**ORIGINAL ARTICLE**

**Fibroblast growth factor 10 haploinsufficiency causes chronic obstructive pulmonary disease**

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**ABSTRACT**

**Background** Genetic factors influencing lung function may predispose to chronic obstructive pulmonary disease (COPD). The fibroblast growth factor 10 (FGF10) signalling pathway is critical for lung development and lung epithelial renewal. The hypothesis behind this study was that constitutive FGF10 insufficiency may lead to pulmonary disorder. Therefore investigation of the pulmonary functions of patients heterozygous for loss of function mutations in the FGF10 gene was performed.

**Methods** The spirometric measures of lung function from patients and non-carrier siblings were compared and both groups were related to matched reference data for normal human lung function.

**Results** The patients show a significant decrease in lung function parameters when compared to control values. The average FEV1/IVC quota (FEV1%) for the patients is 0.65 (80% of predicted) and reversibility test using Terbutalin resulted in a 3.7% increase in FEV1. Patients with FGF10 haploinsufficiency have lung function parameters indicating COPD. A modest response to Terbutalin confirms an irreversible obstructive lung disease.

**Conclusion** These findings support the idea that genetic variants affecting the FGF10 signalling pathway are important determinants of lung function that may ultimately contribute to COPD. Specifically, the results show that FGF10 haploinsufficiency affects lung function measures providing a model for a dosage sensitive effect of FGF10 in the development of COPD.

**INTRODUCTION**

Chronic obstructive pulmonary disease (COPD) is a complex disease and one of the leading causes of death in developed countries. The disease results from major remodelling of the airspace and changes in the lung epithelium.1 The simplest classification of COPD is related to the degree of airflow obstruction, regardless of the underlying mechanism. In contrast to asthma, which is defined by a reversible airflow obstruction, COPD is characterised by irreversible and usually progressive obstruction with an abnormal inflammatory response to noxious gases or particles.2 3 The most important risk factor for the development of COPD is cigarette smoking, but only a minority of smokers will develop the disease, suggesting contributions of genetic factors.4 This is supported by the demonstration of familial aggregation of lung function measures as well as the clustering of COPD within families.5 Genetic studies in humans and in animal models suggest that genes associated with lung developmental processes are also implicated in COPD.1 4 6 Lung development requires branching morphogenesis, a process that is dependent on cell–cell communication between the mesenchyme and the lung epithelium. Both lung development and lung homeostasis depend on precise molecular signalling involving different fibroblast growth factors (FGF), sonic hedgehog (SHH), bone morphogenetic protein (BMP), transforming growth factor β (TGFβ), retinoic acid, the wingless (WNT) related family of proteins, and various transcription factors.7 8 Similar mechanisms are required to maintain and regenerate the lung epithelium from progenitor cells located both in the epithelium and in the mesenchyme along the airways.9 Thus, factors and molecular pathways involved in the differentiation and renewal of the lung epithelium provide candidate mechanisms in pulmonary disease.10 11

FGF10 is expressed in the mesenchyme and its signalling is mediated by the receptor FGFR2b required for the development of many branched organs including lungs, thyroid, pituitary, lacrimal, and salivary glands.12 13 During early lung development, FGF10 signalling is tightly regulated by the retinoic acid dependent network through either WNT (via DKK1) or TGFβ-1 signalling.6 In the developing lung FGF10 is regulated by SHH signalling and other members of the FGF family.14 15 Furthermore, decreased FGF10-FGFR2 signalling alters the expression of SHH and BMP4, as well as FGF10 itself, establishing a feedback loop.16 The importance of the FGF10 pathway during lung initiation and development has been established in model systems after targeted disruption of the corresponding genes or by ectopic over-expression of the proteins.17 20 The most profound disruption of lung morphogenesis is observed in Fgfr2 null mice embryos showing failure to form lung buds.21 22 In the adult lung, FGF10 has been identified as a survival factor, and exogenous Fgf10 expression in mice decreases induced alveolar epithelial cell DNA damage, apoptosis, and the extent of induced lung fibrosis.23 24 Furthermore, FGF10 is coordinately expressed with BMP4 during hyperoxia injury repair.25 In vitro studies have recently shown that soluble FGF10 and hepatocyte growth factor (HGF) can replace the requirement for mesenchymal support in lung epithelial formation, allowing for clonal expansion and self-renewal.26 These data suggest a model in which the adult lung contains FGF10 dependent cells, giving rise to airway and alveolar epithelial cell lineages.
In humans, heterozygous mutations in the genes encoding FGF10 or FGFRR2b result in autosomal dominant aplasia of lacrimal and salivary glands (ALSG) and lacrimo-auriculo-dento-digital syndrome (LADD), respectively. No lung phenotype has been described in these patients, with the exception of a male LADD patient presenting with complex pulmonary malformations.\textsuperscript{28–32} FGF10 and FGFRR2b are essential for lung initiation, lung development, and lung epithelial proliferation in experimental systems, and we hypothesised that haploinsufficiency for FGF10 affects lung function that may ultimately result in the development of COPD.\textsuperscript{33} To examine this hypothesis, we measured the pulmonary functions in patients with loss of function mutations in the FGF10 gene as well as their non-carrier siblings. Spirometric values from both groups were compared with reference data for pulmonary functions.\textsuperscript{34} Lung function parameters were also measured in a mice model heterozygous for \textit{Fgf10} in order to validate a pulmonary phenotype. Our results indicate that \textit{FGF10} haploinsufficiency affects lung function that may ultimately lead to the development of COPD.

**METHODS**

**Patients**

Twelve patients with ALSG and heterozygous loss of function mutations in \textit{FGF10} were included in this study. The patients belong to two previously described families segregating a deletion (n=10) or a nonsense mutation (n=2) in the \textit{FGF10} gene, respectively.\textsuperscript{29} The age range of patients was from 13–69 years, with a mean age of 39 years. Six patients were male and six female. Six of the 12 patients were diagnosed with ‘asthma’ before this study and four patients had milder allergies. Two were smokers (<one package a day), three patients stopped smoking 3–11 years ago, and seven patients never smoked (supplementary table S1). None of the patients had any known lung malformation or abnormal tendency for airway/lung infections. Three non-carrier siblings (age 30, 33 and 34 years, one male and two females, respectively) from one family were included as ‘internal’ controls. Informed consent was obtained from all adult participants or parents of children who participated in this study under a protocol approved by the Regional Ethical Committee.

**Mice**

Mice heterozygous for the Fgf10 gene\textsuperscript{22} were housed and bred in the animal core facility of the Biomedical Centre, Uppsala University under pathogen-free conditions. In total, 12 haploinsufficient (Fgf10 +/−) mice and 12 wt (Fgf10 +/+ ) mice (C57BL/6) were used in this study at 13–17 weeks of age. Genotyping was performed using a primer against the neo-cassette together with a specific mouse Fgf10 primer (available upon request). The study followed a protocol approved by the Regional Ethical Committee for animal experiments.

**Pulmonary function**

Lung function measurements of study participants were performed by trained technicians and according to the American Thoracic Society (ATS) or European Respiratory Society (ERS) criteria.\textsuperscript{35} Lung function in mice were investigated using the eSpiro Forced Maneuvers System (FMS) (EMMS, Hants, UK). \textit{Human}: We measured forced expiratory volume during 1 s (FEV\text sub{1}) and vital capacity (VC) using a computerised pneumotachograph spirometer. Inspiratory vital capacity (IVC) was measured and total lung capacity (TLC) calculated with a volume plethysmograph (Jaeger Masterscreen body; Hoechberg, Germany). FEV\text sub{1} was determined after treatment with terbutaline. \textit{Mouse}: Mice were anaesthetised (0.2 ml/20 g mouse, intraperitoneally, with ketamine, xylazine, acepromazine, and saline, 2:1:0.3:6.6 ml). The eSpiro FMS was used to assess the lung function parameters forced expiratory volume in 75 ms (FEV\text sub{75}), forced vital capacity (FVC), inspiratory capacity, and TLC.

**Reference data**

Normal reference values for FEV\text sub{1} were obtained from a published dataset of spirometry from childhood to old age.\textsuperscript{34} The references for normal human lung function are based on measures from an ethnically matched control population (sex, age, height). Individual reference values (predicted values) for TLC were calculated according to the following formulae: TLC (l) males=(7.956* (height (m))−6.948, RSD=0.77, TLC (l) females=(7.107* (height (m))−6.45, RSD=0.83.\textsuperscript{35} The reference values are comparable to previously published values from the Swedish population.\textsuperscript{27–29} The lower limit of the normal (LLN) range was calculated by subtracting 1.645 * RSD (residual SD) from the predicted value.

**Statistics**

Spirometry data from \textit{FGF10} heterozygous patients were compared with the predicted reference values using paired Student t test assuming equal variance, as well as to the values obtained from three siblings without \textit{FGF10} mutations using two-tailed Student t test assuming equal variance. Data from the healthy siblings were compared with predicted reference values using two-tailed Student t test assuming equal variance to validate the use of the prediction equations. Airway obstruction was classified according to the ATS/ERS 2005 (obstruction FEV\text sub{1}/VC <LLN) and the GOLD 2006 (FEV\text sub{1}/VC <0.70) standards.\textsuperscript{30,32} The use of a single cut-off for FEV\text sub{1}/VC of 0.70 (GOLD 2006) to define the limits of normal of lung function, without considering age, may result in a false negative diagnosis of obstruction before age 45–50 years as well as a false positive diagnosis above this age because of the effect of age on lung volumes. A method used to minimise this potential misclassification, as supported by both the ATS and the ERS, is the LLN values for FEV\text sub{1}/VC (ATS/ERS 2005).\textsuperscript{41} This is based on the normal distribution and classifies the bottom 5% of the healthy population as abnormal. We classified the patients according to both standards (supplementary table S2). The influence of smoking on pulmonary functions (FEV\text sub{1}, IVC, and FEV\text sub{1}/IVC according to Tiffeneau index)\textsuperscript{42} was assessed with two-way ANOVA and two-tailed Student t test, assuming equal variance, by grouping individuals as non-smokers, current smokers, and former smokers. The group of non-smokers was compared to current smokers and former smokers; non-smokers and former smokers were compared to current smokers, and non-smokers were compared to current smokers.

The data from wt (Fgf10 +/+ ) mice were compared to heterozygous (Fgf10 +/−) littermates using two-tailed Student t test assuming equal variance. Comparison of lung measures between the two groups of mice was performed with and without normalisation to body weight (BW), lung weight (LW) or lung capacity (total lung capacity TLC). Values of p<0.05 were considered significant.

**RESULTS**

**Lung function in human subjects**

The \textit{FGF10} heterozygous patients showed a significant decrease in IVC (83% of predicted; p<0.001), FEV\text sub{1} (68% of predicted; p<0.001), and FEV\text sub{1}/IVC quota (FEV\text sub{1}/VC%; 0.65; p<0.001), but not in TLC (95% of predicted; p=0.12) (table 1 and table S2).
Significant differences in FEV$_1$ and FEV$_1$/IVC were also observed when comparing FGF10 heterozygous patients and the three non-carrier siblings used as controls (table 2). The values of the non-carrier siblings were similar to their predicted values, validating them as ‘internal’ controls in the study (table 1). Consistent with pulmonary obstructive disease, the average FEV$_1$/IVC was reduced (<0.7 and <LLN) and the FEV$_1$ (<70% of predicted) and a normal, or slightly reduced IVC (IVC ≥80% of predicted; table 1). Reversibility test using terbutaline gave a significant increase in FEV$_1$ (3.7% of predicted value, p=0.0013; table 2). This was still subnormal when compared to the predicted reference value (2.70 ±0.75 vs 3.63 ±0.60 (LLN 2.37 ±0.50); tables 1 and 2). Smoking, or having a history of smoking, did not correlate with lower IVC, FEV$_1$ or FEV$_1$/IVC in our group of patients (supplementary tables S1 and 2). Individual data revealed two individuals (MS90 and GA80) classified as COPD according to the GOLD standard (supplementary table S2). On the other hand, the two oldest individuals (IA57 and BA45) were classified as COPD according to the GOLD standard but could be defined as false negatives when applying the ATS/ERS 2005 standard (supplementary table S2). On the other hand, the two oldest individuals (IA57 and BA45) were classified as COPD according to the GOLD standard but could be defined as false positive using the ATS/ERS 2005 standard. Individual BA45 was “borderline” normal with an FEV$_1$/IVC of 0.65 (LLN 0.64), while individual IA57 could be classified as having a restrictive phenotype with a normal FEV$_1$/IVC (0.66 >LLN 0.62), due to a very low IVC (2.86 <LLN 3.46), and a low TLC (6.05 <LLN 6.11). Individual LA41 had an obstructive disease (FEV$_1$/IVC 0.627 <LLN 0.652) with restrictive components (IVC 3.35 <LLN 3.71, and TLC 5.11 <LLN 6.27), indicating a mixed phenotype.

### Lung function in mouse model

Fgf10 +/-- mice showed a significant decrease in FEV$_{75}$, FVC, as well as a decrease in the FEV$_{75}$/FVC quota (FEV$_{75}$%). The mice also had a decreased lung size (TLC) as well as body and lung weight compared to wild type (wt) male littermates. However, the lung/body weight ratio was similar when comparing Fgf10 +/-- and Fgf10 +/+ mice. To eliminate a possible effect of body and lung size, mice data were normalised to lung capacity, body weight, lung weight, and by comparing weight matched mice. When comparing weight matched Fgf10 +/-- and Fgf10 +/+ mice we observed a slight decrease in lung weight (supplementary table S3). Lung morphology and histopathology (gemma staining) appeared normal in Fgf10 +/+ mice (supplementary figure 1).

### DISCUSSION

Independent genome-wide association studies (GWAS) have recently identified single nucleotide polymorphisms (SNPs) near the hedgehog interacting protein (HHIP) locus on chromosome 4 associated with airflow obstruction. HHIP is a critical negative regulator of the hedgehog pathway, including the ligand sonic hedgehog (SHH). Overexpression of Shh in mouse lung rudiments reduces transcription of the Fgf10 gene. Similarly to mice deficient for Fgf10, branching morphogenesis is greatly impaired in mice deficient for Hhip and it has been hypothesised that the lung phenotype in Hhip deficient mice results from hyperactive Shh followed by inhibition of Fgf10 expression. This may explain the morphological similarities in mouse models deficient for either Hhip or Fgf10 and supports altered Fgf10 expression as a candidate mechanism in COPD. Interestingly, the Fgf10 downstream target, STAT V3, has recently been associated with COPD, providing yet another link to Fgf10 signalling. The results of our study show that haploinsufficiency for Fgf10 is associated with reduced pulmonary function consistent with COPD. FGF10 insufficient patients show a non-reversible airway obstruction when compared with both predicted reference values and siblings with normal FGF10.
alleles. These findings provide a direct connection between FGF10 expression and human lung function. Patients with FGF10 insufficiency, also associated with the ALSG syndrome, have a significantly reduced FEV1 (68% of predicted) and an average FEV1/FVC of 0.65 (normally above 0.70–0.75) consistent with obstructive lung disease. This is supported by a normal TLC and a disproportionally and slightly reduced IVC (83% of predicted). In COPD, both FEV1 and IVC are below predicted levels, but the FEV1/FVC ratio is normal or increased.40 The modest increase in FEV1 (5.7% of predicted value) after terbutaline treatment in our patients indicates that the obstructive symptoms are irreversible. A reversible obstruction (eg, asthma) is defined by an increase of at least 12% in FEV1 in response to terbutaline treatment.49 The degree of COPD in our group of patients is classified as moderate or stage II COPD (FEV1/FVC < LLN and 60% <FEV1 <69% of predicted).41

To validate our results from spirometric measures in patients with FGF10 haploinsufficiency we studied the pulmonary functions in Fgf10 heterozygous mice. Even though FGF10 deficient mice show a reduction in pulmonary functions similar to those observed in our patients, these results are most probably due to the smaller size of the Fgf10 +/- mice. Studies of pulmonary functions in mice indicate that direct comparisons between mice and human for key variables such as FEV and FVC are less useful.50 The mouse lung shares general structure and physiological mechanisms with the human lung but with some differences. Mice have fewer submucosal glands restricted to the upper trachea and only six to eight branches before reaching the terminal bronchiole.51 Furthermore, the respiratory bronchioli—structures shown to be involved in the development of emphysema—are absent in the mouse.52 These differences may, in part, explain the differences in spirometric measurements between mice and man.

Despite tremendous efforts to understand the genetics behind COPD, most factors remain obscure with rare exceptions such as ∆s1-antitrypsin deficiency.53–56 Our results support the idea that genetic variants affecting the FGF10 signalling pathway are important determinants of lung function that may ultimately contribute to COPD. Specifically, we show that FGF10 haploinsufficiency affects lung function measures, providing a model for a dosage sensitive effect of FGF10 in the development of COPD.

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Competing interests None.

Ethics approval The human studies were approved by the local ethics committee in Uppsala University Hospital, the Swedish Society for Medical Research, and Futurum academy of science and healthcare, Jönköping County Council.

Contributors JK wrote the manuscript, performed statistical analyses and interpreted the data. CB, JB, HHF, and CSB performed the experiments. PB and BB participated in study design and data interpretation. ND designed the study and edited the manuscript.

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