosyak L, Kurinov I, Zuo H, Sturchler E, Cheng TC, Subramanyam P, Brown AP, Brennan SC, Mun HC, et al. (2016) Structural mechanism of ligand activation in human calcium-sensing receptor. *eLife* **5**:e13662.
- Gregory KJ, Hall NE, Tobin AB, Sexton PM, and Christopoulos A (2010) Identification of orthosteric and allosteric site mutations in M₂ muscarinic acetylcholine receptors that contribute to ligand-selective signaling bias. *J Biol Chem* **285**:7459–7474.
- Gregory KJ, Nguyen ED, Malosh C, Mendenhall JL, Zic JZ, Bates BS, Noetzel MJ, Squire EF, Turner EM, Rook JM, et al. (2014) Identification of specific ligand-receptor interactions that govern binding and cooperativity of diverse modulators to a common metabotropic glutamate receptor 5 allosteric site. *ACS Chem Neurosci* **5**:282–295.
- Gregory KJ, Nguyen ED, Reiff SD, Squire EF, Stauffer SR, Lindsley CW, Meiler J, and Conn PJ (2013) Probing the metabotropic glutamate receptor 5 (mGlu₅) positive allosteric modulator (PAM) binding pocket: discovery of point mutations that engender a “molecular switch” in PAM pharmacology. *Mol Pharmacol* **83**:991–1006.
- Gregory KJ, Noetzel MJ, Rook JM, Vinson PN, Stauffer SR, Rodriguez AL, Emmitte KA, Zhou Y, Chun AC, Felts AS, et al. (2012) Investigating metabotropic glutamate receptor 5 allosteric modulator cooperativity, affinity, and agonism: enriching structure-function studies and structure-activity relationships. *Mol Pharmacol* **82**:860–875.
- Langmead CJ, Fry VA, Forbes IT, Branch CL, Christopoulos A, Wood MD, and Herdon HJ (2006) Probing the molecular mechanism of interaction between 4-*n*-butyl-1-[4-(2-methylphenyl)-4-oxo-1-butyl]-piperidine (AC-42) and the muscarinic M₁ receptor: direct pharmacological evidence that AC-42 is an allosteric agonist. *Mol Pharmacol* **69**:236–246.
- Leach K, Conigrave AD, Sexton PM, and Christopoulos A (2015) Towards tissue-specific pharmacology: insights from the calcium-sensing receptor as a paradigm for GPCR (patho)physiological bias. *Trends Pharmacol Sci* **36**:215–225.
- Leach K and Gregory KJ (2017) Molecular insights into allosteric modulation of class C G protein-coupled receptors. *Pharmacol Res* **116**:105–118.
- Leach K, Gregory KJ, Kufareva I, Khajehali E, Cook AE, Abagyan R, Conigrave AD, Sexton PM, and Christopoulos A (2016) Towards a structural understanding of allosteric drugs at the human calcium-sensing receptor. *Cell Res* **26**:574–592.
- Leach K, Loiacono RE, Felder CC, McKinzie DL, Mogg A, Shaw DB, Sexton PM, and Christopoulos A (2010) Molecular mechanisms of action and in vivo validation of an M4 muscarinic acetylcholine receptor allosteric modulator with potential antipsychotic properties. *Neuropsychopharmacology* **35**:855–869.
- Leach K, Sexton PM, and Christopoulos A (2007) Allosteric GPCR modulators: taking advantage of permissive receptor pharmacology. *Trends Pharmacol Sci* **28**:382–389.
- Leach K, Sexton PM, Christopoulos A, and Conigrave AD (2014) Engendering biased signalling from the calcium-sensing receptor for the pharmacotherapy of diverse disorders. *Br J Pharmacol* **171**:1142–1155.
- Leach K, Wen A, Cook AE, Sexton PM, Conigrave AD, and Christopoulos A (2013) Impact of clinically relevant mutations on the pharmacoregulation and signaling bias of the calcium-sensing receptor by positive and negative allosteric modulators. *Endocrinology* **154**:1105–1116.
- Manglik A and Kruse AC (2017) Structural basis for G protein-coupled receptor activation. *Biochemistry* **56**:5628–5634.
- May LT, Leach K, Sexton PM, and Christopoulos A (2007) Allosteric modulation of G protein-coupled receptors. *Annu Rev Pharmacol Toxicol* **47**:1–51.
- Mayr B, Schnabel D, Dörr HG, and Schöfl C (2016) Genetics in endocrinology: gain and loss of function mutations of the calcium-sensing receptor and associated proteins: current treatment concepts. *Eur J Endocrinol* **174**:R189–R208.
- Miedlich SU, Gama L, Seuwen K, Wolf RM, and Breitwieser GE (2004) Homology modeling of the transmembrane domain of the human calcium sensing receptor and localization of an allosteric binding site. *J Biol Chem* **279**:7254–7263.
- Motulsky H and Christopoulos A (2004) *Fitting Models to Biological Data Using Linear and Nonlinear Regression. A Practical Guide to Curve Fitting*, Oxford University Press, New York.
- Nemeth EF and Goodman WG (2016) Calcimimetic and calcilytic drugs: feats, flops, and futures. *Calcif Tissue Int* **98**:341–358.
- Nemeth EF, Steffey ME, Hammerland LG, Hung BC, Van Wagenen BC, DelMar EG, and Balandrin MF (1998) Calcimimetics with potent and selective activity on the parathyroid calcium receptor. *Proc Natl Acad Sci USA* **95**:4040–4045.
- Petrel C, Kessler A, Dauban P, Dodd RH, Rognan D, and Ruat M (2004) Positive and negative allosteric modulators of the Ca²⁺-sensing receptor interact within overlapping but not identical binding sites in the transmembrane domain. *J Biol Chem* **279**:18990–18997.
- Petrel C, Kessler A, Maslah F, Dauban P, Dodd RH, Rognan D, and Ruat M (2003) Modeling and mutagenesis of the binding site of Calhcx 231, a novel negative allosteric modulator of the extracellular Ca²⁺-sensing receptor. *J Biol Chem* **278**:49487–49494.
- Ray K and Northup J (2002) Evidence for distinct cation and calcimimetic compound (NPS 568) recognition domains in the transmembrane regions of the human Ca²⁺ receptor. *J Biol Chem* **277**:18908–18913.
- Zhang C, Zhang T, Zou J, Miller CL, Gorkhali R, Yang JY, Schillmiller A, Wang S, Huang K, Brown EM, et al. (2016) Structural basis for regulation of human calcium-sensing receptor by magnesium ions and an unexpected tryptophan derivative co-agonist. *Sci Adv* **2**:e1600241.

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