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Research Article

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Prostaglandin Protects Isolated Guinea Pig Chief Cells against Ethanol Injury via an Increase in Diacylglycerol

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Abstract

We studied cellular processes activated by prostaglandins (PG) that are involved in the protection of gastric chief cell injury estimated in terms of dye exclusion test, release of lactate dehydrogenase (LDH), or ^{51}Cr from prelabeled chief cells. Pretreatment of chief cells with 3×10^{-6} M PGE₂ or PGE₁ at 37°C and pH 7.4 for 15 min maximally reduced not only ethanol- but also taurocholic acid-caused LDH release from chief cells. PGs equipotently stimulated increases in the accumulation of diacylglycerol and cyclic AMP without elevating intracellular Ca²⁺ concentrations in gastric chief cells. The rank order of the potency was equal to that of PGs to reduce the injury. Pretreatment of chief cells with synthetic 1-oleoyl-2-acetyl-*sn*-glycerol (OAG) or 12-*o*-tetradecanoyl phorbol 13-acetate (TPA) reduced the injury of chief cells, while 4 α -phorbol 12,13-didecanoate, an inactive phorbol ester, failed to reduce the injury and 1-(5-isoquinolinylnsulfonyl)-2-methylpiperazine (H7) blocked the protective action of PGE₂. On the other hand, forskolin and dbcAMP had no effect on ethanol-caused LDH release and diacylglycerol formation in chief cells. These results suggest that PGE₂ and PGE₁ possess the direct protective action against ethanol- or taurocholic acid-caused injury in chief cells, presumably through the activation of the diacylglycerol/protein kinase C signaling pathway. (*J. Clin. Invest.* 1990. 86:1897-1903.) Key words: prostaglandin • lactate dehydrogenase

Introduction

Prostaglandins (PG)¹ have been shown to possess a variety of actions, such as the stimulation of mucous and bicarbonate secretion, mucosal blood flow, and ion transport and the inhi-

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1. Abbreviations used in this paper: [Ca²⁺]_i, intracellular free Ca²⁺ concentration; CCh, carbachol; CCK8, COOH-terminal octapeptide of cholecystokinin; dbcAMP, dibutyryl cyclic AMP; DG, 1,2-*sn*-diacylglycerol; 16,16dmPGE₂, 16,16 dimethyl prostaglandin E₂; fura-2/AM, fura-2 acetoxymethyl; H7, 1-(5-isoquinolinylnsulfonyl)-2-methylpiperazine; IBMX, isobutyl-1-methylxanthine; LDH, lactate dehydrogenase; OAG, 1-oleoyl-2-acetyl-*sn*-glycerol; PDBu, phorbol 12,13-dibutyrate; 4 α PDD, 4 α -phorbol 12,13-didecanoate; PG, prostaglandin; TPA, 12-*o*-tetradecanoylphorbol 13-acetate.

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tion of gastric acid secretion in the stomach (1-3). Some of the effects may contribute to reduce injury to the gastric mucosa caused by various noxious agents lumenally applied in humans and experimental animals (4-6). This PG action, which seems to be independent of their inhibitory action on gastric acid secretion, is referred to as cytoprotection (7). However, in addition to the mechanisms proposed, recently available evidence has suggested that PGs may also preserve the viability of not only gastric glands but also a homogeneous population of gastric cells exposed to noxious agents in vitro (8-10). Because PGs can directly interact with and protect gastric cells in the absence of mucus, blood flow, or bicarbonate, it is reasonable to speculate that PGs exert their protective effects presumably through activating intracellular processes. However, so far pretreatment of mucous-producing cells with dbcAMP has been shown not to reduce taurocholic acid-induced injury of the cells (9).

Accordingly, to clarify the mechanism by which PGs protect gastric cells against noxious agents-induced injury, we evaluated the effects of PGs on ethanol- or taurocholic acid-caused injury of isolated chief cells prepared from guinea pig stomach. We examined whether PGs stimulate not only cyclic AMP accumulation but also any other intracellular signals in gastric chief cells and clarified what cellular events activated by PGs are related to their protective effects on chief cell injury.

Methods

Materials. $^{45}\text{CaCl}_2$, ^{51}Cr , [^3H]arachidonic acid, and cAMP assay kit were obtained from Amersham Corp. (Arlington Heights, IL). [^3H]Glycerol was obtained from ICN K&K Laboratories Inc. (Plainview, NY). 16,16-Dimethyl prostaglandin E₂ (16,16dmPGE₂) was donated from Upjohn Co. (Kalamazoo, MI). Other PGs were donated from Ono Pharmaceutical Co. Ltd. (Japan). Collagenase (type 1), trypsin inhibitor (type 1s), 1-oleoyl-2-acetyl-*sn*-glycerol (OAG), forskolin, secretin, dibutyryl cAMP (dbcAMP), fura-2 acetoxymethyl (fura-2/AM), bovine hemoglobin, 12-*O*-tetradecanoylphorbol 13-acetate (TPA), Ca²⁺ ionophore A23187, 1-(5-isoquinolinylnsulfonyl)-2-methyl piperazine (H7), 4 α -phorbol 12,13-didecanoate (4 α PDD), porcine pepsinogen, isobutyl-1-methylxanthine (IBMX), and taurocholic acid (sodium salt) were purchased from Sigma Chemical Co. (St. Louis, MO). COOH-terminal octapeptide of cholecystokinin (CCK8) was donated from Squibb Institute Inc. (Princeton, NJ). Percoll was obtained from Pharmacia Fine Chemicals (Piscataway, NJ). Protein assay dye reagent concentrate was obtained from Bio-Rad Laboratories (Richmond, CA). All other materials used were of the highest grade available.

Preparation of isolated guinea pig gastric chief cells. The isolated guinea pig gastric chief cells were prepared as described previously (11, 12). The fundus and corpus of the stomach removed from a male guinea pig (Hartley strain, 250-300 g) were minced and digested for 30 min at 37°C in a incubation medium containing 0.1% collagenase. The gastric chief cell-enriched fraction was obtained after centrifugation of the unfractonated gastric cells dispersed in Percoll solution at 30,000 g for 15 min. For experiments, the chief cells were suspended and incu-

bated at 37°C in an oxygenated incubation medium constituting of 128 mM NaCl, 1.1 mM MgCl₂·6H₂O, 4.7 mM KCl, 1.28 mM CaCl₂·2H₂O, 1 mM Na₂ HPO₄, 0.5% BSA, 10 mM Hepes, 2% MEM amino acid (50 times concentrated), 0.1% glucose and 0.01% soybean trypsin inhibitor, at pH 7.4 (HR buffer).

LDH determination. Isolated chief cells were pretreated for 15 min at 37°C in HR buffer containing either PG, other agents or 0.025% ethanol as a solvent unless otherwise indicated. Chief cells were then incubated in HR buffer containing indicated volume of ethanol or taurocholic acid at 37°C for 1 h. The LDH released into the supernatant was determined by using a spectrophotometric LDH assay kit of Nihon Syoji using pyruvate as the substrate. LDH release during the incubation was expressed as a percentage of total LDH content present in the chief cells before the incubation.

Dye exclusion test. Gastric chief cells were pretreated with PGs and incubated with a noxious agent as described above. The washed chief cells were then suspended in 1 ml of PBS and thereafter 0.1 ml trypan blue solution (0.4 g/dl) was added directly to the cell suspension according to the method of Phillips (13). The number of stained or nonstained cells was counted within 10 min.

⁵¹Cr release assay. Isolated chief cells were incubated in HR buffer containing 10 μCi/ml of ⁵¹Cr for 2 h. The labeled cells were pretreated with PGs and incubated with ethanol for 1 h. Thereafter, ⁵¹Cr radioactivity of the cells and of the supernatant was counted with a gamma counter. The percentage of ⁵¹Cr released per sample was expressed by the method previously reported (14).

Pepsinogen measurement. Chief cells were incubated at 37°C for 30 min in the presence of test agents. Thereafter pepsinogen released into the medium was measured by the method of Anson and Mirsky using acid-denatured hemoglobin as previously described (15). Pepsinogen release was expressed as a percentage of total pepsinogen content present in the chief cells before the incubation. Pepsinogen content in chief cells was measured by the method above using porcine pepsinogen (3,800 peptic units/mg) as a standard.

Measurement of intracellular free Ca²⁺ concentration ([Ca²⁺]_i). [Ca²⁺]_i of isolated chief cells was measured by the method utilizing the Ca²⁺-selective fluorescence indicator fura-2 essentially as described previously (11).

Measurement of initial Ca²⁺ influx rate. Initial Ca²⁺ influx rate into chief cells was measured according to the methods of Mauger et al. as described previously (16). The cells were incubated in the presence of ⁴⁵CaCl₂ (2 μCi/ml) with test agents for 30, 90, 150, and 210 s at 37°C. The ⁴⁵Ca²⁺ associated with the cells was increased linearly within the range of these time points (data not shown). The initial Ca²⁺ influx rate was calculated by using a slope of the linear regression line of Ca²⁺ uptake.

Measurement of ⁴⁵Ca²⁺ efflux from chief cells. Chief cells were prelabeled with ⁴⁵Ca²⁺ (2 μCi/ml) by incubation in HR buffer at pH 7.4 for 1 h. Thereafter, the cells were incubated with test agents for indicated periods, centrifuged, and counted for the radioactivity (17).

Measurement of cyclic AMP accumulation. Chief cells suspended in HR buffer at pH 7.4 were incubated with PGs in the presence of 0.1 mM IBMX at 37°C. Intracellular cAMP was extracted by sonicating and boiling the chief cells in 0.02 N HCl containing 5 mM EDTA as described previously (18). Cyclic AMP extracted was measured by a radioimmunoassay procedure.

Assay of 1,2-sn-diacylglycerol (DG) formation. Gastric chief cells were incubated with either [³H]arachidonic acid (2 μCi/ml) in BSA-free HR buffer or [³H]glycerol (15 μCi/ml) in usual HR buffer at 37°C for 2 h. The cells were then washed, PGs or other agents were added into the cell suspensions, and the incubation was continued for the desired time interval. The reaction was stopped by the addition of 1.88 ml cold chloroform/methanol (2:1, vol/vol) followed by the addition of 0.62 ml chloroform and the same volume of distilled water. To separate [³H]DG by thin layer chromatography, dried lipids (dissolved in chloroform/methanol, 90:10) and authentic standards were spotted on silica gel plates activated at 110°C for 60 min and developed in the solvent system of hexane/diethylether/acetic acid (70:30:10) twice. Ra-

dioactive spots were identified by phosphomolybdic acid staining, and the radioactivity in each area was determined by liquid scintillation counting (19, 20).

Statistical analysis. All values are given mean±SE. The Student's *t* test was used for statistical evaluation. Comparisons involving more than two groups were performed by an analysis of variance. With these analysis, an associated probability (*P* value) of < 5% was considered statistically significant. All experiments were performed at least four times in triplicate unless otherwise indicated.

Results

We examined the effects of varying concentrations of PGE₂ on ethanol-caused injury of chief cells. The incubation of chief cells with 8% ethanol for 1 h caused LDH release to 20.3±0.3% of total content as compared with 3.6±0.2% of total content in the absence of ethanol. However, PGE₂ pretreatment lessened the LDH release in a dose-dependent manner with the maximal reduction to 15.1±0.6% at 3 × 10⁻⁶ M. A half-maximal dose (ED₅₀) of the inhibitory effect of PGE₂ on LDH release was 1.8 × 10⁻⁷ M. PGE₂ pretreatment for 15 min also dose-dependently reduced 5 mM taurocholic acid-caused LDH release from 21.9±0.7% to 15.9±0.6% with an ED₅₀ of 1.6 × 10⁻⁷ M. When the chief cell viability was estimated by the percentage of trypan blue-stained cells, PGE₂ similarly reduced their percentage from 56.5±0.2% to 36.0±0.3% with an ED₅₀ of 3.2 × 10⁻⁷ M. A similar dose response curve was also observed when the viability was estimated by ⁵¹Cr release (Fig. 1). Therefore, we evaluated chief cell viability by measuring the release of LDH in the following experiments.

The protective effect of PGE₂ against ethanol-caused injury was reduced when chief cells were pretreated at either acid or alkaline pH or at reduced temperature. PGE₂ maximally preserved chief cell viability when the cells were pretreated at 37°C and pH 7.4 (Table I). These results suggest that PGE₂ protects chief cells from ethanol injury only at physiological conditions.

As compared with PGE₂, other PGs such as 16,16dm-PGE₂, and PGE₁ also significantly reduced ethanol-caused LDH release to 14.8±0.7% and 15.8±0.8%, respectively. PGA₂

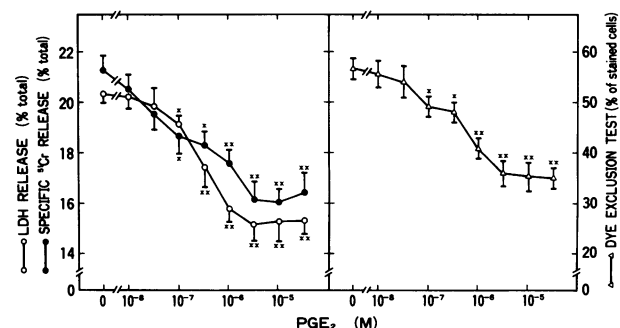


Figure 1. Effect of PGE₂ pretreatment on ethanol-caused LDH release, ⁵¹Cr release, and inability of chief cells to exclude trypan blue. Isolated chief cells were incubated with indicated concentrations of PGE₂ at 37°C for 15 min and thereafter incubated with 8% ethanol for 1 h. Chief cell injury caused by ethanol was estimated in terms of LDH release (○), ⁵¹Cr release (●), and dye exclusion test (Δ) as described in Methods. Chief cell samples were prepared in triplicate in each experiment and results shown are mean±SE of four separate experiments. Values indicated (**P* < 0.05, ***P* < 0.01) are significantly different from a respective control.

Table 1. Effects of PGE₂ Pretreatment at Different pH or Temperature on Ethanol-caused LDH Release

Temperature	4°C		16°C		37°C
PGE ₂ pretreatment					
(-)	22.9±0.9		22.2±0.7		20.5±0.6
(+)	22.4±1.0		19.0±0.8		15.1±0.7
Reduction of LDH release (%)		6.6±2.3		14.2±2.0	26.5±1.7
pH	6.0	6.8	7.4	7.8	8.2
PGE ₂ pretreatment					
(-)	22.0±0.9	20.9±0.8	20.2±0.6	21.3±0.7	22.7±0.7
(+)	19.2±1.0	15.9±0.8	14.9±0.5	17.8±0.6	20.1±0.7
Reduction of LDH release (%)	12.6±1.8	23.5±1.4	26.4±1.6	16.6±1.3	11.4±1.7

Chief cells were pretreated with 3×10^{-6} M PGE₂ either at the indicated temperature at pH 7.4 or at the indicated pH at 37°C for 15 min. Chief cells were subsequently incubated with 8% ethanol for 1 h at 37°C and pH 7.4. Thereafter LDH released into the medium was measured. Chief cell samples were prepared in triplicate in each experiment and results shown are mean±SE of four separate experiments.

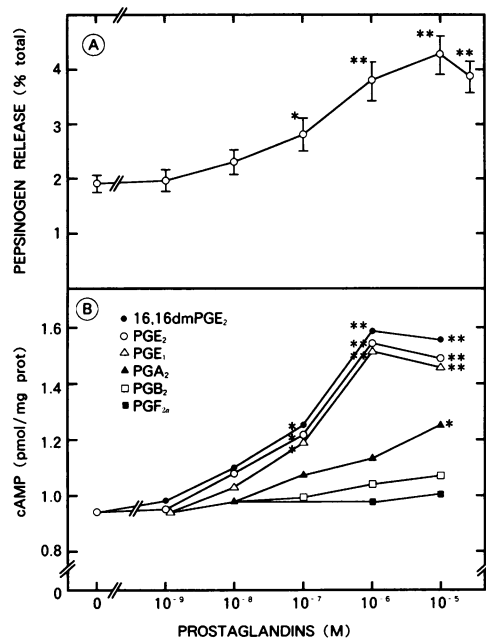


Figure 2. Effect of PGs on pepsinogen release and cAMP accumulation. (A) Chief cells were incubated with indicated concentrations of PGE₂ at 37°C for 30 min, and thereafter pepsinogen released into the medium was measured. Chief cell samples were prepared in triplicate in each experiment and results shown are mean±SE of four separate experiments. Chief cells in each sample ($5.4 \pm 1.7 \times 10^5$ cells) contain 19.7 ± 2.1 μg of pepsinogen before the incubation. (B) Chief cells were incubated with indicated concentrations of (●) 16,16dmPGE₂, (○) PGE₂, (Δ) PGE₁, (▲) PGA₂, (□) PGB₂, and (■) PGF_{2α} at 37°C for 30 min. Cyclic AMP extracted from chief cells was measured by radioimmunoassay. Chief cell samples were prepared in triplicate in each experiment and results shown are mean values of four separate experiments. Each SE less than 8% of mean values was omitted for clarification. Values indicated (**P* < 0.05, ***P* < 0.01) are significantly different from a control value.

reduced LDH release to $19.4 \pm 0.6\%$, but the reduction was not statistically significant. PGB₂ and PGF_{2α} had no effect on ethanol-caused LDH release.

PGE₂ not only reduced ethanol-caused LDH release but also stimulated pepsinogen secretion dose dependently. The maximal pepsinogen response observed at 10^{-5} M was $4.3 \pm 0.4\%$ total corresponding to 0.85 ± 0.08 μg pepsinogen with the calculated ED₅₀ of 2.4×10^{-7} M (Fig. 2A). The effects of PGE₂ at varying concentrations of up to 10^{-5} M on pepsinogen secretion and ethanol-caused chief cell injury showed a similar proportional concentration dependency. Therefore, we examined what cellular processes are involved in PG-induced biological actions in chief cells. 16,16dmPGE₂, PGE₂, and PGE₁ stimulated a dose-dependent increase in cAMP accumulation with the maximal response at 10^{-6} M and ED₅₀ of 1.0×10^{-7} , 1.1×10^{-7} and 1.2×10^{-7} M, respectively. PGA₂ had only a weak stimulatory effect and PGB₂ and PGF_{2α} had no effect on cAMP accumulation (Fig. 2B). We further examined whether pretreatment of chief cells with dbcAMP or forskolin reduces ethanol-caused LDH release from chief cells. However, ethanol-caused increase in LDH release was not reduced by the pretreatment with 10^{-5} M forskolin or 10^{-4} M dbcAMP at all (data not shown).

By contrast with the effects of PGs on cAMP, 3×10^{-6} M PGE₂ caused no significant change in $[Ca^{2+}]_i$ of chief cells (data not shown). Furthermore, 3×10^{-6} M PGE₂ did not affect a basal rate of initial Ca^{2+} influx of 0.7 ± 0.3 nmol/min per mg of chief cell protein. On the other hand, CCK8 or CCh stimulated an increase in initial Ca^{2+} influx rate to 4.8 ± 0.5 or 6.6 ± 0.4 nmol/min per mg protein.

We next examined the possibility that PGs stimulate an increase in DG accumulation, thereby exerting their protective effects on chief cells. Incubation of chief cells with 3×10^{-6} M PGE₂ for 15 s to 30 min resulted in a biphasic increase in DG accumulation in chief cells. DG response to PGE₂ reached to

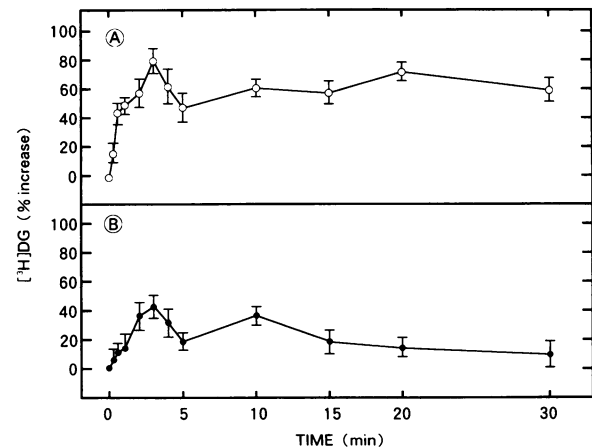


Figure 3. Effect of PGE₂ on [³H]DG formation. Chief cells pre-labeled with [³H]glycerol (○) or [³H]arachidonic acid (●) were incubated with 3×10^{-6} M PGE₂ at 37°C for indicated time periods. Thereafter the reaction was stopped as described in Methods. DG generated was collected in an organic phase, resolved, and separated by TLC. Values were expressed as percent increase over the basal value (mean values, $1,658 \pm 121$ dpm, [³H]arachidonic acid; $1,884 \pm 144$ dpm, [³H]glycerol). Samples were prepared in triplicate in each experiment and results shown are mean±SE of four separate experiments.

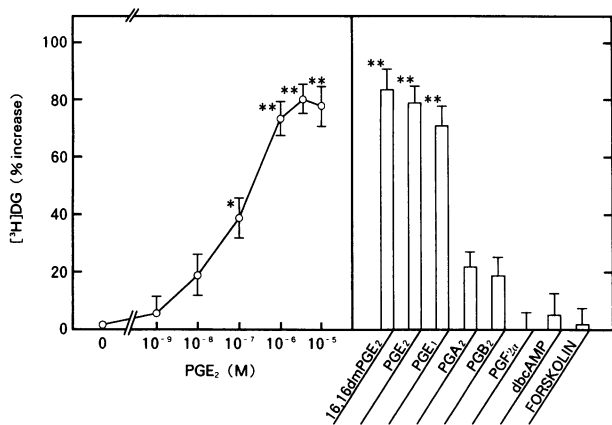


Figure 4. Effect of varying doses of PGs, dbcAMP, and forskolin on [³H] DG formation. Chief cells were pre-labeled with [³H]glycerol and thereafter incubated with indicated agents at 37°C for 3 min. (A) Effects of varying doses of PGE₂ on DG generation. (B) Effects of 3 × 10⁻⁶ M PGs, 10⁻⁴ M dbcAMP, and 10⁻⁵ M forskolin. Chief cell samples were prepared in triplicate in each experiment, and results shown are mean ± SE of four separate experiments. Values indicated (*P < 0.05, **P < 0.01) are significantly different from control.

the first peak value at 3 min with 80 ± 4% increase above a basal level in [³H]glycerol-labeled chief cells. PGE₂ further stimulated an increase in DG accumulation with the second peak at 20 min. DG accumulation was consistently sustained above a base level for at least 30 min. A biphasic increase in DG accumulation in a similar time course was also observed in response to 3 × 10⁻⁶ M PGE₂ in chief cells pre-labeled with [³H]arachidonic acid (Fig. 3).

This increase in DG accumulation at 3 min was dependent on either doses of PGE₂ or PGs used. The maximal increase in DG in response to PGE₂ was observed at 3 × 10⁻⁶ M with an ED₅₀ of 1.0 × 10⁻⁷ M (Fig. 4). 16,16dmPGE₂ or PGE₁ at 3 × 10⁻⁶ M almost equally stimulated an increase in DG accumulation, but PGA₂, PGB₂, or PGF₂α at 3 × 10⁻⁶ M did not significantly stimulate the increase in DG accumulation. Furthermore, 10⁻⁴ M dbcAMP or 10⁻⁵ M forskolin had no stimulatory effect on DG accumulation (Fig. 4).

Therefore, we investigated the role of DG/protein kinase C signaling pathway for the protection of chief cells from ethanol

injury by PGs. Pretreatment of chief cells for 15 min with varying doses of OAG, a synthetic diacylglycerol, which has been shown to activate protein kinase C directly (21), reduced ethanol-caused LDH release from chief cells in a dose-dependent manner (Fig. 5 A). TPA or PDBu, when added instead of OAG into pretreatment medium also reduced ethanol-caused LDH release dose dependently. By contrast, 4αPDD, an inactive phorbol ester (22), did not cause any reduction of ethanol-caused LDH release (Fig. 5 B). TPA or PDBu only at a high dose of 50 nM also failed to reduce ethanol-caused LDH release. When chief cells were pretreated with PGE₂ in the presence of H7, a protein kinase C inhibitor (23), H7 reversed the reduction of ethanol-caused LDH release by PGE₂ (Fig. 5 C).

When chief cells were pretreated with varying doses of OAG plus 10⁻⁴ M dbcAMP, dbcAMP did not affect the protective action of OAG or the chief cell injury at all (Fig. 6 A). On the other hand, simultaneous pretreatment of chief cells with OAG plus 10⁻⁷ M Ca²⁺ ionophore A23187 significantly reversed the reduction of ethanol-caused LDH release by OAG (Fig. 6 B). Even if chief cells were pretreated with varying doses of OAG plus 3 × 10⁻⁶ M PGE₂, additive effects on the reduction of LDH release were not observed (Fig. 6 C). Pretreatment of chief cells with 10⁻⁷ M CCK8, another pepsinogen releasing agonist which has been shown to stimulate polyphosphoinositides breakdown in chief cells (17), also maximally reduced LDH release from a control value of 20.3 ± 0.3% to 18.2 ± 0.5%, but the reduction was approximately one-third of that obtained with 10⁻⁴ M OAG or 3 × 10⁻⁶ M PGE₂.

Because Ca²⁺ ionophore blocked the protective effect of OAG on chief cell injury, we considered the possibility that OAG exerts the protective effect by increasing Ca²⁺ efflux from chief cells. 2.5 × 10⁻⁵ M OAG or 3 × 10⁻⁶ M PGE₂ significantly stimulated Ca²⁺ efflux from chief cells pre-labeled with ⁴⁵Ca²⁺ at 5–15 min of incubation. On the other hand, 10⁻⁴ M dbcAMP did not affect Ca²⁺ efflux from chief cells at all (Fig. 7).

Discussion

In the present study by using isolated gastric chief cells, we confirmed previous reports which suggest that PGs directly

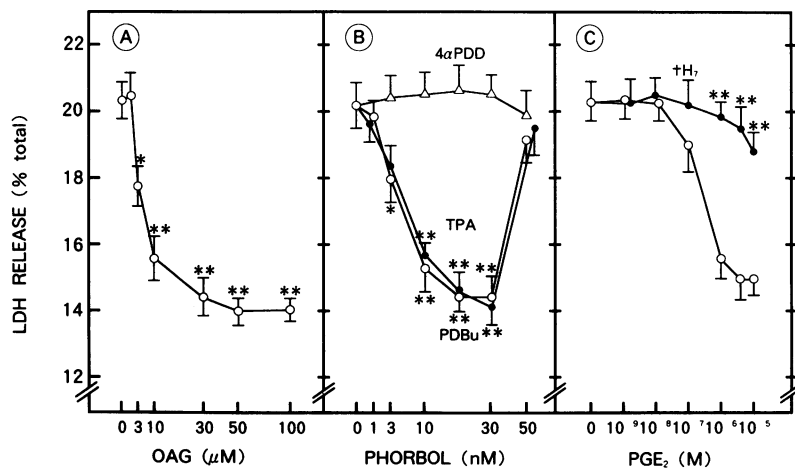


Figure 5. Effects of agents which may affect protein kinase C activity on ethanol-caused LDH release from chief cells. Isolated chief cells were pretreated with varying concentrations of (A) OAG (○), (B) phorbols (□] PDBu, [●] TPA, [Δ] 4αPDD) or (C) PGE₂ in the presence (●) or absence (○) of 5 × 10⁻⁵ M H7 at 37°C for 15 min and subsequently incubated with 8% ethanol. Thereafter, LDH released into the medium was measured. Chief cell samples were prepared in triplicate in each experiment and results shown are mean ± SE of four separate experiments. In A and B, values indicated (*P < 0.05, **P < 0.01) are significantly different from a control value obtained without OAG or phorbols. In C, values indicated (**P < 0.01) are significantly different from those without H7 at respective PGE₂ concentrations.

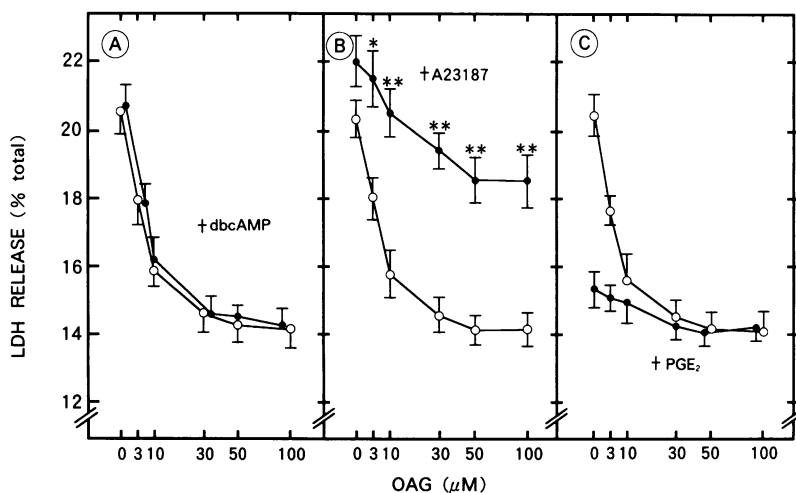


Figure 6. Combined effects of OAG with either dbcAMP, A23187, or PGE₂ on ethanol-caused LDH release. Chief cells were pretreated with varying concentrations of OAG (○) plus either (A) 10⁻⁴ M dbcAMP (●), (B) 10⁻⁷ M A23187 (●), or (C) 3 × 10⁻⁶ M PGE₂ (●) at 37°C for 15 min. The cells were subsequently incubated with 8% ethanol for 1 h. Chief cell samples were prepared in triplicate in each experiment and results shown are mean ± SE of four separate experiments. In B, values indicated (*P < 0.05, **P < 0.01) are significantly different from respective control values obtained without A23187.

protect gastric cells from injury caused by noxious agents. Furthermore, we extended the previous studies *in vitro* and provided new data on the possible mechanism by which PGs protect chief cells directly. In the present study, PGE₂ dose dependently reduced ethanol-caused increases in percentage of trypan blue-stained chief cells and release of cytoplasmic enzyme LDH and ⁵¹Cr from prelabeled chief cells. The methods we used to evaluate the degree of chief cell injury quantitatively limit the estimation of cellular viability and function. However, previous studies have also suggested by using a scanning electron microscopy that the measurement of leaked LDH is a sensitive and accurate index of ethanol-caused injury to the gastric chief cell plasma membranes and PGE₂ pretreatment preserves the structural integrity of not only plasma

membranes but also subcellular organelles in most chief cells exposed to ethanol (8, 24). Therefore, present results suggest that our methods utilized detected chief cell injury and PGE₂, at least in part, effectively protected ethanol- or taurocholic acid-caused injury of chief cells.

We showed in the present study that the maximal protective effect was observed when chief cells were pretreated with 3 × 10⁻⁶ M PGE₂ at 37°C and pH 7.4, but the protection was reduced when the pretreatment was carried out at 4°C. These results suggest that PGs may activate intracellular biochemical events thereby causing the protection of chief cells. Several lines of evidence suggest that PGs exert their protective effects on chief cells presumably through the activation of diacylglycerol/protein kinase C signaling pathway.

First, PGE₂ stimulated an increase in the accumulation of diacylglycerol in gastric chief cells prelabeled with [³H]arachidonic acid or [³H]glycerol in a dose-dependent manner. The ED₅₀ of PGE₂ to cause the protection of chief cells is very close to the ED₅₀ of PGE₂ to generate diacylglycerol. Furthermore, the rank order of potency of various PGs to generate diacylglycerol in chief cells is parallel with that of various PGs in reducing ethanol-caused injury. It is well known that diacylglycerol-activated protein kinase C plays a crucial role for a variety of biological actions in a number of cell types (25).

Second, not only OAG but also TPA, a phorbol ester which has been shown to activate protein kinase C apparently by substituting for diacylglycerol (21, 22), effectively reduced chief cell injury in a dose-dependent manner. On the other hand, 4αPDD, an inactive phorbol ester which does not bind to or activate protein kinase C (22), did not show any protective effect on ethanol-caused injury. Furthermore, H7 reduced the protective action of PGE₂, suggesting that protein kinase C may be involved in the action of PGE₂. Although at present we do not know yet why TPA at a very high dose failed to protect chief cell injury, TPA itself at high doses may be cytotoxic.

Third, although PGs stimulated an increase in cAMP accumulation in gastric chief cells, cAMP-dependent signaling pathway does not appear to be involved in PGs protection of chief cell injury. When chief cells were pretreated with forskolin at 10⁻⁵ M which has been shown to stimulate an increase in the accumulation of cAMP in chief cells (16), it failed to re-

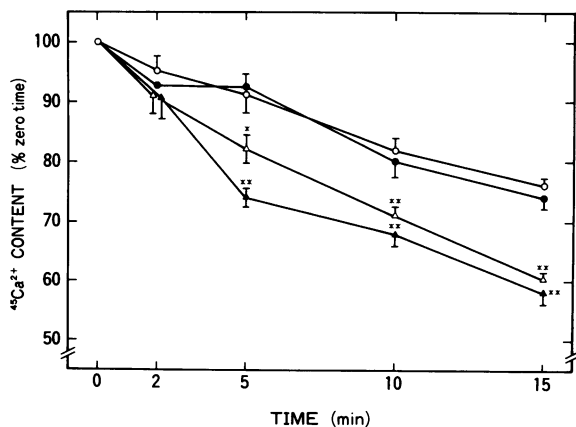


Figure 7. Effects of PGE₂, OAG or dbcAMP on ⁴⁵Ca²⁺ efflux from chief cells prelabeled with ⁴⁵Ca²⁺. Chief cells prelabeled with ⁴⁵Ca²⁺ were incubated with 3 × 10⁻⁶ M PGE₂ (Δ), 2.5 × 10⁻⁵ M OAG (▲), or 10⁻⁴ M dbcAMP (●) for the indicated periods. (○) Control values. Triplicate chief cell samples were then centrifuged and the radioactivity associated with the cells was counted. The ⁴⁵Ca²⁺ efflux was expressed as the percent radioactivity of the cells before the incubation. Results shown are mean ± SE of four separate experiments. Values indicated (*P < 0.05, **P < 0.01) are significantly different from control values at respective time points.

duce the extent of chief cell injury caused by ethanol. DbcAMP at 0.1 mM also did not show any protective effect on chief cell injury.

Fourth, PGs did not stimulate any increase in cytosolic free Ca^{2+} concentrations and initial Ca^{2+} influx rate in gastric chief cells. On the other hand, either PGE_2 or OAG stimulates Ca^{2+} efflux at 5–15 min of incubation. Moreover, Ca^{2+} ionophore, when added into preincubation medium in the presence of OAG, partially inhibited the protective effect of OAG on ethanol-caused chief cell injury. Therefore, an increase in $[\text{Ca}^{2+}]_i$ in chief cells may negatively affect the protective action of OAG against chief cell injury and OAG-induced chief cell protection may be due to its effect on Ca^{2+} efflux. However, further works are clearly required to clarify whether PGE_2 - or OAG-induced increase in Ca^{2+} efflux is really involved in the protection of chief cells exposed to noxious agents. Of interest is the observation that although CCK8 causes polyphosphoinositide breakdown thereby producing inositol phosphate and diacylglycerol in gastric chief cells (16), CCK8 at a maximally effective concentration for pepsinogen secretion does not protect chief cells from ethanol-caused injury to the extent which PGs do. An increase in $[\text{Ca}^{2+}]_i$ stimulated by CCK8 may also negatively affect the endogenously generated diacylglycerol-provoked protection.

In general, diacylglycerol has been shown to be derived from the polyphosphoinositide breakdown which is considered to be stimulated by phospholipase C activation by extracellular signals (26). However, in the present study, we were not able to observe any increase in $[\text{Ca}^{2+}]_i$ in chief cells. We have previously shown that Ca^{2+} mobilization can be observed in gastric chief cells when polyphosphoinositide breakdown is occurred by the stimulation of cholecystokinin (16). Therefore, PGs seem not to stimulate polyphosphoinositide breakdown to generate diacylglycerol in chief cells, although we can not completely exclude the possibility that diacylglycerol is generated in part from hydrolysis of inositol containing phospholipids (27). At present, we can not account for how PGs stimulate an increase in the accumulation of diacylglycerols in chief cells, but these data may be consistent with the observations that in some cells not all of the stimulated diacylglycerol can be derived from the hydrolysis of polyphosphoinositide (20, 28). The mechanism by which PGs stimulate an increase in DG accumulation in chief cells is currently under investigation.

One note of caution that we must consider in interpreting our findings is that the cell preparations used for the present study consisted of 95% chief cells in our best preparation. Accordingly, it is possible that the observed changes in diacylglycerol accumulation occurred in non-chief cells contaminating our preparations. However, the correlation between alterations in these parameters and PG-induced change in pepsinogen secretion suggests that the events are closely linked, probably within chief cells. Confirmation of this linkage requires additional studies in chief cells or other cell preparations of greater purity.

In conclusions, besides various effects of PGs in vivo, PGE_2 and PGE_1 seem to possess the direct protective action against ethanol- or taurocholic acid-caused injury in guinea pig chief cells, presumably through the activation of diacylglycerol/protein kinase C signaling pathway.

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