Reduction of bleomycin induced lung fibrosis by candesartan cilexetil, an angiotensin II type 1 receptor antagonist

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Background: Signalling of angiotensin II via angiotensin II type 1 receptor (AT1) promotes cardiac and renal fibrosis, but its role in lung fibrosis is little understood. Using a rat bleomycin (BLM) induced model of pulmonary fibrosis, we examined the expression of AT1 in the lung and the effect of an AT1 antagonist on pulmonary fibrosis.

Methods: Adult male Sprague-Dawley rats were given 0.3 mg/kg BLM intratracheally. Two days earlier they had received 10 mg/kg/day of the AT1 antagonist candesartan cilexetil mixed in the drinking water. AT1 expression in the lungs was examined by immunohistochemistry and immunoblot methods. The effect of the AT1 antagonist on pulmonary fibrosis was studied by analysis of bronchoalveolar lavage (BAL) fluid, histopathology, and hydroxyproline assay.

Results: Immunohistochemical studies showed overexpression of AT1 in inflammatory immune cells, alveolar type II cells, and fibroblasts. A quantitative assay for AT1 showed that AT1 expression was significantly upregulated in cells from BAL fluid after day 3 and in the lung homogenates after day 21. Candesartan cilexetil significantly inhibited the increase in total protein and albumin, as well as the increase in total cells and neutrophils in BAL fluid. On day 21 candesartan cilexetil also ameliorated morphological changes and an increased amount of hydroxyproline in lung homogenates. In addition, BLM increased the expression of transforming growth factor (TGF)-β in BAL fluid on day 7; this increase was significantly reduced by candesartan cilexetil.

Conclusion: AT1 expression is upregulated in fibrotic lungs. Angiotensin II promotes lung fibrosis via AT1 and, presumably, in part via TGF-β1.

Angiotensin II, which is mainly generated by angiotensin converting enzyme (ACE) and chymase, is a peptide that plays a crucial role in regulating blood pressure and sodium homeostasis via specific receptors including angiotensin II type 1 receptor (AT1). Candesartan cilexetil is an AT1 selective antagonist and is widely used for the treatment of hypertension. Recent studies have shown that angiotensin II is also closely associated with tissue injury and fibrogenesis in circulatory organs. The administration of candesartan cilexetil significantly attenuates injury and fibrosis in injured heart models in vivo.

The process of fibrosis is also important in the lung since its progression may result in respiratory failure and a poor prognosis in patients with various types of interstitial lung disorders. The progression of fibrosis is closely related to a complicated network consisting of many cytokines, chemical mediators, growth factors, and biophysical peptides derived from inflammatory immune cells, endothelial cells, and alveolar type II cells. As a member of this network, a mediator in the lung is a key multi-functional cytokine in many fibrotic diseases. Angiotensin II is closely associated with TGF-β1 production in cardiac and renal fibrosis, which suggests that angiotensin II leads to fibrosis in these models through acceleration of TGF-β1 synthesis via AT1. TGF-β1 also acts as a principal fibrogenic mediator in the lung. The aims of this study were (1) to clarify whether the expression of AT1 in the lung increases with the progression of fibrosis and, if so, whether candesartan cilexetil attenuates the fibrosis and (2) to determine how it is associated with the cytokine network including TGF-β1.

Using a rat model of pulmonary fibrosis generated by BLM with/without candesartan cilexetil, we examined the relationship between morphological changes, hydroxyproline content, number of inflammatory immune cells in BAL fluid, and the expression of TGF-β1 and AT1 in the lung tissues.
METHODS

Animals and preparation of BLM induced pulmonary fibrosis

Adult male Sprague-Dawley rats weighing 200–220 g (Sankyo Lab, Tokyo, Japan) were anaesthetised and 0.3 mg/kg BLM (Nippon Kayaku, Tokyo, Japan) in 0.5 ml sterile saline or 0.5 ml of sterile saline (vehicle control) were administered intratracheally. Two days before the administration some rats received 10 mg/kg/day candesartan cilexetil (Takeda Chemical Industries, Tokyo, Japan) dissolved in the drinking water. Rats were divided into four groups as follows: BLM alone, BLM + candesartan, saline alone, and saline + candesartan. The rats were killed on days 0, 1, 3, 7, 14, or 21 (n = 4–6 for each group and time point). The lungs were lavaged five times with 7 ml saline and the lavage fluids were centrifuged at 250g for 10 minutes to separate the cells and the supernatants. A total cell count was performed with a haemocytometer and differential cell counts were performed on cytospin preparations stained with Wright-Giemsa. Two hundred cells were counted for determination of the differential cell count. The supernatants, resuspended pellets, and lavaged lung tissues were stored at −30°C.

Histopathology

Rats from each group were killed on days 0, 3, 7, 14, 21 and 10% buffered formalin was intratracheally instilled into the lungs. Lung tissue was embedded in paraffin and lung sections were stained with haematoxylin and eosin for examination by light microscopy.

Immunohistochemistry

Immunohistochemical analysis of AT1 using specific antibody was performed by the avidin-biotin complex (ABC) technique. Paraffin embedded lung sections were dewaxed with xylene, hydrated, and heated at 121°C for 15 minutes in a steam steriliser in 10 mM citrate buffer for antigen retrieval. Endogenous peroxidase activity was quenched by the addition of 0.3% (vol/vol) hydrogen peroxide in methanol for 30 minutes. The slides were first blocked with 1% goat serum in phosphate buffered serum (PBS) for 30 minutes. Polyclonal rabbit anti-human AT1 antibody (Santa Cruz Biotech, Santa Cruz, CA, USA) was used for the immunostaining. For negative controls, non-specific rabbit IgG was used. The antibodies were used at a dilution of 1:200 (AT1) or 1:1000 (anti-AT1) in PBS. The sections were incubated for 60 minutes with the antibody, then washed with PBS over a period of 15 minutes. Incubation with primary antibody (anti-AT1 polyclonal antibody diluted 1:1000) for 1 hour. The membrane was then subjected to a chemiluminescence detection system (Amersham Pharmacia Biotech, Buckinghamshire, UK) and exposed to x ray film. Densitometric analysis was performed using the NIH image program (http://rsb.info.nih.gov/nih-image/).

Hydroxyproline assay

Frozen lung tissues were homogenised by a polytron tissue homogeniser in saline containing 0.1 M phenylmethylsulfonyl fluoride. The homogenised sample (n = 6) was hydrolysed in 6 N HCl and the hydroxyproline concentration was determined according to the method of Kivirikko et al.

Protein and albumin assay

The protein concentration in BAL fluid was measured using the bicinchoninic acid (BCA) method (Pierce Chemical, Rockford, IL, USA) (n = 4–5). The albumin concentration in BAL fluid was measured using brom cresol green (BCG) solution (Sigma Co, St Louis, MO, USA) (n = 4 or 5).

Measurement of TGF-β1 in BAL fluid

BAL fluid samples underwent acidification to convert the latent TGF-β1 to an active form. 1 N HCl was used followed by neutralisation with 1 N NaOH. Total (latent and active form) TGF-β1 levels were measured using a quantitative sandwich enzyme immunoassay kit (Promega, Madison, WI, USA). All samples were assayed in duplicate (n = 4–6).

Statistical analysis

The results are expressed as mean (SD) with mean difference and 95% confidence intervals (CI) between groups and from day 0. Statistical analysis was carried out by one way or two way analysis of variance (ANOVA) followed, when differences were significant, by appropriate post hoc tests using the Tukey-Kramer test. p values of <0.05 were considered to be statistically significant.

RESULTS

Histopathology

After intratracheal administration of BLM, the histopathological changes in the lung were evaluated on days 0, 3, 7, 14, and 21. On day 3 infiltration of neutrophils and macrophages into the alveolar walls and spaces was seen, together with thickening of the alveolar interstitium. These changes were most prominent on day 7. On days 14 and 21 the number of inflammatory cells was small while fibroblasts were numerous, and a mild collagenous deposit with interstitial thickening was observed in the alveolar walls (fig 1C). Daily oral administration of candesartan cilexetil resulted in moderate amelioration of inflammatory cell infiltration by day 7, together with a marked reduction in interstitial thickening by BLM on day 21 (fig 1D).

Immunohistochemistry of AT1

The expression of AT1 in the lungs of the four groups was immunohistochemically analysed on day 21. In the lungs of rats in the saline alone group specific immunoreactivity was observed in alveolar macrophages, alveolar type II cells, endothelial cells, and bronchial epithelial cells (fig 2A). AT1

night. For dot blot analysis, 1 µg of cells from the BAL fluid was mounted on a nitrocellulose membrane and air dried. Both membranes were blocked by PBS containing 2% skimmed milk and 0.1% Tween-20 for 1 hour. The membranes were washed in 0.1% Tween-20 in PBS twice for 5 minutes each. Incubation with primary antibody (anti-AT1 polyclonal antibody diluted 1:1000 by Tween-20-PBS) was performed for 1 hour at room temperature. After washing the membranes were incubated with secondary antibody conjugated with horseradish peroxidase (anti-rabbit IgG 1:1000 dilution) for 1 hour. The membrane was then subjected to a chemiluminescence detection system (Amersham Pharmacia Biotech, Buckinghamshire, UK) and exposed to x ray film. Densitometric analysis was performed using the NIH image program (http://rsb.info.nih.gov/nih-image/).
positive capillary endothelial cells, interstitial macrophages, and fibroblasts were also observed in fibrotic lesions in lungs from the BLM alone group (fig 2B). The intensity and extent of immunoreactivity did not differ between the BLM alone group and the BLM + candesartan group (data not shown). Negative controls were unstained in all specimens (fig 2C and D).

Figure 1  Histopathological findings on day 21 after administration of bleomycin (BLM). The groups given saline alone (A) and saline + candesartan (B) had a normal alveolar structure. The group given BLM alone (C) had marked alveolar thickening and diffuse fibrotic changes with cellular infiltration, while the group given BLM + candesartan (D) had reduced alveolitis and fibrotic changes. Bar = 50 μm.

Figure 2  Immunohistochemical study of angiotensin II type 1 receptor (AT1) in (A, C) the saline alone group on day 0, and (B, D) the BLM alone group on day 21. Immunostaining was performed using (A, B) polyclonal rabbit anti-human AT1 antibody and (C, D) non-specific rabbit IgG. Normal lung section showed specific immunoreactivity in alveolar macrophages, alveolar type II cells, endothelial cells, and bronchial epithelial cells (A). Injured lung sections showed labelling of infiltrated capillary endothelial cells and fibroblasts in the thickened alveolar interstitium (B). The negative controls were unstained in all specimens (C, D). Bar = 25 μm.
Comparative quantification of AT1

The level of AT1 expression in the BLM alone group was measured in lung homogenates and BAL fluid cells by Western blot and dot blot analysis, respectively. In lung homogenates on days 0, 3, 7, 14, and 21, anti-AT1 antibody recognised the protein with an apparent molecular mass of 46 kDa, which is identical to AT1 reported by Harrison-Bernard et al.24 The band on the left in each panel shows the control on day 0. BLM increased the expression of AT1 on days 14 and 21 compared with day 0 (n = 5). (B) Dot blot analysis of AT1 expression in BAL fluid cells. BLM administration resulted in an increase in the expression of AT1 which was maximal on day 3 (n = 5).

Hydroxyproline content of lung tissue

A significant increase in the concentration of hydroxyproline in lung tissue was observed on days 14 and 21 in the BLM alone group but no such increase was seen in the saline alone group. The concentrations for all four groups on day 21 are shown in fig 5. The levels were found to be significantly different in the four groups (p<0.01, one way ANOVA). In the BLM alone group the increase in hydroxyproline content (4.83 (0.97) mg/lung) was almost double that in the saline alone group (2.35 (0.71) mg/lung, mean difference 2.49, 95% CI 1.53 to 3.45, p<0.05) and the saline + candesartan group (2.46 (0.41) mg/lung, mean difference 2.38, 95% CI 1.36 to 3.39, p<0.05). This increase was significantly blocked by candesartan cilexetil (BLM + candesartan group: 3.28 (0.57) mg/lung, mean difference 1.56, 95% CI 0.59 to 2.53, p<0.05).

BAL fluid cells

Recovery rates of BAL fluid ranged from 83% to 100% and were not significantly different between the four groups. Neither the saline alone group nor the saline + candesartan group showed any significant difference in total cell numbers or differential cell counts during the 21 days. A significant difference in total cell number, neutrophils, and alveolar macrophages was seen between the BLM alone group and the BLM + candesartan group (p<0.01, p<0.01, p<0.01, respectively, two-way ANOVA; fig 6). A significant difference was not seen in the differential cell counts or the total cell numbers when the saline alone group was compared with the other groups.

Figure 3  Typical immunoblot profiles of angiotensin II type 1 receptor (AT1) in rats following intratracheal administration of bleomycin (BLM), n = 5. (A) Western blot analysis of AT1 expression in lung homogenates at days 0, 3, 7, 14, and 21 after BLM. Anti-AT1 polyclonal antibody recognised the protein with an apparent molecular mass of 46 kDa, which is identical to AT1 reported by Harrison-Bernard et al.24 The band on the left in each panel shows the control on day 0. BLM increased the expression of AT1 on days 14 and 21 compared with day 0 (n = 5). (B) Dot blot analysis of AT1 expression in BAL fluid cells. BLM administration resulted in an increase in the expression of AT1 which was maximal on day 3 (n = 5).

Figure 5  Comparison of hydroxyproline concentration in lung homogenates on day 21 in the four groups (n = 6): saline alone, saline + candesartan cilexetil, bleomycin (BLM) alone, and BLM + candesartan cilexetil. The concentrations were found to be significantly different in the four groups (p<0.01, one way ANOVA). The BLM alone group had a significantly increased hydroxyproline concentration on day 21. Candesartan cilexetil significantly reduced the increase in hydroxyproline content caused by the administration of BLM. Data are shown as mean (SD). *p<0.05 v saline alone group; †p<0.05 v BLM + candesartan group.

Figure 4  Change in density of angiotensin II type 1 receptor (AT1) expression after administration of bleomycin (BLM) in rats (n = 5). The densitometric intensity of each immunoreactive band or spot exposed to x ray film was analysed using the NIH imaging program. Significant differences between time points were seen in both lung homogenates and BAL fluid cells (p<0.01, p<0.01, respectively, one way ANOVA). In lung homogenates (A) the increase in AT1 expression began on day 14 and reached a peak on day 21. In BAL fluid cells (B) AT1 expression had increased rapidly by day 3 and then declined gradually until day 21. Data were corrected to the mean value for day 0 and are shown as mean (SD). *p<0.05 compared with day 0.
also seen between time points from day 0 to day 21. A post hoc test showed that BLM induced a significant increase in mean (SD) total cell counts on days 3, 7, and 14 (day 3: 462 (126) × 10⁶/ml, mean difference 359 × 10⁶, 95% CI 224 × 10⁶ to 495 × 10⁶, p < 0.05; day 7: 589 (127) × 10⁶/ml, mean difference 486 × 10⁶, 95% CI 350 × 10⁶ to 623 × 10⁶, p < 0.05; day 14: 396 (92.9) × 10⁶/ml, mean difference 294 × 10⁶, 95% CI 191 × 10⁶ to 397 × 10⁶, p < 0.05) compared with baseline levels at day 0 (102 (36.3) × 10⁶/ml; fig 6A). The maximal cell number was observed on day 7. Candesartan cilexetil significantly inhibited the increase on day 7 (342 (35.0) × 10⁶/ml, mean difference 247 × 10⁶, 95% CI 108 × 10⁶ to 383 × 10⁶, p < 0.05). Alveolar macrophages (C) increased from day 3 and peaked on day 14; this increase was reduced with candesartan cilexetil.

ANOVA). Total cell counts (A) and neutrophils (B) increased significantly and peaked on day 7. These increases were significantly inhibited by BLM (BLM and the effect of candesartan cilexetil; n = 4 or 5). Animals given BLM alone and those treated with BLM + candesartan showed a significant difference in the time course in total cell count, neutrophils, and alveolar macrophages (p < 0.01, p < 0.01, p < 0.01, respectively, two way ANOVA). Total cell counts (A) and neutrophils (B) increased significantly and peaked on day 7. These increases were significantly inhibited by candesartan cilexetil. Alveolar macrophages (C) increased from day 3 and peaked on day 14; this increase was reduced with candesartan cilexetil. Data were obtained from five rats and are shown as mean (SD) values. *p < 0.05 versus BLM alone; †p < 0.05 versus BLM + candesartan group; ‡p < 0.05 versus BLM + candesartan group; §p < 0.05 versus BLM + candesartan group.

Figure 6  Change in (A) total cell count, (B) neutrophils, and (C) alveolar macrophages in BAL fluid during the 21 days following administration of bleomycin (BLM) and the effect of candesartan cilexetil (n = 4 or 5). Animals given BLM alone and those treated with BLM + candesartan showed a significant difference in the time course in total cell count, neutrophils, and alveolar macrophages (p < 0.01, p < 0.01, p < 0.01, respectively, two way ANOVA). Total cell counts (A) and neutrophils (B) increased significantly and peaked on day 7. These increases were significantly inhibited by candesartan cilexetil. Alveolar macrophages (C) increased from day 3 and peaked on day 14; this increase was reduced with candesartan cilexetil. Data were obtained from five rats and are shown as mean (SD) values. *p < 0.05 versus BLM alone; †p < 0.05 versus BLM + candesartan group; ‡p < 0.05 versus BLM alone on day 0.

**TGF-β₁ levels in BAL fluid**

TGF-β₁ levels in BAL fluid supernatants were measured using ELISA kits (fig 8). There was a significant difference between the time course in TGF-β₁ levels in the BLM alone and BLM + candesartan groups (p < 0.01 and p < 0.01, respectively, two way ANOVA). TGF-β₁ concentrations in the BLM alone group were significantly increased on day 7 (129 (16.3) pg/ml) compared with day 0 (27.2 (12.6) pg/ml, mean difference 102, 95% CI 83.4 to 121, p < 0.05). Candesartan cilexetil significantly inhibited the increase on day 7 (94.9 (32.1) pg/ml, mean difference 34.5, 95% CI 1.78 to 67.2, p < 0.05).
DISCUSSION

This study has shown that the expression of AT1 is upregulated in BLM induced pulmonary fibrosis and that the AT1 antagonist candesartan cilexetil inhibits the synthesis of TGF-β1 and ameliorates the pulmonary fibrosis. These results suggest that signalling of angiotensin II via AT1 promotes BLM induced pulmonary fibrosis.

This is the first report to show the detailed localisation of AT1 in the rat lung, although the presence of AT1 in human lung has been reported. AT1 in rat lung was localised on alveolar macrophages, alveolar type II cells, vascular smooth muscle cells, endothelial cells, and fibroblasts. These cells are believed to play an important role in the process of fibrosis. The localisation of AT1 in normal rat lung, as shown in fig 2, was the same as that in human lung as described in a previous paper. AT1 expression in the lung increased markedly after intratracheal administration of BLM, which is known to generate fibrosis, suggesting that angiotensin II as a ligand of AT1 is involved in pulmonary fibrosis. The quantitative assay for AT1 showed that the time course of intensity of AT1 expression in BAL fluid cells was different from that in lung homogenates. AT1 increased as early as day 3 in BAL fluid cells, while in the lung homogenates the increase did not take place until days 14–21. This time lag may be due to the different cellular components of these samples. Neutrophils and alveolar macrophages may contribute to the increase in AT1 in BAL fluid cells while capillary endothelial cells, interstitial macrophages, and fibroblasts may be the major sources of the increase in the lung homogenates.

Lung fibroblasts are known to proliferate under the stimulation of angiotensin II via AT1. Since the fibroblast is a major cell component involved in the synthesis of collagen fibres, the increase in AT1 in this cell may be critically important for the formation of fibrosis. An increased number of these cells was histopathologically observed on day 21, and the expression of AT1 in the lung homogenates continued to increase for 21 days after BLM instillation, suggesting that AT1 mediated fibrosis occurs progressively. If chronic types of lung injury such as idiopathic pulmonary fibrosis also maintain a high level of AT1 expression, candesartan cilexetil should be administered over a long period in order to achieve a satisfactory therapeutic effect and, moreover, an agent inhibiting the gene expression of AT1 should be developed for use as a novel therapeutic strategy.

Activated neutrophils in the alveolar space release toxic radical oxygen species and various proteolytic enzymes which result in severe tissue injury. Many chemotactic factors and adhesion molecules including E-selectin and interleukin (IL)-8 are involved in the migration and accumulation of neutrophils in the bronchoalveolar space. The administration of candesartan cilexetil significantly reduced neutrophil accumulation in the bronchoalveolar space, suggesting that angiotensin II may play a crucial role in the acceleration of neutrophil chemotaxis in the BLM injured lung. Previous studies support this suggestion. Angiotensin II generates E-selectin on endothelial cells and promotes leucocyte adhesion to endothelial cells; these effects are inhibited by the AT1 antagonist losartan. The AT1 antagonist also inhibits the expression of another adhesion molecule, Mac-1, on neutrophils. Angiotensin II is itself a potent factor in promoting neutrophil migration; it induces the synthesis of chemotactic factors in endothelial cells and quinapril, an ACE inhibitor, reduces the synthesis of IL-8 in macrophages in an arteriosclerosis model. These findings and the results of the present study therefore show that the renin-angiotensin II system is involved in neutrophil accumulation in the BLM injured lung.
IL-8 is known to be not only a neutrophil chemotactic chemokine but also a pathogenic mediator of pulmonary fibrosis. Since neutrophils are a major source of IL-8, its overproduction by increased neutrophils in the BLM injured lung is likely to enhance the progression of fibrosis.

TGF-β1 is the most important factor in mediating pulmonary fibrosis. Patients with idiopathic pulmonary fibrosis and some animal models of pulmonary fibrosis have shown increased TGF-β1 production. TGF-β1 is mainly produced by alveolar macrophages. Our study showed that TGF-β1 levels in BAL fluid increased from day 3 and peaked on day 7 when the increase was significantly inhibited by treatment with an AT1 antagonist. These results suggest that angiotensin II promotes pulmonary fibrosis directly or indirectly via the acceleration of TGF-β1 synthesis. Similar results have been reported in animal models of cardiac and renal fibrosis. In vitro studies have also shown that angiotensin II increases TGF-β1 synthesis in cultured cardiac fibroblasts, vascular smooth muscle cells, bronchial smooth muscle cells, and renal mesangium cells. These reports, together with the results of our study, show that angiotensin II plays, in part, a role in accelerating the formation of fibrosis mediated by TGF-β1, which is partially inhibited by AT1 antagonists. Since the major source of TGF-β1 in the BLM induced fibrotic lungs seems to be alveolar macrophages, it is necessary to evaluate the expression of AT1 by these cells and the subsequent intracellular signal transduction pathway covering the production of TGF-β1.

Administration of BLM increased vascular permeability of the lung. Not only BLM itself but also the toxic radical oxygen species and various proteolytic enzymes cause injury to endothelial cells or other structural components of the alveolar-capillary membrane which results in increased vascular permeability. The total protein and albumin content of the BAL fluid are ordinal markers of vascular permeability. As shown in fig 7, AT1 antagonist suppressed the acceleration of vascular permeability caused by BLM. Moreover, analysis of BAL fluid showed that the proportion of albumin to total protein in the BLM + candesartan group on day 7 was significantly higher than in the BLM alone group. This suggests that candesartan cilexetil may reduce vascular permeability. The total protein and albumin content of albumin to total protein in the BLM of albumin to total protein in the BLM alone was significantly higher than in the BLM alone group. This suggests that candesartan cilexetil may reduce vascular permeability. The total protein and albumin content of albumin to total protein in the BLM alone group. This suggests that candesartan cilexetil may reduce vascular permeability.

In conclusion, our results indicate that AT1 expression is upregulated in BLM induced lung fibrosis and AT1 antagonist prevents the accumulation of inflammatory cells and development of lung fibrosis. Angiotensin II promotes lung injury and fibrosis via AT1 and, presumably, in part via TGF-β1.

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REFERENCES

LUNG ALERT

New evidence for the Dutch hypothesis?


This study, designed to evaluate risk factors for the persistence and relapse of asthma, followed children born in Dunedin, New Zealand between April 1972 and March 1973. A total of 613 subjects (51.4% of the total study population) underwent spirometry, bronchial provocation, allergy testing, and questionnaires at roughly 2 year intervals between the ages of 9 and 26. Overall, one in four had persistent or relapsing wheeze at the age of 26. Wheezing was reported at some point by 72.6% of the population. The risk of persistent/relapsing wheeze was increased by the presence of airway hyperresponsiveness and house dust mite allergy; female subjects were more likely to have persistent wheeze. The chance of relapsing wheeze was greater, the earlier its onset. Compared with the rest of the population, in male subjects with persistent wheeze the fall in the FEV$_1$/FVC ratio was, on average, 6.8% and in female subjects it was 4.7%. However, the slopes of the decline in lung function were similar for all groups, indicating that impairment in lung function existed before the first measurement at the age of 9.

Given this longitudinal decline in lung function, these findings may be considered to support the “Dutch hypothesis” that falling lung function is due to endogenous factors such as airway hyperresponsiveness leading to airway remodelling and the development of COPD. However, data for FEV$_1$ as a percentage of predicted was not presented, and only 51.4% of subjects attended all the visits. Clearly, more long term data are needed in this interesting area of study.

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Can inactivated Mycobacterium vaccae vaccine help protect HIV positive patients against tuberculosis?


Inactivated vaccines have better safety profiles than live vaccines in HIV positive subjects, and previous research has shown that immunological responses to non-tuberculous mycobacteria provide protection against tuberculosis (TB).

This study compared the response of both HIV positive (n = 39) and HIV negative (n = 10) individuals with prior BCG immunisation to five doses of heat inactivated Mycobacterium vaccae (MV) vaccine with a control. Lymphocyte proliferation (LPA) and interferon gamma (IFN-$

\gamma$) responses to MY and Mycobacterium tuberculosis (MTB) antigens were measured at intervals before, during, and after the course. The immune response to MV increased significantly after immunisation and was still evident at >1 year. There was a less significant response to MTB antigen in HIV positive subjects. No significant change was seen in the CD4 cell counts or viral loads of HIV positive patients before and after immunisation. There were no major adverse effects.

Inactivated MV vaccine appears to boost BCG primed immunity in HIV positive individuals which may provide protection against the development of active TB. Further large scale trials of the efficacy of this regimen are currently in progress.

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