Increased expression of 5-hydroxytryptamine_{2A/B} receptors in idiopathic pulmonary fibrosis: a rationale for therapeutic intervention

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ABSTRACT
Background Idiopathic pulmonary fibrosis (IPF) has a poor prognosis and limited responsiveness to available treatments. It is characterised by epithelial cell injury, fibroblast activation and proliferation and extracellular matrix deposition. Serotonin (5-hydroxytryptamine; 5-HT) induces fibroblast proliferation via the 5-HT_{2A} and 5-HT_{2B} receptors, but its pathophysiological role in IPF remains unclear. A study was undertaken to determine the expression of 5-HT receptors in IPF and experimental lung fibrosis and to investigate the effects of therapeutic inhibition of 5-HT_{2A/B} signalling on lung fibrosis in vivo and in vitro.

Methods and results Quantitative RT-PCR showed that the expression of 5-HT_{2A/B} and 5-HT_{2B} was significantly increased in the lungs of patients with IPF (n=12) and in those with non-specific interstitial pneumonia (NSIP, n=6) compared with transplant donors (n=12). The expression of 5-HT_{2B} was increased specifically in IPF lungs but not in NSIP lungs. While 5-HT_{2A} protein largely localised to fibroblasts, 5-HT_{2B} localised to the epithelium. To assess the effects of 5HT_{2A/B} inhibition on fibrogenesis in vivo, mice were subjected to bleomycin-induced lung fibrosis and treated with the 5-HT_{2A/B} antagonist terguride (or vehicle) in a therapeutic approach (days 14–28 after bleomycin). Terguride-treated mice had significantly improved lung function and histology and decreased collagen content compared with vehicle-treated mice. Functional in vitro studies showed that terguride is a potent inhibitor of transforming growth factor β_{1} or WNT3a-induced collagen production.

Conclusion The studies revealed an increased expression of 5-HT_{2A/B} specifically in IPF. Blockade of 5-HT_{2A/B} signalling by terguride reversed lung fibrosis and is thus a promising therapeutic approach for IPF.

INTRODUCTION
Idiopathic pulmonary fibrosis (IPF) is a progressive and fatal lung disease of unknown origin. 1,2 It is characterised by alveolar epithelial cell damage, increased deposition of extracellular matrix (ECM) in the lung interstitium, enhanced fibroblast/myofibroblast proliferation and activation and, ultimately, distortion of normal lung architecture and loss of respiratory function. 3,4 The hallmark lesions of IPF are fibroblast foci, sites featuring activated myofibroblasts mainly responsible for altered ECM deposition in the lung interstitium. 3 The pathogenesis of IPF includes inappropriate alveolar regeneration in response to repetitive epithelial injuries perpetuated by dysregulated wound repair in response to local inflammation. 5,6 While the initial trigger of this disease is most likely represented by repetitive epithelial injuries of unknown origin, the interstitial fibroblast/myofibroblast represents the key effector cell responsible for ECM deposition in patients with IPF and in animal models of lung fibrosis such as the bleomycin-induced mouse model. 7,8 It is currently unclear, however, which mediators and pathways are involved in disease initiation and progression. While soluble cytokines such as transforming growth factor (TGF) β_{1}, interleukin (IL)-1β or WNT ligands have been assigned a pathogenic role in IPF and experimental models thereof, 9,10 therapeutic options neutralising their activity have not been studied in appropriate clinical trials.

Serotonin (5-hydroxytryptamine, 5-HT) exhibits well-characterised regulatory functions in multiple physiological systems. 13,14 It is involved in the regulation of cell migration, proliferation, cytokine production and vasoregulation. 15,16 The multiple actions of 5-HT are mediated by seven different 5-HT receptor subtypes (5-HT_{1A} to 5-HT_{2C}). 17 Importantly, 5-HT and its receptors 5-HT_{2A} and 5-HT_{2B} have been implicated in fibrotic disorders such as retinopathy or pleural fibrosis, carcinoid heart disease and liver fibrosis. 18,19 In addition, cumulative evidence suggests an important biological role for 5-HT signalling in the lung, supported by increased levels of 5-HT and active signalling thereof in asthma and pulmonary hypertension. 20,21 In addition, recent evidence has suggested that the 5-HT_{3A} antagonists ketanserin and SB215505 reduced collagen mRNA and protein levels in the bleomycin-induced mouse model. 22 Accordingly, this evidence demands a detailed characterisation of 5-HT receptor expression in IPF and analysis to determine whether 5-HT_{2A/B} antagonists represent a therapeutic option in IPF, particularly since 5-HT_{2B} antagonists are already in clinical use for other diseases. 24,25 Among these, terguride (1,1-diethyl-3-(6-methyl-5a-ergolino)urea) is a potent antagonist of 5-HT_{2A} and HTR_{2B}, a partial dopaminergic agonist and an α_{2}-adrenoceptor agent. 24,26 Terguride is clinically approved, well-tolerated and has been shown to be efficacious in the treatment of ovulation disorders due to hyperprolactinaemia. 27
The expression and localisation of the 5-HT system in IPF lungs was investigated and compared with non-specific interstitial pneumonia (NSIP) and donor lungs. We performed a therapeutic approach in vivo using the mouse model of bleomycin-induced lung fibrosis to determine whether 5-HTR2A/B antagonism by terguride is a suitable therapeutic option for IPF.

MATERIAL AND METHODS

Human tissue and cells

Lung tissues were obtained from 12 patients with IPF with a histological pattern of usual interstitial pneumonia (UIP) (4 women, 8 men; mean±SD age 57±9 years), 6 patients with non-specific interstitial pneumonia (NSIP, fibrotic phenotype) (4 women, 2 men; mean±SD age 55±9 years) and 12 control subjects (transplant organ donors; 6 women, 6 men; mean±SD age 40±10 years) after explantation. Detailed patient characteristics are shown in table S1 in the online supplement. Human ATII cells were isolated as previously described.36 The purity and viability of ATII cell preparations was consistently >90% and >95%, respectively. Freshly isolated human ATII cells were used for gene expression analysis.

Animals

Pathogen-free 6–8-week-old female C57BL/6N mice (Charles River Laboratories, Sulzfeld, Germany) weighing 18–22 g were used throughout the study. All experiments were performed in accordance with the guidelines of the ethics committee of University of Giessen School of Medicine and approved by the local authorities. Mice had ad libitum access to water and rodent laboratory chow. Bleomycin sulphate (Laboratories Almirall SA, Barcelona, Spain) was dissolved in sterile 0.9% saline and applied as a single dose of 1.5 U/kg body weight in a total volume of 80 μl by orotracheal application. Control mice received 80 μl saline only. Mice were treated with terguride 14 days after bleomycin application. As an internal control, a treatment arm with daily administration of 50 mg/kg imatinib was included in the study. Treatment with imatinib was started immediately following instillation of bleomycin. The detailed treatment scheme is shown in figure S4 in the online supplement.

Immunohistochemistry

Human lungs were submerged in 4% (w/v) paraformaldehyde after explantation and processed for paraffin embedding.29 30 Immunohistochemical staining of sections (5 μm) was performed according to the manufacturer’s protocol (Histostain Plus Kit; Zymed Laboratories/Invitrogen, Carlsbad, California, USA). Antigen retrieval was performed in 6.5 mM sodium citrate pH 6.0 in a pressure cooker, after which endogenous peroxidase activity was quenched with 3% (v/v) hydrogen peroxide for 20 min. The following antibodies were used in the study: 5HTR2A (polyclonal, sc-50396, dilution 1:200; Santa Cruz Biotechnologies, California, USA); 5HTR3B (polyclonal, ab12926, dilution 1:200; Abcam, Cambridge, UK). An antibody against 5HTR3B (polyclonal, HPA012867, dilution 1:100, Sigma-Aldrich, St Louis, Missouri, USA) was also used and staining was performed using the Vectastain Fast Red Chromogen Kit (Zymed Laboratories/Invitrogen). For the negative control, an isotype-matched non-specific antibody was used (ab2410, dilution 1:200, Abcam).

Further experimental procedures

The detailed procedures for the determination of terguride levels, cell culture, quantitative RT-PCR, collagen assay, lung function and determination of the fibrotic score are provided in the online supplement.

Statistical analysis

All ΔCt values obtained from quantitative real-time PCR (qRT-PCR) were analysed for normal distribution with the Shapiro-Wilk test using assignment of a normal distribution with p>0.05 and confirmed using quantile-quantile plots. The means of indicated groups were compared using the two-tailed Student t test or one-way analysis of variance (ANOVA) with Tukey HSD post hoc test for studies with more than two groups. The results were considered statistically significant when p<0.05.

RESULTS

Expression analysis of 5-HT receptors and the 5-HT transporter 5-HTT in donor and IPF lung specimens

Initially we analysed the expression of 5-HT receptors and the 5-HT transporter 5-HTT in lung tissue samples from controls (transplant donors) and from patients with NSIP or IPF using qRT-PCR. As shown in figure 1, all investigated 5-HT receptors except 5HTR2C were expressed in normal as well as diseased lung tissue. 5-HTR1A, 5-HTR1B (figure 1A), 5-HTR2A and 5-HTR3B (figure 1B) were markedly upregulated in IPF lungs (mean±SEM log-fold change 2.36±0.43, 2.14±0.46, 3.74±0.41 and 1.60±0.52, respectively). 5-HTT was expressed at high levels in control lungs; it was dramatically downregulated in IPF (log-fold change –4.05±0.71, figure 1C) but not in NSIP lungs (fibrotic NSIP). Interestingly, in NSIP lung specimens 5-HTR2A and 5-HTT were not differentially regulated compared with control subjects but 5-HTR1A, 5-HTR1B and 5-HTR3B were also upregulated (figure 1). The increased expression of 5-HTR2A thus presented a specific event in UIP.

Localisation of 5-HTR2A and 5-HTR2B in donor and IPF lung specimens

Following the initial results, we focused our study on protein localisation of 5-HTR2A and 5-HTR2B. In order to localise cell types capable of 5-HT signalling via these 5-HTR, we assessed the distribution of 5-HTR2A and 5-HTR2B in donor and IPF lung tissue sections by immunohistochemistry (figures 2 and 3, respectively). 5-HTR2A was mainly expressed in bronchial and vascular smooth muscle cells; in IPF, additional staining in interstitial fibroblasts was observed (figure 2, arrows). As shown in figure 3 and figure S2 in the online supplement, 5-HTR2B was expressed largely in endothelial cells in control tissue. In IPF lungs, 5-HTR2B staining was mainly observed in bronchial and alveolar epithelial cells (figure 3A, arrowheads). To further corroborate these results we quantified the cell-specific gene expression patterns of 5-HTR2A and 5-HTR2B in primary human ATII cells derived from patients with IPF or transplant donors using qRT-PCR. As shown in figure 3B, there was a significant increase in 5-HTR2B mRNA expression in IPF ATII cells whereas the 5-HTR2A level was not significantly different in human ATII cells from patients with IPF and transplant donors.

Administration of the 5-HTR2A/B antagonist terguride in vivo

We next assessed whether inhibition of 5-HTR2A/B represented an effective therapeutic option in lung fibrosis using mice subjected to orotracheal instillation of bleomycin, a well-accepted experimental model of lung fibrosis.31 First we analysed the expression of 5-HT receptors and 5-HTT in experimental lung fibrosis. As shown in figure S1 in the online supplement, 5-HTR2A and 5-HTR2B were significantly upregulated 7 and
14 days after bleomycin injury, which is in agreement with previous reports.23 Next we administered the clinically approved 5-HTR2A/B antagonist terguride to mice subjected to bleomycin. In order to ensure that terguride partitions into lung tissue, we next determined the concentrations of terguride in plasma and lung tissue under our experimental conditions (see figure S3 in online supplement). One hour after intraperitoneal administration of 1.2 mg/kg terguride the plasma concentrations of terguride ranged from 2.06–20.26 nM. In perfused lung tissue, comparable terguride concentrations to plasma were detected in all animals independent of the degree of fibrosis (figure S3 in online supplement). This indicated that terguride rapidly equilibrated in lung tissues and during the time course of bleomycin-induced lung fibrosis. Treatment with terguride was well tolerated and no side effects were observed throughout the study. The body weights of the mice were not significantly different between bleomycin-treated mice, independent of treatment with vehicle or terguride (see figure S3 in online supplement).

**Effects of 5-HTR2A/B antagonism on lung fibrosis in vivo**

To determine whether 5-HTR2A/B antagonism affected the progression of lung fibrosis we initiated treatment with terguride 14 days after bleomycin instillation (referred to as ‘therapeutic approach’). Mice were treated with two different doses of terguride by intraperitoneal application twice daily until day 28. As an internal control, a treatment arm with daily administration of 50 mg/kg imatinib was included in the study.32 Treatment with imatinib was started immediately after instillation of bleomycin (referred as ‘preventive approach’). The detailed treatment scheme is shown in figure S4 in the online supplement. We investigated the effects of 5-HTR2A/B antagonism on lung fibrosis in vivo.

![Figure 1](image1.png) **Figure 1** mRNA expression profile of serotonin (5-HT) receptors (A) 5-HTR1A,B, (B) 5-HTR2A,B and (C) the 5-HT transporter (5-HTT) in lung samples from controls (transplant donors) and patients with non-specific interstitial pneumonia (NSIP) and idiopathic pulmonary fibrosis/usual interstitial pneumonia (IPF/UIP) assessed by quantitative real-time PCR. The results are derived from 12 controls, 6 patients with NSIP and patients with 12 IPF/UIP and are presented as single values per lung sample. *p<0.05.

![Figure 2](image2.png) **Figure 2** Expression and localisation of the serotonin (5-HT) receptor 5-HTR2A in lung tissue from controls (transplant donors) and patients with idiopathic pulmonary fibrosis (IPF). Immunohistochemical staining was performed on tissue sections of donor lungs (left panels) and IPF lungs (right panels). Stainings are representative of four independent experiments using at least three different donor or IPF lung tissues. Scale bars indicate 100 μm. Arrows indicate positive staining of fibroblasts in IPF.
antagonism by terguride in vivo on key features of lung fibrosis such as impaired respiratory function, destroyed lung architecture and increased collagen deposition.

The assessment of lung compliance in vivo showed slight but significant improvements in lung function of terguride-treated but not imatinib-treated mice (figure 4A). We next assessed the degree of lung fibrosis by determination of the total hydroxyproline content (figure 4B) and fibrotic score (figure 4C). As shown in figure 4B, mice subjected to 5-HTR2A/B antagonism by terguride had significantly less total collagen than vehicle-treated controls (bleomycin+vehicle: 148±22%, bleomycin+terguride 0.4: 122±27%, bleomycin+terguride 1.2: 91±21%, n=10 each). These findings were corroborated by the fact that 5-HTR2A/B antagonism also led to a dose-dependent decrease in the fibrotic score (bleomycin+vehicle: 5.1±0.3, bleomycin +terguride 0.4: 1.9±0.6, bleomycin+terguride 1.2: 1.46±0.2 compared with saline-treated controls, n=10 each; figure 4C). The effects of terguride on collagen content as well as the fibrotic score were comparable to those of imatinib treatment (preventive approach). Finally, immunohistochemistry of vehicle- and terguride-treated mouse lungs confirmed a marked attenuation of fibrosis with less ECM deposition and restored lung architecture (figure 4D).

**Effects of 5-HTR2A/B antagonism on lung fibrosis in vitro**

Fibroblasts are the key effector cells in lung fibrosis and are responsible for enhanced ECM deposition. We therefore assessed the effects of terguride on lung fibroblasts stimulated by the profibrotic mediators transforming growth factor (TGF)-β1 or WNT3a. Both, TGF-β1 and WNT3a treatment of human lung fibroblasts (HFL-1) led to a significant induction of the ECM components type I collagen Col1a1 as assessed by qRT-PCR (figure 5A). Moreover, these changes were confirmed at the total collagen level as assessed by Sircol assay (figure 5B). Most importantly, terguride treatment resulted in a significant decrease in TGF-β1-induced as well as WNT3a-induced collagen production at the mRNA (TGF-β1: 2.61±0.65, TGF-β1+terguride 1000: 0.26±0.77, WNT3a: 3.99±0.55, WNT3a+terguride 1000: 0.39±0.22; figure 5A) and protein level (TGF-β1: 1.54±0.29, TGF-β1+terguride 1000: 0.71±0.28, WNT3a: 1.28±0.16, WNT3a+terguride 1000: 0.96±0.12, figure 5B).

**DISCUSSION**

In this study we found altered expression of 5-HTR in IPF, suggesting that 5-HT signalling is altered in IPF. We analysed lung tissue samples from patients with IPF, NSIP and controls (transplant donors) and observed a marked upregulation of...
5-HTR1A/B and 5-HTR2A/B but a dramatic downregulation of 5-HTT in IPF. Interestingly, in NSIP lung specimens 5-HTR2A was not differentially regulated compared with control subjects, which suggests this is a specific event in UIP.

We demonstrated that inhibition of 5-HTR2A and 5-HTR2B resulted in marked attenuation of bleomycin-induced lung fibrosis and thereby may be a suitable therapeutic approach in lung fibrosis. These findings are of special interest as IPF has a poor prognosis owing to limited responsiveness to currently available therapies.

5-HT and its receptors 5-HTR2A and 5-HTR2B have been implicated in fibrotic disorders such as retroperitoneal fibrosis, nephropathies, carcinoid heart disease and liver fibrosis. Cumulative evidence of the biological role of 5-HT in the lung has emerged, such as controlling vasoreactivity or broncho-reactivity. Under disease conditions such as asthma, increased levels of 5-HT correlate with the clinical status and pulmonary function, and several studies have indicated of the involvement of type I collagen mRNA level of type I collagen fibroblasts. Using qRT-PCR analysis, we demonstrated that inhibition of 5-HTR2A and 5-HTR2B resulted in marked attenuation of bleomycin-induced lung fibrosis and thereby may be a suitable therapeutic approach in lung fibrosis. These findings are of special interest as IPF has a poor prognosis owing to limited responsiveness to currently available therapies.

Figure 4 Lung function and collagen content after terguride treatment in experimental lung fibrosis. Mice were subjected to saline or bleomycin instillation and treated with terguride (ter). (A) Lung compliance measurements were obtained on days 14, 21 and 28. After 28 days the lungs were processed for (B) hydroxyproline measurement, (C) determination of the fibrotic score in lung sections stained with haematoxylin and eosin or (D) histological assessment using haematoxylin and eosin staining of lung sections. Scale bars indicate 50 μm.

Figure 5 Effect of terguride (ter) treatment on transforming growth factor (TGF)-β1-induced or WNT3a-induced type I collagen α1-expression in fibroblasts. (A) Using qRT-PCR analysis, the mRNA level of type I collagen α1 was assessed in human lung fibroblasts after stimulation with TGF-β1 (2 ng/ml) or WNT3a (100 ng/ml) for 12 h as indicated (n=4). (B) Human lung fibroblasts were stimulated with TGF-β1 (2 ng/ml) or WNT3a (100 ng/ml) for 24 h and the total collagen content was measured using the Sircol collagen assay (n=4).

5-HT leads to vasoconstriction and vascular proliferation under hypoxic conditions, and enhanced 5-HTR2B expression seems to be centrally involved in the development of this disease.

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between 5-HTT and fibrogenesis.\textsuperscript{41} Ablation of 5-HTT reduces the levels of 5-HT because platelet 5-HT is known to contribute 99\% of 5-HT in the blood. Since 5-HTT is also the predominant pathway of 5-HT clearance by lung cells, an increase in the local 5-HT concentration in the lung with concomitant increases in binding to and signalling through 5-HT\textsubscript{3} receptors may be the consequence.\textsuperscript{41} Downregulation of 5-HTT and upregulation of 5-HT\textsubscript{2A} and 5-HT\textsubscript{2B} may lead to increased 5-HT levels and enhanced receptor availability, respectively, and thus synergistic regulation of fibrogenesis. A detailed investigation of the underlying mechanisms remains elusive owing to limitations of investigative tools. In particular, the assessment of free and unbound 5-HT in nanomolar concentrations in blood is influenced by artificial activation of platelets and is therefore difficult to assess and interpret in mouse models.

We assessed the potential antifibrotic effects of 5-HT\textsubscript{2A/B} antagonism using a therapeutic approach in the mouse model of bleomycin-induced lung fibrosis. We administered terguride, which is clinically approved for the treatment of ovulation disorders due to hyperprolactinemia and hyperprolactaemic pituitary adenoma.\textsuperscript{27} Terguride belongs to the ergoline class of compounds which have been shown to have a high affinity to several 5-HT receptor isoforms. In contrast to most ergoline derivatives including ergotamine, pergolide, cabergoline and bromocriptine, terguride is a potent 5HT\textsubscript{2A} and 5HT\textsubscript{2B} receptor antagonist.\textsuperscript{24} Chronic treatment with ergoline derivatives such as pergolide and, to a lesser extent, cabergoline or with anorectic drugs such as fenfluramine has been associated with retroperitoneal, pleural and pericardial fibrosis as well as valvular heart disease.\textsuperscript{42} These drugs exhibit 5-HT\textsubscript{2B} receptor agonist properties. In contrast, terguride acts as a non-surmountable antagonist on 5-HT\textsubscript{2B}. Interestingly, Hauso et al recently showed that terguride treatment prevented 5-HT-induced heart valve disease and hypertrophy in rats.\textsuperscript{44}

We have shown that administration of terguride at two different doses in a therapeutic approach had potent dose-dependent therapeutic effects in the mouse model of bleomycin-induced lung fibrosis. This model has been widely used for drug efficiency studies over the years, but most compounds have been tested in a preventive rather than a therapeutic approach. The preventive approach interferes mostly with the inflammatory and early fibrogenic response whereas the therapeutic approach acts on the progression of fibrosis, better reflecting the clinical situation in IPF.\textsuperscript{51} We showed that treatment with terguride in a therapeutic fashion led to a significant improvement in respiratory function, collagen deposition, lung architecture and survival. This suggests that terguride may have anti-remodelling properties. Previous evidence has indicated that 5-HT binding to 5-HT\textsubscript{2A} exerts mitogenic properties via enhanced TGF-\protect\beta\textsubscript{1} expression. This mechanism seems to be regulated via protein kinase C and extracellular signal-regulated kinase.\textsuperscript{19} In addition, 5-HT\textsubscript{2B} has been reported to interact with the platelet-derived growth factor receptor pathway to regulate cell cycle progression.\textsuperscript{45} Both receptors may provide possible pathways which lead to the reverse actions of 5-HT\textsubscript{2A/B} antagonism by terguride. We have further corroborated the interaction between 5-HTR and TGF-\protect\beta\textsubscript{1} signalling, demonstrating that 5-HT is involved in collagen production in human lung fibroblasts induced by TGF-\protect\beta\textsubscript{1}.

Our findings are further validated by a recent study addressing the role of 5-HT signalling in experimental lung fibrosis. Fabre et al\textsuperscript{23} reported increased lung 5-HT levels and 5-HT\textsubscript{2A} and 5-HT\textsubscript{2B} expression during fibrogenesis in the bleomycin model of lung fibrosis, which we have confirmed in this study. Using a preventive approach and generic 5-HT\textsubscript{2A/B} antagonists in the bleomycin model, they also demonstrated attenuation of lung fibrosis with the inhibitors ketanserin (a 5-HT\textsubscript{2A} inhibitor) and SB258505 (a 5-HT\textsubscript{2B} inhibitor).\textsuperscript{23} Our results are consistent with these findings, thus confirming the relevance of 5-HT signalling in lung fibrosis and its role as a therapeutic target. In addition, we have advanced the understanding of the role of 5-HT in the pathophysiology of IPF by (1) characterising and quantifying in detail the expression of 5-HT receptors in human IPF/UIP, NISP and control tissues; (2) demonstrating the therapeutic feasibility of terguride in the experimental model of lung fibrosis using therapeutically relevant conditions, thereby providing a rationale for translational research of a well-characterised and safe drug in pulmonary fibrosis; and (3) by investigating in detail the biological mechanisms involved in the therapeutic effect of terguride—for example, by demonstrating a significant role of 5-HT\textsubscript{2A/B} in collagen turnover induced by the profibrotic mediators TGF-\protect\beta\textsubscript{1} and WNT\textsubscript{3a}.

In summary, we present strong evidence for an important role of 5-HT signalling via 5-HT\textsubscript{2A/B} in lung fibrosis in vivo and in vitro, and the possibility of selectively and safely modulating this pathway in vivo which may provide a possible therapeutic approach for patients with IPF.


SUPPLEMENT MATERIAL AND METHODS

Cell culture

The human lung fibroblasts (HFL-1) [German Collection of Microorganisms and Cell Cultures (DSMZ)] were maintained in DMEM containing 10% FCS and cultured in a humidified atmosphere of 5% CO₂ at 37 °C. For stimulation experiments cell were serum-starved for 24 h before treatment. Cells were treated with TGF-β1 (2 ng/ml) or WNT3a (100ng/ml) for 12 h or 24 h as indicated, and collagen content was determined by qRT-PCR and Sircol collagen assay.

Reverse transcription and quantitative real-time PCR

Total RNA extraction, cDNA synthesis, and quantitative (q)RT-PCR were performed using the primers listed in Table S2. Under identical cycling conditions, all primer sets worked with similar efficiencies to obtain simultaneous amplification in the same run, as described before24. Sequences were taken from GeneBank, all accession numbers are denoted. Hydroxymethylbilane synthase (HMBS) and hypoxanthine phosphoribosyltransferase 1 (HPRT1) for mouse and human, respectively and both ubiquitously and equally expressed genes that are free of pseudogenes, were used as reference genes in all qRT-PCR reactions. Relative transcript abundance is expressed as ΔCt value (ΔCt = Ctreference – Cttarget), where higher ΔCt values indicate higher transcript abundances, and negative ΔCt values represent genes that are less expressed compared with the reference gene. The fold-change of the transcript levels in IPF/NSIP versus control can be estimated by 2ΔΔCt, where ΔΔCt values are calculated as ΔCtpulmipf – ΔCtcontrol. Positive 2ΔΔCt values indicate upregulation, negative values indicate downregulation of a target gene.
Collagen assay

Total collagen content was determined using the Sircol Collagen Assay kit (Biocolor). Equal amounts of protein lysates were added to 1 ml of Sircol dye reagent, followed by 30 min of mixing. After centrifugation at 10,000 × g for 10 min, the supernatant was carefully aspirated and 1 ml of alkali reagent was added. Samples and collagen standards were then read at 540 nm in a spectrophotometer (Bio-Rad). Collagen concentrations were calculated using a standard curve with acid-soluble type 1 collagen.

Determination of terguride levels in the mouse

Terguride levels were determined in mouse plasma and lung tissue samples by liquid chromatography-mass spectrometry (LC-MS). In brief, two lung samples each were pooled and homogenised in phosphate buffer (100 mmol/L). A 200µl sample of lung homogenate or plasma was then precipitated with acetonitrile (400µl) and centrifuged. The supernatant was dried in a vacuum centrifuge, resuspended in phosphate buffer (200µl), and quantitated by LC-MS using terguride standards, which were prepared on the respective matrices used for LC-MS. Proterguride was applied as an internal standard. LC-MS was carried out on a Waters 2795 Alliance Quattro Micro (Micromass/Waters) using a Luna 3µC18 Phenomenex column with a column temperature of 30°C. A gradient with a trinary mobile phase system consisting of water and 0.1% formic acid (mobile phase A), acetonitrile (mobile phase B), and 2% formic acid in water (mobile phase C) at a flow rate of 0.4 ml/min was applied.

Lung function

Compliance was measured invasively in anaesthetized animals with the use of a commercially available system (Hugo Sachs Elektronik-Harvard apparatus, March-Hugstetten, Germany). Briefly, animals were anaesthetized by intraperitoneal injection with ketamine/xylazine (20 µl ketamine/20 µl xylazine/40 µl NaCl). After deep anaesthesia was achieved, animals were
tracheotomised, intubated and artificially ventilated with room air using the above mentioned system. Lung compliance was measured for the next at least ten minutes, after which animals were euthanized by a dose of pentobarbital (100-150 mg/kg body weight). All lung tissues were excised and snap frozen, or placed in 4% (w/v) paraformaldehyde and processed for paraffin embedding.

Assessment of lung fibrosis

Lung paraffin sections were stained with hematoxylin and eosin. For the quantitative histological analysis, a numeric fibrotic scale was used (Ashcroft score)\(^26\). Briefly, the grade of lung fibrosis was scored on a scale from 0 to 8 by examining more than 25 successive fields at a magnification of ×100 in a blinded fashion. The mean of all scores obtained from each field was employed as the fibrotic score of the specimen. To avoid observer bias, two experienced observers interpreted the images independently in a blinded fashion, and the mean of the observers' findings was considered to be the fibrotic score of the specimen.

Hydroxyproline assay

The whole collagen content of lungs was assessed by determining the hydroxyproline levels using HPLC. Briefly, lungs were homogenized in PBS, dried, and hydrolyzed in 6 N HCl at 110 °C for 24 h. Aliquots were added to 1.4% chloramine T (Sigma), 10% n-propanol, and 0.5 M sodium acetate, pH 6.0. After 20 min of incubation at RT, 1 M p-dimethylaminobenzaldehyde (Sigma) in 70% n-propanol / 20% perchloric acid was added and the sample incubated at 65°C. The absorbance was measured at 550 nm and the amount of hydroxyproline in each sample calculated against a standard curve.
SUPPLEMENT TABLES

Table S1. Characteristics of patients. VC = vital capacity, TLC = total lung capacity, DLCO/VA = diffusing capacity of the lung for CO per unit of alveolar volume (all in % predicted), PaO2/CO2 = partial pressure of O2 /CO2 in the arterial blood.

<table>
<thead>
<tr>
<th>No.</th>
<th>Diagnosis</th>
<th>Gender</th>
<th>Age (yr)</th>
<th>VC (%)</th>
<th>DLCO/VA (%)</th>
<th>TLC (%)</th>
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Table S2. Primer sequences and amplicon sizes for human and mouse tissues.

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SUPPLEMENT FIGURE LEGENDS

**Figure S1**
The mRNA expression profile of 5-HT-receptors (5-HTR) and the serotonin transporter 5-HTT in lung samples from bleomycin- or saline-treated mice 7 or 14 days after injury, as indicated. The mRNA levels of 5-HTR<sub>1a,1b</sub>, 5-HTR<sub>2a-c</sub>, and the serotonin transporter 5-HTT were assessed by quantitative real-time PCR (qRT-PCR). Results are derived from n = 4 per group and presented as fold change compared with the respective saline control, * p < 0.05.

**Figure S2**
Expression and localization of 5-HTR<sub>2B</sub> in lung tissues. Additional immunohistochemical stainings of donor and IPF patients using the HPA012867 antibody, as described in detail in Material and Methods. Scale bars indicate 100µm.

**Figure S3**
Application of the 5-HTR<sub>2A/B</sub> antagonist terguride in vivo. (A) The chemical structure of terguride. (B) Mice were subjected to a single inhalative instillation of bleomycin. After 7, 14, or 21 days, the 5-HTR<sub>2A/B</sub> antagonist terguride was applied intraperitoneally (i.p.) and the concentration of terguride in plasma and lung tissues were determined. (C) Body weight was assessed at the indicated time points.

**Figure S4**
The treatment scheme for the therapeutic approach in vivo. Mice were subjected to a single inhalative application of bleomycin, which led to the development of lung fibrosis by day 14 - 21 after an initial inflammatory phase (day 3 - 7). Treatment with terguride was initiated 14 days after bleomycin instillation, using the indicated concentrations, via intraperitoneal (i.p.) application twice daily until day 28. As an internal control, a treatment arm with daily
administration of 50 mg/kg BW imatinib was included in the study. Treatment of imatinib was started immediately following instillation of bleomycin (n=10 for each group).
Figure S1

- 5-Htr1a
- 5-Htr1b
- 5-Htr2a
- 5-Htr2b
- 5-Htr2c
- 5Htt

Fold change (2^ΔΔCt)

Bleomycin 7d  n=4
Bleomycin 14d  n=4

* * *
Figure S2

Donor

IPF

5-HTR2B
Figure S3

A

B

C

Effect of Bleomycin and Terguride on body weight (g) in different treatment groups:

- **Bleomycin + Terguride 0.4 mg/kg**
- **Bleomycin + Terguride 1.2 mg/kg**
- **Saline**

Plasma terguride concentration [pmol/ml] vs. days:

- **ctrl**
- **7d**
- **14d**
- **21d**

Terguride concentration in lung tissue [pmol/g wet weight] vs. days:

- **Plasma**
- **Lung**

Body weight (g) vs. days for different treatment groups:

- **Saline**
- **Bleomycin**
- **Bleomycin + Terguride 0.4 mg/kg**
- **Bleomycin + Terguride 1.2 mg/kg**
Figure S4

Bleomycin 1.5 U/kg

Non-treated (n=10)

Terguride 0.4 mg/kg (n=10)

Terguride 1.2 mg/kg (n=10)

Imatinib 50 mg/kg (n=10)

0    7    14    21    28 days

saline (n=10)