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https://doi.org/10.15017/18383
Binding of phospholipase C-related but catalytically inactive protein to phosphatidylinositol 4,5-bisphosphate via the PH domain

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A R T I C L E   I N F O

Article history:
Received 20 February 2009
Accepted 9 March 2009
Available online 17 March 2009

Keywords:
Calcium
Inositol trisphosphate
Phosphoinositide
Phospholipase C
Pleckstrin homology domain

A B S T R A C T

A well-known protein module regulating molecular interactions is the pleckstrin homology (PH) domain whose best-characterised ligand is phosphoinositide. In the present study, we analysed the PH domain from PRIP (phospholipase C-related but catalytically inactive protein, comprising types 1 and 2) regarding phosphatidylinositol 4,5-bisphosphate [PtdIns(4,5)P2] binding employing a variety of binding assays. The PH domains prepared from PRIP-1 and -2 showed similar binding profiles to soluble ligands in vitro and showed similar plasma membrane localisation to that of PLC-δ1; however, the PH domain with the N-terminal extension of PRIP-1 but not PRIP-2 showed even distribution throughout the cytoplasm, indicating that the N-terminal extension of PRIP-1 inhibited binding to PtdIns(4,5)P2 present in the plasma membrane. A chimeric molecule of PLC-δ1 PH domain with the N-terminal extension of PRIP-1 exhibited similar localisation to PRIP-1 PH domain with the N-terminal extension. Binding assay to liposomes containing various concentrations of PtdIns(4,5)P2 revealed that the PH domain of PLC-δ1 bound steeply to the maximum, even at a concentration of 1.2 mol%, whereas the PH domains from PRIP-1 and -2 bound depending on the concentration up to 5 mol%. We also performed binding experiments using saponin-permeabilised PC12 cells. PH domains from PRIP increased the binding to cells preincubated with the brain cytosol extract in the presence of ATP, during which PtdIns(4,5)P2 were probably synthesised. The binding of PH domain with the following EF hand motifs showed Ca2+-dependent binding. These results indicate that the PH domain of PRIP binds to PtdIns(4,5)P2 present in the plasma membrane, depending on the concentrations of the lipid ligand and Ca2+, suggesting that PRIP might play physiological roles in events involved in the changes of these parameters, probably including Ins(1,4,5)P3.

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1. Introduction

Phospholipase C-related, but catalytically inactive protein (PRIP-1) was first identified in the brain cytosol fraction as a novel d-myoinositol 1,4,5-trisphosphate [Ins(1,4,5)P3] binding protein and was tentatively named p130 based on molecular size [1]. Subsequent molecular cloning studies revealed that the molecule is similar to phospholipase C-α1 but catalytically inactive, which is the reason for the revised name [2–5]. In an attempt to explore the biological function of PRIP-1 in relation to the binding to Ins(1,4,5)P3 via its pleckstrin homology (PH) domain [6,7], we performed serial experiments, first using COS-1 cells stably over-expressing PRIP-1 [8] and then cultured neurons prepared from PRIP-1 knock-out mice [9]. The results showed that both cells over-expressing and little-expressing PRIP-1 produced a reduced Ins(1,4,5)P3-mediated Ca2+ increase in cells, probably for different reasons [8,9], indicating that the presence of an appropriate amount of PRIP-1 is needed to produce right Ins(1,4,5)P3-mediated Ca2+ signalling.

PRIP-1 was also isolated from the membrane fraction of the brain with much the same molecular size as that from the cytosol fraction [5], indicating no lipid modification for localisation in the membrane fraction, but association to the membrane constituents. The most feasible constituent of molecules bearing PH domains would be phosphatidylinositol 4,5-bisphosphate [PtdIns(4,5)P2], which is a minor but key phospholipid mainly on the cytoplasmic leaflet of the plasma membrane. We then performed cellular experiments using COS-1 cells transfected with genes for a variety of deletion mutants of PRIP-1 to examine whether the PH domain is implicated in membrane localisation by binding to PtdIns(4,5)P2, and drew the tentative conclusion that the membrane association of PRIP-1 indeed occurred, but the association via the binding of the PH domain to PtdIns(4,5)P2 was partial [10]. Subsequently, in vitro experiments using the liposomes containing PtdIns(4,5)P2 clearly indicated that the PH domain of PRIP-1 binds PtdIns(4,5)P2, and further Ca2+ at physiological concentration enhances the binding when the recombinant PH domain molecule used contains EF hand motifs, like those of PLC-δ [11,12].
We also observed that full-length PRIP-1 was mainly localised in the cytoplasm [6,8], but the isolated PH domain (amino acid residues 82–298) was mainly seen at the surface membrane [8,13]. We further examined the recombinant molecule comprising N-terminal extension (residues 24–81) plus the PH domain regarding localisation, resulting in the main location in the cytoplasm. These results indicate that the N-terminal region of PRIP-1 preceding the PH domain would prohibit it from associating with the surface membrane. Furthermore, an isoform of PRIP-1 with relatively broad tissue distribution, including the brain, was also reported [14,15], indicating that PRIP comprises types 1 and 2.

PtIns(4,5)P2 has been receiving constant attention because it regulates a wide variety of processes, including exocytic and endocytic membrane trafficking [16–20], ion channel and transporter function [21], enzyme activation [22], and protein recruitment [23–26]. In order to extend our experiments in exploring the biological function of PRIP, probably through competing for PtIns(4,5)P2 with other molecules involved in the functions described above, it would be important to reevaluate whether the PH domain is involved in the membrane association of PRIP via binding to PtIns(4,5)P2.

In the present study, we examined the cellular localisation and binding to PtIns(4,5)P2-containing liposome of the PH domains derived from PRIP-2 as well as PRIP-1, and further evaluated the effect of N-terminal extension on the localisation of the PH domain. Finally, we examined whether the association is indeed regulated by PtIns(4,5)P2 via binding by the PH domain, employing semi-intact cellular experiments.

2. Materials and methods

2.1. Materials

[1H]Ins(1,4,5)P3 (specific radioactivity: 8510 Ci/mmol) was purchased from PerkinElmer (Waltham, MA). Ins(1,4,5)P3 and short-chain (C8, water-soluble) PtIns(4,5)P2 and natural PtIns(4,5)P2 were obtained from Cell Signal Inc. (Lexington, KY). Glutathione-Sepharose 4B and pGEX vectors were from GE Healthcare (Uppsala, Sweden). pEGFP-N1 vector was from Clontech (Palo Alto, CA). Phosphatidylincholine (PC) was purchased from Sigma (St. Louis, MO). Monoclonal antibody against GST was from Santa Cruz Biotechnology (Santa Cruz, CA). Other reagents used were of the highest grade available.

2.2. DNA constructs, recombinant protein expression and purification

The PH domain of human PLC-δ1 (residues 1–140), and its mutant R40L was described previously [27,28]. The long version of PH domain (1–298) of PRIP-1 (1PHL) was amplified from full-length PRIP-1 construct using the primers 5′-TAAAGGCACTATCATGAGGGCGCGG-3′ and 5′-ATGTCGACCGCAGGTAGTATTGATTTTCTTGGC-3′. The short version (1PHS, 74–298) was amplified using the primers 5′-TAGAATTCGACCCAACATTGGAGGGCGGCGG-3′ and 5′-ATGTCGACCGCCAGCACTAC-3′ and 5′-ATGTCGACCGCTAGTATTGATTTTCTTGGC-3′. The long version of the PH domain (1–325) of PRIP-2 (2PHL) was amplified from full-length PRIP-2 using the primers 5′-TAAAGGCACTATCATGAGGGCGCGG-3′ and 5′-ATGTCGACCGCAGGTAGTATTGATTTTCTTGGC-3′. The short version (2PHS, 102–325) was amplified using the primers 5′-TAGAATTCGACCCAACATTGGAGGGCGGCGG-3′ and 5′-ATGTCGACCGCTAGTATTGATTTTCTTGGC-3′. After the digestion with EcoRI and SalI, the product was ligated into the pEGFP-N1 plasmid (CLONTECH) and pGEX-4T3 plasmid digested with the same enzymes. The chimeric molecule of the PLC-δ1-PH domain (1N-6PH) was created by connecting the N-terminal PRIP-1 (1–81 residues) with PLC-δ1PH (1–142 residues). The mutant form of 1PHS (1PHSm) was also generated in plasmids pGEX-4T3 and pEGFP-N1 by site-directed mutagenesis (Quick Change; Stratagene) using the primers 5′-CTCTCAGCATCTACAATCAATTTTCTACCCATGACAC-3′ and 5′-GTGTCTGAGGGTTTCTACCCATGACAC-3′ for R134Q.

For purification of recombinant PH domains conjugated with GST, Escherichia coli BL-21 (DE3) was transformed with pGEX-4T3 constructs. The bacterial cells were grown up to 0.4 of absorbance at 600 nm at 37 °C and then with 250 μM isopropyl -β-D-thiogalactopyranoside (IPTG) at 18–20 °C for an additional 12–14 h. Bacterial lysate was prepared by sonication in a lysis buffer [50 mM Tris–HCl (pH 8.0), 300 mM NaCl, 1 mM EDTA, 1 mM DTT and protease inhibitor cocktail containing 5 μg/ml pepstatin A, 10 μM leupeptin, 1.7 μg/ml aprotinin, and 50 μM 4-amidinophenylmethanesulfonyl fluoride hydrochloride], followed by rotation after the addition of 1% Triton X-100 for 20 min. Purification was achieved using Glutathione-Sepharose 4B beads. After extensive washing with a lysis buffer without protease inhibitor cocktail, the proteins were eluted with 20 mM reduced glutathione in a lysis buffer without protease inhibitor cocktail, but containing 1 mM DTT. Purity was checked by staining with Coomassie Brilliant Blue (CBB) in SDS-polyacrylamide gel electrophoresis (SDS-PAGE). The recombinant molecules of interest were dialyzed against the solution for assay for more than 6 h, followed by centrifugation at 100,000 g for 30 min before use.

2.3. Binding assay of recombinant proteins to [3H]Ins(1,4,5)P3

The binding to [3H]Ins(1,4,5)P3 (0.9 nM; 0.37 kBq radioactivity) of each recombinant protein (8 pmol) was performed in a reaction mixture (0.4 ml) containing 50 mM Tris–HCl (pH 8.3), 0.2% Triton X-100, and 2 mM EGTA. The reaction mixture was incubated on ice for 15 min, followed by the addition of 50 μl of 10 mg/ml bovine g-globulin and 0.5 ml of 30% (w/v) polyethylene glycol 6000. The precipitate formed after centrifugation at 15,000 rpm for 5 min was dissolved in 0.5 ml of 0.1 N NaOH and then counted for radioactivity as an emulsion with 5 ml scintillation cocktail. Non-specific binding in the presence of 100 μM unlabelled Ins(1,4,5)P3 was also measured (approximately 200–250 dpm) and subtracted from the values measured in its absence (range 3500–5800 dpm), enabling calculation of the specific binding.

2.4. Binding assay of recombinant proteins to liposomes

Liposomes composed of phosphatidylcholine (PC) or with PtIns(4,5)P2 were made according to the method described [29]. Briefly, PC alone or with PtIns(4,5)P2 at different concentrations dissolved in chloroform solution was dried under nitrogen to make a lipid film in a glass tube. A solution (buffer A) containing 25 mM Hepes/NaOH buffer (pH 7.4), 100 mM NaCl and 1 mM EDTA was added to make a final lipid concentration of 20 mM and the mixture was sonicated twice until uniform turbidity was achieved so that small unilamellar vesicles were formed. Stock liposomes thus made were kept in a refrigerator until use (for approximately a month). The phospholipid vesicles (final conc. of total lipid was 1 mM) were mixed with purified recombinant proteins in 50 μl buffer B [20 mM Tris–HCl (pH 7.5), 150 mM NaCl, 1 mM EDTA and 1 mM DTT]. The mixture was incubated on ice for 10 min, followed by centrifugation at 100,000 g for 30 min. The supernatant was carefully removed, and mixed with 5× concentrated sample buffer for SDS-PAGE to a final volume of 50 μl. The pellet was dissolved in SDS buffer up to the same volume (50 μl). Both the supernatant and the pellet were analysed by SDS-PAGE, followed by staining with CBB. Densitometric analyses were performed on a film obtained with an NIH image analyser. Bovine serum albumin was used as the standard protein to assess the linearity of protein concentration after staining with “Quick CBB” (Wako Pure Chemical Co. Ltd., Tokyo, Japan) for 30 min.

2.5. Binding assay of recombinant proteins to permeabilised PC12 cells

The binding assay of recombinant proteins to permeabilised PC12 cells was performed as described [30]. PC12 cells were permeabilised
using 0.05% saponin on ice for 15 min, followed by vigorous washing with KGl buffer [20 mM Hepes/KOH (pH 7.2), 120 mM potassium glutamate, 20 mM potassium acetate, 2 mM EGTA]. Cells suspended in a KGl buffer (in the absence of Mg-ATP) were left at room temperature for 20 min for consistent results. Permeabilised PC12 cells were then incubated with rat brain extract (0.5 mg/ml) prepared as described [31] in the presence of Mg-ATP at 30 °C for 10 min, and then washed once briefly. The cells (8 x 10⁵ cells) were incubated with the recombinant molecule (GST-conjugated) of interest for 30 min on ice in a total volume of 200 μl, followed by separation of the mixture by rapid centrifugation (30 s, 21,600 g) through 0.0625% sucrose in KGl buffer (200 μl). Supernatants were carefully withdrawn, tubes were wiped carefully, and cell pellets were resuspended in SDS sample buffer, followed by separation by SDS-GAGE, transfer to a polyvinylidene fluoride membrane and analysis by Western blotting using anti-GST antibody for the experiments and anti-β-actin antibody as a reference.

2.6. Transfection of cells for confocal microscopy

Madin–Darby canine kidney (MDCK) cells were plated onto 14-mm diameter circular glass coverslips at a density of 4 x 10⁴ cells/well in a 12-well plate and cultured in Dulbecco’s modified Eagle’s medium (DMEM) supplemented with 10% (v/v) fetal bovine serum for a day. The cells were then transiently transfected with plasmid DNAs (1 μg) using Lipofectamine2000 and OPTI-MEM (Invitrogen) according to the manufacturer’s protocol. Twenty-four hours after transfection, the cells were fixed with 4% paraformaldehyde and observed by a confocal laser scanning microscope (Zeiss: LSM510-Meta).

3. Results and discussion

3.1. Constructs prepared from PRIP-1 and -2, and their cellular localisation

Fig. 1A is a schematic representation of the constructs used in the present study. A long version of the PH domain from PRIP-1 (amino acid residue 1–298, abbreviated as 1PHL) and -2 (amino acid residue 1–325, 2PHL) contained the N-terminal extension preceding the PH domain, while the short version (amino acid residues 74–298 and 102–325 for 1PHS and 2PHS, respectively) lacked it. The PH domain of PLC-δ1 (δPH, amino acid residue 1–142) was employed as a reference since it is well recognised to localise at the surface membrane by direct binding to PtdIns(4,5)P₂ [27,32–34]. In this case, to test whether the N-terminal extension preceding the PH domain seen in the PRIP family can influence the localisation of the PH domain, a chimera was also constructed incorporating the N-terminal extension from PRIP-1 (1N-δPH). All constructs were conjugated with EGFP (enhanced green fluorescent protein) for mammalian cells or GST (glutathione S-transferase) for the E. coli expression system at the C-terminus for visualisation or binding assay, respectively. Fig. 1B shows the CBB staining pattern of purified samples, indicating that the gel also contained the mutant of 1PHS in which R134 was mutated to Q (1PHSm), which corresponds to critical R40 in PLC-δ1 [27,34]. Fig. 1C depicts the domain organisation of PRIP and PLC-δ1 for references.

We first examined the binding properties of these molecules isolated by the E. coli expression system to Ins(1,4,5)P₃ and PtdIns(4,5)P₂ in a solution. Each molecule examined, except for 1PHS, exhibited binding ability to [³H]Ins(1,4,5)P₃ which was displaced by unlabelled Ins(1,4,5)P₃ and water-soluble (short-chain, C8)-PtdIns(4,5)P₂ in a dose-dependent manner (Fig. 2). The values of IC₅₀ indicate the gross binding affinity; those of Ins(1,4,5)P₃ and PtdIns(4,5)P₂ were as follows in nM: 1PHS (40 and 110), 1PHL (30 and 90) or 2PHL (40 and 90) and 2PHS (40 and 90), respectively. Collectively, the affinity to Ins(1,4,5)P₃ appeared to be about 3-fold higher than to PtdIns(4,5)P₂, but there was little difference between short and long versions, and between types 1 and 2.

MDCK cells were transfected with expression plasmids to examine the cellular localisation of these PH domains. As shown in Fig. 3, molecules 1PHS, 2PHS and 2PHL fused with GFP were mainly found close to the surface membrane, the pattern of which is similar to that of 1PHL and -2.
observed with δPH, whereas molecule 1PHL were mainly found in the cytoplasm. 1PHSm also showed a similar pattern to 1PHL (results not shown). A distinct difference between the long and short versions was thus observed with the PH domain of PRIP-1. These observations indicate that the binding affinities of these PH domains assessed by displacement of bound [3H]Ins(1,4,5)P₃ by Ins(1,4,5)P₃ and PtdIns(4,5)P₂ in a solution do not necessarily explain the difference in cellular localisation.

The N-terminal region composed of residues 1–73 from PRIP-1, but not the corresponding region (residue 1–101) of PRIP-2, seems to cause the difference by prohibiting the PH domain (74–298) from coming close to the surface membrane. To examine whether this is applicable to δPH, which binds both PtdIns(4,5)P₂ and Ins(1,4,5)P₃ with similar affinity (see Fig. 4B) [1,6,27], the chimera abbreviated as 1N-δPH in Fig. 1A was expressed in MDCK cells. As shown in Fig. 4A, cytoplasmic fluorescence in a diffuse fashion appeared to be increased by incorporating the N-terminal extension of PRIP-1. This does not appear to be caused by changes in binding properties to the ligands in a solution; the displacement profile of bound [³H]Ins(1,4,5)P₃ by Ins(1,4,5)P₃ and short-chain (C₈)-PtdIns(4,5)P₂ was not different between δPH and 1N-δPH (Fig. 4B), indicating that N-terminal extension does not influence the binding profile to soluble ligands, but might provide steric hindrance or physically interact with other molecules inside cells, thus leading to the inhibition of δPH to bind PtdIns(4,5)P₂ present in the surface membrane. Similar results were observed when a chimera abbreviated as 1N-2PHS (N-terminal extension of PRIP-1 connected to type 2 PH short version) was used (results not shown).

The appearance of the isolated PH domain in intact cells represented by 1PHS and 2PHL, shown in Fig. 3, seemed to be attributed to the direct association with PtdIns(4,5)P₂ present at the inner leaflet of the surface membrane because 1PHSm exhibited cytoplasmic localisation, and the binding to reconstituted liposomes containing PtdIns(4,5)P₂ increased as the concentration increased, as shown in Fig. 5. These observations were similar to those observed with δPH, but the affinity of δPH to PtdIns(4,5)P₂ present in liposomes appeared

![Fig. 2. Binding of PH domains to [³H]Ins(1,4,5)P₃ and the displacement by Ins(1,4,5)P₃ and PtdIns(4,5)P₂. [³H]Ins(1,4,5)P₃ bound to PH domain was displaced by various concentrations of unlabelled Ins(1,4,5)P₃ (solid lines) or C₈ (water-soluble) PtdIns(4,5)P₂ (dotted lines). All recombinant molecules assayed ((A) 1PHS, 1PHL and (B) 2PHS, 2PHL, see Fig. 1A) were conjugated with GST for purification [see Fig. 1B for purity]. Thin dotted line indicates 50% displacement. Results are shown as the mean ± SE of four independent experiments.](image)

![Fig. 3. Subcellular localisation of PH domains fused with GFP in MDCK cells. (A) MDCK cells grown on a glass coverslip were transiently transfected with each construct fused with GFP, followed by fixation with 4% paraformaldehyde. Images were obtained using a confocal laser scanning microscope (Zeiss: LSM510-Meta) and are typical images from more than 20 independent experiments. (B) GFP-fused PH domains expressed in MDCK cells were analysed for a lack of severe hydrolysis inside cells by Western blotting using anti-GFP antibody.](image)
membrane was detached toward the cytosol in response to cellular containing PtdIns(4,5)P2, to a similar extent as 1PHS and 2PHL. localised mainly inside cells was also capable of binding to liposomes liposomes was steep in clear contrast to the shallow pro

by varying the concentration of Ins(1,4,5)P3, we observed that 1PHL (4,5)P2 (probably the concentration at even distribution, see below) inhibition of 1PHS and 1PHL to liposomes containing 1.2 mol% PtdIns

domain of PRIP-1, but not PRIP-2, provides some interference in the

inhibition was more sensitive to increasing concentrations of Ins(1,4,5)P3 (solid lines) and C8-PtdIns(4,5)P2 (dotted lines) of [3H]Ins(1,4,5)P3 bound to δPH or 1N-δPH. For the purity of the protein samples used, see Fig. 1B. Thin dotted line indicates 50% displacement. Results are the mean of two independent experiments.

to be higher than the other PH domains (PH domains from PRIP), since the binding profile to increasing concentrations of PtdIns(4,5)P2 in liposomes was steep in clear contrast to the shallow profiles observed with the other PH domains (Fig. 5B). It was noteworthy that 1PHL localised mainly inside cells was also capable of binding to liposomes containing PtdIns(4,5)P2, to a similar extent as 1PHS and 2PHL. Furthermore, 1PHSm induced decreased binding to liposomes containing an increasing concentration of PtdIns(4,5)P2, but still showing binding over non-specific binding levels (Fig. 5B). This can probably be attributed to electrostatic interactions between anionic phospholipids [PtdIns(4,5)P2] and basic amino acids (R131, R132 and K133) remaining in the mutant. These results probably indicate that the PH domains of PRIP-1 and -2 would directly associate with PtdIns(4,5)P2 present in reconstituted lipid bilayers, and in the cellular surface membrane, such as δPH, and further, together with those of Figs. 2 and 3, indicate that the N-terminal extension preceding the PH domain of PRIP-1, but not PRIP-2, provides some interference in the association under cellular conditions. When we compared the binding inhibition of 1PHS and 1PHL to liposomes containing 1.2 mol% PtdIns (4,5)P2 (probably the concentration at even distribution, see below) by varying the concentration of Ins(1,4,5)P3, we observed that 1PHL was more sensitive to increasing concentrations of Ins(1,4,5)P3 (Supplementary Fig. 1). This observation might partly explain the difference in cellular localisation, despite little difference in binding profiles of solutions and liposomes. PRIP-1 under cellular physiological conditions might be regulated in its membrane association by a subtle change in the concentration of Ins(1,4,5)P3.

In the previous experiment [13], however, we observed a distinct difference between δPH and 1PHS; δPH found at the surface membrane was detached toward the cytosol in response to cellular stimulation, probably by decreasing PtdIns(4,5)P2 concentration caused by hydrolysis, whereas 1PHS remained present around the plasma membrane even upon cellular stimulation. We here again observed that 1PHS, 2PHS and 2PHL were hardly detached from the surface membrane of MDCK cells upon cellular stimulation with ATP, a purinergic agonist (data not shown). Ins(1,4,5)P3 production in HeLa cells stimulated with ATP was measured, with the result that the production of Ins(1,4,5)P3 was not inhibited by 1PHS, but was inhibited in HeLa cells expressing δPH, probably because PtdIns(4,5)P2 complexed with the PH domain could not be a good substrate for hydrolysis [13]. From these results we had tentatively assumed the possibility that 1PHS seen at the plasma membrane is only present just beneath the plasma membrane without direct binding to PtdIns (4,5)P2. However, this possibility might have to be amended, as the PH domain of PRIP-1 and -2 directly associates with PtdIns(4,5)P2 present in liposomes and the cell surface membrane, as shown in Figs. 3 and 5, although the affinity is apparently lower than that of δPH, as seen in Fig. 5B. The concentration of PtdIns(4,5)P2 has been estimated to be 30–160 μM depending on cell types under the assumption of even distribution [35–37] or ~1 mol% (in a different unit) for total plasma membrane of PC12 cells with asymmetrical leaflet distribution [35–37], but that in the microdomain for dense-core vesicle docking, for instance, was estimated to be ~6 mol% equivalent up to ~1 mM [38]. Taken together, we might assume that the PH domains from PRIP-1 and -2 with lower affinity to PtdIns(4,5)P2 bind to the surface membrane restricted to sites abundant with the lipid, while that from PLC-δ1 could bind PtdIns(4,5)P2 present anywhere in the surface membrane, which could be a good substrate for PLC, since the reported Km value is around 50–100 μM [32,33].

3.2. Membrane association of PH domain using permeabilised cells

PC12 cells have been widely used for the exocytosis assay of dense-core vesicles containing norepinephrine, the process of which requires
the preceding synthesis of PtdIns(4,5)P\textsubscript{2} to enable vesicles and plasma membranes to develop competence for Ca\textsuperscript{2+}-triggered fusion [31,39–43]. So here we employed PC12 cells permeabilised with saponin for analysis. Permeabilised cells were preincubated with the brain cytosol extract for 10 min at 30 °C in the presence of Mg-ATP, during which PtdIns(4,5)P\textsubscript{2} were probably synthesised [39,40], and then washed once briefly, followed by incubation with recombinant molecules of interest at 50 nM for 10 min on ice, and separation by centrifugation and analysis of bound molecules by Western blotting. Blots are typical of 3 independent experiments, and the graph below shows the summary, represented as relative to the binding with no soluble ligand. 2PHL gave similar results to that of 1PHS. (C) Effect of the treatment with PLC-δ1. Permeabilised PC12 cells preincubated with brain extract plus ATP were incubated with recombinant PLC-δ1 (0.1 mg/ml) at 30 °C for 10 min at free Ca\textsuperscript{2+} concentration of 1 μM (indicated as "Ca\textsuperscript{2+} +") or less than 0.01 μM (indicated as "Ca\textsuperscript{2+} −"). After washing, binding assays were performed as described above. Blots are the typical of three independent experiments, and the graph shows the summary, represented as relative to the binding with no treatment. *: p<0.05.
was examined. The constructs for PH domain prepared in this study contained EF hand motifs (see Fig. 1A and B). 1PHS was mainly used here. As shown in Fig. 7, the binding of the PH domain with EF hand motifs (1PHS) increased in a Ca²⁺-dependent manner, while δPH lacking the following EF hand motifs showed no Ca²⁺-dependent increase of binding, results similar to those observed using liposomes containing PtdIns(4,5)P₂. The impact of the Ca²⁺ effect appeared bigger under the unprimed conditions, but was still observed under primed conditions, indicating the physiological relevance of this Ca²⁺ effect.

Acknowledgements

This work was funded by Grants-in-Aid for Scientific Research from the Ministry of Education, Culture, Sports, Science and Technology of Japan (to HT, TK and MH), The Uehara Memorial Foundation (to MH) and The Japan Diabetes Association (to TK). JG is a recipient of the Iwadare Scholarship and MF is a research fellow from the Ministry of Education, Culture, Sports, Science and Technology of Japan.

Appendix A. Supplementary data

Supplementary data associated with this article can be found, in the online version, at doi:10.1016/j.cellsig.2009.03.008.

References

Supplementary Figure 1. Gao et al
Supplementary Figure 2. Gao et al