Angiotensin II does not stimulate proliferation of rat thyroid PC Cl3 cell line

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Abstract

In PC Cl3 cells, a rat thyroid cell line, angiotensin (Ang II) activates the atypical protein kinase C- ζ (PKC- ζ) and the extracellular signal-regulated kinase (ERK) pathways. We here studied the Ang II effects on PC Cl3 cell proliferation. It was found that Ang II: (1) induced the phosphorylation of protein kinase B (PKB), (2) induced the growth-related early gene c-fos expression, (3) enhanced the cyclin E and p27^{kip} expression, (4) had no effects on Cdk2, and (5) did not affect the transition from G0/G1 to S phase. Inhibition of phosphoinositide-3kinase by LY294002 further increased the effect of Ang II on p27^{kip} induction, whilst PKCs inhibition by GF109203X decreased such effect. The role of PKC- ζ was recognized by the use of a synthetic myristoylated peptide with sequences based on the endogenous PKC- ζ

pseudosubstrate and by PKC- ζ downregulation using the small interfering RNA (siRNA). Insulin had a replicating effect on PC Cl3 cells, induced the phosphorylation of PKB, decreased p27^{kip} expression and had no effect on the PKC- ζ cytosol-to-membrane translocation. PC Cl3 cell proliferation was induced more potently by simultaneous stimulation with insulin and Ang II than by stimulation with insulin alone, and the effect on p27^{kip} expression was similar to that obtained with insulin only. These observations demonstrate that in PC Cl3 cells Ang II causes a block in G1 phase, although both ERK and PKB pathways are activated, and this effect may be due to the upregulation of p27^{kip} and PKC- ζ operativity.

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Introduction

Angiotensin II (Ang II) was originally discovered as a potent vasoconstrictor; however, research in the past several decades has provided ample evidence that Ang II modulates cell growth (Gill & Simonian 1977, Geisterfer *et al.* 1988, Nafilan *et al.* 1989, Bunkenburg *et al.* 1992). Depending on the specific cell type and presence of other growth factors, Ang II induces proliferation, hypertrophy, apoptosis or differentiation.

The Ang II receptor AT1, is expressed by rat thyroid (Montiel & Jimenez 1998) and a relationship between thyroid function and several components of the renin-angiotensin system has been established (Montiel *et al.* 1984, Ruiz *et al.* 1987, Catanzaro 1995). Recently, our laboratory has shown that PC Cl3 cells, a rat thyroid cell line, express a functional AT1 receptor that regulates the Na⁺/K⁺ATPase activity and activates the extracellular signal-regulated kinases 1 and 2 (ERK1/2) pathway through the activity of the atypical protein kinase C- ζ (Marsigliante *et al.* 2003). A direct role of ERK1/2 has been demonstrated in Ang II-induced DNA synthesis and proliferation of cultured rat aortic smooth muscle cells (Marrero *et al.* 1997).

Since both the activity of the Na⁺/K⁺ATPase and the activation of the ERK pathway are greatly involved in the

proliferation of various cells (Ferrandi *et al.* 2004, Muscella *et al.* 2004), the purpose of this study was to investigate first whether Ang II affects PC Cl3 cell proliferation. To this end, we analyzed the effects of Ang II on the transition from G0/G1 to S phase, on cell number and metabolic activity and, finally, on total cell protein content.

Clearly, Ang II induced proliferation that requires complete cell progression through the various steps of the cell cycle. In contrast, cells undergoing Ang II-mediated hypertrophy, apoptosis, and differentiation are arrested in the G1 phase (Wolf & Wenzel 2003).

The proliferation of mammalian cells is controlled by an intricate network of biochemical pathways that ensure that each cell-cycle event is performed correctly and in the proper sequence. Growth factor-induced signals are required for progression through the G1 phase and must converge, in late G1, on the cell-cycle engine to ensure the commitment of cells to enter S phase (Pardee 1989). The regulation of G1 progression and G1/S transition is governed, at least in part, by the concerted action of cyclin-dependent kinases (Cdks) and their regulatory cyclin subunits (Draetta 1994, Sherr 1994, Grana & Reddy 1995). Biochemical and genetic data indicate that cyclin E-Cdk2 activity is essential for entry into S phase (Ohtsubo *et al.* 1995, Krude *et al.* 1997). The activity of Cdks is regulated by a combination of

mechanisms. These include the synthesis of cyclin and Cdk, the assembly of these proteins into complexes and the interaction with Cdk inhibitory proteins (Morgan 1995). The Cdk2 inhibitor p27^{kip} has been shown to play an important role in cell-cycle regulation. High levels of p27^{kip}, present in quiescent cells, have been shown to decline upon mitogen induction (Sherr & Roberts 1995). This decrease in p27^{kip} appears to be critical in enabling the cells to enter the cell cycle.

Another important step in response to proliferative signal is the transient transcriptional activation of immediate early genes associated with cell division, such as the proto-oncogenes *c-fos*, *c-jun*, and *egr-1* (Herschman 1991). The protein products of these genes are involved in transcriptional control and lead to a cascade of gene activation mediating the growth response. Indeed, it has been shown that Ang II also increases the expression of several growth related genes *in vitro*, including *c-fos* (Taubman *et al.* 1989, Nafilan *et al.* 1990, Rosenberg & Hostetter 1993). Ang II has been shown to activate phosphatidylinositol 3-kinases (PI3K) in various cell types (Saward & Zahradka 1997, Dugourd *et al.* 2003). PI3K are lipid kinases that phosphorylate the 3'-OH group of the inositol ring in inositol phospholipids after activation by a variety of extracellular stimuli. Several downstream targets of PI3K have been identified, such as the serine/threonine kinase Akt, also known as protein kinase B (PKB). Once phosphorylated, Akt activates various proteins involved in many cellular responses, including cell survival and growth promotion (Datta *et al.* 1999, De Vita *et al.* 2000). Recently, it has been shown that Akt is phosphorylated and activated by the AT1 receptor in a PI3K-dependent manner in various cell types (Eguchi *et al.* 1999, Takahashi *et al.* 1999, Yang & Raizada 1999, Garcia-Caballero *et al.* 2001, Rolz *et al.* 2002).

The major task of the present study was to investigate the mechanisms underlying the proliferative response of PC Cl3 cells to Ang II. To this end we examined the effects of Ang II on p27^{kip} and other cell-cycle proteins, on *c-fos* expression and Akt/PKB activity in PC Cl3 cells.

Materials and Methods

Materials

Fetal bovine serum (FBS) and glutamine were from Euroclone (Paignton, UK). Hydrocortisone, transferrin, L-glycyl-histidyl-lysine, and somatostatin were from MP Biomedicals (Irvine, CA, USA). Coon's modified Ham's F12 medium, bovine serum albumin (BSA), and other reagents were from Sigma-Aldrich Co. The mitogen-activated extracellular kinase (MEK) inhibitor PD98059, the protein kinase C (PKC) inhibitor GF109203X, and the PI3K inhibitor LY294002 were obtained from Sigma Chemical Co. Myristoylated PKC- ζ pseudosubstrate peptide (Myr-SIYRRGARRWRKL, ζ -PS) was obtained from Calbiochem-Novabiochem (Schwalbach, Germany).

Cell culture

Several different batches of the PC Cl3 cells were used in the experiments and the cells were grown for 40 passages approximately. Each experiment was performed on four different cell preparations, with at least three replicates in a single experiment. PC Cl3 rat thyroid cells were grown in Coon's modified Ham's F12 medium, supplemented with 5% FBS, containing L-glutamine (2 mM), gentamicin (50 mg/ml) and six-hormone mixture consisting of insulin (1 μ g/ml), hydrocortisone (3.62 μ g/ml), transferrin (5 μ g/ml), L-glycyl-histidyl-lysine (20 ng/ml), somatostatin (10 ng/ml), and thyroid-stimulating hormone (TSH; 1 mU/ml). The cells were maintained in a water-saturated atmosphere of 5% CO₂ and 95% air at 37 °C. The cells were passaged every 7–8 days, and the culture medium was changed every 2–3 days.

Cell proliferation assay

The conversion of MTT (3-(4,5-dimethylthiazol-2-yl)-2,5-diphenol tetrazolium bromide) by PC Cl3 cells was used as an indicator of cell number as described by Mosmann (1983), with some modification as described previously (Muscella *et al.* 2002). This method measures the reduction of MTT by active mitochondria, which results in a colorimetric change measured at optical density 570. PC Cl3 cells were grown in 96-well plates for 1 day in Coon's modified Ham's F12 medium, supplemented with 5% FBS, containing a mixture of six hormones. The media were replaced with hormone-free media, supplemented with 0.5% FBS ('starved cells'), for 48 h to induce quiescence. Ang II (100 nM) or insulin (100 nM) or Ang II and insulin in combination (used in a single dose of 100 nM) were added to each well, except for one strip, used as control. After 24, 48, and 72 h, MTT (250 μ g/ml) was added to the cells for a 4-h incubation and cells were lysed in acidified isopropanol. The plates were subsequently read on a spectrophotometer at 570 nm (Bio-Rad). The results were expressed as percent viability of each well in respect of control.

Cell count

PC Cl3 cells were seeded at 5×10^4 cells/well on 24-well plates, and cells were counted in a Thomas cell-chamber 24, 48 and 72 h after treatment as described above.

Cell-cycle analysis

Cell cycle was evaluated by staining cellular DNA with the DNA intercalator propidium iodide (PI) using a semiautomatic procedure (DNA-Prep Reagents Kit; Beckman Coulter Inc., Fullerton, CA, USA), followed by analysis on an EPICS XL flow cytometer.

In brief, 1×10^6 cells, starved for 48 h in fresh medium without hormones and with 0.5% FBS, and then stimulated with agonists for 18 h. Then cells were harvested by trypsinization, pelleted, washed twice with free calcium

and magnesium PBS, fixed, permeabilized and stained by addition of 100 µl detergent reagent lysing and permeabilizing reagent (LPR), followed by 1 ml PI solution. After mixing, samples were incubated for 30 min at room temperature and analyzed by flow cytometry. Cell-cycle data analysis was performed on 20 000 events.

Western blot analysis

To obtain whole protein cell extracts for western blot analysis of Akt/PKB, c-fos, cyclin E, Cdk2, and p27^{kip} PC Cl3 cells were scraped in the following buffer A (mM): 20 Tris-HCl, pH 8, containing 100 NaCl, 2 EDTA, 2 N₃VO₄, 2% Nonidet P-40, and also containing a mixture of protease inhibitors, including 0.5 phenylmethylsulfonyl fluoride, 10 µg/ml leupeptin, and 10 µg/ml pepstatin. Cells were then passed several times through a 20-gauge syringe and centrifuged at 16 000 g for 20 min at 4 °C. Proteins were determined with the Bio-Rad protein assay kit 1, using lyophilized BSA as standard.

For the separation of PKC-ζ between the cytosol and membrane, after washing with cold PBS, PC Cl3 cells were scraped into the following buffer (mM): 20 Tris-HCl, pH 7.5, containing 250 sucrose, 2 EGTA, 2 EDTA, 10 MgCl₂, 2 dithiothreitol, and protease inhibitors, and after two freeze and thaw cycles, cells were centrifuged at 900 g for 15 min at 4 °C and the resultant pellet was resuspended and centrifuged once more. The pooled supernatants were centrifuged at 100 000 g for 60 min to obtain cytosolic and membrane fractions. The pellet at 100 000 g was solubilized in buffer B (mM): 20 Tris-HCl, pH 7.5, 150 NaCl, 1 EGTA, 1 EDTA, and protease inhibitors, containing 2% Nonidet P-40.

Equal amounts of proteins (50 µg) from homogenates were loaded and separated on 10% SDS gels by electrophoresis and transferred to a nitrocellulose membrane. The sheet was blocked with 3% non-fat dry milk in buffered saline before incubation for 1 h at 25 °C with 2 µg/ml of anti body (Ab) to Akt/PKB (Sc-7985-R), c-fos (Sc-52), cyclin E (Sc-481), Cdk2 (Sc-6248), p27^{kip} (Sc-1641) and PKC-ζ isoform (Sc-876) (all from Santa Cruz Biotechnology, Santa Cruz, CA, USA) in blocking solution. The proteins were detected with goat anti-rabbit or anti-mouse IgG conjugated with peroxidase (Santa Cruz Biotechnology), using enhanced chemiluminescence (ECL) (Amersham Life Sciences Inc.). The intensity of the bands was quantified by scanning densitometry using the NIH Image 1.62 software (National Institutes of Health, Bethesda, MD, USA).

Total protein content

PC Cl3 cells were plated in six-well plates (1 × 10⁴ cells/well), starved for 48 h, and stimulated with a daily dose of Ang II. Total protein was determined by the method of Lowry *et al.* (1951) after 24, 48 and 72 h.

Design and preparation of small interfering RNA (siRNA)

siRNAs were prepared by an *in vitro* transcription method. Initially, four siRNA-ζ target sites specific to rat PKC-ζ mRNA, as determined by blast analysis, were chosen. For each siRNA, sense and antisense templates were designed based on each target sequence and partial T7 promoter sequence. All template oligonucleotides were chemically synthesized and PAGE purified. *In vitro* transcription, annealing, and purification of siRNA duplexes were performed using the protocol supplied with the T7 RiboMAX Express RNAi System (Promega). Briefly, 2 µg approximately of each single-strand (ss) transcription template was first annealed with the T7 promoter and filled in by Klenow DNA polymerase to form double-strand transcription templates. For preparation of each siRNA duplex, transcription reactions were first performed with separated antisense and sense templates using the T7 RNA polymerase provided with the kit, and then annealed to form siRNA duplexes. Then, the siRNA duplex was treated with DNase and RNase to remove the extranucleotides of transcribed siRNA to meet the structural 3VUU overhang and 5' phosphate requirement.

PKC-ζ immunoblotting was performed 24 h posttransfection to determine the efficiency of siRNA incorporation in PC Cl3 and to measure PKC-ζ expression. Quantitative analysis of PKC-ζ expression, as measured by intensity of PKC-ζ immunoreactivity, in siRNA-ζ- or siRNA-NS-transfected PC Cl3 revealed a higher reduction in PKC-ζ expression with the following sense RNA sequence: 5'-GCUGAGAUCUGUAUCGCUC-3'.

siRNA transfection

PC Cl3 cells (50–70% confluence) were transfected with siRNA duplexes using the protocol supplied with the CodeBreaker siRNA transfection reagent (Promega). Briefly, transfection reagent was first diluted into Coon's modified Ham's F12 medium without serum and antibiotics for about 15 min, and then the siRNA duplex was added to the medium to form a lipid-siRNA complex. Following an additional 15-min incubation, transfection was initiated by adding the lipid-siRNA complex to six-well plates. The final concentration of siRNA was 30 nM.

Statistical analysis

Experimental points represent the mean ± s.d. of 4–6 replicates. Statistical analysis was carried out using Student's *t*-test for unpaired samples and ANOVA. When indicated, *post hoc* tests (Bonferroni and Dunn) were also performed.

A *P* value of <0.05 was considered to be statistically significant.

Figure 1 Proliferative effects of Ang II and insulin in PC Cl3 cells. The histograms show cell cycle analysis of PC Cl3 cells grown in complete culture medium (containing FBS and 6 hormones growth factor mixture) (A), starved for 48 h (G₀/G₁ synchronization) (B), starved for 48 h and then treated for 18 h with Ang II 100 nM (C) and insulin 100 nM as positive control (D). The table shows the mean $\pm$ s.d. of the percentage distribution of the cells in the different phases of the cell cycle, obtained in other independent experiments ($n=4$) performed on different cell preparations. Statistical analysis was carried out using Student's *t*-test for unpaired samples: $P<0.01$ between starved and insulin-treated PC Cl3 cells; $P>0.05$ between starved and Ang II-treated PC Cl3 cells; $P<0.01$ between Ang II and insulin-treated PC Cl3 cells (E). The effect of insulin and Ang II on cell proliferation is additive. Starved PC Cl3 cells were treated without or with Ang II or Insulin or incubated with insulin and Ang II in combination agonists for the indicated time. Proliferation was analyzed by a spectrophotometric MTT assay as described in Materials and Methods. The data are mean $\pm$ s.d. of four different experiments run in eight replicates and are presented as percent proliferation of control. ANOVA: $P<0.0001$. Values with shared letters are not significantly different according to Bonferroni/Dunn *post hoc* tests.

Results

Effect of Ang II on PC Cl3 cell proliferation

Proliferation was analyzed by a spectrophotometric 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenol-2H-tetrazolium bromide (MTT) assay and confirmed by direct cell counts. The basal metabolic activity was unchanged in the presence of Ang II and Ang II had no replicating effect on starved PC Cl3 cells (Fig. 1E). The same results were obtained by direct cell counting (data not shown). As a positive control for the proliferation studies insulin was used, which is known to be a powerful mitogen for PC Cl3 cells and to stimulate the phosphorylation of ERK1/2 (Florio *et al.* 2001). Indeed, insulin produced mitogenic effects in PC Cl3 cells (Fig. 1D and E). PC Cl3 cell proliferation was induced more potently by simultaneous stimulation with insulin (100 nM) and Ang II (100 nM) than by stimulation with insulin alone (Fig. 1E). The growth characteristics of PC Cl3 cell line has been studied by means of flow cytometry analysis. The histograms depict the cell-cycle distribution of PC Cl3 cells grown in medium supplemented with FBS and six hormones mixture (Fig. 1A) and after 48 h starvation (Fig. 1B). More than 90% of PC Cl3 cells appeared blocked in G0/G1 phases and Ang II (100 nM) was not able to increase the G1/S progression in the cell cycle (Fig. 1C).

Ang II (100 nM) did not increase the protein/cell number ratio (control cells = 188.5 ± 7.4 pg/cell; cells treated with 100 nM Ang II = 191 ± 8.7 pg/cell), indicating that Ang II did not cause hypertrophy as in other cell types.

Effects of Ang II on cyclin E, Cdk2 and p27^{kip} expressions

Cyclin E, Cdk2 but not p27^{kip} were expressed in starved PC Cl3 cells. Treatment with 100 nM Ang II caused a small but significant increase in cyclin E expression, which reached a maximum at 20 h (Fig. 2B), had caused no effect on Cdk2 (Fig. 2C) and a significant increase in p27^{kip} after 4-h incubation with Ang II, and reaching a maximum at 20 h (Fig. 2A).

Ang II induces phosphorylation of PKB/Akt

Since Ang II is known to increase, in some cells, the phosphorylation state of Akt/PKB (Takahashi *et al.* 1999), we studied the effects of Ang II treatment on PKB/Akt. Stimulation of PC Cl3 cells with Ang II increased the phosphorylation state of PKB/Akt in a time and concentration dependent manner. Ang II caused the phosphorylation of Akt/PKB within 3 min, reached a maximum at 1 h and remained unchanged for up to 24 h (Fig. 3A). The maximum effect was achieved at 100 nM of Ang II (Fig. 3B).

Role of PI3K and MEK on p27^{kip} induction

Since the ERK 1/2 and PI3K pathways are independently activated, and implicated in Ang II-mediated DNA synthesis and cell number increase in various cells (Dugourd *et al.* 2003),

we further examined the effect of the PI3K inhibitor LY294002 and of the MEK inhibitor PD98059 on Ang II-induced p27^{kip} activation. Pre-treatment of PC Cl3 cells with 1–100 μ M LY294002 enhanced the Ang II-stimulated phosphorylation of p27^{kip} (Fig. 4), while pre-treatment of PC Cl3 cells with 1–25 μ M PD98059 did not have any significant effect on the Ang II-stimulated phosphorylation of p27^{kip} (Fig. 4).

Role of PKC- ζ on p27^{kip} induction

Since, we showed previously that Ang II activates ERK pathway through the activity of atypical PKC- ζ , PC Cl3 cells were also pre-treated with the PKC inhibitor GF109203X. As shown in Fig. 4, GF109203X greatly inhibited the induction of p27^{kip} at the highest concentration used; since GF109203X has a half-maximal inhibitory constant (IC₅₀) for PKC- ζ isozyme > 1.5 μ M (whereas of 110 nM or lower for all the other PKC isoforms (Martiny-Baron *et al.* 1993), which are not activated by Ang II in these cells) these results suggest that a atypical PKC isoform is involved in the induction of p27^{kip} by Ang II.

PC Cl3 cells were also pre-incubated with 10, 50, and 100 μ M cell permeable ζ -PS for 60 min. ζ -PS did not affect the basal p27^{kip} but provoked a dose-dependent inhibition of Ang II-induced p27^{kip} activation (Fig. 5A).

We next examined the effect of the downregulation of PKC- ζ isoform by siRNA, on p27^{kip} induction. Preliminary experiments by western blotting using antibodies against PKC- ζ , demonstrated the efficient downregulation induced by siRNA (Fig. 5B). Densitometric analysis revealed a 76% reduction in endogenous PKC- ζ protein expression following transfection with 30 nM siRNA- ζ for 24 h (Fig. 5C).

The siRNA inhibition of PKC- ζ prevent the Ang II-mediated increase of p27^{kip} (Fig. 5B).

These results confirmed that atypical PKC- ζ is involved in the induction of p27^{kip} by Ang II.

Ang II induces c-fos expression in PC Cl3

Western blot analysis using an antibody that recognized the c-fos protein showed that in quiescent PC Cl3 cells Ang II stimulates the expression of this protein in a dose-dependent way (1–1000 nM; Fig. 6). The expression of c-fos proto-oncogene protein peaked at 30 min. On the basis of these results further experiments were carried out using 30-min incubation time for protein detection.

The effects of insulin on PKC- ζ , ERK1/2, PKB, and p27^{kip} in PC Cl3 cells

As reported above (Fig. 1A), insulin has a replicating effect on PC Cl3 cells. Noteworthy, the proliferation of PC Cl3 cells incubated with insulin and Ang II in combination (used in a single dose of 100 nM) appeared further increased. We therefore studied the effects of 100 nM insulin on the same

Figure 2 Effects of Ang II on the expression level of (A) p27^{kip}, (B) cyclin E and (C) Cdk2 in PC Cl3 cells. Starved PC Cl3 cells were stimulated with 100 nM Ang II for different times over a 24 h period. Equal amounts of lysate protein were resolved by SDS-gel electrophoresis on 10% acrylamide gel and transferred to polyvinylidene difluoride (PVDF) membrane. The membrane was probed with anti-p27^{kip}, anti-cyclin E and anti-Cdk2 antibodies and the proteins visualized by chemiluminescence detection. To confirm equal protein loading in western blot, the membranes were re-probed with β -actin antibody. Densitometry analysis of bands obtained in other independent experiments ($n=4$) performed on different cell preparations are represented as histograms. Values with shared letters are not significantly different according to Bonferroni/Dunn *post hoc* tests. CycE, cyclin E.

Figure 3 The Ang II-induced phosphorylation of Akt in PC Cl3 cells. (A) Starved PC Cl3 cells were treated without or with 100 nM Ang II for the indicated periods. (B) Starved cells were incubated with increasing concentration of Ang II for 60 min. The cell extracts were electrophoretised through 10% SDS-PAGE gel and analyzed by western blotting using anti-phospho-Akt (p-Akt) antibody, and after stripping with anti-total (phosphorylated plus unphosphorylated) Akt (Akt) antibody. Representative immunoblots are shown. Densitometry analysis of bands obtained in other independent experiments ($n=4$) performed on different cell preparations, are represented as histograms. Values with shared letters are not significantly different according to Bonferroni/Dunn *post hoc* tests.

proteins, which are targets of the Ang II action. As shown in Fig. 7 PC Cl3 cells incubated with 100 nM insulin showed a significant decrease of p27^{kip}, a small but significant increase in cyclin E expression, no effects on Cdk2 (Fig. 7A) and the

Figure 4 Role of PI3K, ERK1/2 and PKC- ζ on the effects of Ang II on p27^{kip} protein. Starved PC Cl3 cells were pre-treated for 45 min with LY294002 (LY), PD98059 and GF109203X (GX) and then stimulated with 100 nM Ang II for 20 h. Cells were also treated for 45 min with 100 μ M LY294002 and 10 μ M GF109203X only. The resolved proteins were immunoblotted with anti-p27^{kip} and the proteins visualized by chemiluminescence detection. To confirm equal protein loading in western blot, the membranes were re-probed with β -actin antibody. Densitometry analysis of bands obtained in other independent experiments ($n=4$) performed on different cell preparations, are represented as histograms. Values with shared letters are not significantly different according to Bonferroni/Dunn *post hoc* tests.

Figure 5 Role of PKC- ζ on p27^{kip} induction. (A) Starved PC Cl3 cells were pre-treated with increasing concentrations of a synthetic myristoylated pseudosubstrate of PKC- ζ (ζ -PS) for 60 min and then stimulated with 100 nM Ang II for 20 h. (B) Cells were also transfected with siRNA- ζ and then stimulated without or with 100 nM Ang II for 20 h. The resolved proteins were immunoblotted with anti-p27^{kip} and the proteins visualized by chemiluminescence detection. (C) Analysis of PKC- ζ expression in siRNA- ζ transfected or not PC Cl3 cells. To confirm equal protein loading in western blot, the membranes were reprobed with β -actin antibody. The figure is representative of three independent experiments. C, control.

phosphorylation of PKB (Fig. 7B); finally, insulin did not provoke translocation of PKC- ζ from the cytosol to cellular membranes (Fig. 7D). Interestingly, the effect of the simultaneous stimulation with insulin and Ang II on p27^{kip} expression (Fig. 7C) and on PKC- ζ translocation (Fig. 7D) was similar to the effect obtained with insulin alone.

Discussion

In the present study, we have investigated the growth-promoting effects of Angiotensin II (Ang II) in PC Cl3 cells since, as previously shown (Marsigliante *et al.* 2003), Ang II stimulates the phosphorylation of the extracellular signal-

Figure 6 The Ang II-induced expression of c-fos protein in PC Cl3 cells. (A) Starved PC Cl3 cells were treated without or with 100 nM Ang II for the indicated periods. (B) Cells were incubated with increasing concentration of Ang II for 30 min in serum-starved medium. The cell extracts were electrophoresed through 10% SDS-PAGE gel and analyzed by western blotting using anti-c-fos. The same results were obtained in other independent experiments ($n=4$) performed on different cell preparations. Representative immunoblots are shown. Values with shared letters are not significantly different according to Bonferroni/Dunn *post hoc* tests.

regulated kinases 1 and 2 (ERK1 and 2). ERKs are involved in the activation of nuclear transcription factors, such as the c-fos, c-jun, and c-myc protooncogenes (Wagstaff *et al.* 2000, Buckley *et al.* 2001, Ekokoski *et al.* 2001). c-fos and c-jun proteins associate to form homo- and hetero-dimer AP-1 complexes that bind to specific DNA sequences of target genes. The regulatory region of the c-fos gene contains several *cis*-acting elements, which are the target of a variety of interdependent signal transduction pathways, such as PKA, PKC, calmodulin (CAM) kinase, and mitogen activated protein (MAP) kinase cascades. c-fos is induced by all studied (co)mitogenic stimuli, including TSH (cAMP), insulin/insulin-like growth factor-I, phorbol esters, serum and growth factors in FRTL-5 (Colletta *et al.* 1986, Isozaki & Kohn 1987), and Wistar rat thyroid (WRT) cells (Tominaga *et al.* 1994), and in dog (Deleu *et al.* 1999) and human thyrocytes (Huber *et al.* 1991, Heinrich & Kraiem 1997). c-fos expression has been claimed to be required for TSH-dependent proliferation of Fisher rat thyroid low serum

Figure 7 Effects of insulin on (A) the expression level of p27^{kip}, cyclin E (Cyc E) and Cdk2, (B) the phosphorylation state of Akt, (D) the PKC- ζ translocation from cytosol (cyt) to membrane (mem) in PC Cl3 cells. (C) Effects of the simultaneous stimulation with insulin and Ang II on expression level of p27^{kip}, cyclin E and Cdk2. (A and B) Starved PC Cl3 cells were stimulated with 100 nM insulin for the indicated periods. Equal amounts of lysate protein were resolved by SDS-gel electrophoresis on 10% acrylamide gel and transferred to PVDF membrane. The membranes were probed with (A) anti-p27^{kip}, anti-cyclin E, anti-Cdk2 or (B) anti-phospho-Akt (p-Akt), and after stripping with anti-total (phosphorylated plus unphosphorylated) Akt (Akt) antibodies and the proteins visualized by chemiluminescence detection. To confirm equal protein loading in western blot, the membranes were reprobed with β -actin antibody. (C) Starved PC Cl3 cells were incubated with insulin and Ang II in combination (used in a single dose of 100 nM) and equal amounts of lysate protein were resolved by SDS-gel electrophoresis on 10% acrylamide gel and transferred to PVDF membrane. The membranes were probed with anti-p27^{kip}, anti-cyclin E and anti-Cdk2 antibodies. To confirm equal protein loading in western blot, the membranes were reprobed with β -actin antibody. (D) Starved PC Cl3 cells were treated with insulin (100 nM) or with insulin (100 nM) in combination with Ang II (100 nM) for the indicated periods and cytosol (cyt) and membrane (mem) fractions were used for immunoblotting with PKC- ζ specific antibody. Each figure is representative of four independent experiments performed on different cell preparations.

(FRTL)-5 cells (Foti *et al.* 1990), and the activation of AT1 receptors has been shown to induce, in other cell systems, the expression of c-fos (Muscella *et al.* 2002).

The results of the present study showed that Ang II activated ERK signal transduction pathway and induced c-fos expression without acting as a mitogen in PC Cl3 cells. Since, in some cell types, such as vascular smooth muscle cells, the Ang II-induced proliferation appeared mediated by the stimulation of ERK1/2 cascade (Marrero *et al.* 1997) together with the activation of PI3K (Saward & Zahradka 1997), we consequently investigated the Ang II-induced Akt activation also in PC Cl3 cells. The results demonstrated that Ang II induced Ser-437 phosphorylation of Akt, an activation that is not sufficient for proliferation of PC Cl3 cells. However, although both ERK and PI3K/Akt pathways are activated, the effects Ang II in PC Cl3 proliferation are not evident. Since progression through the cell cycle is controlled by a series of cyclin and Cdk complexes that may be inactivated by Cdk inhibitors, to further determine the role of Ang II in PC Cl3 cells, we studied the expression of cyclin E, Cdk₂ and the Cdk-inhibitor p27^{kip} in PC Cl3 cells stimulated with Ang II. Compared with the cells grown in serum-free medium, Ang II treatment enhanced the expression of p27^{kip} protein; this may cause a block in the G1 phase of Ang II-stimulated PC Cl3 cells, which are not able to progress into the S phase. It is interesting to speculate that G1 phase entry coupled with subsequent cell-cycle arrest may be a general mechanism for cellular hypertrophy. Nevertheless, Ang II did not increase the protein/cell number ratio indicating that it did not cause hypertrophy as in other cell types.

Thus these observations prompted us to further investigate the relationship between PI3K, ERK1/2 and p27^{kip}. We took advantage of the well-characterized PI3K inhibitor LY294002 and of the MEK inhibitor PD98059. We found that LY294002 increases p27^{kip} induction via Ang II, while PD98059 had no effects (Fig. 4). This may be due to the fact that inhibition of the PI3K/Akt pathway resulted in loss of p27^{kip} from cyclin D1 complexes, a shift of p27^{kip} to cyclin E-Cdk2 both of which precede G1 arrest (Liang *et al.* 2002). Since insulin is able to stimulate the proliferation of PC Cl3 cells and activates the same pathways that are activated by Ang II, other than the cytosol-to-membrane translocation of PKC- ζ , we examined the effects of the inhibition of this upstream activator of ERK1/2. We found that GF109203X greatly inhibited p27^{kip} induction via Ang II, suggesting that PKC- ζ may reduce p27^{kip} ubiquitination and degradation.

Furthermore, the role of PKC- ζ was proved by the use of its pseudosubstrate that efficiently inhibited the Ang II response on p27^{kip} induction and also by PKC- ζ down-regulation using the siRNA (Fig. 5).

However, the exact signal transduction pathways by which Ang II causes cell-cycle arrest are unknown, while it is clear that Ang II activates multiple signaling pathways, including PKC- ζ , MAPK and Akt that control cellular growth responses. In tubular cells Ang II leads to phosphorylation of the mitogen-activated protein kinases ERK1/2 that phosphorylate p27^{kip} at serine

residues. The p27^{kip} phosphorylation at serine/theonine residues leads, in contrast to tyrosine phosphorylation, to an increased stability and less degradation through the ubiquitin pathway (Wolf & Wenzel 2003). Therefore, it is possible that Ang II-stimulated PKC- ζ activation has a similar role in the processes that regulate p27^{kip} stability, although we can only speculate at this point. Consistent with the higher proliferative effects of co-stimulation with insulin and Ang II, it would be reasonable to postulate that an additional functional pathway exists by which insulin can modulate the stability of p27^{kip} at least through the inhibition of the Ang II-activated PKC- ζ pathway.

In summary, the results of the present study provide demonstration that (1) in PC Cl3 cells Ang II stimulates proliferation, but at the same time it stimulates growth inhibitory systems causing a block in G1 phase, although both MAPK and PI3K/Akt pathways are active and (2) this effect may be, at least in part, due to the upregulation of p27^{kip} controlled by the activity of PKC- ζ .

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