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CAPRI and RASAL impose different modes of information processing on Ras due to contrasting temporal filtering of Ca\(^{2+}\)

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Introduction

Ras is a key player in cell signaling with a central role in cancer. Oncogenic mutants are locked in the GTP-bound conformation resistant to GTPase-activating proteins (GAPs). We recently discovered closely related Ca\(^{2+}\)-triggered GAPs; RASAL (Allen et al., 1998; Walker et al., 2004) and CAPRI (Lockyer et al., 2001). Each contains tandem C2 domains (C2A and C2B) and a GAP-related domain (GRD) flanked by a pleckstrin homology (PH) domain and Bruton’s tyrosine kinase (Btk) motif. RASAL is a Ca\(^{2+}\) sensor responding in-phase to repetitive Ca\(^{2+}\) signals by associating with the plasma membrane and deactivating Ras (Walker et al., 2004). Here, we compare the behavior of CAPRI with that of RASAL and PKC\(_\gamma\) after G protein–coupled receptor stimulation. Studies of conventional PKC isoforms have resolved a frequency-dependent activation mechanism reliant on oscillatory associations with the plasma membrane in concert with fluctuations in intracellular Ca\(^{2+}\) (via the C2 domain) and DAG (via the C1 domain) (Oancea and Meyer, 1998; Violin et al., 2003). RASAL also tracks Ca\(^{2+}\) oscillations (Walker et al., 2004). However, CAPRI is a novel low-pass filter for Ca\(^{2+}\), dependent on predominantly Ca\(^{2+}\)-independent interactions between the plasma membrane and the PH domain after receptor activation.

Results and discussion

Spatial and temporal regulation of CAPRI by agonist-evoked Ca\(^{2+}\) signals

RASAL and PKC\(_\gamma\) are sensors of repetitive Ca\(^{2+}\) signals (Oancea and Meyer, 1998; Violin et al., 2003; Walker et al., 2004), and we wondered if CAPRI would behave similarly. We tested responses of GFP-CAPRI, GFP-RASAL, and GFP-PKC\(_\gamma\) to histamine stimulation of HeLa cells. This demonstrated long-term association of GFP-CAPRI with the plasma membrane at supra-maximal doses of agonist (Fig. 1 A and B). Half-maximal dissociation back to the cytosol was >280 s, contrasting with half-maximal dissociation of 17 s for GFP-RASAL and 13 s for GFP-PKC\(_\gamma\) (Fig. 1 C). A 10-μM dose...
evoked similar CAPRI translocation to the membrane as 100 μM histamine, but with a faster rate of dissociation (Fig. 1 B; half-maximal dissociation 88 s).

The novel kinetics of CAPRI translocation led us to test responses in other cell lines (Fig. S1, A and B; available at http://www.jcb.org/cgi/content/full/jcb.200504167/DC1). During measurements of RASAL- and PKCγ-expressing HEK293 cells, rapid oscillations between the cytosol and the plasma membrane could be readily detected (Fig. S1 A). 33% of RASAL-expressing cells displayed sinusoidal oscillations (n = 12), compared with 38% of PKCγ-expressing cells (n = 16). Evidence for oscillations in CAPRI translocation was absent. In fact, during >100 live-cell imaging experiments in multiple in vitro cell lines including CHO, HeLa, HepG2, HEK293, and COS-7 with a variety of Ca2+-mobilizing agonists (carbachol, histamine, and ATP), we failed to observe oscillatory interactions of GFP-CAPRI with the membrane (unpublished data). The orientation of the fluorescent protein had no influence (Fig. S1 C).

CAPRI does not suppress Ca2+ oscillations, but is refractory to them

Dawson first demonstrated that Ca2+ release could be stimulated by GTP and blocked by nonhydrolyzable GTP analogues (Dawson, 1985). Evidence has since accumulated that small GTPases are involved in the regulation of Ca2+ entry (Bird and Putney, 1993). A direct role for oncogenic Ras in regulating Ca2+ spike frequency has been shown in NIH3T3 cells (Lang et al., 1991). We therefore wondered whether CAPRI expression altered Ca2+ responses. In an earlier study, we generated a series of CHO cell lines stably expressing CAPRI or a CAPRI GRD mutant (R473S; GAP dead) (Lockyer et al., 2001). To determine if CAPRI expression had any impact on agonist-stimulated Ca2+ signals, we monitored cytosolic free Ca2+ in single cells. Supra-maximal ATP stimulation produced similar averaged Ca2+ peak–plateau transients between wild-type or mutant cells showing that Ras GAP function was not interfering with maximal Ca2+ responses with this agonist and cell type (Fig. 2 A). It did not exclude the possibility that CAPRI inhibits Ca2+ oscillations at lower agonist doses. To determine if CAPRI expression blocked Ca2+ oscillations in HeLa cells, we performed sequential imaging of GFP-CAPRI and Fura-2 (Fig. 2, B and C). As shown in Fig. 2 B, during sinusoidal Ca2+ signals there was a prolonged recruitment of CAPRI over a period of 500 s. This contrasted with the in-phase translocations of GFP-RASAL recently measured (Walker et al., 2004) and studies reported for other C2 domain–containing Ca2+ sensors such as conventional PKCs (Oancea and Meyer, 1998; Violin et al., 2003). Ca2+ spikes induced by a low dose of histamine generated weak, or typically more transient, CAPRI translocation (Fig. 2 C). At a submaximal histamine dose (1 μM) this was difficult to detect; by conventional confocal imaging only 5% of GFP-CAPRI–expressing cells displayed clear translocations (n = 60 cells), compared with 88% at 50 μM histamine (n = 75). During sequential GFP-CAPRI and Fura-2 imaging we detected no significant differences in spike periodicity and peak dye emission between transfected or nontransfected cells.
in the same field (unpublished data). Sequential confocal imaging of Fluo-4 and RFP-CAPRI also demonstrated that CAPRI does not faithfully track histamine-evoked changes in cytosolic Ca\(^{2+}\) (Fig. 2 D). As shown in Fig. 2 E, the translocation of GFP-CAPRI to the plasma membrane was dependent on agonist, was not cumulative, and displayed reversibility. A repetitive application of 50 \(\mu\)M histamine for 60 s followed by wash-out of agonist, followed by restimulation for 60 s as shown (average \(n = 3\) cells). (F) Translocation of CAPRI-YFP and CFP-RASAL induced by 50 \(\mu\)M ATP in HeLa cells and stimulated with ATP to prove that CAPRI and RASAL show different responses to the same stimulus, in the same cell (Fig. 2 F). Where RASAL oscillated, CAPRI translocated in a sustained manner.

**CAPRI translocation and Ca\(^{2+}\) entry**

We examined the role of extracellular Ca\(^{2+}\) for agonist-induced recruitment by stimulating GFP-CAPRI-expressing cells in the presence or absence of Ca\(^{2+}\). In histamine-stimulated HeLa cells the peak translocation of CAPRI was not significantly.

Figure 2. **CAPRI is refractory to Ca\(^{2+}\) oscillations.** (A) Ca\(^{2+}\) mobilization induced by application of 50 \(\mu\)M ATP in CHO cell lines. Black = parental CHO T cell response (\(n = 35\) cells); blue = CHO-CAP6 clone stably expressing CAPRI (\(n = 57\) cells); red = CHO-MUT13 clone stably expressing CAPRI (R473S) (\(n = 23\) cells). (B) Representative trace of the change in Fura-2 emission (black trace) and GFP-CAPRI translocation (blue trace) from a single cell displaying sinusoidal Ca\(^{2+}\) oscillations after 50 \(\mu\)M histamine. (C) Representative trace of the change in Fura-2 emission (black trace) and GFP-CAPRI translocation (blue trace) from a single cell displaying baseline Ca\(^{2+}\) spikes after 5 \(\mu\)M histamine. (D) Ca\(^{2+}\) mobilization in HeLa cells (black trace) after 50 \(\mu\)M histamine with sequential RFP-CAPRI translocation (blue trace, average \(n = 5\) experiments) (E) Reversibility of GFP-CAPRI translocation by histamine. Change in Fura-2 emission indicated in black, relative CAPRI translocation in blue. Cells stimulated with 50 \(\mu\)M histamine for 60 s, followed by wash-out of agonist, followed by restimulation for 60 s as shown (average \(n = 3\) cells). (F) Translocation of CAPRI-YFP and CFP-RASAL induced by 50 \(\mu\)M ATP in HeLa cells.
affected by the absence of Ca\(^{2+}\) entry (Fig. S2, A and B; available at http://www.jcb.org/cgi/content/full/jcb.200504167/DC1). However, it was clear that the rate of dissociation from the membrane was accelerated compared with controls. The response contrasted with oscillatory translocations of GFP-RASAL (Fig. S2 C) (Walker et al., 2004). Dissociation of CAPRI from the membrane under Ca\(^{2+}\)-free conditions took 230 s, significantly longer than the cytosolic Ca\(^{2+}\) signal drop (Fig. S2 B). This showed that the maintenance of CAPRI at the plasma membrane was only partially dependent on Ca\(^{2+}\) entry.

A significant lag time in dissociation compared with the decrease in cytosolic Ca\(^{2+}\) (which frequently oscillated) indicated that Ca\(^{2+}\)-independent interactions after Ca\(^{2+}\)-induced translocation were critical for maintenance at the plasmalemma.

Role of the tandem C2 domains and PH domain for membrane translocation

We showed previously that the C2A and C2B domains of CAPRI in tandem (C2AB) are necessary and sufficient for sensing an increase in cytosolic Ca\(^{2+}\) (Lockyer et al., 2001). GFP-C2AB has a tendency to concentrate in the nucleoplasm. As shown in Fig. 3 A, ionomycin caused the rapid translocation of GFP-C2AB to the nuclear membrane in a Ca\(^{2+}\)-dependent manner. Translocation to the plasma membrane was also detected in cells where cytosolic GFP-C2AB was sufficiently high. It seemed likely that electrostatic and lipid headgroup interactions were necessary for C2 domain–driven translocation, as is the case for other C2 domain–containing Ca\(^{2+}\) sensors (Rizo and Sudhof, 1998). In support of a role for electrostatic interactions in dictating membrane specificity, we noted that histamine failed to induce the nuclear translocation of GFP-C2AB but did cause translocation of cytosolic protein to the plasma membrane (Fig. 3 A). Ionomycin generates a sustained, high amplitude Ca\(^{2+}\) elevation, so it seems likely that the GFP-C2AB domain loses membrane specificity under chronically high Ca\(^{2+}\) conditions.

Do the C2AB domains mediate the long-term association with the plasmalemma? In HeLa cells expressing GFP-C2AB, histamine induced a transient association with the plasma membrane (Fig. 3 B). This showed that the GRD/PH/Btk domains were involved in mediating sustained translocation. We tested this further with chimeras of CAPRI and RASAL made by C2AB domain swapping. We found the CAPRI/RASAL chimera to be a sensitive tracker of repeti-
Figure 4. **TIRFM.** (A) GFP-CAPRI translocation induced by 100 μM histamine monitored by TIRFM (representative experiment for n = 4 cells). (B) GFP-RASAL translocation induced by 100 μM histamine monitored by TIRFM (representative experiment for n = 3 cells). (C) Representative experiment of GFP-CAPRI translocation in HeLa cells stimulated with 1 μM histamine (n = 3 cells). Arrows indicate mini-oscillations in one trace; similar oscillations are also detectable in other cells as shown. (D) Representative trace of sequential Fura-2 (black trace; single wavelength 380-nm excitation) and TIRF imaging of GFP-CAPRI (blue trace). Mini-oscillations correlate with cytosolic Ca^{2+} spikes.

Published data) or after ionomycin treatment as previously reported (Lockyer et al., 2001). It therefore seems likely that the PH domain provides additional binding energies to the membrane surface and accounts for Ca^{2+}-independent interactions with the plasmalemma. Possible ligands include inositol lipids (Lockyer et al., 1997, 1999) and Go subunits (Jiang et al., 1998). However, it is increasingly apparent that most PH domains are targeted to membranes by multiple factors; highly selective binding to a single phosphoinositide, or protein, is the exception to the rule (Lemmon, 2004). The CAPRI PH domain operates as a membrane-binding module in conjunction with the action of the C2 domains; the fact that recruitment is agonist dependent (Fig. 2 E) suggests that a signaling intermediate(s) is interacting with the PH domain at the membrane. The affinity must be insufficient to translocate ΔC2-CAPRI but is important when combined with Ca^{2+}-triggered, C2 domain–dependent membrane binding.

**Total internal reflection fluorescence microscopy detects CAPRI oscillations**

Total internal reflection fluorescence microscopy (TIRFM) offers superior observation of fluorescent molecules at the cell surface. Imaging GFP-CAPRI responses to histamine by TIRFM showed CAPRI translocation to be peak–plateau (Fig. 4 A; Video 1, available at http://www.jcb.org/cgi/content/full/jcb.200504167/DC1). The initial spike of translocation was more apparent than by Nipkow confocal imaging (Fig. 1). GFP-RASAL exhibited rapid oscillations in parallel experiments (Fig. 4 B; Video 2).

Conventional confocal or wide-field fluorescence microscopy had indicated that GFP-CAPRI translocations were inconsistent, and more transient, at low doses of histamine (Fig. 2 C). However, by TIRFM we observed 100% of GFP-CAPRI–expressing cells responding to 1 μM histamine (n = 15), highlighting the sensitivity of the technique. Interestingly, we could detect mini-oscillations superimposed on the plateau translocation response at early time points (Fig. 4 C), and by combining sequential wide-field imaging of Fura-2 (nonratiometrically) and RFP-CAPRI, we could show that mini-oscillations correlated with individual Ca^{2+} spikes (Fig. 4 D).

**Ca^{2+}-triggered Ras deactivation by CAPRI monitored in live cells**

In previous studies we were unable to reconstitute an in vitro GAP assay to monitor CAPRI function with temporal resolution (Lockyer et al., 2001). We therefore used the Ras-binding domain from Raf-1 tagged to GFP (GFP-RBD) as a reporter of active Ras on cell membranes (Walker and Lockyer, 2004). In a proportion of nonstarved cells cotransfected with H-Ras the RBD localized to the plasma membrane of HeLa cells (Fig. 5 A), or to the plasma membrane and Golgi apparatus of COS-7 cells (Fig. 5 B). By measuring the increase in cytosolic fluorescence as the RBD dissociated from the membrane (indicating Ras-GTP turnover), we determined the spatial and temporal kinetics of Ca^{2+}-triggered CAPRI GAP activity in transfected cells in a similar approach to our recent analysis of RASAL (Walker et al., 2004).
To ensure the GFP fusion protein of CAPRI retained GAP activity, we cotransfected HeLa cells with GFP-CAPRI and an HcRed version of the RBD (Fig. 5 A). In nonstarved cells the RBD reporter decorated the plasma membrane and was intensely concentrated in the nucleus (we have masked the nucleoplasm in Fig. 5 A for clarity). Histamine-evoked Ca\textsuperscript{2+} signals induced the translocation of GFP-CAPRI to the plasma membrane. This led to the synergistic dissociation of the HcRed-RBD, indicating Ras deactivation. Rather than use HcRed-RBD, which accumulates in the nucleus, we measured the kinetics of CAPRI-dependent Ras deactivation using GFP-RBD. This has been extensively characterized in COS cells for measuring Ras activation (Bondeva et al., 2002; Chiu et al., 2002). GAP function was tested by cotransfection with CAPRI, H-Ras, and GFP-RBD (Fig. 5 B). After ATP stimulation the GFP-RBD fluorescence was lost from membrane ruffles (but not the Golgi), leading to an increase in fluorescence in the cytosol (Fig. 5 B). At late time points it was apparent that Ras activity slowly recovered at the plasmalemma. This is best appreciated by viewing Video 3 (available at http://www.jcb.org/cgi/content/full/jcb.200504167/DC1). Reactivation of Ras may have been a consequence of residual serum factors inducing guanine nucleotide exchange factor stimulation because cells were not starved before imaging. Loss of fluorescence from the plasma membrane was not observed in cells cotransfected with a GAP-dead mutant of CAPRI (R473S) (Lockyer et al., 2001), or CAPRI with V12 H-Ras (unpublished data). The experiments confirmed that CAPRI has little basal GAP activity in resting cells and is acutely regulated by Ca\textsuperscript{2+} mobilization. The assays also showed that CAPRI specifically deactivates Ras at the plasma membrane.

We analyzed the kinetics of GFP-RBD dissociation from the plasma membrane in CAPRI-expressing HeLa or COS-7 cells stimulated with histamine or ATP, respectively. COS-7 cells not transfected with CAPRI (Fig. 5 C), or cotransfected with CAPRI R473S (unpublished data), showed no dissociation.
of the RBD from the membrane. HeLa cells stimulated with histamine displayed a more sustained deactivation of Ras (Fig. 5 D) than COS-7 cells stimulated with ATP (Fig. 5 C). The data match the kinetics of GFP-CAPRI translocation by correlative experiments in Fig. 5 (E and F) or directly as in Fig. S3 (available at http://www.jcb.org/cgi/content/full/jcb.200504167/DC1). CAPRI has no detectable GAP activity at resting Ca\(^{2+}\), suggesting it exists in a closed conformation in the cytosol. We believe that conformational changes must occur to facilitate the activation of Ras GAP function upon translocation. Using GFP-RBD and H-Ras to transiently transflect cells has allowed us to follow the sustained Ca\(^{2+}\)-triggered GAP activity of CAPRI in real time.

The fact that we can generate mini-oscillations of GFP-CAPRI is significant. It is likely that some GFP-CAPRI tracked cytosolic Ca\(^{2+}\) oscillations because membrane-binding sites were saturated. This indicates there is a threshold for plasma membrane ligands(s) that engages the PH domain. At higher intensity stimulation, excess ligand must be available to trap GFP-CAPRI, rendering the protein refractory to cytosolic Ca\(^{2+}\) fluctuations. Assuming that endogenous levels of CAPRI are lower than that achievable by overexpression, it seems highly unlikely that endogenous CAPRI oscillates during repetitive Ca\(^{2+}\) signals induced by G protein–coupled receptor activation because the membrane ligands for the PH domain would be in excess. This contrasts with RASAL, which is highly dependent on the C2 domains for membrane interactions according to our domain-swapping experiments. Thus, CAPRI is able to integrate the Ca\(^{2+}\)-triggered translocation with a second signal provided by the PH domain. Each Ca\(^{2+}\)-triggered GAP filters the Ca\(^{2+}\) signal differentially; this may convey alternative modes of information to regulate Ras and cell function.

### Materials and methods

#### Constructs

GFP-CAPRI and deletion constructs were made as previously described (Lockey et al., 2001). GFP-RASAL was made as previously described (Walker et al., 2004). GFP-RASAL HcRed-RBD was generated by PCR using GFP-RBD (human Ras-1 residues 51–131) as a template with primers containing HindIII and BamHI overhangs for cloning into pHcRed-C1 (BD Biosciences). RFP-CAPRI was generated by replacing eGFP with mRFP by PCR using monomeric RFP template. GFP-CAPRI (W664A) was generated by PCR using the Dplp mutagenesis method with primers 5′-GAAATGAGCT-3′ and 5′-GCAAGCGAGCAGACTGCGTATTAGCTCATTCT-3′. GFP-CAPRI/RASAL was generated by PCR to produce a chimera of Met-1 to Leu-275 of CAPRI and Leu-274 to Pro-804 of RASAL. GFP-RASAL/CAPRI was generated by PCR to produce a chimera of Met-1 to Leu-273 of RASAL and Leu-276 to Thr-803 of CAPRI.

#### Live cell imaging

For Ca\(^{2+}\) imaging experiments, cells were loaded with 1 μM Fura-2 AM or 1 μM Fluo-4 AM (Molecular Probes, Inc.) in KH buffer (10 mM Hepes, 118 mM NaCl, 4.7 mM KCl, 10 mM glucose, 1.2 mM KH\(_2\)PO\(_4\), 2.4 mM NaHCO\(_3\), 1.2 mM MgCl\(_2\), and 1.3 mM CaCl\(_2\), pH 7.4). Ca\(^{2+}\)-free KH buffer omitted CaCl\(_2\) for 0.5 mM EGTA. Imaging was performed using a microscope (Diaphot; Nikon) and a Fluor 40× 1.3 NA oil immersion objective (Nikon). Fluorescence was collected using an LSR Rainbow filter (Olympus) equipped with a M20 illumination system, dual-port illuminator, Hamamatsu Orca ER camera, and 488 nm/532 nm solid-state lasers. This was attached to a microscope (Olympus) with a Plan Fluor 40× 1.3 NA oil immersion objective. Dual GFP/HcRed, dual GFP/mRFP, and Fluo-4/mRFP analysis was performed on a confocal microscope (LSM 510 Meta; Carl Zeiss MicroImaging, Inc.). Fluorescence was collected using an LSR Rainbow filter (Olympus) equipped with a M20 illumination system, dual-port illuminator, Hamamatsu Orca ER camera, and 488 nm/532 nm solid-state lasers. This was attached to a microscope (Olympus) with a Plan Fluor 40× 1.3 NA oil immersion objective. Dual GFP/HcRed, dual GFP/mRFP, and Fluo-4/mRFP analysis was performed on a confocal microscope (LSM 510 Meta; Carl Zeiss MicroImaging, Inc.). Fig. 5 A was processed in Adobe Photoshop version 7 as detailed in the figure legend. All other images were processed in CorelDraw version 11 for brightness, contrast, and intensity.

The fluorescence intensity at each time point (F\(_t\)) and the maximum fluorescence intensity of the same area (F\(_{\text{max}}\)) was calculated. The Relative Translocation parameter F\(_t\)/F\(_{\text{max}}\) was calculated. The Relative Translocation parameter is F\(_{\text{max}}\)/F\(_t\). TIRFM was performed using a Biosystems cell Imaging Station (Olympus) equipped with an M20 illumination system, dual-port illuminator, Hamamatsu Orca ER camera, and 488 nm/532 nm solid-state lasers. This was attached to a microscope (Olympus) with a Plan Apo N 60×/1.45 NA oil TIRFM objective. Dual-port illumination enabled combined TIR and wide-field illumination for the fast and synchronized alternation of Fura-2 (380-nm nanometric measurement) and RFP imaging. The ratio of the fluorescence intensity at each time point (F\(_t\)) and the minimum fluorescence intensity of the same area (F\(_{\text{min}}\)) was calculated. The Relative Translocation parameter is 1 – F\(_{\text{min}}\)/F\(_t\).

#### Online supplemental material

Details of cell culture and transfection are available online at http://www.jcb.org/cgi/content/full/jcb.200504167/DC1.

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