

**Supplementary Table S1.** Additional entry criteria not described within the text

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**Description of inclusion criteria**

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- For women of childbearing potential: agreement to remain abstinent (refrain from heterosexual intercourse) or use two adequate methods of contraception, including at least one method with a failure rate of <1% per year, during the treatment period and for at least 3 months after the final follow-up visit
  - For men: agreement to remain abstinent (refrain from heterosexual intercourse) or use contraceptive measures and agreement to refrain from donating sperm
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**Description of exclusion criteria**

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- Clinical evidence of active infection
  - Any condition that is likely to result in death in the 12 months after the start of screening
  - Lung transplantation anticipated or any planned significant surgical intervention
  - Known hypersensitivity to the active substance or any excipient of either pirfenidone or nintedanib
  - Hepatic impairment and/or severe renal impairment
  - History of GI tract perforation, unstable or deteriorating cardiac or pulmonary disease (other than IPF), long QT syndrome, alcohol or substance abuse, use of any tobacco product, etc
  - Bleeding risk
  - Pregnancy or lactation
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GI, gastrointestinal; IPF, idiopathic pulmonary fibrosis; QT interval, time between Q and T waves.