Activated Nuclear Metabotropic Glutamate Receptor mGlu5 Couples to Nuclear Gq/11 Proteins to Generate Inositol 1,4,5-Trisphosphate-Mediated Nuclear Ca²⁺ Release *

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Running Title: Nuclear mGlu5 Couples to Gq/11/PI-PLC/IP3 Pathway

Recently we have shown that the metabotropic glutamate 5 (mGlu5) receptor can be expressed on nuclear membranes of heterologous cells or endogenously on striatal neurons where it can mediate nuclear Ca²⁺ changes. Here, pharmacological, optical, and genetic techniques were used to show that upon activation, nuclear mGlu5 receptors generate nuclear inositol 1,4,5-trisphosphate (IP3) in situ. Specifically, expression of an mGlu5 F767S mutant in HEK293 cells which blocks Gq/11 coupling or introduction of a dominant negative Gα₉ construct in striatal neurons prevented nuclear Ca²⁺ changes following receptor activation. These data indicate that nuclear mGlu5 receptors couple to Gq/11 to mobilize nuclear Ca²⁺. Nuclear mGlu5-mediated Ca²⁺ responses could also be blocked by the phospholipase C (PLC) inhibitor, U73122, the phosphatidylinositol (PI) PLC inhibitor 1-O-octadecyl-2-O-methyl-sn-glycero-3-phosphorylcholine (ET-18-OCH₃), or by using siRNA targeted against PLCβ1 demonstrating that PI-PLC is involved. Direct assessment of inositol phosphate production using a PIP2/IP3 'biosensor' revealed for the first time that IP3 can be generated in the nucleus following activation of nuclear mGlu5 receptors. Finally, both IP3 and ryanodine receptor blockers prevented nuclear mGlu5-mediated increases in intranuclear Ca²⁺. Collectively, this study shows that like plasma membrane receptors, activated nuclear mGlu5 receptors couple to Gq/11 and PLC to generate IP3-mediated release of Ca²⁺ from Ca²⁺-release channels in the nucleus. Thus the nucleus can function as an autonomous organelle independent of signals originating in the cytoplasm and that nuclear mGlu5 receptors play a dynamic role in mobilizing Ca²⁺ in a specific, localized fashion.

Many cells produce Ca²⁺ signals both in the cytoplasm as well as the nucleus. Nuclear Ca²⁺ is thought to play a vital role in a variety of nuclear functions such as cell division, proliferation, protein import, apoptosis and gene transcription (1). Nuclear Ca²⁺ may be mobilized from a number of sources including diffusion of cytosolic Ca²⁺ waves through nuclear pore complexes (2), release into the nucleoplasm from the nuclear lumen (3,4), or release from the so-called nucleoplasmic reticulum, invaginations of the nuclear envelope into the nucleoplasm itself (5). Presumably Ca²⁺ release from the lumen of the nuclear membrane and/or the nucleoplasmic reticulum would either amplify Ca²⁺ signals arriving via the nuclear pore complex and/or independently generate nucleoplasmic Ca²⁺ transients. Such a model of nuclear Ca²⁺ release is bolstered by data documenting the presence and activation of the inositol 1,4,5-trisphosphate receptors (IP3Rs1) and ryanodine receptors (RyRs) on inner nuclear membranes and the nucleoplasmic reticulum (3,4,6-9). Thus these Ca²⁺-release channels are perfectly poised to independently regulate nucleoplasmic Ca²⁺ levels.

Despite the importance of IP₃Rs and RyRs in controlling the release of nuclear Ca²⁺, very little information is available regarding how these receptors themselves are activated. One of the most prevalent models is one in which IP₃ produced in the cytoplasm diffuses into the nucleus thereby activating nuclear IP₃Rs (10). This premise is supported by studies showing that extracellular insulin-like growth factor-1 (IGF-1) activates its own receptor as well as phospholipase C (PLC) to stimulate IP₃ mobilization and the ensuing activation of both cytoplasmic and nuclear IP₃Rs (11,12). Alternatively, IP₃ might be synthesized within the nucleus itself. This hypothesis is consistent with reports showing that the nucleus contains all of the enzymatic machinery necessary to synthesize IP₃ (13). Moreover, the nucleus also has the components necessary for the production of the RyR agonists, cyclic ADP-ribose (cADPr; 14) or nicotinic acid adenine dinucleotide phosphate (NAADP; 7). Thus, the nucleus could potentially function as an
independent signaling unit controlling Ca\textsuperscript{2+} signals in a specific, localized fashion.

How might nuclear IP\textsubscript{3} be generated? Typically, cell surface G-protein coupled receptors (GPCRs) associated with G\textsubscript{q}-like proteins (G\textsubscript{q} and its close homolog, G\textsubscript{11}; 15) activate phospholipases such as PLC to hydrolyze phosphatidylinositol 4,5-bisphosphate (PIP\textsubscript{2}) leading to IP\textsubscript{3} production. GPCRs coupling with G\textsubscript{i/o}-like proteins can also increase IP\textsubscript{3} levels via dissociated \(\beta\)/\(\gamma\) subunit activation of PLC (16,17). These same mechanisms might also exist in the nucleus where several GPCRs have been recently localized not only to the plasma membrane but also intracellularly on nuclear membranes (18-20). For example, receptors known to couple to G\textsubscript{q/11} such as the prostaglandin EP receptor EP\textsubscript{1}, as well as the platelet-activating factor (PAF) and lysophosphatidic acid (LPA-1) receptors have been found on the nuclear envelope (21). Similarly, we have shown that the Group 1 metabotropic glutamate receptors, mGlu1 and mGlu5, both of which couple to G\textsubscript{q/11}, are endogenously expressed on the inner nuclear membranes of many different types of neurons (18,19,22). G\textsubscript{q} itself is present on nuclear membranes (23). Moreover, we have previously shown that a GFP-tagged G\textsubscript{q} protein co-localizes with mGlu5 and the inner nuclear membrane protein, lamin B\textsubscript{2}, in heterologous cell types (18). Thus, signaling systems are in place that could generate the requisite second messengers within the nucleoplasm itself.

Surprisingly, few studies have directly examined signal transduction pathways associated with nuclear GPCRs. Given the importance of nuclear Ca\textsuperscript{2+} signaling and the unlikelihood, at least in neurons, that IP\textsubscript{3} could diffuse long distances from its presumed site of synthesis in neuronal processes, it seems reasonable to propose that IP\textsubscript{3} is generated \textit{in situ} via the actions of GPCRs located on the inner nuclear membrane. Here we show that, like plasma membrane receptors, activated nuclear mGlu5 receptors couple to G\textsubscript{q/11} and phosphatidylinositol (PI)-PLC in mGlu5-expressing HEK cells as well as striatal nuclei to generate IP\textsubscript{3}-mediated release of Ca\textsuperscript{2+} via Ca\textsuperscript{2+} release channels in the nucleus. Taken together, these data point to a novel mode of nuclear Ca\textsuperscript{2+} generation, independent of cytosolic Ca\textsuperscript{2+}, mediated through activated nuclear GPCRs.

**EXPERIMENTAL PROCEDURES**

**Materials**—Quisqualate, 2-methyl-6-(phenylethynyl)-pyridine (MPEP), 7-hydroxyiminocyclopropen[b]chromen-1a-carboxylic acid ethyl ester (CPCCOEt), [1-(6-[17 beta-3-methoxyestra-1,3,5-(10) triene-17-y]) amino/hexyl] 1H-pyrrole-2,5-dione (U73122), O-(Octahydro-4,7-methano-1H-inden-5-y) carbonopotasium dithioate (D609), and 2-Aminoethoxydiphenylborane (2-APB) were purchased from Tocris Cookson Inc., Ellisville, MO. Phosphatidylinositol3-kinase (PI3K) inhibitor LY294002 was from Cell Signaling Technology, Inc., Beverly, MA, [1-(4-aminophenyl)-3-methylcarbamyl-4-methyl-3,4-dihydro-7,8-methyleneedioxy-5H-2,3-benzodiazepine] (GYKI53655), was from Research Biochemicals Inc., Natick, MA and myo-[2-\textsuperscript{3}H(N)]inositol (specific activity = 20.0 Ci/mmol) was from American Radiolabeled Chemicals, Inc., Saint Louis, MO. Optiprep\textsuperscript{TM} (60% Iodixanol) was purchased from Accurate Chemical and Scientific Corporation, Westbury, NY, and Xestospongin C was from Calbiochem, San Diego, CA. Unless otherwise indicated all other chemicals were from Sigma, Saint Louis, MO.

**Cell Culture and Plasmids**—HEK293 cells and the mGlu5 stable HEK cell line were maintained as described (18,24). Primary striatal cultures were prepared and maintained exactly as described (19). The mGlu5 mutant (F767S) was generated by recombinant polymerase chain reaction (25) using sense and antisense primers containing the relevant phenylalanine to serine mutation and wild type mGlu5 construct (24) as template. The mutation was confirmed by sequencing. The mGlu5 mutant F767S stable HEK cell line was generated using standard transfection techniques followed by repetitive rounds of limiting dilution (24). The construct containing the pleckstrin homology domain of PLC\textsubscript{δ1} fused to Enhanced Green Fluorescent Protein (pEGFP-C1-PLC\textsubscript{δ1}-PH), was a gift from Dr. T. Meyer, Stanford University, Stanford, CA (26). pDsRed2 encoding DsRed2, a red fluorescent protein, was obtained from Clontech Laboratories, Inc., Mountain View, CA and subcloned into the pcDNA3 vector. The dominant negative G\textsubscript{αq} construct (pcDNA3.1+G\textsubscript{αq} containing mutations Q209L/D277N) was obtained from University of
RNA Interference- On-Target plus siRNA duplex targeting PLCβ1 (5'-CAGUUGAAUUGUCGAAUAUU -3' and 5'-PUAUUCGACAAUUCAACUGU -3') and On-Target plus scrambled siControl non-targeting siRNA #1 (sequence of the sense strand 5'-UGGUUUACAUGUCGACUAA -3') were purchased from Dharmacon, Inc., Chicago, IL. Neurons were transfected with either the PLCβ1 siRNA or scrambled siRNA or co-transfected along with DsRed2 construct as a transfection marker (2 µg siRNA + 200 ng pcDNA3-DsRed2 construct per 35-mm dishes) using Lipofectamine 2000 (Invitrogen, Carlsbad, CA) as described (28). The efficacy of knockdown was confirmed by immunocytochemistry and western blot analysis 48 and 72 h post-transfection.

Subcellular Fractionation and Preparation of Nuclei- Plasma membrane and nuclear fractions were prepared from HEK cells as described (18). Nuclei were prepared from postnatal day 10 (P10) rat striata as described (19) and further purified using a 25%-30%-35% (w/v) iodixanol gradient as per manufacturer’s instructions. Aliquots from each fraction were used for gel electrophoresis and membrane binding. Protein concentrations of each fraction were determined using the Bradford assay (Bio-Rad, Richmond, CA). Purity of nuclei was assessed by loading HEK cells or striatal neurons with the cytoplasmic mitochondrial-selective stain MitoTracker Deep Red 633 as well as the DNA specific vital dye, HOECHST 33258 as described (29). After a 20 min pre-incubation, subcellular fractionation was performed and nuclear staining was monitored.

Immunocytochemistry and Western Blot Analysis- Cells for immunocytochemical and confocal microscopic studies were grown on poly-D-lysine-treated glass coverslips, 8-well chamber slides, and/or dishes with 35-mm glass grids (MatTek, Ashland, MA). Fixation, blocking, and antibody incubation were as described (24). Primary antibodies included affinity-purified anti-C-terminal mGlu5 (1:250; 30), rabbit polyclonal anti-mGlu5 (1:200; Upstate/Millipore, Charottesville, VA), and monoclonal anti-lamin B2 (1:100; Zymed Laboratories, San Francisco, CA). Anti-PLCβ1 (1:300 of mouse monoclonal, K92) was a gift from Dr. Pann-Ghill Suh (Pohang University of Science and Technology, South Korea). Secondary antibodies were as described (18,19,24). Proteins obtained from subcellular fractionation were resuspended in lysis buffer (150 mM NaCl, 1 mM EDTA, 0.1% SDS, 1% NP40, 0.5% sodium deoxycholate, 50 mM TrisHCl pH 7.5 and protease inhibitors Complete Tablets; Roche Applied Science, Indianapolis, IN). Lysates were subjected to SDS-PAGE, blotted as described (19) and probed with anti-mGlu5 (1:1000), anti-lamin B2 (1:2000), monoclonal anti-Na/K ATPase (α-6F, 1:1000; Development Studies Hybridoma Bank, University of Iowa, Iowa City), monoclonal anti-PLCβ1 (1:200, D-8: sc-5291; Santa Cruz Biotechnology, Inc., Santa Cruz, CA) and monoclonal anti-β-actin (1:5000; Sigma). Horseradish peroxidase conjugated with goat anti-rabbit IgG (1:2000; Cell Signaling Technology, Inc.) or anti-mouse IgG (1:2000; Sigma) was used in conjunction with enhanced chemiluminescence (Amersham Biosciences, Piscataway, NJ) to detect the signal. Densitometric analyses were performed using the Molecular Dynamic’s Storm 860 Imager (Amersham Biosciences) together with associated software.

[^H]Quisqualate Binding and Uptake-[^H]Quisqualate (specific activity = 26.0 Ci/mmol) was made by SRI International. Nuclear and plasma membrane fractions were used in binding assays as described (24).[^H]Quisqualate uptake was performed as described (19).

Fluorescent Measurements of Ca^{2+} in Intact Cells and Isolated Nuclei- For whole cell measurements, mGlu5 wild type or mutant expressing HEK cells were washed with serum-free medium (SFM), incubated with Oregon Green 488 BAPTA-1 AM (Molecular Probes) and imaged as described (18). Similarly, primary cultured striatal neurons transiently transfected with DsRed2 or co-transfected with DNGαq/DsRed2, scrambled siRNA/DsRed2 or PLCβ1 siRNA/DsRed2 (10:1 ratio in all co-transfections) were prepared as described (19). To measure Ca^{2+} changes in individual nuclei, nuclei from HEK cells or rat P10 striatal tissues were prepared and processed as described (18,19). Drugs at 100x concentration were added to the side of the dish and allowed to diffuse at room temperature. Extra- and intracellular buffers used...
for Ca\textsuperscript{2+} measurements on intact cells or isolated nuclei were as described (19). Following image collection, cells/nuclei were fixed, stained with anti-mGlu5, lamin B\textsubscript{2} and/or PLC\textbeta{}1, and field relocated.

Single Cell and Nuclear Imaging of IP\textsubscript{3}—Dissociated striatal neurons were transiently transfected with pEGFP-C1-PLC\textdelta{}1-PH (2 µg 35-mm dishes\textsuperscript{−1}). After 24-48 h cells were washed with Neurobasal medium (NB; Invitrogen) and used for monitoring IP\textsubscript{3} generation in real time. Nuclei prepared from mGlu5/HEK cells transiently transfected with pEGFP-C1-PLC\textdelta{}1-PH, were imaged as described (18,19). To monitor IP\textsubscript{3} generation and Ca\textsuperscript{2+} changes simultaneously, nuclei were loaded with Calcium Crimson AM (Molecular Probes).

Confocal Microscopy and Data Analysis—All Ca\textsuperscript{2+} measurements and IP\textsubscript{3} imaging were done using laser scanning confocal microscopes FluoView 500 and/or FluoView 1000 (Olympus, Center Valley, PA) as described previously (18,19). Simultaneous IP\textsubscript{3} and Ca\textsuperscript{2+} measurements were done using bandpass barrier filter settings at 483-536 nm for EGFP and 590-680 nm for Calcium Crimson (31). Analysis of changes in GFP fluorescence in striatal neurons or HEK nuclei, in real time, was performed as described (32). Increases in IP\textsubscript{3} were detected by measuring the translocation of EGFP from the plasma membrane to the cytosol or from nuclear membrane to the nucleoplasm. Images were processed with MetaMorph (version 5.0.7) Professional Image Analysis software. Data are presented as ratio of change in fluorescent intensity at a given time to initial fluorescence and expressed as percentage (ΔF/F\textsubscript{o}, %).

\[ ^3\text{H}-\text{IP} \] Measurements—IP was measured in cells expressing mGlu5 as described (33,34). For measuring IP levels in nuclei, cells were treated as described (33,34), nuclei were prepared and then resuspended in intracellular buffer (in mM: 125 KCl, 2 KH\textsubscript{2}PO\textsubscript{4}, 2 MgCl\textsubscript{2}, 0.3 CaCl\textsubscript{2}, 10 D-glucose, 1 ATP and 40 HEPES pH 7.0) before being processed further (34). Data calculated as the ratio of labeled IP or IP\textsubscript{3} to total radioactivity (inositol, IP\textsubscript{1}, IP\textsubscript{2} and IP\textsubscript{3}) were expressed as percentage of response in unstimulated HEK cells or nuclei expressing mGlu5.

RESULTS

mGlu5 Receptors are Localized at both the Plasma and Nuclear Membranes—We have previously shown that mGlu5 is expressed on plasma membranes as well as on many intracellular membranes including nuclear membranes (18,19). To confirm and extend these studies, we expressed N-terminal HA-tagged mGlu5 in HEK cells and used anti-HA antibody and differential permeabilization techniques to verify the presence of mGlu5 on plasma and nuclear membranes (supplemental Fig. S1). In other studies immunogold EM with antibodies directed against the C-terminal portion of mGlu5 was used to show that immunogold particles were abundantly associated with ER and nuclear membranes (supplemental Fig. S1). Previously we showed that agonists such as glutamate and quisqualate reach nuclear receptors via both sodium-dependent transporters and cystine glutamate exchangers (19). Here, we confirm that quisqualate is taken up by HEK cells, that the mGlu5 agonist, DHPG is an impermeable agonist, and that the drugs LY395053 and LY367366 are impermeable antagonists (35) (supplemental Fig. S2). Thus, as an impermeable agonist, DHPG cannot activate nuclear mGlu5 receptors in isolated nuclei derived from mGlu5 expressing HEK cells (mGlu5/HEK) or endogenously expressed on striatal nuclear membranes whereas quisqualate can (supplemental Figs. S3E-H). Finally, using differential labeling with vital dyes such as MitoTracker which is excluded from the nucleus and Hoechst, which is retained in the nuclear compartment, we provide visible evidence of the purity of these nuclear preparations (supplemental Figs. S3A-D). Therefore mGlu5 receptors are present and functional on nuclear membranes in heterologous cells as well as endogenous striatal nuclei.

Nuclear mGlu5 Couples to Nuclear G\textsubscript{q/11} Proteins—Certain nuclear GPCRs appear to mediate nucleoplasmic Ca\textsuperscript{2+} changes via pertussis toxin sensitive pathways suggesting a G\textsubscript{q/11}-driven mechanism (21). However, we have previously shown that in HEK293 cells stably expressing mGlu5, pertussis toxin does not affect mGlu5-mediated cytoplasmic or nuclear Ca\textsuperscript{2+} oscillations (18), thus we hypothesized that nuclear mGlu5 couples to nuclear G\textsubscript{q/11} proteins to activate downstream signaling components. To test this
idea, an mGlu5 mutant was constructed in which phenylalanine at position 767 in the third intracellular loop was replaced with serine (F767S), leading to the loss of G-protein coupling (36,37). Like wild type mGlu5, the F767S mutant was localized on both plasma and nuclear membranes where it co-localized with the nuclear membrane marker, lamin B2 (Figs. 1A,B). Similarly, nuclei isolated from mGlu5 or F767S stable HEK cell lines co-expressed either receptor together with lamin B2 confirming expression on nuclear membranes (Figs. 1C,D). Using subcellular fractionation, both wild type and mutant receptors were clearly expressed on plasma membrane as well as nuclear fractions as indicated by the membrane-specific markers, Na+K+ATPase and lamin B2, respectively (Figs. 1E,F). Thus immunocytochemical analysis and western blotting techniques demonstrated that the expression pattern and protein levels of wild type and mutant mGlu5 in HEK cells were similar.

To determine whether the F767S mutant exhibited similar binding characteristics to wild type receptors, binding studies using [3H]quisqualate as a radioligand were performed on both nuclear and plasma membrane fractions. Dissociation constants (Kd) for wild type and mutant mGlu5 plasma membrane receptors were 385.4±54.6 and 492.8±36.1 nM, respectively. The Kd's for wild type and mutant mGlu5 nuclear receptors were 576.6±15.9 and 752.7±43.5 nM respectively. The total number of mGlu5 binding sites (Bmax) as revealed from non-linear regression analysis was 896.9±49.4 fmol/mg for wild type mGlu5 plasma membrane receptors, 1285.5±128.9 fmol/mg for mutant mGlu5 plasma membrane receptors, 638.7±54.9 fmol/mg for wild type nuclear receptors and 1060.7±133.2 fmol/mg for mutant mGlu5 nuclear receptors. HEK cells expressing empty vector did not exhibit significant specific binding to [3H]quisqualate (Kd > 2.0 mM). Thus, like wild type nuclear mGlu5 receptors (18), F767S binds agonist and appears to be correctly folded in HEK cells. Data pooled across four experiments indicates that about 51.7%±3.5 of mGlu5 receptors are on plasma and intracellular membranes and that 48.3%±3.5 of mGlu5 receptors are present on nuclear membranes derived from mGlu5/HEK cells.

To test whether F767S could mediate Ca2+ changes, HEK cells stably expressing wild type or mutant mGlu5 were loaded with the Ca2+ indicator Oregon Green BAPTA-1AM. Esterified Oregon Green BAPTA-1AM is hydrolyzed within the nucleus such that it is retained for at least 30 min (38,39). As shown previously (18), bath application of glutamate induced Ca2+ oscillations in both cytoplasm and nucleoplasm of wild type cells that were inhibited by the membrane-permeable mGlu5-specific antagonist MPEP (Fig. 2A). In contrast, no such oscillations were observed in cells expressing F767S (Fig. 2B). More directly, agonist induced Ca2+ oscillations in nuclei derived from wild type cells whereas no such induction was observed in nuclei isolated from F767S cells (Fig. 2C versus 2D). The quantitation of data from 3-5 independent experiments is shown in Figs. 2E,F. Thus although the F767S mutant bound ligand and was expressed on the same membranes as the wild type receptor, it was completely inactive in either intact cells or isolated nuclei. Given that the F767S mutation prevents G-protein coupling, and that pertussis toxin was ineffective in blocking mGlu5-mediated nuclear Ca2+ oscillations (18) as well as that HEK293 cells do not express Go (36), these data demonstrate that mGlu5-induced Ca2+ oscillations in the nucleus are a consequence of functional nuclear Gq/11 coupling.

Nuclear mGlu5 Stimulates Nuclear PI-PLC- On the plasma membrane, activation of Gq/11-coupled receptors leads to PLC activation and PIP2 hydrolysis. Because both the substrate PIP2 (40) as well as PLC (β1 isoform; 41) are known to be in the nucleus (13), we tested whether nuclear mGlu5 mediates nuclear Ca2+ changes due to PLC stimulation. Using the same experimental paradigm, mGlu5-expressing nuclei were treated with glutamate to induce real time nuclear Ca2+ oscillations prior to bath application of the PLC inhibitor U73122 (3 μM) or its inactive analog U73343 (3 μM). This concentration was chosen since others have shown that 3 μM U73122 specifically inhibits PLC activity without affecting intracellular Ca2+ levels (27,42). As shown in Figs. 3A,F, the active analog U73122 inhibited glutamate-induced Ca2+ oscillations, whereas the inactive analog, U73343, did not (Figs. 3B,F). Moreover, pre-treatment with U73122 (3 μM for 120 s) prevented glutamate-induced Ca2+ oscillations in mGlu5-expressing HEK nuclei.
whereas pre-treatment with either U73343 or the drug vehicle, 0.1% DMSO, did not prevent Ca\(^{2+}\) oscillations (n > 10; data not shown).

Given that U73122 inhibits all types of PLCs, we further tested whether PI-PLC or phosphatidylincholine PLC (PC-PLC) was involved. To do so, mGlu5-responding nuclei were treated either with the PI-PLC inhibitor ET-18-OCH\(_3\) or the PC-PLC selective inhibitor D609 (42). Only the PI-PLC inhibitor blocked mGlu5-mediated nuclear Ca\(^{2+}\) oscillations (Figs. 3C,D,F). Finally, because studies have shown that IGF-1 induces nuclear Ca\(^{2+}\) transients through G\(_{q/11}\) subunits and PLC as well as via PI3 kinase (PI3K; 12), we tested whether the PI3K inhibitor LY294002 could block agonist-induced Ca\(^{2+}\) responses. Bath application of LY294002 had no effect on mGlu5-mediated nuclear responses (Figs. 3E,F). Taken together, these data reveal that like plasma membrane receptors, nuclear mGlu5 receptors also couple to G\(_{q/11}\) and PLC. Inasmuch as all of the experiments were done in acutely isolated nuclei, it is clear that all of the necessary enzymatic machinery is available for these responses independent of cellular components.

Nuclear mGlu5 Generates Nuclear IP\(_3\)-

Due to the transient and readily metabolizable nature of IP\(_3\), it can be difficult to accurately measure in the cytosol let alone the nucleoplasm. Indeed, only one study has reported IP\(_3\) changes in isolated nuclei (33). Moreover, mGlu5 is known to exhibit high constitutive activity in heterologous cell types (43). Not surprisingly then, biochemical assays to measure agonist-induced changes in inositol phosphates (IP) or more specifically, IP\(_3\), revealed small yet significant increases in IP and IP\(_3\) levels as compared to untreated controls. Specifically, 41 and 21% increases in IP levels were observed in glutamate treated mGlu5/HEK cells and nuclei, respectively (cytoplasmic IP\(_3\) levels were normalized at 100.0\(+\)0.1 without glutamate and 141.2\(+\)5.3 in the presence of 10 \(\mu\)M glutamate. Nuclear IP levels were 100.0\(+\)0.3, in the absence of glutamate and 120.8\(+\)5.2 in its presence; n = 3, *, p < 0.05). Separate experiments examining IP\(_3\) changes revealed an approximately 25% increase in glutamate treated mGlu5/HEK cells and about a 15% increase in isolated mGlu5/HEK nuclei (cytoplasmic IP\(_3\) levels were normalized at 100.0\(+\)2.5 in the absence of glutamate whereas glutamate-treated cytoplasmic IP\(_3\) levels were 125.0\(+\)5.2. IP\(_3\) levels in isolated nuclei were 100.0\(+\)1.3 in untreated controls and 115.3\(+\)3.2 for glutamate-treated; n = 3, *, p < 0.05). Consistent with the notion that mGlu5 is constitutively active, the inverse agonist, MPEP, which locks the receptor into its inactive state (44), reduced basal IP levels by approximately 5-fold in the absence (19.2\(+\)0.7) or presence of glutamate (23.5\(+\)7.6). Moreover, IP levels were about 17% in F767S/HEK cells regardless of treatment.

To circumvent the limitations of the biochemical assay, we used a well-established construct ‘pEGFP-C1-PLC\(_{\delta 1}\)-PH’ in which the pleckstrin homology (PH) domain of PLC\(_{\delta 1}\) with its high affinity for the polar group of PIP2 has been tagged with GFP (26,45). This probe is bound to PIP2 in the plasma membrane and the increase in IP\(_3\) is indicated by the translocation of the fusion protein from the plasma membrane to the cytoplasm. Because this probe not only depends upon IP\(_3\) but also on the PIP2 concentration in the plasma membrane, it is perhaps more aptly referred to as a PIP2/IP\(_3\) biosensor (46). Therefore, mGlu5/HEK cells were transiently transfected with the PIP2/IP\(_3\) biosensor, nuclei were isolated, and GFP-expressing nuclei were imaged in real time (Fig. 4). Under basal conditions the PIP2/IP\(_3\) biosensor is located at the inner nuclear membrane due to its affinity for PIP2 (Fig. 4A, panel 2). Upon glutamate treatment and PLC activation, the biosensor moved off the membrane into the nucleoplasm owing to its 20-fold higher affinity for IP\(_3\) (Fig. 4A, remaining panels; 45). Glutamate-induced biosensor responses were oscillatory and completely blocked by MPEP (Figs. 4A,B). Isolated HEK nuclei expressing the PIP2/IP\(_3\) biosensor or F767S mutant HEK nuclei co-expressing the biosensor never showed this response (data not shown). When analyzed by expressing the peak increases in nucleoplasmic GFP as compared to basal fluorescence, significant changes in F/F\(_0\) were observed (Fig. 4C).

To demonstrate more clearly that nuclear mGlu5-induced IP\(_3\) leads to Ca\(^{2+}\) changes in the nucleus, we loaded mGlu5/HEK nuclei expressing the PIP2/IP\(_3\) biosensor with the Ca\(^{2+}\) fluorophore, Calcium Crimson-AM. Bath application of glutamate leads to marked biosensor movement, which was concomitant with the rise in Ca\(^{2+}\) in...
mGlu5/HEK nuclei (Figs. 4D,E). Collectively, these data demonstrate that activated nuclear mGlu5 receptors generate nuclear IP3.

**Nuclear mGlu5-mediated Ca2+ Changes Originate from IP3R and RyR**- Numerous studies have shown that upon activation IP3R and RyR located on the inner nuclear membrane can release Ca2+ from the nuclear lumen into the nucleoplasm (7,9). Therefore, we used specific IP3R and RyR inhibitors to test the hypothesis that nuclear mGlu5 receptors coupled to Gq/11 and PLC activate inner nuclear membrane Ca2+ channels to mediate Ca2+ changes. Glutamate-induced Ca2+ oscillations in isolated mGlu5/HEK nuclei were blocked by both the IP3R inhibitor, 2-APB (100 μM) and the RyR antagonist, ryanodine (100 μM) (Fig. 5). Pretreating with either drug prevented the induction of mGlu5 responses in nuclei isolated from the mGlu5/HEK cell line (not shown). Another IP3R antagonist, the highly specific Xestospongin C (2 μM; 47,48) also inhibited glutamate-induced Ca2+ oscillations in isolated mGlu5/HEK nuclei by approximately 75% (Figs. 5B,D). Thus, mGlu5-mediated Ca2+ responses arise via IP3R and RyR channels.

**Striatal Nuclear mGlu5 also Uses the Gq/11/PI-PLC/IP3 Pathway to Mediate Nuclear Ca2+ Changes**- To extend these findings to a more physiological system, we examined nuclear mGlu5 signal transduction pathways in dissociated striatal neurons or in nuclei acutely isolated from striatal tissue that we have previously shown to express functional nuclear mGlu5 receptors (19). The role of Gq/11 in mediating mGlu5 responses was determined by transfecting striatal neurons with dominant-negative Gαq (DNGαq) together with DsRed2 or with DsRed2 only. Two days later neurons were loaded with Oregon Green BAPTA-1 AM, imaged to acquire base-line Ca2+ changes and then treated with 10 μM quisqualate. As quisqualate also activates AMPA channels and mGlul receptors, it was bath-applied in the presence of 5 μM GYKI53655, an AMPA antagonist, and 20 μM CPCCOEt, an mGlul antagonist. Consistent with previous results (19), neurons transfected with DsRed2 alone showed Ca2+ increases in both the cytoplasm as well as nucleus consisting of two phases, an initial rapid rise followed by a sustained plateau. Both responses could be terminated by addition of MPEP (Figs. 6A,C). In contrast, neurons co-transfected with DNGαq and DsRed2 failed to show similar changes in cytoplasmic or nuclear Ca2+ levels (Figs. 6B,D). Specifically, there was approximately an 80% reduction in the nuclear Ca2+ levels in cells co-transfected with DsRed2 and DNGαq as compared to DsRed2 only following quisqualate treatment (Fig. 6E). Cell surface mGlu5 receptors served as an internal positive control of DNGαq efficacy; hence cytoplasmic Ca2+ levels were also reduced by 80-85% in cells co-transfected with DsRed2 and DNGαq versus DsRed2-only following quisqualate treatment (Fig. 6). For further support of a predominant role of Gq/11 in mGlu5-mediated nuclear Ca2+ increases, striatal cultures were pretreated with pertussis toxin for 18 h. Like mGlu5/HEK cells, pertussis toxin did not affect striatal mGlu5-mediated cytoplasmic or nuclear Ca2+ responses ruling out a Gαi/o-mediated response (n > 15; data not shown).

To determine whether PLCβ was involved in nuclear mGlu5-mediated Ca2+ responses, siRNA targeted against PLCβ1, the most prevalent nuclear isoform (41), was used to knockdown expression. Western blotting of striatal lysates prepared 48 and 72 h after siRNA transfection showed that there was at least a 60-70% knockdown of PLCβ1 using the targeted siRNA versus a scrambled control (Figs. 7A,B). Immunostaining of transfected cultures further revealed a knockdown of PLCβ1 following the introduction of PLCβ1-specific siRNA versus the scrambled control (Figs. 7C,D last panels). Scrambled siRNA together with DsRed2 had no effect on agonist-induced mGlu5 Ca2+ responses (Figs. 7C,E,G) whereas neurons co-transfected with PLCβ1 siRNA exhibited a 78% reduction in cytoplasmic and an 83% reduction in nuclear Ca2+ following quisqualate treatment (Figs. 7D,F,G). Pharmacological support of these data comes from acutely isolated striatal nuclei experiments in which nuclei were loaded with the Ca2+ fluorophore followed by quisqualate to induce a sustained nuclear Ca2+ response. This response could be blocked by MPEP (19; not shown), U73122 (Figs. 8A,F), or ET-18-OCH3 (Figs. 8C,F) but not U73343 (Figs. 8B,F), D609 (Figs. 8D,F) or LY294002 (Figs. 8E,F). Thus, PI-PLC but not PC-PLC or PI3K are required for nuclear Ca2+...
signaling following quisqualate activation of mGlu5 nuclear receptors.

To test whether mGlu5-induced striatal nuclear Ca\textsuperscript{2+} changes were mediated via IP\textsubscript{3} generation, neurons were transfected with the PIP2/IP3 biosensor. Images taken from a single optical plane revealed that GFP fluorescence was associated with both cell surface (Fig. 9; blue arrows) and nuclear membranes (Fig. 9; red arrows) when compared with the corresponding transmitted light images (Fig. 9A) or with the lamin B2 staining (Fig. 9B). These results are consistent with the HEK data (Fig. 4) and with reports that PLC\textsubscript{δ1} accumulates in nuclear compartments (49,50). Upon quisqualate stimulation, the PIP2/IP3 biosensor moved away from either membrane exhibiting a sustained fluorescent increase akin to observed Ca\textsuperscript{2+} changes (Figs. 9C,D). Peak increases in cytoplasmic or nucleoplasmic fluorescence expressed as a percent of basal fluorescence showed similar responses (Fig. 9E). To ensure that these were mGlu5-mediated responses, cultures were fixed, stained with an anti-mGlu5 antibody, and subsequently field-relocated (not shown). Moreover, quisqualate-mediated Ca\textsuperscript{2+} responses could be blocked by 2-APB, Xestospongin C or ryanodine (Fig. 10). Taken together, these data strongly support a model in which nuclear mGlu5 receptors lead to the activation of G\textsubscript{q/11}, PLC, and IP\textsubscript{3} to generate changes in nuclear Ca\textsuperscript{2+} levels.

The basic signaling components ascribed to nuclear mGlu5 receptors are supported by a number of observations. First, mutations that block mGlu5 coupling to G\textsubscript{q/11} prevented cytoplasmic and nuclear Ca\textsuperscript{2+} changes in heterologous cells and their isolated nuclei despite normal agonist binding (Fig. 2). Second, when expressed in striatal neurons, dominant negative G\textsubscript{q} abolishes mGlu5-mediated nuclear Ca\textsuperscript{2+} increases (Fig. 6). Third, low concentrations of the widely used PLC inhibitor, U73122, blocked mGlu5-mediated increases of nuclear Ca\textsuperscript{2+} in both mGlu5/HEK and striatal nuclei (Figs. 3,8). Moreover, knockdown of PLC\textsubscript{β1} in striatal neurons significantly reduced mGlu5-mediated nuclear Ca\textsuperscript{2+} responses as well (Fig. 7). Finally, in situ IP\textsubscript{3} production was revealed following mGlu5 activation in both heterologous and striatal nuclei using a sensitive optical PIP2/IP3 biosensor approach (Figs. 4,9). Taken together, these data strongly support a model in which nuclear mGlu5 receptors lead to the activation of G\textsubscript{q/11}, PLC, and IP\textsubscript{3} to generate changes in nuclear Ca\textsuperscript{2+} levels.

The traditional idea that GPCRs signal only from the cell surface is gradually being refined by studies showing that even internalized receptors can serve as scaffolds for signaling molecules (55) or, more directly, intracellular receptors can couple to various intracellular G proteins. Thus if mechanisms exist by which a receptor might be activated, signaling molecules are available to transmit the signal. Receptor activation might be accomplished in a variety of ways. For example, a large number of GPCRs such as the prostaglandin, PAF, and LPA receptors, whose ligands are bioactive lipids derived from membrane hydrolysis, are located on nuclear membranes (review, 21). As ligand-generating enzymes are also present on nuclear membranes and because such ligands readily diffuse through lipid bilayers, PGE\textsubscript{2}, PAF and LPA can easily activate their cognate receptors. In contrast, mGlu5 or mGlu1 ligand binding domains are within the nuclear lumen such that agonists must traverse both the cell surface lipid bilayer as well as the outer nuclear membrane for receptor activation (18,19,22). Mechanistically agonist
transport is achieved via the sodium-dependent glutamate transporter and/or the cystine, glutamate xCT exchanger (19,22). Although other mechanisms might also lead to intracellular mGlu5 or mGlu1 receptor activation, direct transfer of ligand is at least one effective means of delivering agonist to such receptors.

There appear to be many different mechanisms involved in mobilizing Ca\(^{2+}\) from storage in the nuclear envelope. Many of the described nuclear GPCRs couple to several different G proteins including G\(_{q/11}\) and G\(_{i/o}\). To date, G\(_{i/o}\)-mediated signaling pathways seem to predominate regardless of whether a particular GPCR couples to G\(_{q/11}\) at the plasma membrane. For instance, although PGE2 EP1 receptors couple to G\(_{q/11}\) at the cell surface, nuclear EP1 receptors generate nucleoplasmic Ca\(^{2+}\) signals in a pertussis toxin sensitive fashion (56,57). Similarly, although the LPA-1 receptor is known to interact with G\(_{i/o}\), G\(_{q/11}\) and G\(_{12/13}\) proteins at the cell surface, it is also G\(_{i/o}\)-coupled on nuclear membranes (20). IGF-1 stimulation is also known to mobilize Ca\(^{2+}\) from nuclear stores. Results suggest that extracellular IGF-1 activates IGF-1R, which in turn activates a pertussis toxin-sensitive G protein. The latter stimulates PI3K and subsequently PLC to generate cytoplasmic/perinuclear IP\(_3\) which diffuses into the nuclear lumen (11,12). The possibility that cytoplasmic IP\(_3\) diffuses into the nucleus via nuclear pore complexes can be ruled out here because the present experiments utilize pure isolated nuclei. Thus, for nuclear mGlu5 receptors, signal transduction appears to be via the same canonical G\(_{i/o}\)/PLC/IP\(_3\) pathway that is also found at the plasma membrane (58).

Interestingly, both Ca\(^{2+}\) channels, IP\(_3\)R and RyR, contribute to the mGlu5-mediated nuclear Ca\(^{2+}\) rises since adding their respective antagonists blocked the Ca\(^{2+}\) changes. Although activation of IP\(_3\)Rs is consistent with numerous studies showing that mGlu5 generates IP\(_3\), participation of RyR channels in the same process is intriguing. One possibility is that IP\(_3\) mediated Ca\(^{2+}\) release leads to the activation of the RyR. In contrast, ryanodine may directly inhibit IP\(_3\)-mediated Ca\(^{2+}\) signals (59). Which of these models, direct or indirect inhibition of IP\(_3\)-evoked Ca\(^{2+}\) release, holds true for mGlu5-mediated Ca\(^{2+}\) changes remains to be tested. Finally, since IP\(_3\)Rs are present on the inner nuclear membrane, the translocation of IP\(_3\) from its site of synthesis into the nucleoplasm as inferred from the PIP2/IP\(_3\) biosensor experiments, is puzzling (Fig. 4A). Conceivably, IP\(_3\) generated at the nuclear membrane might translocate to the nucleoplasmic reticulum that also expresses IP\(_3\)Rs in order to further release nuclear Ca\(^{2+}\) in specialized sub domains (6,60). In support of this idea, translocation of the biosensor was never uniform in neuronal nuclei; biosensor movement was skewed towards one side of the nucleus or another (Fig. 9C). Thus, spatial constraints, diffusion characteristics and/or intranuclear buffering capabilities may regulate specific local Ca\(^{2+}\) signals in nuclear sub domains.

It is widely believed that the amplitude, duration, and/or frequency of Ca\(^{2+}\) fluctuations can differentially regulate downstream effectors such as transcriptional regulators, coactivators and/or modifying enzymes (61-63). In the nucleus differential regulation of Ca\(^{2+}\) is crucial since blocking increased nuclear Ca\(^{2+}\) prevents Ca\(^{2+}\)-induced developmental processes and/or long term plasticity (62,64-66). For example, using an in vivo genetic approach, Limback-Stokin et al. (67) demonstrated that nuclear Ca\(^{2+}\)signaling pathways, not cytoplasmic, were responsible for converting short term memory into long term memory. Clearly, one consequence of nuclear mGlu5 activation is prolonged nuclear Ca\(^{2+}\) responses (19, studies herein). Presumably, sustained mobilization of nuclear Ca\(^{2+}\) activates different signaling pathways including nuclear Ca\(^{2+}\)/calmodulin-activated kinase IV, a kinase known to phosphorylate the cAMP response element binding protein (CREB; 68). Indeed, we have previously shown that agonist treatment can directly activate CREB in mGlu5-expressing isolated striatal nuclei (19). CREB may, in turn, lead to de novo gene transcription. Although these notions remain to be tested, activation of nuclear receptors creates yet one more way in which a cell can nuance its responses to a given stimuli.

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† The abbreviations used are: IP3R, inositol 1,4,5-trisphosphate receptors; GPCRs, G-protein coupled receptors; mGlu5, metabotropic glutamate receptor 5; PLC, phospholipase C; PI, phosphatidylinositol; PIP2, phosphatidylinositol 4,5-bisphosphate; ER, endoplasmic reticulum; HEK, human embryonic kidney; EGFP, enhanced green fluorescent protein; PH, pleckstrin homology; MPEP, 2-methyl-6-(phenylethynyl)pyridine; CPCCOEt, 7-(hydroxyimino)cyclopropan[β]chromen-1-carboxylate ethyl ester; BAPTA, 1,2-bis(2-aminophenoxy)ethane-Ν,Ν,Ν',Ν'-tetraacetic acid; AMPA, α-amino-3-hydroxy-5-methyl-4-isoxazolepropionic acid; 2-APB, 2-aminoethoxydiphenyl borate; CREB, cAMP-response element-binding protein.

FIGURE LEGENDS

Fig. 1. mGlu5 wild type and F767S mutant exhibit nuclear localization. Stable HEK cell lines expressing either wild type mGlu5 or the F767S mutant, were fixed, permeabilized, and processed for immunocytochemistry using receptor-specific antibodies as well as anti-lamin B2. Cells were analyzed by confocal microscopy to detect receptor localization (red) or lamin B2 distribution (green). Photographs represent single optical sections of 0.4 µm merged such that yellow indicates co-localization of the specific antigens. Receptors expressed by intact cells (A, B; TL, transmitted light) or their isolated nuclei (C, D) exhibited pronounced co-localization with lamin B2. Subcellular fractionation of HEK cell lines expressing either wild type mGlu5 (E) or the F767S mutant (F) shows that wild type or mutant receptors can be detected in fractions containing either the nuclear (N) or plasma membranes (PM). Equal amounts of protein (30 µg) from each fraction were separated on reducing SDS gels and transferred to nylon membranes (T, total cell lysates). The same blot was sequentially probed with antibodies against mGlu5, the inner nuclear marker, lamin B2, and the plasma membrane marker, Na\(^+\)K\(^+\)-ATPase.

Fig. 2. Nuclear mGlu5 couples to nuclear G\(_{q/11}\) protein. Representative traces are shown of cytoplasmic (grey line) or nuclear (black line) Ca\(^2+\) responses from wild type mGlu5 expressing HEK cells (A) or the F767S cell line (B) as well as their respective nuclei (C, D) following bath addition of 10 µM glutamate (Glu) and 1 µM MPEP when indicated and scanned at 5.4 s/scan. Oscillations are represented as the fractional change in fluorescence relative to the basal value. Compiled data from maximum response of initial peak (ΔF/F\(_{0}\), %) from either wild type cells or nuclei (E) or F767S cells or nuclei (F) from n \(>\) 30 cells, from three to five independent experiments. *, p < 10\(^{-4}\) when compared with Ca\(^2+\) responses from wild type mGlu5/HEK cells and nuclei treated with glutamate only.

Fig. 3. Nuclear mGlu5 stimulates nuclear PI-PLC. A-E, representativ traces of nuclear (black line) Ca\(^2+\) responses in isolated mGlu5-expressing HEK nuclei represented as the fractional change in fluorescence relative to the basal level. Isolated nuclei were treated with 10 µM glutamate and the specified antagonists bath applied when indicated by the arrow; U73122 (PLC inhibitor, 3 µM), U73343
neurons were transiently co-transfected with scrambled siRNA and DsRed2 (0.005 when compared with PLCβ1 siRNA transfected neurons. The data shown are compiled from three independent experiments. *, p < 0.001 when compared with Ca2+ responses from nuclei treated with glutamate only.

**Fig. 4. Nuclear mGlu5 generated IP3 production can be measured using PIP2/IP3 biosensor.** A, 1st panel, transmitted light image of selected nucleus; remaining panels, time-lapse imaging of mGlu5/HEK nucleus transiently transfected with PIP2/IP3 biosensor treated at the indicated times (seconds) with 10 μM glutamate or 1 μM MPEP. Red circle corresponds to area measured for representative trace shown in B, where Glu-mediated oscillations are represented as the fractional change in fluorescence relative to the basal IP3 levels. C, compiled data from maximum response of initial peak (ΔF/F₀, %) for nuclear responses from n = 5, from three experiments. *, p < 0.001 when compared with baseline IP3 responses. D, nuclei isolated from mGlu5/HEK stable cell line transiently transfected with the PIP2/IP3 biosensor were loaded with Calcium Crimson-AM in order to simultaneously measure Ca2+ and biosensor changes. Shown is a representative trace of nuclear Ca2+ (red line) and PIP2/IP3 biosensor (green line) responses in an isolated nucleus treated with 10 μM glutamate and 1 μM MPEP as indicated by the arrow. E, compiled data from maximum response of initial peak (ΔF/F₀, %) for nuclear IP3 (green) and Ca2+ (red) responses from n = 5, from three independent experiments. *, p < 0.001 when compared with baseline IP3 response; **, p < 0.01 when compared with baseline Ca2+ response.

**Fig. 5. Glutamate-activated nuclear mGlu5-mediated Ca2+ changes can be abrogated by IP3R and RyR blockers.** A-C, representative traces of nuclear (black line) Ca2+ responses in isolated mGlu5-expressing nuclei represented as the fractional change in fluorescence relative to the basal level. Isolated nuclei were treated with 10 μM glutamate and the indicated antagonists; 2-APB (IP3R modulator, 100 μM), Xestospongin C (IP3R antagonist, 2 μM) and Ryanodine (RyR antagonist, 100 μM). D, compiled data from maximum response of initial peak (ΔF/F₀, %) for nuclear responses from n = 20 for 2-APB and Ryanodine and n = 14 for Xestospongin C, from three independent experiments. *, p < 0.001 when compared with Ca2+ responses from nuclei treated with glutamate only.

**Fig. 6. Endogenous mGlu5 receptors expressed on striatal neurons couple to the Gq family of G-proteins.** DIV12 striatal neurons transiently transfected with DsRed2 (A) or co-transfected with dominant negative Goαi and DsRed2 (B) were loaded with Oregon Green BAPTA-1AM (2nd panels) and quisqualate (Quis; 10 μM) was bath-applied (3rd panels). Last panels, images of post hoc identified mGlu5-positive cells. Representative traces are shown of quisqualate-mediated cytoplasmic (blue line) or nuclear (red line) Ca2+ responses from a control neuron (DsRed2 transfected; C) or a neuron co-transfected with dominant negative Goαi and DsRed2 (D). MPEP was added as indicated (1 μM; black line). E, compiled data from maximum response of initial peak (ΔF/F₀, %) from either control cells (n = 22) or dominant negative Goαi transfected neurons (n = 20) from four independent experiments. *, p < 0.0001 when compared with Ca2+ responses from DsRed2 transfected control cells.

**Fig. 7. Knockdown of PLCβ1 leads to reduction in striatal mGlu5-mediated Ca2+ changes.** A, primary striatal neurons were transiently transfected with either scrambled siRNA (S) or PLCβ1 siRNA (P). At the times indicated, cells were lysed and proteins were separated on reducing SDS gels and transferred to nylon membranes. The same blot was sequentially probed with antibodies against PLCβ1 (Santa Cruz) and β-actin. B, the relative abundance of PLCβ1 was measured by western blotting; the efficiency of knockdown is expressed in arbitrary units compared to the PLCβ1 levels in scrambled siRNA transfected neurons. The data shown are compiled from three independent experiments. *, p < 0.005 when compared with PLCβ1 protein levels in scrambled siRNA transfected cells. DIV12 striatal neurons were transiently co-transfected with scrambled siRNA and DsRed2 (C) or with PLCβ1 siRNA
and DsRed2 (D). Two days later, neurons were loaded with Oregon Green BAPTA-1AM (2nd panels) and quisqualate (Quis; 10 µM) was bath-applied (3rd panels). Representative traces are shown for cytoplasmic (blue line) or nuclear (red line) Ca$^{2+}$ responses from scrambled (E) or PLCβ1 siRNA (F) following quisqualate and subsequently, 1 µM MPEP administration (black line). Imaged neurons were post hoc identified using mGlu5 (4th panels) or PLCβ1 antibodies (last panels). G, compiled data from maximum response of initial peak ($\Delta F/F_o$, %) from either scrambled ($n=13$) or PLCβ1 ($n=17$) siRNA transfected neurons from three independent experiments. **, $p<0.0001$ when compared with Ca$^{2+}$ responses from control siRNA and DsRed2 transfected cells.

**Fig. 8. Endogenous mGlu5 receptors expressed on striatal nuclei mediate Ca$^{2+}$ changes via PI-PLC pathway.** P10 striatal nuclei were acutely isolated, loaded with Oregon Green BAPTA-1AM, and imaged in real time. A-E, quisqualate-mediated (10 µM) representative traces of nuclear (black line) Ca$^{2+}$ responses treated with indicated antagonists bath applied as noted. F, compiled data from maximum response of initial peak ($\Delta F/F_o$, %) for nuclear Ca$^{2+}$ responses induced by quisqualate alone or together with specified antagonists from $n>8$ in all cases, from three to four independent experiments. *, $p<0.001$ when compared with Ca$^{2+}$ responses from nuclei treated only with quisqualate.

**Fig. 9. Activation of striatal mGlu5 receptors generates nuclear IP$_3$.** DIV12 striatal cultures were transiently transfected with the PIP2/IP$_3$ biosensor. A, single optical section of 0.25 µm showing transmitted light image (TL, 1st panel), biosensor fluorescence (PIP2/IP$_3$, 2nd panel), and the merged image (3rd panel). B, single optical section of 0.25 µm showing biosensor fluorescence alone (green; 1st panel), lamin B$_2$ immunofluorescence (red; 2nd panel), and the merged image (3rd panel) on nuclear (red arrow) and plasma membranes (blue arrow). C, 1st panel, single optical section of 0.25 µm showing biosensor distribution on nuclear (red arrow) and plasma membranes (blue arrows); remaining panels, translocation of biosensor following 10 µM quisqualate application as indicated. Circles correspond to areas measured for representative trace shown in D (cytoplasmic blue or nuclear red) where biosensor changes are represented as the fractional change in fluorescence relative to the basal levels of fluorescence. E, compiled data from maximum response of initial peak ($\Delta F/F_o$, %) for cytoplasmic (blue) or nuclear (red) responses from $n=5$, from three separate experiments. *, $p<0.01$ when compared with baseline responses.

**Fig. 10. Striatal mGlu5 receptors release nuclear Ca$^{2+}$ via Ca$^{2+}$ release channels.** A-C, representative traces of nuclear (black line) Ca$^{2+}$ responses in isolated striatal nuclei represented as the fractional change in fluorescence relative to the basal level. Isolated nuclei were treated with 10 µM quisqualate and indicated antagonists. D, Bar graph shows compiled data from maximum response of initial peak ($\Delta F/F_o$, %) for nuclear responses from $n>10$ for either antagonist, from three independent experiments. After treatment with antagonists the Ca$^{2+}$ responses were significantly different when compared with Ca$^{2+}$ responses in presence of quisqualate only (*, $p<0.001$).

**Fig. 11. Proposed model of signal transduction pathway associated with nuclear mGlu5 receptors.** Endogenous striatal nuclear mGlu5 receptors, like plasma membrane receptors, couple to the $G_{q11}$/PI-PLC/IP$_3$ pathway. EAAT: sodium-dependent transporter, xCT: cystine-glutamate transporter, DAG: diacyl glycerol, ER: endoplasmic reticulum, NR: nucleoplasmic reticulum.
Figure 1

Wild Type mGlur5

Intact Cells

A TL  WT  Lamin B2  WT/Lamin B2

Mutant F767S mGlur5

Intact Cells

B TL  F767S  Lamin B2  F767S/Lamin B2

Isolated Nuclei

C TL  WT  Lamin B2  WT/Lamin B2

D TL  F767S  Lamin B2  F767S/Lamin B2

E T  PM  N

mGlur5

Na+ K+ ATPase

Lamin B2

F T  PM  N

mGlur5

Na+ K+ ATPase

Lamin B2
Figure 4

A

B

C

D

E

10 \mu m

\Delta F/F_o (%) vs. sec

\Delta F/F_o (%) vs. sec

\Delta F/F_o (%) vs. Glu

\Delta F/F_o (%) vs. Glu

Glu

MPEP

Glu

MPEP

Glu

MPEP

Glu

MPEP

Glu

MPEP

0

50

100

0

50

100
Figure 5

(A) Glu → 2-APB

(B) Glu → Xestospongin

(C) Glu → Ryanodine

(D) Graph showing the effects of various treatments on ΔF/F₀ (%)

| Treatment       | Glu | 2-APB | Xestospongin | Ryanodine |
|-----------------|-----|-------|--------------|-----------|
|                  | +   | +     | +            | -         |
|                  | +   | +     | -            | +         |
|                  | +   | -     | -            | +         |

* indicates statistical significance.
Figure 7

A

| 48 | 72 h |
|----|------|
| S  | P    | S  | P    |
| ![Image](image1.png) | ![Image](image2.png) | ![Image](image3.png) | ![Image](image4.png) |

PLCβ1

β-actin

B

Relative abundance (arbitrary units)

| 48 | P    | S  | 72 h |
|----|------|----|------|
| ![Image](image5.png) | ![Image](image6.png) | ![Image](image7.png) | ![Image](image8.png) |

C

Scrambled siRNA

DsRed2

Oregon Green

Quis

mGlu5

PLC

D

PLC siRNA

10 µm

10 µm

E

F

G

ΔF/F₀ (%)

ΔF/F₀ (%)

ΔF/F₀ (%)

Scrambled siRNA

PLC siRNA

Quis

MPEP

Quis

MPEP

** **
Figure 9

A

TL  PIP2/IP$_3$  TL / PIP2/IP$_3$

B

PIP2/IP$_3$  Lamin B$_2$  PIP2/IP$_3$ / Lamin B$_2$

C

Quis

D

E

$\Delta F/F_0$ (%) vs. time (sec)

$\Delta F/F_0$ (%) vs. Quis

* significant difference
Figure 11
Activated nuclear metabotropic glutamate receptor mGlu5 couples to nuclear Gq/11 proteins to generate inositol 1,4,5-trisphosphate-mediated nuclear Ca2+ release

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