Welcome to EU-IPFF’s Newsletter March 2022

Sponsor News

Welcome to a new member of the EU-IPFF

Editorial by President of EU-IPFF Steve Jones

Welcome to our first newsletter of 2022 which looks back on 2021 and details some of our plans for 2022. We look forward to sharing news on the impact of COVID-19 on our patient communities, the progress we have made in the fight against rare diseases, and the work we are doing to support and empower patients across Europe.

I hope you all enjoy this newsletter and will continue to support us in our mission. Whether you are a patient, carer, or healthcare professional please do the work by the EURoDis, European Pulmonary Fibrosis Patient Summit, which will take place virtually on 3rd-5th November 2022.

As a federation dedicated to improving the lives of the people affected by pulmonary fibrosis, this e-newsletter seeks to provide you with updates on our activities, events, and the latest developments in the field. It is a platform to inform the broader community about the disease and its impact on patients and their loved ones and health care professionals.

The EU-IPFF have a strong commitment to supporting the PF community and working with all stakeholders to bring about a future where more patients have access to proven treatments and a possible cure.

Steve Jones

New webinar series to be launched on 28th March

The EU-IPFF is proud to host a series of six webinars on topics such as clinical trials and research and patients participating in clinical trials benefit from receive regular updates on the latest developments and insights into the disease.

Keep PF research by taking part in clinical trials

This is Living Campaign

The EU-IPFF are using what we can now understand about the disease to inform the broader community about the impact of PF and its symptoms. We are also working with our partners to raise awareness and support the This is Living campaign which is aimed at outreach to patients with chronic illness. A global effort its aims to empower and inspire patients and to inform the broader community about the disease.

Patient Organisations across the world are supporting the This is Living campaign by reaching out to patients with PF to give their experiences and insights into their disease.

Dr. Simon Hart

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