
REVIEW

Molecular Structure of Phospholipase D and Regulatory Mechanisms of Its Activity in Plant and Animal Cells

Y. S. Kolesnikov¹, K. P. Nokhrina², S. V. Kretynin¹, I. D. Volotovskii³,
J. Martinec⁴, G. A. Romanov⁵, and V. S. Kravets^{1*}

¹*Institute of Bioorganic Chemistry and Petrochemistry, National Academy of Sciences of Ukraine, ul. Murmanskaya 1, 02094 Kiev, Ukraine; fax: +38 (044) 573-2552; E-mail: kravets@bpci.kiev.ua*

²*Plant Biotechnology Institute, National Research Council of Canada, 110 Gymnasium Place, Saskatoon, SK, S7N 0W9, Canada; E-mail: Kateryna.Nokhrina@nrc-cnrc.gc.ca*

³*Institute of Biophysics and Cell Engineering, National Academy of Sciences of Belarus, ul. Akademicheskaya 27, 220072 Minsk, Belarus; E-mail: volot@biobel.bas-net.by*

⁴*Institute of Experimental Botany, Academy of Sciences of the Czech Republic, Rozvojova 263, 16502 Prague 6, Czech Republic; fax: +420 (225) 106-456, E-mail: martinec@ueb.cas.cz*

⁵*Institute of Plant Physiology, Russian Academy of Sciences, ul. Botanicheskaya 35, 127276 Moscow, Russia; E-mail: gar@ippras.ru*

Received February 1, 2011

Revision received March 23, 2011

Abstract—Phospholipase D (PLD) catalyzes hydrolysis of phospholipids with production of phosphatidic acid, which often acts as secondary messenger of transduction of intracellular signals. This review summarizes data of leading laboratories on specific features of organization and regulation of PLD activity in plant and animal cells. The main structural domains of PLD (C2, PX, PH), the active site, and other functionally important parts of the enzyme are discussed. Regulatory mechanisms of PLD activity are characterized in detail. Studies associated with molecular design, analysis, and synthesis of new nontoxic substances capable of inhibiting different PLD isoenzymes *in vivo* are shown to be promising for biotechnology and medicine.

DOI: 10.1134/S0006297912010014

Key words: phospholipase D, domains, calcium, lipids, G-proteins, protein kinases, protein–protein interactions

Phospholipase D (EC 3.1.4.4) (PLD) is an enzyme widely occurring in plants [1, 2] and animals [3, 4]; it catalyzes hydrolysis of the phosphodiester bond in structural phospholipids (phosphatidylcholine, phosphatidylethanolamine, etc.) with production of phosphatidic acid and low molecular weight water-soluble compounds of choline or ethanolamine type [5]. Phosphatidic acid is considered to be not only a necessary structural element of membranes but also an important secondary messenger of hormone and stress signaling cascades [6, 7] and as a regulator of membrane transport [8]. Phosphatidic acid can also be generated by diacylglycerol kinases [9].

PLD was found and cloned in plants, and the model plant *Arabidopsis thaliana* has been shown to have 12

PLD genes (see reviews [10, 11]). PLD is very sensitive to external and internal factors, and its diversity is favorable for fine coordination of enzyme activity with the state of cell metabolism. PLD is actively studied on different organizational levels in both plants and animals.

This review analyzes the data on PLD of plants and animals and also gives the authors' concept about the involvement of PLD in transmission of signals of such important phytohormones as cytokinins. In particular, known regulatory mechanisms of PLD activity are analyzed, and comparative characteristics of these enzymes in cells of different origin are presented. A wide spectrum of factors known to play key roles in PLD regulation is considered, the organization of the enzyme is analyzed, its role in the regulation of cell metabolism is determined, and main directions for studies on lipid signaling systems are described.

Studies in biochemistry and molecular genetics were favorable for determination of the role of PLD in plant

Abbreviations: Gα, α-subunit of heterotrimeric G-protein; PKC, protein kinase C; PLD, phospholipase D; PtdIns(4,5)P₂, phosphatidylinositol-4,5-bisphosphate.

* To whom correspondence should be addressed.

and animal cells [12]. There are data on the involvement of PLD in malignancies [13], angiogenesis [14], leukocyte adhesion, and chemotaxis [15]; the functions of the enzyme are considered in platelets [16], nerve cells [17], myocardium [18], lungs [19], in transport of intracellular membranes [20, 21], and signal transmission in animals [6, 22–25] and plants [2, 11]. Moreover, the therapeutic potential of PLD inhibitors has been pointed out [4]. However, the published data on the regulatory mechanisms of PLD activity in plants and animals and on the involvement of PLD in signal transduction have not been analyzed yet. The purpose of this review is to fill this gap and to inform readers about the current status researches in this field.

PLD of animal cells is encoded by two genes and exists in two forms: PLD1 and PLD2 [26, 27]. PLD1 (120 kDa) is mainly present on endomembranes of the cell and is transported to plasma membrane under the influence of extracellular stimuli. PLD2 (106 kDa) is located on the plasma membrane, has low *in vivo* activity, and is moderately stimulated by known activators of PLD1. Plants are found to have many genes of various PLD isoenzymes that encode high molecular weight multidomain proteins with molecular weights of 90–125 kDa. In *Arabidopsis* plants there are 12 genes of PLD: PLD α (3), β (2), γ (3), δ , ϵ , and ζ (2) [2]; in rice there are 17 genes of PLD: PLD α (8), β (2), δ (3), κ , ζ (2), and ϕ [28]. PLD α 3 [29], PLD δ [11], PLD ζ 1 [2], and PLD ϵ [30] are located mainly on the plasma membrane, PLD ζ 2 is associated with the tonoplast membrane [31], PLD α 1 shuttles between the cytosol and membranes [2], whereas PLD γ 1 is located on intracellular membranes [2]. The location of PLD β is not strictly determined [2]. PLD α 1 is the most active and PLD ζ 1 and PLD ϵ are the least active isoenzymes in plants. In the absence of activators and calcium in the incubation medium all PLD isoenzymes of plant cells are inactive *in vitro*.

SPECIFIC FEATURES OF PRIMARY STRUCTURE OF PLD

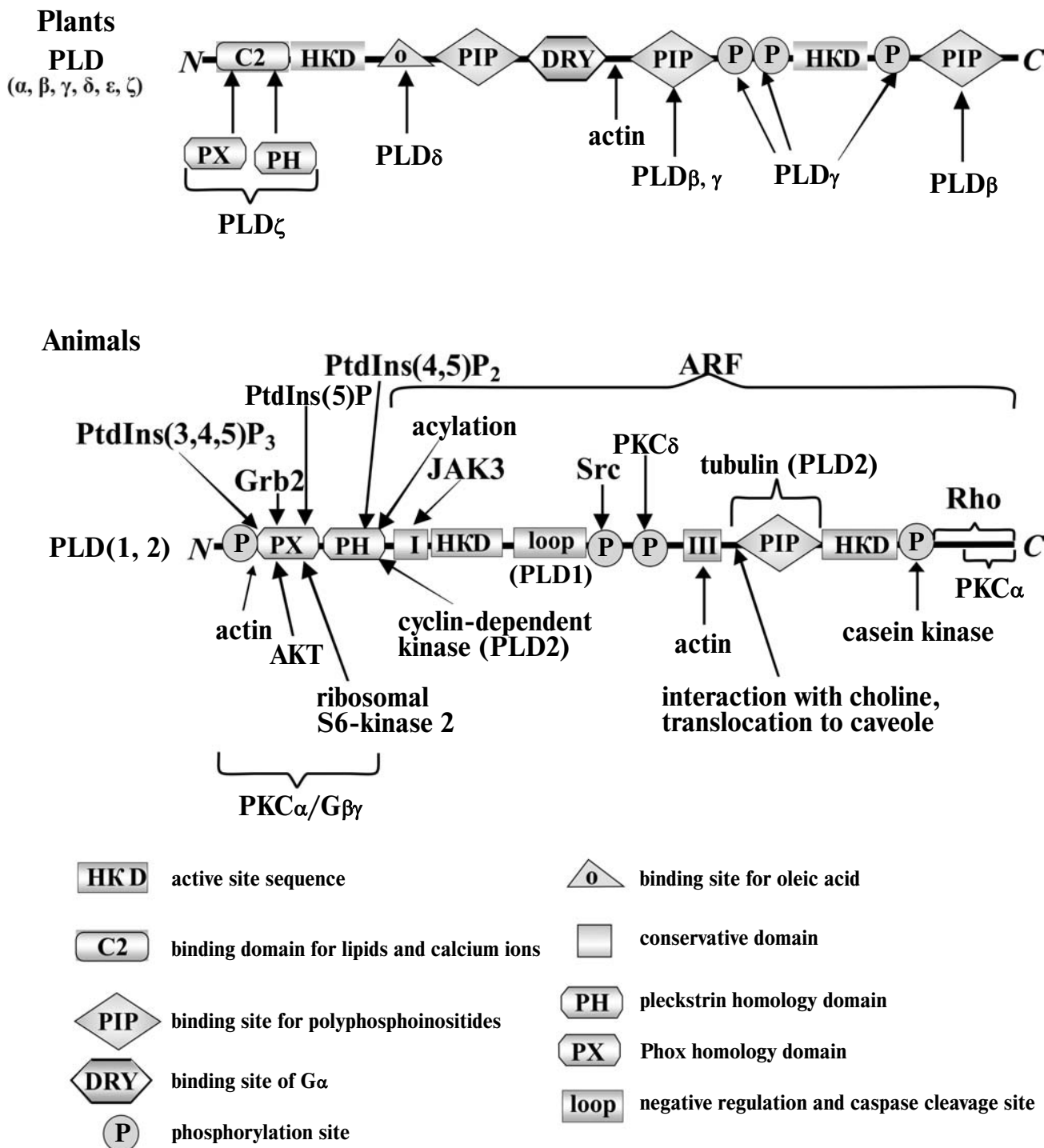
Active site of PLD. PLDs from animal and plant cells have a modular structure where sites responsible for catalysis are surrounded by regulatory sequences (figure). The active site of PLD consists of four conservative amino acid sequences (I–IV) among which II and IV are especially conservative in various organisms. Motifs II and IV contain the sequence HxKxxxxDxxxxxxGSxN (abbreviated HKD, where x are nonconservative amino acids), and due to this sequence PLD can catalyze hydrolysis of phospholipids [32]. Residues of histidine (H), lysine (K), and aspartic acid (D) are directly involved in hydrolysis of phosphodiester bonds of phospholipids. The histidine residue acts as a nucleophile that attacks the phosphorus atom of the substrate [32–34]. The catalytic site of PLD is

formed when the HKD sequences are in the correct proximity. The PLD-catalyzed hydrolysis of the phosphodiester bond is supposed to occur in two stages by the so-called ping-pong mechanism: successive attack of the substrate by imidazole residues of histidines, each of which is a component of the catalytic sequences of HKD. As a result, the phosphatidic acid residue of the cleaved phospholipid makes transient covalent bond with the imidazole group of histidine of PLD producing a short-lived intermediate that can be easily cleaved on attack by a water [35, 36].

Lipid-binding PLD sequences. Lipids can modulate PLD activity in plant and animal cells. In particular, some phospholipids usually act not as substrates but as cofactors of the enzyme. The PLD activity depends on binding of the cofactor phosphatidylinositol-4,5-bisphosphate (PtdIns(4,5)P₂). In animals this is explained by the presence within the active site (between motifs III and IV) of a conservative sequence that includes residues of basic and aromatic amino acids [37, 38]. This sequence ensures high levels of PLD1 and PLD2 activities and also promotes the translocation of PLD1 to membranes in response to extracellular signals [39, 40].

The amino acid sequence of PLD enriched by basic and hydrophobic residues is involved in binding polyphosphoinositides [37]. This structure and its inverted repeat (RxxxxKxRR and RKxRxxxxR) in PLD β of *Arabidopsis* and tomato surround the C-terminal sequence HKD [2, 41]. PLD β 1 contains all four residues of basic amino acids (mainly Arg) that bind polyphosphoinositides, whereas one of them in PLD γ and PLD ζ is replaced by a nonpolar or acidic residue. This structure in PLD δ has two conservative residues of basic amino acids, whereas in PLD α and PLD ϵ there is only one such residue and the other residues are replaced by nonpolar or acidic ones [2, 11]. In addition to inverted repeats that bind polyphosphoinositides (called PBR2), PLD β also contains another binding site for these lipids (PBR1) that is located near the N-terminal HKD motif [2]. PBR1 is believed to be an independent binding site of PtdIns(4,5)P₂ and some other phosphoinositides; it consists of highly conservative amino acid residues and contains many side chains of lysine, arginine, and histidine residues. The specific binding of PtdIns(4,5)P₂ with them plays an extremely important role in the regulation of plant PLD β by this phospholipid because it promotes conformational rearrangements strengthening the enzyme association with the substrate [2, 41, 42]. PLD α and PLD δ contain amino acid substitutions in the abovementioned binding site of PtdIns(4,5)P₂ [2]. The phosphate group in the 4'-OH position of the inositol ring of PtdIns(4,5)P₂ plays a key role in formation of ionic bonds with lysine residues K437 and K440 of the PLD β molecule [2].

Domain C2 interacting with lipid membranes depending on the Ca²⁺ concentration is another type of phosphoinositide-binding sequences of plant PLDs. This



Features of phospholipase D structure of animal and plant (*Arabidopsis*) cells. DRY, binding site of α -subunit of the heterotrimeric G-protein; $G\beta\gamma$, β - and γ -subunits of the heterotrimeric G-protein; I and III, conservative domains; PtdIns(5)P, phosphatidylinositol-5-phosphate; PtdIns(4,5)P₂, phosphatidylinositol-4,5-bisphosphate; PtdIns(3,4,5)P₃, phosphatidylinositol-3,4,5-trisphosphate

domain is not strictly specific to PtdIns(4,5)P₂. At low level of Ca²⁺ negatively charged PtdInsP and PtdIns(4,5)P₂ molecules attract positively charged amino acid residues R and K in the C2 domain and thus favor the binding with substrates of PLD α of *Arabidopsis* and

catalysis by PLD β and PLD γ [2, 43]. Binding sites of PtdIns(4,5)P₂ in domain C2 are also found in PLD α of cabbage [44], poppy [45], and tomatoes [46]. The C2 domain in tomato PLD α binds with high affinity phosphatidic acid but not phosphatidylcholine, but associa-

tion with phosphoinositides is markedly weakened on increase in the level of their phosphorylation [46].

Domain PX. Animal PLDs and plant PLD ζ contain the *N*-terminal domain PX (named after the protein p47phox that is a subunit of NADPH oxidase of phagocytes), which includes a conservative sequence enriched in proline residues [2, 28]. The PX domain of PLD1 selectively binds PtdIns(5)P, thus promoting the translocation of the enzyme to membranes of endocytic vesicles [39], and also interacts with PtdIns(3)P and PtdIns(3,4)P₂ on the membranes [47]. This domain contains two binding sites of membrane phospholipids: one of them is highly specific to PtdIns(3,4,5)P₃, PtdIns(3)P, PtdIns(5)P, and other phosphoinositides, whereas the other also binds phosphatidic acid, phosphatidylserine, and other negatively charged lipids. The concurrent binding of phospholipids with the abovementioned sites synergistically increases the affinity of domain PX for the membranes [48, 49]. The PX domain of PLD1 plays an important role in regulation of the enzyme translocation into endocytosis sites on the membranes under the influence of extracellular signals [40, 49]. In PLD1 and PLD2 this domain can stimulate the GTPase activity of the protein dynamin [50] and ensure the binding with PLD2 and activation of phospholipase C γ 1 [51], tyrosine kinase Syk [52], and protein kinase C ζ [53].

Domain PH. PLD1 and PLD2 of animals [38] and also PLD ζ of plants [2, 28] include the domain PH (pleckstrin homology domain) responsible for specificity of localization of these enzymes in cells. The PH domain is located near the *C*-terminus of the PX domain. Deletions or point mutations in the domain of PLD1 and PLD2 do not influence the activity *in vitro* [47, 54], but inactivate the enzymes *in vivo* due to disturbance of their localization in cells [38, 40]. The PH domain of PLD2 binds PtdIns(4,5)P₂ with moderate affinity but with high selectivity [38]. Deletion of this domain in PLD1 reduces the binding of this phosphoinositide [47], but did not influence the *in vitro* activation of PLD1 and PLD2 by phosphoinositides [37, 38]. This suggests the presence in the PLD structure of other sites of association with PtdIns(4,5)P₂. The PH domain of PLD1 and PLD2 contains conservative cysteine residues (Cys240 and Cys241) that can be palmitoylated and thus ensure the binding with the membranes independently of phosphoinositides [55–57], which stabilizes the interaction of the enzyme with the lipid bilayer [56]. Palmitoylation promotes the translocation of PLD1 into endocytosis sites on the membranes under the influence of an extracellular stimulus [40] and the translocation of PLD2 from early endosomes onto the plasma membrane [38].

Loop region. The primary structure of PLD1 of animal cells is uniquely characterized by the presence of a loop-shaped domain consisting of 116 amino acid residues (a.a.). Insertion of this sequence into PLD2 does not change the activity of this isoenzyme. The loop-

shaped domain of PLD1 contains a site of negative regulation, which possibly is responsible for the low basal *in vitro* activity of PLD1 as compared to that of PLD2 [54]. This conservative sequence belonging to PLD1 also contains sites of cleavage by caspases. The *in vitro* action of caspases on this sequence weakens the sensitivity of PLD1 to protein kinase C but increases its sensitivity to small GTPases [58].

Sites of Ca²⁺ binding. Based on specific features of the primary structure of the *N*-terminal domains binding lipids, PLD are subdivided onto C2-PLD (PLD α , PLD β , PLD γ , PLD δ , and PLD ϵ of plants) and PX/PH-PLD (PLD ζ of plants and PLD1/2 of animals) (figure). The presence of *N*-terminal conservative domain C2 consisting of 130 a.a. that binds Ca²⁺ and phospholipids is a unique feature of the majority of plant PLDs [2]. The binding of Ca²⁺ with the domain C2 provides for the PLD translocation to substrate on the membrane and induces changes in the enzyme conformation [46]. A direct binding of Ca²⁺ by three-looped sequences of the C2 domain [11] has been shown for PLD α and PLD β of *Arabidopsis* [2]. Calcium forms a coordination bond with four or five acidic amino acid residues in the calcium-binding loops (CBL) of the PLD domain C2. The main role in coordination of Ca²⁺ in the C2 domain of PLD β , γ , and δ belongs to two aspartic acid residues in the Ca²⁺-binding loop 3 (CBL3), the only Asn residue in the Ca²⁺-binding loop 2 (CBL2), and Asp in the Ca²⁺-binding loop 1 (CBL1) [2, 43]. In PLD α two of these amino acids are replaced in CBL1 and CBL2 by basic or neutral residues [11, 43], whereas the C2 domain of PLD ϵ does not have any amino acid binding with calcium [30]. In PLD α of other plants coordination bonds with Ca²⁺ are formed by asparagine and aspartic acid [59], glutamine and glutamic acid [41], and also by their combination [42]. The above-described amino acid substitutions are responsible for the low affinity of PLD α and PLD ϵ for Ca²⁺, and therefore these isoenzymes hydrolyze phospholipids only in the presence of high concentrations of Ca²⁺ [2] or at low pH increasing the affinity for calcium. The Ca²⁺ binding has been also observed in the active site of PLD α [60] and PLD β , where it is stimulated by phosphatidylserine [61].

Other functional regions of PLD. PLDs of animal cells also contain other functional regions on the *N*- and *C*-termini. In particular, the *N*-terminal amino acid sequence of human PLD1 (residues 1–49 and 216–318 including the PH domain) plays a key role in the stimulation of this enzyme by protein kinase C α [62]. Moreover, the sequence consisting of 155 *N*-terminal amino acids of human PLD1, especially Ser2, is involved in the interaction with polymerized F-actin [63]. The actin-binding domain of plant PLD located between catalytic subdomains in the conservative domain III contains asparagine and threonine residues that are directly involved in the actin binding [64]. The *C*-terminus of PLD can regulate

the enzyme localization in the cells, contribute to its interaction with small GTPases of the RhoA family, and also play an extremely important role in stabilizing the active conformation of the catalytic site of PLD [65, 66].

In plants, an amino acid sequence DRY located on the *N*-terminal end of the HKD C-subdomain is responsible for the binding of PLD α 1 with the heterotrimeric G-protein subunit G α . Except for PLD ζ and PLD γ 2, this sequence is also present in other PLD isoenzymes of *Arabidopsis* [67]. Potential phosphorylation sites of serine, threonine, and tyrosine residues are found in the primary structure of PLD γ of *Brassica oleracea* and also in their homologs of poppy PLD α [45, 68].

REGULATORY MECHANISMS OF PLD ACTIVITY

Activity of PLD is controlled by hormones, neurotransmitters, and other physiologically active molecules acting on the enzyme through specific regulatory mechanisms. In such case PLD is often contributes to signal transduction through production of phosphatidic acid as a second messenger. In particular, PLD is involved in transduction of signals of classic plant hormones such as abscisic acid [11] and cytokinin [69].

Regulation of PLD activity by calcium ions. Calcium plays an important role as a modifier of PLD activity in plant cells, for example under the influence of cytokinin [70] and also as a cofactor of PLD isoenzymes containing the C2 domain [2, 43, 61]. Calcium ions activate PLD α and PLD β through binding with the C2 domain. This leads to conformational changes in the enzymes that result in strengthening the substrate binding and weakening the association with PtdIns(4,5)P $_2$ [2]. Calcium also binds to the active site of PLD β , which increases its affinity for the activator (PtdIns(4,5)P $_2$) in comparison with affinity for the substrate (phosphatidylcholine) [61]. In turn, the binding of PtdIns(4,5)P $_2$ with the active site of the enzyme increases its affinity for the substrate [2]. In plants there are PLDs with activities depending on millimolar or micromolar concentration of Ca $^{2+}$ *in vitro*.

Plant PLDs are active *in vitro* in the presence of Ca $^{2+}$ concentrations of 10–100 mM and hydrolyze phospholipids independently of the presence in the substrate mixture of lipid cofactors (PtdInsP or PtdIns(4,5)P $_2$). In particular, Ca $^{2+}$ (1–100 mM) *in vitro* activates PLD α isolated from cabbage [60], strawberry [42], *Vigna sinensis* [71], sunflower [72], and poppy [45]. The maximal activity of PLD α 1 in *Arabidopsis* is observed at the Ca $^{2+}$ concentration of 25 mM. On using pure phosphatidylcholine as a substrate, the activity of this enzyme is stimulated at Ca $^{2+}$ concentrations of 20–100 mM [2]. At the Ca $^{2+}$ concentration of 50 mM, the other PLD isoenzyme, PLD ϵ , is also active [30]. Such high concentrations of Ca $^{2+}$ activate those plant PLD isoenzymes that have the C2 domain with low affinity for Ca $^{2+}$ [2].

Plant PLDs are also stimulated by micromolar concentrations of Ca $^{2+}$ [5]. In the case of PLD β , this is due to a high affinity of the C2 domain for Ca $^{2+}$ [2]. Activities of PLD β and PLD γ of *Arabidopsis* are maximal at 50 μ M Ca $^{2+}$. PLD γ is less stimulated by Ca $^{2+}$ than PLD β and PLD α . Increase in Ca $^{2+}$ concentration (>50 μ M) is associated with a decrease in the activities of PLD γ and PLD β [2, 43]. Activities of the PLD γ subtypes PLD γ 1 and PLD γ 2 depend on Ca $^{2+}$ concentration similarly [73]. Moreover, PLD α of castor bean in acidic medium (pH 4.5–5.0) is also stimulated *in vitro* at low Ca $^{2+}$ concentration (50 μ M), but on pH increase the active concentration of Ca $^{2+}$ increases to millimolar level [2]. The activity of isoenzyme PLD δ of *Arabidopsis* is observed *in vitro* at both micro- and millimolar Ca $^{2+}$. The activity of PLD δ in the presence of an activator (oleic acid) is maximal at 100 μ M Ca $^{2+}$ [11]. PLD ϵ is also active *in vitro* at micromolar Ca $^{2+}$ concentrations, but only in the presence of oleic acid [30].

Plant cells also contain calcium-independent PLDs: PLD ζ 1 of *Arabidopsis* [2] and PLD of mustard seeds [74]. Moreover, Ca $^{2+}$ does not markedly change the *in vitro* activity of purified PLD1 from animal cells [26].

Modification of PLD activity by lipids. PLDs are regulated by various lipids, and among them polyphosphoinositides that are considered to be especially important modifiers of cytoskeleton dynamics, membrane transport, and enzyme activity [75]. PLDs sensitive to these phospholipids are involved in the transduction of intracellular signals of hormones, in particular of cytokinins in plants [76–78]. In animal cells the catalytic activities of PLD1 and PLD2 are sensitive to PtdIns(4,5)P $_2$, which acts as their cofactor [26]. The binding of PtdIns(4,5)P $_2$ with PLD1 and especially with PLD2 is important for the stimulation of these enzymes [37, 55, 79]. PLD2 is allosterically activated by phosphatidylinositol-4,5-bisphosphate *in vitro* [37]. Optimal concentrations of this lipid provide for a positive effect of activators (proteins Rho, ARF1, and PKC α) on PLD1 *in vitro* promoting the PLD binding with a lipid bilayer [80]. The enzyme of PtdIns(4,5)P $_2$ biosynthesis, phosphatidylinositol-4-phosphate-5-kinase I α , promotes an increase in PLD activity in Cos7 cells. The PLD2 activity increases under conditions of a simultaneous expression of the genes encoding PLD and the abovementioned phosphoinositide kinase [79]. In comparison with other phosphoinositides, PtdIns(3,4,5)P $_3$ binds to residue Arg179 in the PX domain of PLD1 [49], stimulates the enzyme activity [49, 81], and also promotes the translocation of PKC ζ into the cell nucleus [82] leading to the activation of the nuclear PLD.

In plants polyphosphoinositides are important cofactors of PLD β , PLD γ , and PLD ζ , and in the absence of the formers these isoenzymes are inactive *in vitro*. PLD β of *Arabidopsis* displays the most pronounced specificity. This isoenzyme is stimulated *in vitro* over a wide

range of PtdIns(4,5)P₂ concentrations [11]. PLDβ is less sensitive *in vitro* to other phosphatidylinositol phosphates [2]. PtdIns(4)P and PtdIns(4,5)P₂ are additive in increasing the activities of PLDβ and PLDγ [2, 43]. Unlike PLDγ2, PLDγ1 is the most sensitive to PtdIns(4,5)P₂ [73]. PLDα does not depend on polyphosphoinositides in the presence of optimal Ca²⁺ concentrations but is stimulated by these lipids *in vitro* in acidic medium in the presence of micromolar Ca²⁺ concentrations [2, 11]. The activity of PLDδ of *Arabidopsis* depends on PtdIns(4,5)P₂ because this lipid increases the affinity of the enzyme active site for the substrate [11, 83]. The PLDζ depends only on PtdIns(4,5)P₂ [2], whereas PtdIns(3,4)P₂ is a cofactor of PLDε at micromolar concentrations of Ca²⁺ [30]. Thus, PtdInsP₂ acts in *Arabidopsis* as a cofactor of PLDβ, PLDγ, PLDε, and PLDζ. Note that PLD activity independent of PtdInsP₂ has been detected in mustard seeds [74].

Other lipids can be negative regulators of PLD. In particular, C₂-, C₆-, C₈- and long-chain ceramides inhibit the activity of PLD by affecting its interactions with protein activators [84], acting on the catalytic site of PLD or on its binding with PtdInsP₂ [85, 86]. Lysophospholipids also specifically inhibit PLD *in vitro* independently of PtdInsP₂ and the small GTPase Arf. They directly bind to PLD and noncompetitively inhibit the enzyme in animals [87] and plants [88].

Unsaturated fatty acids, products of lipid cleavage, are believed to be activators of animal PLD2 *in vitro*. This enzyme can be stimulated by fatty acids synergistically with PtdIns(4,5)P₂ [89, 90]. PLDδ and PLDε of *Arabidopsis* under *in vitro* conditions depend on oleic acid [11, 30], but this may be associated with modification of properties of the lipid bilayer with which these isoenzymes are associated [83]. The mechanism of the inhibitory effect *in vitro* of N-acetyethanolamines (N-lauroylethanolamine, N-myristoylethanolamine) on the activity of PLDα from *Arabidopsis*, castor bean, and cabbage [91] is unknown.

Regulation of PLD activity by G-proteins. Animal cells contain specific protein factors that are important for activation of PLD. These factors are small monomeric GTPases of the Rho and ARF families of the Ras superfamily. Rho GTPases play an important role in the regulation of various processes in animal cells, including transcription, cell cycle, secretion, cell transformation, and cytoskeleton rearrangements [92]. Small GTPases of the Rho family are also considered as modulators of PLD activity. Some of these proteins — Rac1, Cdc42, and especially RhoA — allosterically activate PLD1 increasing its catalytic activity by lowering the Michaelis constant [80]. In the presence of nonhydrolyzable analogs of GTP, small GTPases Rac1, Cdc42, and RhoA activate PLD1 [26, 93, 94] and PLD2 [95] *in vitro*. RhoA [96, 97], Rac1 [98], and Cdc42 [99] are PLD1 activators *in vivo*. The C3-exoenzyme from *Clostridium botulinum*, which

inhibits the functioning of Rho proteins via their ADP-ribosylation, decreases the activity of PLD1 [82].

The binding of RhoA with the C-terminus of PLD1 [33, 80, 100] depends on residues of basic amino acids of the sequence 946-962 of the enzyme and on activation site I of the RhoA protein. A similar site of Cdc42 also plays an important role in the mechanism of binding and activation of PLD1 [101]. Other important binding sites for RhoA on the C-terminus of human PLD1 are residues Ile870, Gln975, and Asp999 [102]. The Rho proteins interact with PLD1 only in an active GTP-bound state [103]. Some Rho GTPases (RhoA, Rac1, and Cdc42) stimulate PLD1 synergistically with ARF1, but not with PKCα [80]. Although RalA GTPases cannot directly stimulate PLD *in vitro* and *in vivo*, they play an important role in the activation of PLD1 and PLD2 [104, 105] by forming a complex with Rho and ARF proteins [106, 107]. This evidences in favor of the synergic influence of two monomeric GTPases on PLD activity.

PLD activity also depends on some cytosolic GTPases representing ADP-ribosylation factors (ARF). The PLD1 activity *in vitro* increases 10-15-fold on incubation with ARF proteins (ARF1, ARF3, ARF4, ARF5, and ARF6) [26, 47, 108], whereas the activity of PLD2 increases 1.5-2-fold (but independently of GTP) [54]. ARF1 and ARF6 activate PLD1 *in vivo* [109, 110], whereas ARF6 activates PLD2 [111]. Comparatively to ARF6, ARF1 preferentially activates PLD1 *in vitro* [112], whereas ARF6 unlike ARF1 markedly stimulates PLD2 *in vivo* [108]. ARF1 stimulates PLD1 *in vitro* synergistically with Rho proteins (RhoA, Rac1, Cdc42) [80, 112] and PKCα [80]. Synergistic effect of ARF1 and Rho was observed only at optimal concentrations of PtdInsP₂ [80].

The translocation of ARF to membranes plays an important role in the mechanism of activation of PLD. The translocation of ARF6 stimulates PLD1 [113] and PLD2 [108] *in vivo*. ARF proteins (exemplified by ARF1) are very important as in their absence no PLD activation occurs. ARF1 activates PLD1 *in vitro* by increasing the catalytic constant without changes in the K_m value [80]. The replacement of GDP by GTP in ARF6 is important for activation of PLD [114], and this explains the stimulatory effect of guanine nucleotide exchange factors on the activity of PLD1 *in vivo* [115, 116]. The N-terminus of PLD2 (amino acids 1-308) is essential for modulation of this enzyme activity by ARF. The first 73 N-terminal amino acids of ARF are involved in the activation of PLD1 [117], whereas the sequence of amino acids 35-94 plays an important role in the interaction of these proteins [118]. Thus, small GTPases of the Rho and ARF families are key activators of PLD in animals.

Both α- and β/γ-subunits of heterotrimeric G-proteins (G_q, G_i, G₁₂) contribute to the PLD activation by various hormones in animal cells, although this process occurs *in vivo* with direct involvement of proteins Rho, protein kinase C [119, 120], ARF6, or tyrosine kinase Src

[121, 122]. The direct influence of heterotrimeric G-proteins results in inactivation of PLD. Combination of G β 1 γ 1 subunits suppresses PLD1 and PLD2, whereas G β 1 γ 1 and G β 2 γ 2 inhibit PLD1 stimulated *in vitro* and *in vivo*. These protein complexes interact with the N-terminal region of PLD within PX and PH domains (amino acid residues 3-311) [123]. In plant cells the binding of PLD α with the G-protein α -subunit (G α) in the GDP-form suppresses PLD activity *in vitro* [67, 124]. Addition of GTP into the system restores the PLD activity. In this case the interaction of PLD α 1 with G α stimulates the GTPase activity of G α leading to dissociation of the PLD α –G α complex [67]. Therefore, the inactive heterotrimeric G-protein is thought to suppress the membrane-associated PLD α , whereas stimulation of the G-protein by an extracellular signal promotes PLD α activation. Hence heterotrimeric G-proteins are thought to be important regulators of PLD in plant and animal cells.

Phosphorylation as a mechanism of PLD regulation.

Phosphorylation/dephosphorylation of proteins at specific amino acid residues plays an important role in mechanisms of signal transduction, regulation of enzyme activities, and formation of protein complexes in both plant and animal cells [125-127]. Mainly Ser, Thr, His, Asp, and Tyr residues are phosphorylated. The Tyr phosphorylation is most frequently studied in this field [127].

The stimulation of PLD in the cells by phorbol esters (in particular, by phorbol 12-myristate 13-acetate (PMA), which is a chemical analog of diacylglycerol, activator of protein kinase C) and also its suppression by specific inhibitors confirm the role of PKC as a key factor of PLD regulation *in vivo* [128]. PKC α and PKC β 1 stimulate PLD [129, 130], but PKC δ and PKC ϵ can act as both positive and negative regulators of PLD1 and PLD2 *in vivo*. PKC δ can displace PLD1 from the complex with PKC α [131, 132]. Results of *in vitro* studies have shown that human and rat PLD1 are activated under the influence of PKC α , PKC β 1, and PKC β 2 independently of their kinase activity [26, 47, 130]. The activation of PLD through protein–protein interactions can also be realized by atypical PKCs: PKC ι (PLD2) [133] and PKC ζ (PLD1, PLD2) [82, 134].

Phorbol esters as PKC activators induce the PLD1 binding with PKC α that leads to activation of PLD1 [130, 135]. PLD2 can also bind PKC α [136] and PKC δ [137]. The N- and C-terminal sequences of PLD are involved in the activation of PLD1 on binding protein kinase C α [80]. PKC α decreases the Michaelis constant and increases the catalytic constant of the PLD1 reaction. Unlike Rho, PKC α activates PLD1 *in vitro* synergistically with Arf1. This occurs only at optimal concentrations of PtdInsP $_2$ [80]. The binding of active PKC α with PLD2 also stimulates PLD2 *in vivo*, whereas the phosphorylation of PLD2 *in vitro* in many Ser and Thr sites by PKC α inactivates it [136, 138]. However, the phosphorylation of Thr566 under the influence of PKC δ is accompanied by

activation of PLD2 [131, 137]. Phosphorylation by PKC α and PKC β of PLD2 at Ser and Thr leads to translocation of PLD2 to membranes and its activation *in vivo* [139]. However, PKC α and PKC β phosphorylate and as a result inactivate PLD1 *in vitro* [140]. The phosphorylation of PLD1 at Thr under the influence of PKC α suppresses PLD1 *in vivo* [141], and the translocation of PLD1 to plasma membranes and its resulting activation are due to phosphorylation of the enzyme by protein kinase C α at Ser2, Thr147, and Ser561 [142].

Activation of growth factor receptors stimulates soluble protein tyrosine kinases of the Src family. Inhibitors of Src are known to suppress the PLD2 activation in PC12 cells caused by depolarization [143]. Members of the Src family p60Src and p56Lyn mediate PLD activation in platelets [144]. In contrast to PLD1, PLD2 is phosphorylated on Tyr and is activated by other tyrosine kinases of the Src family – Fyn and Fgr [145]. However, tyrosine kinase v-Src phosphorylates PLD1 and PLD2 on Tyr in COS-7 cells without influencing their activity [146]. Thyroid gland carcinoma protein kinase RET/PTC also directly phosphorylates PLD2 on Tyr without its activation [147].

The PLD2 isoform expressed as a constitutively active enzyme in various cells [148] is a tyrosine-phosphorylated protein *in vivo* [149] and *in vitro* [150]. Its phospholipase activity is regulated dually by phosphorylation/dephosphorylation and also can induce cell proliferation with involvement of Tyr179 and Tyr511 [150]. Mutations of this and other tyrosine residues – Tyr296 and Tyr415 – result in a decrease in the PLD2 activity [151], whereas the phosphorylation level of each of them catalyzed by different enzymes can influence the functioning of PLD2. The phosphorylation of PLD2 at Y296 in the PH domain by the epidermal growth factor receptor provides for the low activity of the enzyme *in vitro* only in tumor cells. However, tyrosine kinases JAK3 and Src phosphorylating PLD2 at Tyr415 and Tyr511, respectively, are responsible for the high activity of this enzyme *in vitro* in less invasive cells and thus prevent the negative effect of the Tyr296 phosphorylation. The Tyr511 residue is a dual site for regulation of the PLD2 activity. Dephosphorylation at Tyr511 and Tyr296 by tyrosine phosphatases CD45 and PTP1B stimulates PLD, whereas dephosphorylation at Tyr415 decreases the activity of this enzyme *in vitro* [150]. Phosphorylated Tyr179 and Tyr511 in PLD2 promote its binding with the SH2 domain of the adapter protein Grb2, which ensures a high lipase activity of the enzyme in COS-7 cells [151]. Thus, it is concluded that nonreceptor tyrosine kinases/phosphatases modulate PLD activity through direct phosphorylation/dephosphorylation of this enzyme on Tyr residues.

PLD activity sharply increases in tumor cell lines in response to stress caused by removal of serum. Dephosphorylation of Tyr179 and Tyr511 in PLD2 can

act as a signal directed to decrease the activity of PLD together with the accompanying proliferative potential capable of preventing cell malignization [151, 152]. The increased expression and activity of PLD have been found in cells of many human tumors, which suggests a functional association of the PLD activity with cancer signals and tumorigenesis [153]. Results of these studies have allowed new nontoxic PLD inhibitors to be created as promising antitumor drugs [153, 154].

Ser/Thr-protein kinases also play an important role in the regulation of PLD in animal cells. The mechanism of activation of PLD by casein kinase II (PLD2) [155], cyclin-dependent protein kinase 5 (PLD2) [156], ribosomal S6-kinase 2 (PLD1) [157], and also by AMP-activated protein kinase (PLD1) [158] is largely due to phosphorylation of the specific residues Ser134 [156], Thr147 (in the PX domain) [157], and Ser505 [158]. However, it is still unclear whether the activation of PLD is caused by protein kinase Rho dependent on the Rho protein [159] or by calcium/calmodulin-derived protein kinase II [144] phosphorylation of the enzyme and whether the PLD2 phosphorylation on the active Ser/Thr protein kinase AKT (Thr175 in the PX domain) [160] or by protein kinase A in the case of PLD1 [161] changes the enzyme activity. It is also unclear whether the activation of PLD [162] is associated with the enzyme phosphorylation on Ser and Thr by MAP kinases p38, ERK1, and ERK2 [163–165] or is caused only by protein–protein interactions.

Incubation of plasma membrane from cells of the plant *B. oleracea* with acidic phosphatase *in vitro* decreases the activity of PtdInsP₂-dependent PLD by 80%, whereas inactivation of the phosphatase partially restores the activity of the PLD. Moreover, PLD γ of this plant was shown to be recognized by antibodies against phosphoserine/threonine/tyrosine [68]. The presence of phosphorylation sites in the structure of PLD γ of *B. oleracea* [68] determines the role of this posttranslational modification in the regulation of PLD activity in plants.

Influence of cytoskeleton and light on PLD activity.

The cytoskeletal proteins are modifiers of PLD. PLD activity depends on the degree of actin polymerization. In the presence of active monomeric GTPases, monomeric G-actin inactivates PLD1 in the membrane fractions or negatively influences this enzyme obtained by immunoprecipitation [166, 167]. However, G-actin stimulates a purified recombinant enzyme PLD1 expressed in cells of insects [63]. A highly purified polymerized F-actin stimulates PLD1 *in vitro* but only in the presence of activators — G-proteins ARF and Rac1 [2, 63]. The translocation of PLD1 to actin filaments is caused by phosphorylation of the Ser2 residue on the PLD N-terminus by protein kinase C [63].

The polymerized F-actin stimulates plant PLD activity, whereas monomeric G-actin suppresses it [2]. In tobacco cells PLD β 1 occurs to be a partner of actin, and

the interaction with actin is provided for by Asn323 and Thr382 of the phospholipase [64]. Thus, the interaction of actin and PLD β 1 creates a positive feedback system in which F-actin activates phospholipase D and lipase-produced phosphatidic acid, in turn, activates F-actin in sites of its interaction with cell membranes [2, 64].

The activity of animal PLD2 is suppressed *in vitro* by β -actin, which prevents the access of PLD2 to the substrate [166]. The protein α -actinin that links actin filaments suppresses PLD2 *in vitro* by affecting its binding with ARF1 [167]. The negative effect of β -actin is inhibited in the presence of ARF1, whereas the binding of β -actin with PLD2 is prevented by association of α -actinin with this enzyme [166]. The phosphorylated form of cofilin (a protein promoting actin polymerization and depolymerization) specifically binds and stimulates the basal activity of PLD1 *in vitro* by promoting its translocation to the plasma membrane [168]. Gelsolin (an actin-capping protein) suppresses PLD1 and PLD2 *in vitro* [169], but in the presence of nucleotide triphosphate binds to PLD and activates it [170]. The actin-binding protein fodrin [171] and also dimeric and trimeric spectrin suppress PLD activity and are involved in local regulation of PLD by actin [171]. However, the effect of two latter proteins might be caused by disturbance of the interaction of PLD1 with PtdInsP₂ [172].

Tubulin monomers were shown to inhibit PLD2 of animal cells due to direct and reversible binding with the enzyme [173]. In plants, stabilizers and destabilizers of microtubules stimulate the *in vivo* accumulation of phosphatidic acids produced by PLD [174]. Therefore, it is concluded that cytoskeleton proteins can act as regulators of PLD activity in plant and animal cells providing the required level and location of phosphatidic acid in the cell.

Plant PLDs are known to be so-called dark-induced proteins (din-proteins). The role of a photoregulatory receptor is played by phytochrome: both white and red lights suppress PLD activity in etiolated plantlets, whereas far red light restores it (a reversible photo-effect) [175, 176]. This indicates a possible involvement of the phytochrome receptor in the regulation of PLD activity. Moreover, the photo-dependence of PLD activity is associated with photosynthesis, because diuron stimulates and glucose suppresses, respectively, the enzyme activity in green and etiolated oat plantlets.

Involvement of PLD in hormone signaling. PLDs are involved in perception and intracellular transduction of hormonal signals in both animals and plants. In animals PLDs are activated by hormones (growth hormone, vasopressin, gonadotropin-releasing hormone, etc.) and various neurotransmitters (histamine, bradykinin, noradrenaline, etc.) [177]. Interesting data were recently obtained in studies on transmembrane receptors of animal hormones. Phospholipase D was shown to be a key enzyme for many of them, being involved in their secretion, endo-

Properties and role of different PLD isoenzymes in regulation of plant cell metabolism

PLD isoenzyme	Molecular weight, kDa	Substrate specificity	Cofactors			Regulatory mechanisms			pH-optimum	Localization in chromosomes		Sensitivity to hormones and stresses of PLD mutants (↑ – increase, ↓ – decrease)	
			calcium	PtdInsP ₂	oleic acid	phosphorylation	G-proteins	actin		<i>A. thaliana</i> L.	<i>Oryza sativa</i> L.	knockout/knockdown	over-expression
PLD α	91.8 [2]	PC > PE [188]	mM, μ M [2, 11]	no [2, 11]			yes [67]	yes [64]	5.5-6.5 [11]	III, I, V [2]	I, III, V, VI, IX [28]	ABA (↓) [28, 181-183, 189, 190], Et (↓) [182], Glc (↓) [191], Ss (↑) [29, 192, 193], Os (↑) [29, 193], Dr (↑) [193-195], Fz (↓) [196-199], W (↓) [200, 201]	ABA (↑) [183], Glc (↑) [191], Ss (↓) [29], Os (↓) [29], Dr (↓) [183]
PLD β	109 [2]	PC = PE [188]	μ M [43]	yes [43]			yes [67]	yes [2, 64]	7 [43]	II, IV [43]	III, X [28]	ABA (↓) [28], G (↑) [28], Bs (↓) [195], Ss (↑) [28]	
PLD γ	96 [2]	PE > PC [188]	μ M [2, 73]	yes [2, 73]		yes [68]	yes [67]		6.5-7.5 [2, 73]	IV [2]			
PLD δ	97.7 [11]	PE > PC [2, 83]	μ M, mM [11]	yes [11]	yes [11]		yes [67]	yes [64]	7 [11]	IV [2]	III, VII, IX [28]	Oxs (↑) [202], Fz (↑) [198, 199], Ss (↑) [193]	Fz (↓) [199]
PLD ϵ		PC = PE > PG [30]	mM, μ M [30]	yes [30]	yes [30]		yes [67]			I [11]		N starvation (↑), Os, Ss (↑) [30]	N starvation (↓), Os, Ss (↓) [30]
PLD ζ	124 (ζ 1) [2]; 118 (ζ 2) [31]	PC [2]	no [2]	yes [2]			no [67]		7 [2]	III [2]	I, V [28]	IAA (↓) [184], transport of IAA (↓) [184, 185], P starvation (↑) [203, 204]	transport of IAA (↑) [184, 185]
PLD κ											II [28]		
PLD ϕ											VI [28]		

Note: ABA, abscisic acid; Bs, biotic stress; G, gibberellin; Fz, freezing; Dr, drought; IAA, indolylacetic acid; Oxs, oxidative stress; Os, osmotic stress; W, wound; Ss, salt stress; P, phosphorus; Et, ethylene; N, nitrogen; PC, phosphatidylcholine; PG, phosphatidylglycerol; PE, phosphatidylethanolamine.

cytosis, and signaling. For example, the parathyroid hormone 1 type receptor (PTH1R) upon interaction with the ligand increased PLD activity, PLD1 being involved in the internalization and intracellular traffic while PLD2 in receptor endocytosis [178]. The subclass of formyl peptide receptors (FPR1) stimulated by an agonist also can activate PLD, and the phospholipase plays an important role in endocytosis, membrane recycling, and receptor signaling [179, 180]. PLD2 acting in plasma membrane similarly promotes endocytosis of angiotensin II receptor type 1 [40].

The involvement of PLD in the transduction of abscisic acid signal has been shown in plants [28, 181, 182]. Moreover, there are data on the involvement of PLD in the action of gibberellins [28] and in auxin transport [184, 185] (table). PLD also plays a role in the activation by cytokinins of the primary response genes [77, 186, 187], and a rapid increase in PLD activity was observed upon cytokinin treatment [69, 76]. The PLDs involved in transduction of signals of different phytohormones are also different. The abscisic acid signal is transduced with the involvement of phospholipase D isoforms α and partially β (table) [28, 181], whereas the cytokinin signals are transduced not with the PLD α -isoform but more likely with the involvement of PLD γ [76]. Thus, PLD is an important and specific factor in the signal transduction of various hormones in both animals and plants.

PLD is a widespread enzyme involved in the fine regulation of metabolism of cells with different types of organization. This enzyme is necessary for growth and development of animals and plants, especially under stress and light. The primary structure of PLD is known in detail, and regulatory mechanisms of activities of these enzymes and their interaction with other signaling pathways in cell are intensively studied. It has been shown that the activity of known isoforms of animal PLDs is regulated by serine/threonine and tyrosine protein kinases, GTP-binding proteins (ARF and Rho), the cytoskeleton, polyphosphoinositides, protein-protein interactions, and hormones. In plants PLDs are directly regulated by Ca^{2+} , polyphosphoinositides, heterotrimeric G-proteins, cytoskeleton, some phytohormones, and possibly by protein kinases. However, mechanisms of *in vivo* regulation of individual PLD enzymes are studied insufficiently, especially in plants, despite the established biological role for many molecular PLD forms (table). In the majority of biological processes PLD activity is influenced by a number of concurrent regulators, and their action can be additive, synergic, or opposing. New data on the role of PLD in the regulation of proliferation and in carcinogenesis has stimulated intensive searches for new inhibitors of PLDs as promising antitumor preparations. To better understand the functional role of PLD isoforms in cells, *in vivo* studies are important, and this can be favored by molecular design, analysis, and synthe-

sis of new nontoxic substances inhibiting different PLD isoforms *in vivo*.

This work was supported by the State Foundation for Basic Research of the Ukraine (projects Nos. F40.4/081 and F41.4/042), National Academy of Sciences of Ukraine (No. 18-10), National Academy of Sciences of Ukraine and Academy of Sciences of Czech Republic (No. 5/1-2011), Czech Science Foundation (No. 501/11/1654), the State Foundation for Basic Research of Belarus (B09K-061, B07K-058 and B05K-108), the Russian Foundation for Basic Research (projects 11-04-90491-Ukr. and 11-04-00614), and the Russian Academy of Sciences Presidium program "Molecular and Cell Biology".

REFERENCES

1. Bargmann, B. O., and Munnik, T. (2006) *Curr. Opin. Plant Biol.*, **9**, 515-522.
2. Hong, Y., Zhang, W., and Wang, X. (2010) *Plant Cell Env.*, **33**, 627-635.
3. Rudge, S. A., and Wakelam, M. J. O. (2009) *Biochim. Biophys. Acta*, **1791**, 856-861.
4. Vitale, N. (2010) *Curr. Chem. Biol.*, **4**, 244-249.
5. Oblozinsky, M., Bezakova, L., Mansfeld, J., and Ulbrich-Hofmann, R. (2011) *Phytochemistry*, **72**, 160-165.
6. Wang, X., Devaiah, S. P., Zhang, W., and Welti, R. (2006) *Progr. Lipid Res.*, **45**, 250-278.
7. Kravets, V. S., Kolesnikov, Y. S., Kuznetsov, V. V., and Romanov, G. A. (2008) *Russ. J. Plant Physiol.*, **55**, 568-577.
8. Idkowiak-Baldys, J., Baldus, A., Raymond, J. R., and Hannun, Y. A. (2009) *J. Biol. Chem.*, **284**, 22322-22331.
9. Arisz, S. A., Testerink, C., and Munnik, T. (2009) *Biochim. Biophys. Acta*, **1791**, 869-875.
10. Zhang, W., Yu, L., Zhang, Y., and Wang, X. (2005) *Biochim. Biophys. Acta*, **1736**, 1-9.
11. Li, M., Hong, Y., and Wang, X. (2009) *Biochim. Biophys. Acta*, **1791**, 927-935.
12. Kravets, V. S., Kolesnikov, Y. S., Kretynin, S. V., Kabachevskaya, E. M., Lyakhovich, G. V., Bondarenko, O. M., Volotovskii, I. D., and Kukhar, V. P. (2010) *Biopolymers and Cell*, **26**, 175-186.
13. Foster, D. A. (2006) *Curr. Sign. Trans. Ther.*, **1**, 295-303.
14. Hamik, A., and Jain, M. K. (2010) *Blood*, **116**, 1194-1196.
15. Gomez-Cambronero, J., Di Fulvio, M., and Knappek, K. (2007) *J. Leukocyte Biol.*, **82**, 272-281.
16. Vorland, M., Thoursen, V. A. T., and Holmsen, H. (2008) *Platelets*, **19**, 582-594.
17. Kanaho, Y., Funakoshi, Y., and Hasegawa, H. (2009) *Biochim. Biophys. Acta*, **1791**, 898-904.
18. Tappia, P. S., Dent, M. R., and Dhalla, N. S. (2006) *Free Rad. Biol. Med.*, **41**, 349-361.
19. Cummings, R., Parinandi, N., Wang, L., Usatyuk, P., and Natarajan, V. (2002) *Mol. Cell. Biochem.*, **235**, 99-109.
20. Roth, M. G. (2008) *Traffic*, **9**, 1233-1239.
21. Bader, M.-F., and Vitale, N. (2009) *Biochim. Biophys. Acta*, **1791**, 936-941.

22. Lee, C. S., Kim, K. L., Jang, J. H., Choi, Y. S., Suh, P.-G., and Ryu, S. H. (2009) *Biochim. Biophys. Acta*, **1791**, 862-868.
23. Cazzolli, R., Shemon, A. N., Fang, M. Q., and Hughes, W. E. (2006) *IUBMB Life*, **58**, 457-461.
24. Becker, K. P., and Hannun, Y. A. (2005) *Cell. Mol. Life Sci.*, **62**, 1448-1461.
25. Weernink, P. A. O., Han, L., Jakobs, K. H., and Schmidt, M. (2007) *Biochim. Biophys. Acta*, **1768**, 888-900.
26. Hammond, S. M., Jenco, J. M., Nakashima, S., Cadwallader, K., Gu, Q., Cook, S., Nozawa, Y., Prestwich, G. D., Frohman, M. A., and Morris, A. J. (1997) *J. Biol. Chem.*, **272**, 3860-3868.
27. Colley, W. C., Sung, T., Roll, R., Jenco, J., Hammond, S. M., Altshuller, Y., Bar-Sagi, D., Morris, A. J., and Frohman, M. A. (1997) *Curr. Biol.*, **7**, 191-201.
28. Li, G., Lin, F., and Xue, H.-W. (2007) *Cell Res.*, **17**, 881-894.
29. Hong, Y., Pan, X., Welti, R., and Wang, X. (2008) *Plant Cell*, **20**, 803-816.
30. Hong, Y., Devaiah, S. P., Bahn, S. C., Thamasandra, B. N., Li, M., Welti, R., and Wang, X. (2009) *Plant J.*, **58**, 376-387.
31. Yamaryo, Y., Dubots, E., Albrieux, C., Baldan, B., and Block, M. A. (2008) *FEBS Lett.*, **582**, 685-690.
32. Ponting, C. P., and Kerr, I. D. (1996) *Protein Sci.*, **5**, 914-922.
33. Sung, T. C., Roper, R., Zhang, Y., Rudge, S. A., Temel, R., Hammond, S. M., Morris, A. J., Moss, B., Engebrecht, J., and Frohman, M. A. (1997) *EMBO J.*, **16**, 4519-4530.
34. Lerchner, A., Mansfeld, J., Kuppe, K., and Ulbrich-Hofmann, R. (2006) *Protein Eng. Des. Sel.*, **19**, 443-452.
35. Gottlin, E. B., Rudolph, A. E., Zhao, Y., Mathews, H. R., and Dixon, J. E. (1998) *Proc. Natl. Acad. Sci. USA*, **95**, 9202-9207.
36. Orth, E. S., Brandao, T. A., Souza, B. S., Pliego, J. R., Vaz, B. G., Eberlin, M. N., Kirby, A. J., and Nome, F. (2010) *J. Am. Chem. Soc.*, **132**, 8513-8523.
37. Sciorra, V. A., Rudge, S. A., Prestwich, G. D., Frohman, M. A., Engebrecht, J., and Morris, A. J. (1999) *EMBO J.*, **18**, 5911-5921.
38. Sciorra, V. A., Rudge, S. A., Wang, J., McLaughlin, S., Engebrecht, J., and Morris, A. J. (2002) *J. Cell Biol.*, **159**, 1039-1049.
39. Du, G., Altshuller, Y. M., Vitale, N., Huang, P., Chasserot-Golaz, S., Morris, A. J., Bader, M. F., and Frohman, M. A. (2003) *J. Cell Biol.*, **162**, 305-315.
40. Du, G., Huang, P., Liang, B. T., and Frohman, M. A. (2004) *Mol. Biol. Cell*, **15**, 1024-1030.
41. Pinhero, R. G., Almquist, K. C., Novotna, Z., and Paliyath, G. (2003) *Plant Physiol. Biochem.*, **41**, 223-240.
42. Yuan, H., Chen, L., Paliyath, G., Sullivan, A., and Murr, D. P. (2005) *Plant Physiol. Biochem.*, **43**, 535-547.
43. Pappan, K., Qin, W., Dyer, J. H., Zheng, L., and Wang, X. (1997) *J. Biol. Chem.*, **272**, 7055-7061.
44. Schaffner, I., Rucknagel, K.-P., Mansfeld, J., and Ulbrich-Hofmann, R. (2002) *Eur. J. Lipid Sci. Technol.*, **104**, 79-87.
45. Lerchner, A., Mansfeld, J., Schaffner, I., Schops, R., Beer, H. K., and Ulbrich-Hofmann, R. (2005) *Biochim. Biophys. Acta*, **1737**, 94-101.
46. Tiwari, K., and Paliyath, G. (2011) *Plant Physiol. Biochem.*, **49**, 18-32.
47. Hodgkin, M. N., Masson, M. R., Powner, D., Saquib, K. M., Ponting, C. P., and Wakelam, M. J. O. (2000) *Curr. Biol.*, **10**, 43-46.
48. Stahelin, R. V., Ananthanarayanan, B., Blatner, N. R., Singh, S., Bruzik, K. S., Murray, D., and Cho, W. (2004) *J. Biol. Chem.*, **279**, 54918-54926.
49. Lee, J. S., Kim, J. H., Jang, I. H., Kim, H. S., Han, J. M., Kazlauskas, A., Yagisawa, H., Suh, P. G., and Ryu, S. H. (2005) *J. Cell Sci.*, **118**, 4405-4413.
50. Lee, C. S., Kim, I. S., Park, J. B., Lee, M. N., Lee, H. Y., Suh, P.-G., and Ryu, S. H. (2006) *Nat. Cell. Biol.*, **8**, 477-484.
51. Jang, I. H., Lee, S., Park, J. B., Kim, J. H., Lee, C. S., Hur, E. M., Kim, I. S., Kim, K. T., Yagisawa, H., Suh, P. G., and Ryu, S. H. (2003) *J. Biol. Chem.*, **278**, 18184-18190.
52. Lee, J. H., Kim, Y. M., Kim, N. W., Kim, J. W., Her, E., Kim, B. K., Kim, J. H., Ryu, S. H., Park, J. W., Seo, D. W., Han, J. W., Beaven, M. A., and Choi, W. S. (2006) *Blood*, **108**, 956-964.
53. Kim, J. H., Kim, J. H., Ohba, M., Suh, P. G., and Ryu, S. H. (2005) *Mol. Cell. Biol.*, **25**, 3194-3208.
54. Sung, T. C., Zhang, Y., Morris, A. J., and Frohman, M. A. (1999) *J. Biol. Chem.*, **274**, 3659-3666.
55. Sugars, J. M., Celtek, S., Manifava, M., Coadwell, J., and Ktsitakis, N. T. (1999) *J. Biol. Chem.*, **274**, 30023-30027.
56. Sugars, J. M., Celtek, S., Manifava, M., Coadwell, J., and Ktsitakis, N. T. (2002) *J. Biol. Chem.*, **277**, 29152-29161.
57. Xie, Z., Ho, W., and Exton, J. H. (2001) *J. Biol. Chem.*, **276**, 9383-9391.
58. Riebeling, C., Bourgoin, S., and Shields, D. (2008) *Biochim. Biophys. Acta*, **1781**, 376-382.
59. Simoes, I., Mueller, E.-C., Otto, A., Bur, D., Cheung, A. Y., Faro, C., and Pires, E. (2005) *FEBS J.*, **272**, 5786-5798.
60. Stumpe, S., Konig, S., and Ulbrich-Hofmann, R. (2007) *FEBS J.*, **274**, 2630-2640.
61. Pappan, K., Zheng, L., Krishnamoorthi, R., and Wang, X. (2004) *J. Biol. Chem.*, **279**, 47833-47839.
62. Kook, S., and Exton, J. H. (2005) *Cell. Signal.*, **17**, 1423-1432.
63. Farquhar, M. J., Powner, D. J., Levine, B. A., Wright, M. H., Ladds, G., and Hodgkin, M. N. (2007) *Cell. Signal.*, **19**, 349-358.
64. Pleskot, R., Potocky, M., Pejchar, P., Linek, J., Bezvoda, R., Martinec, J., Valentova, O., Novotna, Z., and Zarsky, V. (2010) *Plant J.*, **62**, 494-507.
65. Hughes, W. E., and Parker, P. (2001) *Biochem. J.*, **356**, 727-736.
66. Liu, M., Gutowski, S., and Sternweis, P. C. (2001) *J. Biol. Chem.*, **276**, 5556-5562.
67. Zhao, J., and Wang, X. (2004) *J. Biol. Chem.*, **279**, 1794-1800.
68. Novotna, Z., Linek, J., Hynek, R., Martinec, J., Potocky, M., and Valentova, O. (2003) *FEBS Lett.*, **554**, 50-54.
69. Kravets, V. S., Kretynin, S. V., Kolesnikov, Y. S., Getman, I. A., and Romanov, G. A. (2009) *Doklady Biochem. Biophys.*, **428**, 264-267.
70. Kravets, V. S., Kolesnikov, Y. S., Kretynin, S. V., and Romanov, G. A. (2010) *Ukrain. Bot. J.*, **67**, 189-194.
71. Ali, Y. B., Carriere, F., and Abousalham, A. (2007) *Protein Expr. Purif.*, **51**, 162-169.
72. Moreno-Perez, A. J., Martinez-Force, E., Garces, R., and Salas, J. J. (2010) *J. Plant Physiol.*, **167**, 503-511.

73. Qin, C., Li, M., Qin, W., Bahn, S. C., Wang, C., and Wang, X. (2006) *Biochim. Biophys. Acta*, **1761**, 1450-1458.
74. Khatoon, H., Talat, S., and Younus, H. (2007) *Int. J. Biol. Macromol.*, **40**, 232-236.
75. Munnik, T., and Vermeer, J. E. (2010) *Plant Cell Environ.*, **33**, 655-669.
76. Kravets, V. S., Kolesnikov, Ya. S., Kretynin, S. V., Getman, I. A., and Romanov, G. A. (2010) *Physiol. Plant.*, **138**, 249-255.
77. Amini, A., Glevarec, G., Andreu, F., Reverdiau, P., Rideau, M., and Creche, J. (2008) *J. Plant Growth Regul.*, **27**, 394-399.
78. Tarasova, O. V., and Medvedev, S. S. (2008) *Vestnik St.-Petersburg Univ.*, **3**, 85-90.
79. Divecha, N., Roefs, M., Halstead, J. R., D'Andrea, S., Fernandez-Borga, M., Oomen, L., Saquib, K. M., Wakelam, M. J. O., and D'Santos, C. (2000) *EMBO J.*, **19**, 5440-5449.
80. Henage, L. G., Exton, J. H., and Brown, H. A. (2006) *J. Biol. Chem.*, **281**, 3408-3417.
81. Brizuela, L., Rabano, M., Gangoiti, P., Narbona, N., Macarulla, J. M., Trueba, M., and Gomez-Munoz, A. (2007) *J. Lipid Res.*, **48**, 2264-2274.
82. Gayral, S., Deleris, P., Laulagnier, K., Laffargue, M., Salles, J.-P., Perret, B., Record, M., and Breton-Douillon, M. (2006) *Circ. Res.*, **99**, 132-139.
83. Qin, C., Wang, C., and Wang, X. (2002) *J. Biol. Chem.*, **277**, 49685-49690.
84. Mansfield, P. J., Carey, S. S., Hinkovska-Galcheva, V., Shayman, J. A., and Boxer, L. A. (2004) *Blood*, **103**, 2363-2368.
85. Singh, I. N., Stromberg, L. M., Bourgoin, S. G., Sciorra, V. A., Morris, A. J., and Brindley, D. N. (2001) *Biochemistry*, **40**, 11227-11233.
86. Webb, L. M., Arnholt, A. T., and Venable, M. E. (2010) *Mol. Cell Biochem.*, **337**, 153-158.
87. Ryu, S. B., and Palta, J. P. (2000) *J. Lipid Res.*, **41**, 940-944.
88. Ryu, S. B., Karlsson, B. H., Ozgen, M., and Palta, J. P. (1997) *Proc. Natl. Acad. Sci. USA*, **94**, 12717-12721.
89. Kim, J. H., Kim, Y., Lee, S. D., Lopez, I., Arnold, R. S., Lambeth, J. D., Suh, P., and Ryu, S. H. (1999) *FEBS Lett.*, **454**, 42-46.
90. Massenburg, D., Han, J. S., Liyanage, M., Patton, W. A., Rhee, S. G., Moss, J., and Vaughan, M. (1994) *Proc. Natl. Acad. Sci. USA*, **91**, 11718-11722.
91. Austin-Brown, S. L., and Chapman, K. D. (2002) *Plant Physiol.*, **129**, 1892-1898.
92. Karlsson, R., Pedersen, E. D., Wang, Z., and Brakebusch, C. (2009) *Biochim. Biophys. Acta*, **1796**, 91-98.
93. Min, D. S., and Exton, J. H. (1998) *Biochem. Biophys. Res. Commun.*, **248**, 533-537.
94. Hodgkin, M. N., Clark, J. M., Rose, S., Saquib, K., and Wakelam, M. J. O. (1999) *Biochem. J.*, **339**, 87-93.
95. Subra, C., Grand, D., Laulagnier, K., Stella, A., Lambeau, G., Paillasse, M., De Medina, P., Monsarrat, B., Perret, B., Silvente-Poirot, S., Poirot, M., and Record, M. (2010) *J. Lipid Res.*, **51**, 2105-2120.
96. Salvador, G. A., and Giusto, N. M. (2006) *Exper. Eye Res.*, **83**, 202-211.
97. Everett, P. B., and Senogles, S. E. (2010) *J. Neurochem.*, **112**, 963-971.
98. Momboisse, F., Lonchamp, E., Calco, V., Ceridono, M., Vitale, N., Bader, M. F., and Gasman, S. J. (2009) *Cell. Sci.*, **122**, 798-806.
99. Yoon, M. S., Cho, C. H., Lee, K. S., and Han, J. S. (2006) *Biochem. Biophys. Res. Commun.*, **347**, 594-600.
100. Yamazaki, M., Zhang, Y., Watanabe, H., Yokozeki, T., Ohno, S., Kaibuchi, K., Shibata, H., Mukai, H., Ono, Y., Frohman, M. A., and Kanaho, Y. (1999) *J. Biol. Chem.*, **274**, 6035-6038.
101. Walker, S., Wu, W., Cerione, R. A., and Brown, H. A. (2000) *J. Biol. Chem.*, **275**, 15665-15668.
102. Cai, S., and Exton, J. (2001) *Biochem. J.*, **355**, 799-785.
103. Genth, H., Schmidt, M., Gerhard, R., Aktories, K., and Just, I. (2003) *Biochem. Biophys. Res. Commun.*, **302**, 127-132.
104. Ljubicic, S., Bezzi, P., Vitale, N., and Regazzi, R. (2009) *PLoS ONE*, **4**, 1-10.
105. Corrotte, M., Nyguen, A. P., Harlay, M. L., Vitale, N., Bader, M. F., and Grant, N. G. (2010) *J. Immunol.*, **185**, 2942-2950.
106. Xu, L., Frankel, P., Jackson, D., Rotunda, T., Boshans, R. L., D'Souza-Schorey, C., and Foster, D. A. (2003) *Mol. Cell. Biol.*, **23**, 645-654.
107. Vitale, N., Mawet, J., Camonis, J., Regazzi, R., Bader, M. F., and Chasserot-Golaz, S. (2005) *J. Biol. Chem.*, **280**, 29921-29928.
108. Hiroyama, M., and Exton, J. H. (2005) *J. Cell Biochem.*, **95**, 149-164.
109. Begle, A., Tryoen-Toth, P., de Barry, J., Bader, M. F., and Vitale, N. (2009) *J. Biol. Chem.*, **284**, 4836-4845.
110. Garcia, A., Zheng, Y., Zhao, C., Toschi, A., Fan, J., Shraibman, N., Brown, H. A., Bar-Sagi, D., Foster, D. A., and Arbiser, J. L. (2008) *Clin. Cancer Res.*, **14**, 4267-4274.
111. Mitchell, R., Robertson, D. N., Holland, P. J., Collins, D., Lutz, E. M., and Johnson, M. S. (2003) *J. Biol. Chem.*, **278**, 33818-33830.
112. Perez-Mansilla, B., Ha, V. L., Justin, N., Wilkins, A. J., Carpenter, C. L., and Thomas, G. M. H. (2006) *Biochim. Biophys. Acta*, **1761**, 1429-1442.
113. Ma, W.-N., Park, S.-Y., and Han, J.-S. (2010) *Exp. Mol. Med.*, **42**, 456-464.
114. Muralidharan-Chari, V., Clancy, J., Plou, C., Romao, M., Chavrier, P., Raposo, G., and D'Souza-Schorey, C. (2009) *Curr. Biol.*, **19**, 1875-1885.
115. El Azreq, M. A., Garceau, V., Harbour, D., Pivot-Pajot, C., and Bourgoin, S. G. (2010) *J. Immunol.*, **184**, 637-649.
116. Bach, A. S., Enjalbert, S., Comunale, F., Bodin, S., Vitale, N., Charrasse, S., and Gauthier-Rouviere, C. (2010) *Mol. Biol. Cell.*, **21**, 2412-2424.
117. Zhang, G. F., Patton, W. A., Lee, F. J., Liyanage, M., Han, J. S., Rhee, S. G., Moss, J., and Vaughan, M. (1995) *J. Biol. Chem.*, **270**, 21-24.
118. Liang, J. O., Sung, T. C., Morris, A. J., Frohman, M. A., and Kornfeld, S. (1997) *J. Biol. Chem.*, **272**, 33001-33008.
119. Du, G., Altshuller, Y. M., Kim, Y., Han, J. M., Ryu, S. H., Morris, A. J., and Frohman, M. A. (2000) *Mol. Biol. Cell*, **11**, 4359-4368.
120. Xie, Z., Ho, W. T., Spellman, R., Cai, S., and Exton, J. H. (2002) *J. Biol. Chem.*, **277**, 11979-11986.
121. Ushio-Fukai, M., Alexander, R. W., Akers, M., Lyons, P. R., Lassegue, B., and Griendling, K. K. (1999) *Mol. Pharmacol.*, **55**, 142-149.

122. Le Stunff, H., Dokhac, L., Bourgoin, S., Bader, M. F., and Harbon, S. (2000) *Biochem. J.*, **352**, 491-499.
123. Preininger, A. M., Henage, L. G., Oldham, W. M., Yoon, E. J., Hamm, H. E., and Brown, H. A. (2006) *Mol. Pharmacol.*, **70**, 311-318.
124. Lein, W., and Saalbach, G. (2001) *Biochim. Biophys. Acta*, **1530**, 172-183.
125. Ubersax, J. A., and Ferrell, J. E. (2007) *Nature Rev. Mol. Cell Biol.*, **8**, 530-541.
126. Lee, H. K., Yeo, S., Kim, J. S., Lee, J. G., Bae, Y. S., Lee, C., and Baek, S. H. (2010) *Mol. Pharmacol.*, **78**, 478-485.
127. Fedina, E. O., Karimova, F. G., and Tarchevsky, I. A. (2006) *Biochemistry (Moscow)*, **71**, 423-429.
128. Shemon, A. N., Sluyter, R., and Wiley, J. S. (2007) *Immunol. Cell Biol.*, **85**, 68-72.
129. Eldar, H., Ben-Av, P., Schmidt, U. S., Livneh, E., and Liscovitch, M. (1993) *J. Biol. Chem.*, **268**, 12560-12564.
130. Jin, J. K., Kim, N. H., Lee, Y. J., Kim, Y. S., Choi, E. K., Kozlowski, P. B., Park, M. H., Kim, H. S., and Min, D. S. (2006) *Neurosci. Lett.*, **407**, 263-267.
131. Meacci, E., Nuti, F., Catarzi, S., Vasta, V., Donati, C., Bourgoin, S., Bruni, P., Moss, J., and Vaughan, M. (2003) *Biochemistry*, **42**, 284-292.
132. Gorshkova, I., He, D., Berdyshev, E., Usatuyk, P., Burns, M., Kalari, S., Zhao, Y., Pendyala, S., Garcia, J. G., Pyne, N. J., Brindley, D. N., and Natarajan, V. (2008) *J. Biol. Chem.*, **283**, 11794-11806.
133. Mwanjewe, J., Spitaler, M., Ebner, M., Windegger, M., Geiger, M., Kampfer, S., Hofmann, J., Uberall, F., and Grunicke, H. H. (2001) *Biochem. J.*, **359**, 211-217.
134. Parmentier, J.-H., Pavicevic, Z., and Malik, K. U. (2006) *Am. J. Physiol. Heart Circ. Physiol.*, **290**, 46-54.
135. Hu, T., and Exton, J. H. (2004) *J. Biol. Chem.*, **279**, 35702-35708.
136. Chen, J.-S., and Exton, J. H. (2004) *J. Biol. Chem.*, **279**, 22076-22083.
137. Chae, Y. C., Kim, K. L., Ha, S. H., Kim, J., Suh, P.-G., and Ryu, S.-H. (2010) *Mol. Cell. Biol.*, **30**, 5086-5098.
138. Chen, J. S., and Exton, J. H. (2005) *Biochem. Biophys. Res. Commun.*, **333**, 1322-1326.
139. Kim, S., Choi, M. S., and Choi, M.-U. (2005) *Exp. Mol. Med.*, **37**, 418-426.
140. Min, D. S., Kim, E., and Exton, J. (1998) *J. Biol. Chem.*, **273**, 29986-29994.
141. Hu, T., and Exton, J. H. (2003) *J. Biol. Chem.*, **278**, 2348-2355.
142. Han, J. M., Kim, Y., Lee, J. S., Lee, C. S., Lee, B. D., Ohba, M., Kuroki, T., Suh, P.-G., and Ryu, S. H. (2002) *Mol. Biol. Cell*, **13**, 3976-3988.
143. Banno, Y., Nemoto, S., Murakami, M., Kimura, M., Ueno, Y., Ohguchi, K., Hara, A., Okano, Y., Kitade, Y., Onozuka, M., Murate, T., and Nozawa, Y. (2008) *J. Neurochem.*, **104**, 1372-1386.
144. Martinson, E. A., Scheible, S., and Presek, P. (1994) *Cell Mol. Biol.*, **40**, 627-634.
145. Choi, W. S., Hiragun, T., Lee, J. H., Kim, Y. M., Kim, H. P., Chahdi, A., Her, E., Han, J. W., and Beaven, M. A. (2004) *Mol. Cell. Biol.*, **24**, 6980-6992.
146. Ho, W.-T., Xie, Z., Zhao, Z. J., and Exton, J. H. (2005) *Cell. Signal.*, **17**, 691-699.
147. Kim, Y.-R., Byun, H. S., Won, M., Park, K. A., Kim, J. M., Choi, B. L., Lee, H., Hong, J. H., Park, J., Seok, J. H., Kim, D. W., Shong, M., Park, S.-K., and Hur, G. M. (2008) *BMC Cancer*, **8**, 1-11.
148. Xu, L., Shen, Y., Joseph, T., Bryant, A., Luo, J. Q., Frankel, P., Rotunda, T., and Foster, D. A. (2000) *Biochem. Biophys. Res. Commun.*, **273**, 77-83.
149. Hui, L., Zheng, Y., Yan, Y., Bargonetti, J., and Foster, D. A. (2006) *Oncogene*, **25**, 7305-7310.
150. Henkels, K. M., Peng, H.-J., Frondorf, K., and Gomez-Cambronero, J. (2010) *Mol. Cell. Biol.*, **30**, 2251-2263.
151. Henkels, K. M., Short, S., Peng, H.-J., Di Fulvio, M., and Gomez-Cambronero, J. (2009) *Biochem. Biophys. Res. Commun.*, **389**, 224-228.
152. Foster, D. A. (2009) *Biochim. Biophys. Acta*, **1791**, 949-955.
153. Su, W., Chen, Q., and Frohman, M. A. (2009) *Future Oncol.*, **5**, 1477-1486.
154. Kang, D. W., Min, G., Park, D. Y., Hong, K. W., and Min, D. S. (2010) *Exper. Mol. Med.*, **42**, 555-564.
155. Ahn, B.-H., Min, G., Bae, Y.-S., and Min, D. S. (2006) *Exper. Mol. Med.*, **38**, 55-62.
156. Lee, H. Y., Jung, H., Jang, I. H., Suh, P.-G., and Ryu, S. H. (2008) *Cell. Signal.*, **20**, 1787-1794.
157. Zeniou-Meyer, M., Liu, Y., Begle, A., Olanish, M., Hanauer, A., Becherer, U., Rettig, J., Bader, M. F., and Vitale, N. (2008) *Proc. Natl. Acad. Sci. USA*, **105**, 8434-8439.
158. Kim, J. H., Park, J. M., Yea, K., Kim, H. W., Suh, P. G., and Ryu, S. H. (2010) *PLoS One*, **5**, 1-13.
159. Schmidt, M., Vob, M., Oude Weernink, P. A., Wetzels, J., Amano, M., Kaibuchi, K., and Jakobs, K. H. (1999) *J. Biol. Chem.*, **274**, 14648-14654.
160. Di Fulvio, M., Frondorf, K., and Gomez-Cambronero, J. (2008) *Cell. Signal.*, **20**, 176-185.
161. Jang, M. J., Lee, M. J., Park, H. Y., Bae, Y. S., Min, D. S., Ryu, S. H., and Kwak, J. Y. (2004) *Exper. Mol. Med.*, **36**, 172-178.
162. Patel, S., Djerdjouri, B., Raoul-Des-Essarts, Y., Dang, P. M.-C., El-Benna, J., and Perianin, A. (2010) *J. Biol. Chem.*, **285**, 32055-32063.
163. Paruch, S., El-Benna, J., Djerdjouri, B., Marullo, S., and Perianin, A. (2006) *FASEB J.*, **20**, 142-144.
164. Varadharaj, S., Steinhour, E., Hunter, M. G., Watkins, T., Baran, C. P., Magalang, U., Kuppusamy, P., Zweier, J. L., Marsh, C. B., Natarajan, V., and Parinandi, N. L. (2006) *Cell. Signal.*, **18**, 1396-1407.
165. Wakelam, M. J. O., McNee, G. S., and Rudge, S. A. (2010) *Adv. Enzyme Regul.*, **50**, 57-61.
166. Lee, S., Park, J. B., Kim, J. H., Kim, Y., Kim, J. H., Shin, K. J., Lee, J. S., Ha, S. H., Suh, P. G., and Ryu, S. H. (2001) *J. Biol. Chem.*, **276**, 28252-28260.
167. Park, J. B., Kim, J. H., Kim, Y., Ha, S. H., Kim, J. H., Yoo, J., Du, G., Frohman, M. A., Suh, P., and Ryu, S. H. (2000) *J. Biol. Chem.*, **275**, 21295-21301.
168. Han, L., Stope, M. B., de Jesus, M. L., Oude Weernink, P. A., Urban, M., Wieland, T., Roskopf, D., Mizuno, K., Jakobs, K. H., and Schmidt, M. (2007) *EMBO J.*, **26**, 4189-4202.
169. Banno, Y., Fujita, H., Ono, Y., Nakashima, S., Ito, Y., Kuzumaki, N., and Nozawa, Y. (1999) *J. Biol. Chem.*, **274**, 27385-27391.
170. Steed, P. M., Nager, S., and Wennogle, L. P. (1996) *Biochem. J.*, **35**, 5229-5237.

171. Lukowski, S., Lecomte, M. C., Mira, J. P., Marin, P., Gautero, H., Russomarie, F., and Geny, B. (1996) *J. Biol. Chem.*, **271**, 24164-24171.
172. Lukowski, S., Mira, J. P., Zachowski, A., and Geny, B. (1998) *Biochem. Biophys. Res. Commun.*, **248**, 278-284.
173. Chae, C., Lee, S., Lee, H. Y., Heo, K., Kim, J. H., Kim, J. H., Suh, P. G., and Ryu, S. H. (2005) *J. Biol. Chem.*, **280**, 3723-3730.
174. Pejchar, P., Pleskot, R., Schwarzerova, K., Martinec, J., Valentova, O., and Novotna, Z. (2008) *Cell Biol. Int.*, **32**, 554-556.
175. Kabachevskaya, E. M., Liakhnovich, G. V., and Volotovskii, I. D. (2004) *Russ. J. Plant Physiol.*, **51**, 769-773.
176. Kabachevskaya, E. M., Liakhnovich, G. V., Kisel, M. A., and Volotovskii, I. D. (2007) *J. Plant Physiol.*, **164**, 108-110.
177. Liscovitch, M., Czarny, M., Flucci, G., and Tang, X. (2000) *Biochem. J.*, **345**, 401-415.
178. Garrido, J. L., Wheeler, D., Vega, L. L., Friedman, P. A., and Romero, G. (2009) *Mol. Endocrinol.*, **23**, 2048-2059.
179. Brandenburg, L. O., Konrad, M., Wruck, C., Koch, T., Pufe, T., and Lucius, R. (2008) *Neurosciences*, **156**, 266-276.
180. Brandenburg, L. O., Seyferth, S., Wruck, C. J., Koch, T., Rosenstiel, P., Lucius, R., and Pufe, T. (2009) *Mol. Membr. Biol.*, **26**, 371-383.
181. Zhang, W., Qin, C., Zhao, J., and Wang, X. (2004) *Proc. Natl. Acad. Sci. USA*, **101**, 9508-9513.
182. Fan, L., Zheng, S., and Wang, X. (1997) *Plant Cell*, **9**, 2183-2196.
183. Sang, Y., Zheng, S., Li, W., Huang, B., and Wang, X. (2001) *Plant J.*, **28**, 135-144.
184. Li, G., and Xue, H.-W. (2007) *Plant Cell*, **19**, 281-295.
185. Mancuso, S., Marras, A., Muguai, S., Schlicht, M., Zarsky, V., Li, G., Song, L., Xue, H. W., and Baluska, F. (2007) *Plant Signal Behav.*, **2**, 240-244.
186. Romanov, G. A., Getman, I. A., and Schmulling, T. (2000) *Plant Growth Regul.*, **32**, 337-344.
187. Romanov, G. A., Kieber, J. J., and Schmulling, T. (2002) *FEBS Lett.*, **515**, 39-43.
188. Pappan, K., Austin-Brown, S., Chapman, K. D., and Wang, X. (1998) *Arch. Biochem. Biophys.*, **353**, 131-140.
189. Liu, P. F., Chang, W. C., Wang, Y. K., Chang, H. Y., and Pan, R. L. (2008) *Biochim. Biophys. Acta*, **1779**, 164-174.
190. Zhang, Y., Zhu, H., Zhang, Q., Li, M., Yan, M., Wang, R., Wang, L., Welti, R., Zhang, W., and Wang, X. (2009) *Plant Cell*, **21**, 2357-2377.
191. Hong, Y., Pan, X., Welti, R., and Wang, X. (2008) *Plant Signal Behav.*, **3**, 1099-1100.
192. Yu, L., Nie, J., Cao, C., Jin, Y., Yan, M., Wang, F., Liu, J., Xiao, Y., Liang, Y., and Zhang, W. (2010) *New Phytol.*, **188**, 762-773.
193. Bargmann, B. O. R., Laxalt, A. M., ter Riet, B., van Schooten, B., Merquiol, E., Testerink, C., Haring, M. A., Bartels, D., and Munnik, T. (2009) *Plant Cell Physiol.*, **50**, 78-89.
194. Mane, S. P., Vasquez-Robinet, C., Sioson, A. A., Heath, L. S., and Grene, R. (2007) *J. Exper. Bot.*, **58**, 241-252.
195. Yamaguchi, T., Kuroda, M., Yamakawa, H., Ashizawa, T., Hirayae, K., Kurimoto, L., Shinya, T., and Shibuya, N. (2009) *Plant Physiol.*, **150**, 308-319.
196. Rajashekar, C. B., Zhou, H. E., Zhang, Y., Li, W., and Wang, X. (2006) *J. Plant Physiol.*, **163**, 916-926.
197. Welti, R., Li, W., Li, M., Sang, Y., Biesiada, H., Zhou, H.-E., Rajashekar, C. B., Williams, T. D., and Wang, X. (2002) *J. Biol. Chem.*, **277**, 31994-32002.
198. Li, W., Wang, R., Li, M., Li, L., Wang, C., Welti, R., and Wang, X. (2008) *J. Biol. Chem.*, **283**, 461-468.
199. Li, W., Li, M., Zhang, W., Welti, R., and Wang, X. (2004) *Nat. Biotechnol.*, **22**, 427-433.
200. Bargmann, B. O., Laxalt, A. M., Riet, B., Testerink, C., Merquiol, E., Mosblech, A., Reyes, A. L., Pieterse, C. M., Haring, M. A., Heilmann, I., Bartels, D., and Munnik, T. (2009) *Plant Cell Environ.*, **32**, 837-850.
201. Wang, C., Zien, C. A., Afithile, M., Welti, R., Hildebrand, D. F., and Wang, X. (2000) *Plant Cell*, **12**, 2237-2246.
202. Zhang, W., Wang, C., Qin, C., Wood, T., Olafsdottir, G., Welti, R., and Wang, X. (2003) *Plant Cell*, **15**, 2285-2295.
203. Cruz-Ramirez, A., Oropeza-Aburto, A., Razo-Hernandez, F., Ramirez-Chavez, E., and Herrera-Estrella, L. (2006) *Proc. Natl. Acad. Sci. USA*, **103**, 6765-6770.
204. Li, M., Qin, C., Welti, R., and Wang, X. (2006) *Plant Physiol.*, **140**, 761-770.