

# Nintedanib: A Novel Therapeutic Approach for Idiopathic Pulmonary Fibrosis

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**Idiopathic pulmonary fibrosis (IPF) is a chronic, progressive, fibrotic lung disease with no clear etiology and few therapeutic options. Growth factors that act as mediators in the development of this disease might be important therapeutic targets. Nintedanib is a triple-tyrosine kinase inhibitor and a potent antagonist of growth factors such as platelet-derived growth factor, vascular endothelial growth factor, and basic fibroblast growth factor, and it is currently evaluated in clinical trials as a potential IPF therapy. Treatment with nintedanib may slow decline in lung function, decrease the frequency of exacerbations, and improve quality of life in subjects with IPF. This observation, together with extensive safety and pharmacokinetic data from studies of nintedanib in malignancy, led the way for the clinical development of this drug in IPF. Observations from clinical trials, together with the preclinical data, suggest that nintedanib may become an important therapeutic option for individuals with IPF. High-dose nintedanib improved the quality of life, slowed the progression of lung fibrosis and the decline of lung function, and reduced the rate of exacerbations in individuals with mild and moderate IPF. This is a short review based on the available data (September 2013) on nintedanib. Key words: interstitial pneumonias; idiopathic pulmonary fibrosis; growth inhibitors; mediators; BIBF 1120; nintedanib. [Respir Care 2014;59(9):1–•. © 2014 Daedalus Enterprises]**

## Introduction

Idiopathic pulmonary fibrosis (IPF; also called cryptogenic fibrosing alveolitis) is a specific form of chronic, progressive, fibrosing interstitial pneumonia of unknown cause that occurs in adults and is limited to the lungs. It is associated with the histopathologic and/or radiologic pattern of usual interstitial pneumonia. In the past, treatment was aimed at minimizing inflammation and slowing the progression from inflammation to fibrosis. However, the underlying lesion in IPF may be more fibrotic than inflammatory, explaining why few subjects respond to anti-inflammatory therapies, and prognosis remains poor.<sup>1,2</sup>

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No therapy has been proven to be efficacious, and management generally includes some combination of supportive care (eg, supplemental oxygen, pulmonary rehabilitation), consideration for participation in clinical trials introducing the use of novel drugs, consideration for lung transplant evaluation, and identification and treatment of possible comorbidities.<sup>1,3</sup> Supportive care may be preferred as the sole treatment option for many patients given the lack of proven therapy. Prevention of gastroesophageal reflux and recurrent microaspiration may slow disease progression. A number of other agents (eg, anticoagulation,<sup>4</sup> colchicine,<sup>5</sup> cyclophosphamide,<sup>6</sup> endothelin receptor antagonists,<sup>7</sup> interferon gamma-1b,<sup>8</sup> methotrexate,<sup>9</sup> cyclosporine,<sup>10</sup> penicillamine,<sup>11</sup> imatinib,<sup>12</sup> etanercept,<sup>13</sup> N-acetylcysteine,<sup>14</sup> prednisone/azathioprine/N-acetylcysteine combination<sup>15,16</sup>) have been used in the past in either case series or clinical trials. The process of fibrosis is driven by innumerable growth factors and their downstream intracellular signaling pathways. It is therefore not surprising that multiple cell signaling pathways have been identified as potential therapeutic targets in IPF.<sup>2,17</sup> When evaluated in a clinical study of subjects with IPF, imatinib given once daily at a dosage of 600 mg/d for 96 weeks demon-

strated no significant benefit over placebo in slowing disease progression or lung function impairment.<sup>12</sup> Imatinib, sometimes referred to by its investigational name STI-571, is a tyrosine kinase inhibitor used in the treatment of multiple cancers, most notably Philadelphia chromosome-positive (Ph<sup>+</sup>) chronic myelogenous leukemia. Like all tyrosine kinase inhibitors, imatinib works by preventing a tyrosine kinase enzyme, in this case BCR-ABL, from phosphorylating subsequent proteins and initiating the signaling cascade necessary for cancer development, thus preventing the growth of cancer cells and leading to their death by apoptosis. Because the BCR-ABL tyrosine kinase enzyme exists only in cancer cells and not in healthy cells, imatinib works as a form of targeted therapy: only cancer cells are killed through the drug's action. In this regard, imatinib was one of the first cancer therapies to show the potential for such targeted action and is often cited as a paradigm for research in cancer therapeutics. Imatinib is used to treat chronic myelogenous leukemia, gastrointestinal stromal tumors, and a number of other malignancies. One study demonstrated that imatinib mesylate was effective in patients with systemic mastocytosis, including those who had the D816V mutation in c-Kit.<sup>18</sup> However, since imatinib binds to tyrosine kinases when they are in the inactive configuration and the D816V mutant of c-Kit is constitutively active, imatinib does not inhibit the kinase activity of the c-Kit D816V mutant. Experience has shown, however, that imatinib is much less effective in patients with this mutation, and patients with the mutation compose nearly 90% of cases of mastocytosis.<sup>19-21</sup>

Pirfenidone shows promise and is the only licensed treatment currently available for individuals with IPF.<sup>3,22-25</sup> Pirfenidone has antifibrotic and anti-inflammatory properties in various *in vitro* systems and animal models of fibrosis. A number of cell-based studies have shown that pirfenidone reduces fibroblast proliferation, inhibits transforming growth factor beta-stimulated collagen production, and reduces the production of fibrogenic mediators such as transforming growth factor beta. Pirfenidone has also been shown to reduce production of inflammatory mediators such as tumor necrosis factor alpha and interleukin-1beta in both cultured cells and isolated human peripheral blood mononuclear cells. These activities are consistent with the broader antifibrotic and anti-inflammatory activities observed in animal models of fibrosis. A recent Cochrane review concluded that pirfenidone appears to improve progression-free survival and, to a lesser effect, pulmonary function in patients with IPF.<sup>23</sup> In Europe, pirfenidone is indicated for the treatment of mild-to-moderate IPF. It was approved by the European Medicines Agency in 2011. It was approved for use in Japan in October 2008 and in India in 2010.

## Preclinical Data

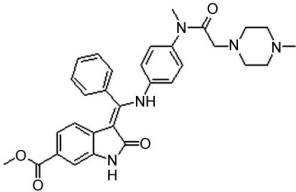
Nintedanib (or intedanib), formally known by the development code BIBF 1120 (or BIBF-1120 or BIBF1120), is a potent, oral, small-molecule, intracellular inhibitor of the following receptor tyrosine kinases: platelet-derived growth factor (PDGF) receptors alpha and beta (IC<sub>50</sub> = 59 and 65 nmol/L); vascular endothelial growth factor (VEGF) receptors 1, 2, and 3 (IC<sub>50</sub> = 13–34 nmol/L); and fibroblast growth factor (FGF) receptors 1, 2, and 3 (IC<sub>50</sub> = 137–108 nmol/L).<sup>26</sup> Nintedanib competitively binds the ATP-binding site of its target receptor tyrosine kinase.<sup>27</sup> The exact receptor-binding kinetics are not completely understood; however, it has been shown that nintedanib causes sustained (> 32 h) inhibition of VEGF receptor (VEGFR)-2 phosphorylation.<sup>26</sup> At present, only limited data have been published on the effect of BIBF on fibrogenesis either *in vitro* or *in vivo* in animal models. The main published work assessing tyrosine kinase receptor inhibition in fibrosis utilized an analog of nintedanib (BIBF 1000) and found that 50 mg of BIBF 1000/kg, when dosed both prophylactically and therapeutically, inhibits the development of fibrosis in the rat bleomycin model.<sup>28</sup>

Although the exact role of angiogenesis in the fibrotic process is still debated, increased angiogenic activity (secondary to an imbalance in angiogenic and angiostatic factors) is seen in human fibrotic lung tissue.<sup>29</sup> PDGF and FGF act synergistically to induce endothelial cell proliferation and contribute to pericyte recruitment and stability of blood vessel walls.<sup>30</sup> Early administration of a VEGFR-2 inhibitor in the murine bleomycin model attenuates fibrosis, microvessel formation, and bronchoalveolar lavage inflammatory cell counts.<sup>31</sup> *In vitro*, nintedanib inhibits VEGF-, PDGF-, and FGF-mediated proliferation of the 3 cell types contributing to angiogenesis: endothelial cells, vascular smooth muscle cells, and pericytes (EC<sub>50</sub> = 10–79 nmol/L).

## Clinical Data

Nintedanib is an orally available 6-methoxycarbonyl-substituted indolinone (derived from indoline) (Table 1) that acts as a multiple-receptor tyrosine kinase inhibitor and that has recently shown promising results in a phase-2 trial in IPF.<sup>32,33</sup> Originally developed as a cancer treatment, nintedanib acts by simultaneously inhibiting 3 receptor families implicated in angiogenesis: PDGF, VEGF, and FGF.<sup>26</sup> Single angiogenic inhibitors have had limited success in clinical practice due to redundancy in angiogenic pathways. Multiple-target receptor tyrosine kinase inhibitors, including nintedanib, have been developed to address this problem.<sup>27</sup> VEGF was found to be overexpressed by tumor cells in response to tissue hypoxemia and was reported to increase proliferation and migration of

Table. The Identity of Nintedanib

CAS number: 656247-17-5 UNII: G6HRD2P839	Systemic Name	Trade Name	Route	Formula	Molecular Mass	Appearance	Purity
	Methyl (3Z)-3- {[(4-{methyl[(4- methylpiperazin-1-yl) acetyl]amino}phenyl) amino](phenyl)methylidene)- 2-oxo-2,3-dihydro-1H- indole-6-carboxylate	Vargatef (Boehringer Ingelheim)	Oral and intravenous	C <sub>31</sub> H <sub>33</sub> N <sub>5</sub> O <sub>4</sub>	539.6248 g/mol	Yellow	> 99%

endothelial cells and to inhibit their apoptosis.<sup>34,35</sup> PDGF is another growth factor that was found to be involved in the pathogenesis of various proliferative disorders such as tumors or pulmonary fibrosis, in which it was found to stimulate proliferation of smooth muscle cells as well as fibrogenesis.<sup>34,36</sup> FGF was demonstrated to induce proliferation of smooth muscle cells, myofibroblasts, and fibroblasts, and its expression was found to be up-regulated in IPF.<sup>35</sup> PDGF, VEGF, and FGF are also critical profibrotic mediators that have been shown to play a role in driving the development of fibrosis.<sup>28</sup> The role of VEGF in IPF is contradictory. A heterogeneity of vascular remodeling in IPF has been reported, with increased vascular density in areas with low-grade fibrosis and decreased vascular density in the most extensively fibrotic lesions.<sup>37</sup> It has been shown that there is increased expression of VEGF in capillary endothelial cells and alveolar type II epithelial cells in highly vascularized alveolar septa. In contrast, fibroblasts and leukocytes in fibrotic lesions are weakly immunoreactive with VEGF, suggesting a possible role for VEGF in the vascular heterogeneity of IPF.<sup>37</sup> The question according to these findings is whether the increase in vascular density observed in the least fibrotic areas is actively a consequence of the development of the fibrogenic process or represents a compensatory mechanism.<sup>38</sup> The role of VEGF in this process needs to be clarified further. Plasma VEGF concentrations do not differ between patients with IPF and controls. However, baseline plasma levels of VEGF<sup>39</sup> are significantly related to the extent of parenchymal involvement in high-resolution computed tomography, and patients with IPF who develop progressive disease have significantly higher baseline levels of VEGF.<sup>39</sup> In contrast, bronchoalveolar lavage fluid concentrations of VEGF are significantly depressed in patients with IPF<sup>40-42</sup> and correlate with diffusing capacity of the lung for carbon monoxide (D<sub>LCO</sub>). The latter correlation possibly reflects the diminished epithelial surface area versus the diminished gene expression or intraluminal secretion of VEGF.<sup>42</sup>

### Pharmacology

In vitro, nintedanib is rapidly metabolized by hepatocytes via ester cleavage to form the metabolite BIBF 1202.

In healthy volunteers, BIBF 1202 was detected along with the parent compound 15 min after nintedanib administration, with maximum concentration for BIBF 1202 being reached at 2.5 h. The authors speculate that the rapid appearance of the metabolite results from first-pass metabolism in the intestinal wall.<sup>43</sup> The major route of metabolite elimination is via the liver and through excretion in feces. There is minimum excretion in the urine.<sup>44</sup> The hepatic metabolism of nintedanib is (mainly) CYP450-independent.<sup>45,46</sup> There have been no reported drug-drug interactions in clinical trials on malignancy.

### Safety

Data from studies on malignancy show that the most frequent adverse events reported in the TOMORROW trial (To improve pulMONaRy fibROSis With BIBF 1120) were diarrhea (27%), nausea (14.5%), and vomiting (7.7%).<sup>32</sup> Serious adverse event rates were similar between groups (subjects vs controls), although the discontinuation rate due to adverse events was highest in the 150 mg twice daily group at 30.6%. This rate is in comparison to 25.9% in the placebo group and 14.0% in the 100 mg twice daily group. The majority of the reported diarrhea cases were mild or moderate. The rate of gastrointestinal side effects rose with increasing doses of nintedanib, with 8.2% of subjects in the maximum dosing arm reporting serious diarrhea compared to 1.2 and 0% in the 50 mg twice daily and placebo arms, respectively. Only 12 subjects (across all arms) discontinued the drug due to diarrhea.

Elevation of liver enzymes (reversible) occurred more frequently in subjects receiving 300 mg of nintedanib/d compared to placebo. There were, however, only 9 clinically important episodes of liver enzyme elevation (3 times the upper limit of the normal range) across all of the treatment arms. Only 2 subjects needed to discontinue the drug due to hepatotoxicity, and there were no cases of drug-induced liver failure. All liver function tests normalized with either reduction or discontinuation of nintedanib. Ongoing safety studies include an extension of the TOMORROW trial (ClinicalTrials.gov identifier NCT01170065) and a study, in Japanese subjects, of nintedanib given together with pirfenidone (identifier NCT01417156).<sup>47,48</sup>

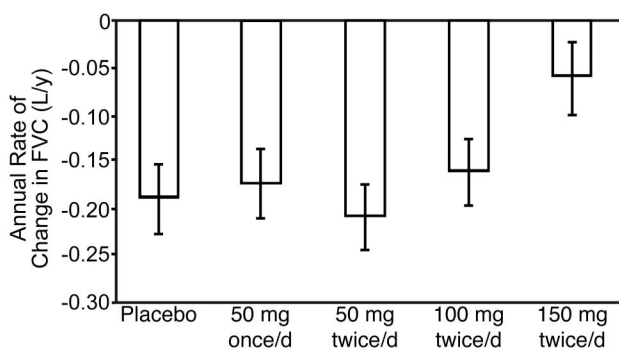


Fig. 1. Annual rate of change in FVC following the administration of nintedanib (TOMORROW trial). Data are expressed as mean  $\pm$  SD.

### Clinical Trials

The TOMORROW study was a 12-month, randomized, placebo-controlled, phase-2 study evaluating the efficacy and safety of 4 nintedanib doses (50 mg once daily, 50 mg twice daily, 100 mg twice daily, 150 mg twice daily) in subjects with IPF.<sup>33</sup> Subjects were at least 40 y old with a predefined IPF diagnosis of < 5 y prior to the study screening. Eligible subjects had FVC  $\geq$  50%, predicted  $D_{LCO}$  30–79%, and  $P_{aO_2} \geq$  55 mm Hg. Concomitant oral therapy with prednisone (or equivalents) at  $\leq$  15 mg was administered if the patient was stable during the 8 weeks previous to study enrollment.<sup>32</sup> The primary efficacy end point was represented by the FVC decline rate, and the secondary end points included changes from baseline in FVC,  $D_{LCO}$ ,  $S_{pO_2}$ , total lung capacity, exercise capacity, St George Respiratory Questionnaire (SGRQ) scores, incidence of acute exacerbations, and overall mortality. The highest BIBF 1120 dose was associated with the most significant therapeutic effect on the lung function decline compared to placebo, with the drug reducing the annual rate of lung function decline by 68.4% compared to the placebo group. The highest dose of BIBF 1120 therapy was also associated with a lower percentage of subjects exhibiting a significant reduction of FVC (of > 10% or > 200 mL) compared to the placebo group (23.8 vs 44%,  $P = .004$ ). Unlike the placebo, BIBF 1120 preserved the total lung capacity ( $-0.24$  vs  $0.12$  L,  $P < .004$ ) (Fig. 1). Mean change from baseline in  $S_{pO_2}$  was  $-0.2\%$  with BIBF 1120 and  $-1.3\%$  with placebo ( $P = .02$ ). The highest dose therapy was also associated with a lower percentage of significant desaturation (> 4% reduction from baseline in resting  $S_{pO_2}$ ) over the study period (3.6% for BIBF 1120 and 11.0% for placebo,  $P = .03$ ). BIBF 1120 did not exert a significant therapeutic benefit on  $D_{LCO}$  and on the exercise capacity compared to placebo. Health-related quality of life evaluated with the SGRQ was found to be significantly improved with the highest BIBF dose compared to placebo. The difference was also clinically important: mean change

of  $-0.66$  points with the active treatment compared to 5.46 points with placebo ( $P = .007$ ). The most significant therapeutic benefit was detected in the SGRQ score: 9.6-unit improvement with BIBF 1120 compared to placebo ( $P = .003$ ).<sup>49</sup> The highest BIBF 1120 dose was associated with a significant reduction in the incidence of acute exacerbations compared to placebo (2.4 vs 15.7 per 100 subject years,  $P = .02$ ). The 150- and 100-mg BIBF 1120 doses were also associated with a trend toward a lower mortality rate compared to placebo ( $P = .04$  for 100 mg and  $P = .06$  for 150 mg). No significant differences in terms of mortality rates in the treatment groups compared to placebo were reported. Two phase-3 studies aimed at evaluating the efficacy and safety of BIBF 1120 at 150 mg twice daily for 52 weeks in subjects with IPF are also planned. These 2 studies have similar end points, which include the annual decline rate in FVC, quality of life, time to first exacerbation, and overall survival.<sup>50</sup>

### Future Directions

The results of the TOMORROW study were sufficiently positive for Boehringer Ingelheim (who introduced BIBF 1120) to undertake 2 parallel phase-3 registration studies of nintedanib in IPF (ClinicalTrials.gov identifiers NCT01335464 and NCT01335477). These studies, IMPULSIS I and II, are identical 52-week trials of nintedanib administered at 150 mg twice daily compared to placebo. As with the TOMORROW study, the primary end point is annual rate of decline in FVC (expressed in mL over 52 weeks). Target recruitment was  $\sim$ 550 subjects for each study. Both trials were initiated in April 2011, and it is anticipated that the final study visit will be conducted in the first half of 2014. It is to be hoped, therefore, that results will be available later in 2014.

The TOMORROW study represents an important advance in the treatment of IPF.<sup>43,51,52</sup> Compared with other treatment approaches, including the administration of antioxidants such as N-acetylcysteine and of pirfenidone, the nonspecific suppression of the inflammatory response with systemic glucocorticoids and potent immunosuppressive agents such as azathioprine and cyclophosphamide, and the use of antifibrotic cytokines such as interferon gamma-1b, phosphodiesterase-5 inhibitors, and endothelin receptor antagonists, the beneficial effects of BIBF 1120 on the selected population seem to be far better. Newer inhibitors of fibrogenic pathways now being developed may have the potential to produce even more effective treatments that selectively target fibrogenic pathways without affecting the immune and inflammatory responses.

### Summary

IPF is a debilitating, progressive, and possibly fatal disease. Encouraging data suggest that targeting PDGF, VEGF,



and FGF receptors slows the development of fibrosis. The multiple targeting of these 3 mediators by the novel tyrosine kinase inhibitor nintedanib has led to promising clinical trial results. Preclinical and clinical data demonstrated that high-dose oral nintedanib is able to slow the progression of lung fibrosis and the decline of lung function and reduce the rate of exacerbations in individuals with mild and moderate IPF. Importantly for subjects with this chronic progressive disease, nintedanib improves the quality of life compared with placebo, and it is safe to use. The results of more clinical trials with nintedanib are eagerly awaited to determine the role of nintedanib in the management of IPF.

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