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Activated CaMKII α binds to the mGlu₅ metabotropic glutamate receptor and modulates calcium mobilization

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Running title: CaMKII modulates mGlu₅ by interaction with the CTD

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Abbreviations: CaMKII, Calcium Calmodulin Dependent Protein Kinase II; CA-CaMKII, constitutively active CaMKII; CaM, Calmodulin; CaMKAP, CaMKII Associated Protein; CTD, C-terminal Domain; D₂R, D2 Dopamine Receptor; D₃R, D3 Dopamine Receptor; GPCR, G protein-coupled receptor, GST, Glutathione S-Transferase, GST-mGlu_{5a}-CTD, GST tagged mGlu_{5a} construct containing residues 827-964; KO, knock-out; LTD, long term depression; LTP, long term potentiation; mGlu, metabotropic glutamate receptor; mGlu₁, metabotropic glutamate receptor 1; mGlu₅, metabotropic glutamate receptor 5; PKA, protein kinase A; PKC, protein kinase C; WT, wild-type.

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Abstract

Ca²⁺ calmodulin dependent protein kinase II (CaMKII) and metabotropic glutamate receptor 5 (mGlu₅) are critical signaling molecules in synaptic plasticity and learning memory. Here, we demonstrate that mGlu₅ is present in CaMKII α complexes isolated from mouse forebrain. Further *in vitro* characterization showed that the membrane-proximal region of the C-terminal domain (CTD) of mGlu_{5a} directly interacts with purified Thr286-autophosphorylated (activated) CaMKII α . However, the binding of CaMKII α to this CTD fragment is reduced by the addition of excess Ca²⁺/calmodulin or by additional CaMKII α autophosphorylation at non-Thr286 sites. Furthermore, *in vitro* binding of CaMKII α is dependent on a tribasic residue motif Lys-Arg-Arg (KRR) at residues 866-868 of the mGlu_{5a}-CTD, and mutation of this motif decreases the co-immunoprecipitation of CaMKII α with full-length mGlu_{5a} expressed in heterologous cells by about 50%. The KRR motif is required for two novel functional effects of co-expressing constitutively active CaMKII α with mGlu_{5a} in heterologous cells. First, cell-surface biotinylation studies showed that CaMKII α increases the surface-expression of mGlu_{5a}. Second, using Ca²⁺ fluorimetry and single-cell Ca²⁺-imaging, we found that CaMKII α reduces the initial peak of mGlu_{5a}-mediated Ca²⁺ mobilization by about 25%, while doubling the relative duration of the Ca²⁺ signal. These findings provide new insights into the physical and functional coupling of these key regulators of postsynaptic signaling.

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Introduction

The ability of excitatory glutamatergic synapses to undergo dynamic changes in strength, termed synaptic plasticity, is critical for many behaviors. It is well established that glutamate activation of diverse ionotropic and metabotropic receptors is critical for short- and long-term control of many neuronal responses (Niswender and Conn, 2010), and that these responses require a complex and incompletely understood network of signaling proteins. Among the 7 members of the metabotropic glutamate (mGlu) receptor family, mGlu₁ and mGlu₅ specifically couple through G $\alpha_{q/11}$ to stimulate multiple signaling pathways, including phosphoinositide hydrolysis and mobilization of intracellular Ca²⁺ stores. These receptors have long been implicated in multiple forms of long-term depression (LTD) that require new protein synthesis (Huber et al., 2001; Oliet et al., 1997; Palmer et al., 1997) or increased endocannabinoid signaling (Luscher and Huber, 2010). Despite many similarities, mGlu₁ and mGlu₅ can be differentially regulated by various mechanisms and have been shown to have different neuronal roles. For instance, in hippocampus, mGlu₁ increases the frequency of spontaneous inhibitory post synaptic currents while mGlu₅ potentiates NMDA receptor currents (Mannaioni et al., 2001). In particular, mGlu₅ has been specifically implicated in a number of neuropsychiatric disorders including addiction, schizophrenia, Fragile X Syndrome, Obsessive Compulsive Disorder, and Alzheimer's Disease (Ade et al., 2016; Foster and Conn, 2017; Grueter et al., 2008; Hu et al., 2014; Michalon et al., 2012; Ronesi et al., 2012).

Like mGlu₅, Ca²⁺/calmodulin (CaM)-dependent protein kinase II α (CaMKII α) is a key signaling protein in dendritic spines. CaMKII α is activated by Ca²⁺/CaM binding and undergoes autophosphorylation at Thr286 (Baucum et al., 2015; Miller et al., 1988; Mukherji et al., 1994; Rich and Schulman, 1998; Yang and Schulman, 1999). Thr286 autophosphorylation increases

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the affinity for $\text{Ca}^{2+}/\text{CaM}$ and stabilizes the kinase in a constitutively active conformation. This constitutive activity is essential for normal synaptic plasticity in many brain regions (Coultrap et al., 2014; Giese et al., 1998; Jin et al., 2015; Mockett et al., 2011; Shonesy et al., 2014; Silva et al., 1992a; Silva et al., 1992b; Zhou et al., 2007) including mGlu_{1/5}-dependent long term depression (LTD) in the hippocampus (Huber et al., 2001; Mockett et al., 2011). Interestingly, both CaMKII α - and mGlu₅-knockout mice display deficits in learning and memory and hippocampal synaptic plasticity (Huber et al., 2001; Jia et al., 1998; Simonyi et al., 2005). Although both mGlu₅ and CaMKII are critical to many forms of plasticity, a functional link between the two has not been widely investigated.

The intracellular C-terminal domains (CTDs) of mGlu₁ and mGlu₅ have emerged as important loci for regulation by protein binding and phosphorylation (Enz, 2012; Mao and Wang, 2016). While there are two splice variants of mGlu₅ (mGlu_{5a} and mGlu_{5b}), most studies have focused on mGlu_{5a} and few pharmacological differences between splice variants have been identified (Joly et al., 1995; Minakami et al., 1995; Romano et al., 1996). The mGlu₅-CTD has been shown to bind to a number of different proteins including $\text{Ca}^{2+}/\text{CaM}$ and Homer to regulate cell surface expression of the receptor (Choi et al., 2011; Lee et al., 2008; Roche et al., 1999; Saito et al., 2002). Protein kinase A (PKA) and protein kinase C (PKC) also regulate mGlu₅ surface expression through phosphorylation of the CTD (Mao et al., 2008; Uematsu et al., 2015). It was recently reported that CaMKII can bind to the CTD and intracellular loop 2 of both mGlu₁ and mGlu₅ (Jin et al., 2013b; Raka et al., 2015) and that CaMKII modulates mGlu₅ agonist-induced internalization and ERK1/2 activation (Raka et al., 2015). Here, we identify three basic residues (Lys⁸⁶⁶-Arg⁸⁷⁷-Arg⁸⁸⁸) in the membrane proximal region of the mGlu_{5a}-CTD that are essential for a direct interaction with activated CaMKII α , and provide novel insights into multiple factors that

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modulate the interaction. We also show that CaMKII binding to the CTD is important for the regulation of mGlu₅ surface expression and Ca²⁺ mobilization. These data provide novel insights into the molecular basis and function of the mGlu₅-CaMKII interaction that may be involved in synaptic plasticity.

Materials and methods

DNA constructs

The GST-mGlu_{5a}-CTD expression construct was created by PCR amplification of the region encoding residues 827-964 of mGlu_{5a} (NP_058708.1) using primers

5'CTGGAAGTTCTGTTCCAGGGGCCCGGATCCAAACCGGAGAGAAAT 3' (Forward) and

5' GCCGCAAGCTTGTCGACGGAGCTCGAATTCTTAGGTCCCAAAGCGCTT 3' (Reverse)

and inserting the product into BamHI/EcoRI sites of pGEX6P using a sequence and ligation independent cloning (SLIC) cloning protocol (Li and Elledge, 2012).

The pCGN plasmid to express WT mGlu_{5a} with an N-terminal HA-tag was made by amplifying the entire rat mGlu_{5a} coding sequence (forward primer:

5'TGACGTGCCTGACTATGCCTCTAGAATGGTCCTTCTGTTGATCCT3'; reverse primer:

5' ACTCACCTGAAGTTCTCAGGATCCTCACAACGATGAAGAACTCT3') and inserting

the fragment into XbaI and BamHI restriction sites of the empty pCGN plasmid (a gift from Dr. Winship Herr, Université de Lausanne, Switzerland, Addgene plasmid ID 53308).

The K⁸⁶⁶RR⁸⁶⁸ mutation to AAA in mGlu_{5a} was generated by site-directed mutagenesis of the pGEX6P or pCGN constructs (see above) using a Quick Change protocol (Agilent) with the following primers

5'GGGTTTCCCCAGAGGAGCCGGCGGCGGCCACAGGTTGACTAGGCTGCT3'

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(Forward) and

5'AGCAGCCTAGTCAACCTGTGGGCGCCGCGGCTCCTCTGGGGAAACCC3'

(Reverse).

We used pcDNA3.1 constructs to express untagged and mApple-tagged WT-CaMKII α and a constitutively active T286D/T305A/T306A triple mutant of CaMKII α (CA-CaMKII α), as previously described (Jalan-Sakrikar et al., 2012; Jiao et al., 2008; Stephenson et al., 2017). In the CA-CaMKII α , the phospho-mimetic T286D mutation results in constitutive CaMKII α activity and the phospho-null T305A/T306A mutations prevent CaMKII α phosphorylation at these sites, which interferes with binding of Ca²⁺/CaM and α -actinin (Jalan-Sakrikar et al., 2012).

Recombinant protein purification

Expression and purification of recombinant mouse CaMKII α has been described before (McNeill and Colbran, 1995). To express GST-tagged proteins, pGEX6P-1 plasmids were transformed into BL21(DE3) bacteria cells. Cells were grown in LB media at 37°C to reach OD~0.6. Cells were cooled to room temperature and IPTG (0.2 mM) was then added to induce protein expression for 12-16 hours. Inducing protein expression at room temperature substantially reduced protein degradation seen when proteins were expressed at 37°C. Expressed proteins were purified using Pierce Glutathione Agarose beads (Cat. #16101) following manufacture's instruction. Eluted proteins were then dialyzed in 10 mM HEPES pH 7.5, 25 μ M PMSF, 62.5 μ M Benzamidine, 62.5 μ M EDTA, 0.1% Triton X-100 overnight with one buffer change.

CaMKII α autophosphorylation

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Purified mouse CaMKII α was autophosphorylated under two different conditions. Typically, CaMKII α was incubated with 50 mM HEPES, pH 7.5, 10 mM Mg(CH₃-COO)₂, 0.5 mM CaCl₂, 2 μ M CaM, 40 μ M ATP on ice for 90 s before addition of EDTA and EGTA (20 mM final) to terminate phosphorylation by chelation of Mg²⁺ and Ca²⁺. Similar conditions were previously shown to result in the selectively autophosphorylation of Thr286 (McNeill and Colbran, 1995). Where indicated, identical autophosphorylation reactions were incubated for 10 minutes at 30°C to perform a more extensive phosphorylation at several additional sites (Baucum et al., 2015).

GST pulldown and CaM binding competition

Purified GST-mGlu₅-CTD (1 μ M) and CaMKII α (62.5 nM; pre-autophosphorylated as indicated in figure legends) were incubated at 4°C in GST-pulldown buffer (50 mM Tris-HCl pH 7.5; 150 mM NaCl; 1% (v/v) Triton X-100) with either 2 mM EGTA or 2.5 mM CaCl₂ plus 10 μ M calmodulin, as indicated. An aliquot (5%) of each incubation was saved as an input sample. After 1 h, pre-washed glutathione agarose beads (Pierce Cat. #16101) (15 μ l of a 50:50 slurry) were added and incubation was continued at 4°C for an additional 1 h. The beads were then separated by centrifugation (2000 x g, 30 s), and washed three times with GST-pulldown buffer containing either 2 mM EGTA or 2.5 mM CaCl₂, respectively. Beads were then incubated at 4°C with GST pulldown buffer containing 20 mM glutathione, adjusted to pH 8.0, for 10 min. After centrifugation, eluted proteins were transferred to a new tube, mixed with 4X SDS-PAGE buffer and heated for 10 minutes at 90°C prior to SDS-PAGE and Western Blot analysis.

Cell culture, transfection and immunoprecipitation

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HEK293A cells (Invitrogen Catalog #R70507) were maintained in complete DMEM supplemented with 5% FBS, 2 mM L-glutamine, 20 mM HEPES, 0.1 mM Non-Essential Amino Acids, 1 mM sodium pyruvate, at 37 °C in a humidified incubator containing 5% CO₂ and 95% O₂. Vectors encoding mApple-CaMKII α (WT or CA) and mGlu_{5a} (3 μ g DNA each) or empty vector controls (3 μ g) were co-transfected into one 10-cm dish of 60-70% confluent HEK293A cells using 3 μ l of Fugene 6 (Promega Catalog # E2691) per μ g of DNA. About 48 hours later, cells were lysed in 50 mM Tris-HCl pH 7.5, 150 mM NaCl, 1 mM EDTA, 1 mM EGTA, 1 mM DTT, 0.5% NP-40 (v/v), 0.5% deoxycholate (v/v), 0.2 mM PMSF, 1 mM benzamidine, 10 μ g/ml leupeptin, 10 μ M pepstatin, and 1 μ M microcystin. Cell lysates were cleared by centrifugation (10 min at 12,000 x g), and a 30 μ L sample of the input was saved for SDS-PAGE. The remaining supernatant was incubated at 4 °C for 1 hour with rabbit anti-HA antibodies and 20 μ l prewashed Dynabeads Protein A (Thermo-Fisher, Cat. #10001D, 50% v/v). Beads were isolated magnetically, washed three times using lysis buffer and eluted using 2X Laemmli sample buffer for 10 minutes at room temperature prior to SDS-PAGE and Western Blotting.

Biotinylation and cell surface expression

Transfected HEK293A cells (see above) were placed on ice, the media was gently removed and the cells were immediately washed two times using ice cold PBS. Cells were then scraped into ice-cold PBS, transferred to a 1.5 mL tube, centrifuged at 4 °C (500 x g; 3 min), and gently resuspended in 1 mL of cold PBS containing 2 mg of EZ-Link Sulfo-NHS-SS-Biotin. After gently rocking for 1 hour, excess reagent was quenched by addition of 50 mM Tris HCl pH 8.0, and cells were centrifuged and washed again in 1 mL of 50 mM Tris HCl. Cells were then suspended in 1 mL of ice-cold lysis buffer (25 mM Tris HCl, pH 7.4, 150 mM NaCl, 1% NP-40,

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0.5% sodium deoxycholate containing 0.2 mM PMSF, 1 mM benzamidine, 10 µg/ml leupeptin, and 10 µM pepstatin), and incubated on ice for 30 min. Insoluble material was removed by centrifugation (16,000 x g; 10 min, 4°C) and a 30 µL aliquot of the supernatant was saved for an input sample for SDS-PAGE (Cho et al., 2014). The remaining supernatants were mixed for 1 hour at 4°C with magnetic NeutrAvidin beads (30 µL; 50% slurry). The beads were separated magnetically and washed three times with lysis buffer. Biotinylated proteins were dissociated from the beads in SDS sample buffer containing 150 mM DTT for 10 minutes at room temperature. The biotinylated and total protein samples were analyzed by Western blotting for mGlu5.

Immunoblotting and semi-quantitative analysis

Since heating samples results in aggregation of full-length mGlu5 protein, all samples that were blotted for the full-length receptor were eluted in SDS sample buffer containing 150 mM DTT for 10 minutes at room temperature before SDS-PAGE. SDS-polyacrylamide gels were transferred to nylon-backed nitrocellulose membranes in 10 mM CAPS buffer. After blocking in TTBS (50 mM Tris-HCl, pH 7.5, 0.1% (v/v) Tween 20, 150 mM NaCl) containing 5% nonfat milk, membranes were incubated for either 2 h at room temperature for purified protein studies or overnight at 4 °C in HEK293A and brain lysate samples with primary antibodies diluted in TTBS with 5% milk. Membranes were washed 5 times in TTBS and incubated for 1 h at room temperature with secondary antibodies conjugated to horseradish peroxidase (Promega or Santa Cruz Biotechnology), or infrared dyes (LiCor Biosciences, Lincoln, NE) diluted in TTBS with 5% milk. Antibody signals were visualized via enzyme-linked chemiluminescence using the Western Lightening Plus-ECL, enhanced chemiluminescent substrate (PerkinElmer) and

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visualized using Premium X-ray Film (Phenix Research Products). Images were quantified using ImageJ software. Secondary antibodies conjugated to infrared dyes (LI-COR Biosciences) were used for development with an Odyssey system (LI-COR Biosciences).

Antibodies

The following antibodies were used for immunoblotting at the indicated dilutions: Total CaMKII α (Thermo Catalog # MA1-048 1:5000) and p-Thr286 CaMKII α (Santa Cruz Biotechnology Catalog # sc-12886-R, 1:3000), mGlu₅-specific antibody (Millipore, Catalog # AB5675, 1:3000), rabbit anti-HA (Santa Cruz, Catalog #sc805, 5 μ L for immunoprecipitation), goat-GST antibody (Abcam Catalog # ab181652, 1:10,000).

Secondary antibodies: HRP-conjugated anti-rabbit (Promega catalog #W4011, 1:3000), HRP-conjugated anti-mouse (Promega catalog #W4021, 1:3000), and HRP-conjugated anti-goat (Santa Cruz Biotechnology catalog #sc-2056, 1:3000), IR dye-conjugated donkey anti-rabbit 800CW (LI-COR Biosciences catalog #926–32213, 1:10,000), and IR dye-conjugated donkey anti-mouse 680LT (LI-COR Biosciences catalog #926–68022, 1:10,000).

Mice

CaMKII-KO mice were generated in the Vanderbilt Transgenic Mouse Core as a by-product of published CRISPR/Cas9 mediated experiments directed at creating a knock-in E183V mutation of CaMKII α (Stephenson et al., 2017). We selected a founder containing a deletion of 11-base pairs (TGCTGAGGAAG) from exon 8, leading to a frame shift and early translational termination. The knockout of CaMKII α was confirmed by Western Blot. Primers used to genotype the CaMKII α knockout mice are: (Forward) 5'GATACCTCTCCCCAGAAGGAC3', (Reverse) 5'TGCAGTGGTAAGGAGTGGTG3' for wild-type and (Forward)

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5'GGACAGTACAACCCCAGCTT3' and (Reverse) 5'CCCGTACGGGGTCCTTCCTCA3' for knockout generating a 206bp band for WT, 351bp band for KO, and 557bp band for all mice and the CaMKII α -KO was confirmed by immunoblotting brain lysates. All mice were on a mixed B6D2 (C57BL/6J (B6) x DBA/2J (D2)) background and were housed (2–5 per cage) on a 12 h light-dark cycle with food and water *ad libitum*. Wild-type and knockout experimental mice (littermates) were generated using a HETXHET breeding strategy. All animal procedures were approved by the Vanderbilt University Institutional Animal Care and Use Committee in accordance with the National Institutes of Health *Guide for the care and use of laboratory animals*.

Mouse brain tissue preparation and immunoprecipitation

Both male and female mice were anesthetized with isoflurane, decapitated, and forebrains were quickly dissected. Half of a forebrain (cut along the mid-line) was homogenized using at least 20 strokes with a Dounce homogenizer in 1.5 mL of an isotonic buffer containing 150 mM KCl, 50 mM Tris-HCl, 1 mM DTT, 1% (v/v) Triton X-100, 1% sodium deoxycholate, 0.2 mM PMSF, 1 mM benzamidine, 10 μ g/ml leupeptin, 10 μ M pepstatin, and 1 μ M microcystin. The homogenate was rotated end-over-end at 4°C for 30 min and then centrifuged at 10,000 \times g for 30 min to remove insoluble material. A 30 μ L input sample was saved before CaMKII α (MA1-048) antibody and 20 μ L magnetic Protein G beads (Invitrogen Catalog #10003D) was added to 1 mL of homogenate and rotated end over end for 3-4 hours. Beads were separated magnetically and washed three times with homogenization buffer. Immunoprecipitated complexes were eluted using 2X Lamelli Sample Buffer containing 150 mM DTT for 10 minutes at room temperature. CaMKII and mGlu5 immunoprecipitation was analyzed by immunoblot analysis.

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Ca²⁺ Imaging in 96-well Plates

A FlexStation II liquid handler/plate reader (Molecular Devices) was used for intracellular Ca²⁺ measurements in HEK293A cells stably expressing low amounts of the rat mGlu_{5a} receptor (293A-5α^{LOW} cells) as previously described (Gregory et al., 2012; Hammond et al., 2010; Noetzel et al., 2012). The cells were maintained at 37°C in complete DMEM supplemented with 10% fetal bovine serum, 2 mM l-glutamine, 20 mM HEPES pH 7.5, 0.1 mM nonessential amino acids, 1 mM sodium pyruvate, antibiotic/antimycotic solution (Invitrogen), and 500 µg/ml G418, in a humidified incubator containing 5% CO₂/95% O₂. For experiments, 10 cm dishes were transfected with 3 µg of mApple control vector (mApp) or 3 µg mApp-CaMKIIα (wild type or CA; see above). The following day, cells were transferred to clear-bottomed, black-walled, poly(d-lysine)-coated 96-well plates (BD BioCoat, Bedford, MA) (3 × 10⁴ cells per well) in DMEM containing 10% dialyzed FBS, 20 mM HEPES, 1 mM sodium pyruvate, and incubated overnight at 37°C in 5% CO₂. Approximately 24 hours later, medium was manually removed and replaced with Hanks' balanced salt solution containing 20 mM HEPES, 2.5 mM probenecid, and 2 µM Fluo-4/acetoxymethyl ester dye, pH 7.4, and plates were incubated for 30 min (37°C, 5% CO₂). This medium was manually removed and replaced with 40 µl of calcium assay buffer (Hanks' balanced salt solution, 20 mM HEPES, and 2.5 mM probenecid, pH 7.4). Glutamate additions were performed after a 30 s baseline to construct concentration-response curves (CRCs) and plates were monitored for a total of 120 sec using an excitation wavelength of 488 nm, an emission wavelength of 525 nm, and a cutoff wavelength of 515 nm. Data were collected with SoftMax Pro (Molecular Devices) then transformed and agonist concentration-response curves were fitted to a four-parameter logistic equation with GraphPad Prism. For area under the

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curve measurements, time parameters were set to measure the area under the curve between the timepoints of 10-60 seconds to capture the initial Ca^{2+} peak. Raw values were generated by SoftMax Pro and normalized to the max response of control cells.

Single Cell Ca^{2+} Imaging

HEK293A cells were transiently transfected to express full-length mGlu_{5a} (WT or AAA: 0.3 μg DNA) and either mApple or mApple-CA-CaMKIIα (3 μg DNA). The following day, transfected cells were plated in clear glass-bottomed, poly(D-lysine)-coated 29 mM dishes (Cellvis D29-10-1.5-N) (5×10^4 cells) in DMEM containing 10% dialyzed FBS, 20 mM HEPES, 1 mM sodium pyruvate, and 1% Pennicillin/Streptomycin (Gibco) and incubated overnight at 37°C in 5% CO₂. The day of the experiment, cells were incubated in media supplemented with 2 μM Fura-2 acetoxymethyl ester (Molecular Probes) for 20 minutes at 37°C in 5% CO₂, and then transferred to Ca^{2+} imaging solution (150 mM NaCl, 5 mM KCl, 2 mM CaCl₂, 2 mM MgCl₂, 10 mM glucose and 10 mM HEPES pH 7.5 (~313 mOsm)). After incubation for 20 minutes at 37°C in 5% CO₂, fluorescence imaging was performed using a Nikon Eclipse TE2000-U microscope equipped with an epifluorescence illuminator (Sutter, Inc), a CCD camera (HQ2; Photometrics Inc), and Nikon Elements software. Cells were perfused at 37°C at a flow of 2 mL/min with Ca^{2+} imaging solution. First, the field of view was imaged using 568 nm excitation to detect cells expressing either mApple-CA-CaMKII or mApp-PCDNA3.1 control. Then, ratios of emitted fluorescence (at 510 nm) from mApple positive cells were measured following excitation at 340 nm and 380 nm (F340/F380); ratios were measured every 3 seconds for a 1 min baseline period. Then the cells were treated with 100 μM glutamate (added to the Ca^{2+} imaging solution) for 10 min during which the FURA2 F340/F380 ratios were collected every 3 s. Relative changes in

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Ca²⁺ of the mApple expressing cells were analyzed using Nikon Elements software. The F340/F380 ratios of each cell were normalized to the first F340/F380 ratio acquired for that cell during the baseline period ($F/F_0 = (340/380 \text{ value}) / (\text{baseline } 340/380 \text{ value})$) and then analyzed using Clampfit software (Molecular Devices, Sunnyvale, California). Peak Ca²⁺ responses for all cells were aligned (at 45 seconds), a ROUT test was first used to identify outliers in the maximal $\Delta F/F_0$ values for all cells within each experimental group, and then $\Delta F/F_0$ values for each time point were averaged together for each dish of cells. The maximal Ca²⁺ response (Max Peak) was defined as the average of all the peak $\Delta F/F_0$ values on an experimental day. An average trace for each day of experiments was generated to calculate the half-life of each condition. To compare half-lives of the Ca²⁺ signals the decline of the average $\Delta F/F_0$ values in each dish were normalized to the average max peak in that dish, and then fitted to a non-linear one phase exponential decay fit constrained to $y_0 = 1$ using Graphpad Prism software (version 6.0). We determined $\Delta F/F_0$ and half lives in 5 independent experiments (transfections) on separate days (19-124 cells per condition per day) and tested for differences using a Student's T-Test. All values are presented as the mean \pm SEM.

Results

Mouse forebrain lysates contain CaMKII α -mGlu₅ complexes

In order to confirm that mGlu₅ specifically associates with CaMKII in the brain, we incubated forebrain lysates from WT or CaMKII α -KO mice with a CaMKII α -specific monoclonal antibody or a control IgG. The resulting immune complexes were isolated and then immunoblotted for mGlu₅ and CaMKII α . Note that the mGlu₅ antibody used for these studies recognizes an epitope

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that is shared by the two known mGlu₅ splice variants, mGlu_{5a} and mGlu_{5b}, which are differentially expressed during development (Minakami et al., 1995; Romano et al., 1996). Input samples from WT and CaMKII α -KO tissue prepared in parallel contain similar levels of the monomeric and dimeric forms of mGlu₅ (Figure 1), although the ratio of monomeric and dimeric species varied between independent experiments (not shown). CaMKII α complexes isolated from WT mouse forebrain contained both monomeric and dimeric forms of mGlu₅, with the ratio of these forms reflecting variability in the ratio detected in the inputs. However, very little mGlu₅ could be detected in IgG control complexes isolated from WT tissue, or in CaMKII α complexes isolated from CaMKII α -KO tissue (Figure 1). Thus, mGlu₅ is a *bona fide* component of the CaMKII α complexes present in mouse brain lysates.

CaMKII α directly binds to the mGlu₅ C-terminal domain

Like most prior studies, we chose to use the mGlu_{5a} splice variant for our molecular studies. It was previously reported that residues 827-964 of the mGlu_{5a}-CTD bind to inactive CaMKII α , but that CaMKII autophosphorylation disrupted the interaction (Jin et al., 2013b). In order to confirm this finding, we generated a GST-tagged mGlu_{5a} CTD construct containing residues 827-964 (GST-mGlu_{5a}-CTD) for use in glutathione agarose co-sedimentation experiments. Initial studies detected weak binding of inactive CaMKII α to GST-mGlu_{5a}-CTD that was not consistently above background binding to a GST negative control (data not shown). Therefore, we systematically tested interactions of GST-mGlu_{5a}-CTD with CaMKII α in various activation states.

Purified CaMKII α was autophosphorylated in the presence of Ca²⁺/CaM *in vitro* at either 4°C or at 30°C. Similar total levels of Thr286 autophosphorylation were detected by immunoblotting

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following incubation at either 4°C or at 30°C (Figure 2A), but the 30°C autophosphorylation reduced the electrophoretic mobility of CaMKII α . These observations are consistent with prior studies showing that incubation at 4°C results in selective Thr286 autophosphorylation (McNeill and Colbran, 1995), whereas incubation at 30°C allows for extensive autophosphorylation at several other sites (Baucum et al., 2015).

We then performed glutathione-agarose cosedimentation experiments to test the interaction of GST-mGlu₅-CTD with CaMKII α in these different activation states (after terminating the autophosphorylation reactions by chelating metal ions with excess EGTA and EDTA). The selective Thr286-autophosphorylation (4°C) protocol resulted in a robust enhancement of CaMKII α binding to GST-mGlu₅-CTD relative to the non-phosphorylated kinase, but this interaction was substantially reduced following more extensive *in vitro* phosphorylation at 30°C (Figure 2B). The short exposure times used for development of these immunoblots failed to detect weak binding of inactive CaMKII α to GST-mGlu₅-CTD. In combination, these data show that while activation and Thr286 autophosphorylation of CaMKII α strongly enhances binding to the mGlu_{5a} CTD, the interaction can be weakened by autophosphorylation at additional non-Thr286 sites.

Binding of activated CaMKII α to the GST-mGlu_{5a} CTD is disrupted by Ca²⁺/CaM

Ca²⁺/CaM binds to residues 889-917 within the CTD of mGlu_{5a} with important functional consequences (Choi et al., 2011; Lee et al., 2008; Minakami et al., 1997). Moreover, it was previously reported that excess Ca²⁺/CaM disrupts the binding of inactive CaMKII α to the mGlu_{5a}-CTD (Jin et al., 2013b). Therefore, we tested whether excess Ca²⁺/CaM also disrupts the binding of activated CaMKII α to GST-mGlu_{5a}-CTD. Thr286-autophosphorylated CaMKII α (4°C

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protocol) robustly binds to GST-mGlu_{5a}-CTD, as noted above, but this interaction was essentially eliminated by inclusion of excess Ca²⁺/CaM in the binding assay (Figure 2C). Thus, binding of activated CaMKII α to the mGlu_{5a}-CTD is also blocked by Ca²⁺/CaM, suggesting that multiple Ca²⁺ sensitive proteins are involved in the regulation of mGlu₅ signaling.

Identification of a CaMKII α -binding determinant in the mGlu₅-CTD

As an initial approach to identify key CaMKII α binding determinants in the mGlu_{5a}-CTD, we compared residues 827-964 of mGlu₅ with CaMKII α -binding domains that have been previously identified in other proteins. Our lab recently showed that activated CaMKII α binds to the N-terminal domains of Cav1.2 and Cav1.3 L-type voltage gated Ca²⁺ channels, and that this interaction is disrupted by mutation of three basic residues (Arg⁸³-Lys-Arg⁸⁵) to alanine (Wang et al., 2017). Similar tri-basic residue motifs are also present within CaMKII α binding domains that have been previously identified in the intracellular loops of the D₂ and D₃ dopamine receptor (Liu et al., 2009; Zhang et al., 2014), and the mGlu₁-CTD (Jin et al., 2013a). Notably, the CaMKII α -binding fragment of the mGlu_{5a}-CTD also contains a tribasic residue motif (residues Lys⁸⁶⁶-Arg⁸⁶⁷-Arg⁸⁶⁸) (Figure 3A). We found that substituting alanines for Lys⁸⁶⁶-Arg⁸⁶⁷-Arg⁸⁶⁸ in the mGlu_{5a}-CTD essentially abolished the binding of activated CaMKII α to GST-mGlu₅-CTD *in vitro* (Figure 3B). These data identify a key determinant for CaMKII-binding to the CTD of mGlu_{5a}.

CaMKII α activation increases CaMKII α -mGlu_{5a} association in heterologous cells

In order to better understand the interaction of CaMKII α with full-length mGlu_{5a} we conducted co-immunoprecipitation experiments from lysates of transfected HEK293A cells. We first tested

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the hypothesis that CaMKII α activation would increase the association with full-length mGlu_{5a}, as with *in vitro* binding of CaMKII α to GST-mGlu_{5a}-CTD. We expressed mApple-tagged WT CaMKII α in the absence or presence of mGlu_{5a} with an N-terminal HA-epitope tag in HEK293A cells. Prior to HA immunoprecipitation, the cell lysates were split into two aliquots and pre-incubated with either excess EGTA and EDTA or with Ca²⁺/CaM, Mg²⁺, ATP, and phosphatase inhibitors to stimulate CaMKII α autophosphorylation. CaMKII activation in the lysates resulted in a robust increase in autophosphorylation at Thr286, without the large shift in electrophoretic mobility that was observed following autophosphorylation of purified CaMKII α at 30°C (Figure 4A). HA-immunoprecipitation from the two preincubated lysates yielded similar amounts of the monomeric and dimeric species of HA-mGlu_{5a}, but CaMKII α activation resulted in a statistically significant ~3-fold increase in the amount of co-immunoprecipitated CaMKII α (Figure 4A). These data show that full-length mGlu_{5a} preferentially interacts with activated WT CaMKII α .

Association of activated CaMKII α with full-length mGlu_{5a} requires Arg⁸³-Lys-Arg⁸⁵

We next investigated whether the association of activated CaMKII α with full-length mGlu_{5a} involves the CTD. In order to avoid complications that might arise from preincubating cell lysates to activate WT-CaMKII α , we used an mApple-tagged constitutively active T286D/T305A/T306A triple mutant of CaMKII α (mApple-CA-CaMKII α); the phospho-mimetic T286D mutation results in constitutive CaMKII α activity and the phospho-null T305A/T306A mutations prevent CaMKII α phosphorylation at these sites, which interferes with binding of Ca²⁺/CaM and α -actinin (Jalan-Sakrikar et al., 2012). The mApple-CA-CaMKII α was expressed alone, or co-expressed with either HA-mGlu_{5a} or HA-mGlu_{5a}-AAA (with Lys⁸⁶⁶-Arg⁸⁶⁷-Arg⁸⁶⁸

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mutated to alanines). HA-immunoprecipitation from cell lysates confirmed a robust association of mApple-CA-CaMKII α with WT mGlu_{5a} that was partially (~50%) reduced by the triple alanine mutation in the CTD (Figure 4B). These data demonstrate that the Lys⁸⁶⁶-Arg⁸⁶⁷-Arg⁸⁶⁸ residues in the mGlu_{5a}-CTD play an important role in the association of activated CaMKII α with the full-length mGlu₅ receptor.

CaMKII α increases basal mGlu_{5a} surface expression

Since the CTD is known to modulate mGlu₅ cell surface expression and consequently mGlu₅ signaling, we investigated the effect of CaMKII α on the cell-surface expression of full-length mGlu_{5a}. Intact HEK293A cells expressing mGlu₅ with or without mApple-CA-CaMKII α were incubated with Sulfo-NHS-SS-Biotin to biotinylate all surface-expressed proteins. Streptavidin-conjugated magnetic beads were then used to isolate cell-surface proteins from cell lysates. Immunoblotting of total cell lysates and isolated cell-surface proteins revealed that the co-expression of mApple-CA-CaMKII α increased the proportion of mGlu_{5a} expressed on the cell-surface by 3.0 ± 0.7 -fold (SEM) under basal conditions ($p=0.036$; one-sample t-test vs. hypothetical value of 1) (Figure 5). In order to determine whether CaMKII α interaction with the mGlu_{5a} CTD is important for this effect we examined the cell surface expression of mGlu_{5a}-AAA, in which Lys⁸⁶⁶-Arg⁸⁶⁷-Arg⁸⁶⁸ in the CTD were replaced with alanines. In the absence of co-expressed CaMKII, the surface expression of mGlu_{5a}-AAA was not significantly different from those of WT mGlu_{5a} (1.6 ± 0.6 -fold (SEM); $n=5$; $p=0.35$; one-sample t-test vs. hypothetical value of 1). Moreover, the co-expression of mApple-CA-CaMKII α had no effect on cell-surface expression of mGlu_{5a}-AAA. These data demonstrate that interaction with the mGlu_{5a}-CTD is necessary for CaMKII α -mediated increases in mGlu_{5a} cell-surface expression.

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CaMKII α reduces mGlu_{5a}-stimulated peak Ca²⁺ mobilization

To investigate the effect of CaMKII α on mGlu_{5a} signaling, we measured glutamate-induced Ca²⁺ mobilization in populations of 293A-5a^{LOW} cells that stably express mGlu_{5a} and were transiently transfected to co-express mApple or mApple-tagged CaMKII α (either WT or CA). A similar fraction of the total cells expressed detectable levels of mApple-tagged WT- or CA-CaMKII α in each transfection (typically ~60%). After loading glutamate-starved cells with fluo-4-AM, a fluorescent Ca²⁺ indicator, we measured fluorescence responses of total cell populations to increasing glutamate concentrations (0.01-100 μ M) (Figure 6A). An overlay of raw traces from cells expressing mApple, mApple-WT-CaMKII α , or mApple-CA-CaMKII α in a representative experiment is shown in Figure 6B. Peak Ca²⁺ responses (increased fluorescence) at each glutamate concentration were expressed as a ratio to the maximum response to a saturating concentration of glutamate (100 μ M) in mApple-expressing control cells for each individual experiment, and then data were averaged across 5 independent experiments. Glutamate increased the peak fluorescence in a concentration-dependent manner, with an apparent EC₅₀ of 0.38 \pm 0.03 μ M in control cells, similar to previous analyses (Hammond et al., 2010; Schoepp et al., 1999). The glutamate response was unaffected by co-expression of mApple-WT-CaMKII α , but the co-expression of mApple-CA-CaMKII α reduced peak Ca²⁺ responses at the highest concentrations of glutamate by approximately 20%, without affecting the apparent EC₅₀ (Figure 6C). As an alternative measure of Ca²⁺ responses, we determined the area under the curve of the initial Ca²⁺ peak at the highest glutamate concentration. There was no difference in area under the curve between cells expressing mApple or mApple-CaMKII α -WT, but the co-expression of mApple-CA-CaMKII α significantly reduced the area under the curve (Control, 109.2 \pm 2.7; WT,

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101.3±7.4; CA, 82.6±4.1. One-way ANOVA, $p=0.011$, $F=6.280$. Sidak's *post hoc* test for multiplicity adjusted p values: WT vs. control, $p=0.51$, CA vs. control, $p=0.0073$) (data not shown). Since mApple-CA-CaMKII α is expressed in only a fraction of the cell population in each well, the measured reductions in maximal Ca²⁺ responses presumably under-estimate the actual impact of expressing CA-CaMKII α in each cell. However, these data cannot differentiate whether this effect reflects decreased Ca²⁺ mobilization within each cell or a decrease in the fraction of responsive cells. Nevertheless, the data indicate that the co-expression of CA-CaMKII α but not WT-CaMKII α can reduce mGlu_{5a}-stimulated peak Ca²⁺ mobilization.

CaMKII α prolongs mGlu_{5a}-mediated Ca²⁺ signaling

To address caveats associated with studies investigating the effects of mApple-CA-CaMKII α on Ca²⁺ mobilization in 293A-5a^{LOW} cells, we also examined Ca²⁺ mobilization in single HEK293A cells transfected to express full-length mGlu_{5a} with either mApple alone (control) or mApple-CA-CaMKII α . After loading all cells with Fura-2-AM, a ratiometric Ca²⁺ indicator, single cells were selected for analysis based on the presence of mApple as a marker of transfection.

Application of 100 μ M glutamate to cells co-expressing soluble mApple or mApple-CA-CaMKII α with WT mGlu_{5a} produced an initial peak of Fura-2 fluorescence followed by highly variable changes of fluorescence over the next 10 min (Figure 7A). In a majority of cells in each group (53-68%) Ca²⁺ signals waned over time, sometimes with a secondary shoulder, but subpopulations of the cells displayed clear Ca²⁺ oscillations that either returned to baseline between oscillations (10-21%) or were super-imposed on a more sustained Ca²⁺ elevation (18-25%) (Supplemental Figure 1A-C). However, the percentage of WT mGlu_{5a} cells exhibiting Ca²⁺ oscillations was unaffected by the co-expression of mApple-CA-CaMKII α (Supplemental Figure

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1D). Since it is unclear whether oscillating and non-oscillating cells have different physiological effects, we developed an approach to analyze the responses of all cells (both oscillating and non-oscillating) across 5 independent experiments, revealing that the initial peak fluorescence was significantly reduced ($p=0.009$) in cells expressing mApple-CA-CaMKII α vs. cells expressing mApple alone (Figure 7B), consistent with data from stably transfected cell populations (Figure 6). Moreover, the Ca²⁺ signal was relatively prolonged in cells expressing mApple-CA-CaMKII α vs. cells expressing mApple alone, as reflected by a statistically-significant increase in the half-life of the fluorescence signal (Figure 7A inset, 7C). We also analyzed responses in subsets of the cells within each population that exhibited at least three baseline Ca²⁺ oscillations (Supplemental Figure 2A). There was no statistically significant difference in the total number of mGlu_{5a}-mediated Ca²⁺ oscillations between transfection conditions (Supplemental Figure 2B). However, co-expression of mApple-CA-CaMKII α reduced the relative rate of decay of peak Ca²⁺ signals in successive oscillations (Supplemental Figure 2C). Co-expression of mApple-CA-CaMKII α also increased the frequency of Ca²⁺ oscillations, as reflected by a reduction of the inter-event intervals (Supplemental Figure 2D). In combination, these data indicate that CaMKII α can reduce the amplitude of initial mGlu_{5a}-dependent Ca²⁺ mobilization while extending the relative duration of Ca²⁺ signals and increasing the frequency of oscillations when they are present.

To test the hypothesis that CaMKII α binding to the mGlu_{5a}-CTD is necessary for the modulation of Ca²⁺ mobilization, we examined the effect of co-expressing mApple-CA-CaMKII α with mGlu_{5a}-AAA, in which Lys⁸⁶⁶-Arg⁸⁶⁷-Arg⁸⁶⁸ in the CTD were replaced with alanines (Figure 7D). This CTD mutation had little effect on glutamate-stimulated Ca²⁺ mobilization in cells expressing mApple. Moreover, the co-expression of mApple-CA-CaMKII α with mGlu_{5a}-AAA

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had no statistically significant effect on either the initial peak (Figure 7E) or the duration (Figure 7F) of the glutamate-stimulated Ca^{2+} signal relative to control cells expressing mApple alone. Furthermore, the CTD mutation had no statistically significant effect on the responses of cells displaying baseline Ca^{2+} oscillations (Supplementary Figure 1D), but abrogated the CaMKII-dependent modulation, as reflected by a lack of effect on the peak height decay of successive Ca^{2+} oscillations (Supplemental Figure 2C) and the Ca^{2+} oscillation frequency (Supplemental Figure 2D). These data indicate that binding to the mGlu_{5a}-CTD is important for both the increase of initial peak Ca^{2+} signals and for the prolonged Ca^{2+} signaling induced by co-expression of CA-CaMKII α .

Discussion

In a previous report, the membrane proximal region of the mGlu_{5a}-CTD was shown to bind *inactive* CaMKII (Jin et al., 2013b). Here we extend these findings by further characterizing the physical and functional relationship between these key regulators of synaptic transmission. We confirmed that CaMKII α and mGlu₅ specifically interact in mouse brain. However, our data show that mGlu_{5a}-CTD residues 827-964 bind more strongly to CaMKII α in an active, Thr286-autophosphorylated conformation, but that this interaction is disrupted by excess Ca^{2+} /CaM or by robust CaMKII autophosphorylation at additional undefined sites. Furthermore, our data indicate that CaMKII binding to the CTD exerts complex effects on mGlu_{5a} surface expression and downstream Ca^{2+} mobilization.

There is a growing appreciation that specific physiological actions of CaMKII are modulated in part through dynamically-regulated interactions with CaMKII-associated proteins (CaMKAPs). Several CaMKAPs preferentially interact with activated conformations of CaMKII; these

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CaMKAPs can be sub-classified based on differences between the amino acid sequences of their CaMKII-binding domains. CaMKII-binding domains in the NMDA receptor GluN2B subunits and calcium channel $\beta 1$ and $\beta 2$ subunits resemble the CaMKII regulatory domain (Grueter et al., 2008; Strack et al., 2000). In contrast, the amino acid sequence of a CaMKII-binding domain in densin has similarity with a naturally occurring CaMKII inhibitor protein (CaMKIIN) (Jiao 2011). Here, we show here that the binding domain for activated CaMKII in the mGlu_{5a} CTD does not resemble these CaMKAPs. Rather, this novel interaction requires three basic residues (Lys⁸⁶⁶-Arg⁸⁶⁷-Arg⁸⁶⁸), similar to the recently identified interaction of activated CaMKII with the N-terminal domains of L-type voltage-gated Ca²⁺ channels (Wang et al., 2017). Interestingly, triple basic residue motifs can also be identified in CaMKII-binding domains of other GPCRs, including intracellular loops of the G α_i -coupled D₂ and D₃ dopamine receptors (Zhang et al., 2014; Liu et al., 2009), and the CTD of the mGlu₁ receptor (Jin et al., 2013a; Jin et al., 2013b), which also couples to G $\alpha_{q/11}$ (Figure 3). Thus, it will be interesting to investigate the role of these triple basic residue motifs in CaMKII binding to additional GPCRs.

One unusual aspect of CaMKII binding to the mGlu_{5a}-CTD is that, while the *in vitro* interaction requires CaMKII α activation and Thr286 autophosphorylation, additional autophosphorylation at non-Thr286 sites following incubation at 30°C reduces the binding. Our recent proteomics analyses of purified CaMKII α autophosphorylated *in vitro* using a similar 30°C protocol detected 17 autophosphorylation sites, in addition to Thr-286 (Baucum et al., 2015). Presumably the autophosphorylation at one or more of these non-Thr286 sites interferes with *in vitro* CaMKII α binding to mGlu_{5a}. While this is a potentially interesting finding, parallel proteomics analyses of CaMKII isolated from mouse brain failed to detect phosphorylation at many of these *in vitro* sites (Baucum et al., 2015). However, it is possible that this observation explains why Jin and

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colleagues found that autophosphorylated CaMKII did not bind to mGlu_{5a} *in vitro* (Jin et al., 2013b) because their autophosphorylation reactions were incubated at 30°C.

Our data show that CaMKII α activation enhances the association with full-length mGlu_{5a}, and that this interaction involves the Lys⁸⁶⁶-Arg⁸⁶⁷-Arg⁸⁶⁸ motif in the CTD (Figure 3). However, triple alanine substitution of CTD residues 866-868 reduced the interaction by only ~50%, suggesting that CaMKII may interact with additional regions in mGlu_{5a} or bind to the receptor through an indirect interaction. Indeed, a CaMKII interaction with the second intracellular loop of mGlu₅ has been previously reported (Raka et al., 2015), although we have been unable to detect direct binding of purified CaMKII to a GST fusion protein containing the mGlu₅ second intracellular loop (data not shown). Although our data cannot preclude a role for a secondary or indirect interaction, our analyses in heterologous cells indicate that CaMKII α interaction with the CTD is critical for several novel functional effects of CaMKII on mGlu_{5a} signaling. First, we show here that CaMKII α can increase cell surface expression of mGlu_{5a}. Second, we found that CaMKII α has complex effects on mGlu_{5a}-dependent Ca²⁺ mobilization. As noted previously, mGlu₅ activation can induce temporally-diverse intracellular Ca²⁺ responses in heterologous cells and in neurons (Flint et al., 1999; Jong and O'Malley, 2017; Kim et al., 2005; Mao and Wang, 2003b; Uematsu et al., 2015). The co-expression of CaMKII had little effect on the proportion of cells exhibiting different oscillatory or non-oscillatory response patterns (Supplementary Figure 1). However, we found that the co-expression of CA-CaMKII α reduces the amplitude of the initial peak Ca²⁺ signals (Figure 6C, Figure 7B), but prolongs the duration of the Ca²⁺ signals (Figure 7C, Supplementary Figure 2C) in both the total responding cell population or only in cells that exhibit baseline Ca²⁺ oscillations. The co-expression of CA-CaMKII α also increases the frequency of baseline Ca²⁺ oscillations (Supplementary Figure 2D). All of these effects are

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prevented by the triple alanine substitution for Lys⁸⁶⁶-Arg⁸⁶⁷-Arg⁸⁶⁸ in the CTD (Figure 7D-F, Supplementary Figure 2C,D). Presumably, the effect of CaMKII α to increase basal cell-surface expression contributes to the prolongation of Ca²⁺ signaling, but the mechanisms underlying the reduced initial peak Ca²⁺ signal, observed in both stable 293A-5a^{LOW} cell populations and in single transiently-transfected cells, remains unclear. Taken together, our data show that binding of CaMKII α can play an important role in modulating cellular responses to mGlu_{5a} activation. Further examination into the contribution of these mechanisms in synaptic plasticity and neuronal Ca²⁺ signaling are warranted in future studies.

Interestingly, cell surface expression of mGlu_{5a} is also modulated by direct binding of Ca²⁺/CaM to the CTD, similar to the effects of CaMKII α binding to the CTD reported herein, and Ca²⁺/CaM also prolongs mGlu₅-mediated Ca²⁺ signaling (Lee et al., 2008). The Ca²⁺/CaM binding domain involved in mediating these effects is located 30-40 residues C-terminal to the tribasic residue motif that is critical for CaMKII binding. Nevertheless, we found that Ca²⁺/CaM competes for binding of activated CaMKII to the mGlu_{5a}-CTD *in vitro* (Figure 2C). Taken together, our data suggest an intriguing model in which the binding of calmodulin might confer a relatively transient Ca²⁺-dependent modulation of mGlu_{5a} surface expression and signaling, but that increased CaMKII α autophosphorylation at Thr286 would result in sustained binding to the CTD and longer-term modulation of mGlu_{5a} surface expression and Ca²⁺ mobilization. Since Thr286 autophosphorylation of CaMKII is sensitive to changes in the source, duration, or frequency of Ca²⁺ signals originating from multiple channels (Pasek et al., 2015), such as those occurring during synaptic plasticity, as well as to the regulated activities of protein phosphatases, this may provide a mechanism for cross-talk with other signaling pathways.

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As noted above, CaMKII has also been shown to interact with a membrane proximal region in the CTDs of mGlu₁ (Jin et al., 2013a), and the CaMKII-binding domain in mGlu₁ contains a tri-basic residue motif, similar to the motif we have here identified as being critical for CaMKII binding to the mGlu_{5a}-CTD. However, CaMKII was shown to desensitize mGlu₁ signaling whereas we found that CaMKII prolongs mGlu₅ signaling. This apparently differential modulation of mGlu₁ and mGlu₅ by CaMKII may contribute to their distinct neuronal roles (Mannaioni et al., 2001; Valenti et al., 2002; Volk et al., 2006). Interestingly, the effects of CaMKII on mGlu₁ signaling are mediated in part by phosphorylation at Thr871, which lies within the CaMKII-binding domain. Therefore, it will be interesting to investigate whether phosphorylation is required for the effects of CaMKII on mGlu₅ signaling, as well as the physical interaction demonstrated here.

The effects of CaMKII α on mGlu_{5a} must also interface with the known modulation of mGlu₅ signaling by other mechanisms. Prior studies have shown that several protein kinases modulate mGlu₅ via the CTD. For example, PKC phosphorylates Ser901 in the mGlu_{5a} CTD to inhibit Ca²⁺/CaM binding and antagonize the aforementioned modulation by Ca²⁺/CaM (Lee et al., 2008). In addition, PKA phosphorylates Ser870 in mGlu_{5a}, prolonging Ca²⁺ mobilization, similar to the effects of CaMKII reported here, and enhancing ERK activation (Uematsu et al., 2015). However, it was previously reported that CaMKII reduces mGlu₅-stimulated ERK1/2 activation and increases agonist-induced mGlu₅ internalization (Raka et al., 2015). It is possible that the enhanced agonist-induced internalization in part results from the increased basal surface expression reported herein (Figure 5). Although the mechanistic relationships between these different modes of mGlu₅ regulation remain to be more clearly established, the convergence of Ca²⁺/CaM, CaMKII, PKA and PKC actions within an ~60 amino acid region in the long CTD

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(345 amino acids) suggests that the actions of mGlu_s are tightly controlled across different time frames, presumably fine-tuning neuronal responses such as different forms of synaptic plasticity.

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Participated in research design: Marks, Shonesy, Wang, Niswender, Colbran

Conducted experiments: Marks

Contributed new reagents or analytic tools: Marks, Wang, Stephenson, Niswender

Performed data analysis: Marks, Shonesy

Wrote or contributed to the writing of the manuscript: Marks, Shonesy, Colbran

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Legends for Figures:

Figure 1. *Co-immunoprecipitation of mGlu5 with mouse forebrain CaMKII α* . Solubilized fractions from WT or CaMKII α -KO mouse forebrain were immunoprecipitated using CaMKII α -specific (α) or control (IgG) antibodies, as indicated. Inputs and immune complexes were analyzed by immunoblotting: mGlu5 was detected only in immune complexes isolated from WT tissue using the CaMKII α antibody. Open and closed arrowheads indicate dimeric and monomeric species of mGlu5. Representative of 3 similar experiments.

Figure 2. *CaMKII autophosphorylation at Thr286 enhances binding to the mGlu5 C-terminal domain*. **A. Autophosphorylation of purified CaMKII α** . Purified CaMKII α was incubated with $Mg(C_2H_3O_2)_2$, $CaCl_2$, CaM, and ATP for either 90 s at 4°C or 10 min at 30°C and samples were immunoblotted for total or phospho-Thr286 CaMKII. While quantitative analysis (right) indicated that there was a similarly robust Thr286 autophosphorylation using these two conditions, the 10 min/30°C incubation resulted in a substantial reduction in electrophoretic mobility due to phosphorylation at additional unidentified sites. Data are plotted as the mean \pm SEM (n=3) and analyzed using a one-way ANOVA (p=0.0167, F=8.746, R square = 0.7446) with Sidak's *post-hoc* test for multiplicity adjusted *p* values: Control vs. 4°C, p=0.031. Control vs. 30°C, p=0.035. 4°C vs. 30°C, p=1.00 **B. The GST-mGlu5 α -CTD binds CaMKII α following selective autophosphorylation at Thr286**. GST-mGlu5 α -CTD was incubated with purified CaMKII α that had been preincubated as in panel A, and complexes were isolated using glutathione agarose. Immunoblot analyses revealed that CaMKII α binding to the CTD was strongly enhanced by selective Thr286 autophosphorylation at 4°C, but that the autophosphorylation of additional sites on CaMKII at 30°C substantially reduced binding. Data are plotted as the mean \pm SEM (n=3) and

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were analyzed using a one-way ANOVA ($p=0.005$, $F=2.477$, R square = 0.829) with Sidak's *post-hoc* test for multiplicity adjusted p values: Control vs. 4°C, $p=0.009$. 4°C vs. 30°C, $p=0.011$. Control vs. 30°C, $p=1.00$. **C. Binding of activated CaMKII α to GST-mGlu_{5a}-CTD is disrupted by Ca²⁺/CaM.** Purified CaMKII α was autophosphorylated for 90 s at 4°C (see Panel A) and then incubated with GST-mGlu_{5a}-CTD in the absence or presence of excess Ca²⁺/CaM (see Methods). Complexes were isolated using glutathione-agarose and then immunoblotted as indicated. Data are plotted as the mean \pm SEM; excess Ca²⁺/CaM significantly reduced CaMKII α binding ($p < 0.0001$ relative to theoretical value of 1.00 by one-sample t-test; $n=4$).

Figure 3. Identification of CaMKII-binding determinants in the mGlu_{5a}-CTD. **A. Alignment of part of the mGlu_{5a}-CTD with amino acid sequences surrounding known CaMKII-binding domains.** Tribasic residue motifs (highlighted) were identified within CaMKII-binding domains from other proteins, as well as within the CaMKII binding fragment in the CTD of mGlu_{5a}. Mutation of R⁸³K⁸⁴R⁸⁵ to AAA in the Cav1.3 N-terminal domain (NTD) disrupts the binding of CaMKII (Wang et al., 2017). The red and blue fonts indicate residues in each domain that are identical and homologous, respectively, with residues in the mGlu_{5a} sequence. Underlined residues in the mGlu₁-CTD and the D₂ dopamine receptor (IL3: third intracellular loop) demark the sequences of synthetic peptides that were shown to compete for CaMKII binding (Jin et al., 2013a; Zhang et al., 2014). **B. Mutation of the tribasic residue motif in the mGlu_{5a} CTD disrupts CaMKII binding.** Thr286 autophosphorylated CaMKII α (90 s/4°C protocol) was incubated with GST-mGlu_{5a}-CTD (WT or with a K⁸⁶⁶R⁸⁶⁷R⁸⁶⁸ to AAA mutation) and complexes were analyzed as in Fig. 1. The K⁸⁶⁶R⁸⁶⁷R⁸⁶⁸/AAA mutation essentially abolishes CaMKII binding Data are plotted as the mean \pm SEM ($p = 0.003$ by a one sample t-test; $n=4$).

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Figure 4. Role of the CTD in full-length mGlu_{5a} binding to activated CaMKII α . **A.** *CaMKII activation enhances interaction with full length mGlu₅.* Solubilized fractions of HEK293A cells expressing HA-tagged mGlu_{5a} and/or mApple-tagged WT CaMKII α (as indicated above lanes) were pre-incubated with Ca²⁺/CaM, MgAc₂, and ATP in the presence or absence of excess EDTA (–/+ activation, respectively) and then immunoprecipitated using antibodies to the HA epitope. Lysates and immune complexes were analyzed by immunoblotting, as indicated. CaMKII α activation results in robust Thr-286 autophosphorylation, which increases CaMKII α association with HA-mGlu_{5a}. Data are plotted as the mean \pm SEM (p=0.043; one-sample t-test; n=4). Open and closed arrowheads indicate dimeric and monomeric species of mGlu_{5a}. **B.** *CaMKII association with full-length mGlu_{5a} is disrupted by mutation of the CTD tribasic residue motif.* Solubilized fractions of HEK293A cells expressing HA-mGlu₅ (WT or with the K⁸⁶⁶R⁸⁶⁷R⁸⁶⁸/AAA mutation) and mApple-tagged CA-CaMKII α were immunoprecipitated using antibodies to the HA epitope. Lysates and the immune complexes were analyzed by immunoblotting, as indicated. The K⁸⁶⁶R⁸⁶⁷R⁸⁶⁸/AAA mutation reduced the association of CA-CaMKII α with HA-mGlu₅. Data are plotted as the mean \pm SEM (p=0.028; one sample t-test; n=4).

Figure 5. CaMKII enhances the cell-surface expression of mGlu_{5a} via interaction with the CTD. Cell surface biotinylation analyses of HEK293A cells expressing mGlu_{5a} (WT or with K⁸⁶⁶R⁸⁶⁷R⁸⁶⁸/AAA mutation in the CTD) with either mApple or mApple-tagged CA-CaMKII α . The co-expression of CA-CaMKII α increased steady-state surface expression levels of WT mGlu_{5a} (p=0.036; one sample t-test; n=6), but not of the K⁸⁶⁶R⁸⁶⁷R⁸⁶⁸/AAA mutant. Data are plotted as the mean \pm SEM (p = 0.569; one sample t-test; n=5).

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Figure 6. CaMKII α regulates mGlu_{5a}-stimulated Ca²⁺ mobilization in 293A-5a^{LOW} cells. Time courses of intracellular Ca²⁺ responses to glutamate were measured by changes in Fluo-4 fluorescence in stable 293A-5a^{LOW} cells in 96 well plates. **A.** Time courses of Ca²⁺ responses. Example of calcium responses to increasing glutamate concentrations collected in a row of 8-wells. **B.** Overlay of individual Ca²⁺ responses to increasing concentrations of glutamate (labeled by colors in Panel A) from 293A-5a^{LOW} cells transiently transfected to express mApple control, mApple-CaMKII α -WT or mApple-CA-CaMKII α from a representative experiment. **C.** Concentration response curves. Initial peak Ca²⁺ responses ($\Delta F/F_0$) at each concentration were normalized to the maximal glutamate-stimulated response in control (mApple-transfected) cells within each experiment. Normalized Ca²⁺ responses are plotted as the mean \pm SEM (n=5 experiments) as a function of glutamate concentration. The expression of CaMKII α -WT had no impact on the Ca²⁺ response curve but the expression of CA-CaMKII α reduced peak Ca²⁺ responses (multiple comparisons two-way ANOVA: Sources of Variation: CaMKII p<0.0001, Interaction p=0.029. Tukey's *post hoc* test for multiplicity adjusted *p* values: mApple vs. WT, *p* = 0.926, mApple vs. CA-CaMKII α , *p* <0.0001. WT vs. CA-CaMKII α , *p*=0.0002). The inset table shows the maximum response (Max), EC₅₀ (μ M), and Hill coefficient (\pm SEM) obtained by fitting the data in GraphPad Prism.

Figure 7. CaMKII α binding to the CTD is required for modulation mGlu_{5a}-stimulated Ca²⁺ mobilization. HEK293A cells were transiently transfected to express mGlu_{5a} (WT or K⁸⁶⁶RR⁸⁶⁸/AAA) with either mApple or mApple-CA-CaMKII α for single-cell Fura-2 Ca²⁺

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imaging (see Methods). **A, D.** *Representative data from a single experiment.* Averaged normalized changes in fluorescence from 58-114 cells ($\Delta F/F_0$: mean \pm SEM) expressing mGlu_{5a}-WT (**A**) or mGlu₅-K⁸⁶⁶R⁸⁶⁷R⁸⁶⁸/AAA (**D**) in the presence (blue lines) or absence (red lines) of mApple-CA-CaMKII α . The inset graphs show line-fits for time courses of the decline of Ca²⁺ signals from the peak $\Delta F/F_0$ under each condition. **B, C, E, F.** *Summary data.* The bar graphs depict mean \pm SEM values for peak Ca²⁺ signals ($\Delta F/F_0$) (**B, E**) and half-lives for the decline in Ca²⁺ signals (**C, F**) with super-imposed data points from each experiment (n=5). Expression of constitutively-active mApple-CA-CaMKII decreases the peak Ca²⁺ signal but increases the half-life of the Ca²⁺ signal with mGlu_{5a}-WT (**B**, p=0.009. **C**, p=0.001), but has no significant effect on the mGlu_{5a}-K⁸⁶⁶R⁸⁶⁷R⁸⁶⁸/AAA mutant that disrupts CaMKII binding to the CTD (**E**, p=0.155. **F**, p=0.415). Paired Student's t-tests were used for statistical comparisons in each panel.

Figure 1

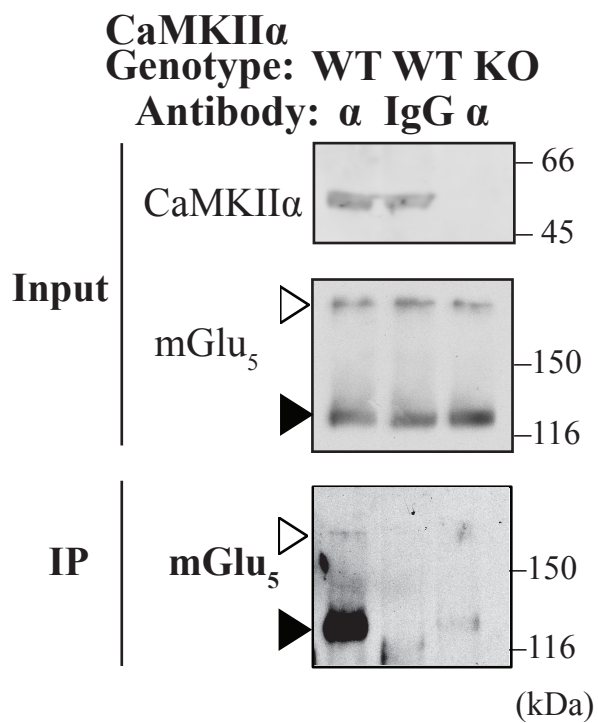
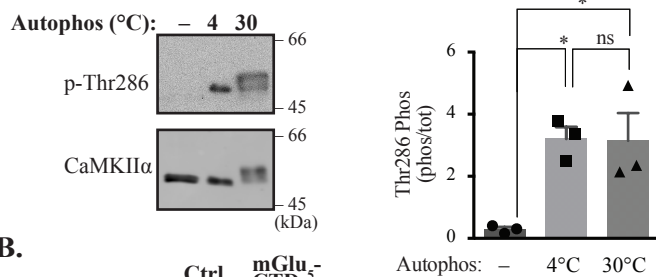
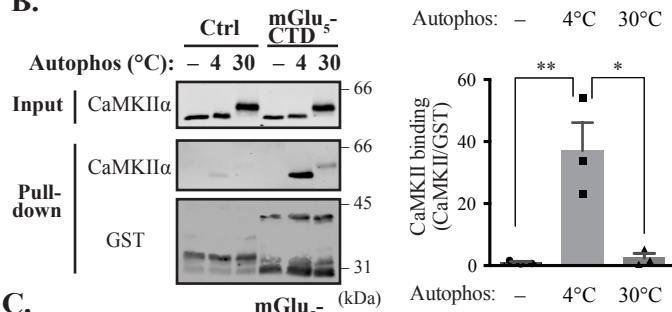


Figure 2

A.



B.



C.

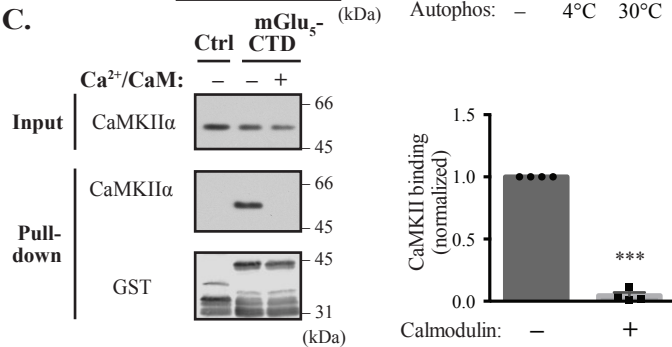


Figure 3

A.

mGlu ₅ -CTD	847	GDGKSSSAASRSSLVNLWKR	RGSSGETLSSNGKSV	882	
mGlu ₁ -CTD	861	GDGK--LPCRSNTFLNIFRR	KPGAGNANSNGKSV	891	
Ca _v 1.2-NTD	67	SAGNATIST-VSSTQRKR	QQYKPKKQGSTTA	97	
Ca _v 1.3-NTD	68	TM-STSAAPPVGSLSQRKR	QQYAKSKKQGNSSN	99	
D ₃ R-IL3	203	VTVLVYARIY-IVLRQRKR	ILLTRQNSQCISIRP	234	
D ₂ R-IL3	345	TMPNGKTRTS-LKTMSRRKLS	QQEKKATQMLA	376	

B.

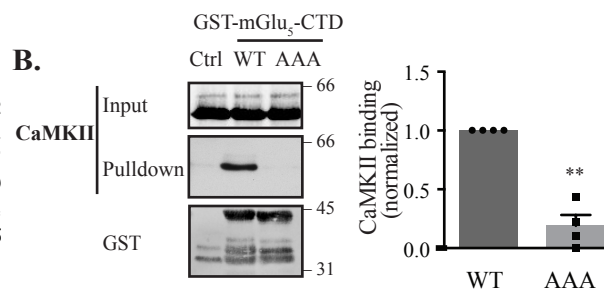
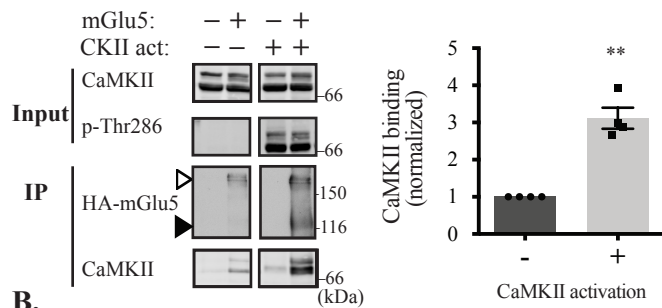


Figure 4

A. IP: HA



B.

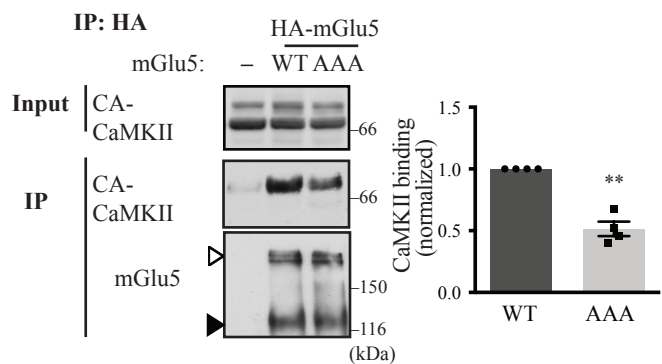


Figure 5

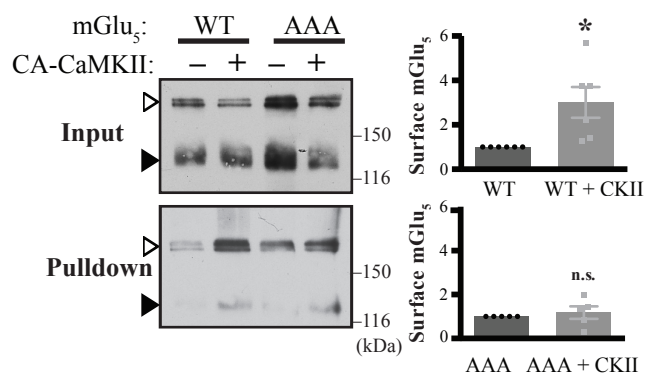


Figure 6

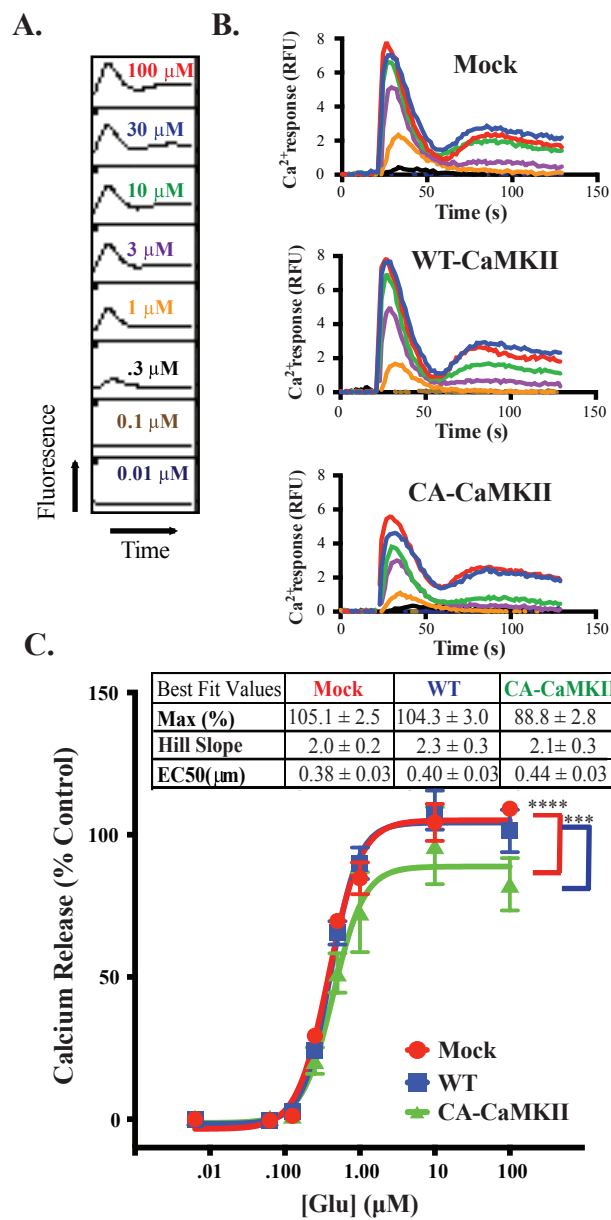
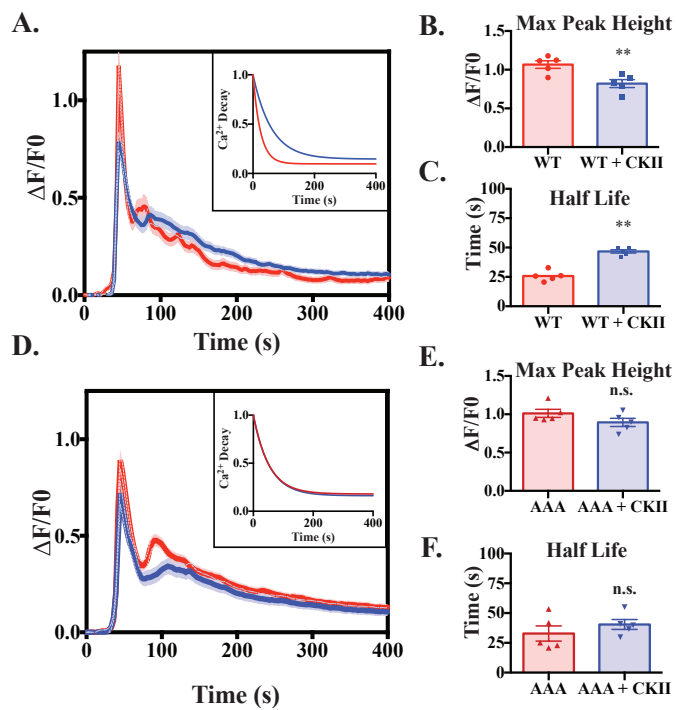


Figure 7



Supplementary Figures 1 and 2

Activated CaMKII α binds to the mGlu₅ metabotropic glutamate receptor and modulates calcium mobilization

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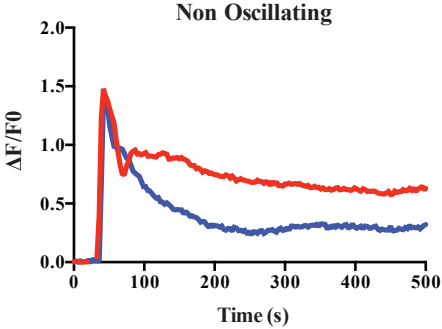
Supplemental Figure 1. Analysis of the variability of glutamate-induced mGlu5 Ca^{2+} responses in transfected HEK293A cells (from data summarized in Fig. 7). Responses of individual cells could be divided into two main categories: **A. *Non-oscillators***: An initial peak of Ca^{2+} that is sustained or returns to baseline over time (blue), in some cases with second shoulder (red), with no clear oscillations. **B. *Oscillating cells***: The initial Ca^{2+} peak decays in an oscillatory pattern (up to ~25 oscillations in 10 min) that may (blue) or may not (red) return to baseline between successive oscillations. **C. *Distribution of responding cells between non-oscillating or oscillating categories for each transfection condition***. The ratio of non-oscillating to oscillating HEK293A cells was not affected by co-expression of mApple-CA-CaMKII α with WT mGlu5a (p=0.28 by Fisher's exact test) or by mutation of Lys⁸⁶⁶-Arg⁸⁶⁷-Arg⁸⁶⁸ in the CTD to alanines in mGlu5a-AAA (p=0.62).

Supplementary Figure 2. Analyses of Ca^{2+} responses in baseline-oscillating HEK293A cells (blue trace in Fig. S1B) expressing mGlu5-WT or mGlu5-AAA with either mApple or mApple-CA-CaMKII. **A.** Overlays of ten representative Ca^{2+} responses for each transfection condition from cells selected for ≥ 3 baseline oscillations. **B.** There were no statistically significant differences in the total number of oscillations recorded over 10 min between transfection conditions (total numbers of baseline-oscillating cells: mGlu5-WT, 78; mGlu5-WT+CKII, 62; mGlu5-AAA, 38; mGlu5-AAA+CKII, 21. One-way ANOVA: p = 0.08). The error bars depict the range between the minimum (3) and maximum number of oscillations, boxes indicate the 25-75th percentile, lines within each box indicate the median, and the "+" sign within each box indicates the mean. **C.** Peak responses for the first 5 oscillations were normalized to the first Ca^{2+} peak and plotted as the mean \pm SEM for each transfection condition. Co-expression of mApple-CA-CaMKII

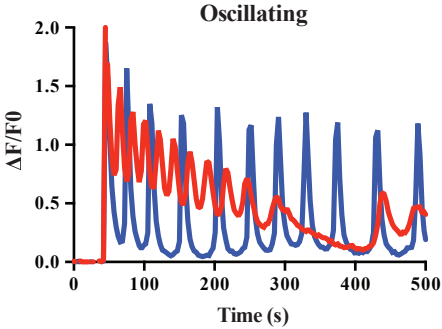
significantly slows the rate of decay of successive peak Ca^{2+} responses in cells expressing mGlu_{5a}-WT, but has no effect in cells expressing mGlu_{5a}-AAA (2-way repeated measures ANOVA. CaMKII effect, $p < 0.0001$, Sidak's test for multiplicity adjusted p values: mGlu_{5a} vs mGlu_{5a}-WT+CaMKII $p = 0.0003$, mGlu_{5a} vs. mGlu_{5a}-AAA $p > 0.999$, mGlu_{5a}-AAA vs mGlu_{5a}-AAA + CaMKII $p > 0.999$). **D.** Cumulative probability curves for mean inter-event intervals in baseline-oscillating cells. Co-expression of mApple-CA-CaMKII significantly decreases inter-event intervals between Ca^{2+} -oscillations in cells expressing mGlu_{5a}-WT but not mGlu_{5a}-AAA (Kruskal Wallis test $p = 0.006$, Dunn's test for multiplicity adjusted p values: WT-mGlu_{5a} vs WT-mGlu_{5a} + CaMKII $p = 0.003$, WT-mGlu_{5a} vs mGlu_{5a}-AAA $p > 0.999$, mGlu_{5a}-AAA vs mGlu_{5a}-AAA + CKII $p > 0.999$).

Supplemental Figure 1

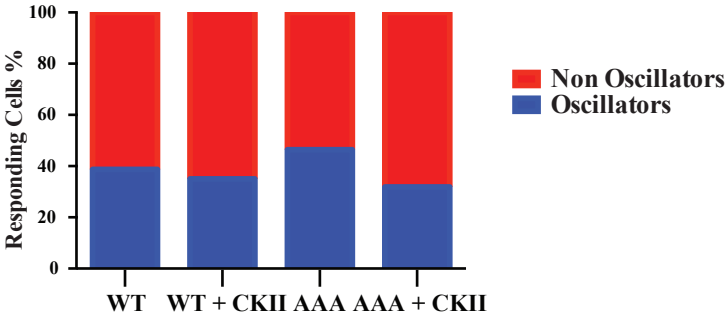
A.



B.

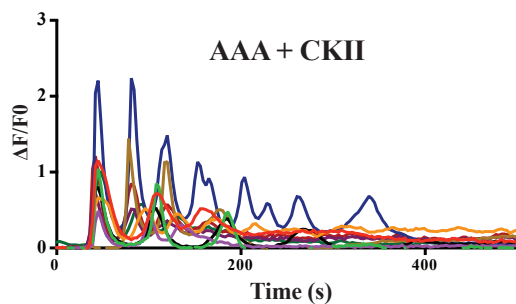
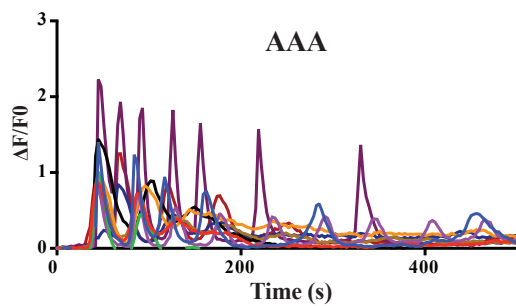
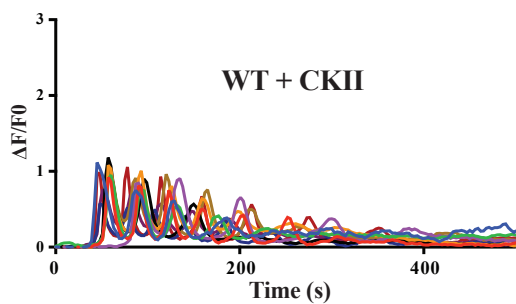
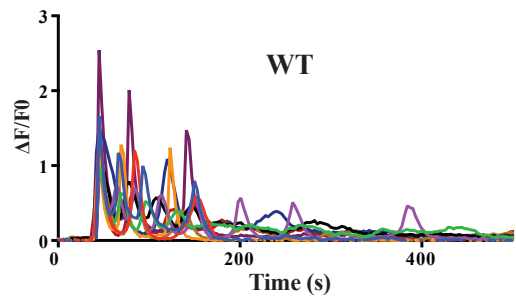


C.

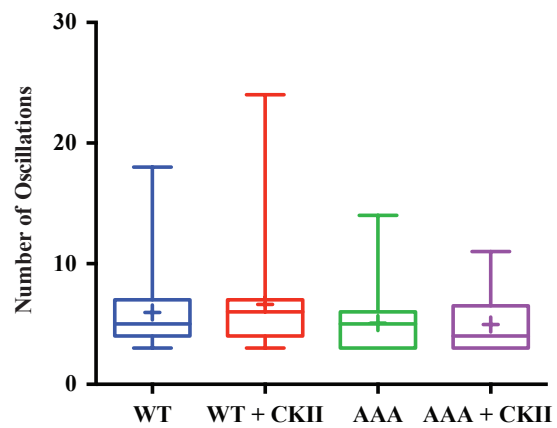


Supplemental Figure 2

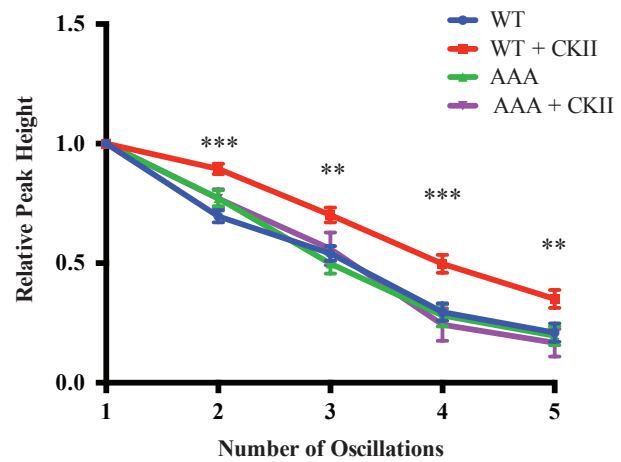
A.



B.



C.



D.

