



# Action for Pulmonary Fibrosis

We're Action for Pulmonary Fibrosis. We exist to help find a cure for pulmonary fibrosis and for everyone affected by the disease to have a better future. We provide support and information, raise awareness, campaign and educate to improve access to the highest standard of care for people living with all forms of pulmonary fibrosis. We're committed to funding research to drive the search for a cure.

As you will be aware, Covid has significantly exacerbated problems with timely diagnosis and vital care for people with PF. Forthcoming changes (such as the wider range of people becoming eligible for anti-fibrotic treatment) will bring significantly more PF patients to the attention of specialist care. This is a positive development and a huge logistical problem which services currently feel unequipped to tackle.



We are inviting ILD professionals to come together in a round table discussion; sharing your perspectives, experience and insight. We will challenge ourselves to do our best thinking. We are stronger together.

evidence  
workforce  
data  
virtual care  
collaborate  
anti-fibrotics  
expert  
rehabilitation  
vital  
patient voice  
partnerships  
hopeful  
mental health  
research  
pathways  
palliative care  
determined

*What are your hopes and fears for Pulmonary Fibrosis patients in 2022 and beyond?*

*What are you aware of that is already being done to improve, mitigate, or prepare?*

*What data, evidence or insight do you have? What do you need?*

*How might we approach these issues creatively, boldly, collaboratively and nationally to find and work toward potential solutions?*

## We'd like to invite you to be part of the conversation.

Join us for a round table discussion on Zoom  
Tuesday 18 January 2022, 7.45-9.15pm

RSVP to Louise Wright, Chief Executive of Action for Pulmonary Fibrosis,  
on [louise@actionpulmonaryfibrosis.org](mailto:louise@actionpulmonaryfibrosis.org)