

Utility of nintedanib for severe idiopathic pulmonary fibrosis: a single-center retrospective study [Response to letter]

This article was published in the following Dove Press journal:
Drug Design, Development and Therapy

Mitsuhiro Abe¹
Kenji Tsushima^{1,2}
Koichiro Tatsumi¹

¹Department of Respiriology, Graduate School of Medicine, Chiba University, Chuo-ku, Chiba city, Chiba 260-8670, Japan; ²Department of Pulmonary Medicine, International University of Health and Welfare, School of Medicine, Narita city, Chiba 286-8686, Japan

Dear editor

In our study, we observed that even in Severe Group of patients with interstitial pulmonary fibrosis (IPF), nintedanib administration suppressed the reduction in the forced vital capacity (FVC) (Figure 3).¹ However, the frequency of side effects tended to be more (Table 2) and the prognosis was significantly worse in Severe Group than in Mild Group (Figure 5).

Orsatti et al pointed out three limitations of our research.² First, the two study groups presented clinically meaningful differences at baseline. However, we found that even in Severe Group, FVC reduction was suppressed by nintedanib administration ($p=0.029$) (Figure 3). The INPULSIS-ON trial revealed that nintedanib administration in IPF patients with a more advanced functional impairment may have the same beneficial effect on FVC reduction as among patients with less severe impairment. However, we believe that it is important to demonstrate a beneficial effect among patients with severe IPF in the real world.

Second, a higher mortality rate among patients with severe IPF is expected independent of the pharmacological intervention; lower FVC and diffusing capacity of the lungs for carbon monoxide (D_{LCO}) seen at baseline among patients with severe IPF were associated with a worse prognosis. Clearly, patients with more severe disease have a worse prognosis. We suggested in our paper that patients with severe disease should start treatment earlier considering the poor prognosis.

Furthermore, our study revealed that patients with severe IPF had a greater FVC decline than those with mild-to-moderate IPF in the year preceding nintedanib administration. A more severe decline in FVC over time has also been associated with a worse prognosis. Hence, even in patients with severe IPF who experienced a more severe decline in FVC, we believe that it is important to clarify the usefulness of nintedanib in the real world.

We agree that it is important to conduct a multi-center prospective clinical trial. However, it may not often be feasible to compare treatment effect with a control group in the real world; retrospective research is also important in such circumstances.

Disclosure

KTs has received lecture fee from Boehringer Ingelheim. The other authors report no conflicts of interest in this communication.

Correspondence: Mitsuhiro Abe
Department of Respiriology, Graduate School of Medicine, Chiba University, 1-8-1 Inohana, Chuo-ku, Chiba 260-8670, Japan
Tel +8 143 226 2576
Email mthrsngm@chiba-u.jp

References

1. Abe M, Tsushima K, Sakayori M, et al. Utility of nintedanib for severe idiopathic pulmonary fibrosis: a single-center retrospective study. *Drug Des Devel Ther*. 2018;12:3369–3375.
2. Orsatti L, Fortea J, Quaresma M. Utility of nintedanib for severe idiopathic pulmonary fibrosis: a single-center retrospective study 45 [Letter]. *Drug Des Devel Ther*. 2019;13: 1177—1178.

Drug Design, Development and Therapy

Dovepress

Publish your work in this journal

Drug Design, Development and Therapy is an international, peer-reviewed open-access journal that spans the spectrum of drug design and development through to clinical applications. Clinical outcomes, patient safety, and programs for the development and effective, safe, and sustained use of medicines are a feature of the journal, which has also

been accepted for indexing on PubMed Central. The manuscript management system is completely online and includes a very quick and fair peer-review system, which is all easy to use. Visit <http://www.dovepress.com/testimonials.php> to read real quotes from published authors.

Submit your manuscript here: <https://www.dovepress.com/drug-design-development-and-therapy-journal>