

Phospholipase D is activated and phosphorylated by casein kinase-II in human U87 astrogloma cells

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Abbreviations: CKII, casein kinase II; PKC, protein kinase C; PLD, phospholipase D; PtdBut, phosphatidylbutanol

Abstract

Elevated expression of protein casein kinase II (CKII) stimulated basal phospholipase D (PLD) activity as well as PMA-induced PLD activation in human U87 astrogloma cells. Moreover, CKII-selective inhibitor, emodin and apigenin suppressed PMA-induced PLD activation in a dose-dependent manner as well as basal PLD activity, suggesting the involvement of CKII in the activation of both PLD1 and PLD2. CKII was associated with PLD1 and PLD2 in co-transfection experiments. Furthermore, CKII induced serine/threonine phosphorylation of PLD2 *in vivo*, and the multiple regions of PLD2 were phosphorylated by CKII *in vitro* kinase assay using glutathione S-transferase-PLD2 fusion protein fragments. Elevated expression of CKII or PLD increased cell proliferation

but pretreatment of cells with 1-butanol suppressed CKII-induced cell proliferation. These results suggest that CKII is involved in proliferation of U87 cells at least in part, through stimulation of PLD activity.

Keywords: casein kinase II; cell proliferation; phospholipase D; phosphorylation

Introduction

Protein kinase CKII is a ubiquitous and highly conserved Ser/Thr kinase, found in various subcellular compartments in eukaryotes. It exists as a constitutively active tetramer that contains two catalytic subunits, α or α' (37-44 kDa), and two regulatory β subunits (24-28 kDa) (Allende and Allende, 1995; Pinna and Meggio, 1997; Litchfield, 2002). The two catalytic subunits are highly homologous, but the α' subunit has a unique required role in spermatogenesis (Xu *et al.*, 1999). CKII phosphorylates serine or threonine in acidic domains, with (S/T)XX(D/E) being the canonical motif (Hrubey and Roach, 1990; Marin *et al.*, 1992). CKII was postulated to contribute to tumorigenesis because its activity is enriched in many human solid tumors, transformed cell lines, and rapidly proliferating tissues (Ahmed, 1994; Daya-Makin *et al.*, 1994). Dysregulated expression of CKII in cells can be oncogenic, as transgenic expression of CKII α can promote lymphoma (Seldin and Leder, 1995; Kelliher *et al.*, 1998; Landesman-Bollag *et al.*, 1998) and breast cancer (Landesman-Bollag *et al.*, 2001). In addition, it has been demonstrated that CKII catalyzes the phosphorylation of several oncogene products such as Myc, Myb, Jun, Fos, p53, and Rb (Pinna, 1990; Issinger, 1993; Allende and Allende, 1995). These observations suggest that CKII plays a critical role in cell proliferation and oncogenesis.

Phospholipase D (PLD) catalyzes the hydrolysis of phosphatidylcholine (PC), the major membrane phospholipids, to form phosphatidic acid (PA) and choline (Exton, 1998). PA can alter the activities of many enzymes and proteins (Andresen *et al.*, 2002) and can be further metabolized to diacylglycerol (DAG) and lysophosphatidic acid by PA phosphohydrolase and phospholipase A2, respectively. Thus, PLD influences many important intracellular events via producing these downstream products. PLD activity is regulated by many stimuli such as cytokine, growth factors, hormones, neurotrans-

mitters, and other molecules involved in extracellular communication (Exton, 1998). PA is generally recognized as the signaling product of PLD and functions as an effector in multiple physiological process including cell proliferation, differentiation, secretion, and migration. In mammals, two isoforms of PLD, PLD1 and PLD2, have been cloned and are being characterized for regulation and cellular function (Frohman and Morris, 1999). However, segregated roles of the two PLD isoforms in cellular responses are still poorly understood. Activation of PLD occurs through interactions of the ARF and Rho families as well as protein kinase C (PKC) (Frohman and Morris, 1999). The relative contribution of these factors to PLD activation is highly dependent on the cell type and signaling model examined. Consistent with a complex set of regulatory requirements, PLD appears to contain several different post-translational modifications. We have reported that PLD can be phosphorylated on tyrosine residues by c-Src and the combination of H₂O₂ and vanadate (Min *et al.*, 1998; Ahn *et al.*, 2003; Kim *et al.*, 2004b). In addition, PLD1 is phosphorylated on serine and threonine residue by PKC (Min *et al.*, 1998; Kim *et al.*, 2000). In the present study, we report that CKII is involved in PMA-induced PLD1 and PLD2 activation, phosphorylates PLD2 and interacts with both PLD1 and PLD2. Our results suggest that CKII is involved in proliferation of U87 cells at least in part, through stimulation of PLD activity

Materials and Methods

Reagents

Glutathione-sepharose 4B was from Amersham Bioscience Biotech. Anti-phospho-ser/thr antibody was from Zymed. The antibody to CKII- α was from Santa Cruz. Emodin, apigenin and PMA were from Sigma. Calphostin C was purchased from Biomol Research Laboratories (Plymouth Meeting, PA). A polyclonal antibody that recognizes both PLD1 and PLD2 was generated as previously described (Min *et al.*, 2001). Phosphatidylbutanol (PtdBut) standard was from Avanti Polar Lipid. [9, 10-³H]myristate was purchased from PerkinElmer Life Sciences. Silica gel 60 A thin layer chromatography plates were from Whatman. Horseradish peroxidase-conjugated anti-mouse IgG and anti-rabbit IgG were from Kirkegaard & Perry Lab (Gaithersburg, MD). The ECL Western blotting detection kit was from Amersham Bioscience. Human CKII was purified homogeneity via four chromatography steps using DEAE-cellulose, phosphocellulose, heparin-agarose, and gel filtration from the lysates of *E. Coli* that expressed both α and β subunits of CKII (Kim *et al.*, 1998).

Cell culture and transfections

U87 MG human astrogloma was obtained from the American Type Culture Collection (Rockville, MD). The cells were maintained in DMEM (Life Technologies, Inc) supplemented with 10% (v/v) fetal bovine serum under 5% CO₂. U87 cells were transiently transfected for 40 h with expression plasmid using LipofectAMINE (Invitrogen) according to manufacturer's instructions. U87 cells stably overexpressing PLD isozyme were obtained by transfection, using LipofectAMINE. Transfected cells were selected with G418 (700 μ g/ml) for 21 days at 37°C. At that time antibiotics-resistant colonies were pooled and expanded for further analysis under selective conditions.

PLD activity assay

PLD activity was assessed by measuring the formation of [³H] phosphatidylbutanol (PtdBut), the product of PLD-mediated transphosphatidylation, in the presence of 1-butanol. After 20 h transfection with LipofectAMINE, U87 cells in 6-well plates were serum-starved in the presence of 1 μ Ci/ml [³H]myristic acid. After overnight starvation, the cells were washed three times with 5 ml of phosphate-buffered saline (PBS) and pre-equilibrated in serum-free DMEM for 1 h. For the final 10 min of preincubation, 0.3% butan-1-ol was included. At the end of the preincubation, cells were treated with agonists for the indicated times. The extraction and characterization of lipids by thin-layer chromatography were performed as previously described (Kim *et al.*, 2004a).

Construction and preparation of GST fusion proteins

GST fusion proteins of PLD2 fragments were constructed as described previously (Kim *et al.*, 2002). The PCR products were digested with the restriction enzymes and inserted into pGEX-4T1 vector (Amersham Pharmacia Biotech). *E. coli* BL21 cells were transformed with the individual expression vectors encoding the GST fusion protein, and after harvesting the cells, GST fusion proteins of PLD2 fragments were purified as described previously (Kim *et al.*, 2002).

Immunoprecipitation

Cells were washed twice with ice-cold phosphate-buffered saline and then lysed in the extraction buffer (20 mM Hepes, pH 7.2, 1% Triton X-100, 1% sodium deoxycholate, 0.2% SDS, 200 mM NaCl, 1 mM Na₃VO₄, 1 mM NaF, 10% glycerol, 10 μ g/ml leupeptin, 10 μ g/ml aprotinin, 1 mM phenylmethylsulfonyl fluoride). The resulting cell lysates was spun at 15,000 $\times g$ in a Eppendorf microcentrifuge for 10 min at 4°C to pellet the unbroken cells. The supernatant

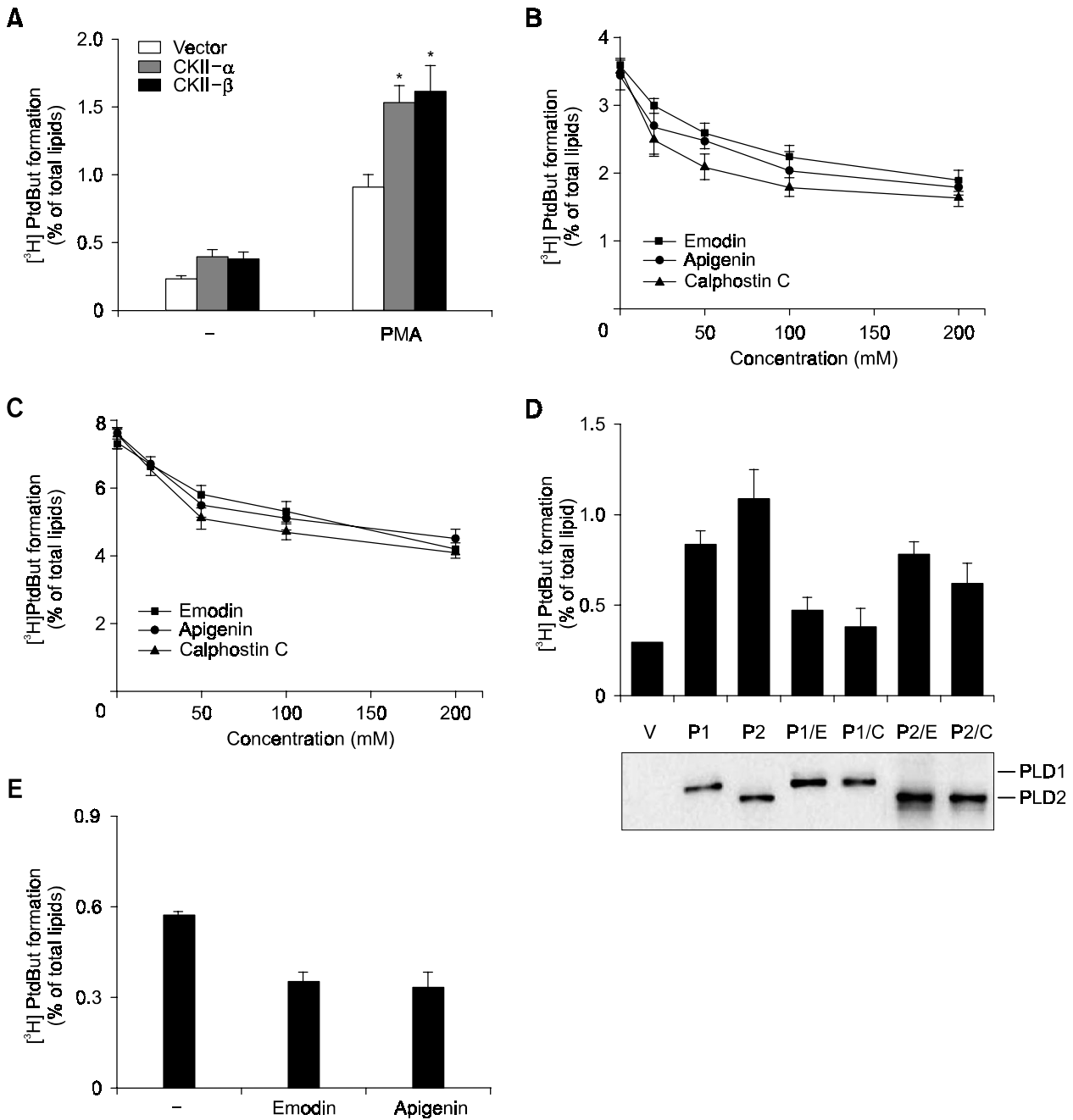


Figure 1 CKII is involved in PMA-induced PLD activation. (A) U87 cells were transiently transfected with CKII- α or β , labeled with [³H]myristic acid, and treated with 100 nM PMA for 1 h. **P* < 0.05 compared to cells transfected with vector and treated with PMA. U87 cells overexpressing PLD1 (B) and PLD2 (C) were labeled with [³H]myristic acid, preincubated with the indicated concentrations of emodin, apigenin, or calphostin C and stimulated with PMA for 1h. (D) U87 cells over-expressing PLD1 (P1), PLD2 (P2) or vector were treated with or without emodin (E, 100 μ M) and calphostin C (C, 100 μ M). PLD expression was analyzed by Western blot using antibody to PLD which recognizes both PLD1 and PLD2. (E) U87 cells were incubated with or without emodin (100 μ M) and apigenin (100 μ M). The radioactivity incorporated into phosphatidylbutanol was measured as described in Materials and Methods. Results are means \pm S.D of three independent experiments.

was then precleared for 30 min with preimmune IgG and protein A sepharose at 4°C with rocking. Protein concentrations were determined using Bradford method with bovine serum albumin as a standard (4).

Equal protein aliquots of precleared cell lysates (1 mg) were incubated with the indicated antibodies and 40 μ l of a 1:1 slurry of protein A sepharose beads for 4 h at 4°C. The immune complexes were

collected by centrifugation and washed five times with a buffer (20 mM Tris, pH 7.5, 1 mM EDTA, 1 mM EGTA, 150 mM NaCl, 2 mM Na₃VO₄, 10% glycerol and 1% Nonidet P-40) and resuspended in sample buffer. Immune complexes were boiled in SDS-sample buffer.

Western blotting

Proteins samples were analyzed by SDS-polyacrylamide gel electrophoresis and were transferred to a nitrocellulose membrane. The blots were then blocked with 5% non-fat milk and incubated with appropriate primary antibodies followed by incubation with horseradish peroxidase-conjugated secondary antibody. Immunoreactive bands were detected using enhanced chemiluminescence.

In vitro kinase assay

Kinase assays were performed using 250 ng of GST-PLD2 fragments as substrate. GST-PLD2 fragment were incubated with 500 ng of purified CKII in 30 μ l of kinase buffer (20 mM Tris-HCl, pH 7.4, 20 μ M ATP, and 5 mM MnCl₂ containing 10 μ Ci [γ -³²P]ATP) at 37°C for 10 min. Kinase reactions were terminated by addition of SDS-PAGE loading buffer with β -mercaptoethanol, boiled for 5 min, separated by SDS-PAGE, and exposed to film.

Cell proliferation assay

Cells in 96-well plates were seeded in DMEM supplemented with 10% fetal bovine serum and incubated with emodin or apigenin at 37°C for 24 h. Afterwards, cells were incubated for 2 h with CellTiter 96 Aqueous (MTS) solution, absorbance of coloured MTS products in the supernatant was determined on ELISA reader at 495 nm.

Results and Discussion

CKII is involved in PMA-induced PLD activation

Many substrates of CKII are involved in signal transduction pathways (Blanquet, 2000). In most cases, phosphorylation of these substrates occurs following their interaction with the kinase. However, despite some advances, the significance of these events remains to be clarified or is still quite enigmatic. It has been suggested that CKII might have potentially important roles in neuronal function such as neuronal differentiation and proliferation (Blanquet, 2000). It was demonstrated that PKC stimulation via phorbol ester treatment increases CKII activity (Sanghera *et al.*, 1992; Silva-Neto *et al.*, 2002). PKC is also known to stimulate PLD activity (Frohman

and Morris, 1999). Recently, it has been reported that CKII interacts with PLD1 (Ganley *et al.*, 2001). Thus, we investigated whether CKII affects agonist-induced PLD activation in human U87 astrogloma cells. CKII- α or β subunit was transiently transfected in U87 astrogloma cells, and the cells were stimulated with or without 4 β -phorbol 12-myristate 13-acetate (PMA). Interestingly, both CKII- α and β subunit stimulated basal PLD activity as well as PMA-induced PLD activation (Figure 1A). To confirm that CKII is involved in PMA-induced PLD activation, we used CKII-specific inhibitors, emodin and apigenin. Moreover, we used U87 cells expressing PLD1 and PLD2 to examine which isozyme of PLD activity is affected by CKII. As shown in Figure 1B and C, These inhibitor suppressed in a dose-dependent manner PMA-induced PLD activation and showed similar potency in inhibiting PLD activity. Calphostin C, PKC inhibitor as a positive control, inhibited activation of PLD by PMA. Emodin and apigenin suppressed PLD1 or PLD2 activity as well as basal PLD activity (Figure 1D, E) and. These drugs did not affect the expression of PLD isozymes. These results suggest that CKII is involved in PMA-induced PLD1 or PLD2 activation as well as basal PLD activity.

CKII interacts with PLD1 and PLD2

Little is known about CKII regulation except for the well-characterized multiple dimerization combinations between the two catalytic subunits (α and α') and the regulatory subunit (β) that maximizes its kinase activity (Allende and Allende, 1998). However, recent studies indicate that CKII may form hetero-complexes with unrelated kinases and potentially modify its function (Bren *et al.*, 2000). Atypical PKC directly interact with CKII, and hence potentially regulates its activity (Bren *et al.*, 2000). To examine whether the interaction of CKII with PLD is involved in PLD activation by CKII, we tried to perform co-transfection experiments in U87 cells using its expression vector. After transfection, cell lysates were prepared and subjected to immunoprecipitation using an antibody directed against PLD or CKII- α , and monitored by successive immunoblotting for CKII- α or PLD. The data presented in Figure 2 show that anti-PLD antibody and anti-CKII- α antibody, but not non-immune IgG (NG), immunoprecipitates CKII- α , and PLD, respectively. These results suggest that CKII- α associates with both PLD1 and PLD2.

CKII phosphorylates PLD2

It has been reported that CKII phosphorylates PLD1 and this phosphorylation event does not influence PLD1-mediated hydrolysis of phosphatidylcholine *in vitro* (Ganley *et al.*, 2001). In this study, we found

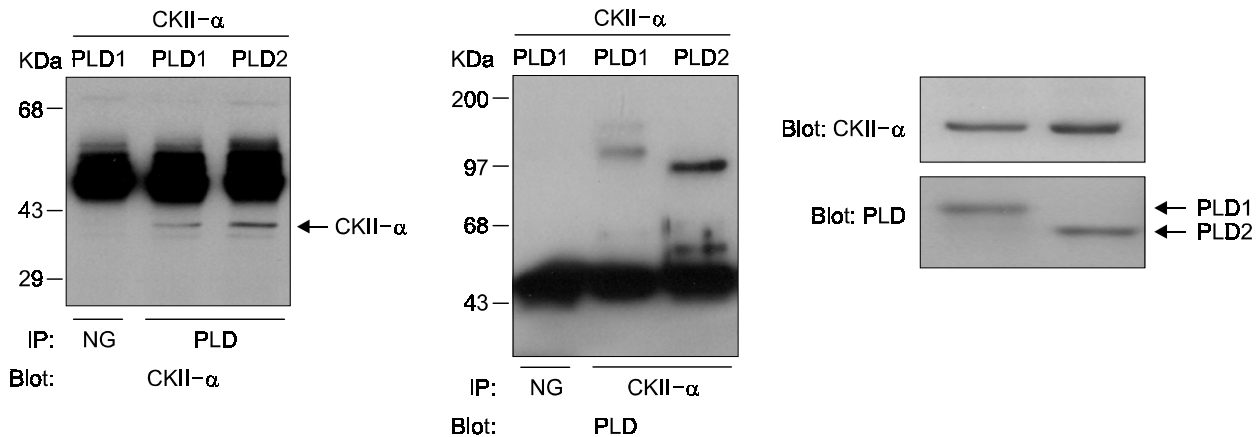


Figure 2. CKII associates with PLD isozymes. U87 cells were co-transfected with PLD1 or PLD2 and CKII- α . Cell lysates were immunoprecipitated with non-immune IgG (NG), anti-PLD or anti-CKII- α antibodies and immune complexes were subjected to immunoblotting using anti-CKII- α or anti-PLD antibodies, respectively. Expression of PLDs and CKII- α were determined by immunoblotting. Data are representative of three experiments.

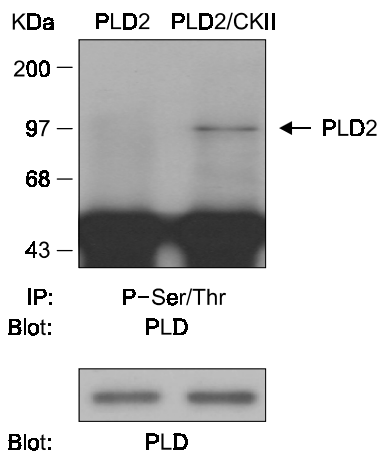


Figure 3. PLD2 is phosphorylated by CKII- α in co-transfection experiments. U87 cells were transiently transfected with only PLD2 or CKII and PLD2, and the cell lysates were immunoprecipitated with anti-phospho-Ser/Thr antibody. The immune complexes were analyzed by immunoblotting using anti-PLD antibody. PLD expression were determined by immunoblotting using anti-PLD antibody.

that inhibitors of CKII suppressed PMA-induced PLD1 and PLD2 activation *in vivo* as shown in Figure 1B and C. These results suggest that regulation of PLD activity by CKII *in vitro* can be different from that of *in vivo*. Next, we tried to examine whether CKII can phosphorylate PLD2. Transfection with only PLD2 expression vector or co-transfection with CKII- α and PLD2 cDNA was performed in U87 cells, and the lysates were immunoprecipitated with anti-phospho-serine/threonine antibody, and analyzed by immunoblotting using anti-PLD antibody. As shown in Figure 3, anti-phospho-serine/threonine antibody

immunoprecipitated PLD2 in the cells co-transfected with both expression vectors, but not in the cells transfected with only PLD2, suggesting that PLD2 is phosphorylated by CKII. In addition, we analyzed the phosphorylation site(s) of PLD2 by CKII. To identify the PLD2 sequence phosphorylated by CKII, we constructed the GST fusion proteins shown in Figure 4A. *In vitro* kinase assays using purified GST-PLD2 fragments and CKII holoenzyme were performed. GST-PLD2 fragments were incubated with CKII holoenzyme in kinase buffer containing [γ - 32 P]ATP. The reactions were separated by SDS-PAGE, and autoradiography was performed. Multiple regions of PLD2 were phosphorylated *in vitro* by CKII (Figure 4B). To examine whether GST-PLD2 fragment is specifically phosphorylated by CKII, GST-PLD2 F5 fragment was used for *in vitro* kinase assay in the presence or absence of apigenin. As shown in Figure 4C, apigenin inhibited CKII-induced phosphorylation of PLD2 fragment, suggesting the phosphorylation of PLD2 by CKII. These results suggest that PLD2 is phosphorylated by CKII *in vivo* and *in vitro*. Future studies are needed to identify the amino acid residues and roles of PLD2 phosphorylated by CKII.

CKII inhibitors suppress proliferation of U87 astrogloma cells

It has been reported that CKII activity increases when quiescent cells are stimulated to proliferate (Issinger, 1993; Allende and Allende, 1995; Pinna and Meggio, 1997). Studies performed using selective inhibitors also illustrate that CKII is required during various stages of cell cycle progression (Ford *et al.*, 2000). Moreover, CKII activity levels are elevated in a num-

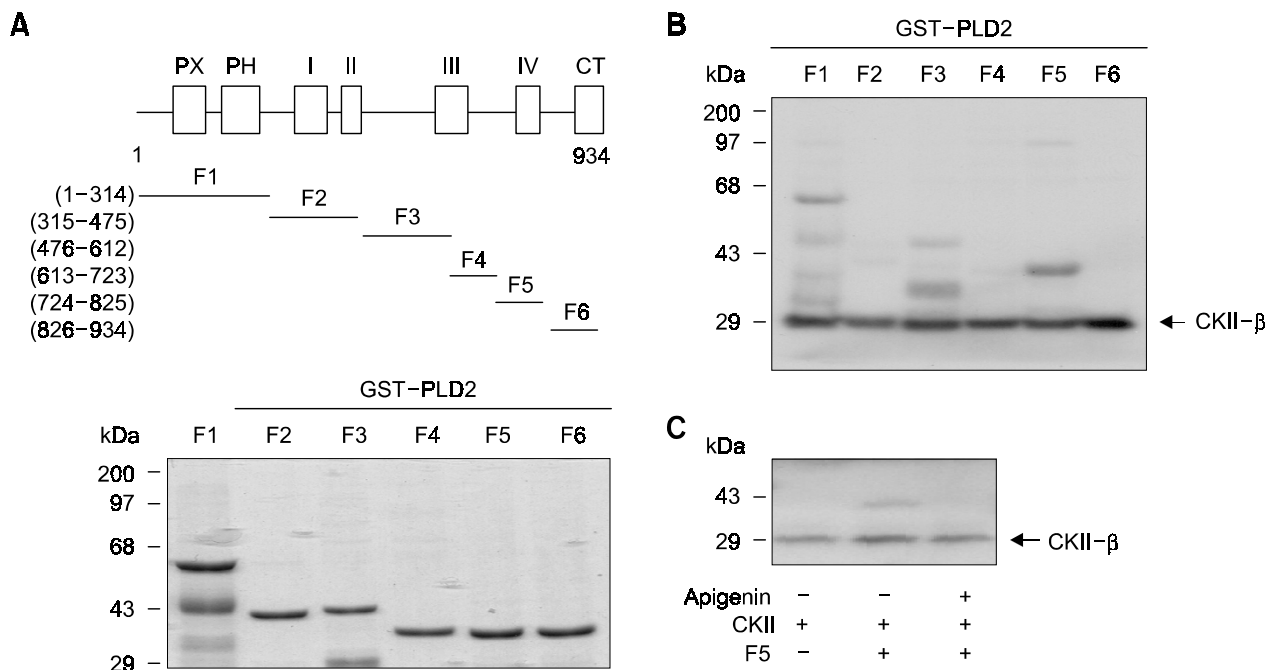


Figure 4. Multiple regions of PLD2 are phosphorylated by CKII *in vitro* kinase assay. (A) Primary structure and individual domains of PLD2. Boxes indicate regions of highly conserved sequences in PLD isozymes (top). PX, phox domain; PH, pleckstrin homology domain; CR I, II, III, IV, conserved region I, II, III, IV; CT, carboxyl terminus. Purified GST-fusion proteins containing different regions of PLD2 were subjected to SDS-PAGE followed by Coomassie Brilliant Blue staining (bottom). (B) GST-PLD2 fragments were incubated with CKII holoenzyme in kinase buffer containing [γ - 32 P]ATP at 37°C for 10 min. (C) GST-PLD2 F5 fragment was incubated with or without apigenin in the presence of CKII holoenzyme in kinase buffer. The kinase reactions were terminated by addition of SDS loading with β -mercaptoethanol buffer, boiled for 5 min, and separated by SDS-PAGE. The gels were dried and exposed to film to visualize all bands.

ber of tumors and leukemic cells. Collectively, these studies implicate CKII as an important component of signaling pathways involved in cell cycle progression and transformation. PLD also has emerged as a critical regulator of cell proliferation and survival, and prevents apoptosis (Foster, 2003). It is therefore possible that the interaction of CKII with PLD, and activation of PLD by CKII may impact cell functions. Thus, we tried to examine the possibility that blocking of PLD activity by CKII inhibitors might be involved in the suppression of cell proliferation. To assess the effect of CKII on proliferation of U87 astrogloma cells, we used the CellTiter 96 Aqueous proliferation assay that is based on metabolic conversion of a tetrazolium compound, MTS, to a coloured product by living cells. The absorbance intensity of the MTS product is directly proportional to the number of viable cells in culture. As shown in Figure 5, transfection of CKII or PLD2 significantly increased absorbance, but treatment of CKII inhibitors (emodin, apigenin) and 1-butanol which block PA production by PLD, decreased absorbance compared with that of control cells. In addition, 1-butanol suppressed CKII-induced cell proliferation. These results suggest that CKII is involved in proliferation of U87 cells at least in

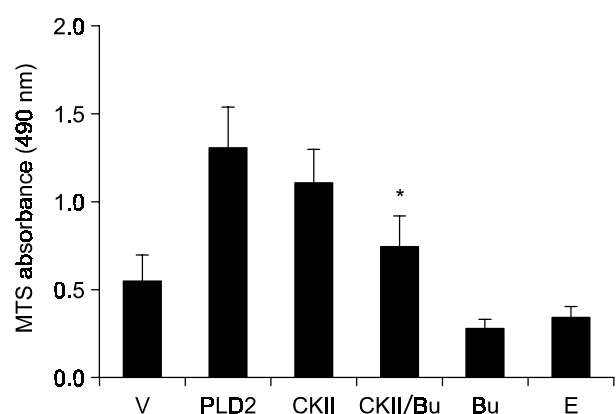


Figure 5. CKII is involved in proliferation of U87 cells in part, through stimulation of PLD activity. U87 cells were transfected with PLD2 or CKII- α and treated with or without 1-butanol (0.3%), emodin (50 μ M) for 24 h. The number of viable cells was determined by the CellTiter Aqueous (MTS) reagent. Differences in absorbance at 495 nm were plotted as relative changes in the number of live cells. * $P < 0.05$ compared to CKII cells. Results are means \pm S.D. of three independent experiments.

part, through stimulation of PLD activity

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