Dysregulation of elastin expression by fibroblasts in pulmonary emphysema: role of cellular retinoic acid binding protein 2

L Plantier, C Rochette-Egly, D Goven, A Boutten, M Bonay, G Lesèche, M Fournier, B Crestani, J Boczkowski

ABSTRACT

Background: All-trans retinoic acid (ATRA) stimulates elastin synthesis by lung fibroblasts and induces alveolar regeneration in animal models of pulmonary emphysema. However, ATRA treatment has had disappointing results in human emphysema. It was hypothesised that a defect in the ATRA signalling pathway contributes to the defect of alveolar repair in the human emphysematous lung.

Methods: Fibroblasts were cultured from the lung of 10 control subjects and eight patients with emphysema. Elastin and retinoic acid receptor (RAR)-β mRNA were measured in those cells in the presence of incremental concentrations of ATRA. RXRs, retinoic X receptors (RXRs) and cellular retinoic acid binding protein (CRABP) 1 and 2 mRNAs were measured as well as CRABP2 protein content. The effect of CRABP2 silencing on elastin and RAR-β expression in response to ATRA was measured in MRC5 lung fibroblasts.

Results: ATRA at 10⁻⁸ M and 10⁻⁹ M increased median elastin mRNA expression by 182% and 126% in control but not in emphysema fibroblasts. RAR-β mRNA expression was induced by ATRA in control as well as emphysema fibroblasts. RXRs and CRABP1 mRNAs were similarly expressed in control and emphysema fibroblasts while CRABP2 mRNA and protein were lower in emphysema fibroblasts. CRABP2 silencing abrogated the induction of elastin but not RAR-β expression by ATRA in MRC5 fibroblasts.

Conclusion: Pulmonary emphysema fibroblasts fail to express elastin under ATRA stimulation. CRABP2, which is necessary for elastin induction by ATRA in MRC-5 cells, is expressed at low levels in emphysema fibroblasts. This alteration in the retinoic acid signalling pathway in lung fibroblasts may contribute to the defect of alveolar repair in human pulmonary emphysema. These results are the first demonstration of the involvement of CRABP2 in elastin expression.

Pulmonary emphysema is a chronic degenerative lung disease characterised by an imbalance between alveolar destruction and repair which results in the progressive destruction of pulmonary alveoli and chronic respiratory failure. Lung fibroblasts and myofibroblasts play a major role in the course of pulmonary repair processes, notably through the secretion of elastin, an essential component of the pulmonary extracellular matrix.

Signalling by retinoic acid, the main active metabolite of vitamin A, is of particular importance for the development, maintenance and repair of pulmonary alveoli, as assessed by the following arguments: firstly, elevation of retinoic acid levels in the lung is a stimulus for the alveologenesis phase of lung development; secondly, all-trans retinoic acid (ATRA) induces the expression of elastin in lung fibroblasts; thirdly, vitamin A deficiency leads to an emphysema-like phenotype in the lung of adult rats; finally, the systemic administration of ATRA has been reported to abrogate elastase induced emphysema in adult rats and mice.

Reticoid acid exerts its effects by binding two families of nuclear receptors, the retinoic acid receptors (RAR-α, β and γ) and the retinoid X receptors (RXR-α, β and γ), which translocate to the nucleus on binding where they act as transcription factors. The binding of retinoic acid to RARs and the transcriptional activity of RXRs are greatly enhanced by a 15 kDa cytosolic protein, cellular retinoic acid binding protein 2 (CRABP2).

In light of those elements, we hypothesised that an alteration in the retinoic acid signalling pathway might contribute to the defect of alveolar repair that is observed in human pulmonary emphysema. To explore this hypothesis, we focused on elastin production by lung fibroblasts. We first determined whether ATRA induced elastin and RAR-β mRNA expression in lung fibroblasts cultured ex vivo from human control and emphysematous lung samples. Then, expression of RAR-α, RAR-β, RAR-γ, RXR-α, RXR-β, RXR-γ, CRABP1 and CRABP2 was determined in those cells. As we found a selective reduction in CRABP2 expression in fibroblasts from emphysematous patients, we determined whether suppression of CRABP2 expression in lung fibroblasts using a siRNA strategy abolished the induction of elastin expression by retinoic acid.

MATERIALS AND METHODS

Lung samples

The study was approved by the ethics committee of Paris-Bichat University Hospital, Paris, France. Patients gave informed consent.

Emphysema patients

Fibroblasts were cultured from lung samples from eight patients with severe pulmonary emphysema undergoing lung volume reduction surgery (n = 5) or lung transplantation (n = 5). Median age of the patients was 58 years (interquartile range (IQR) 55, 58.5). All patients were smokers or ex-smokers (53 pack-years, IQR 30, 58) and had normal plasma α₁ antitrypsin levels. Emphysema was diagnosed in
the presence of an obstructive ventilatory disorder and over-distension on lung function tests associated with characteristic chest CT and histological findings, and the absence of any associated lung disease was verified. The median total lung capacity of patients with emphysema was 127% predicted (IQR 122, 129) and median forced expiratory volume in 1 s 28% predicted (IQR 18, 35).

Control patients
Fibroblasts were cultured from lung samples from 10 patients undergoing lung surgery for cancer. The age of the control subjects (68 years, IQR 65, 71) was not different from that of patients with emphysema (p = 0.13). Lung samples were taken from an uninvolved segment, and the absence of emphysema was verified microscopically. Five patients were active or past smokers (36 pack-years, IQR 30, 40) and five were never-smokers. The median total lung capacity of the control patients was 98% predicted (IQR 86, 114) and their median forced expiratory volume in 1 s was 90% predicted (IQR 75, 107).

Isolation of pulmonary fibroblasts
Pulmonary fibroblasts were cultured from lung explants, as previously described.12 Fibroblasts were cultured with DMEM culture medium (Gibco/Invitrogen, Cergy-Pontoise, France) with 10% fetal calf serum (Fetalclone 2; Hyclone, Logan, Utah, USA), 100 UI/ml penicillin G, 100 μg/ml streptomycin sulphate and 0.25 μg/ml amphotericin B (Gibco/Invitrogen). Cells were maintained at 37°C with 5% CO₂ and were used at passage 5. To confirm the fibroblastic nature of cultured cells, all cell cultures were evaluated immunocytochemically at passage 5. All cells stained positive with anti-vimentin, anti-desmin and anti-prolyl-4-hydroxylase antibodies (DakoCytomation, Trappes, France). Three of the 10 control cultures and all emphysema cultures were stained positive with anti-vimentin, anti-desmin and anti-prolyl-4-hydroxylase antibodies (DakoCytomation, Trappes, France). Three of the 10 control cultures and all emphysema cultures contained 5–10% of smooth muscle actin positive cells. Staining with antibodies directed against smooth muscle myosin heavy chain-1, pancytokeratin and CD31 was always negative.12

Modulation of elastin and RAR-β mRNA expression by retinoic acid in pulmonary fibroblasts
Pulmonary fibroblasts (100 000/well) were seeded in 6 well plates and cultured to 80% confluence. They were then incubated with incremental concentrations of ATRA (10⁻⁶ M to 10⁻¹⁰ M) for 72 h. ATRA was dissolved in dimethylsulfoxide (DMSO), and cells cultured with 1% DMSO were used as controls. Total RNA was extracted with the Nucleospin extraction system (Macherey-Nagel, Hoerdt, France) and reverse transcribed into complementary DNAs (cDNAs) with MMLV retro transcrptase (Invitrogen) according to the manufacturer’s instructions. Quantitative real time PCR using a SybrGreen fluorochrome (Sigma, St Quentin-Fallavier, France) was performed with a Mx3000P thermocycler (Stratagene, La Jolla, California, USA) to quantify elastin and RAR-β cDNAs as well as ubiquitin C (UBC) cDNA as an endogenous control.13 cDNA copy numbers were expressed relative to a standard prepared from pooled lung fibroblast cDNAs that were used for all experiments. Amplification specificity was verified by agarose gel electrophoresis and melting curves.

<table>
<thead>
<tr>
<th>Gene</th>
<th>Forward primer</th>
<th>Reverse primer</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ubiquitin C</td>
<td>CACTTGTCCTGGCTGCTGA</td>
<td>TTTTTGGGAAAGGCAAACAGT</td>
</tr>
<tr>
<td>Elastin</td>
<td>GAGCTTTTGCTGGAAATCCCA</td>
<td>GGCAGTCTTCTGGTGCTAG</td>
</tr>
<tr>
<td>CRABP1</td>
<td>AGGCGCTACGCGCACTTT</td>
<td>AATTTCGAGCACTGCTGGAAG</td>
</tr>
<tr>
<td>CRABP2</td>
<td>CAAACAGGAGGAGCAACTTTCCTAC</td>
<td>CTCTTCCTCATACTTGAAGTT</td>
</tr>
<tr>
<td>RAR-α</td>
<td>CCTCTGAGCAAGTCTGACACT</td>
<td>GTCCAGATCAGCAGCATCA</td>
</tr>
<tr>
<td>RAR-β</td>
<td>TTAAGTACGAGGATTTGTCAAAG</td>
<td>GGTTAAGGCGCTGAGAAAGT</td>
</tr>
<tr>
<td>RAR-γ</td>
<td>GAGGCTGTTGGTTGAGCTCTAAAT</td>
<td>TCTCTAGTGGTCGTTTGTCTCA</td>
</tr>
<tr>
<td>RXR-α</td>
<td>AGGGCTGAGGGGAAGGTT</td>
<td>AGGAAGGTTCAAAGGGTGTT</td>
</tr>
<tr>
<td>RXR-β</td>
<td>CGTTCATCGGCTTAAGT</td>
<td>TCTCTAGGAAAGGATCGCAT</td>
</tr>
<tr>
<td>RXR-γ</td>
<td>GTGCAACAGTGTACAGGTCACCA</td>
<td>CGGGAGGATGTACAGGTCACCAATCCCG</td>
</tr>
</tbody>
</table>

CRABP, cellular retinoic acid binding protein; RAR, retinoic acid receptor; RXR, retinoic X receptor.

Determination of the intracellular content of fibroblasts in RAR, RXR, CRABP1 and CRABP2 mRNA in the absence of stimulation
RAR-α, RAR-β, RAR-γ, RXR-α, RXR-β, RXR-γ, CRABP1 and CRABP2 mRNAs were quantified by reverse transcription-PCR in unstimulated control and emphysema fibroblasts, as described above. Primers sequences are listed in table 1.

Determination of the intracellular content of CRABP2 protein in fibroblasts
Fibroblasts at passage 5 were cultured to confluence in 75 cm² flasks (Corning, Schiphol-Rijk, The Netherlands). Cells were rinsed twice with phosphate buffered saline (Gibco/Invitrogen) and proteins were extracted with Cytobuster Protein Extraction Reagent (Novagen, Madison, USA) according to the manufacturer’s protocol. Cellular proteins (25 μg) were separated by sodium dodecyl sulphate–polyacrylamide gel electrophoresis, transferred onto a PVDF membrane (Biorad, Marne-la-coquette, France) and incubated with a mouse monoclonal primary antibody binding CRABP2 (SCRA-3B5)14 diluted 1/1000 for 1 h at room temperature. A mouse monoclonal antibody binding β-actin (A-5316; Sigma) was used as an endogenous control. Detection was performed with peroxidase conjugated sheep anti-mouse (NA 951) and donkey anti-rabbit (NA954) antibodies and ECL reagent (Amersham, Little Chalfont, UK). The Bio-Vision system (Fisher Bioblock Scientific, Illkirch, France) was used for densitometric quantification of protein bands.
with either 20 nM of anti-CRABP2 siRNA (sense: 5'-GCG CAC CAC AGA GAT TAA CTT CAA G-3') or 20 nM scramble control RNA (sense 5'-GCG CAC CGA GAA TTA TTC ACA CAA G-3'; Stealth Technology, Invitrogen) using Lipofectamine 2000 (Invitrogen), according to the manufacturer's protocol, and were incubated with 10% fetal bovine serum. Fresh medium, fetal bovine serum, RNAs and Lipofectamine 2000 were added every 3 days until cellular subconfluence was obtained. Transfected cells were then treated with incremental concentrations of ATRA, and elastin, RAR-β and UBC mRNAs were quantified as described above. The CRABP2 mRNA and protein content in siRNA and control RNA treated cells was determined by RT-PCR and western blotting, as described above.

**Statistical analysis**

Data are expressed as median and interquartile range (IQR), expressed as the values for the 25th and 75th centiles.
0.58, 1.74) was lower than that of control fibroblasts (0.915, IQR 0.14, 0.79, p = 0.05) (fig 2B, C).

ATRA failed to induce elastin mRNA expression in MRC-5 fibroblasts treated with a siRNA targeting CRABP2 mRNA while RAR-β induction was preserved

To determine whether the low expression of CRABP2 could explain the absence of elastin induction by ATRA, we tested whether CRABP2 gene silencing affected elastin and RAR-β induction by ATRA in the MRC-5 pulmonary fibroblast cell line.

Median CRABP2 mRNA was reduced by 97.6% in cells treated with the anti-CRABP2 siRNA compared with cells treated with the control RNA (p = 0.045, fig 3A). Median content in CRABP2 protein of siRNA treated cells was reduced by 95.5% compared with that of control RNA treated cells (p = 0.045, fig 3B). The anti-CRABP2 siRNA had no additional cytotoxicity compared with the control RNA.

In MRC-5 fibroblasts treated with the control RNA, exposure to ATRA at all concentrations induced an increase in the median elastin/UBC mRNA ratio which reached a maximum of 511% at an ATRA concentration of $10^{-7}$ M. Induction of elastin gene expression by ATRA was completely abrogated in cells treated with the anti-CRABP2 siRNA (fig 3C). In the presence of ATRA at any concentration, the elastin/UBC mRNA ratio was superior in MRC-5 cells treated with the control RNA compared with cells treated with the anti-CRABP2 siRNA.
Airway biology

In contrast, CRABP2 silencing did not abrogate the induction of RAR-β mRNA by ATRA in those cells. In siRNA as well as in control RNA treated cells, ATRA at a concentration of 10^{-7} M and above induced an elevation in the RAR-β mRNA content of cells (fig SD).

DISCUSSION

The main results of this study are as follows: (1) retinoic acid induced elastin mRNA expression in control but not in emphysema lung fibroblasts while it induced RAR-β mRNA in both groups, (2) this discrepancy was associated with decreased levels of CRABP2 mRNA and protein in emphysema fibroblasts and (3) silencing of CRABP2 expression in the MRC5 human lung fibroblast cell line reproduced the phenotype of emphysema fibroblasts, as it resulted in the loss of elastin mRNA but not RAR-β induction by retinoic acid. These results are the first demonstration of the involvement of CRABP2 in elastin expression and strongly suggest that the lack of elastin induction by retinoic acid in emphysema fibroblasts is related to the low expression of CRABP2 in those cells.

In our experiments, ex vivo culturing of lung fibroblasts implicated selection bias towards cells able to proliferate outside of their normal environment. Moreover, whether the reduction in CRABP2 expression by cultured emphysema fibroblasts was related to a global reduction of its expression in lung fibroblasts in this disease or reflected the loss of a particular subgroup of CRABP2 expressing lung fibroblasts cannot be determined from our study. Interestingly, skin fibroblasts express different levels of CRABP2 depending on the site of their isolation, as subcutaneous fat fibroblasts express much lower levels of CRABP2 than their dermis derived counterparts. Regulation of CRABP2 expression is incompletely understood but seems to involve the RARs. However, alteration of this pathway seems unlikely in the present study as expression of RAR mRNAs was similar in fibroblasts from emphysematous and control patients. The AP-2 transcription factor and protein kinase C have also been involved in the regulation of CRABP2 expression, but their role in pulmonary emphysema is unknown. Whatever the mechanism of the reduced CRABP2 expression in fibroblasts from patients with emphysema, the present results are in line with previous reports showing that the phenotype of lung fibroblasts is deeply altered in the emphysematous lung, as those cells have been shown to express markers of cellular senescence, to have a reduced proliferation rate and to secrete low amounts of hepatocyte growth factor, a key mitogen for alveolar epithelial cells, while they fail to link the endothelial and epithelial compartments of the lung through direct intercellular contacts, as is observed in the normal lung.

In our experiments, ATRA increased the elastin mRNA content in fibroblasts in a non-dose dependent manner as the maximal effect was observed at ATRA concentrations of 10^{-6} M in primary cells and 10^{-7} M in MRC-5 cells. We do not know the reason for this lack of a dose dependent effect. However, a similar phenomenon was described in human primary skin fibroblasts in which ATRA potentiated 8-bromo-cAMP induced hepatocyte growth factor. In those cells, the maximal effect of ATRA was obtained at a concentration of 10^{-7} M.

It may be argued that the different RAR-β mRNA expression that was observed between control and emphysema fibroblasts in the presence of 10^{-7} M to 10^{-6} M ATRA may have contributed to the different response of those cells to ATRA with regard to elastin expression. However, this hypothesis seems unlikely as RAR-β has been consistently shown to inhibit alveolar septation, of which elastin synthesis is an essential step. Moreover, in addition to the silencing experiments performed in MRC-5 cells, demonstration that CRABP2 deficiency accounted for the reduced elastin synthesis in emphysema fibroblasts would have been strengthened by the transfection of those cells with an expression vector that increases CRABP2 to the level of controls. This experiment could not be performed in this study as primary cells could not survive in the presence of the transfection reagents.

Elastic fibres, an essential component of the lung extracellular matrix as they provide the elasticity needed for cyclic ventilation, are extremely stable and undergo very limited renewal and remodelling in healthy individuals. Their degradation is a hallmark of human emphysema as well as of numerous animal models of this disease. While elastic fibre repair has not been demonstrated to date in the human emphysematous lung, data obtained from animal studies point towards a potentially essential role of the lack of elastic fibre regeneration and repair in the constitution of emphysema lesions. Treatment of mice with beta-aminopropionitrile, a chemical inhibitor of elastin cross linking, leads to aggravated emphysema after elastase instillation, while mice expressing low levels of elastin develop more severe emphysema than wild-type mice after exposure to cigarette smoke. From a pathophysiological point of view, demonstration that fibroblasts cultured from human emphysematous lung have an impaired capacity to express elastin mRNA under retinoic acid stimulation may help explain the defect in alveolar maintenance and repair in this disease. It must be noted that our study was limited to the exploration of elastin mRNA and that differences in the post-transcriptional regulation of elastin production may have been overlooked.

Given the beneficial effect of ATRA in animal models of emphysema, the therapeutic potential of this molecule was evaluated in human patients with emphysema. Disappointingly, a 3 month course of ATRA could not reverse the functional and morphological changes associated with emphysema, despite obtaining high plasma ATRA levels. Our results may provide an explanation for the lack of a therapeutic effect of ATRA in humans with emphysema. CRABP2 is associated with loss of elastin gene expression and the regeneration and repair in the constitution of emphysema have not been demonstrated to date in the human emphysematous lung, data obtained from animal studies point towards a potential role of the lack of elastic fibre repair in the constitution of emphysema lesions. Treatment of mice with beta-aminopropionitrile, a chemical inhibitor of elastin cross linking, leads to aggravated emphysema after elastase instillation, while mice expressing low levels of elastin develop more severe emphysema than wild-type mice after exposure to cigarette smoke. Thus from a pathophysiological point of view, demonstration that fibroblasts cultured from human emphysematous lung have an impaired capacity to express elastin mRNA under retinoic acid stimulation may help explain the defect in alveolar maintenance and repair in this disease. It must be noted that our study was limited to the exploration of elastin mRNA and that differences in the post-transcriptional regulation of elastin production may have been overlooked.

Overall, our results indicate that low expression of CRABP2 is associated with loss of elastin gene expression by retinoic acid in fibroblasts cultured from human emphysematous lung, which may participate in the lack of alveolar repair and may help explain the lack of a therapeutic effect of ATRA in this disease.

Acknowledgements: We thank Dr Nadira Houhou (Service de Virologie, Hôpital Bichat) for kindly providing the MRC-5 cells used in this study.

Funding: LP was supported by a research fellowship from INSERM (Poste d’Accueil) and a grant from the Société de Pneumologie de Langue Française (SPLF). JB was supported by INSERM and Assistance Publique-Hôpitaux de Paris (Contrat d’Interface). Part of this project was supported by a grant from the Chancellerie des Universités (Legs Poix).

Competing interests: None.

Ethics approval: The study was approved by the ethics committee of Paris-Bichat University Hospital, Paris, France.
REFERENCES


Scadding-Morrison Davies Joint Fellowship in Respiratory Medicine 2009

This fellowship is available to support visits to medical centres in the United Kingdom or abroad for the purpose of undertaking studies related to respiratory medicine. Applications are invited from medical graduates practising in the United Kingdom, including consultants and irrespective of the number of years in that grade. There is no application form but a curriculum vitae should be submitted together with a detailed account of the duration and nature of the work and the centres to be visited, confirming that these have agreed to provide the facilities required. Please state the sum of money needed for travel and subsistence. A sum of up to £20 000 can be awarded to the successful candidate, or the sum may be divided to support two or more applications. Applications should be sent to Dr I A Campbell, Secretary to the Scadding-Morrison Davies Fellowship, Llandough Hospital, Penarth, Vale of Glamorgan CF64 2XX by 31 January 2009.
Dysregulation of elastin expression by fibroblasts in pulmonary emphysema: role of cellular retinoic acid binding protein 2

L Plantier, C Rochette-Egly, D Goven, A Boutten, M Bonay, G Lesèche, M Fournier, B Crestani and J Boczkowski

Thorax 2008 63: 1012-1017 originally published online July 11, 2008
doi: 10.1136/thx.2007.093302

Updated information and services can be found at:
http://thorax.bmj.com/content/63/11/1012

These include:

References
This article cites 30 articles, 8 of which you can access for free at:
http://thorax.bmj.com/content/63/11/1012#BIBL

Email alerting service
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Topic Collections
Articles on similar topics can be found in the following collections
- Pulmonary emphysema (25)
- Chemotherapy (183)

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/