ORIGINAL ARTICLE

Exploration of a potent PI3 kinase/mTOR inhibitor as a novel anti-fibrotic agent in IPF

Paul F Mercer,1 Hannah V Woodcock,1 Jessica D Eley,1 Manuela Platé,1 Michal G Sulikowski,1 Pascal F Durrenberger,1 Linda Franklin,1 Carmel B Nanthakumar,2 Yim Man,2 Federica Genovese,3 Robin J McAnulty,1 Shuying Yang,2 Toby M Maher,4 Andrew G Nicholson,4 Andy D Blanchard,2 Richard P Marshall,2 Pauline T Lukey,2 Rachel C Chambers1

ABSTRACT

Rationale

Idiopathic pulmonary fibrosis (IPF) is the most rapidly progressive and fatal of all fibrotic conditions with no curative therapies. Common pathomechanisms between IPF and cancer are increasingly recognised, including dysfunctional pan-PI3 kinase (PI3K) signalling as a driver of aberrant proliferative responses. GSK2126458 is a novel, potent, PI3K/mammalian target of rapamycin (mTOR) inhibitor which has recently completed phase I trials in the oncology setting. Our aim was to establish a scientific and dosing framework for PI3K inhibition with this agent in IPF at a clinically developable dose.

Methods

We explored evidence for pathway signalling in IPF lung tissue and examined the potency of GSK2126458 in fibroblast functional assays and precision-cut IPF lung tissue. We further explored the potential of IPF patient-derived bronchoalveolar lavage (BAL) cells to serve as pharmacodynamic biosensors to monitor GSK2126458 target engagement within the lung.

Results

We provide evidence for PI3K pathway activation in fibrotic foci, the cardinal lesions in IPF. GSK2126458 inhibited PI3K signalling and functional responses in IPF-derived lung fibroblasts, inhibiting Akt phosphorylation in IPF lung tissue and BAL derived cells with comparable potency. Integration of these data with GSK2126458 pharmacokinetic data from clinical trials in cancer enabled modelling of an optimal dosing regimen for patients with IPF.

Conclusions

Our data define PI3K as a promising therapeutic target in IPF and provide a scientific and dosing framework for progressing GSK2126458 to clinical testing in this disease setting. A proof-of-mechanism trial of this agent is currently underway.

Trial registration number

NCT01725139, pre-clinical.

INTRODUCTION

Idiopathic pulmonary fibrosis (IPF) represents one of the most rapidly progressive fibrotic conditions with a median survival of 3.5 years from diagnosis,1 worse than many cancers. The last decade has seen an exponential rise in clinical trial activity in IPF with agents which exert pleiotropic effects, such as pirfenidone (Esbriet) and nintedanib (Ofev) recently granted approval as the first agents to slow disease progression.2 3 However, there still remains an urgent need to develop therapeutic strategies which halt rather than slow disease progression.

Fibrotic foci, the sentinel lesions in IPF, comprise fibroblasts and α smooth muscle actin positive (αSMA+) myofibroblasts embedded in a type I collagen-rich matrix. These lesions are thought to represent the leading edge of the relentless fibrotic response and arise as a result of a highly abnormal wound healing response characterised by dysregulated epithelial/mesenchymal crosstalk and a complex network of proliferation and differentiation signals received from a remodelled and homeostatically degenerate microenvironment4–7 (reviewed in8). Parallels between IPF and cancer are increasingly recognised, with active signalling pathways, transcriptomic and miRNA profiles and epigenetic signatures showing a considerable degree of overlap with lung cancer.9–11 Moreover, increased 18F-fluorodeoxyglucose (FDG) uptake, a marker of

Key messages

What is the key question?

▸ Is there a scientific rationale for the use of a clinically advanced pan-PI3 kinase (PI3K)/mTOR inhibitor in idiopathic pulmonary fibrosis (IPF)?

What is the bottom line?

▸ We provide a comprehensive scientific and dosing framework for targeting PI3 kinase signalling in IPF with the pan-PI3 kinase/mTOR inhibitor, GSK2126458, based on the exploration of novel functional endpoints in patient-derived cells and precision cut lung slices.

Why read on?

▸ This study provides key data to support the progression of this agent to a proof-of-mechanism trial in patients with IPF and further proposes a roadmap for the repositioning of existing oncology agents as novel therapeutic agents in the IPF setting with minimal requirement for the additional use of experimental animals.

To cite: Mercer PF, Woodcock HV, Eley JD, et al. Thorax Published Online First: [please include Day Month Year] doi:10.1136/thoraxjnl-2015-207429

Copyright Article author (or their employer) 2016. Produced by BMJ Publishing Group Ltd (& BTS) under licence.
aerobic glycolysis in cancer cells, was recently reported in the lungs of patients with IPF. Together these observations support the notion that a number of pharmacological agents developed for oncology might be explored as potential therapeutic agents in the setting of IPF.

Class 1 PI3K form an important oncogenic signalling node, integrating signals from a variety of inputs including tyrosine kinase receptors, G-protein coupled receptors and activated Ras, to mediate a variety of cellular processes, including cell cycle progression, growth and proliferation, metabolic and synthetic pathways in addition to inflammatory responses. There is evidence that class 1 PI3K isoforms, most notably p110, exhibit increased expression in IPF tissue and fibroblast lines, with signalling activated downstream of several key profibrotic growth factors implicated in IPF, including platelet-derived growth factor and transforming growth factor (TGF)-β. Evidence suggests that dysregulated PI3K signalling due to low activity of the tumour suppressor, phosphatase and tensin homologue (PTEN) may contribute to the profibrotic phenotype of IPF myofibroblasts, and moreover that inhibition of class 1 PI3K has anti-fibroproliferative effects in vitro and in tumour necrosis factor α induced fibrosis in vivo.

GSK2126458 is a potent, highly selective pyridilsulfonamide inhibitor of class 1 isozymes of PI3K in addition to mammalian target of rapamycin (mTOR). Originally developed in the oncology setting, GSK2126458 is currently (at the time of writing) being evaluated in a phase I open-label dose escalation trial in subjects with refractory solid tumours or lymphoma.

The aim of the present study was to evaluate the scientific and dose rationale for re-positioning GSK2126458 as a potential novel therapeutic agent for IPF. We explore evidence for PI3K pro-fibrotic signalling in IPF lung tissue in combination with performing in vitro pharmacological assessment of GSK2126458 and related development compounds in lung IPF and non-IPF control fibroblasts. We also assess inhibitor effects using novel functional endpoints in precision-cut IPF tissue slices, and further evaluate pAkt levels in IPF bronchoalveolar lavage (BAL) cells as a potential pharmacodynamic (PD) biomarker to monitor target engagement in future clinical studies. Integration of these data with pharmacokinetic (PK) data for GSK2126458 from oncology trials enabled modelling of an optimal dosing regimen for IPF and the progression of GSK2126458 to an IPF proof-of-mechanism (PoM) study.

METHODS

Patient material
All human samples were obtained with informed signed consent and with research ethics committee approval (10/H0504/9, 10/H0720/12 and 12/EM/0058). Patients with IPF were diagnosed in accordance with current international guidelines. Tissue for lung slice experiments was obtained from Asterand Europe (Royston, UK) in compliance with the UK Human Tissue Act 2004.

Primary fibroblast and bronchial epithelial cell culture
Primary human lung fibroblasts (LFs) were grown from explant cultures of IPF (IPF-LF) or non-IPF control (C-LF) lung tissue, and cultured as previously described. Primary human bronchial epithelial cells (HBECs) were isolated from regions of normal airway from IPF or non-IPF control lung tissue as previously described. See online supplementary material.

Immunohistochemistry
Immunostaining for pAkt, pAkt, αSMA and CD68 was conducted on 4 μm formalin-fixed paraffin-embedded serial sections of human lung biopsy material from n=12 patients with IPF, using the avadin–biotinylated HRP enzyme complex method (Vector Laboratories). Confocal dual-immunofluorescence microscopy for these markers was also performed on representative fibrotic foci. See online supplementary material.

Akt phosphorylation and collagen synthesis in precision-cut lung slices
Precision cut slices of IPF lung, 8 mm, were cultured for 24 h in DMEM supplemented with 0.4% FCS (5% CO2/100% humidity) prior to incubation with the PI3K inhibitors Compound 1 or GSK2126458. The effect of short-term exposure (2 h) to Compound 1 on Akt phosphorylation was assessed by Milliplex Map (Millipore) bead assay of lung slice homogenates prepared in ice cold PhoshoSafe buffer (Millipore). In separate experiments, the effect of GSK2126458 on levels of the collagen formation marker P1NP in tissue supernatant was assessed by proprietary competitive ELISA (Nordic Bioscience) as previously described. Slices were incubated with inhibitor for up to 5 days prior to harvest. See online supplementary material.

Fibroblast and BALF fluid cell phospho-proteins
The effect of GSK2126458 on FCS-induced Akt phosphorylation in C-LFs (n=2) and IPF-LFs (n=5) was assessed by Meso Scale Discovery (MSD) system. Phosphorylation events (pAkt and pSMAD2) in C-LFs were also assessed by Western blot. Densitometric analysis of Western blots was performed using ImageQuant TL v8.1 software (GE Healthcare). The intensity of bands in phospho-protein bands was divided by the intensity of the total protein following stripping and re-probing. Values were expressed as a ratio of the positive (stimulated) control. The effect of GSK2126458 on bronchoalveolar lavage fluid (BALF) cell pellet obtained from patients with IPF (n=6) undergoing routine diagnostic BAL was also assessed by MSD. See online supplementary material.

Fibroblast proliferation
The effect of GSK2126458 on FCS-induced proliferation in control (n=2) and IPF-derived primary human LFs (n=5) was assessed by MTS assay. Fibroblast proliferation was also assessed by 5-ethyl-2′-deoxyuridine (EdU) incorporation into DNA of fibroblast and BALF fluid cell phospho-proteins

Determination of collagen 1 gene expression and Pro-collagen synthesis
Coll1A1 gene expression in confluent fibroblast monolayers following TGFβ stimulation was assessed by RT-PCR following TRIzol extraction as per the manufacturer’s instructions (Thermo Fisher). RNA was DNase treated using a DNafree kit (Ambion) and in real-time RT-PCR conducted using the MESA GREEN qPCR MasterMix Plus for SYBR Assay (EugeneTec). Detailed methods are provided in the online supplementary material. Fibroblast pro-collagen production was assessed by high-performance liquid chromatography (HPLC) quantitation of hydroxyproline in supernatants from confluent fibroblast
monolayers, as previously described. Collagen biosynthesis and deposition was also assessed by molecular crowding assay modified from previously described methods using the IN Cell Analyzer 6000 high content imaging system (GE Healthcare Life Sciences). See online supplementary material.

RESULTS
Active PI3K signalling within IPF fibrotic foci
We first performed DAB-based immunohistochemistry (IHC) to explore evidence of functional PI3K signalling in areas of active fibrosis in IPF lung biopsy material (representative sections from Figure 1).

Akt phosphorylation co-localised within fibrotic foci, macrophages and bronchial epithelium in the idiopathic pulmonary fibrosis (IPF) lung. Immunofluorescence for α smooth muscle actin (αSMA, green, A and D), Akt phosphorylated at serine 473 (pAktS473, red, B) and threonine 308 (pAktT308, red, E) is shown co-localised (C and F) within fibrotic foci (A–F). Immunofluorescence for CD68 (green, G and J), pAktS473 (red, H), pAktT308 (red, K) is shown co-localised (I and L) within macrophages in airspaces (panels G–L). In panels M–R immunofluorescence for pAktS473 (red, N), pAktT308 (red, Q) is shown localised to unstained cells morphologically characteristic of bronchial epithelium (N, Q, O and R). Cells morphologically characteristic of airway smooth muscle are evident (green, M and P). DAPI-stained nuclei are highlighted in blue. 25 μm scale bars are shown.
n=12 patients shown), assessing the phosphorylation state of the critical downstream kinase, Akt. Online supplementary figure S1 shows prominent immunoreactivity for phosphorylated AktS473 and AktT308 localised to classic spindle-shaped, αSMA+ myofibroblasts and epithelium overlying fibrotic foci and cells morphologically characteristic of capillary endothelium (see online supplementary figure S1A,C,G). CD68+ macrophages residing in airspaces also exhibited signals for pAktS473 and pAktT308 (see online supplementary figure S1A,C,G). In contrast to myofibroblasts in fibrotic foci, airway smooth muscle cells demarcated by αSMA+ signal showed little evidence of Akt phosphorylation, while prominent phosphorylation was observed in the bronchial epithelium (see online supplementary figure S1B,D,F). A more detailed assessment of the cellular localisation of pAkt was provided by confocal dual immunofluorescence (figure 1). These studies confirmed the localisation of both pAktS473 and pAktT308 to αSMA+ myofibroblast subsets, however a differential staining pattern was observed in the overlying epithelium which preferentially exhibited signal for pAktS473 (figure 1A–F). In the bronchial epithelium, pAktS473 and pAktT308 showed differential subcellular distribution, with pAktS473 distributed apically to basolaterally throughout the cell layer, in comparison to pAktT308 which was predominantly apically located (figure 1M–R). αSMA+ airway smooth muscle was negative for pAkt. CD68+ macrophages exhibited clear but heterogeneous staining pattern for pAktS473 and pAktT308 (figure 1G–L).

Studies performed in live cultured precision-cut lung slices obtained from a patient with IPF confirmed prominent Akt phosphorylation in lung tissue and this signal was reduced in a concentration-dependent manner following incubation with a closely related analogue of GSK2126458, Compound 1 (figure 2) (see online supplementary figure S2 for compound structures and online supplementary table S1 for compound affinity data).

GSK2126458 inhibits Akt phosphorylation in IPF BALF cells

IHC studies (figure 1 and online supplementary figure S1) revealed that macrophages within IPF alveolar spaces were highly immunoreactive for pAkt. We therefore reasoned that since macrophages are a prominent cell type in IPF BALF, this observation could be exploited to develop a PD biosensor to monitor in vivo inhibition of PI3K signalling by GSK2126458.

BALF differential cell counts confirmed that macrophages formed a prominent cell type (n=6, see online supplementary table S5). Akt phosphorylation was observed for BALF cells obtained for all patients and incubation with GSK2126458 resulted in a concentration-dependent reduction in pAkt levels (figure 4 and see online supplementary table S6, geometric mean IC50=0.58 nM (95% CI 0.40 to 0.86)). The inhibitory profile of GSK2126458 was further found to be maintained in BALF cells re-suspended in ice-cold saline for up to 35 min following incubation with the drug (data not shown).

Integrated PK/PD models to guide dose selection of GSK2126458 in patients with IPF

To develop a dose rationale for GSK2126458 for future IPF clinical trials, the data from these in vitro studies were integrated with human PK data for GSK2126458 from ongoing clinical trials in patients with advanced solid tumours (http://www.gsk-clinicalstudyregister.com/study/112826#rs), according to the scheme outlined in figure 5. Stochastic model simulations were undertaken, providing predictions for pharmacological engagement over time for a range of doses. According to the model, the maximum tolerated dose of 2.5 mg twice a day dosing, GSK2126458 would likely provide enough exposure to significantly impact on the key PD endpoints, including Akt phosphorylation.
Figure 3  GSK2126458 inhibits Akt phosphorylation and proliferation in normal and idiopathic pulmonary fibrosis (IPF) primary human lung fibroblasts. Control (C-LF, n=2, A) or IPF lung fibroblasts (IPF-LF, n=5, C) were pre-incubated with increasing concentrations of GSK2126458. Following 30 min stimulation with 20% FCS, pAktS473 signal was normalised to total Akt and expressed as % maximum response (maximum asymptote) and is shown as mean±SEM of n=3 replicate wells per condition. A representative Western blot is shown visualising the effect of GSK2126458 on FCS-induced phosphorylation of AktS473 in a representative C-LF line (E). Subconfluent control (C-LF, n=2, B) or IPF lung fibroblasts (IPF-LF, n=5, D) were pre-incubated with increasing concentrations of GSK2126458. Following 72 h stimulation with 10% FCS, fibroblast proliferation was measured using an MTS assay at 490 nm. Data were expressed as % maximum response (maximum asymptote) and are shown as mean±n=6 replicate wells per condition. Incorporation of EdU (5-ethynyl-2′deoxyuridine) into the DNA of sub-confluent fibroblasts was also used as a measured of proliferation following 72 h of FCS stimulation (C-LF, n=5, dashed lines; IPF-LF, n=2, solid lines (F)). IC50 values (panels A–D and F) were calculated using four-parameter non-linear regression, and representative curves are shown for each fibroblast line. FCS stimulated and unstimulated DMSO vehicle control fibroblast data points are indicated as (+) and (−), respectively (panels A–F). DMSO was maintained at 0.1% for all conditions. Activation of fibroblast caspase 3 and 7 was assessed following incubation with increasing concentrations of GSK2126458 in the presence of 10% FCS by Caspase Glo (G). FCS-stimulated (+) vehicle control (0.1% DMSO) fibroblasts are indicated. Data are expressed as fold relative to baseline cells (no FCS, 0.1% DMSO) and shown as mean±SEM of n=3 replicate wells per condition. **p<0.0001 for differences in raw data values from FCS-stimulated controls (two-way ANOVA, Tukey’s multiple comparisons test).
phosphorylation and fibroblast proliferation in the lungs of patients with IPF (table 1).

Effect of GSK2126458 on epithelial cell apoptosis

The observation that the epithelium also displays a prominent pAkt signal lead us to further investigate the potential effect of GSK2126458 on epithelial cell apoptosis. Figure 6 illustrates that caspase 3,7 activity was induced in primary HBECs at concentrations of ≥3 μM (n=2 control donors, n=1 IPF donor).

GSK2126458 inhibits pro-fibrotic responses to TGFβ1 in primary human LFs in vitro

Progressive, excessive and disorganised deposition of collagen and other extracellular matrix (ECM) proteins underlies the gross remodelling leading to loss of lung function in IPF, with TGFβ1 implicated as a critical mediator of this process.5 Recent reports have further suggested the involvement of the PI3K signalling axis in SMAD-independent TGFβ1 signalling.15 28 We therefore also explored the potential of GSK2126458 to interfere with TGFβ1-induced fibroblast collagen production.

We first defined the kinetics of PI3K activation in relation to canonical SMAD signalling following TGFβ1 (1 ng/ml) stimulation. Figure 7A, B illustrates that exogenous TGFβ1 induces rapid phosphorylation of SMAD2 peaking at 2 h, with pAktS473 exhibiting a slower kinetic peaking at 12–24 h (figure 7D). Delayed Akt phosphorylation is in agreement with previous reports, suggesting the potential involvement of an autocrine mediator.15 Of note in our experiments, this peak precedes the peak of collagen (COL1A1) gene expression (36 h; figure 7C).

Table 1 Predicted pharmacodynamics of GSK2126458 in idiopathic pulmonary fibrosis patients by dose (Max: maximum inhibition (%), Min: minimum inhibition (%))

| Dose (mg), twice daily | pAkt/Akt Proliferation BAL |
|-----------------------|---------------------------|
|                       | Max | Min | Max | Min | Max | Min |
| 0.25                  | 14.3 | 8.3 | 24.5 | 19.2 | 24.2 | 12.4 |
| 1                     | 35.8 | 23.2 | 36.4 | 30.4 | 62.5 | 45.7 |
| 2                     | 49.6 | 34.9 | 42.5 | 37.1 | 76  | 65.2 |
| 2.5                   | 54.1 | 38.7 | 44.6 | 39.6 | 79  | 69.7 |

BAL, bronchoalveolar lavage.
We next assessed the effect of GSK2126458 on SMAD2 and Akt phosphorylation at 12 h after the addition of exogenous TGFβ3. Figure 7E shows that GSK2126458 had no influence on SMAD2 phosphorylation but inhibited TGFβ3-induced Akt phosphorylation in a concentration-dependent manner, with maximum inhibition obtained at 10 nM (figure 7E,F).

We next characterised the potency of GSK2126458 for inhibiting TGFβ3-induced collagen deposition in fibroblasts, using a novel high-content imaging based molecular crowding assay.27 Figure 8A illustrates that GSK2126458 attenuated TGFβ3-induced collagen deposition in fibroblasts, with an IC_{50} of 132.5 nM (95% CI 65.5 to 268), and leads to a reduction in cell number with an IC_{50} of 1.66 μM (95% CI 0.51 to 5.43). Induction of caspase 3,7 in molecular crowded conditions is consistent with concentrations of GSK2126458 which cause reduction in cell count, suggesting apoptosis in crowded conditions may be responsible for the observed reduction in cell number (see online supplementary figure S3). Collectively these data indicate a marked separation between the inhibitory effects of GSK2126458 on collagen deposition and induction of apoptosis. For comparison, the selective TGFβRI (ALK5) inhibitor, SB525334, inhibited TGFβ3-induced collagen deposition with an IC_{50} of 200.5 nM (95% CI 159 to 252; figure 8B). Images capturing the effects of inhibitors on primary fibroblasts isolated from IPF and control lungs in this assay are also shown (figure 8C).

Pan-PI3K/mTOR inhibition reduces collagen formation markers in ex vivo slices of IPF lung tissue

Our final aim was to explore evidence for a direct functional link between PI3K/mTOR signalling and matrix formation in this disease setting by monitoring the release of the human procollagen 1 amino-terminal peptide (P1NP) in IPF tissue slices using a competitive ELISA based on a proprietary monoclonal antibody.25 Figure 8D shows that the related GSK2126458 analogue, Compound 1, reduced P1NP levels released into lung slice supernatants, with maximal inhibition observed at >100 nM; the effect of a Alk5 inhibitor, SB525334 (10 μM), is also shown.

DISCUSSION

PI3K represents a key oncogenic signalling node, implicated in a range of cellular processes from inflammation to cancer. More recent evidence also suggests a role for this pathway in fibrosis, however to the best of our knowledge there are currently no PI3K agents in clinical development for IPF or any other fibrotic condition. The aim of the current study was to further explore the rationale for targeting PI3K signalling in IPF and to define the pharmacological profile of a potent and clinically advanced PI3K/mTOR inhibitor, GSK2126458, as a novel anti-fibrotic.

PI3K mediates the conversion of phosphatidylinositol 4,5 bisphosphate (PIP2) to phosphatidylinositol 3,4,5 trisphosphate (PIP3) at the plasma membrane, inducing the recruitment and phosphorylation of Akt at threonine 308 (pAkt^{T308}) and serine 473 (pAkt^{S473}) to initiate downstream signalling. The initial focus for our PI3K pathway validation studies in IPF was therefore to perform detailed IHC studies to determine Akt phosphorylation in IPF biopsy tissue. These studies showed demonstrable immunoreactivity for pAkt^{S473} associated with activated myofibroblasts within IPF fibrotic foci, as previously suggested.18 We further provide evidence for pAkt^{T308} immunostaining associated with these myofibroblasts and thus provide the first evidence of full Akt activation in situ in this disease context. Immunofluorescent confocal co-localisation studies further provide the first evidence of cell-specific differential phosphorylation of Akt: whereas pAkt^{T308} immunostaining predominates in myofibroblasts and a thin band of cells located at the epithelial–mesenchymal border of fibrotic foci, pAkt^{S473} immunostaining was evident in both the epithelium and myofibroblasts. Moreover, heterogeneous distribution and phosphorylation of Akt were observed in bronchial epithelium and macrophages in the IPF lung. Differential phosphorylation of Akt has not been extensively studied; however in cancer, it has been proposed to influence the interactions of Akt with its substrates and thereby influence cell motility and invasion.29 30 Of interest, in non-small cell lung cancer, elevated pAkt^{T308} relative to pAkt^{S473} is associated with poor prognosis.31 Future studies to determine the implications of differential phosphorylation of Akt in the context of IPF are beyond the scope of this article but may shed important light on the role of Akt in this disease setting. Confidence in PI3K as a target pathway was further extended by functional studies in IPF-derived lung tissue. Akt phosphorylation was readily detectable in precision-cut cultured IPF lung slices and this signal was inhibited by pan-PI3K/dual mTOR inhibition in a concentration-dependent manner.

Subsequent proliferation studies revealed that GSK2126458 inhibited serum-induced Akt phosphorylation and proliferation at nanomolar potency with no difference in compound potency observed. Of note, this anti-proliferative profile of GSK2126458 is comparable to that observed in previous in vitro oncology studies using the BT474 and T47D breast cancer lines.21 While pan-PI3K/mTOR inhibitors offer the advantage of overcoming functional redundancy between class I isoforms,32 and blocking potential crosstalk and feedback of compensatory mechanisms through inhibition of three key nodes (PI3K, mTORC1 and mTORC2), the prospect for mechanism-based on-target toxicities for this class of inhibitor, in particular with respect to inducing apoptosis, are well documented.33 In our studies, using primary LFs and bronchial epithelial cells, the anticipated pro-apoptotic effects associated with inhibiting...
PI3K/mTOR signalling were not manifest until concentrations of GSK2126458 were significantly higher than those required for exerting demonstrable anti-fibroproliferative and pro-apoptotic effects. This window is encouraging and suggests there is considerable potential for GSK2126458 to be administered at concentrations that would primarily impact on deleterious fibroblast functional responses in IPF.

Assessment of drug PDs in the lung of patients with IPF is challenging due to limitations associated with tissue access. Our IHC observation of pAkt immunostaining being associated with macrophages within airspaces raised the possibility that cells harvested from IPF BALF could serve as a potential PD biosensor for monitoring effective pharmacological engagement of mechanism; namely, pAkt inhibition in the lungs of patients recruited to early PoM studies. To this end, a BALF cell processing protocol was developed which involved minimal cell handling by assaying pAktS473 in whole BALF cell pellets on the basis that macrophages were a prominent cell type present. These data would also provide additional evidence from primary patient-derived lung cells to further support a dose rationale for GSK2126458. Interestingly, BALF cell pellets from patients with IPF exhibited pAktS473 activation without the need for exogenous stimulation and were sensitive to GSK2126458 inhibition, at sub-nanomolar concentrations.

Recent studies have implicated PI3K signalling in promoting M2 polarisation in IPF-derived BAL macrophages, raising the possibility that...
modulating macrophage biology might itself provide an additional benefit to GSK2126458 treatment in this disease context.

To build a model for dose prediction in IPF, pharmacological data obtained from fibroblast signalling and mitogenic assays were integrated with BALF cell PI3K phosphorylation data and existing PK data from previous human oncology trials. The model simulations predicted that within the typical clinical dose range (2–3 mg twice daily), GSK2126458 would significantly impact fibroblast PI3K signalling and proliferation in the lungs of patients with IPF.

The decision to progress GSK2126458 to a PoM trial in IPF was predicated on an appropriate dose rationale for this compound as a fibroblast anti-proliferative agent, based on the widespread evidence linking this pathway to cell growth and proliferative responses in multiple cell types and disease settings.38 However, in terms of key IPF pathomechanisms, we

---

**Figure 8** Pan-PI3 kinase (PI3K)/mammalian target of rapamycin (mTOR) inhibition attenuates transforming growth factor (TGF)-β1-induced collagen production in human lung fibroblasts (LFS), and collagen formation markers in idiopathic pulmonary fibrosis (IPF) lung tissue. Confluent LFs were incubated with increasing concentrations of GSK2126458 (A) or SB525334 (B) and stimulated for 48 h with TGFβ1 (1 ng/ml) and collagen biosynthesis assayed by molecular crowding assay (0.1% DMSO vehicle was constant for all experimental conditions). Data are expressed as mean fluorescent intensity (n=4 reads per well) and cell counts obtained from DAPI counterstaining (blue channel). Data are shown as mean±SEM of n=3 replicate wells per condition. IC50 values were calculated using four-parameter non-linear regression. Replicate experiments were carried out on human lung fibroblasts (HLFs) from a non-IPF donor and representative curves are shown. Images from each cell treatment are shown (C). 8 mm slices of IPF lung were cultured for 24 h in DMEM supplemented with 0.4% FCS and treated with vehicle, 10 μM SB525334 (open circles) or increasing concentrations of Compound 1 (inhibitor) for 0–3 days (filled grey circles) and further from 3 to 5 days (filled black circles) following media change. P1NP levels were measured in harvested supernatants (D). Data are shown as ±SEM of n=4 replicate wells per condition; *p<0.05, **p<0.01 (two-way ANOVA with Bonferroni post-hoc testing).
were also interested in determining the potential of this agent to inhibit TGFβ-induced myofibroblast collagen production, based on limited literature reports suggesting the involvement of PI3K in SMAD-independent TGFβ1 signalling. These studies revealed that GS2126458 was highly effective at attenuating TGFβ3-induced myofibroblast collagen deposition at concentrations around the 100 nM range.

Our final aim was to directly link PI3K/mTOR signalling to pro-fibrotic collagen formation in IPF lung tissue using an ex vivo human tissue model. Type I collagen is the most represented extracellular protein in the lungs and its dysregulated turnover is a hallmark of fibrosis. During maturation of type I collagen, soluble pro-collagen is processed to its insoluble form by carboxy and amino terminal endopeptidase cleavage, generating peptide fragments such as the N-terminal P1NP which can be monitored as a sensitive biomarker of collagen formation. Our studies revealed that P1NP was released from ex-vivo IPF precision-cut lung slices for up to 5 days in culture and further that PI3K/mTOR inhibition attenuated this process. To the best of our knowledge, these studies represent the first direct demonstration of a functional link between PI3K/mTOR signalling and ECM formation in this human disease setting. It is noteworthy that in isolated fibroblasts and in lung tissue slices, the pan-PI3K/mTOR inhibitors exhibited comparable efficacy to the selective TGFβRII (Alk–5) inhibitor, SB25334. This raises the possibility that GS2126458 could interfere with the fibrogenic effects of TGFβ1 signalling at clinically attainable doses, while potentially sparing its essential homeostatic and tumour suppressor functions. This is a particularly important consideration in patients with IPF who are at a heightened risk of developing lung cancer.

In conclusion, we have considerably extended the scientific rationale for targeting PI3K signalling in IPF using patient-derived cells and lung tissue. We provide evidence for active PI3K signalling in the fibrotic foci of IPF patients together with data that GS2126458 attenuates serum-induced proliferation and TGFβ1-induced collagen production in IPF fibroblasts. Population modelling and integration of preclinical biomarker data with human PK data has enabled us to develop a dosing framework which should safely and effectively engage PI3K signalling at clinically attainable doses, while potentially sparing its essential homeostatic and tumour suppressor functions. This is a particularly important consideration in patients with IPF who are at a heightened risk of developing lung cancer.

Acknowledgements The authors gratefully acknowledge Sharada Karaman (UCL), Susan Boyce, Alicia-Edmenson Cook, Milda Berahoe, Simon Peace and the Oncology Metabolism DPU (GSK) and Morten Karsdal (Nordic Biosciences) for their contribution to this study and preparation of this manuscript.

Contributors Conception and design: PFM, HVW, CBN, RJMa, TMM, ADB, RPM, PTL and RCC. Data acquisition: PFM, IDE, LF, MGS, PD, MP, YM and FG. Analysis and interpretation: PFM, CBN, SY, AGN, ADB, RPM, PTL and RCC.

Funding This work was funded by a GlaxoSmithKline collaborative framework agreement with University College London (UCL). The authors also acknowledge support from the Medical Research Council UK (MR/K020478/1), the Biotechnology and Biological Research Council (BB/L502133/1), the National Institute for Health Research University College London Hospitals Biomedical Research Centre and the National Institute for Health Research Royal Brompton Hospital Biomedical Research Unit.

Competing interests RCC declares that this work was funded as part of a collaborative framework agreement awarded to RCC by GlaxoSmithKline. CBN, YM, SY, ADB, RPM and PTL are employees of GlaxoSmithKline with a commercial interest in the area of investigation. FG is an employee of Nordic Biosciences with a commercial interest in proprietary assay development for biomarker discovery.

Patient consent Obtained.

Ethics approval UCLH and Imperial College London Ethics Committees.

Provenance and peer review Not commissioned; externally peer reviewed.

Open Access This is an Open Access article distributed in accordance with the terms of the Creative Commons Attribution (CC BY 4.0) license, which permits others to distribute, remix, adapt and build upon this work, for commercial use, provided the original work is properly cited. See: http://creativecommons.org/licenses/by/4.0/

REFERENCES

1 Lee AS, Mira-Avendano I, Ryu JH, et al. The burden of idiopathic pulmonary fibrosis: an unmet public health need. Respi Med 2014;108:955–67.
2 Riechel I, de Bois RM, Raghu G, et al. Efficacy and safety of nintedanib in idiopathic pulmonary fibrosis. N Engl J Med 2014;370:2071–82.
3 King TE Jr, Bradford WZ, Castro-Bernardini S, et al. A phase 3 trial of pirfenidone in patients with idiopathic pulmonary fibrosis. N Engl J Med 2014;370:2083–92.
4 Datta A, Scotton CJ, Chambers NC. Novel therapeutic approaches for pulmonary fibrosis. Br J Pharmacol 2011;163:70–7.
5 Akhurst RJ, Hata A. Targeting the TGFβ signalling pathway in disease. Nat Rev Drug Discov 2012;11:790–811.
6 Horan GS, Wood S, Ona Y, et al. Partial inhibition of integrin alpha(β)6 prevents pulmonary fibrosis without exacerbatng inflammation. Am J Respir Crit Care Med 2008;177:56–65.
7 Parker MW, Rossi D, Peterson M, et al. Fibrotic extracellular matrix activates a pro-fibrotic positive feedback loop. J Clin Invest 2014;124:1622–35.
8 Daniels CE, Jett JR. Does interstitial lung disease predispose to lung cancer? Curr Opin Pulm Med 2005;11:431–7.
9 Rabinovich EI, Kapetanaki MG, Steinfeld I, et al. Global methylation patterns in idiopathic pulmonary fibrosis. PLoS ONE 2012;7:e53770.
10 Selman M, Pardo A. Revealing the pathogenic and aging-related mechanisms of the enigmatic idiopathic pulmonary fibrosis... an integral model. Am J Respir Crit Care Med 2014;189:1611–72.
11 Vancheri C. Common pathways in idiopathic pulmonary fibrosis and cancer. Eur Respir Rev 2013;22:265–72.
12 Groves AM, Win T, Sreenan NJ, et al. Idiopathic pulmonary fibrosis and diffuse parenchymal lung disease: implications from initial experience with 18F-FOG PET/CT. J Nucl Med 2009;50:528–45.
13 Vanhaesebroeck B, Guillermot-Guibert J, Graupera M, et al. The emerging mechanisms of isoform-specific PI3K signalling. Nat Rev Mol Cell Biol 2010;11:329–41.
14 Conte E, Gilli E, Fruciano M, et al. PI3K p110γ overexpression in idiopathic pulmonary fibrosis lung tissue and fibroblast cells: in vitro effects of its inhibition. Lab Invest 2013;93:566–76.
15 Horowitz IC, Lee DY, Waghry M, et al. Activation of the pro-survival phosphatidylinositol 3 kinase/AKT pathway by transforming growth factor-beta1 in mesenchymal cells is mediated by p38 MAPK-dependent induction of an autocrine growth factor. J Biol Chem 2004;279:1359–67.
16 Kavanagh WM, Klippel A, Escobedo JA, et al. Modification of the 85-kilodalton subunit of phosphatidylinositol-3 kinase in platelet-derived growth factor-stimulated cells. Mol Cell Biol 1992;12:3415–24.
17 White ES, Attras RG, Hu B, et al. Negative regulation of myofibroblast differentiation by PTEN (Phosphatase and Tensin Homolog Deleted on chromosome 10). Am J Respir Crit Care Med 2006;173:112–21.
18 Xia H, Diebold D, Nho R, et al. Pathological integrin signaling enhances proliferation of primary lung fibroblasts from patients with idiopathic pulmonary fibrosis. Exp Med 2008;205:1659–72.
19 Conte E, Fruciano M, Fagone E, et al. Inhibition of PI3K prevents the proliferation and differentiation of human lung fibroblasts into myofibroblasts: the role of class I P110 isoforms. PLoS ONE 2011;6:e24663.
20 Le Cras TD, Kofragner TR, Davidson C, et al. Modification of PI3K by PX-866 prevents transforming growth factor-alpha-induced pulmonary fibrosis. Am J Pathol 2010;176:679–86.
21 Knight SD, Adams ND, Burgess JL, et al. Discovery of GS2126458, a Highly Potent Inhibitor of PI3K and the Mammalian Target of Rapamycin. ACS Med Chem Lett 2010;1:39–43.
22 Raghuv G, Collard HR, Egan JJ, et al. An official ATS/ERS/JRS/ALAT statement: idiopathic pulmonary fibrosis: evidence-based guidelines for diagnosis and management. Am J Respir Crit Care Med 2011;183:788–824.
23 Keerthisingam CB, Jenkins RG, Harrison NK, et al. Cyclooxygenase-2 deficiency results in a loss of the anti-proliferative response to transforming growth factor-beta in human fibroctic lung fibroblasts and promotes...
bleomycin-induced pulmonary fibrosis in mice. *Am J Pathol* 2001;158:1411–22.

24 Crowley C, Klanrit P, Butler CR, et al. Surface modification of a POSS-nanocomposite material to enhance cellular integration of a synthetic bioscaffold. *Biomaterials* 2016;83:283–93.

25 Leeming DJ, Larsen DV, Zhang C, et al. Enzyme-linked immunosorbent serum assays (ELISAs) for rat and human N-terminal pro-peptide of collagen type I (PINP)—assessment of corresponding epitopes. *Clin Biochem* 2010;43:1249–56.

26 Chambers RC, Laurent GJ, Westergren-Thorsson G. Cadmium inhibits proteoglycan and procollagen production by cultured human lung fibroblasts. *Am J Respir Cell Mol Biol* 1998;19:498–506.

27 Chen CZ, Peng YX, Wang ZB, et al. The Scar-in-a-Jar: studying potential antibacterial compounds from the epigenetic to extracellular level in a single well. *Br J Pharmacol* 2009;158:1196–209.

28 Hong M, Wilkes MC, Penheiter SG, et al. Non-Smad transforming growth factor-β signaling regulated by focal adhesion kinase binding the p85 subunit of phosphatidylinositol 3-kinase. *J Biol Chem* 2011;286:17841–50.

29 Bozulic L, Hemmings BA. PIKKing on PKB: regulation of PKB activity by phosphorylation. *Curr Opin Cell Biol* 2009;21:256–61.

30 Islam MR, Jones SJ, Macluskey M, et al. Is there a pAkt between VEGF and oral cancer cell migration? *Cell Signal* 2014;26:1294–302.

31 Vincent EE, Elder DJ, Thomas EC, et al. Akt phosphorylation on Thr308 but not on Ser473 correlates with Akt protein kinase activity in human non-small cell lung cancer. *Br J Cancer* 2011;104:1755–61.

32 Foukas LC, Berenjeno IM, Gray A, et al. Activity of any class IA PI3K isoform can sustain cell proliferation and survival. *Proc Natl Acad Sci USA* 2010;107:11381–6.

33 Fruman DA, Rommel C. PI3K and cancer: lessons, challenges and opportunities. *Nat Rev Drug Discov* 2014;13:140–56.

34 Gibbons MA, Mackinnon AC, Ramachandran P, et al. Ly6Chi monocytes direct alternatively activated profibrotic macrophage regulation of lung fibrosis. *Am J Respir Crit Care Med* 2011;184:569–81.

35 Murray LA, Rosada R, Moreira AP, et al. Serum amyloid P therapeutically attenuates murine bleomycin-induced pulmonary fibrosis via its effects on macrophages. *PloS ONE* 2010;5:e9683.

36 Prasse A, Pechkovsky DV, Toews GB, et al. A vicious circle of alveolar macrophages and fibroblasts perpetuates pulmonary fibrosis via CCL18. *Am J Respir Crit Care Med* 2006;173:781–92.

37 Stahl M, Schupp J, Jäger B, et al. Lung collagens perpetuate pulmonary fibrosis via CD204 and M2 macrophage activation. *PloS ONE* 2013;8:e81382.

38 Marone R, Cmiljanovic V, Giese B, et al. Targeting phosphoinositide 3-kinase: moving towards therapy. *Biochim Biophys Acta* 2008;1784:159–85.

39 Anderton MJ, Mellor HR, Bell A, et al. Induction of heart valve lesions by small-molecule ALK5 inhibitors. *Toxicol Pathol* 2011;39:916–24.