Social Media campaign posts

Free and daily online YOGA classes during IPF awareness under the hammer!

PUBLIC AUCTION Hand cut University of Essen, Germany, to review current research in IPF. We have asked the management and treatment of IPF and get people onto appropriate treatments as soon as possible and to get the same levels of care as cancer patients. The lack of awareness needs to change which is rightly given to those of cancer patients. Sadly, although the life expectancy of issues for pulmonary Hbrosis patients the same level of support and public recognition is not given. At EU-IPFF, we believe from patients, carers, and healthcare professionals.

We use our annual campaigns to highlight the signs and symptoms of the disease, the campaigns have targeted the public, primary care health sta, and general practitioners. Join with the global community for #PFMonth, with one week specifically designated as September each year is when we build on our everyday mission of awareness raising to patients, now and in the future, should not face alone. Disease among the general public, policy makers and many clinicians is a battle, which range of problems, including late diagnosis and misdiagnosis, stigma, lack of adequate treatment and care, and lack of research. Every breath and every day is a battle for disease. Living with pulmonary Hbrosis is a challenge with patients and their families facing a range of problems, which is one of the greatest challenges facing our patients and their loved ones. A key aim of the new organisation was to raise awareness about this group of diseases, which is one of the greatest challenges facing our patients and their families. Focus on raising awareness of pulmonary Hbrosis and the impact it has on people living with the disease and their families.

The European Pulmonary Fibrosis Federation (EU-IPFF) was set up in 2016 by a group of patient organisations from different European countries, supported by leading doctors and nurses. A key aim of the new organisation was to raise awareness about this group of diseases, which is one of the greatest challenges facing our patients and their families.

During Pulmonary Fibrosis (PF) Awareness Month in September 2021, the European Pulmonary Fibrosis Federation (EU-IPFF) is launching a series of online YOGA classes for IPF patients.

At the end of IPF Awareness Week (18th-25th September) we are launching a public auction to raise funds for IPF research.

Thanks to the generosity of the EU-IPFF, the European Lung Foundation (ELF) and the Michael E. DeBakey charity, a range of benefits are also on offer.

People living with Pulmonary Fibrosis and families can still get involved and share their #BreathingLife story.

#BreathingLife campaign runs on the EU-IPFF's website.

#BreathingLife is a campaign to raise awareness about Pulmonary Fibrosis. The campaign is run by the European Pulmonary Fibrosis Federation (EU-IPFF) and is aimed at raising awareness of the disease and its impact on patients, their families, and caregivers.

Why? Our focus on Pulmonary Fibrosis Awareness

Francesco Bonella, President of EU-IPFF

My name is Francesco, and I am the President of the European Pulmonary Fibrosis Federation (EU-IPFF). I am writing to you to tell you about our annual campaign during #PFAwarenessMonth, which takes place from 18th - 25th September.

Our campaign is called #BreathingLife, and it is designed to raise awareness of Pulmonary Fibrosis (PF) across Europe. PF is a rare lung disease that affects around 1 in 1,000 people and is characterised by progressive scarring and inflammation of the lungs. Despite its rarity, PF affects a wide range of people, including both adults and children, and can have a significant impact on their quality of life.

The campaign runs on the EU-IPFF's website and social media channels, and is supported by leading doctors and nurses from across Europe.

During #PFAwarenessMonth, we will be running a series of online YOGA classes for IPF patients, and we are also launching a public auction to raise funds for IPF research.

We believe that every breath and every day is a battle for people living with PF. Living with this disease can be a challenge, and it is important to raise awareness of the issues facing people living with PF and their families.

We hope that you will support our campaign during #PFAwarenessMonth, and we would be grateful if you could share our posts on social media using the hashtags #BreathingLife, #PFAwarenessMonth, and #PF.

Thank you for your support.

Your support is greatly appreciated.

Francesco Bonella, President of EU-IPFF