Vitamin A inhibits pancreatic stellate cell activation: implications for treatment of pancreatic fibrosis

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Background and aims: Activated pancreatic stellate cells (PSCs) are implicated in the production of alcohol induced pancreatic fibrosis. PSC activation is invariably associated with loss of cytoplasmic vitamin A (retinol) stores. Furthermore, retinol and ethanol are known to be metabolised by similar pathways. Our group and others have demonstrated that ethanol induced PSC activation is mediated by the mitogen activated protein kinase (MAPK) pathway but the specific role of retinol and its metabolites all-trans retinoic acid (ATRA) and 9-cis retinoic acid (9-RA) in PSC quiescence/activation, or its influence on ethanol induced PSC activation is not known. Therefore, the aims of this study were to (i) examine the effects of retinol, ATRA, and 9-RA on PSC activation; (ii) determine whether retinol, ATRA, and 9-RA influence MAPK signalling in PSCs; and (iii) assess the effect of retinol supplementation on PSCs activated by ethanol.

Methods: Cultured rat PSCs were incubated with retinol, ATRA, or 9-RA for varying time periods and assessed for: (i) proliferation; (ii) expression of α smooth muscle actin (α-SMA), collagen I, fibronectin, and laminin; and (iii) activation of MAPKs (extracellular regulated kinases 1 and 2, p38 kinase, and c-Jun N terminal kinase). The effect of retinol on PSCs treated with ethanol was also examined by incubating cells with ethanol in the presence or absence of retinol for five days, followed by assessment of α-SMA, collagen I, fibronectin, and laminin expression.

Results: Retinol, ATRA, and 9-RA significantly inhibited: (i) cell proliferation, (ii) expression of α-SMA, collagen I, fibronectin, and laminin, and (iii) activation of all three classes of MAPKs. Furthermore, retinol prevented ethanol induced PSC activation, as indicated by inhibition of the ethanol induced increase in α-SMA, collagen I, fibronectin, and laminin expression.

Conclusions: Retinol and its metabolites ATRA and 9-RA induce quiescence in culture activated PSCs associated with a significant decrease in the activation of all three classes of MAPKs in PSCs. Ethanol induced PSC activation is prevented by retinol supplementation.

Pancreatic fibrosis is a common histopathological feature of alcoholic chronic pancreatitis. Fibrogenesis involves a specific cell type in the pancreas, namely the pancreatic stellate cell (PSC). Recent studies have demonstrated that PSCs become activated (as indicated by proliferation, increased expression of the cytoskeletal protein α smooth muscle actin (α-SMA), and synthesis of extracellular matrix proteins) when cultured on plastic or when exposed to ethanol and its oxidative metabolite acetaldehyde or to factors known to be upregulated during pancreatic injury such as proinflammatory cytokines, growth factors, and oxidant stress.

PSCs in their quiescent (non-activated) state store retinol as lipid droplets in their cytoplasm. Activation of PSCs is invariably associated with loss of the retinol containing droplets from their cytoplasm. Studies in other cell systems have established that retinol is a compound essential to normal cell biology. In particular, metabolites of retinol, all-trans retinoic acid (ATRA) and 9-cis retinoic acid (9-RA), have been shown to mediate a number of cellular functions, including proliferation, differentiation, and protein synthesis. Metabolism of retinol has been best studied in the liver. It has been demonstrated in hepatic stellate cells (HSCs) that retinol enters the cell bound to cellular retinol binding protein. It is then either esterified to retinyl ester for storage or converted to retinoldehyde by retinol dehydrogenase (RolDH) and subsequently to retinoic acid by retinaldehyde dehydrogenase (RALDH). Two forms of retinoic acid, ATRA and 9-RA, serve as ligands for two families of nuclear receptors: retinoic acid receptors (RARs, β, and γ) and retinoid X receptors (RXRs, β, and γ). RARs bind to ATRA with high affinity whereas 9-RA is a bifunctional ligand which can bind to and activate both RARs and RXRs. Following ligand binding, these compounds interact with cis acting DNA sequences called retinoic acid responsive elements in the promoter regions of target genes, thereby regulating gene expression. It is of interest that studies with HSCs have demonstrated that on exposure to activating factors, levels of retinoic acid and expression of their receptors is decreased. Moreover, maintenance of the quiescent phenotype of HSCs has been shown to be dependent on adequate levels of retinoic acid and its receptors in the cells. A recent study by Jaster and colleagues has reported decreased proliferation and collagen synthesis in PSCs on exposure to ATRA. However, there has been no comprehensive study in the literature to date examining the effects of retinol and both of its metabolites (ATRA and 9-RA) on PSC function. A number of observations suggest a link between metabolism of ethanol and retinol. Ethanol and its oxidative metabolism...
metabolite acetaldehyde are key activating factors of PSCs. Oxidation of ethanol to acetaldehyde is mediated by the enzyme alcohol dehydrogenase (ADH). Acetaldehyde is then further oxidised to acetic acid via aldehyde dehydrogenase. It has been established that the retinol metabolising enzymes RolDH and RALDH belong to the same family of alcohol and acetaldehyde metabolising dehydrogenases, respectively. Interestingly, studies have shown that rats chronically exposed to ethanol display significantly reduced retinoic acid levels in the liver. This is thought to be due to competitive inhibition by ethanol of retinol metabolism by RolDH and ADH as these enzymes can utilise both ethanol and retinol as substrates. Thus retinoic acid depletion may be an important mechanism by which ethanol promotes stellate cell activation.

In recent studies, we and others have identified the mitogen activated protein kinase (MAPK) pathway as a major cell signalling pathway mediating ethanol induced and growth factor induced PSC activation. MAPKs (which include extracellular regulated kinases 1 and 2 (ERK1/2), N terminal c-Jun kinase (JNK), and p38 kinase) play a major role in regulating protein synthesis in mammalian cells. MAPKs modulate the activity of a number of downstream transcription factors and protein kinases by phosphorylation, thereby controlling gene expression and cell behaviour. Activation of MAPKs is a reversible process; they are inactivated by a group of protein phosphatases known as MAP kinase phosphatases (MKPs). These are dual specificity phosphatases which have been shown to exert their effect by dephosphorylating tyrosine and threonine residues on MAPKs, thereby leading to their inactivation. MAPK phosphatase 1 (MKP-1), a member of the MKP family, has recently been demonstrated to inactivate all three classes of MAPKs. It is interesting to note that recent studies have shown that MKP-1 expression is increased on exposure to retinoic acid thereby resulting in a decrease in MAPK activity. However, whether retinol influences the MAPK signalling pathway or MKP-1 expression in stellate cells (whether from the pancreas or liver) is not known.

Therefore, the aims of this study were to: (i) examine the effect of retinol and its metabolites ATRA and 9-RA on PSC activation, (ii) determine whether retinol influences MAPK signalling and MKP-1 expression in PSCs, and (iii) assess the influence of retinol on ethanol treated PSCs.

METHODS

Cell culture reagents, protease type XIV, all-trans retinol, ATRA, and 9-RA were purchased from the Sigma Chemical Company (St Louis, Missouri, USA). Primers for RolDH II, RARα, RXRα, and RXRβ were obtained from Sigma Genosys Australia Pty Ltd. Sources of antibodies were as follows: a-SMA, fibronectin, and laminin—Sigma; phospho-ERK1/2 and total ERK1/2, total p38 kinase and total JNK—Cell Signaling Technology (Beverly, Massachusetts, USA); phospho-p38 kinase and phospho-JNK—Promega (Sydney, Australia); MKP-1, RARα, RXRα, and RXRβ—Santa Cruz Biotechnology (Santa Cruz, California, USA); collagen type I—Rockland Immunochromicals (Gilbertsville, Pennsylvania, USA); and secondary goat antirabbit antibody—Dako (Sydney, Australia).

Isolation and culture of PSCs

Rat PSCs were isolated by density gradient centrifugation, as detailed previously. This technique results in a pure preparation of PSCs, as evidenced by positive staining for stellate cell selective markers and negative staining for possible contaminants such as endothelial cells and macrophages. The yield of PSCs is 1.5–2 million cells per rat pancreas.) Cells were cultured in Iscove’s modified Dulbecco’s medium (IMDM) supplemented with 10% fetal bovine serum at 37°C in a humidified 95% air/5% CO2 atmosphere. All experiments were performed with culture activated cells (passages 1–3) with the exception of experiments to assess the effect of retinol on freshly isolated (quiescent) cells.

Expression of retinol dehydrogenase and RAR and RXR receptor mRNA using RT-PCR

Expression of mRNA for RolDH II and the retinoic acid receptors RARα, RXRα, and RXRβ was examined using reverse transcriptase-polymerase chain reaction (RT-PCR). Primer sequences were as follows:

- Rat RolDH II
  - Forward primer: 5’ ACC TGG CAT CTT ATC TGA AA 3’
  - Reverse primer: 5’ AGT CGA GTC AGC CTT GAG TA 3’
- Rat RARα
  - Forward primer: 5’ CCC AGC CAC CAT TGA GAC 3’
  - Reverse primer: 5’ TAC ACC ATG TTC TTT AGG ATC 3’
- Rat RXRβ
  - Forward primer: 5’ TCA ACT CCA CAG TGT CAC GTG G 3’
  - Reverse primer: 5’ TAA ACC CCA TAG TGC TTC CC 3’

Treatment with test factors

Treatment of PSCs with retinol and retinoic acid

Culture activated PSCs (passages 1–3) were exposed to retinol (10 μM), ATRA (10 μM), or 9-RA (10 μM) for varying periods of time. Medium was changed every 48 hours. All manipulations were performed in subdued light. Cells incubated with culture medium with an equivalent amount of vehicle—DMSO 0.1% (for ATRA and 9-RA experiments) or DMSO 0.025% + 8 mM ethanol (for retinol experiments)—served as controls. Cell viability was assessed by trypan blue exclusion studies.

Treatment of PSCs with ethanol and retinol

To determine the effect of retinol on PSCs treated with ethanol, cells were incubated with 10 μM retinol in the presence or absence of 50 mM ethanol for five days. Medium was changed every 24 hours. At the end of the incubation period, cell lysates were collected and expression of α-SMA, collagen I, fibronectin, and laminin was assessed by western blotting. Cell viability was assessed as described previously.

Assessment of PSC function

Cell proliferation

PSC proliferation was assessed by measuring incorporation of [3H] thymidine into cellular DNA, as previously described. In addition, cell counts were performed using a haemocytometer. Results are expressed as a percentage of control.

α-Smooth muscle actin and extracellular matrix protein expression

Levels of α-SMA, collagen I, fibronectin, and laminin in PSCs were determined by western blotting of cell lysates, as detailed previously, using a monoclonal mouse anti-α-SMA (1:200) or polyclonal rabbit anticallogen I (1:1000), antifibronectin.
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Accession of RAR and RXR protein expression in PSCs

Assessment of RAR and RXR protein expression in PSCs

Detection of MKP-1 expression

Treatment of freshly isolated PSCs with retinol

Figure 1 (A, B) Expression of retinol dehydrogenase II (RolDH II), retinoic acid receptor (RAR), and retinoid X receptor (RXR) in pancreatic stellate cells (PSCs). (A) Total RNA was extracted from three separate cell preparations and analysed for RolDH II expression by reverse transcriptase-polymerase chain reaction (RT-PCR). An RNA ladder was run in lane 1 (Std) to determine the size of the PCR products observed. Lane 2 contained rat liver cDNA as a positive control. Lane 3 contained a PCR product for RolDH II from rat PSCs. Cont, control. (B) RARα, RXRα, and RXRβ expression was analysed in total RNA obtained from three separate cell preparations by RT-PCR. Lane 1 contained an RNA ladder (Std) to determine the size of the PCR products observed. Lanes 2, 3, and 4 contained PCR products for RARα, RXRα, and RXRβ from rat PSCs.

Figure 2 Effect of retinol and its metabolites on pancreatic stellate cell (PSC) proliferation. (A) [3H] thymidine incorporation studies and (B) cell count data demonstrating that incubation of PSCs with retinol (Rol) decreased PSC proliferation significantly when compared with controls (Cont) after 72 hours and that this effect was sustained over 96 hours. All-trans retinoic acid (ATRA) and 9-cis retinoic acid (9-RA) also significantly decreased PSC proliferation after 48 hours when compared with controls and this effect was sustained over 96 hours (n = 3 separate cell preparations; ***p<0.001).

Detection of MAPK activation

Using a polyclonal rabbit anti-MKP-1 antibody (1:1000). To determine the role of MKP-1 in regulating the action of ATRA on MAPKs, we examined whether any observed inhibition of MAPKs by ATRA could be prevented by the protein phosphatase inhibitor sodium orthovanadate (SV), a potent inhibitor of protein tyrosine phosphatases, which include MKP-1.11 12 The influence of this inhibitor on PSC function (cell proliferation and extracellular matrix protein expression) was also assessed. PSCs were treated with 10 μM ATRA in the presence or absence of 1 μM SV for 48 hours. At the end of the incubation period, cell counts were performed using a haemocytometer and cell lysates were collected for assessment of MAPK activation and extracellular matrix protein expression by western blotting, as described above.

Effect of retinol and its metabolites on pancreatic stellate cell proliferation, showing significant differences in [3H] thymidine incorporation and cell counts between control (Cont) and treated groups.*p<0.05, **p<0.01, ***p<0.001.

A

B

Detection of MAPK activation

ERK1/2, p38 kinase, and JNK activation in culture activated PSCs exposed to retinol (10 μM), ATRA (10 μM), or 9-RA (10 μM) for 4, 24, and 48 hours was assessed by western blotting of cell lysate proteins, as described previously.17 26 To confirm that total MAPK levels were unchanged by the treatments, aliquots of the above cell lysates were subjected to western blotting analysis using rabbit polyclonal antibodies against ERK1/2, p38 kinase, or JNK (1:1000), which recognise both the phosphorylated and non-phosphorylated forms of the enzymes; the blots were then analysed by densitometry.

To determine whether retinol could prevent/retard the transition of freshly isolated (quiescent) PSCs to their activated phenotype, freshly isolated cells were incubated in IMDM supplemented with 10% fetal bovine serum in the presence or absence of 1 μM SV for 24 hours. The influence of SV on the expression of MAPKs in PSCs was assessed using western blotting analysis with rabbit polyclonal antibodies against ERK1/2, p38 kinase, or JNK (1:1000).

Assessment of RAR and RXR protein expression in PSCs

RAR and RXR expression in PSCs treated with 10 μM ATRA or 10 μM 9-RA for 4, 24, and 48 hours was assessed by western blotting using polyclonal rabbit antibodies against RARα, RXRα, or RXRβ (1:2000).

Detection of MKP-1 expression

MKP-1 expression in PSCs incubated with retinol, ATRA, or 9-RA for 3, 6, and 24 hours was assessed by western blotting using polyclonal rabbit antibodies against MKP-1 (1:1000), and antiammonio (1:1000) antibodies.29 30 A goat antimouse antibody (1:2000) or a goat antirabbit antibody (1:2000) was used as the secondary antibody. Expression of α-SMA, collagen I, fibronectin, and laminin was detected by the enhanced chemiluminescence (ECL) technique using the Amersham’s enhanced ECL kit, and quantified using densitometry (BioRad GelDoc Image One software).

Effect of retinol and its metabolites on pancreatic stellate cell (PSC) proliferation. (A) [3H] thymidine incorporation studies and (B) cell count data demonstrating that incubation of PSCs with retinol (Rol) decreased PSC proliferation significantly when compared with controls (Cont) after 72 hours and that this effect was sustained over 96 hours. All-trans retinoic acid (ATRA) and 9-cis retinoic acid (9-RA) also significantly decreased PSC proliferation after 48 hours when compared with controls and this effect was sustained over 96 hours (n = 3 separate cell preparations; ***p<0.001).

A

B

RESULTS

Expression of retinol dehydrogenase II and retinoic acid receptors in PSCs

Using RT-PCR, rolDH II, RARα, RXRα, and RXRβ expression was observed in PSCs (fig 1A). PCR products were 398 bp, 195 bp, 165 bp, and 175 bp, respectively, in agreement with previous reports.26 27

Retinol and its metabolites ATRA and 9-RA induce PSC quiescence

Cell proliferation

Retinol, ATRA, and 9-RA reduced the proliferation of culture activated PSCs (n = 4 separate cell preparations), as assessed by thymidine incorporation into cellular DNA as well as by direct cell counts. There was a significant reduction after 72 hours of incubation with retinol and this effect was sustained over 96 hours (fig 2). Proliferation was also significantly reduced in the presence of ATRA and 9-RA (fig 2). Notably, this reduction was observed at an earlier time point (48 hours) than that produced by retinol and this effect was sustained over 96 hours. We confirmed that the observed reduction in PSC proliferation was not due to cytotoxicity of retinol, ATRA, or 9-RA, as evaluated by phase contrast microscopy and trypan blue exclusion studies (results not shown).

α-SMA expression

Quiescence of PSCs after incubation with retinol (n = 3 separate cell preparations) was also demonstrated by significantly reduced α-SMA expression after 72 hours, an effect that was sustained over 96 hours (fig 3). Similar results were obtained with ATRA and 9-RA (fig 3). ATRA was also associated with reduced α-SMA expression at the earlier time point of 48 hours.

Extracellular matrix protein expression

Extracellular matrix protein expression was significantly reduced in PSCs exposed to retinol or its metabolites (n = 3 separate cell preparations; fig 4A–C). After 48 hours of incubation, retinol significantly decreased collagen I, fibronectin, and laminin expression and this effect was sustained over 72 hours. Similar results were obtained for ATRA and 9-RA.

Retinol and its metabolites ATRA and 9-RA influence MAPK activation in PSCs

ERK1/2 activation

Incubation of PSCs with retinol (n = 4 separate cell preparations) caused a significant decrease in ERK1/2 activation at 4 hours; this decrease was sustained over 48 hours (fig 5). Similarly, ERK1/2 activation was reduced in the presence of ATRA and 9-RA (fig 5). Western blotting of control and treated cell lysates for total ERK demonstrated that total ERK expression was unchanged by retinol, ATRA, and 9-RA treatment. (Densitometry data expressed as per cent of control (mean (SEM)): at 4 h retinol 100.5 (5.8), ATRA 99.6 (4.6); 9-RA 93.7 (3.6); at 24 h retinol 96.7 (9.2), ATRA 92.0 (8.9); 9-RA 102.7 (5.1); at 48 h retinol 107.75 (10.6), ATRA 111.5 (6.5); 9-RA 105.0 (7.2).) Thus these compounds appear to have a specific effect on ERK1/2 phosphorylation.

p38 kinase activation

Retinol significantly decreased p38 kinase activation after 48 hours of incubation (n = 4 separate cell preparations; fig 6). Both ATRA and 9-RA also significantly reduced p38 kinase activation. However, this reduction was observed at the earlier time points of 4 and 24 hours and was sustained over 48 hours (fig 6). Total p38 kinase expression was unchanged by the treatments. (Densitometry data expressed as per cent of control (mean (SEM)): at 4 h retinol 102.8 (3.9), ATRA 94.2 (3.96); 9-RA 90.57 (4.8); at 24 h retinol 93.07 (2.8), ATRA 98.6 (0.48); 9-RA 99.2 (2.8); at 48 h retinol 90.55 (4.86), ATRA 95.4 (3.0); 9-RA 102.6 (5.8)).

JNK activation

In contrast with the effect on ERK and p38 kinase activation, incubation of PSCs with retinol or 9-RA had no effect on JNK 2 (p54) activation (n = 4 separate cell preparations; fig 7). However, ATRA significantly decreased JNK 2 (p54) activation after 4 hours and this decrease was sustained over 48 hours (fig 7). Total JNK expression was not affected by retinol, ATRA, or 9-RA treatment. (Densitometry data expressed as per cent of control (mean (SEM)): at 4 h retinol 95.4 (10.0), ATRA 89.3 (8.7); 9-RA 103.7 (6.9); at 24 h retinol
Effect of retinol, ATRA, and 9-RA on MKP-1 expression

ATRA significantly increased MKP-1 expression in PSCs (n = 3 separate cell preparations) at 6 hours and this increase was sustained over 24 hours (fig 8). In contrast, retinol and 9-RA had no effect on MKP-1 expression (results not shown).

Effect of the protein phosphatase inhibitor sodium orthovanadate on MAPK expression, cell proliferation, and extracellular matrix protein expression in ATRA treated PSCs

To confirm whether MKP-1 plays a role in mediating the effect of ATRA on MAPK activation in PSCs, cells were treated with the protein phosphatase inhibitor SV (n = 3 separate cell preparations). As shown in fig 9, the ATRA induced decrease in activation of ERK1/2, p38 kinase, and JNK 2 was completely prevented in the presence of SV, suggesting that MKP-1 may mediate the ATRA induced inhibition of MAPK activation. SV treatment of PSCs also prevented the ATRA induced decrease in: (i) PSC proliferation (data expressed as per cent of control (mean (SEM)): control 100, ATRA 44.74 (8.72) (p < 0.01 ATRA v control), SV 85.35 (16.5), ATRA+SV 98.03 (26.8) (p < 0.01 ATRA v control)) and (ii) collagen I, fibronectin and laminin expression (fig 10), further supporting the concept that the effects of ATRA on PSC function may be mediated via the MAPK pathway. Trypan blue exclusion studies confirmed cell viability in the presence of SV (results not shown).
Effect of retinol on ethanol induced PSC activation

In order to determine whether retinol supplementation could prevent ethanol induced PSC activation, α-SMA expression was assessed (n = 3 separate cell preparations; fig 12). As expected, ethanol alone significantly increased α-SMA expression confirming our previously published results. Retinol alone significantly reduced α-SMA expression confirming our results described in fig 3. Importantly, retinol also significantly reduced α-SMA expression in PSCs treated with ethanol. Of particular interest was the finding that in the presence of ethanol, retinol was unable to fully exert its inhibitory effect when compared with retinol alone (compare Rol with E50+Rol, fig 12).

Extracellular matrix protein expression

As illustrated in fig 13, retinol significantly inhibited expression of all three extracellular matrix proteins in PSCs (n = 3 separate cell preparations) and also prevented the
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ethanol induced increase in collagen I, fibronectin, and laminin expression in the cells. Interestingly, as with \( \alpha \)-SMA expression, retinol was unable to fully exert its inhibitory effect on collagen I and laminin expression in the presence of ethanol, compared with retinol alone.

**Effect of retinol on freshly isolated PSCs**

Incubation of freshly isolated PSCs with retinol for five days significantly decreased expression of both \( \alpha \)-SMA and fibronectin in cells compared with controls (Fig 14A, B).

**DISCUSSION**

This study provides novel data demonstrating that vitamin A (retinol) induces quiescence in both freshly isolated and culture activated PSCs. We have also shown that the metabolites of vitamin A (ATRA and 9-RA) inhibit the activation of culture activated PSCs. Notably, we have found that the above effects are associated with a significant decrease in activation of all three classes of MAPKs, suggesting that vitamin A induced quiescence is mediated via the MAPK pathway. Another novel finding of this study (of relevance to alcohol induced fibrosis) is the prevention of ethanol induced PSC activation by retinol. To the best of our knowledge, this aspect has not been previously studied in pancreatic or hepatic stellate cells.

The present study has also provided evidence indicating that PSCs possess the ability to convert retinol into its active metabolites. We have shown that cultured PSCs express the enzyme RolDH II, an enzyme which is essential for the conversion of cellular retinol to retinoic acid. Our findings also suggest the presence of a functional retinoic acid signalling pathway in PSCs. We have identified mRNA for RAR and RXR are transcriptionally activated by ATRA and 9-RA. The observed induction of receptor protein expression by their ligands in PSCs concurs with a previous report in cardiomyocytes, showing that both RAR and RXR are transcriptionally activated by ATRA and 9-RA.

The signalling pathways responsible for regulating PSC activation have been the focus of much interest in recent years, as therapeutic targeting of relevant signalling molecules may enable inhibition of PSC activation and prevention of fibrogenesis. Studies by our group and others have demonstrated that all three classes (ERK1/2, JNK, and p38 kinase) of the MAPK pathway play an important role in mediating PSC activation on exposure to profibrogenic factors. The ERK1/2 pathway has been shown to mediate PSC proliferation by increasing the activity of the transcription factor AP-1. Recently, our group and others have shown that the p38 kinase pathway plays a major role in mediating \( \alpha \)-SMA expression in culture activated and ethanol treated PSCs. Masamune and colleagues reported that both p38 kinase and JNK play an important role in mediating extracellular matrix protein synthesis in PSCs. Given the above, it would be reasonable to speculate that inhibition of PSC activation by vitamin A may be mediated by inhibition of...
the MAPK pathway. The results of our study support this concept. We have demonstrated for the first time that retinol, ATRA, and 9-RA significantly decrease activation of ERK1/2 and p38 kinase in culture activated PSCs after 4 hours of incubation and this decrease was sustained over 48 hours. In addition, ATRA (but not retinol or 9-RA) significantly decreased JNK 2 (p54) activation at 4, 24, and 48 hours. Our results with ATRA concur with those reported in human bronchial cells and hepatocytes.23–26 Both ERK1/2 and JNK MAPKs are known to regulate the activity of the transcription factor AP-1, which in turn is known to be an essential factor for the regulation of cell proliferation.22 On activation, AP-1 binds to a DNA sequence motif in target genes, thereby regulating their expression.27 A number of studies in other cell types have shown that increased AP-1 activity is essential in regulating cell proliferation (see review).28 It is of interest to note that on exposure to vitamin A, AP-1 activity is decreased in a number of different cell types, and this is thought to be one of the mechanisms whereby vitamin A inhibits cell proliferation.29

The mechanisms of MAPK inhibition (inactivation) by vitamin A are not yet determined. While activation of MAPK depends on phosphorylation of its tyrosine and threonine residues by upstream kinases, recent studies indicate that inactivation of MAPK is secondary to dephosphorylation via MKPs.23 MKP-1 has been shown to inactivate all three classes of MAPKs.24 Interestingly, studies in other cell types have reported that treatment with vitamin A increases expression of MKP-1 which in turn results in a decrease in MAPK activity.25 29 42 The current study has demonstrated that ATRA (but not retinol or 9-RA) induces expression of MKP-1 in PSCs. This induction is evident after 6 hours of incubation and is sustained over 24 hours. We have further demonstrated that the protein phosphatase inhibitor SV (known to inhibit phosphatases such as MKP-1) prevents the inhibition of MAPK activation in PSCs treated with ATRA. These findings support the concept that an increase in MKP-1 activity may play a role in the observed decrease in MAPK activation in PSCs exposed to ATRA. The mechanisms responsible for the retinol and 9-RA induced decrease in MAPK activation are unclear (given that these compounds did not increase MKP-1 activity in PSCs). It is possible that these compounds induce the activity of other phosphatases of the MAPK phosphatase family, as has been reported by Palm-Leis and colleagues30 in cardiomyocytes.

It is now generally accepted that ethanol induced PSC activation plays a major role in alcoholic pancreatic fibrosis.31 In view of our findings that retinol induces quiescence of PSCs in culture, we examined the effect of retinol on ethanol treated PSCs. Our results have shown that retinol prevents ethanol induced PSC activation, as evidenced by a significant

Figure 9  Effect of sodium orthovanadate (SV) on mitogen activated protein kinase (MAPK) activation in all-trans retinoic acid (ATRA) treated cells. Representative western blots and densitometry analysis showing a significant decrease in activation of all three classes of MAPK in pancreatic stellate cells (PSCs) treated with ATRA for 24 hours (ERK1/2, extracellular regulated kinases 1 and 2, p38 kinase; and JNK 2, c-Jun N terminal kinase 2). This decrease was prevented in the presence of SV (*p<0.02, **p<0.001, ATRA v control (Cont); tp<0.02, ttp<0.003, ATRA v ATRA+SV; n=3 separate cell preparations). Total MAPK levels were unchanged by the treatments.

Figure 10  Effect of sodium orthovanadate (SV) on all-trans retinoic acid (ATRA) induced inhibition of extracellular matrix protein expression in pancreatic stellate cells (PSCs). Representative western blots and densitometry analysis showing a significant decrease in collagen I, fibronectin, and laminin expression in PSCs treated with ATRA for 48 hours and prevention of this decrease in the presence of SV (*p<0.02, **p<0.03 ATRA v control (Cont); tp<0.04, ttp<0.02, ATRA v ATRA+SV; n=3 separate cell preparations).
decrease in α-SMA expression and, more importantly, a significant decrease in the expression of the extracellular matrix proteins collagen I, fibronectin, and laminin which are major components of fibrous tissue. In the course of these studies, an interesting observation was made with respect to the effect of retinol on PSC activation in the presence and absence of ethanol. In the presence of ethanol, retinol was found to be unable to exert its full inhibitory effect on

Figure 11 (A, B) Effect of all-trans retinoic acid (ATRA) and 9-cis retinoic acid (9-RA) on retinoic acid receptor (RAR) and retinoid X receptor (RXR) receptor expression in culture activated pancreatic stellate cells (PSCs). (A) Representative western blots and densitometry analysis showing a significant increase in RARα protein expression in PSCs treated with ATRA for 4, 24, and 48 hours (p < 0.04, p < 0.03, p < 0.02; n = 3 separate cell preparations). Cont, control. (B) Representative western blots and densitometry analysis showing a significant increase in RXRα and RXRβ protein expression in PSCs treated with 9-RA for 4, 24, and 48 hours (p < 0.04, p < 0.02, p < 0.03, p < 0.004, p < 0.005; n = 3 separate cell preparations). Note that both RXRα and RXRβ were represented by a number of protein bands with increased intensity, suggesting the presence of multiple activated isoforms.

Figure 12 Effect of retinol on ethanol induced α smooth muscle actin (α-SMA) expression in pancreatic stellate cells (PSCs). (A) Representative western blot showing α-SMA expression in PSCs incubated with culture medium alone (Cont) or with 50 mM ethanol (E50) in the presence or absence of retinol (Rol) for five days. (B) Densitometry analysis of western blots showing a significant increase in α-SMA expression in PSCs when treated with ethanol (n = 3 separate cell preparations; **p < 0.01) compared with controls. Rol significantly decreased basal α-SMA expression (n = 3 separate cell preparations; **p < 0.02 Rol v Cont) and prevented the ethanol induced increase in α-SMA expression in PSCs (n = 3 separate cell preparations; ††p < 0.01 E50 v E50+Rol).

Figure 13 Effect of retinol on ethanol induced extracellular matrix protein synthesis in pancreatic stellate cells (PSCs). (A) Representative western blot showing collagen I, fibronectin, and laminin expression in PSCs incubated with culture medium alone (Cont) or with 50 mM ethanol (E50) in the presence or absence of retinol (Rol) for five days. (B) Densitometry analysis of western blots showing a significant increase in collagen I, fibronectin, and laminin expression in PSCs treated with ethanol compared with controls (n = 3 separate cell preparations; **p < 0.005 E50 v Cont). Rol significantly decreased basal collagen I, fibronectin, and laminin levels (n = 3 separate cell preparations; †p < 0.05 Rol v Cont) and prevented the ethanol induced increase in extracellular matrix protein expression (n = 3 separate cell preparations; ††p < 0.05 E50 v E50+Rol).
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REFERENCES


Figure 14 Effect of retinol on freshly isolated pancreatic stellate cell (PSC) activation. Representative western blots and densitometry analysis showing a significant decrease in expression of (A) α-smooth muscle actin (α-SMA) and (B) fibronectin in freshly isolated PSCs treated with retinol (Rol) for five days compared with controls (Cont) (n = 4 separate cell preparations; ***p < 0.001).


