

1,25(OH)₂ Vitamin D₃ Stimulates Membrane Phosphoinositide Turnover, Activates Protein Kinase C, and Increases Cytosolic Calcium in Rat Colonic Epithelium

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Abstract

The hormonal form of vitamin D, 1,25(OH)₂ vitamin D₃ [1,25(OH)₂D₃], regulates colonic calcium absorption and colonocyte proliferation and differentiation. In this study, we have examined the effect of 1,25(OH)₂D₃ on membrane phosphoinositide turnover, protein kinase C activation, and regulation of intracellular calcium concentration ([Ca²⁺]_i) in isolated rat colonic epithelium. In a concentration-dependent manner, 1,25(OH)₂D₃ stimulated breakdown of membrane phosphoinositides within 15 s, generating diacylglycerol and inositol 1,4,5-triphosphate (IP₃). 1,25(OH)₂D₃ rapidly activated colonic protein kinase C, with maximal translocation of activity from the cytosol to the membrane occurring within 1 min of exposure to the secosteroid. Studies performed in isolated colonocytes with the fluorescent dye fura-2 demonstrated that 10⁻⁸ M 1,25(OH)₂D₃ caused a rapid rise in [Ca²⁺]_i which then transiently decreased before rising to a new plateau value. When these experiments were performed in a calcium-free buffer, an increase in [Ca²⁺]_i was observed, but both the transient and secondary rise were diminished in magnitude, suggesting that 1,25(OH)₂D₃ may stimulate both release of intracellular calcium stores and calcium influx. 1,25(OH)₂D₃ stimulated [³H]thymidine uptake in rat colonocytes, 4 h after an *in vivo* injection. These studies indicate that 1,25(OH)₂D₃ exerts a rapid influence on membrane phosphoinositide metabolism which may mediate certain of the secosteroid's effects on colonocyte calcium transport and proliferation. (*J. Clin. Invest.* 1990. 85:1296–1303.) 1,25(OH)₂ vitamin D₃ • phosphoinositide metabolism • intracellular calcium • protein kinase C • diacylglycerol

Introduction

The hormonally active form of vitamin D, 1,25(OH)₂ vitamin D₃ [1,25(OH)₂D₃],¹ has a well-recognized role in regulating

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1. Abbreviations used in this paper: IP, IP₂, and IP₃, inositol-1-monophosphate, inositol-1,4-bisphosphate, and inositol-1,4,5-triphosphate; KRBG, KRB containing 180 mg% glucose; PI, PIP, and PIPP, phosphatidylinositol, phosphatidylinositol-4-phosphate, and phosphatidylinositol-4,5-bisphosphate.

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calcium absorption and mineral metabolism (1). More recently, it has been established that 1,25(OH)₂D₃ has potent influences on cell proliferation and differentiation in many tissues (2, 3). In several malignant cell lines, including some derived from colon carcinomas, 1,25(OH)₂D₃ has been shown to stimulate differentiation while inhibiting proliferation (4, 5). The effect of 1,25(OH)₂D₃ on growth in normal tissues has been less extensively studied. Birge and Alpers (6) found that repletion of vitamin D-deficient rats rapidly stimulated intestinal mucosal cell proliferation, which was associated with a 20% increase in villus cell number by 32 h. The colonic epithelium is of interest since it manifests both the classic and non-classic actions of vitamin D. Colonic calcium absorption is vitamin D-responsive, and in some circumstances, such as the short bowel syndrome, may be important in calcium homeostasis (7, 8). In addition, the effect of 1,25(OH)₂D₃ on colonocyte growth and differentiation is of particular interest in view of epidemiologic evidence suggesting that sunlight exposure and dietary vitamin D and calcium intake may influence the prevalence of colonic carcinoma (9, 10).

Many of the effects of 1,25(OH)₂D₃ on cellular function can clearly be attributed to its action as a steroidlike hormone. High affinity 1,25(OH)₂D₃ receptors are present in many cell types, including the colon, and interaction of the 1,25(OH)₂D₃-receptor complex with chromatin alters gene expression (11). Attention has focused recently, however, on cellular responses to 1,25(OH)₂D₃ that are very rapid (12) and are not blocked by inhibitors of transcription and translation (13). Several workers have suggested a "liponomic" action of 1,25(OH)₂D₃, where the secosteroid exerts direct effects on cell membranes (13–15). Studies in various cell types have indicated that 1,25(OH)₂D₃ alters membrane phospholipid content (13, 14), fatty acid composition (13, 15), and fluidity (15).

The experiments described in this report demonstrate that 1,25(OH)₂D₃ rapidly stimulates colonic membrane phosphoinositide breakdown, generates inositol 1,4,5-triphosphate (IP₃) and diacylglycerol, and increases colonocyte intracellular calcium concentration ([Ca²⁺]_i). As a consequence of the rise in diacylglycerol content and [Ca²⁺]_i, 1,25(OH)₂D₃ also activates protein kinase C, a known regulator of proliferation and differentiation in many cell systems including the colonic epithelium (16, 17). Moreover, *in vivo* injection of 1,25(OH)₂D₃ stimulates [³H]thymidine incorporation into the DNA of colonocytes.

Methods

Materials. 1,25(OH)₂D₃ was kindly provided by Dr. M. R. Uskokovic (Hoffman-LaRoche Inc., Nutley, NJ), and 25(OH)D₃ was generously supplied by Dr. J. Babcock (Upjohn Co., Kalamazoo, MI). Leupeptin, phenylmethylsulfonyl fluoride, histone (type III-S), phosphatidylserine, phosphatidylinositol standards, DEAE-cellulose, ATP-Na salt, diethylenetriaminepenta-acetic acid, and morpholine propane sulfonic acid (MOPS) buffer salt were purchased from Sigma Chemical Co. (St. Louis, MO). [1-¹⁴C]*n*-butyric acid (15.0 mCi/mmol), [³H]thymidine (2

Ci/mmol), [γ - 32 P]ATP, [3 H]arachidonate, [3 H]myoinositol, [3 H]phosphatidylinositol compounds, and [3 H]inositol phosphate compounds were obtained from New England Nuclear Research Products (Boston, MA). Fura-2 and fura-2 AM (acetoxymethyl ester) were obtained from Molecular Probes (Eugene, OR). Sn-1,2-diacylglycerol kinase was purchased from Lipidex, Inc. (Westfield, NJ), octyl- β -D-glucoside from Boehringer Mannheim Biochemicals (Indianapolis, IN), and cardiolipin and dioleoyl glycerol from Avanti Polar Lipids, Inc. (Pelham, AL).

Isolation of intact colonic epithelium. For the studies of phosphatidylinositol metabolism and protein kinase C activation, an intact rat colonic epithelium preparation was used. The colonic epithelium was isolated as described by Craven et al. (18). Briefly, after anesthesia with pentobarbital (50 mg/kg i.p.), the colon was flushed with 20 ml of Ca/Mg-free Hanks' balanced salt solution at 37°C. A 16-gauge needle was placed in the left ventricle and perfused for 3–4 min at a flow rate of 20 ml/min with 30 mM EDTA in Ca/Mg-free Hanks' solution flushed with 95% oxygen and 5% CO₂ at 37°C. The colon was then excised, gently everted on a glass rod and slid over a 1-ml pipette attached to a rheostat-controlled motor. The epithelium was removed by 5-s bursts of rotation at 1,600 rpm and collected in cold Krebs Ringer bicarbonate buffer, pH 7.4, containing 180 mg% glucose (KRBG). The epithelium was centrifuged at 500 g, washed, and resuspended in the same buffer. Cell viability was assessed by the ability of the colonic crypts to metabolize [14 C]*n*-butyrate to 14 CO₂ (19). This reaction was linear for 2 h. In addition, trypan blue exclusion in > 90% of all cells was observed for at least 2 h.

Treatment of the colonic epithelium with 1,25(OH)₂D₃. The colonic epithelium was incubated in KRBG at 37°C for a total of 30 min. The time course of the effect of 1,25(OH)₂D₃ on protein kinase C was determined by adding the vitamin at different time intervals during the final 15 min of the incubation. To study the dose–response relationship, different concentrations of 1,25(OH)₂D₃ (10^{–12} to 10^{–8} M) were incubated with the colonic epithelium for the last minute of incubation. The control epithelial preparations were treated with an appropriate amount of ethanol vehicle, with the final ethanol concentration never exceeding 0.004%.

Partial purification of protein kinase C. After incubation with 1,25(OH)₂D₃ or vehicle, 20 ml of ice-cold KRBG buffer was added to each tube and the contents immediately centrifuged at 500 g for 5 min. The pellets were then homogenized 15 strokes with a motor-driven teflon pestle in 5 ml of homogenization buffer containing 20 mM Tris-HCl, pH 7.5, 0.5 mM EGTA, 2 mM EDTA, 2.0 mM phenylmethylsulfonyl fluoride, 0.5 mM benzimidazole, 5.0 mM 2-mercaptoethanol, and 10 mg/liter leupeptin (20). The homogenate was subjected to ultracentrifugation at 100,000 g for 60 min and the supernatant S₁ (cytosolic fraction) saved. The pellet was gently resuspended in 5.0 ml of homogenization buffer containing 0.3% Triton X-100 (wt/wt) and left on ice for 60 min before recentrifugation at 100,000 g for 60 minutes. The supernatant S₂ (solubilized membrane fraction) was collected and both S₁ and S₂ fractions were applied to DEAE-cellulose columns (Poly-prep columns, 0.8 × 4 cm, Bio-Rad Laboratories, Richmond, CA) that had been preequilibrated in the homogenizing buffer. In preliminary studies, the columns were washed with 16 ml of buffer and eluted with 32 ml of a linear gradient of 0–0.1 M NaCl in homogenization buffer. As most of the protein kinase C activity eluted in 0.035–0.055 M NaCl, for routine purposes the loaded columns were first washed with 8 ml of homogenizing buffer and then with 4 ml of the same buffer containing 0.02 M NaCl. Finally, protein kinase C was eluted in 2 ml of buffer containing 0.08 M NaCl. The eluted cytosolic or membrane fractions were assayed for protein kinase C activity on the same day.

Assay of protein kinase C. Protein kinase C activity was determined using a histone phosphorylation assay, as described previously (20). The DEAE-purified fractions were incubated in a reaction mixture (final volume 75 μ l) containing (final concentrations) 20 mM Tris-HCl (pH 7.2), 10 mM MgCl₂, 400 μ g/ml histone (type III-S), 1.83 mM CaCl₂ (the actual concentration of unchelated CaCl₂ used was only 1.0 mM, since the contributions of EGTA and EDTA from the column

elution buffer were 0.17 and 0.66 mM, respectively), and 50 μ M [γ - 32 P]ATP (1 μ Ci) with and without 80 μ g/ml of phosphatidylserine. Aliquots of phosphatidylserine (10 mg/ml ethanol) were evaporated under N₂ and sonicated in 20 mM Tris buffer, pH 7.4, before addition to the reaction mixture. Reactions were started by adding 25 μ l of the protein kinase C preparation to 50 μ l of the assay mixture, and the incubations were carried out for 3 min at 30°C. 50 μ l of the assay mixture was blotted onto 2.5 × 2.5-cm phosphocellulose papers (No. P81, Whatman Inc., Clifton, NJ) that had been prewashed in 10% trichloroacetic acid, 2 mM NaH₂PO₄ solution. The papers were then washed by agitating in 250 ml of ice-cold 10% TCA for 12 min and left under running water for 5 min. The filter papers were soaked in 95% ethanol for 5 min, followed by diethyl ether for an additional 5 min, and then air-dried before taking Cherenkov counts. Protein kinase C activity was calculated from the difference in phosphorylation assayed in the presence and absence of phosphatidylserine. Enzyme activity was linear with respect to enzyme concentration in all assays and activity was expressed as picomoles of 32 P per minute per milligram of protein.

Determination of labeled diacylglycerol. The effect of 1,25(OH)₂D₃ on the production of diacylglycerol was determined using colonic epithelial preparations in which membrane phosphoinositides had been prelabeled with [3 H]arachidonate as previously described (16). Colonic epithelium (5 mg of protein/5 ml) was preincubated with 20 μ Ci of [3 H]arachidonate for 1 h at 37°C. In preliminary experiments, we found that the phosphoinositides were the major phospholipid class labeled with [3 H]arachidonate (67% [3 H]phosphoinositides, 16% [3 H]-phosphatidylcholine, 11% [3 H]phosphatidylethanolamine, and 6% [3 H]phosphatidylserine). The epithelium was washed in KRBG containing 10 mg/ml fatty acid free albumin, resuspended in KRBG and incubated at 37°C for 15 min. During this incubation, 1,25(OH)₂D₃ was added for different time intervals as described above. The reaction was terminated by the rapid addition of ice-cold KRBG, and the crypts were centrifuged at 500 g at 4°C for 10 min. The pellet was extracted in chloroform/methanol 2:1 (vol/vol), and the lipid extract was then loaded on to a silicic acid column. Neutral lipids were eluted with chloroform (21), and immediately dried under N₂ to prevent isomerization of 1,2-diglyceride to 1,3-diglyceride. The neutral lipid samples were spotted onto silica gel G coated thin-layer chromatography plates and developed with hexane/ethyl ether/acetic acid 80:20:2 (vol/vol/vol). The lipids were identified with iodine vapor and the spots corresponding to authentic diacylglycerol standard were scraped from the plate, transferred to scintillation vials, and counted.

Measurement of 1,2-diacylglycerol mass. Cell pellets were combined with 10 ml of 2:1 (vol/vol) chloroform/methanol and total cellular lipids extracted as described by Folch et al. (22). An aliquot of this extract was assayed in triplicate for total lipid phosphorus as described by Bartlett (23). Same-day aliquots of lipid extract were also assayed for 1,2-diacylglycerol mass by the diacylglycerol kinase procedure of Preiss et al. (24) with the following modifications. MOPS buffer, pH 6.6, was substituted for imidazole buffer, pH 6.6 (25). For each reaction tube, diacylglycerol kinase (20 mU in 10 μ l) was combined with 50 μ l of reaction buffer (100 mM Na-morpholine propane sulfonic acid, pH 6.6, 100 mM NaCl, 25 mM MgCl₂, 2 mM EGTA) plus 10 μ l of 20 mM dithiothreitol and 10 μ l of 10 mM [γ - 32 P]ATP (5.0 × 10⁴ cpm/nmol), and the mixture was incubated at 25°C for 30 min. as described by Wright et al. (26). Duplicate aliquots of dioleoylglycerol standards or cellular lipid extracts were transferred to 75 × 12-mm capped polyethylene tubes (Sarstedt, Inc., Princeton, NJ) and dried under argon. A 20- μ l aliquot of resuspension buffer (7.5% octyl- β -D-glucoside, 5 mM cardiolipin, 1 mM diethylenetriamine penta-acetic acid) was added, vortexed, and sonicated as described (24). To this, 80 μ l of the enzyme-ATP mixture was added, followed by vortex mixing and incubation at 25°C for 30 min. The reaction was terminated by the addition of 1.67 ml of CHCl₃/MeOH/12 N HCl (66:31:1, vol/vol/vol), followed by 1.67 ml of MeOH/H₂O/CHCl₃ (48:47:3, vol/vol/vol) as described by MacDonald et al. (25), followed by vortex mixing and centrifugation. The upper phase was removed and discarded, and the

lower phase was reextracted with 1.67 ml of MeOH/H₂O/CHCl₃ (48:47:3, vol/vol/vol). After centrifugation and removal of the upper phase, the labeled phosphatidic acid in the lower phase was assayed directly by scintillation counting of an aliquot as justified by Muldoon et al. (27). Standard curves were generated from the assay of known amounts of 1,2-diacylglycerol and data expressed as nanomoles of 1,2-diacylglycerol/100 nmol of lipid phosphorous (mol%) (26).

Determination of labeled inositol phosphates and phosphoinositides. The effect of 1,25(OH)₂D₃ on the colonic content of inositol phosphates and membrane phosphoinositides was determined using colonic epithelial preparations prelabeled with [³H]myoinositol. Colonic epithelium (10 mg protein) was incubated for 2 h in 2 ml of KRBG containing 25 μ Ci of [³H]myoinositol (sp act 12.8 Ci/mmol). In preliminary experiments, we found that 0.21% of [³H]myoinositol was incorporated into the phosphoinositol fraction of total cellular lipid. The epithelium was centrifuged at 500 g for 10 min at 4°C and washed three times with cold KRBG. The crypts were then resuspended in 2 ml of 20 mM HEPES/Tris buffer, pH 7.0, containing 25 mM β -glycerophosphate, 2.0 mM EGTA/Ca⁺, 0.1 M KCl, 5 mM β -mercaptoethanol, 5 mM MgCl₂, 0.02% trypsin inhibitor, and 10 mM LiCl. Use of this buffer was associated with the most reliable results and is similar to other buffers used for the assay of inositol phosphates and phosphoinositides. The suspensions were incubated for 15 min at 37°C and 1,25(OH)₂D₃ was added as above. The reactions were terminated by addition of 0.67 ml 10% perchloric acid and allowed to stand on ice for 15 min. After centrifugation at 500 g, the supernatant was neutralized with 20 mM HEPES/Tris buffer, pH 8.0, before determination of inositol phosphates. The pellet was mixed with 0.5 ml chloroform/methanol/12 N HCl (200:100:0.75, vol/vol/vol) and the extracted phosphoinositides were saved for later analyses. Separation of different species of inositol phosphates was achieved by anion-exchange chromatography on 0.3 ml AG-I \times 8 (HCOO⁻) columns (200–400 mesh) based on the method of Downes et al. (28). After loading, the columns were eluted serially with 1.5 ml each of (a) water, (b) 0.1 M formic acid, 0.2 M ammonium formate, (c) 0.1 M formic acid, 0.4 M ammonium formate, and (d) 0.1 M formic acid, 1 M ammonium formate. Inositol-1-monophosphate (IP), inositol-1,4-bisphosphate (IP₂) and IP₃ eluted in the second, third, and fourth fractions, respectively. The samples were mixed with scintillation fluid and counted. For the separation and quantification of phosphoinositides, 0.2 ml of the acidified pellet extract was treated with 1.5 ml of chloroform/methanol (2:1 vol/vol), vortexed, and centrifuged. The upper phase was discarded and the lower phase was washed two times with 0.75 ml methanol/0.6 N HCl (1:1, vol/vol) and then separated on 1% K-oxalate impregnated silica gel G chromatography plates. Unlabeled phosphoinositides were added as carriers. The plates were developed in chloroform/acetone/methanol/glacial acetic acid/H₂O (40:15:13:12:7 vol/vol/vol/vol/vol) (29). The labeled phosphoinositides including phosphatidylinositol (PI), phosphatidylinositol-4-phosphate (PIP), and phosphatidylinositol-4,5-bisphosphate (PIPP) were visualized by spraying the plates with En³hance (New England Nuclear) and exposing to X-ray film (SM-5) (Eastman Kodak Co., Rochester, NY), using a Kodak X-OMAT enhancing screen, for 3–5 d. Individual phosphoinositide spots were identified by comparison with authentic standards, and quantified by scraping the silica gel from the plate and counting the radioactivity in a liquid scintillation counter.

Isolation of colonocytes. Colonic epithelial cells were isolated as described previously (30). The colon was removed, tied at one end, and rinsed in 0.9% NaCl containing 1.0 mM dithiothreitol. The colon was then filled with 0.1 M phosphate-buffered saline, pH 7.2, containing 1.5 mM EDTA and 0.5 mM dithiothreitol and closed with a Dieffenbach Serrefine clamp (Millers Forge, Plano, TX). The colonic sacs were incubated in 150 ml of phosphate-buffered saline at 37°C for 30 min. Sacs were emptied and cells were collected by centrifugation at 500 g, rinsed in phosphate-buffered saline, and resuspended in the appropriate buffer for measurement of intracellular calcium concentration. Cells were routinely viable for at least 1 h as assessed by trypan blue exclusion.

Measurement of intracellular calcium concentration ([Ca²⁺]_i). Isolated colonocytes were incubated with 1 μ M fura-2 AM for 30 min at 37°C. The cells were collected by centrifugation at 500 g for 5 min at 4°C, and resuspended in buffer containing 145 mM NaCl, 5 mM KCl, 10 mM HEPES, 5 mM MgCl₂, 10 mM D-glucose, pH 7.2 (31). Studies were performed using buffers containing 1 mM CaCl₂, or calcium-free with and without 1 mM EGTA. The kinetics of fluorescence changes in the fura-2-loaded colonocytes were analyzed using an SLM-4800 C spectrofluorometer (SLM Instrument, Inc., Urbana, IL). The cuvette containing the colonocytes was constantly stirred and maintained at 37°C. 1,25(OH)₂D₃ or vehicle was injected directly into the cuvette. Fluorescence was measured at excitation wavelengths of 340 and 380 nm, with emission at 505 nm. Signals from the fluorometer were fed into an IBM-PC (IBM Instruments, Inc., Danbury, CT) with SLM spectrum processor version 3.2 for subsequent analysis. [Ca²⁺]_i was calculated based on the formula of Grynkiewicz et al. (32) assuming the K_d of the Ca²⁺-fura-2 interaction to be 225 nM in the cytosolic environment. The calibration of [Ca²⁺]_i was based on spectra of 1 μ M fura-2 (penta potassium salt) in Ca²⁺-EGTA buffers with free Ca²⁺ values ranging from < 1 to > 10 μ M.

[³H]Thymidine incorporation in rat colonocytes. Rats weighing 250–300 g were injected subcutaneously with either 1,25(OH)₂D₃ (10 ng/100 g body weight) or ethanol vehicle. After 2 h the rats were injected intraperitoneally with 100 μ Ci of [³H]thymidine. 4 h after 1,25(OH)₂D₃ injection the rats were killed and colonocytes prepared as described above. [³H]Thymidine incorporation was measured according to Verbin and Farber (33). DNA was extracted from the precipitate with 10% TCA at 90°C for 10 min and measured as previously described (34).

Statistical analysis. Data were analyzed using Student's *t* test for unpaired data (35).

Results

Effect of 1,25(OH)₂D₃ on colonic phosphoinositides and inositol phosphates. To examine the effect of 1,25(OH)₂D₃ on phosphatidylinositol metabolism, studies were performed using colonic epithelium that was prelabeled with [³H]myoinositol. As shown in Table I, administration of 10⁻⁸ M 1,25(OH)₂D₃ caused a decrease in the membrane content of labeled phosphoinositides within 15 s, with the most prominent response seen in the labeled PIPP content. Concomitantly, there was an increase in the colonic levels of inositol phosphates, with the greatest change seen in the IP₃ content which increased more than twofold at 30 s and 1 min. As indicated in Table II, the effect of 1,25(OH)₂D₃ on phosphoinositide bisphosphate breakdown and IP₃ formation was dose-dependent, with 10⁻⁸ M 1,25(OH)₂D₃ producing a greater effect than 10⁻¹⁰ M.

Effect of 1,25(OH)₂D₃ on colonic diacylglycerol. Colonic epithelium was prelabeled with [³H]arachidonate as shown in Fig. 1. Treatment with 1,25(OH)₂D₃ rapidly increased the colonic [³H]diacylglycerol content, with a maximum response seen at 15–30 s. At later time points, the [³H]diacylglycerol content decreased towards the baseline value. The effect of 1,25(OH)₂D₃ on diacylglycerol levels was concentration-dependent (Fig. 2), as significant increases were seen with both 10⁻¹⁰ and 10⁻⁸ M, but not 10⁻¹² M.

Diacylglycerol mass was measured 90 s after administration of either 10⁻⁸ M 1,25(OH)₂D₃ or ethanol vehicle. Vehicle treated cells had a diacylglycerol content of 0.35 \pm 0.002 mol% compared with 0.57 \pm 0.005 mol% (mean \pm SEM, *n* = 8) in the 1,25(OH)₂D₃-treated cells (*P* < 0.001).

Effect of 1,25(OH)₂D₃ on colonic protein kinase C activity. As shown in Fig. 3, administration of 10⁻⁸ M 1,25(OH)₂D₃

Table I. Time Course of 1,25(OH)₂D₃ Effects on Cellular Phosphoinositides and Inositol Phosphates

	Time					
	0	15 s	30 s	1 min	5 min	15 min
Phosphoinositide						
PI	75.7±0.8	71.4±2.5	67.5±2.7*	70.4±3.0	80.0±3.0	73.6±0.8
PIP	8.1±0.5	7.7±0.3	7.2±0.4	7.5±0.2	7.0±0.5	7.7±0.1
PIPP	7.1±1.0	4.9±0.4	4.2±0.4‡	4.5±0.3*	4.3±0.4*	5.6±0.2
Inositol phosphates						
IP	2.8±2.0	3.2±0.1	4.9±0.4‡	4.2±0.1	3.9±0.1	3.0±0.1
IP ₂	2.5±0.2	3.2±0.1‡	3.0±0.2‡	4.5±0.2‡	2.3±0.1	4.5±1.0
IP ₃	1.8±0.1	2.3±0.3	4.2±0.1‡	4.1±0.1‡	2.4±0.2	3.0±0.2

Colonic epithelial preparations were prelabeled with [³H]myoinositol for 2 h at 37°C. At 2 h, 0.21% of administered myoinositol was incorporated into cellular lipids (4.1 pmol myoinositol). After washing, the epithelial preparations were resuspended in buffer and exposed to 10⁻⁸ M 1,25(OH)₂D₃ for 15 s–15 min. The reactions were terminated by adding 0.67 ml of 10% HClO₄, and after centrifugation the pellets were extracted with 0.5 ml of CHCl₃/CH₃OH/12 N HCl 200:100:0.75 (vol/vol/vol). Individual phosphoinositides including PI, PIP, and PIPP were separated on 1% K-oxalate impregnated silica gel G plates. The spots were visualized by spraying the plates with En³hance and exposing to Kodak X-ray film (SM-5) for 3–5 d. Individual phosphoinositide spots were scraped and counted. The supernatant was neutralized and the inositol phosphates, IP, IP₂, and IP₃ were separated by anion exchange chromatography and assayed by scintillation counting. Data are presented as percentage of total disintegrations (phosphoinositides plus inositol phosphates, ~ 100,000 dpm/experiment) associated with each spot, and given as mean±SEM for six determinations from three separate experiments. *P < 0.05; †P < 0.001; ‡P < 0.02 vs. time 0.

resulted in a rapid activation of colonic protein kinase C. At baseline, 75% of colonic protein kinase C activity was in the cytosol, whereas only 25% was membrane-associated. Within 15 s after 1,25(OH)₂D₃ treatment, there was significant translocation of protein kinase C to the membrane fraction, and peak activation occurred with 1 min of exposure. Subsequently, there was a decrease in membrane-associated protein kinase C to the baseline level in spite of continued exposure to 1,25(OH)₂D₃. Total colonic protein kinase C activity (cytosol + membrane) did not change with 1,25(OH)₂D₃ treatment. Direct addition of 1,25(OH)₂D₃ to either membrane or cytosolic fractions had no effect on protein kinase C activity. Fig. 4 illustrates the concentration dependence of protein kinase C activation by 1,25(OH)₂D₃. All studies were performed with 1 min of 1,25(OH)₂D₃ treatment, and demonstrate a dose-re-

sponse relationship between 10⁻¹² and 10⁻⁸ M 1,25(OH)₂D₃. Exposure of colonic epithelium to the ethanol vehicle or to 25(OH)D₃ at a concentration of up to 10⁻⁶ M did not cause protein kinase C translocation to the membrane fraction. Similar 1,25(OH)₂D₃ induced activation of protein kinase C was seen using the isolated colonocyte preparation (data not shown).

Effect of 1,25(OH)₂D₃ on colonocyte [Ca²⁺]_i. Fig. 5 illustrates the effect of 10⁻⁸ M 1,25(OH)₂D₃ on colonocyte [Ca²⁺]_i, determined using the fluorescent dye fura-2. In buffer containing 1 mM CaCl₂, the baseline [Ca²⁺]_i was 138±10 nM (Fig. 5). The addition of 10⁻⁸ M 1,25(OH)₂D₃ caused a rapid in-

Table II. Concentration Dependence of 1,25(OH)₂D₃ Effects on Cellular Phosphoinositides and Inositol Phosphates

	Vehicle	10 ⁻¹⁰ M	10 ⁻⁸ M
Phosphoinositide			
PI	76.2±0.8	70.9±1.9*	69.0±1.7‡
PIP	8.5±0.4	8.7±0.3	7.3±0.3
PIPP	6.1±0.4	5.5±0.3*	3.8±0.4‡
Inositol phosphate			
IP	2.9±0.2	4.0±0.2*	4.1±0.2*
IP ₂	2.8±0.1	3.8±0.1‡	4.2±0.2*
IP ₃	1.8±0.3	3.7±0.3‡	5.0±0.3‡

Colonic epithelial preparations were prelabeled with [³H]myoinositol as described in Table I, and treated with vehicle or 1,25(OH)₂D₃ (10⁻⁸ and 10⁻¹⁰ M) for 30 s. Membrane phosphoinositides were separated and quantitated as described in Table I. Data are presented as percent of total dpm associated with each spot and given as mean±SEM for six determinations in three separate experiments. *P < 0.05; †P < 0.01 or less compared with vehicle. ‡P < 0.02 compared with 10⁻¹⁰ M.

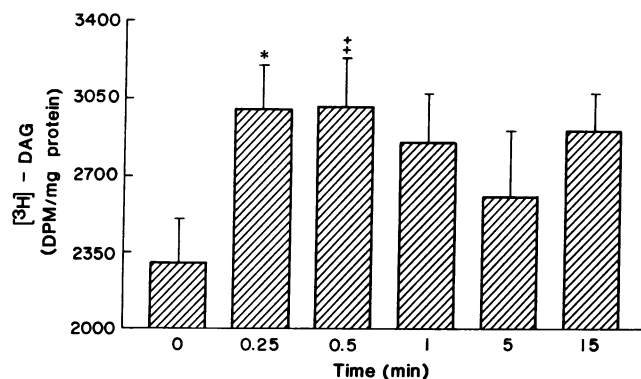


Figure 1. Time course of 1,25(OH)₂D₃-stimulated production of diacylglycerol (DAG). Colonic epithelial preparations were prelabeled with [³H]arachidonate for 1 h at 37°C. After washing, the epithelial preparations were resuspended in KRBG and exposed to 10⁻⁸ M 1,25(OH)₂D₃ for 15 s–15 min. After centrifugation, the pellets were extracted in chloroform/methanol (2:1, vol/vol), and the neutral lipid fraction separated using a silicic acid column. Diacylglycerol was separated by thin-layer chromatography, identified with iodine vapor, scraped from the plate, and counted. Data are presented as disintegrations per minute of DAG per milligram of protein and are given as mean±SEM for six determinations in four separate experiments. *P < 0.005; †P < 0.001 vs. baseline value.

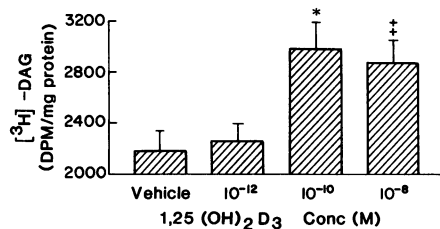


Figure 2. Concentration dependence of 1,25(OH)₂D₃-stimulated production of diacylglycerol (DAG). Colonic epithelial preparations were prelabeled with [³H]arachidonate as described in Fig. 3, and treated with vehicle or 1,25(OH)₂D₃ (10⁻¹² to 10⁻⁸ M) for 30 s as reported in Table II. The amount of [³H]DAG in the membrane pellet was determined as described in Fig. 3. Data are presented as mean±SEM for four to six determinations in three separate experiments. **P* < 0.005; †*P* < 0.001 vs. vehicle.

crease in [Ca²⁺]_i to 315±22 nM after 20–30 s of treatment. This was followed by a decrease in [Ca²⁺]_i, and a subsequent second rise to a new plateau level. Using a lower concentration of 1,25(OH)₂D₃, 10⁻¹⁰ M, the initial transient and plateau values of [Ca²⁺]_i were ~ 25% of that seen with 10⁻⁸ M. When colonocytes were studied in a calcium-free buffer containing 1 mM EGTA, the baseline [Ca²⁺]_i decreased to 65±10 nM. Addition of 10⁻⁸ M 1,25(OH)₂D₃ resulted in a transient increase in [Ca²⁺]_i to 152±5 nM, but no secondary rise was observed and the plateau level was identical to the baseline value. When colonocytes were studied in a calcium-free without EGTA buffer, the baseline [Ca²⁺]_i and response to 1,25(OH)₂D₃ were intermediate between that seen with the 1 mM calcium buffer

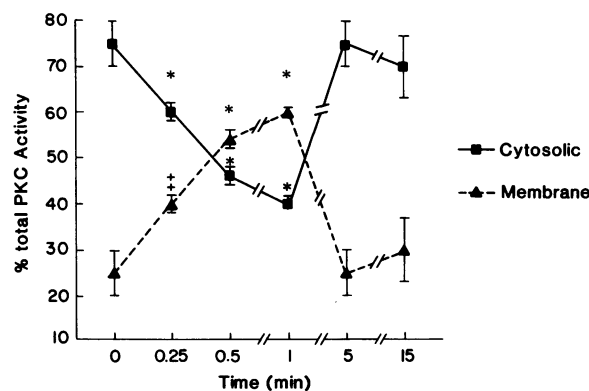


Figure 3. Time course of activation of protein kinase C (PKC) by 1,25(OH)₂D₃. Colonic epithelial preparations were incubated in KRBG at 37°C and treated with 10⁻⁸ M 1,25(OH)₂D₃ for 15 s–15 min. After addition of 20 ml of ice cold KRBG to each tube, the contents were centrifuged, and the pellets then homogenized in 5 ml of buffer. After ultracentrifugation at 100,000 *g* for 60 min, the supernatant S₁ (cytosolic fraction) was saved, and the pellet resuspended in homogenization buffer containing 0.3% Triton X 100 (wt/wt) and left on ice for 60 min. After centrifugation at 100,000 *g* for 60 min, the supernatant S₂ (solubilized membrane fraction) was collected. Protein kinase C in the S₁ and S₂ fractions was partially purified using DEAE-cellulose columns. Protein kinase C activities in the S₁ and S₂ fractions were determined from the differences in phosphorylation of histone (type III-S) measured with and without addition of 80 μg/ml of phosphatidylserine. Data are expressed as percentage of total PKC activity in each fraction. Values given are mean±SEM for 21 determinations in seven separate experiments. **P* < 0.001; †*P* < 0.005 vs. zero time value.

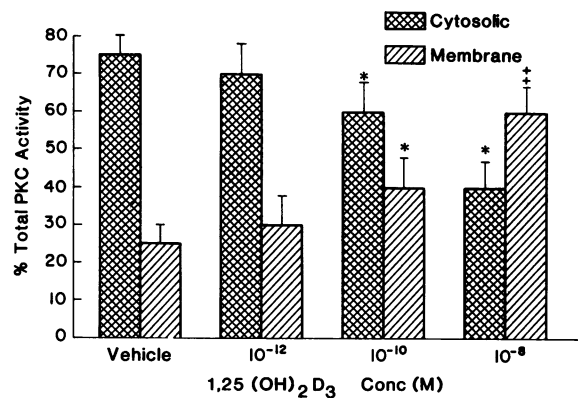


Figure 4. Concentration dependence of 1,25(OH)₂D₃-stimulated activation of protein kinase C (PKC). Colonic epithelial preparations were treated with 1,25(OH)₂D₃ (10⁻¹² to 10⁻⁸ M) or vehicle for 1 min. Cytosolic and solubilized membrane fractions were prepared; protein kinase C was partially purified and assayed as described in Fig. 3. Data are expressed as percentage of total protein kinase C activity in each fraction, and are given as mean±SEM for nine determinations in three separate experiments. **P* < 0.05; †*P* < .001 vs. vehicle.

and the 1 mM EGTA buffer. Addition of ethanol vehicle or 10⁻⁶ M 25(OH)₂D₃ had no effect on colonocyte [Ca²⁺]_i regardless of the buffer calcium concentration used.

Effect of 1,25(OH)₂D₃ on [³H]thymidine incorporation in rat colonocytes. [³H]Thymidine incorporation was measured four hours after a single dose of 1,25(OH)₂D₃. Vitamin D-treated animals incorporated 44% more [³H]thymidine/mg-DNA than vehicle-treated animals (33.5±4.3 vs. 48.4±5.8 pmol of [³H]thymidine/mg of DNA, mean±SEM, *n* = 4, *P* < 0.05).

Discussion

The experiments described in this paper are the first to demonstrate an effect of 1,25(OH)₂D₃ on phosphatidylinositol me-

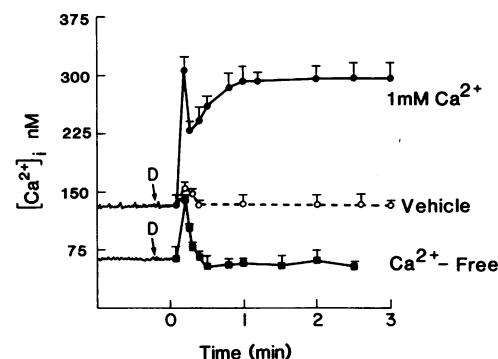


Figure 5. Effect of 10⁻⁸ M 1,25(OH)₂D₃ on [Ca²⁺]_i. Isolated colonocytes were incubated with 1 μM fura-2AM for 30 min at 37°C. After washing, the cells were suspended in buffer containing either 1 mM CaCl₂ (●, ○) or 1 mM K₂H₂ EGTA (■). The kinetics of fluorescence changes were measured at excitation wavelengths of 340 and 380 nm, with emission at 510 nm. [Ca²⁺]_i was calculated based on the formula of Grynkiewicz et al. (32). At the times indicated by the arrows, 10⁻⁸ M 1,25(OH)₂D₃ (●, ■) or ethanol vehicle (○) was injected directly into the cuvette. Data presented are the mean±SD of two to five determinations in three separate experiments.

tabolism, with subsequent activation of protein kinase C and regulation of $[Ca^{+2}]_i$ in intestinal tissue. The responses to $1,25(OH)_2D_3$ occur rapidly, within seconds to minutes, suggesting that they are not the result of the classic steroidlike or genomic action of $1,25(OH)_2D_3$ (36). This mode of action is much slower, inasmuch as $1,25(OH)_2D_3$ interacts with its receptor forming a complex which binds to chromatin and alters gene transcription and eventually protein synthesis (36, 37).

Little information is available on the effect of $1,25(OH)_2D_3$ on membrane PI turnover in various target cells. In a preliminary report, MacLaughlin et al. (38) showed that exposure of keratinocytes to $1,25(OH)_2D_3$ resulted in a two- to fivefold elevation in the level of IP_3 within 30 s and an increase in metabolism of PI to PIPP, the metabolite that is the source of IP_3 . In contrast, Chisholm et al. (39) found no effect of $1,25(OH)_2D_3$ on the level of IP_3 in GH_4C_1 pituitary cells. Baran and Kelly (40) have reported that $1,25(OH)_2D_3$ stimulates phospholipase A_2 -induced deacylation of phosphatidyl-inositol in hepatocyte membranes, generating lysophosphatidyl-inositol which can increase hepatocyte cytosolic calcium levels. In our studies, $1,25(OH)_2D_3$ caused a rapid decrease in the colonic levels of membrane phosphoinositides, and an increase in the cellular content of inositol phosphates, most prominently IP_3 . The increase in colonocyte IP_3 began at 15 s and was significant by 30 s. The rise in IP_3 coincided with the initial rise in $[Ca^{+2}]_i$ suggesting that it may be responsible for the calcium response observed in these cells. This is consistent with the known stimulatory effect of IP_3 on the release of calcium from an endoplasmic reticulum storage pool (41). Activation of the phosphoinositol pathway has also been shown to increase transmembrane calcium cycling by acting upon membrane calcium channels, although the mechanisms for this effect remain uncertain (42).

Our studies have also demonstrated that $1,25(OH)_2D_3$ stimulates diacylglycerol formation. The origin of this diacylglycerol is not entirely clear, however, two observations would tend to support phosphoinositol as the major precursor for diacylglycerol in this study. First, the time course of diacylglycerol formation coincides with the time course of phosphoinositide breakdown. Secondly, phosphoinositides are the major phospholipids labeled when colonic epithelium is exposed to $[^3H]$ arachidonate. This does not, however, exclude the possibility that some of the diacylglycerol formed may have come from other sources such as phosphatidylcholine.

Activation of protein kinase C, as indicated by translocation of the enzyme from the cytosol to the membrane fraction (43), temporally coincides with diacylglycerol formation and the increase in $[Ca]_i$. As has been described in other systems (43), the translocation of protein kinase C to the membrane appears to be transient in spite of continued exposure to $1,25(OH)_2D_3$. This may reflect modification or degradation of the protein kinase C to a form that has altered requirements for diacylglycerol, phosphatidylserine, or calcium, and which may have additional intracellular actions (44). Since protein kinase C appears to play an important role in cell proliferation and differentiation (45, 46), $1,25(OH)_2D_3$ -induced activation of protein kinase C is a potential mechanism by which the secosteroid affects colonocyte growth and maturation. Bile salts and dietary fatty acids, agents which alter colonocyte proliferation and have been implicated in the promotion of colonic carcinoma, have been demonstrated to also activate colonocyte protein kinase C (16–18). The finding that $1,25(OH)_2D_3$ stimulates colonocyte $[^3H]$ thymidine uptake is

consistent with the proliferative effect of protein kinase C as well as the known proliferative effect of vitamin D on intestinal mucosa (6).

Previous studies of the effect of $1,25(OH)_2D_3$ on protein kinase C activity in other cell types have found conflicting results. Ways et al. (47) reported that $1,25(OH)_2D_3$ enhanced phorbol ester-stimulated differentiation and protein kinase C-dependent phosphorylation of cellular proteins in the U937 human monoblastoid cell. $1,25(OH)_2D_3$ increased the protein kinase C level in both the cytosolic and membrane fractions. Martell et al. (48) reported that $1,25(OH)_2D_3$ -induced differentiation of HL-60 cells was inhibited by exposure of the cells to a protein kinase C inhibitor H-7. They demonstrated that protein kinase C activity was increased twofold after 24 h of exposure to $1,25(OH)_2D_3$, and that this response could be blocked by the protein synthesis inhibitor cycloheximide. In contrast, Mezzetti et al. (49) reported that $1,25(OH)_2D_3$ induced differentiation of U937 cells without protein kinase C activation, whereas phorbol ester-induced differentiation of these cells was mediated through protein kinase C. Sasaki et al. (50) observed that $1,25(OH)_2D_3$ decreased the cell growth and enhanced chemical transformation of BALB 3T3 cells without activation of protein kinase C. It appears, therefore, that although $1,25(OH)_2D_3$ stimulates protein kinase C in colonocytes and other cell types, the effects of this secosteroid on cell growth and differentiation are complex and involve other mechanisms in addition to protein kinase C activation.

We have demonstrated that $1,25(OH)_2D_3$ results in a rapid rise in colonocyte $[Ca^{+2}]_i$. The specificity of this response is supported by the observation that exposure of colonocytes to a 100-fold higher concentration of $25(OH)$ vitamin D_3 did not alter $[Ca^{+2}]_i$. When the colonocytes were studied in a calcium-free buffer with or without 1 mM EGTA, $1,25(OH)_2D_3$ caused an increase in $[Ca^{+2}]_i$ but less than that seen in calcium-containing buffers. These results indicate that $1,25(OH)_2D_3$ is either stimulating calcium influx or is causing release of calcium from a cellular pool that is readily exchangeable with extracellular calcium. Since an increase in $[Ca^{+2}]_i$ was seen in calcium-free EGTA containing buffer, $1,25(OH)_2D_3$ must release calcium from intracellular storage sites. The relationship of the phosphoinositol signal transduction pathway to changes in $[Ca^{+2}]_i$ has been extensively studied. Most studies indicate that IP_3 stimulates release of calcium from endoplasmic reticulum (51), although others suggest that IP_3 has effects at the plasma membrane (41). Further work to define the relationship between colonocyte phosphoinositol metabolism and changes in $[Ca^{+2}]_i$ is needed.

A number of previous studies in diverse cell types have also documented an effect of $1,25(OH)_2D_3$ on $[Ca^{+2}]_i$ (38, 39, 52–54). Baran and Milne (52) demonstrated a dose-dependent effect of $1,25(OH)_2D_3$ on hepatocyte $[Ca^{+2}]_i$ that occurred both in the presence and absence of extracellular calcium, indicating that $1,25(OH)_2D_3$ mobilized intracellular calcium pools. Sugimoto et al. (53) demonstrated that as little as 10^{-10} M $1,25(OH)_2D_3$ increased $[Ca^{+2}]_i$ in bovine parathyroid cells, whereas $25(OH)D_3$ and $24,25(OH)_2D_3$ were ineffective. The calcium channel inhibitors verapamil and diltiazem did not block the response to $1,25(OH)_2D_3$. Leiberherr (54) showed that $1,25(OH)_2D_3$, in a concentration as low as 10^{-11} M, transiently increased $[Ca^{+2}]_i$ in mouse osteoblasts within the 1st min of exposure (54). Studies using calcium-free buffers and inhibitors of calcium release from various cellular compartments suggested that the major effect of $1,25(OH)_2D_3$ was to

enhance calcium influx, although $1,25(\text{OH})_2\text{D}_3$ also appeared to release calcium from an endoplasmic reticulum pool and to accelerate calcium efflux. In a preliminary report, MacLaughlin et al. (38) found that 10^{-8} M $1,25(\text{OH})_2\text{D}_3$ increased $[\text{Ca}^{+2}]_i$ twofold in keratinocytes, whereas no response was seen with $25(\text{OH})\text{D}_3$ or vitamin D_3 (38). In their studies, $1,25(\text{OH})_2\text{D}_3$ provoked an increase in $[\text{Ca}^{+2}]_i$ even in a calcium-free buffer. Thus, $1,25(\text{OH})_2\text{D}_3$ appears to alter the $[\text{Ca}^{+2}]_i$ level in different cell types by several mechanisms, including alterations in membrane calcium influx and efflux, release of various intracellular calcium stores, and by potentiating responsiveness to other agents that operate through cytosolic calcium. In colonocytes and most other cells, the response to $1,25(\text{OH})_2\text{D}_3$ is very rapid, consistent with a direct action on cell membranes.

The mechanism by which $1,25(\text{OH})_2\text{D}_3$ stimulates membrane phosphoinositide turnover remain to be elucidated. Previous studies have not identified a plasma membrane vitamin D receptor, thus further research is needed to assess the membrane association of $1,25(\text{OH})_2\text{D}_3$, the role of GTP-binding proteins, and activation of phospholipase C in response to $1,25(\text{OH})_2\text{D}_3$.

The possible relationship among the rapid increase in $[\text{Ca}^{+2}]_i$, protein kinase C activation, and the regulation of transcellular calcium transport also deserves further study. Previous studies have demonstrated enhancement of calcium uptake into intestinal and skeletal muscle cells within 2–5 min of $1,25(\text{OH})_2\text{D}_3$ exposure (55–57), a time period consistent with the rapid effects observed in the present paper. It is possible that $1,25(\text{OH})_2\text{D}_3$ -induced PI breakdown initiates a series of membrane events that increase and ultimately sustain calcium transport, i.e., a “liponomic” mode of action (13). In hepatocytes, $1,25(\text{OH})_2\text{D}_3$ increases hepatocyte phospholipase A_2 activity after 2.5 min of exposure (40), and in renal and intestinal cells, membrane phospholipid composition is altered within 30 min (13, 14). Rasmussen et al. (13) demonstrated that the early effects of $1,25(\text{OH})_2\text{D}_3$ on intestinal brush border membrane lipid composition and calcium uptake were not blocked by inhibitors of transcription and translation, indicating a direct membrane effect. Putkey et al. (58) examined the effect of vitamin D deficiency and essential fatty acid deficiency on enterocyte membrane fluidity lipid composition and calcium flux. Of interest was their finding that the ileum from essential fatty acid and vitamin D deficient chicks failed to respond to vitamin D with an increase in calcium flux. Analysis of lipid composition revealed that these deficient chicks had increased amounts of saturated fatty acids and decreased amounts of linoleic acid. Brasitus et al. (15) showed that $1,25(\text{OH})_2\text{D}_3$ treatment increased the dynamic component of fluidity and corrected the fatty acid composition of small intestinal brush border membranes from vitamin D-deprived rats. These changes occurred within 1–2 h of treatment, temporally preceding changes in duodenal calcium absorption.

Further work must be directed toward identifying specific cellular proteins that are phosphorylated after $1,25(\text{OH})_2\text{D}_3$ -induced protein kinase C activation, and to the elucidation of the functional significance of these target proteins in calcium transport and cell growth and differentiation.

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