### Wednesday, 08. June 2022

Opening Plenary 18:25 - 19:55		R1
Opening Plenar	y	
	Welcome Address	18:25 - 18:25
	Moderator: Isabelle Fajac, FR	
	Welcome Address	18:25 - 18:25
	Moderator: Harm Tiddens, NL	
	Motion is the only permanent condition that exists, it dictates everything	18:25 - 18:25
	Moderator: Harm Tiddens, NL	
	<b>Presentation of the ECFS Award</b>	18:25 - 18:25
	ECFS Award Lecture	18:25 - 18:25
	Speaker: Kris De Boeck, BE	
	Presentation of the Gerd Döring Award	18:25 - 18:25
	The impact of European patient organisations	18:25 - 18:25
	Speaker: Jacquelien Noordhoek, NL	
	Presentation of the CFE Advocacy Award	18:25 - 18:25

ECFS Tomorrow Lounge Session 20:00 - 21:30

**Welcome Reception** 

### Thursday, 09. June 2022

Symposium 08:30 - 10:00 R1 S01 - Symposium 1 - CF Research in the roaring twenties Chair: Damian G Downey, GB Chair: Kors Van der Ent, NL Sense or nonsense of theratyping 08:30 - 08:52 Speaker: Margarida Amaral, PT 08:52 - 09:14 Repair versus modulation: what's best? Speaker: George Retsch-Bogart, US Decentralisation of clinical care and randomised clinical 09:14 - 09:36 trials Speaker: Nicholas Simmonds, GB 09:36 - 10:00 Novel imaging end-points in clinical trials Speaker: Pierluigi Ciet, NL

Symposium 08:30 - 10:00

R2

### S02 - Symposium 2 - New antimicrobial therapy

#### At the end of the session, the participant will be able to:

- Synthetize knowledge on antibiotic combination therapies in the management of people with CF
- Formulate a clear opinion on the value of hyperbaric oxygen therapy
- · Appraise the place of inhaled amikacin in the CF therapeutic arsenal
- $\cdot$   $\,$  Understand the contribution of whole genome sequencing in predicting phenotypic P. aeruginosa resistance

Chair: Aurélie Crabbé, BE

Chair: Geneviève Héry Arnaud, FR

08:30 - 08:52
08:52 - 09:14
09:14 - 09:36
09:36 - 10:00

Symposium 08:30 - 10:00

R3

# ${\it S03}$ - Symposium 3 - Clinical Responses to CFTR modulator therapy from cell to patient At the end of the session, the participant will be able to:

- Describe what represents an effective CFTR modulator medication from a basic science perspective
- · Summarize what response to CFTR modulator therapy clinicians believe to be significant
- Relate objective clinical trial endpoints to symptom and patient related outcomes
- Predict what future care for people with CF who respond to CFTR modulators may look like

Chair: Clémence Martin, FR Chair: Daniel Peckham, GB

man, cb	
What is a positive response to CFTR modulator therapy - a scientist's perspective	08:30 - 08:52
Speaker: Jeffrey Beekman, NL	
What is a positive response to CFTR modulator therapy - a clinician's perspective  Speaker: Barry Plant, IE	08:52 - 09:14
What is a positive response to CFTR modulator therapy - a patient's perspective  Speaker: Audrey Chansard, FR	09:14 - 09:36
Projections for future care for people who respond to CFTR modulator therapy	09:36 - 10:00

Speaker: Jane Davies, GB

Symposium 08:30 - 10:00

R4

# S04 - Symposium 4 - Out in the wilderness: Transplantation and the co-ordination of care At the end of the session, the participant will be able to

- understand the importance of discussing "the elephant in the room" when raising the topic of transplantation with patients. Learning how to discuss the hope of transplantation alongside the real possibility that dying (before; during or after transplantation) is also a reality.
- review best practice what can we learn from different practices across Europe?
- $\cdot$  discuss the importance of ensuring that patients are appropriately reviewed post-transplant and there is someone co-ordinating care between post-transplant physicians and the CF Team.
- Discuss how to deal with anxiety and depression, common issues after transplant

Chair: Dorien Holtslag, NL Chair: Tina D'Hondt, BE

Overview of long-term transplant care in Europe	08:30 - 08:52
Speaker: Espérie Burnet, FR	
Preparing your patient for life and death - striking the balance between hope and defeat  Speaker: Sarah-Jane Mead-Regan, GB	08:52 - 09:14
Sharing the follow-up - best supportive care Speaker: Thomas Kurowski, CH	09:14 - 09:36
Hoping; coping; feeling. Living life after transplant Speaker: Trudy Havermans, BE	09:36 - 10:00

Symposium 08:30 - 10:00

R5

## ${\it S05}$ - Symposium 5 - A whole new world - Physiotherapy management in good responders to modulator therapy: ideas please!

### At the end of the session, the participant will be able to:

- · Adapt to physiotherapy management and their consequences in treating patients who are good responders to modulator therapy.
- · Increase knowledge in mucoactive agents use in good responders to modulator therapy.
- · Identify what are the changes in physiotherapy management during pregnancy in the good responders to modulator therapy. Continue or stop modulator therapy? What are the physiotherapeutic consequences in the different situations?
- $\cdot$   $\,$  Discuss the barriers to physiotherapy treatment in the era of CFTR modulators and formulate goals.

Chair: Marlies Wagner, AT Chair: Jana Plešková, CZ

CF Physiotherapy Management: A time for change?  Speaker: Wytze Doeleman, NL	08:30 - 08:52
Mucoactive agents: to take or not to take, that is the question.  Speaker: Pamela McCormack, GB	08:52 - 09:14
Pregnancy - what are the ch-ch-changes (chances, choices, challenges)  Speaker: Mathilde Legueult, FR	09:14 - 09:36
Adherence; the eternal battle	09:36 - 10:00

Symposium 08:30 - 10:00

R6

# **S06** - Symposium 6 - Understanding the mechanisms of CFTR modulation At the end of the session, the participant will be able to:

Speaker: Klára Benešová, CZ

- $\cdot$  Assess progress towards developing small molecule therapies for premature termination codon variants in CFTR
- Discuss the molecular mechanisms of action of CFTR-targeted therapies
- Compare and contrast the efficacy of CFTR rescue using CFTR-targeted therapies and proteostasis modulators, which target CFTR-interacting proteins

Chair: Miquéias Lopes Pacheco, PT

Chair: Ineke Braakman, NL

eRF1 as a target for readthrough of CFTR nonsense mutation	08:30 - 08:52
Speaker: Steven M. Rowe, US	
Mechanism-based corrector combinations to rescue CFTR misfolding	08:52 - 09:14
Speaker: Ineke Braakman, NL	
Novel targets to rescue the plasma membrane trafficking and stability of CFTR mutants	09:14 - 09:36
Speaker: Carlos M Farinha, PT	
Co-potentiation to maximize the rescue of CFTR gating mutants	09:36 - 10:00

Speaker: Guido Veit, CA

ECFS Tomorrow Lounge Session

10:00 - 10:30

### ECFS Clinical Trials Network Roundup: Now and Ahead

Chair: Damian G Downey, GB

Symposium

10:30 - 12:00 R1

### **S07** - Symposium 7 - Infection beyond CFTR modulators

### At the end of the session, the participant will be able to:

- $\cdot$   $\;$  Synthesise knowledge on the effect of CFTR modulators on both classical CF pathogens and other members of the CF microbiome
- $\cdot$   $\,$  Understand how pathogens such as P. aeruginosa can have an impact on the effect of CFTR modulators
- $\cdot$   $\,$  Understand the changes caused by the effects of CFTR modulators in the clinical management of infection
- $\cdot$  Discuss about potential strategy to combine CFTR modulators and antibiotic therapy to better manage chronic infection in CF.

Chair: Tom Coenye, BE Chair: Tavs Qvist, DK

, –	
Microbial epidemiology of cystic fibrosis in the CFTR modulator era: what has changed?	10:30 - 10:52
Speaker: Geneviève Héry Arnaud, FR	
Does CFTR correction influence the microbiome of the gut and lung in CF patients	10:52 - 11:14
Speaker: Simon Y. Graeber, DE	
Impact of <i>Pseudomonas aeruginosa</i> infection on CFTR function and correction	11:14 - 11:36
Speaker: Emmanuelle Brochiero, CA	
Do CFTR modulators change treatment of infection and clinical outcomes?	11:36 - 12:00

Symposium 10:30 - 12:00

R2

# S08 - Symposium 8 - Monitoring cystic fibrosis in 2022 - Research tools or ready for clinic?At the end of the session, the participant will be able to:

· Evaluate the utility of CT imaging in routine cystic fibrosis care

Speaker: Edith Zemanick, US

- · Appraise the arguments for and against the use of routine CT imaging in cystic fibrosis care
- $\cdot$   $\;$  Appraise the arguments for and against the routine use of multiple breath washout technologies in cystic fibrosis care
- · Question the use of current routine monitoring strategies in cystic fibrosis care

Chair: Michael Fayon, FR Chair: Anna Shawcross, GB

CT is ready for routine clinical practice - PRO Speaker: Harm Tiddens, NL	10:30 - 10:48
CT is not ready for routine clinical practice - CON Speaker: Patrick Flume, US	10:48 - 11:06
Discussion	11:06 - 11:15
Multiple Breath Washout is ready for routine clinical practice across the age groups - PRO	11:15 - 11:33

Speaker: Alexander Horsley, GB

Multiple Breath Washout is not ready for routine clinical practice across the age groups - CON

Speaker: Sanja Stanojevic, CA

**Discussion** 11:51 - 12:00

Symposium 10:30 - 12:00

R3

### S09 - Symposium 9 - Inflammation in early CF lung disease

### At the end of the session, the participant will be able to:

- · Describe the potential cause of early cystic fibrosis lung disease
- · Recognize the role of oxidate stress in early CF pathogenesis
- · Examine the role of inflammatory cells in early CF lung disease
- Summarize techniques to monitor inflammation in clinical and research environments

Chair: Olaf Eickmeier, DE Chair: Robert Gray, GB

What starts CF airway disease? Inciting events, progression and potential reversibility  Speaker: Rabin Tirouvanziam, US	10:30 - 10:52
Measurement of oxidative stress in early CF Speaker: Malcolm Brodlie, GB	10:52 - 11:14
Role of neutrophils and macrophages in early CF Speaker: Luke Garratt, AU	11:14 - 11:36
Monitoring inflammation in early CF Speaker: Hettie Janssens, NL	11:36 - 12:00

Symposium

R4

### S10 - Symposium 10 - Ethics in genetics

At the end of the session, the participant will be able to:

- · Discuss current ethical issues in relation to genetics and CF newborn screening
- · Be aware of the challenges of identifying new, rare, equivocal and single CFTR genetic variants
- Understand how CFTR modulators are impacting CFTR screening practices

Chair: Suzanne van de Vathorst, NL

Chair: Domenico Coviello, IT

10:30 - 12:00

The history and future in ethics of genetic screening (including how to manage incidental findings in general)  Speaker: Eline Bunnik, NL	10:30 - 10:52
What to do with <i>CFTR</i> variants as incidental findings? Speaker: Giuseppe Castaldo, IT	10:52 - 11:14
Genetic counseling in the era of CFTR modulators: towards harmonization?  Speaker: Danieli Salinas, US	11:14 - 11:36
Newborn screening for disease panels: what's next (future of NBS: simultaneous testing of numerous diseases using genetic analysis either as a 2nd tier step after biochemical tests or as a 1st tier step)	11:36 - 12:00

Speaker: Naama Orenstein, IL

Symposium 10:30 - 12:00

R5

# S11 - Symposium 11 - Individual cell cultures as biomarkers of CFTR function At the end of the session, the participant will be able to:

- · Understand methods for CFTR function measurement in patient-derived intestinal organoids
- · Understand methods for CFTR function measurement in patient-derived airway cells
- Explain overlap and differences between intestinal and airway culture models
- Express complementary value of in vitro and in vivo biomarkers of CFTR function

Chair: Jeffrey Beekman, NL

Chair: Anabela Santo Ramalho, BE

Defining diagnostic criteria for cystic fibrosis using intestinal organoids	10:30 - 10:52
Speaker: Francois Vermeulen, BE	
Relations between organoid swelling and long term clinical outcomes	10:52 - 11:14
Speaker: Danya Muilwijk, NL	
2D intestinal organoids for personalized therapy in cystic fibrosis- A performance comparison to donor-matched nasal cells	11:14 - 11:36
Speaker: Liron Birimberg-Schwartz, IL	
CFTR function measurements in primay airway cell cultures of individuals with cystic fibrosis	11:36 - 12:00
Speaker: Iwona Pranke, FR	

Symposium 10:30 - 12:00

R6

### S12 - Symposium 12 - Thinking differently about the GI tract

### At the end of the session, the participant will be able to:

- · To increase knowledge of the role of CFTR in the GI tract
- To be aware of the growing importance of the intestinal microbiome in the management of CF
- $\cdot$   $\,$  To gain a greater understanding of the implications of the screening and management of GI malignancies in an aging CF population

Chair: Frank Bodewes, NL Chair: Anne Munck, FR

CFTR function in the gut - pathophysiology Speaker: Anna Bertolini, NL	10:30 - 10:52
The intestinal microbiome - a silent player?  Speaker: Stephanie Van Biervliet, BE	10:52 - 11:14
GI malignancies - an update Speaker: Denis Hadjiliadis, US	11:14 - 11:36
The effects of CFTR carriers on gastrointestinal phenotype  Speaker: Michael Wilschanski, II.	11:36 - 12:00

Satellite Symposium 12:30 - 14:00

R1

### **Satellite Symposium**

Please find the detailed programme of the Satellite Symposia here.

*Meet the Experts* 12:45 - 13:45

### **Meet the Experts**

Meet the Experts 12:45 - 13:45

## $\it Meet\ the\ Experts$ - Meet the experts 2 - Remote monitoring - Evidence and practical experience

ePoster Corner B

Expert: Christopher Goss, US Expert: Nichola MacDuff, GB

Meet the Experts 12:45 - 13:45

### Meet the Experts - Meet the Experts 3 - Treating the ageing CF patient

ePoster Corner C

Expert: Andrew Jones, GB Expert: Lieven Dupont, BE

*Meet the Experts* 12:45 - 13:45

## $\it Meet\ the\ Experts$ - Meet the Experts 1 - How to manage CFTR modulators in severe patients (FEV1

ePoster Corner A

Expert: Pierre-Régis Burgel, FR Expert: Maarten Ploeger, NL

ECFS Tomorrow Lounge Session

12:45 - 13:45

#### Telemedicine and mental health screening online

Chair: Janne Houben, BE Chair: Pavla Hodkova, CZ Chair: Trudy Havermans, BE Chair: Anna M. Georgiopoulos, US

Chair: Edwina Landau, IL

ePoster Session 14:00 - 15:00

#### **Poster Viewing 1**

P001	CF-NBS start-up in Flanders (Belgium): report of first evaluation after three years	14:00 - 14:00
	Abstract Presenter: Julie Carbonez, BE	
P002	Growth, lung microbiology and structure after implementation of newborn screening in Danish cystic fibrosis patients	14:00 - 14:00
	Abstract Presenter: Marianne Skov, DK	
P003	Wide variability of approach to cystic fibrosis newborn screening across regions of Italy	14:00 - 14:00
	Abstract Presenter: Carlo Castellani, IT	
P004	Evaluation of the efficacy of newborn screening as a tool to identify cystic fibrosis in the Russian Federation	14:00 - 14:00

	Abstract Presenter: Victoria Sherman, RU	
P006	Nephrolithiasis and nephrocalcinosis in infants with cystic fibrosis: a case presentation of two swedish infants  Abstract Presenter: ELisabeth Christiansen, SE	14:00 - 14:00
P007	CFTR genotypes and spermatology in cystic fibrosis patients without CBAVD  Abstract Presenter: Anna Sedova, RU	14:00 - 14:00
P008	L138ins is common <i>CFTR</i> gene mutation in Russian infertile men  Abstract Presenter: Ekaterina Marnat, RU	14:00 - 14:00
P009	Spectrum of CFTR mutations in newly diagnosed cases of cystic fibrosis through newborn screening in the Republic of North Macedonia  Abstract Presenter: Stojka Fustik, MK	14:00 - 14:00
P013	Peptide modulation of COMMD1: corrector and anti- inflammatory effect in cystic fibrosis context? Abstract Presenter: Benjamin Simonneau, FR	14:00 - 14:00
P014	Proof of concept of ionocytes' CFTR content as a novel biomarker for cystic fibrosis diagnosis and follow-up Abstract Presenter: Fabina Ciciriello, IT	14:00 - 14:00
P015	Translational read-through inducing drugs: a study of toxicity in mice models and <i>in vitro</i> safety validation of the specific read-through process  Abstract Presenter: Federica Corrao, IT	14:00 - 14:00
P017	Using chamber measurement of CFTR modulator effects in airway epithelial cells and its correlation to clinical effects of treatment  Abstract Presenter: Bente Aalbers, NL	14:00 - 14:00
P018	People with cystic fibrosis do not show an increased interferon-response transcriptomic signature in nasal epithelial cells  Abstract Presenter: Manu Jain, US	14:00 - 14:00
P019	Monocyte integrin activation as a CFTR targeted drugs evaluation test in cystic fibrosis patients: preliminary analysis  Abstract Presenter: Jessica Conti, IT	14:00 - 14:00
P020	Evaluation of TMEM16A as a modifier of the cystic fibrosis lung phenotype utilising cystic fibrosis patient-specific human-induced pluripotent stem cells  Abstract Presenter: Mark-Christian Jaboreck, DE	14:00 - 14:00
P021	A novel gene delivery approach for CRISPR-Cas9-mediated permanent CFTR correction  Abstract Presenter: Jim Hu, CA	14:00 - 14:00
P022	hCFTRdelR expression and correction of human cystic fibrosis airway epithelia increase with increasing SP-101 and doxorubicin doses	14:00 - 14:00
	Abstract Presenter: Katherine Excoffon, US	

P023	High-resolution imaging reveals impaired lung development in a mouse model of cystic fibrosis-like lung disease  Abstract Presenter: Pinelopi Anagnostopoulou, CH	14:00 - 14:00
	Abstract Fresenter: Finetopi Anagnostopoulou, Cii	
P024	Patient-derived organoids as a model for treatment diagnosis in cystic fibrosis - combinatory treatment with read-through agent, correctors and potentiator on rare mutations	14:00 - 14:00
	Abstract Presenter: Johanna Pott, NL	
P025	Assessment of CFTR modulator combinations in rectal organoids from F508del homozygous patients with cystic fibrosis	14:00 - 14:00
	Abstract Presenter: Senne Cuyx, BE	
P026	Theratyping of the CFTR variant G85E in trans with the complex allele A1006E+V562I using rectal organoids	14:00 - 14:00
	Abstract Presenter: Jessica Conti, IT	
P027	Effects of ivacaftor therapy confirm the results of theratyping using rectal and nasal epithelial cells of a cystic fibrosis patient carrying the ultra-rare CFTR genotype W57G (c.169T>G) / A234D (c.701C>A)	14:00 - 14:00
	Abstract Presenter: Jessica Conti, IT	
P028	Antisense oligonucleotide splicing modulation as a novel cystic fibrosis therapeutic approach for the W1282X nonsense mutation  Abstract Presenter: Yifat Oren, IL	14:00 - 14:00
Dooo		14.00 14.00
P029	Antisense oligonucleotide-based drug development for cystic fibrosis patients carrying the 2789+5 G-to-A splicing mutation	14:00 - 14:00
	Abstract Presenter: Yifat Oren, IL	
P030	Effect of CFTR modulators on lung function, body mass index, and <i>Pseudomonas aeruginosa</i> status in adolescents with cystic fibrosis - one-year follow-up	14:00 - 14:00
	Abstract Presenter: Marina Praprotnik, SI	
P031	Impact of the elexacaftor/tezacaftor/ivacaftor combination regimen on lung function in adults with cystic fibrosis: more than just ${\sf FEV_1}$	14:00 - 14:00
	Abstract Presenter: Doris Dieninghoff, DE	
P032	Sweat chloride values in cystic fibrosis patients on elexacaftor/texacaftor/ivacaftor	14:00 - 14:00
	Abstract Presenter: Nadia Mazzoni, IT	
P033	Anti-inflammatory effects of Kaftrio in adults with cystic fibrosis heterozygous for F508del-CFTR	14:00 - 14:00
	Abstract Presenter: Heledd Jarosz-Griffiths, GB	
P034	Long-term elexacaftor/tezacaftor/ivacaftor CFTR modulation significantly increases lung function and peak power output in people with cystic fibrosis	14:00 - 14:00
	Abstract Presenter: Barlo Hillen, DE	
P035	Elexacaftor/tezacaftor/ivacaftor improves very quickly ${\sf FEV_1}$ and 6MWT very quickly in patients with cystic fibrosis and	14:00 - 14:00

	severe airflow obstruction	
	Abstract Presenter: Laurent MELY, FR	
P036	Exercise performance before and after introduction of elexacaftor/tezacaftor/ivacaftor (Kaftrio) in Scottish children with cystic fibrosis: a two-centre retrospective study	14:00 - 14:00
	Abstract Presenter: Paul D Burns, GB	
P037	Improved aerobic fitness in children on CFTRm triple combination therapy  Abstract Presenter: Uros Krivec, SI	14:00 - 14:00
P038	ELX/TEZ/IVA discontinuation is more common in an older population  Abstract Presenter: Karuna Sapru, GB	14:00 - 14:00
P039	Changes to clinical well-being following initiation of elexacaftor/tezacaftor/ivacaftor in children with cystic fibrosis - single-centre experience	14:00 - 14:00
	Abstract Presenter: Amanda McGrath, GB	
P040	Implementation and preliminary experiences of the HERO-2 real-world research study in patients with cystic fibrosis  Abstract Presenter: Cynthia Brown, US	14:00 - 14:00
P041	Clinical assessment plan for initiation and follow-up of treatment with elexacaftor/tezacaftor/ivacaftor in the danish cystic fibrosis population	14:00 - 14:00
	Abstract Presenter: Christian Leo-Hansen, DK	
P042	The effect of elexacaftor/tezacaftor/ivacaftor on non- pulmonary symptoms in adults with cystic fibrosis Abstract Presenter: Sarah Allgood, US	14:00 - 14:00
P043	Significant reduction in abdominal symptoms assessed with the CFAbd-Score over 6 months of elexacaftor/tezacaftor/ivacaftor - follow-up results from Irish and British cystic fibrosis patients (RECOVER study)	14:00 - 14:00
	Abstract Presenter: Jochen G. Mainz, DE	
P044	Treatment benefits of elexacaftor/tezacaftor/ivacaftor in people with cystic fibrosis carrying non-F508del mutations  Abstract Presenter: Galit Livnat, IL	14:00 - 14:00
P045	The impact of lumacaftor/ivacaftor and tezacaftor/ivacaftor on health outcomes in the setting of severe lung disease due to cystic fibrosis	14:00 - 14:00
	Abstract Presenter: Sanja Stanojevic, CA	
P046	The effectiveness of CFTR modulators in people with cystic fibrosis and rare mutations - a real-world study  Abstract Presenter: Sofia Hanger, GB	14:00 - 14:00
P047	Selection of CFTR modulators for children carrying the genetic variant W1282R  Abstract Presenter: Elena Kondratyeva, RU	14:00 - 14:00
P048	Continuous glucose monitoring in people with cystic fibrosis highlights different glucose tolerance abnormalities	14:00 - 14:00

	according to pancreatic exocrine status.  Abstract Presenter: Luc RAKOTOARISOA, FR	
P049	Question cystic fibrosis again: a refresh of the James Lind Alliance Priority Setting Partnership in cystic fibrosis (CF) Abstract Presenter: Nicola Rowbotham, GB	14:00 - 14:00
P050	Writing with patients, for patients: co-creation of a cystic fibrosis plain language glossary  Abstract Presenter: Kate Hayes, GB	14:00 - 14:00
P051	Improved opportunities for trial participation via the Dutch CF Trial Consortium  Abstract Presenter: Ilonka Paalvast, NL	14:00 - 14:00
P052	Introduction of an innovative 'pill swallowing school' in a cystic fibrosis paediatric population within a tertiary centre Abstract Presenter: Rachel Rowley, GB	14:00 - 14:00
P053	A novel family-focused intervention to support physical activity among children (6-12 years) with cystic fibrosis Abstract Presenter: Owen William Tomlinson, GB	14:00 - 14:00
P054	Severity of COVID-19 infection in cystic fibrosis patients compared to the general population  Abstract Presenter: Chiara Rosazza, IT	14:00 - 14:00
P055	The impact of virtual care on cystic fibrosis disease progression - a prospective multicentre study in children with cystic fibrosis	14:00 - 14:00
	Abstract Presenter: Jakob Medbo, SE	
P056	Acute pulmonary exacerbation markedly weakens beneficial effect of elexacaftor/tezacaftor/ivacaftor on systemic endothelial dysfunction in cystic fibrosis - a single case study  Abstract Presenter: Barbara Salobir, SI	14:00 - 14:00
P057	Monitoring individualised CFTR modulating therapy in an adolescent with cystic fibrosis homozygous for the G85E CFTR variant using in vitro and in vivo methods  Abstract Presenter: Niklas Ziegahn, DE	14:00 - 14:00
P058	Resolution of chronic pneumothorax in an adult patient with severe cystic fibrosis treated with elexacaftor/tezacaftor/ivacaftor	14:00 - 14:00
	Abstract Presenter: Barbara Messore, IT	
P059	Association of <i>Pseudomonas aeruginosa</i> infection stage with lung function trajectory in children with cystic fibrosis  Abstract Presenter: Margaret Rosenfeld, US	14:00 - 14:00
P060	Factors associated with pulmonary functions of cystic fibrosis patients in the National Cystic Fibrosis Patient Registry: a retrospective cohort study	14:00 - 14:00
	Abstract Presenter: Nazan Cobanoglu, TR	
P061	Low $FEV_1$ performance in the 30-34 years of age cohort in Norway: utilising Registry data in the search for causes Abstract Presenter: Audun OS, NO	14:00 - 14:00

P062	An international survey: understanding the health and perspectives of people with cystic fibrosis (CF) not benefitting from CFTR modulators	14:00 - 14:00
	Abstract Presenter: Jennifer Taylor-Cousar, US	
P063	Physiotherapy data for the UK cystic fibrosis Registry - review and re-launch  Abstract Presenter: Lisa Morrison, GB	14:00 - 14:00
P064	Iron deficiency in cystic fibrosis: a prospective study in a modern adult cohort  Abstract Presenter: Quitterie Reynaud, FR	14:00 - 14:00
P065	Cervical cancer screening in women with cystic fibrosis: should we do more?	14:00 - 14:00
	Abstract Presenter: Megan Hadfield, GB	
P066	Changing criteria of paediatric chronic rhinosinusitis indexation in Russian National Cystic Fibrosis Registry  Abstract Presenter: Dmitry Polyakov, RU	14:00 - 14:00
P067	Main characteristics of cystic fibrosis patients: National Patient Registry of Russia (RCFPR) 2020	14:00 - 14:00
	Abstract Presenter: Elena Kondratyeva, RU	
P068	Characteristics of genetic variants of the <i>CFTR</i> gene in the Russian Federation according to the 2020 Registry	14:00 - 14:00
	Abstract Presenter: Anna Voronkova, RU	
P069	Patient-reported outcome measures in children with cystic fibrosis	14:00 - 14:00
P070	Harnessing electronic health records to enhance cbservational research in cystic fibrosis  Abstract Presenter: Nicole Filipow, GB	14:00 - 14:00
P071	The characterisation of a patient cohort that had home spirometry devices and opted into sharing their data with the US Cystic Fibrosis Foundation  Abstract Presenter: Alexander Elbert, US	14:00 - 14:00
P072	SARS Cov2 infection in patients with cystic fibrosis (CF) - a single centre experience  Abstract Presenter: Ani Vidoevska, MK	14:00 - 14:00
P073	Observed impact on admissions during the COVID-19 pandemic of paediatric cystic fibrosis patients in a tertiary hospital setting  Abstract Presenter: Sara Warraich, GB	14:00 - 14:00
P074	Post-COVID-19 condition in children with cystic fibrosis  Abstract Presenter: Anastasia Chernyavskaya, RU	14:00 - 14:00
P075	COVID-19 vaccination uptake in patients with cystic fibrosis  Abstract Presenter: Dion-Emily Manning, GB	14:00 - 14:00
P076	COVID-19 vaccination in children and adolescents with cystic fibrosis - a single centre experience  Abstract Presenter: ALEKSANDRA ZVER, SI	14:00 - 14:00

P077	Vaccination of patients with cystic fibrosis during the COVID-19 pandemic in Croatia  Abstract Presenter: Ivona Markelić, HR	14:00 - 14:00
P078	Adverse effects of Kaftrio in an adult cystic fibrosis clinic	14:00 - 14:00
1070	Abstract Presenter: Douglas McCabe, GB	14.00 - 14.00
P079	Real-life data on the efficacy and safety of tezacaftor/ivacaftor in people living with cystic fibrosis homozygous for F508del and heterozygous for F508del and a residual function mutation  Abstract Presenter: Eef Vanderhelst, BE	14:00 - 14:00
P080	Real-world experience of patient outcomes following one year of elexacaftor/tezacaftor/ivacaftor treatment at a single adult cystic fibrosis centre  Abstract Presenter: Ellen Shiner, GB	14:00 - 14:00
P081	Adverse drug reactions linked to prolonged use of Kaftrio  Abstract Presenter: Michael Kevin Dooney, GB	14:00 - 14:00
P082	Reduced dose Kaftrio can significantly improve respiratory health for those intolerant of the recommended dosing schedule  Abstract Presenter: Michael Kevin Dooney, GB	14:00 - 14:00
P083	Decreased respiratory burst in circulating neutrophils after initiation of CFTR modulator therapy in cystic fibrosis patients with chronic lung infections  Abstract Presenter: Peter Østrup Jensen, DK	14:00 - 14:00
P084	Counteracting inflammation triggered by <i>P. aeruginosa</i> -activated lung-infiltrating Th1/17 cells: a novel approach for precision medicine in cystic fibrosis	14:00 - 14:00
	Abstract Presenter: Moira Paroni, IT	
P085	Evaluation of neutrophil/lymphocyte ratio, platelet/lymphocyte ratio and mean platelet volume in cystic fibrosis (CF) during exacerbation	14:00 - 14:00
	Abstract Presenter: Tugba Sismanlar Eyuboglua, TR	
P086	Serum levels of alpha calcitonin gene-related peptide and vasoactive intestinal peptide as predictors of exacerbation in cystic fibrosis	14:00 - 14:00
	Abstract Presenter: Maha Al-Keilani, JO	
P087	CTN - Lung Clearance Index core facility: quality improvement exercise	14:00 - 14:00
	Abstract Presenter: Clare Saunders, GB	
P088	Ventilation inhomogeneity in patients with cystic fibrosis and pancreatic sufficiency and insufficiency Abstract Presenter: Oded Breuer, IL	14:00 - 14:00
P089	Ventilation heterogeneity in adult patients with cystic fibrosis (CF)  Abstract Presenter: Alexander Chernyak, RU	14:00 - 14:00
P090	Comparison of multiple breath washout and particles in exhaled air for assessment of small airway function in	14:00 - 14:00

	children with cystic fibrosis Abstract Presenter: Annelies M. Zwitserloot, NL	
P091	Correlation of lung function using forced oscillation technique with spirometry in children aged 4-18 years with cystic fibrosis (CF)  Abstract Presenter: SACHIN SINGH, IN	14:00 - 14:00
P092	Exploring the association between hand grip strength (HGS) and lung function (FEV <sub>1</sub> %) in cystic fibrosis: does HGS offer an insight into pulmonary function?  Abstract Presenter: Matthew Taylor, GB	14:00 - 14:00
P093	The effect of probiotic administration on glucose metabolism and metabolomics in cystic fibrosis patients Abstract Presenter: Michal Gur,	14:00 - 14:00
P094	Randomised controlled trial of humanoid robot-based distraction for invasive nursing procedures in children with cystic fibrosis: preliminary results  Abstract Presenter: Sergio Bella, IT	14:00 - 14:00
P095	Effect of hypertonic saline therapy on Lung Clearance Index in preschool children with cystic fibrosis  Abstract Presenter: Almala Pinar Ergenekon, TR	14:00 - 14:00
P096	Tolerability of tigerase (dornase alfa) in children  Abstract Presenter: Vera Shadrina, RU	14:00 - 14:00
P097	Evaluation of the inhaled mannitol tolerance test in children with cystic fibrosis  Abstract Presenter: Vera Shadrina, RU	14:00 - 14:00
P098	The use of mucolytic therapy in cystic fibrosis in the Russian Federation  Abstract Presenter: Vera Shadrina, RU	14:00 - 14:00
P099	Inhaled tobramycin solution hypersensitivity in patients with cystic fibrosis at the Centre for Cystic Fibrosis, Institute for Pulmonary Diseases in Children in North Macedonia	14:00 - 14:00
P100	Abstract Presenter: Marija Atanasova Nadzinska, MK  Beneficial effects and safety of omalizumab during pregnancy in a patient with cystic fibrosis	14:00 - 14:00
P101	Abstract Presenter: Barbara Messore, IT  Microbiological surveillance in cystic fibrosis - an emerging	14:00 - 14:00
1101	multifactorial challenge  Abstract Presenter: Peter Barry, GB	14.00 - 14.00
P102	Prevalence of sensitisation to common inhaled allergens in a Belgian adult cystic fibrosis population and its impact on respiratory outcomes  Abstract Presenter: Henri Marissiaux, BE	14:00 - 14:00
P103	Evaluation of patient opinion about the current and future pharmacy service for an adult cystic fibrosis clinic Abstract Presenter: Douglas McCabe, GB	14:00 - 14:00

P104	Exploring the value of annual chest radiographs in people with cystic fibrosis: an observational study from a single UK centre  Abstract Presenter: Christopher Stovin, GB	14:00 - 14:00
P105	Home spirometry is reliable and of consistent quality in children with cystic fibrosis  Abstract Presenter: Siobhan Carr, GB	14:00 - 14:00
P106	Upper airway disease in adults with cystic fibrosis: a cross- sectional study Abstract Presenter: Saartje Uyttebroek, BE	14:00 - 14:00
P107	Coexistence of cystic fibrosis and metabolic disease: is this a related or coincidental condition?  Abstract Presenter: Halime Nayır Buyuksahin, TR	14:00 - 14:00
P108	Impact of improved nontuberculous mycobacterial culture surveillance strategy in an adult cystic fibrosis cohort Abstract Presenter: Natalie Lorent, BE	14:00 - 14:00
P109	Evaluation of early lung disease in school-age children with cystic fibrosis - a preliminary report  Abstract Presenter: Katarzyna Walicka-Serzysko,	14:00 - 14:00
P110	Trends in asthma diagnosis and usage of asthma medications in children with cystic fibrosis - are we looking for it?	14:00 - 14:00
P111	Abstract Presenter: Anirban Maitra, GB  Atopy and cystic fibrosis: characterisation of a paediatric portuguese population  Abstract Presenter: Susana Castanhinha, PT	14:00 - 14:00

ePoster Session 14:00 - 15:00

ePoster Corner A

### ePoster Session 1 - What have we learned from the psychosocial impacts of COVID-19?

Leader: Trudy Havermans, BE

Leader: Kathryn Braisdell, GB			
Home spirometry and virtual visits in children with cystic fibrosis - the child's perspective	14:00 - 14:06		
Abstract Presenter: Frida Olofsson, SE			
Long-term psychological impact of COVID-19 on adult patients with cystic fibrosis, including transplanted patients	14:06 - 14:12		
Abstract Presenter: Trudy Havermans, BE			
The impact of the COVID-19 lockdown and introduction of precision medication Symkevi and Orkambi on cystic fibrosis patients at Sheffield Children's NHS Foundation Trust, UK (SCFT)	14:12 - 14:18		
Abstract Presenter: Kathryn Braisdell, GB			
Sleep disturbances in children with cystic fibrosis at the beginning and in the first year of the COVID-19 pandemic Abstract Presenter: Tugba Sismanlar Eyuboglu, TR	14:18 - 14:24		
	Home spirometry and virtual visits in children with cystic fibrosis - the child's perspective  Abstract Presenter: Frida Olofsson, SE  Long-term psychological impact of COVID-19 on adult patients with cystic fibrosis, including transplanted patients  Abstract Presenter: Trudy Havermans, BE  The impact of the COVID-19 lockdown and introduction of precision medication Symkevi and Orkambi on cystic fibrosis patients at Sheffield Children's NHS Foundation Trust, UK (SCFT)  Abstract Presenter: Kathryn Braisdell, GB  Sleep disturbances in children with cystic fibrosis at the beginning and in the first year of the COVID-19 pandemic		

EPS1.05	Impact of COVID-19 on mental health among people with cystic fibrosis  Abstract Presenter: Elpis Hatziagorou, GR	14:24 - 14:30
EPS1.06	Evolution of psychological distress during progression of the COVID-19 pandemic in adults with cystic fibrosis Abstract Presenter: Federico Cresta, IT	14:30 - 14:36
EPS1.07	Parental experiences of face-to-face versus virtual cystic fibrosis clinics during the COVID-19 pandemic: questionnaire study  Abstract Presenter: Kieren James Lock, GB	14:36 - 14:42
EPS1.08	Telemedicine clinics for people with cystic fibrosis (CF): experiences 18 months on  Abstract Presenter: Muhammad Haris Mir, GB	14:42 - 14:48
EPS1.09	Exploring associations and indirect effects between mindfulness, psychological flexibility, and well-being in adults with cystic fibrosis: informing future interventions Abstract Presenter: Sophia Kauser, GB	14:48 - 14:54
EPS1.10	Easy Medicines for Burden Reduction And Care Enhancement (EMBRACE): Exploring the views of adults with cystic fibrosis on the current nebulised medicines supply process using homecare deliveries in two UK centres Abstract Presenter: Aoife Lynam, GB	14:54 - 15:00

ePoster Session 14:00 - 15:00

### ePoster Sessions

ePoster Session 14:00 - 15:00

## ePoster Session ${\bf 2}$ - Did some good come from COVID-19? The lasting value of remote treatment

ePoster Corner B Leader: Brenda Button, AU Leader: Donald Urquhart, GB EPS2.01 Home monitoring of airway clearance treatments (ACTs) in 14:00 - 14:06 children and young people with cystic fibrosis Abstract Presenter: Emma Raywood, GB EPS2.02 Comparison of FEV<sub>1</sub> on a remote spirometry device with 14:06 - 14:12 hospital software in an adult cystic fibrosis cohort Abstract Presenter: Paul Wilson, GB EPS2.03 A service evaluation of 'CF THRIVE': an online, student-led 14:12 - 14:18 group for children with cystic fibrosis Abstract Presenter: Madeline Pilbury, GB 14:18 - 14:24 Is tele-physiotherapy as effective as home-visit-guided EPS2.04 physiotherapy among children with cystic fibrosis? Abstract Presenter: Pauline Peftoulidou, GR 14:24 - 14:30 **EPS2.05** Use of BEAM to enhance participation in exercise Abstract Presenter: Aimee Johnson, GB

EPS2.06	"It's good to talk" - development of a virtual international physiotherapy discussion forum  Abstract Presenter: Gemma E Stanford, GB	14:30 - 14:36
EPS2.07	International collaboration and development of a cystic fibrosis physiotherapy-specific telehealth toolkit Abstract Presenter: Lisa Morrison, GB	14:36 - 14:42
EPS2.08	Patient and parent experience of a virtual physiotherapy clinic in children with cystic fibrosis - a pilot study  Abstract Presenter: Linn Hoel, SE	14:42 - 14:48
EPS2.09	Feasibility and acceptability of establishing virtual exercise classes for cystic fibrosis (CF) patients  Abstract Presenter: Ellen Shiner, GB	14:48 - 14:54
EPS2.10	Exercise testing and training in cystic fibrosis clinics in the United Kingdom: a ten-year update  Abstract Presenter: Owen William Tomlinson, GB	14:54 - 15:00

ePoster Session 14:00 - 15:00

### ePoster Session 3 - Pathogenesis and treatment of CF pathogens

ePoster Corner C

Gilpin, GB emanick, US	
Preliminary results of an ongoing study, which determines the prevalence and possible impact of mucoid Staphylococcus aureus on lung disease of people with cystic fibrosis	14:00 - 14:06
Abstract Presenter: Christine Rumpf, DE	
Deciphering the adaptive evolution of successful Pseudomonas aeruginosa lineages within persistent bacterial infections	14:06 - 14:12
Abstract Presenter: Akbar Espaillat, DK	
Using patient-derived airway models to study <i>Pseudomonas</i> aeruginosa colonisation and infection  Abstract Presenter: Signe Lolle, DK	14:12 - 14:18
Does Achromobacter xylosoxidans have specific characteristics that could contribute to its emergence in cystic fibrosis?  Abstract Presenter: Pauline Sorlin, FR	14:18 - 14:24
Induced levofloxacin resistance persists in evolved lineages of <i>Pseudomonas aeruginosa</i> isolates from people with cystic fibrosis after exposure to other inhaled antibiotics  Abstract Presenter: Callum Matthew Sloan, GB	14:24 - 14:30
Isolation and characterisation of nebulised phage for treatment of chronic <i>Pseudomonas aeruginosa</i> (Pa) pulmonary infections in cystic fibrosis (CF) patients <i>Abstract Presenter</i> : Maya Kahan-Hanum, IL	14:30 - 14:36
	Preliminary results of an ongoing study, which determines the prevalence and possible impact of mucoid Staphylococcus aureus on lung disease of people with cystic fibrosis  Abstract Presenter: Christine Rumpf, DE  Deciphering the adaptive evolution of successful Pseudomonas aeruginosa lineages within persistent bacterial infections  Abstract Presenter: Akbar Espaillat, DK  Using patient-derived airway models to study Pseudomonas aeruginosa colonisation and infection  Abstract Presenter: Signe Lolle, DK  Does Achromobacter xylosoxidans have specific characteristics that could contribute to its emergence in cystic fibrosis?  Abstract Presenter: Pauline Sorlin, FR  Induced levofloxacin resistance persists in evolved lineages of Pseudomonas aeruginosa isolates from people with cystic fibrosis after exposure to other inhaled antibiotics  Abstract Presenter: Callum Matthew Sloan, GB  Isolation and characterisation of nebulised phage for treatment of chronic Pseudomonas aeruginosa (Pa) pulmonary infections in cystic fibrosis (CF) patients

EPS3.07	Exploring the therapeutic potential of iminosugars as antibacterial and antibiofilm agents in the treatment of cystic fibrosis lung disease infections  Abstract Presenter: Anna Esposito, IT	14:36 - 14:42
EPS3.08	Inhaled powder tobramycin (TIP): cyclical versus continuous treatment: iBEST Study  Abstract Presenter: Aya Alkhatib, GB	14:42 - 14:48
EPS3.09	Methicillin-sensitive Staphylococcus aureus (MSSA) with inoculum-related reduced susceptibility to cefazolin (CZ) and piperacillin-tazobactam (TZP) in persons with cystic fibrosis (pwCF)  Abstract Presenter: Julianna Svishchuk, CA	14:48 - 14:54
EPS3.10	Opportunistic bacterial pathogens: potential transmissions between the patients and their domestic environment Abstract Presenter: Quentin Menetrey, FR	14:54 - 15:00

ECFS Tomorrow Lounge Session

14:00 - 15:00

Nutrition: Future role of the dietitian in CF

Chair: Sarah Collins, GB Chair: Elizabeth Owen, GB

<i>Workshop</i> 15:00 - 16:30 <b>WS01 - WS01</b> :	Treatment of respiratory infection in cystic fibrosis	R1
Chair: Michal Sh Chair: Michael T	iteinberg, IL	
WS01.01	Antipseudomonal treatment decisions during cystic fibrosis exacerbation management	15:00 - 15:15
	Abstract Presenter: Donald R. VanDevanter, US	
WS01.02	Trends and outcomes of long-term inhaled antimicrobial treatment in people with cystic fibrosis without chronic <i>Pseudomonas aeruginosa</i> infection: an ECFSPR data analysis	15:15 - 15:30
	Abstract Presenter: Michal Shteinberg, IL	
WS01.03	CFTR modulators impact antibiotic susceptibility of Pseudomonas aeruginosa and Staphylococcus aureus Abstract Presenter: Cristina Cigana, IT	15:30 - 15:45
WS01.04	A new weapon against Mycobacterium abscessus Abstract Presenter: Giulia Degiacomi, IT	15:45 - 16:00
WS01.05	Combination of the lytic bacteriophage NP3 with ciprofloxacin has a synergistic effect on biofilms of cystic fibrosis <i>Pseudomonas aeruginosa</i> isolates  Abstract Presenter: Oana Ciofu, DK	16:00 - 16:15
WS01.06	The impact of Orkambi on respiratory cystic fibrosis pathogens  Abstract Presenter: Mahasin Al Shakirchi, SE	16:15 - 16:30

Workshop 15:00 - 16:30 R2 WS02 - WS02: Digestive and liver disease upon elexacaftor/tezacaftor/ivacaftor therapy Chair: Michael Wilschanski, IL Chair: Jaroslaw Walkowiak, PL WS02.01 Abdominal symptoms significantly decline after 24 weeks of 15:00 - 15:15 elexacaftor/tezacaftor/tvacaftor treatment: first results obtained with the cystic fibrosis-specific CFAbd-Score in Germany and the UK Abstract Presenter: Jochen G. Mainz, DE WS02.02 MRI metrics of small bowel water in cystic fibrosis (CF) 15:15 - 15:30 before and after elexacaftor/tezacaftor/ivacaftor: first results from the GIFT-CF3 Study Abstract Presenter: Christabella Ng, GB WS02.03 Investigating changes in liver function tests across the first 15:30 - 15:45 year of elexacaftor/tezacaftor/ivacaftor therapy Abstract Presenter: Nicola Robinson, GB Liver function test abnormalities in cystic fibrosis patients WS02.04 15:45 - 16:00 commenced on ELX/TEZ/IVA in a large adult cystic fibrosis centre over 12 months Abstract Presenter: Daniel Tewkesbury, GB WS02.05 16:00 - 16:15 The use of elexacaftor/tezacaftor/ivacaftor in adult patients with cystic fibrosis and established liver cirrhosis: a case series Abstract Presenter: James Sun, GB WS02.06 16:15 - 16:30 Can liver biopsy guide the management of deranged liver function tests in liver transplant recipients receiving Kaftrio? Abstract Presenter: Phaedra Tachtatzis, GB Workshop 15:00 - 16:30 R3 WS03 - WS03: Modulator therapies: what impact are they having on my life? Chair: Edwina Landau, IL Chair: Majda Oštir, SI WS03.01 Impact of Trikafta on PHQ-9 and GAD-7 scores 15:00 - 15:15 Abstract Presenter: Heather Bruschwein, US WS03.02 Understanding beliefs about elexacaftor/tezacaftor/ivacaftor 15:15 - 15:30 (ETI) therapy in adults living with cystic fibrosis Abstract Presenter: Jamie Duckers, GB WS03.03 Qualitative analyses of the experiences of people with end-15:30 - 15:45 stage cystic fibrosis lung disease receiving ELX/TEZ/IVA Abstract Presenter: Trudy Havermans, BE WS03.04 Psychosocial outcomes from annual review - a comparison 15:45 - 16:00 of young people prescribed CFTR medication and those who were not Abstract Presenter: Steve Jones, GB WS03.05 16:00 - 16:15 Proportion days covered: what does it tell us about adherence to Kaftrio and Kalydeco?

	Abstract Presenter: Amanda Bevan, GB	
WS03.06	Does the use of modulator therapies in adults with cystic fibrosis have a long-term impact on the requirement for intravenous antibiotics	16:15 - 16:30
	Abstract Presenter: Sharon Geoghegan, GB	
<i>Workshop</i> 15:00 - 16:30		R4
<b>WS04</b> - WS04:	There is more to CF than the lung	
Chair: Peter Bar Chair: Dorota Sa		
WS04.01	Do adults with Cystic Fibrosis-Related Diabetes on insulin treatment benefit from using flash glucose monitoring?  Abstract Presenter: Harbinder Sunsoa, GB	15:00 - 15:15
WS04.02	Clinical impact of pathogenic CFTR mutations in paediatric- onset pancreatitis (preliminary analysis from the APPLE study)	15:15 - 15:30
	Abstract Presenter: Márk Félix Juhász, HU	
WS04.03	Pulmonary impacts of the long-term use of proton pump inhibitors to potentialise the effect of pancreatic extracts in children with cystic fibrosis  Abstract Presenter: Sahar Awada, FR	15:30 - 15:45
MICOAOA		15 45 16 00
WS04.04	Impact of planned versus unplanned pregnancy in people with cystic fibrosis  Abstract Presenter: Raksha Jain, US	15:45 - 16:00
WS04.05	What is the scope for colorectal cancer screening in cystic fibrosis? Ten years of experience at a UK cystic fibrosis centre	16:00 - 16:15
	Abstract Presenter: Karuna Sapru, GB	
WS04.06	Neoplastic disease and treatment in cystic fibrosis (CF): a comprehensive single centre case series	16:15 - 16:30
	Abstract Presenter: Alan Anderson, GB	
Symposium 15:00 - 16:30		R5

# SS01 - Special Symposium - Making time for exercise and physical activity At the end of the session, the participant will be able to:

- Describe benefits of exercise and physical activity further to latest data from ACTIVATE-CF study and the updated Cochrane review
- $\cdot$  Evaluate exercise recommendations based on considerations of age, disease severity, patient preference, barriers to exercise
- Summarize the evidence that exercise may replace chest physiotherapy as airway clearance therapy (ACT)
- Discuss different approaches to design studies to examine whether exercise can replace ACT

Chair: Helge Hebestreit, DE Chair: Marcella Burghard, NL

ACTIVATE-CF and Updated Cochrane Review: New perspectives on the benefits of exercise and physical activity

Speaker: Thomas Radtke, CH

15:00 - 15:22

	Exercise recommendations are not 'one size, fits all' Speaker: Mathieu Gruet, FR	15:22 - 15:44
	Exercise as Airway Clearance - Perspectives of the CF community on exercise as replacement for chest physiotherapy  Speaker: Nicola Rowbotham, GB	15:44 - 16:06
	Exercise as Airway Clearance - Routes to designing studies to examine whether exercise can replace ACT Speaker: Zoe Saynor, GB	16:06 - 16:30
Workshop 15:00 - 16:30 WS05 - WS05: U	Inderstanding the impact of CFTR in epithelial biology	R6
Chair: Carlos M F		
Chair: Nicoletta F WS05.01	Pedemonte, IT  Characterisation of the molecular mechanisms underlying PI3Ky-dependent CFTR stability at the plasma membrane Abstract Presenter: Alessandra Murabito, IT	15:00 - 15:15
WS05.02	CFTR rescue by lumacaftor (VX-809) induces an extensive reorganisation of mitochondria in the cystic fibrosis bronchial epithelium  Abstract Presenter: Nicoletta Pedemonte, IT	15:15 - 15:30
WS05.03	Metabolomic impact of the restoration of CFTR activity in the respiratory epithelium  Abstract Presenter: Emmanuelle Bardin, FR	15:30 - 15:45
WS05.04	SARS-CoV-2 infection is reduced in both immortalised and primary CFTR-modulated human bronchial epithelial cells Abstract Presenter: Virginia Lotti, IT	15:45 - 16:00
WS05.05	Proximity profiling of the CFTR interaction landscape in response to Orkambi  Abstract Presenter: Melissa Iazzi, CA	16:00 - 16:15
WS05.06	Novel high-throughput screening tool for monitoring CFTR levels and localisation in cytoplasm membrane with luminescent peptide tag	16:15 - 16:30
	Abstract Presenter: Martin Ondra, CZ	
Workshop 17:00 - 18:30 WS06 - WS06: E	Expanding the use and knowledge on CFTR modulators	R1
Chair: Silke van K Chair: Damian G	Koningsbruggen-Rietschel, DE Downey, GB	
WS06.01	HIT-CF organoid screen with ELX-02 for people with CFTR nonsense mutations as a predictive tool for clinical response  Abstract Presenter: Marlou Bierlaagh, NL	17:00 - 17:15
WS06.02	Impact of one year of treatment with	17:15 - 17:30
	elexacaftor/tezacaftor/ivacaftor on clinical outcomes in	

	people with cystic fibrosis in a real-world setting - the RECOVER study	
	Abstract Presenter: Paul McNally, IE	
WS06.03	Novel CFTR modulator combinations maximise functional rescue of G85E and N1303K in rectal organoids	17:30 - 17:45
	Abstract Presenter: Marjolein Ensinck, BE	
WS06.04	Effects of elexacaftor/tezacaftor/ivacaftor therapy on CFTR function in patients with cystic fibrosis and one or two F508del alleles	17:45 - 18:00
	Abstract Presenter: Simon Y. Graeber, DE	
WS06.05	Trikafta-mediated bicarbonate transport correction in F508del primary cell cultures	18:00 - 18:15
	Abstract Presenter: Agathe Lepissier, FR	
WS06.06	The cystic fibrosis urine bicarbonate challenge test Abstract Presenter: Peder Berg, DK	18:15 - 18:30
<i>Workshop</i> 17:00 - 18:30		R2
<b>WS07 - WS07:</b>	Factors influencing lung function and survival	
Chair: Clémence Chair: Patrick Fl	· ·	
WS07.01	Investigating the effects of dornase alfa and hypertonic saline used in combination on lung function in people with cystic fibrosis	17:00 - 17:15
	Abstract Presenter: Emily Granger, GB	
WS07.02	Lumacaftor/ivacaftor in people with cystic fibrosis: factors predisposing the response and impact on lung function decline	17:15 - 17:30
	Abstract Presenter: Julie Mesinele, FR	
WS07.03	Does changing from lumacaftor/ivacaftor (Orkambi) to tezacaftor/ivacaftor + ivacaftor (Symkevi + ivacaftor) offer any clinical benefits in children with cystic fibrosis (CF)?  Abstract Presenter: Lucy Paskin, GB	17:30 - 17:45
WS07.04	Electronic Home Monitoring of Children with Cystic Fibrosis To Detect and Treat Acute Pulmonary exacerbations and their effect on one-year FEV <sub>1</sub> loss Abstract Presenter: Muruvvet Yanaz, TR	17:45 - 18:00
WS07.05	Investigating the relationship between lung function decline and time to death or lung transplantation, accounting for geographical variability	18:00 - 18:15
	Abstract Presenter: Elrozy Andrinopoulou, NL	
WS07.06	Geographical distance and survival among adult cystic fibrosis lung transplant recipients in the United States  Abstract Presenter: Shivani Patel, US	18:15 - 18:30

Workshop 17:00 - 18:30	Affect has showed in CE or identicle and why?	R3
Chair: Andreas Ju Chair: Lutz Naeh	- <del>-</del>	
WS08.01	Changing epidemiology of cystic fibrosis in Europe from 2010 to 2019: data from the European Cystic Fibrosis Society Patient Registry	17:00 - 17:15
	Abstract Presenter: Elpis Hatziagorou, GR	
WS08.02	Contemporary incidence of cystic fibrosis in Canada and the United States	17:15 - 17:30
	Abstract Presenter: Sanja Stanojevic, CA	
WS08.03	Healthcare-Associated Links in Transmission of Nontuberculous mycobacteria in People with Cystic Fibrosis (HALT NTM): a multi-centre study	17:30 - 17:45
	Abstract Presenter: Jane E. Gross, US	
WS08.04	Real-world data demonstrate clinical response in people with cystic fibrosis (pwCF) who have select residual function (RF) mutations and are treated with ivacaftor (IVA)	17:45 - 18:00
	Abstract Presenter: Mark Higgins, GB	
WS08.05	Parenthood impacts short-term health outcomes in people with cystic fibrosis  Abstract Presenter: Jennifer Taylor-Cousar, US	18:00 - 18:15
WS08.06		10.15 10.20
W 506.06	Investigating associations between air pollution and the severity of cystic fibrosis in Great Britain  Abstract Presenter: Muhammad Saleem Khan, GB	18:15 - 18:30
	, and the second	
Workshop		
17:00 - 18:30	ncluding people with CF and their families in delivering high-qu	R4
W309 - W309. 1	including people with Cr and then lamines in derivering mgn-qu	lanty care
Chair: Jacquelien	Noordhoek, NL	
WS09.01	Make cystic fibrosis research results accessible to all: a French initiative managed by patients and relatives Abstract Presenter: Estelle Ruffier, FR	17:00 - 17:15
MCOO OO	·	17.15 17.20
WS09.02	Setting up an e-learning program for male patients presenting cystic fibrosis infertility  Abstract Presenter: Sophie Ramel, FR	17:15 - 17:30
WS09.03	A review of patients' thoughts, feelings and priorities as evidenced in their responses to the paediatric cystic fibrosis	17:30 - 17:45
	annual review questionnaires  Abstract Presenter: Jill Watkinson, GB	
WS09.04	Psychometric characteristics of the CF Coping Self-Efficacy (CF-CSE) scale	17:45 - 18:00
	Abstract Presenter: Anna M. Georgiopoulos, US	
WS09.05	Using the CFHealthHub digital learning health system to optimise the nebulised medicines supply process in five UK adult cystic fibrosis (CF) centres	18:00 - 18:15

	Abstract Presenter: Sophie Dawson, GB	
WS09.06	A questionnaire to gain an understanding of adherence to inhaled therapies, airway clearance and exercise since commencing the triple combination modulator in cystic fibrosis	18:15 - 18:15
	Abstract Presenter: Paul Wilson, GB	
Workshop 17:00 - 18:30 <b>WS10 - WS10: P</b> Mandy Bryon	Physical health outcomes with or without modulator therapy	R5
Chair: Thomas Ra		
WS10.01	Impact of triple CFTR modulator therapy on airway clearance and nebuliser adherence in adults with cystic fibrosis	17:00 - 17:15
	Abstract Presenter: Louise Warnock, GB	
WS10.02	People with cystic fibrosis on elexacaftor/tezacaftor/ivacaftor therapy demonstrate improved physical activity levels and cardiovascular fitness Abstract Presenter: Thomas Simon FitzMaurice, GB	17:15 - 17:30
WS10.03	Impact of triple CFTR modulator therapy on urinary incontinence symptoms in adults with cystic fibrosis  Abstract Presenter: Alice Midwinter, GB	17:30 - 17:45
WS10.04	Exploring the effects of Kaftrio on the physiotherapists' role and service provision for people with cystic fibrosis: a UK and Republic of Ireland (ROI) survey.  Abstract Presenter: Orla Aisling O'Beirne, GB	17:45 - 18:00
WS10.05	Physical fitness and habitual physical activity in adults with cystic fibrosis - do they improve with elexacaftor/tezacaftor/ivacaftor therapy?  Abstract Presenter: Wolfgang Gruber, DE	18:00 - 18:15
WS10.06	Adherence to nebulised medication in paediatric patients with cystic fibrosis following introduction of modulator therapy	18:15 - 18:30
	Abstract Presenter: Mostin Hu, GB	
<i>Workshop</i> 17:00 - 18:30		R6
	Iow much CF is it? Revisiting the dilemma of CF diagnosis	
Chair: Carlo Cast Chair: Philippe Ro		
WS11.01	Divergent diagnostic assessment of cystic fibrosis and related disorders: an analysis of paediatric cases categorisation by 50 international specialists  Abstract Presenter: Carlo Castellani, IT	17:00 - 17:15

WS11.02	Inconsistent IRT threshold values and CFTR panels in newborn screening for cystic fibrosis across the United States	17:15 - 17:30
	Abstract Presenter: Maryann Riyadh Rehani, US	
WS11.03	Repeated sweat testing in children with cystic fibrosis transmembrane conductance regulator-related metabolic syndrome/cystic fibrosis screen-positive, inconclusive diagnosis (CRMS/CFSPID)	17:30 - 17:45
	Abstract Presenter: Magdalena Postek, PL	
WS11.04	Confirmatory genetic testing for all cystic fibrosis screen positive newborns: a 12-year analysis	17:45 - 18:00
	Abstract Presenter: Steven Zhang, CA	
WS11.05	Clinical, microbiological and functional outcomes of children with inconclusive diagnosis of cystic fibrosis following newborn screening (CFSPID/CRMS)	18:00 - 18:15
	Abstract Presenter: Thomas Perrin, FR	
WS11.06	A 14-year review of cystic fibrosis newborn screening outcomes from a UK regional laboratory  Abstract Presenter: Sarah Driscoll, GB	18:15 - 18:30

Satellite Symposium 19:00 - 20:30

R3

### **Satellite Symposium**

Please find the detailed programme of the Satellite Symposia here.

### Friday, 10. June 2022

*Symposium* 08:30 - 10:00 R1 S13 - Symposium 13 - What's next for CFTR modulators? Chair: Nicholas Simmonds, GB Chair: Dorota Sands, PL Stopping trials in CF patients on CFTR modulators 08:30 - 08:52 Speaker: Gwyneth Davies, GB The value of monitoring CFTR modulator drug levels 08:52 - 09:14 Speaker: Martin Hug, DE Lessons from modulator studies in people with rare residual 09:14 - 09:36 function mutations Speaker: Nicoletta Pedemonte, IT Biomarkers for the personalized assessment of a 09:36 - 10:00

Symposium 08:30 - 10:00

R2

### S14 - Symposium 14 - Lung transplantation - 2022 update

therapeutic response

Speaker: Kors Van der Ent, NL

### At the end of the session, the participant will be able to:

- Examine the current contra-indications to lung transplant in a CF population
- · Evaluate the potential role of CFTR modulators post lung transplantation
- · Discuss the indications and complexities of double organ transplantation in cystic fibrosis
- Identify challenges associated with lung transplantation in the paediatric period

Chair: Kathleen J. Ramos, US Chair: Paul Aurora, GB

Contraindications to lung transplant	08:30 - 08:52
Speaker: Antoine Roux, FR	
<b>CFTR modulator use post lung transplantation</b> Speaker: Kathleen J. Ramos, US	08:52 - 09:14
<b>Double organ transplantation - indications and outcomes</b> Speaker: Lieven Dupont, BE	09:14 - 09:36
Specific challenges with pediatric lung transplantation	09:36 - 10:00

*Symposium* 08:30 - 10:00

R3

### S15 - Symposium 15 - "It's life, Jim, but not as we know it"

Speaker: Paul Aurora, GB

At the end of the session the participants will be able to:

This symposium looks at the changing face of CF – not just because of the advent of CFTR modulators but also the gradual changes that have been happening for some time (improved survival; an aging CF population; bigger adult centres; technological improvements allowing remote monitoring and home working; a global pandemic, a changing job market etc.)

Chair: Lynsay Kinnaird, GB Chair: Ulrike Smrekar, AT

Changing the way we think - coaching and caring for the 08:30 - 08:52

well CF patient.

Speaker: Majda Oštir, SI

Changing the way we think - patient perspectives on 08:52 - 09:14 treatment priorities

Speaker: Jutta Bend, DE

Changing the way we think about the aging CF patient: 09:14 - 09:36

what do we need to know?

Speaker: Yvonne Prins, NL

Changing the way we think - Adjusting and adapting CF 09:36 - 10:00

services

Speaker: Charlotte Dawson, GB

Symposium 08:30 - 10:00

R4

### S16 - Symposium 16 - Nutrition in cystic fibrosis

### At the end of the session, the participant will be able to:

- $\cdot$  gain a greater understanding of nutritional requirements for people with CF in view of advances in medical treatments
- Explore situations when nutritional support might be needed and how it can be delivered
- Gain understanding of how different CFTR modulators influence eating habits and behaviours
- Explore changes in diet and lifestyle that people with CF have adopted in the era of CFTR modulatory therapy
- provide an update on how current apps available in Europe can support the self-management of PERT for people with CF

Chair: Elizabeth Owen, GB Chair: Dimitri Declercq, BE

There is no longer a role for nutrition support in cystic	08:30 - 08:52
fibrosis - True	

Speaker: Darren Sills, GB

There is no longer a role for nutrition support in cystic 08:52 - 09:14 fibrosis - False

Speaker: Monika Mielus, PL

Impact of CFTR modulators on eating habits and behaviour 09:14 - 09:36 (including pt perspective)

Speaker: Sarah Collins, GB

Use of mobile applications to support self-managment of 09:36 - 10:00 PERT (look at future adaptions e.g changing enzyme requirements on modulators)

Speaker: Mieke Boon, BE

Symposium 08:30 - 10:00

R5

## S17 - Symposium 17 - Best of Journal of Cystic Fibrosis, Lancet Respiratory Medicine and European Respiratory Journal

Chair: Patrick Flume, US Chair: Emma Grainger, GB Chair: Marcus Mall, DE

**Introduction** 08:30 - 08:45

JCF - Antisense oligonucleotide splicing modulation as a novel Cystic Fibrosis therapeutic approach for the W1282X Nonsense Mutation	08:45 - 09:05
Speaker: Batsheva Kerem, IL	
Discussant: Donald R. VanDevanter, US	
ERJ - Forskolin-induced Organoid Swelling is Associated with Long-term CF Disease Progression	09:05 - 09:25
Speaker: Danya Muilwijk, NL	
Discussant: Simon Y. Graeber, DE	
LRM - The effect of azithromycin on structural lung disease in infants with cystic fibrosis (COMBAT CF): a phase 3, randomised, double-blind, placebo-controlled clinical trial	09:25 - 09:45
Speaker: Harm Tiddens, NL	
Discussant: Margaret Rosenfeld, US	

Symposium

09:45 - 10:00

08:30 - 10:00 R6

# S18 - Symposium 18 - Enhancing CF registry studies through linkage with other data sources At the end of the session, the participant will be able to:

- $\cdot$   $\,$  Demonstrate knowledge of the challenges and opportunities of linking CF registries with other data sources.
- · Discuss data on environmental factors in lung function decline.

Speaker: Andreas Jung, CH

**Editors'topic** 

- · Recognize how insight from the patient voice can be integrated into CF registries.
- $\cdot$  Describe the challenges and opportunities of linking national CF registries for the purposes of collecting unified data.

Chair: Vincent Gulmans, NL Chair: Uros Krivec, SI

France (Lyon): "Linkage between the French CF Registry and health insurance reimbursement data"  Speaker: Isabelle Durieu, FR	08:30 - 08:52
US/Netherlands: "Linking US registry data with geomarker data: longitudinal associations between environmental factors and lung function decline.  Speaker: Elrozy Andrinopoulou, NL	08:52 - 09:14
UK: "Context is key - weaving patient voice and wider health system insight into the UK CF Registry".  Speaker: Rebecca Cosgriff, GB	09:14 - 09:36
Switzerland/European CF Registry: "Building a COVID-CF registry through linkage between national registries."	09:36 - 10:00

ECFS Tomorrow Lounge Session 10:00 - 10:30

CF plain language glossary launch

*Symposium* 10:30 - 12:00

R1

R2

11:36 - 12:00

### S19 - Symposium 19 - Ageing with cystic fibrosis

### At the end of the session, the participant will be able to:

- · Predict the likely demographics of people with cystic fibrosis into the future
- · Discuss the challenges of cystic fibrosis diabetes and its impact on ageing with cystic fibrosis
- Analyze the role that renal disease may have in the ageing cystic fibrosis population
- $\cdot$  Define the risks associated with obesity in cystic fibrosis and identify challenges associated with cardiovascular health

Chair: Eitan Kerem, IL Chair: Stuart Elborn, GB

The evolving age profile of people with cystic fibrosis - a view to the future	10:30 - 10:52
Speaker: Ruth Keogh, GB	
Cystic fibrosis related diabetes  Speaker: Bruce C. Marshall, US	10:52 - 11:14
Renal disease in cystic fibrosis - an emerging problem?  Speaker: William Plant, IE	11:14 - 11:36
<b>Obesity and cardiovascular health</b> Speaker: Francesco Blasi, IT	11:36 - 12:00

10:30 - 12:00 **S20 - Symposium 20 - Real world data with CFTR modulators** 

Chair: Silke van Koningsbruggen-Rietschel, DE

Chair: Pierre-Régis Burgel, FR

Symposium

Polypharmacy and CFTR modulators in real life	10:30 - 10:52
Speaker: Edwin Brokaar, NL	
<b>Post-authorisation studies: what can we learn from them?</b> Speaker: Lutz Naehrlich, DE	10:52 - 11:14
Ensuring adherence in patients on CFTR modulators	11:14 - 11:36
Speaker: Peter Barry, GB	

CFTR modulators and disease modification: what is the effect on life expectancy?

Speaker: Pierre-Régis Burgel, FR

Symposium 10:30 - 12:00

R3

## ${\it S21}$ - Symposium 21 - The "haves" and the "have nots" - Psychological sequelae of CFTR modulator therapies

#### At the end of the session, the participant will be able to:

- $\cdot$  appreciate the psychological impact of not being able to access CFTR modulator therapy. It will look at those patients who have the right genotypes but aren't able to access the treatment due to financial or political reasons and then will consider those patients who don't have the genotypes necessary to access treatment.
- $\cdot$  appreciate patient expectations of modulator therapies vs patient experience (based on a study looking at the pre-CFTRm counselling given to patients prior to commencement of therapy and what patients wished they had known in hindsight.)
- $\cdot$  Consider the topic of anxiety associated with CFTR modulators. How much is a side-effect and how can we separate this from anxiety relating to the changes CFTR modulators bring.
- apreciate the challenges facing patients whose life expectancy may now be extended what this means for their life choices / prospects / body image / identity.

Chair: Marieke Verkleij, NL Chair: Kathleen Thickett, GB

, -	
Left out in the cold - impact of ineligibility and inaccessibility	10:30 - 10:52
Speaker: Urszula Borawska-Kowalczyk, PL	
Why didn't you warn me? - patient perspectives of modulator therapies	10:52 - 11:14
Speaker: Nichola MacDuff, GB Speaker: Angela Holden, GB	
Anxiety and depression - side effects or mourning missed opportunities?	11:14 - 11:36
Speaker: Anna M. Georgiopoulos, US	
Identity crisis? - Rethinking and reshaping your life	11:36 - 12:00

Symposium 10:30 - 12:00

R4

#### S22 - Symposium 22 - Impact of biofilm in cystic fibrosis

### At the end of the session, the participant will be able to:

· Identify the principal polymicrobial interactions in biofilms in CF

Speaker: Jacquelien Noordhoek, NL

- Appraise the risk of nosocomial infection in people with CF due to biofilm
- Synthetize knowledge on approaches to solving biofilm issues
- · Understand the impact of biofilm-associated pathogens on the immune response

Chair: Jennifer Bomberger, US

Chair: Tom Coenye, BE

Interactions between viruses and bacterial biofilms in the respiratory tract	10:30 - 10:52
Speaker: Catherine Armbruster, US	
Immune responses to <i>Pseudomonas aeruginosa</i> biofilm Infections  Speaker: Claus Moser, DK	10:52 - 11:14
Biofilms in the hospital environment as a risk factor for patient with cystic fibrosis  Speaker: Jane E. Gross, US	11:14 - 11:36
Innovative approaches to treat biofilm-related infections in	11:36 - 12:00

### **CF** patients

Speaker: Tom Coenye, BE

Symposium 10:30 - 12:00

R5

# S23 - Symposium 23 - The changing face of newborn screening for cystic fibrosis At the end of the session, the participant will be able to:

- Discuss current and future challenges to optimizing CF newborn screening programmes
- · Understand the challenge of extended newborn screening panels and optimization of their performance for local populations
- · Be aware of the issues involved when considering extending genetic screening panels

Chair: Jeannette Dankert-Roelse, NL

Chair: Dario Prais, IL

**Towards optimization of the newborn screening algorithm**10:30 - 10:52 **in the Netherlands**Speaker: Karin M. de Winter - de Groot, NL

**Extended variant panels in newborn screening programmes:** 10:52 - 11:14 is it still an option?

Speaker: Denise Kay, US

Screening around the globe: feasibility and challenges in  $$11:14\mbox{ - }11:36$$  LMICs

Speaker: Marco Zampoli, ZA

*Symposium* 10:30 - 12:00

R6

# S24 - Symposium 24 - Advances in gene and cell therapies for cystic fibrosis At the end of the session, the participant will be able to:

- Summarise the state-of-the-art of gene therapy for cystic fibrosis
- · Review the therapeutic potential of different CFTR gene editing strategies
- $\cdot$  Assess understanding of airway epithelial cell populations for the development of stem cell therapies for CF

Chair: Anna Cereseto, IT Chair: Marianne S. Carlon, BE

Pan CFTR mutation correction using CRISPR/Cas9 Speaker: Matthew Porteus, US	10:30 - 10:52
CRISPR genome editing to repair CFTR splicing mutations  Speaker: Giulia Maule, IT	10:52 - 11:14
Antisense oligonucleotide-based drugs to correct premature stop codon and splice variants in cystic fibrosis	11:14 - 11:36
Speaker: Yifat Oren, IL	
Genetically engineered induced pluripotent stem cells (iPSCs) as disease models to test cystic fibrosis therapies	11:36 - 12:00

Satellite Symposium 12:30 - 14:00

R1

#### **Satellite Symposium**

Please find the detailed programme of the Satellite Symposia here.

Speaker: Ulrich Martin, DE

Meet the Experts

12:45 - 13:45

## ${\it Meet the Experts}$ - Meet the experts 6 - Management of drug interactions with CFTR modulators

ePoster Corner C

Expert: Martin Hug, DE Expert: Nicola J. Shaw, GB

Meet the Experts 12:45 - 13:45

## *Meet the Experts* - Meet the experts 4 - 3 CFTR databases (CFTR1, CFTR2 and CFTR-France), 3 complementary sources of information for the clinician

ePoster Corner A

Expert: Caroline Raynal, FR Expert: Karen Raraigh, US Expert: Carlo Castellani, IT

*Meet the Experts* 12:45 - 13:45

# Meet the Experts - Meet the Experts 5 - How to cope with the fast lane and the slow lane in the clinic: caring for patients who are not eligible and those who are eligible for CFTR modulators

ePoster Corner B

Expert: Carsten Schwarz, DE Expert: Maya Kirszenbaum, FR

*Meet the Experts* 12:45 - 13:45

### **Meet the Experts**

ECFS Tomorrow Lounge Session 12:45 - 13:45

#### Physiotherapy in the 20s

Speaker: Gemma E Stanford, GB Discussant: Brenda Button, AU

ePoster Session 14:00 - 15:00

#### **Poster Viewing 2**

P112	Shift of the lung microbiota in patients with cystic fibrosis following antibiotic therapy selected with AtbFinder  Abstract Presenter: Maria Vecherkovskaya, RU	14:00 - 14:00
P113	Microbiology assessment in cystic fibrosis patients on elexacaftor/tezxacaftor/ivacaftor  Abstract Presenter: Nadia Mazzoni, IT	14:00 - 14:00
P114	Airway colonisation by filamentous fungi in patients with cystic fibrosis from a tertiary care hospital in Madrid: a two-year prevalence study	14:00 - 14:00
	Abstract Presenter: Alfonso Pascual del Riquelme, ES	
P115	Comparison of total bacterial and <i>Pseudomonas aeruginosa</i> load during PEx between infrequent and frequent	14:00 - 14:00

	exacerbators  Abstract Presenter: Claire J Houston, GB	
P116	Effects of Symkevi <sup>TM</sup> (tezacaftor/ivacaftor) on the lung and gut microbiota in cystic fibrosis  Abstract Presenter: Ryan Marsh, GB	14:00 - 14:00
P117	Fungal colonisation in patients with cystic fibrosis: preliminary results from a national multicentre study Abstract Presenter: Ainhize Maruri-Aransolo, ES	14:00 - 14:00
P118	Airways respiratory viral infections in cystic fibrosis  Abstract Presenter: Daniela Dolce, IT	14:00 - 14:00
P119	CFTR modulators therapy and lung microbiota diversity in adult cystic fibrosis patients  Abstract Presenter: Olga Voronina, RU	14:00 - 14:00
P120	Metaproteomics profiling of the respiratory microbiota of cystic fibrosis (CF) patients infected by <i>Mycobacterium abscessus</i>	14:00 - 14:00
	Abstract Presenter: Pauline Hardouin, FR	
P122	Identification and characterisation of a Liverpool Epidemic Strain (LES) isolate of <i>Pseudomonas aeruginosa</i> , first collected in 1986 in Calgary, Canada <i>Abstract Presenter</i> : Conrad Izydorczyk, CA	14:00 - 14:00
P123	Role of specialised proresolving lipid mediators in the interaction between Aspergillus fumigatus and cystic fibrosis bronchial epithelial cells  Abstract Presenter: Valerie Urbach, FR	14:00 - 14:00
P124	Collateral sensitivity in multidrug-resistant <i>Pseudomonas</i> aeruginosa  Abstract Presenter: Mette Kolpen, DK	14:00 - 14:00
P125	Genotype-phenotype correlation of triazole-resistant pulmonary aspergillosis in chronic respiratory disease patients	14:00 - 14:00
P126	Abstract Presenter: Renad Aljohani, GB  Pseudomonas aeruginosa infection during long-term suppression treatment with tobramycin inhalation powder (TIP)  Abstract Presenter: Ross P. McCleave, GB	14:00 - 14:00
P127	Haemophilus influenza and antimicrobial resistance in children and young people with cystic fibrosis  Abstract Presenter: Christopher Edwards, GB	14:00 - 14:00
P128	Epidemiological significance of Achromobacter spp. chronic lung infection in patients with cystic fibrosis  Abstract Presenter: Marina Chernukha, RU	14:00 - 14:00
P129	Evidence for a role of Achromobacter xylosoxidans VBNC forms in chronic cystic fibrosis lung infection  Abstract Presenter: Nadia Mazzoni, IT	14:00 - 14:00
P130	Achromobacterspp. phenotypic differences between chronic	14:00 - 14:00
	Tr. r	

	and occasional lung infection in cystic fibrosis  Abstract Presenter: Giulia Maria Saitta, IT	
P131	Clinical impact of Aspergillus fumigatus in children with cystic fibrosis  Abstract Presenter: Giovanna Pisi, IT	14:00 - 14:00
P132	Microbiological monitoring of chronic lung infection with Achromobacter spp. in cystic fibrosis patients Abstract Presenter: Lusine Avetisyan, RU	14:00 - 14:00
P133	Prevalence and factors associated with isolation of Aspergillus from sputum in patients with cystic fibrosis (CF) Abstract Presenter: Elena Gjinovska-Tasevska,	14:00 - 14:00
P134	Prevalence of multi-drug antimicrobial-resistant bacteria in children with cystic fibrosis  Abstract Presenter: Giovanna Pisi, IT	14:00 - 14:00
P135	Comparative characteristics of patients with nontuberculous mycobacteria (NTM) in the Russian Federation according to the 2020 Registry  Abstract Presenter: Elena Kondratyeva, RU	14:00 - 14:00
P138	Biochemical detection of <i>Pseudomonas aeruginosa</i> in sputum and urine from children with cystic fibrosis  Abstract Presenter: Hatice Nur Cömert, DK	14:00 - 14:00
P139	Colistin-rifampicin combination is efficient against biofilms of colistin-resistant <i>Pseudomonas aeruginosa</i> from cystic fibrosis patients  Abstract Presenter: Oana Ciofu, DK	14:00 - 14:00
P140	Eradication treatment of <i>Pseudomonas aeruginosa</i> infections in children with cystic fibrosis  Abstract Presenter: Aynur Gulieva, TR	14:00 - 14:00
P141	Interaction between bacteria in the cystic fibrosis airways Abstract Presenter: Enna E. Gibson, GB	14:00 - 14:00
P142	Upper and lower airways microbiological status in cystic fibrosis patients in stable conditions and in lung transplant patients  Abstract Presenter: Daniela Dolce, IT	14:00 - 14:00
P143	Bronchopulmonary infection/colonisation in Spanish cystic fibrosis patients: preliminary results of a multicentre study Abstract Presenter: Ainhize Maruri-Aransolo, ES	14:00 - 14:00
P144	Impact of the sex steroid hormone estradiol on biofilm formation and phenotype of <i>Pseudomonas aeruginosa</i> isolates from cystic fibrosis patients  Abstract Presenter: Jiwar Al-Zawity, DE	14:00 - 14:00
P145	Monitoring of pulmonary infections in the danish cystic fibrosis cohort during CFTR modulator therapy implementation  Abstract Presenter: Majbritt Jeppesen, DK	14:00 - 14:00
P146	Microbiological diagnostic procedures for respiratory cystic	14:00 - 14:00

	fibrosis samples: results of a survey in Italian laboratories  Abstract Presenter: Daniela Dolce, IT	
P147	In vitro sensitivity of Gram-negative cystic fibrosis isolates to a 4 <sup>th</sup> generation fluoroquinolone Abstract Presenter: Patrick Farrell, GB	14:00 - 14:00
P148	Parental perspectives with sending respiratory tract specimens for microbiology from home in children with cystic fibrosis - experience from a tertiary service provider in the Northwest United Kingdom	14:00 - 14:00
	Abstract Presenter: Anirban Maitra, GB	
P149	The association between the cumulative dose of aminoglycoside exposure and hearing loss in children with cystic fibrosis	14:00 - 14:00
	Abstract Presenter: Cansu Yilmaz Yegit, TR	
P150	Antimicrobial prescribing in people with cystic fibrosis: exploring inhaled antibiotic use for <i>Pseudomonas aeruginosa</i> infections across the ECFS-CTN	14:00 - 14:00
	Abstract Presenter: Callum Matthew Sloan, GB	
P151	Drug sensitivity of <i>Mycobacterium abscessus</i> in patients with cystic fibrosis	14:00 - 14:00
	Abstract Presenter: Elena Kondratyeva, RU	
P152	Trends in intravenous antibiotic prescriptions pre- and post- introduction of Kaftrio in a large UK adult cystic fibrosis centre	14:00 - 14:00
	Abstract Presenter: Jocelyn Choyce, GB	
P153	Hypersensitivity reactions to antibiotics in patients with cystic fibrosis  Abstract Presenter: Aleksandra Kowalik, SE	14:00 - 14:00
P154	Cough swabs are not a suitable alternative to a sputum culture for fungal culture to isolate Aspergillus fumigatus.  Abstract Presenter: Natalie Francis, GB	14:00 - 14:00
P155	Paws for thought: sniffer dogs for infection surveillance in non-sputum-producing people with cystic fibrosis Abstract Presenter: John King, GB	14:00 - 14:00
P156	Patterns of liver dysfunction and the development of a pathway for liver monitoring in adults with cystic fibrosis (CF) on elexacaftor/tezacaftor/ivacaftor (ETI): a quality improvement project  Abstract Presenter: James Sun, GB	14:00 - 14:00
P157	A modified rapid review of gastrointestinal symptoms in people with cystic fibrosis on cystic fibrosis transmembrane regulator modulator therapies  Abstract Presenter: Darren Sills, GB	14:00 - 14:00
P158	What is the impact of elexacaftor/tezacaftor/ivacaftor (ETI) therapy on clinical response of cystic fibrosis patients aged ≥ 40 years and their co-morbidities?  Abstract Presenter: Karuna Sapru, GB	14:00 - 14:00
	Anomati resemen. Karama sapra, OD	

P159	Modifications of anthropometric parameters and body	14:00 - 14:00
1 100	composition after Kaftrio in a group of adolescents and young adults	11100 11100
	Abstract Presenter: Silvia Rigon, IT	
P160	"Seeing a trend" - increasing vitamin A levels on elexacaftor/tezacaftor/ivacaftor therapy Abstract Presenter: David Proud, GB	14:00 - 14:00
P161	Body mass index change in adult patients with cystic fibrosis following the introduction of triple CFTR therapy ivacaftor/tezacaftor/elexacaftor: a regional adult cystic fibrosis centre experience	14:00 - 14:00
	Abstract Presenter: Neil Patel, GB	
P162	Body composition assessment in cystic fibrosis patients on elexacaftor/texacaftor/ivacaftor	14:00 - 14:00
	Abstract Presenter: Veronica Zamponi, IT	
P163	Single-centre experience of changes in nutritional parameters in the first 12 months of elexacaftor/tezacaftor/ivacaftor (ETI / Kaftrio) treatment	14:00 - 14:00
	Abstract Presenter: Joanna Snowball, GB	
P164	Nutritional status and circulating levels of fat-soluble vitamins in cystic fibrosis: cross-sectional analysis and effect of elexacaftor/tezacaftor/ivacaftor	14:00 - 14:00
	Abstract Presenter: Michela Francalanci, IT	
P165	Exploring weight change and subjective experiences on Kaftrio in adults with cystic fibrosis	14:00 - 14:00
	Abstract Presenter: Lidia Sheibani, GB	
P166	"The party is over because I now have a sensible diet" - the experience of people with cystic fibrosis on CFTR modulators	14:00 - 14:00
	Abstract Presenter: Sarah Collins, GB	
P167	Pancreatic enzyme replacement therapy intake and gastrointestinal symptoms in adults with cystic fibrosis: a cross-sectional study in Copenhagen, Denmark	14:00 - 14:00
	Abstract Presenter: Mette Frahm Olsen, DK	
P168	Gastrointestinal (GI) symptoms and their impact on quality of life in cystic fibrosis (CF) patients  Abstract Presenter: Ivana Arnaudova Danevska,	14:00 - 14:00
P169	Psychometric properties of the gastrointestinal symptom tracker self-report measure	14:00 - 14:00
	Abstract Presenter: Sonia Graziano, IT	
P170	An assessment of terminal ileum morphology, using magnetic resonance imaging, in people with cystic fibrosis Abstract Presenter: Grace Lim, GB	14:00 - 14:00
P171	Improved diagnosis and treatment of distal intestinal obstruction syndrome (DIOS) with a simplified treatment routine	14:00 - 14:00
	Abstract Presenter: Marita Gilljam, SE	

P172	Screening for constipation in cystic fibrosis (CF): what are we missing?	14:00 - 14:00
	Abstract Presenter: Jennifer Still, GB	
P173	Increased prevalence of celiac disease in patients with cystic fibrosis: a systematic review and meta analysis  Abstract Presenter: Marcell Imrei, HU	14:00 - 14:00
P174	Effects of proton pump inhibitor treatment on nutritional status and respiratory infection risk in cystic fibrosis: a matched cohort study  Abstract Presenter: CALOGERO SATHYA SCIARRABBA, IT	14:00 - 14:00
P175	How accurate is the glucose management indicator calculated by continuous glucose monitoring compared to the blood HbA1c reading in patients with cystic fibrosis?  Abstract Presenter: Dion-Emily Manning, GB	14:00 - 14:00
P177	Adults with Cystic Fibrosis-Related Diabetes have a significantly elevated rate of gastrointestinal symptoms assessed with the CFAbd-Score  Abstract Presenter: Laura Caley, GB	14:00 - 14:00
	•	
P178	A feasibility assessment of delivering a glycaemic index dietary intervention for managing glucose abnormalities in people with cystic fibrosis	14:00 - 14:00
	Abstract Presenter: Laura Birch, GB	
P179	Glycaemic index dietary intervention in cystic fibrosis: preliminary findings from a feasibility study of dietary manipulation	14:00 - 14:00
	Abstract Presenter: Laura Birch, GB	
P180	Clinical practice versus guidelines for the screening of Cystic Fibrosis-Related Diabetes (CFRD): a French survey of 47 centres	14:00 - 14:00
	Abstract Presenter: Laurence Weiss, FR	
P181	The effects of linagliptin in patients with a diagnosis of Cystic Fibrosis-Related Diabetes	14:00 - 14:00
	Abstract Presenter: Josie Cunningham, GB	
P182	Adiponectin, glucose metabolism and body composition in cystic fibrosis	14:00 - 14:00
	Abstract Presenter: Bibi Uhre Nielsen, DK	
P183	Informative value of fructosamine in diagnosis of carbohydrate metabolism disorders in children with cystic fibrosis	14:00 - 14:00
	Abstract Presenter: Nadezhda Lyabina, RU	
P184	An evaluation of virtual Cystic Fibrosis-Related Diabetes (CFRD) clinics in the adult service during the covid-19 pandemic in Northern Ireland (NI)  Abstract Presenter: Emma Molloy, GB	14:00 - 14:00
P186	High vitamin A in children under 5 years of age Abstract Presenter: Órla Cahill, GB	14:00 - 14:00
P187	Nutritional status and its relation with lung function and	14:00 - 14:00

	vitamin D3 level in cystic fibrosis patients	
	Abstract Presenter: Ivana Arnaudova Danevska,	
P188	Vitamin D status at Bristol Adult CF Centre (BACFC) during the COVID-19 pandemic and following the introduction of the multivitamin Paravit CF	14:00 - 14:00
	Abstract Presenter: Anna Keele, GB	
P189	Assessment of total dietary salt intake: highlighting the need for regular detailed dietetic analysis of total salt intake in Liverpool, UK paediatric cystic fibrosis centre	14:00 - 14:00
	Abstract Presenter: Claire A. Berry, GB	
P190	Features of dosing of enzyme replacement therapy in children with cystic fibrosis in the Russian Federation: cross-sectional study	14:00 - 14:00
	Abstract Presenter: Tatyana Maksimycheva, RU	
P191	Self-compassion explains the positive relationship of mindful eating to uncontrolled and emotional eating in adults with cystic fibrosis	14:00 - 14:00
	Abstract Presenter: Michail Mantzios, GB	
P192	Early adrenal activation in boys with cystic fibrosis may explain impaired final height	14:00 - 14:00
	Abstract Presenter: Gizem Tamer, NL	
P193	Thermal sensation and clinical characteristics of patients with cystic fibrosis during extreme heat conditions	14:00 - 14:00
	Abstract Presenter: Pinelopi Anagnostopoulou, CH	
P194	Evaluation of bone mineral density indicators in children with cystic fibrosis: dynamics of indicators over 15 years  Abstract Presenter: Elena Zhekaite, RU	14:00 - 14:00
P196	Detecting bone disease in patients with cystic fibrosis: influence of genetic and clinical factors over reduced bone mass	14:00 - 14:00
	Abstract Presenter: Tatjana Jakovska Maretti,	
P197	Bone health status over time in people with cystic fibrosis and adherence to assessment of bone health guidelines at University Hospital Limerick	14:00 - 14:00
	Abstract Presenter: Audrey Tierney, IE	
P198	Bone health monitoring and treatment at an adult cystic fibrosis centre	14:00 - 14:00
	Abstract Presenter: Darren Sills, GB	
P199	Bone health and associated factors at an adult cystic fibrosis unit	14:00 - 14:00
D0.00	Abstract Presenter: Darren Sills, GB	44.00 44.00
P200	The prevalence of laryngopharyngeal reflux and sino-nasal symptoms in adults with cystic fibrosis  Abstract Presenter: Stephen Bourke, GB	14:00 - 14:00
P201	Bowel screening for cancer in pre-transplant people with cystic fibrosis and the accuracy of faecal immunochemical testing	14:00 - 14:00

	Abstract Presenter: Aoife Lillis, GB	
P202	CFTR modulation may help refine transplant decisions in lung-liver transplant candidates  Abstract Presenter: Christiane Knoop, BE	14:00 - 14:00
P203	Cystic fibrosis-related diseases at the Centre for Cystic Fibrosis, Institute for Respiratory Diseases in Children in Skopje, Republic of North Macedonia  Abstract Presenter: Sonja Momchilovikj,	14:00 - 14:00
P204	Elexacaftor/tezacaftor/ivacaftor - positive effects on quality of life using the AWESCORE and exercise capacity using the A-STEP in adults with end stage cystic fibrosis  Abstract Presenter: Brenda Button, AU	14:00 - 14:00
P205	Kaftrio improves $VO_{2max}$ in patients with cystic fibrosis Abstract Presenter: Thomas Kent, GB	14:00 - 14:00
P207	Spirometric values and six-minute walking distance in cystic fibrosis patients on elexacaftor/texacaftor/ivacaftor Abstract Presenter: Arianna Peruzzi, IT	14:00 - 14:00
P208	A retrospective observational study of the effects of Kaftrio on the current cohort of patients at Nottingham University Hospitals NUH trust  Abstract Presenter: Kate Hardiman, GB	14:00 - 14:00
P209	Physical fitness and habitual physical activity in children with cystic fibrosis - do they improve with elexacaftor/tezacaftor/ivacaftor therapy?	14:00 - 14:00
P210	Abstract Presenter: Stefanie Dillenhöfer, DE  Association of anaerobic threshold variables from cardiopulmonary exercise testing with the risk for exacerbation in patients with cystic fibrosis	14:00 - 14:00
P211	Abstract Presenter: Natália Evangelista Campos, BR  Ventilatory parameters during incremental exercise in patients with cystic fibrosis	14:00 - 14:00
	Abstract Presenter: Virginia D Alessandro, AR	
P212	The impact of cystic fibrosis-related diabetes on exercise capacity  Abstract Presenter: Richard Porter, GB	14:00 - 14:00
P213	The three-minute step test as a predictor of exacerbations in children and adolescents with cystic fibrosis  Abstract Presenter: Márcio Vinícius Fagundes Donadio, BR	14:00 - 14:00
P214	A comparison of airway clearance devices in adult cystic fibrosis patients: NIPPY Clearway2 versus Intermittent Positive Pressure Breathing (IPPB)  Abstract Presenter: Verity Yarwood, GB	14:00 - 14:00
P215	Airway clearance physiotherapy and health-related quality of life in cystic fibrosis - a substudy of a series of n-of-1 randomised controlled trials	14:00 - 14:00
	Abstract Presenter: Sandra Gursli, NO	
P216	The short-term influence of chest physiotherapy on lung	14:00 - 14:00

	function parameters in children with cystic fibrosis and primary ciliary dyskinesia	
	Abstract Presenter: Mieke Boon, BE	
P217	Exercise as airway clearance therapy (ExACT) in cystic fibrosis: a UK-based e-Delphi survey of patients, caregivers and health professionals	14:00 - 14:00
	Abstract Presenter: Zoe Saynor, GB	
P218	Calibration and cross-validation of accelerometry in children and adolescents with cystic fibrosis  Abstract Presenter: Mayara S. Bianchim, GB	14:00 - 14:00
P219	Establishing a review process for patients using non- invasive ventilation for airway clearance in the St. Bartholomew's cystic fibrosis adult population Abstract Presenter: Georgie Goodwin, GB	14:00 - 14:00
P220	Assessment of use of Manchester musculoskeletal screening tool in cystic fibrosis adults  Abstract Presenter: Nicola Hodgson, GB	14:00 - 14:00
P221	Tension-free vaginal tape - a 15-year review of effect on urinary incontinence in patients with cystic fibrosis  Abstract Presenter: Verity Yarwood, GB	14:00 - 14:00
P222	To assess measured frailty compared to multidisciplinary perception of frailty in the adult cystic fibrosis population Abstract Presenter: Paul Wilson, GB	14:00 - 14:00
P223	Bare Bones - a re-audit of bone health management in patients with cystic fibrosis at St. Vincent's University Hospital	14:00 - 14:00
	Abstract Presenter: Sarah Kelly, IE	
P224	Evolution of dry powder inhaled colistimethate sodium (Colobreathe) in a large adult cystic fibrosis centre  Abstract Presenter: Verity Yarwood, GB	14:00 - 14:00
P225	Long-term tolerability and use of tobipodhaler in people with cystic fibrosis  Abstract Presenter: Zelda Beverley, GB	14:00 - 14:00
P226	Our evolving prescribing practice: inhaled medication prescription changes over five years at the West Midlands Adult Cystic Fibrosis Centre	14:00 - 14:00
	Abstract Presenter: Catherine Brown, GB	
P227	"CF Hero" application as a motivational and therapeutic tool for kids and teenagers with cystic fibrosis Abstract Presenter: Martin Jirásek, CZ	14:00 - 14:00
P228	Using CFHealthHub nebuliser duration data to support continuous improvement by optimising inhaled therapy duration to reduce treatment burden within a digital learning health system  Abstract Presenter: Robert D Sandler, GB	14:00 - 14:00
P229	The introduction of an adherence support clinic in a large UK adult cystic fibrosis centre	14:00 - 14:00

	Abstract Presenter: Jocelyn Choyce, GB	
P230	Time matters: burden of respiratory physiotherapy in adolescents with cystic fibrosis (CF) and their caregivers Abstract Presenter: Chiara Blardone, IT	14:00 - 14:00
P231	Steps Ahead: optimising physical activity in adults with cystic fibrosis - a pilot randomised trial using wearable technology, goal setting and text message feedback  Abstract Presenter: Audrey Tierney, IE	14:00 - 14:00
P232	Feasibility and added value of the computer game "Sparky©" when learning spirometry in preschool children with cystic fibrosis  Abstract Presenter: Myriam Vreys, BE	14:00 - 14:00
P233	Adherence of nebuliser use in children with cystic fibrosis Abstract Presenter: Kieren James Lock, GB	14:00 - 14:00
P234	CoachMois: the experience of an adapted physical activity challenge online at the Roscoff Cystic Fibrosis Centre (Brittany, FR)  Abstract Presenter: Aline DUBOIS, FR	14:00 - 14:00
P235	What influence can educational videos have on daily physical therapy in children, adolescents and young adults with cystic fibrosis? A pilot study  Abstract Presenter: Kevin Cobb, AT	14:00 - 14:00
P236	Commit to Fit - access to gym space for young people with cystic fibrosis in partnership with Helping Hand charity Abstract Presenter: Niju Baby, GB	14:00 - 14:00
P237	Exercise immunology and cystic fibrosis  Abstract Presenter: GAUCHEZ Hugues, FR	14:00 - 14:00
P238	Family participation in cystic fibrosis care  Abstract Presenter: Janne Houben, BE	14:00 - 14:00
P239	The experience of virtual visits and home spirometry in caregivers of children with cystic fibrosis  Abstract Presenter: Viktoria Mellqvist, SE	14:00 - 14:00
P240	Elexacaftor/tezacaftor/ivacaftor treatment improves cystic fibrosis quality of life over multiple domains  Abstract Presenter: Nicola Jane Robinson, GB	14:00 - 14:00
P241	One month with elexacaftor/tezacaftor/ivacaftor in an Italian sample: quality of life and mental health Abstract Presenter: Sonia Graziano, IT	14:00 - 14:00
P242	Young people's views on Kaftrio: a peer research project Abstract Presenter: Eleanor Lee Mindel, GB	14:00 - 14:00
P243	Psychosocial impact of six months of treatment with Symkevi among adult cystic fibrosis patients Abstract Presenter: Sue Braun, BE	14:00 - 14:00
P244	The experiences of taking Kaftrio for cystic fibrosis  Abstract Presenter: Cicely Mathews, GB	14:00 - 14:00

P245	Kaftrio Impact and Side-effect Survey (KISS)	14:00 - 14:00
	Abstract Presenter: Angela Holden, GB	
P246	Improving assessment of palliative care needs among cystic fibrosis children: A Delphi study of the ADAPT-Cystic Fibrosis communication guide  Abstract Presenter: Anna M. Georgiopoulos, US	14:00 - 14:00
P247		14:00 - 14:00
F24/	Dutch translation of the gastrointestinal (GI) symptom tracker for people with cystic fibrosis  Abstract Presenter: Marieke Verkleij, NL	14:00 - 14:00
P248	Parental reports of the oral health-related quality of life and associated factors in children with cystic fibrosis (CF)  Abstract Presenter: Hande Ilgin Sisman, TR	14:00 - 14:00
P249	Awareness of the potential benefits of occupational therapy in a cystic fibrosis multi-disciplinary team Abstract Presenter: Wendy Foo, GB	14:00 - 14:00
P250	Development of a new information resource on fasting during Ramadan and cystic fibrosis  Abstract Presenter: J Carter, GB	14:00 - 14:00
P251	The effect of breathing exercise on the quality of life of 3-12 year-old patients with cystic fibrosis and their families  Abstract Presenter: mine kalyoncu, TR	14:00 - 14:00
P252	A service improvement project: identifying and addressing problems experienced by adults with cystic fibrosis relating to the supply of medicines in primary care	14:00 - 14:00
	Abstract Presenter: Monika Chawla, GB	
P253	Using quality improvement to develop transition to adult care at a UK cystic fibrosis centre  Abstract Presenter: Louise Warnock, GB	14:00 - 14:00
P254	Transition to adult care in children with cystic fibrosis - experience over a decade from a large tertiary centre in the northwest United Kingdom	14:00 - 14:00
D055	Abstract Presenter: Anirban Maitra, GB	14.00 14.00
P255	Transitioning to adult services for children with cystic fibrosis: what matters to all stakeholders?	14:00 - 14:00
	Abstract Presenter: Verena Michael, GB	
P256	Change in the anxiety levels of children with cystic fibrosis and their mothers at the beginning of the COVID-19 pandemic and after one year	14:00 - 14:00
	Abstract Presenter: Tugba Sismanlar Eyuboglu, TR	
P257	Who's talking about cystic fibrosis continued: the effects of the COVID-19 pandemic on the cystic fibrosis online landscape	14:00 - 14:00
	Abstract Presenter: Hisham Ibrahim, IE	

ePoster Session 14:00 - 15:00

**ePoster Sessions** 

14:56 - 14:56

14:00 - 14:06

ePoster Session 14:00 - 15:00

# ePoster Session 6 - Modulators and other approaches: from the lab to the patient

ePoster Corner C

Leader: Isabelle Sermet-Gaudelus, FR Leader: Guido Veit, CA				
EPS6.01	Characterisation of F508del-CFTR rescue by corrector PTI-801	14:00 - 14:07		
	Abstract Presenter: Miquéias Lopes-Pacheco, PT			
EPS6.02	CFTR-modulators improve bicarbonate transport via the CFTR variant D1152H (c.3453G>C) in primary human intestinal epithelial cells	14:07 - 14:14		
	Abstract Presenter: Dora Angyal, NL			
EPS6.03	Enabling RNA-based gene modification in high-throughput screenings using cystic fibrosis patient-derived organoids  Abstract Presenter: Cinthya del Angel Zuvirie, NL	14:14 - 14:21		
EPS6.04	Specialised pro-resolving mediators' biosynthesis by cystic fibrosis airway epithelial cells and their impact on mucociliary clearance	14:21 - 14:28		
	Abstract Presenter: Maëlle Briottet, FR			
EPS6.05	ELX/TEZ/IVA vs. TEZ/IVA in intestinal organoids: analysis of 63 individuals with cystic fibrosis	14:28 - 14:35		
	Abstract Presenter: Eva Furstova, CZ			
EPS6.07	SpliSense's ASO SPL84-23-1 properly distributes and is retained in cystic fibrosis-like mice lungs  Abstract Presenter: Gili Hart, IL	14:35 - 14:42		
EPS6.08	Second-hand smoke exposure reduces the clinical efficacy of ivacaftor in G551D: results from the GOAL study  Abstract Presenter: Steven M. Rowe, US	14:42 - 14:49		
EPS6.09	A discrete choice experiment to quantify the influence of trial features on patients' decisions to join cystic fibrosis clinical trials	14:49 - 14:56		
	Abstract Presenter: Rebecca Dobra, GB			

ePoster Session 14:00 - 15:00

EPS6.10

# ePoster Session 4 - When CF encounters Sars-Cov2 Infection and vaccination

A real-world study evaluating the impact of

adherence in cystic fibrosis

Abstract Presenter: Karen Lester, IE

elexacaftor/tezacaftor/ivacaftor treatment on medication

ePoster Corner A

Leader: Daniel Peckham, GB Leader: Carla Colombo, IT

EPS4.01 The impact of the SARS-CoV-2 pandemic on people with

cystic fibrosis in Ireland: did individuals maintain their key

health outcomes?

Abstract Presenter: Laura Kirwan, IE

EPS4.02	Impact of COVID-19 on the disease course in cystic fibrosis? A Registry-based study	14:06 - 14:12
	Abstract Presenter: Josefien Vandekerckhove, BE	
EPS4.03	Outcomes of COVID-19 in patients with cystic fibrosis in Wales, UK	14:12 - 14:18
	Abstract Presenter: Dawn Lau, GB	
EPS4.04	Evaluation of burden of the COVID-19 pandemic among the cystic fibrosis community in Brno, Czech Republic Abstract Presenter: Miriam Mala, CZ	14:18 - 14:24
EPS4.05	Evolving cystic fibrosis care during COVID-19: a single- centre experience	14:24 - 14:30
	Abstract Presenter: Giulia Spoletini, GB	
EPS4.06	Antibody response to administration of two doses of the BNT162b2 vaccine against SARS-CoV-2 in people with cystic fibrosis	14:30 - 14:36
	Abstract Presenter: Gianfranco Alicandro, IT	
EPS4.07	Covid-19 early vaccination campaign in cystic fibrosis (CF) Italian patients: the experience of the Italian Society for the Study of Cystic Fibrosis (SIFC) Abstract Presenter: Francesca Lucca, IT	14:36 - 14:42
EPS4.08	COVID-19 vaccine uptake in adult people with cystic fibrosis	14:42 - 14:48
21.00	Abstract Presenter: Muhammad Haris Mir, GB	11.12 11.10
EPS4.09	COVID-19 vaccination in households of paediatric cystic fibrosis patients - experience of The Cystic Fibrosis Centre in Zagreb, Croatia	14:48 - 14:54
	Abstract Presenter: Ivan Bambir, HR	
EPS4.10	Evaluation of possible adverse events in relationship to the SARS-CoV-2 vaccination in cystic fibrosis patients with or without lung transplantation: a retrospective single-centre case series	14:54 - 15:00
	Abstract Presenter: Fiorenza Gautschi, CH	
ePoster Session 14:00 - 15:00		
ePoster Session	n 5 - Complementary aspects on CF pathology and treatment	
ePoster Corner	В	
Leader: Carsten Leader: Lieven I		
EPS5.01	Cystic fibrosis (CF) patients chronically infected with Pseudomonas aeruginosa have widespread alterations in both innate and adaptive immunity Abstract Presenter: Mads Lausen, DK	14:00 - 14:06
EPS5.02	Effect of omalizumab on glucocorticoid use and lung function in the treatment of allergic bronchopulmonary aspergillosis in people with cystic fibrosis: a systematic review and meta-analysis	14:06 - 14:12
	Abstract Presenter: J. Stuart Elborn, GB	
EPS5.03	Down-regulation of Aspergillus fumigatus-induced IFN $\beta$ and	14:12 - 14:18

	IFN□□1 expression in cystic fibrosis bronchial epithelial cells is partially rescued by CFTR modulator treatment  Abstract Presenter: Sarah Laverty, GB	
EPS5.04	Chronic exposure to Aspergillus fumigatus leads to cell death and increased eosinophilia in the airways of cystic fibrosis mice  Abstract Presenter: Thomas J Williams, GB	14:18 - 14:25
EPS5.05	Cystic fibrosis chronic rhinosinusitis: implications of automated versus manual sinus CT analysis and patient-reported symptoms  Abstract Presenter: Jennifer Taylor-Cousar, US	14:25 - 14:32
EPS5.06	Implementing tablet-based ototoxicity screening in adults with cystic fibrosis  Abstract Presenter: Shereen Boreland, GB	14:32 - 14:39
EPS5.07	Alkalosis-induced hypoventilation in cystic fibrosis: the importance of efficient renal adaptation  Abstract Presenter: Peder Berg, DK	14:39 - 14:46
EPS5.08	The effects on systemic inflammation of elexacaftor/tezacaftor/ivacaftor  Abstract Presenter: Nicola Robinson, GB	14:46 - 14:53
EPS5.09	Comparison of the Lung Clearance Index in preschool children with primary ciliary dyskinesia and cystic fibrosis Abstract Presenter: Jobst Röhmel, DE	14:53 - 15:00

ECFS Tomorrow Lounge Session

14:15 - 15:15

# Challenging conversations in healthcare: transforming preparedness through simulation

Chair: Gerry Gormley, GB

<i>Workshop</i> 15:00 - 16:30		R1
WS12 - WS12: I	insight into the pathophysiology of CF pulmonary disease	
Chair: Hettie Jan Chair: Isabelle Fa		
WS12.01	Acute pulmonary exacerbations in early cystic fibrosis lung disease are associated with CD3 and PD-1 modulation on lung T cells	15:00 - 15:15
	Abstract Presenter: Vincent Giacalone, US	
WS12.02	BAFF and other soluble factors in airway samples are linked with pathological cystic fibrosis neutrophil phenotype in early childhood	15:15 - 15:30
	Abstract Presenter: Craig J. Schofield, AU	
WS12.03	PD-1 expression on airway macrophages in early cystic fibrosis lung disease coincides with decreased expression of phagocytosis-related markers  Abstract Presenter: Lisa Slimmen, NL	15:30 - 15:45
	·	
WS12.04	Understanding the mechanism of <i>S. aureus</i> killing by neutrophils in the cystic fibrosis airway environment	15:45 - 16:00

	Abstract Presenter: Kayla Fantone, US	
WS12.05	CFTR-TGF $\beta_1$ interaction in inflammatory processes in healthy and CFTR-mutated human bronchial epithelial cells Abstract Presenter: Jan Christoph Thomassen, DE	16:00 - 16:15
WS12.06	Hypertonic saline triggers inflammatory responses in human macrophages	16:15 - 16:30
	Abstract Presenter: Francesca Sposito, GB	
<i>Workshop</i> 15:00 - 16:30		R2
WS13 - WS13: 0	Characterisation of the airway microbiota in cystic fibrosis	
Chair: Tavs Qvist Chair: Pavel Drev		
WS13.01	An invisible threat? <i>Aspergillus</i> -positive cultures and co- infecting bacteria in airway samples	15:00 - 15:15
	Abstract Presenter: Dominic Hughes, GB	
WS13.02	Studying the proteome of respiratory samples from cystic fibrosis patients by shotgun proteomics: differences between exacerbation and clinical stability phases	15:15 - 15:30
	Abstract Presenter: Juan de Dios Caballero, ES	
WS13.03	Real-world relevancy of qPCR for early detection of Pseudomonas aeruginosa infection in people with cystic fibrosis (pwCF)	15:30 - 15:45
	Abstract Presenter: Geneviève Héry-Arnaud, FR	
WS13.04	In vivo lung inflammation induced by $Achromobacter$ spp. clinical isolates with different virulence characteristics	15:45 - 16:00
	Abstract Presenter: Angela Sandri, IT	
WS13.05	Mapping and tackling diversity in antibiotic resistance of sputum isolates in cystic fibrosis	16:00 - 16:15
	Abstract Presenter: Sara Van den Bossche, BE	
WS13.06	Development of antibiotic resistance reveals diverse evolutionary pathways to face the complex and dynamic environment of a long-term treated patient	16:15 - 16:30
	Abstract Presenter: Claudia A Colque, DK	
<i>Workshop</i> 15:00 - 16:30		R3
	Measuring outcomes: where physiotherapy makes a difference	
Chair: Emma Ray Chair: Wolfgang (		
WS14.01	Outcome measures for airway clearance - better the devil you know? Perspectives from adults with cystic fibrosis (CF) Abstract Presenter: Gemma E Stanford, GB	15:00 - 15:15
WS14.02		15.15 15.20
W514.02	Time to first pulmonary exacerbation (PE) in children and adolescents with cystic fibrosis (CF): insights from spirometry, Lung Clearance Index (LCI) and symptoms-limited exercise test (SLET)	15:15 - 15:30
	Abstract Presenter: Alessandra Mariani, IT	

WS14.03	Cystic Fibrosis-Related Diabetes is not associated with maximal aerobic exercise capacity in cystic fibrosis: a cross-sectional analysis of an international multicentre trial (ACTIVATE-CF)	15:30 - 15:45
	Abstract Presenter: Thomas Radtke, CH	
WS14.04	The heart of the matter: is cardiac output a limiting factor for maximal exercise capacity in people with cystic fibrosis?  Abstract Presenter: Marcella Burghard, NL	15:45 - 16:00
WS14.05	Normal fitness data in cystic fibrosis - a scoping review  Abstract Presenter: Owen William Tomlinson, GB	16:00 - 16:15
WS14.06	A retrospective, longitudinal analysis of pulmonary function and peak oxygen uptake in children and adults with cystic fibrosis	16:15 - 16:30
	Abstract Presenter: Hannah Morgan, GB	
<i>Workshop</i> 15:00 - 16:30		R4
	Nutrition status with and without CFTR modulators	
Chair: Monika M Chair: Sarah Coll		
WS15.01	Association of body mass index with clinical outcomes in patients with cystic fibrosis: a systematic review and meta- analysis of 3,100 patients	15:00 - 15:15
	Abstract Presenter: Rita Nagy, HU	
WS15.02	A joint model for lung function and nutritional status decline with recurrent pulmonary exacerbations, death, and lung transplantation using cystic fibrosis patient Registry data	15:15 - 15:30
	Abstract Presenter: Pedro Miranda Afonso, NL	
WS15.03	A mixed-method systematic review of body image in adults with cystic fibrosis (CF)  Abstract Presenter: Darren Sills, GB	15:30 - 15:45
WS15.04	Unanticipated increase in commonly consumed dietary fat when consumed with a new lysophosphatidylcholine-rich nutritional therapy  Abstract Presenter: Virginia Stallings, US	15:45 - 16:00
WS15.05	Impact of refining the U Na/Creat cut-off according to age on the diagnosis of salt depletion in patients with cystic fibrosis	16:00 - 16:15
	Abstract Presenter: Olivia Bauraind, BE	
WS15.06	Fat-soluble vitamin status in F508del homozygous children with cystic fibrosis following commencement of lumacaftor/ivacaftor  Abstract Presenter: Aislinn Kinsella, GB	16:15 - 16:30

Symposium 15:00 - 16:30

R5

# SS02 - SS02: Diagnosing in the liquid world of CFTR related disorders

# At the end of the session, the participant will be able to:

- appreciate the challenges inherent in the diagnostic process of CFTR related disorders
- · use biomarkers of CFTR dysfunction to recognize CFTR related disorders
- · perform an initial clinical evaluation of a person with a CFTR related disorder
- consider benefits and drawbacks of dedicated follow-up programmes for CFTR related disorders

Chair: Kris De Boeck, BE Chair: Carlo Castellani, IT

The CFTR related disorders project: introductory recommendations and work plan	15:00 - 15:22
Speaker: Carlo Castellani, IT	
Biomarkers of CFTR dysfunction in the diagnosis of CFTR related disorders	15:22 - 15:44
Speaker: Isabelle Sermet-Gaudelus, FR	
<b>CFTR related disorders: the initial clinical assessment</b> <i>Speaker</i> : Elke de Wachter, BE	15:44 - 16:06
The CFTR related disorder clinic: advantages and challenges	16:06 - 16:30

Speaker: Nicholas Simmonds, GB

Workshop 15:00 - 16:30

R6

# WS16 - WS16: Beyond CFTR modulators - gene-based and alternative strategies to correct the basic defect

Chair: Jeffrey Beekman, NL Chair: Marianne S. Carlon, BE WS16.01 Correction of the drug-refractory CFTR mutation L227R by 15:00 - 15:15 prime editing Abstract Presenter: Mattijs Bulcaen, BE Base editing strategy to repair the CFTR 2789+5G>A WS16.02 15:15 - 15:30 splicing mutation Abstract Presenter: Simone Amistadi, IT LUNAR®-CF: an mRNA replacement approach for cystic WS16.03 15:30 - 15:45 fibrosis lung disease Abstract Presenter: David Geller, GB WS16.04 Inhalation of SP-101 mediates hCFTRΔR transgene 15:45 - 16:00

expression in the airways of cystic fibrosis and non-cystic fibrosis ferrets

Abstract Presenter: Vethering Exception, US

Abstract Presenter: Katherine Excoffon, US

WS16.05 Identification of drugs activating CFTR-independent fluid 16:00 - 16:15 secretion in nasal organoids based on a high-content screening assay

Abstract Presenter: Gimano Amatngalim, NL

WS16.06 **Development of a high throughput functional screen allows** 16:15 - 16:30

drug repurposing and reveals novel drug candidates for recusing CFTR function in patient-derived organoids with

nonsense mutations

Abstract Presenter: Sacha Spelier, NL

ECFS Tomorrow Lounge Session

15:15 - 16:15

# What psychosocial outcome measures are being used/ found useful in CF?

Chair: Sejal Patel, GB

Chair: Urszula Borawska-Kowalczyk, PL

WS17 - WS17: Clinical and biological updates on CFTR modulators         Chain: David Sheppard, GB       17:00 - 17:15         WS17.01       Therapeutic drug monitoring of elexacaftor, tezacaftor and ivacaftor in adult patients with cystic fibrosis	<i>Workshop</i> 17:00 - 18:30		R1
Chair: David Sheppard, GB		Clinical and biological updates on CFTR modulators	
Ivacaftor in adult patients with cystic fibrosis   Abstract Presenter: Susanne Naehrig, DE	Chair: Barry Plan	nt, IE	
WS17.02   Long-term efficacy of lumacaftor/ivacaftor (LUM/IVA) in children aged two through five years with cystic fibrosis (CF) homozygous for the F508del-CFTR mutation (F/F): a phase 2, open-label extension study   Abstract Presenter: Mirjam Stahl, DE	WS17.01	ivacaftor in adult patients with cystic fibrosis	17:00 - 17:15
pulmonary outcomes in people with cystic fibrosis (pwCF): a long-term real-world study  Abstract Presenter: Christian Merlo, US  WS17.04 CFTR modulation alters the visco-elastic properties of airway mucus in cystic fibrosis patients  Abstract Presenter: Iris Janssens, BE  WS17.05 Elexacaftor/tezacaftor/ivacaftor treatment in people with cystic fibrosis impacts airway progenitor cell function  Abstract Presenter: Nicola Robinson, GB  WS17.06 Impact of elexacaftor/tezacaftor/ivacaftor on lung-recruited neutrophils in cystic fibrosis  Abstract Presenter: Alexandre Cammarata-Mouchtouris, US  Workshop  17:00 - 18:30 R2  WS18 - WS18: Alternative new therapeutic approaches  Chair: Batsheva Kerem, IL  Chair: Margarida Amaral, PT  WS18.01 Manufacturing and device development of SPL84-23-1, an inhaled antisense oligonucleotide, supporting first-inhuman clinical study in cystic fibrosis patients carrying the 3849 mutation  Abstract Presenter: Gili Hart, IL  WS18.02 A PI3Ky mimetic peptide triggers CFTR gating, bronchodilation and reduced inflammation in obstructive	WS17.02	Long-term efficacy of lumacaftor/ivacaftor (LUM/IVA) in children aged two through five years with cystic fibrosis (CF) homozygous for the F508del-CFTR mutation (F/F): a phase 2, open-label extension study	17:15 - 17:30
workshop 17:00 - 18:30 WS18 - WS18 - Manufacturing and device development of SPL84-23-1, an inhaled antisense oligonucleotide, supporting first-inhuman clinical study in cystic fibrosis patients carrying the 3849 mutation Abstract Presenter: Gili Hart, IL  WS18.02 A PI3Ky mimetic peptide triggers CFTR gating, bronchodilation and reduced inflammation in obstructive  Elexacaftor/tezacaftor/ivacaftor treatment in people with cystic fibrosis alternative newathers. 18:00 - 18:15  R20 18:00 - 18:15 - 18:30 18:15 - 18:15 18:15 - 18:30 18:15 - 18:15 18:15 - 18:30 18:15 - 18:15	WS17.03	pulmonary outcomes in people with cystic fibrosis (pwCF): a long-term real-world study	17:30 - 17:45
cystic fibrosis impacts airway progenitor cell function Abstract Presenter: Nicola Robinson, GB  WS17.06 Impact of elexacaftor/tezacaftor/ivacaftor on lung-recruited neutrophils in cystic fibrosis Abstract Presenter: Alexandre Cammarata-Mouchtouris, US  Workshop 17:00 - 18:30 R2  WS18 - WS18: Alternative new therapeutic approaches Chair: Batsheva Kerem, IL Chair: Margarida Amaral, PT  WS18.01 Manufacturing and device development of SPL84-23-1, an inhaled antisense oligonucleotide, supporting first-inhuman clinical study in cystic fibrosis patients carrying the 3849 mutation Abstract Presenter: Gili Hart, IL  WS18.02 A PI3Ky mimetic peptide triggers CFTR gating, bronchodilation and reduced inflammation in obstructive	WS17.04	airway mucus in cystic fibrosis patients	17:45 - 18:00
WS17.06 Impact of elexacaftor/tezacaftor/ivacaftor on lung-recruited neutrophils in cystic fibrosis  Abstract Presenter: Alexandre Cammarata-Mouchtouris, US  Workshop 17:00 - 18:30 R2  WS18 - WS18: Alternative new therapeutic approaches  Chair: Batsheva Kerem, IL Chair: Margarida Amaral, PT  WS18.01 Manufacturing and device development of SPL84-23-1, an inhaled antisense oligonucleotide, supporting first-inhuman clinical study in cystic fibrosis patients carrying the 3849 mutation  Abstract Presenter: Gili Hart, IL  WS18.02 A PI3Ky mimetic peptide triggers CFTR gating, bronchodilation and reduced inflammation in obstructive	WS17.05	cystic fibrosis impacts airway progenitor cell function	18:00 - 18:15
meutrophils in cystic fibrosis  Abstract Presenter: Alexandre Cammarata-Mouchtouris, US  Workshop 17:00 - 18:30 R2  WS18 - WS18: Alternative new therapeutic approaches  Chair: Batsheva Kerem, IL Chair: Margarida Amaral, PT  WS18.01 Manufacturing and device development of SPL84-23-1, an inhaled antisense oligonucleotide, supporting first-inhuman clinical study in cystic fibrosis patients carrying the 3849 mutation  Abstract Presenter: Gili Hart, IL  WS18.02 A PI3Ky mimetic peptide triggers CFTR gating, bronchodilation and reduced inflammation in obstructive			
Workshop 17:00 - 18:30  WS18 - WS18: Alternative new therapeutic approaches  Chair: Batsheva Kerem, IL Chair: Margarida Amaral, PT  WS18.01  Manufacturing and device development of SPL84-23-1, an inhaled antisense oligonucleotide, supporting first-inhuman clinical study in cystic fibrosis patients carrying the 3849 mutation  Abstract Presenter: Gili Hart, IL  WS18.02  A PI3Ky mimetic peptide triggers CFTR gating, bronchodilation and reduced inflammation in obstructive	WS17.06		18:15 - 18:30
17:00 - 18:30  WS18 - WS18: Alternative new therapeutic approaches  Chair: Batsheva Kerem, IL Chair: Margarida Amaral, PT  WS18.01  Manufacturing and device development of SPL84-23-1, an inhaled antisense oligonucleotide, supporting first-inhuman clinical study in cystic fibrosis patients carrying the 3849 mutation  Abstract Presenter: Gili Hart, IL  WS18.02  A PI3Ky mimetic peptide triggers CFTR gating, bronchodilation and reduced inflammation in obstructive		Abstract Presenter: Alexandre Cammarata-Mouchtouris, US	
Chair: Batsheva Kerem, IL Chair: Margarida Amaral, PT  WS18.01  Manufacturing and device development of SPL84-23-1, an inhaled antisense oligonucleotide, supporting first-inhuman clinical study in cystic fibrosis patients carrying the 3849 mutation Abstract Presenter: Gili Hart, IL  WS18.02  A PI3Ky mimetic peptide triggers CFTR gating, bronchodilation and reduced inflammation in obstructive	17:00 - 18:30		R2
WS18.01 Manufacturing and device development of SPL84-23-1, an inhaled antisense oligonucleotide, supporting first-inhuman clinical study in cystic fibrosis patients carrying the 3849 mutation  Abstract Presenter: Gili Hart, IL  WS18.02 A PI3Ky mimetic peptide triggers CFTR gating, bronchodilation and reduced inflammation in obstructive			
inhaled antisense oligonucleotide, supporting first-in- human clinical study in cystic fibrosis patients carrying the 3849 mutation Abstract Presenter: Gili Hart, IL  WS18.02 A PI3Ky mimetic peptide triggers CFTR gating, bronchodilation and reduced inflammation in obstructive			
WS18.02 A PI3Ky mimetic peptide triggers CFTR gating, 17:15 - 17:30 bronchodilation and reduced inflammation in obstructive	WS18.01	inhaled antisense oligonucleotide, supporting first-in- human clinical study in cystic fibrosis patients carrying the	17:00 - 17:15
bronchodilation and reduced inflammation in obstructive		Abstract Presenter: Gili Hart, IL	
	WS18.02	bronchodilation and reduced inflammation in obstructive	17:15 - 17:30

Abstract Presenter: Alessandra Murabito, IT

WS18.03	ETD001: a long-acting inhaled ENaC blocker is well tolerated in humans	17:30 - 17:45
	Abstract Presenter: Paul Russell, GB	
WS18.04	Harnessing CRISPR-Cas9 technology to revert F508del- CFTR defect	17:45 - 18:00
	Abstract Presenter: Anna Cereseto, IT	
WS18.05	Linking the compound database CandActCFTR and CFTR lifecycle map to predict possible active compound combinations  Abstract Presenter: Liza Vinhoven, DE	18:00 - 18:15
WS18.06	Fully automated analysis of airway-artery dimensions on chest-computed tomography in preschool children with cystic fibrosis to evaluate the effect of inhaled hypertonic saline	18:15 - 18:30
	Abstract Presenter: Yuxin Chen, NL	
<i>Workshop</i> 17:00 - 18:30		R3
	Monitoring lung function and structure - where are we?	
Chair: Alexande Chair: Rikke Mu	r Horsley, GB ılvad Sandvik, DK	
WS19.01	Prospective measurement of lung function in 0-4 year-old Danish children with cystic fibrosis	17:00 - 17:15
	Abstract Presenter: Rikke Mulvad Sandvik, DK	
WS19.02	Impact of corrected multiple breath nitrogen washout (MBW) software on assessment of under/unventilated lung units (UVLU) with the MBWShX	17:15 - 17:30
	Abstract Presenter: Christopher Short, GB	
WS19.03	Respiratory impedance analysis at 5Hz may reveal ventilation inhomogeneity in patients with cystic fibrosis Abstract Presenter: Elpis Hatziagorou, GR	17:30 - 17:45
WS19.04	Lung ultrasound in cystic fibrosis bronchiectasis	17:45 - 18:00
	Abstract Presenter: Ioana Mihaela Ciuca, RO	
WS19.05	Arterialised partial pressure of oxygen: an alternative to $FEV_1$ % for tracking cystic fibrosis lung disease in childhood?	18:00 - 18:15
	Abstract Presenter: René Gaupmann, AT	
WS19.06	Validation of airway-artery algorithm to detect and monitor airway disease on chest computed tomography in the ataluren cystic fibrosis cohort  Abstract Presenter: Qianting Lv, NL	18:15 - 18:30
<i>Workshop</i> 17:00 - 18:30		R4
	Complex Psychosocial/Nursing case studies	
Chair: Nichola		
	Borawska-Kowalczyk, PL	17.00 17.15
WS20.01	Empowering self-management using intensive support in a	17:00 - 17:15

	patient with challenging Cystic Fibrosis-Related Diabetes and renal failure	
	Abstract Presenter: Joanna Snowball, GB	
WS20.02	Input supporting a young person with increased cough suppression in the context of COVID-19  Abstract Presenter: Amy Shayle, GB	17:15 - 17:30
WS20.03	What to hope for when there is no hope Abstract Presenter: Edwina Landau, IL	17:30 - 17:45
WS20.04	Enhancing self-compassion and limiting self-criticism in adults with cystic fibrosis decreases the likelihood of hospitalisation  Abstract Presenter: Michail Mantzios, GB	17:45 - 18:00
WS20.05	Cystic fibrosis, lockdown and CFTR modulators - a perfect storm	18:00 - 18:15
	Abstract Presenter: Maggie Hufton, GB	
WS20.06	Co-infection with <i>Nocardia farcinia</i> and SARS-CoV-2 causing the death of an adolescent with cystic fibrosis <i>Abstract Presenter</i> : James Chapman, GB	18:15 - 18:30
Workshop 17:00 - 18:30 <b>WS21 - WS21:</b> I Chair: Francois V	Potpourri of genetics and biomarkers Vermeulen. BE	R5
WS21.02	Cystic fibrosis modifier genes and bacterial infections in Spanish patients	17:15 - 17:30
	Abstract Presenter: Eva Granizo-Rodriguez, ES	
WS21.04	Molecular characterisation of <i>CFTR</i> gene in the Argentinian population: impact of local variants in the sensitivity of genetic testing	17:45 - 18:00
	Abstract Presenter: Carolina Crespo, AR	
WS21.05	Prospective comparison of two sweat test methods Abstract Presenter: Julie Carbonez, BE	18:00 - 18:15
WS21.06	Theß-adrenergic sweat secretion test using the AutoBuSTeD software is a novel, high-sensitive CFTR biomarker for patients with inconclusive CFTR genotype and sweat chloride concentration	18:15 - 18:30
	Abstract Presenter: Sophia Pallenberg, DE	
Workshop 17:00 - 18:30		R6
	Late Breaking Science	
Chair: Jane Davie Chair: Kors Van		
WS22.01	ALPINE2: efficacy and safety of 14-day vs 28-day aztreonam for inhalation solution for <i>Pseudomonas aeruginosa</i> eradication in children and adolescents with cystic fibrosis <i>Abstract Presenter</i> : Francis Gilchrist, GB	17:00 - 17:18

WS22.02	Colonization dynamics of <i>Pseudomonas aeruginosa</i> clinical isolates in a CF-derived airway model	17:18 - 17:36
	Abstract Presenter: Filipa Simões, DK	
WS22.03	The effect of dual-CFTR modulators on airway bacteriology in Copenhagen CF patients	17:36 - 17:54
	Abstract Presenter: Maria Pals Bendixen, DK	
WS22.04	Effect of Elexacaftor/Tezacaftor/Ivacaftor (ELX/TEZ/IVA) on Annual Rate of Lung Function Decline in People with Cystic Fibrosis	17:54 - 18:12
	Abstract Presenter: Tim Lee, GB	
WS22.05	Real-world safety and efficacy of elexacaftor/tezacaftor/ivacaftor (ELX/TEZ/IVA) in people with cystic fibrosis: interim results of a long-term registry-based study	18:12 - 18:30
	Abstract Presenter: Julie K. Bower, US	

## Saturday, 11. June 2022

Symposium 09:00 - 10:30

R1

# S25 - Symposium 25 - What's new with fungi in cystic fibrosis Objectives:

## At the end of the session, the participant will be able to:

- · Synthetize knowledge on the mycobiome in people with cystic fibrosis
- Appraise the changing epidemiology in CF fungal disease induced by CFTR modulators
- · Identify the characteristics of Exophiala infection and its treatment
- · Formulate an opinion on combination of antifungal therapy
- · Understand the relevance of fungal pathogens in people with CF.

Chair: Jean Philippe Bouchara, FR

Chair: Miriam Moffatt, GB

The fungal airway microbiome in cystic fibrosis  Speaker: Françoise Botterel, FR	09:00 - 09:22
What about CF Fungal Disease in the age of CFTR Modulators  Speaker: Amelia Bercusson, GB	09:22 - 09:44
Exophiala spp. in cystic fibrosis Speaker: Jean Philippe Bouchara, FR	09:44 - 10:06
Combination of antifungal therapy in cystic fibrosis  Speaker: Carsten Schwarz, DE	10:06 - 10:30

*Symposium* 09:00 - 10:30

R2

# S26 - Symposium 26 - Women's health

#### At the end of the session, the participant will be able to:

- Identify the difference in outcomes according to gender in cystic fibrosis
- $\cdot$   $\,$  Discuss the current evidence regarding fertility and contraception advice for people with cystic fibrosis
- · Apply best practice advice to the management of pregnancy in cystic fibrosis
- Appraise the current evidence regarding CFTR modulator use in pregnancy

Speaker: Jennifer Taylor-Cousar, US

Chair: Kris De Boeck, BE Chair: Peter Middleton, AU

The issue of the gender gap in CF care  Speaker: Isabelle Fajac, FR	09:00 - 09:22
Fertility and contraception  Speaker: Peter Middleton, AU	09:22 - 09:44
Management of pregnancy in cystic fibrosis  Speaker: Michal Shteinberg, IL	09:44 - 10:06
CFTR modulator therapy and pregnancy - Knowns and unknowns	10:06 - 10:30

Symposium 09:00 - 10:30

R3

10:12 - 10:30

## S27 - Symposium 27 - Master Class in Clinical CF - Case Studies

# At the end of the session, the participant will be able to:

- 1. Describe best practice in challenging clinical cases
- 2. Discuss alternative approaches to management in challenging clinical cases
- 3. Evaluate different management strategies in the clinical care of people with cystic fibrosis
- 4. Synthesize the best management strategy for people with CF who have complex medical challenges

Chair: Peter Barry, GB

Biofilm infection of a central venous port-catheter caused	09:00 - 09:18
by Mycobacterium avium complex in an immunocompetent	
child with cystic fibrosis	
Speaker: Alexandra Kavvalou, DE	
D: (I D If I CD	

Discussant: Ian Balfour-Lynn, GB

# Don't fall asleep, this is the case you are weighting for! 09:18 - 09:36 Speaker: Daniel Tewkesbury, GB

Discussant: Barry Plant, IE

# Intronic Variant Analysis; A Route to Diagnosis 09:36 - 09:54

Speaker: Abbie Stephens, GB Discussant: Silvia Gartner, ES

# Successful individualized dose reduction of elexacaftor/ 09:54 - 10:12 tezacaftor/ ivacaftor in cystic fibrosis patients in response to self-reported anxiety

Speaker: Hisham Ibrahim, IE Discussant: Lieven Dupont, BE

# Inflammation and leucocytoclastic vasculitis in a young cystic fibrosis patient: cause or consequence of severe lung disease?

Speaker: Mariska Delausnay, BE Discussant: Peter Barry, GB

Symposium 09:00 - 10:30 R4

# S28 - Symposium 28 - Liver disease in cystic fibrosis

Chair: Jaroslaw Walkowiak, PL Chair: Carla Colombo, IT

Pathogenesis and clinical manifestations of liver disease in	09:00 - 09:22
paediatrics	

Speaker: Carla Colombo, IT

# Is liver disease in adulthood different? 09:22 - 09:44

Speaker: Philippe Sogni, FR

# Radiological approaches to diagnose and monitor liver 09:44 - 10:06 disease - dilemmas...

Speaker: Dominique Debray, FR

#### **CFTR modulators: a promotor or a cure of CF liver disease?** 10:06 - 10:30

Speaker: Frank Bodewes, NL

*Symposium* 09:00 - 10:30

R5

# S29 - Symposium 29 - Epidemiology of screening and CF genotype

## At the end of the session, the participant will be able to:

- Discuss differences in newborn screening practices across Europe.
- Summarise global patterns in CFTR mutations.
- Describe the role of different CFTR mutations, including rate mutations, on CF diagnosis and disease progression
- Recognise the role of registries in assessing effectiveness on CFTR modulators, including by genotype.

Chair: Andreas Jung, CH Chair: Kevin Southern, GB

Improving the quality and performance of newborn screening for CF across Europe	09:00 - 09:22
Speaker: Jürg Barben, CH	
Impact of nonsense mutations on the evolution of lung disease	09:22 - 09:44
Speaker: Isabelle Sermet-Gaudelus, FR	
Worldwide analysis of CFTR mutations and application to screening	09:44 - 10:06
Speaker: Milan Jr Macek, CZ	
The role of CF Registry in assessing effectiveness of targeted (CFTR modulator) therapy, and impacts by CF	10:06 - 10:28

genotype

Speaker: Elizabeth Cromwell, US

Symposium 09:00 - 10:30

R6

#### S30 - Symposium 30 - Targets for CFTR bypass therapy

## At the end of the session, the participant will be able to:

- Evaluate whether alternative pathways for transmembrane anion transport can be targeted to rehydrate CF airway epithelia
- Summarise progress towards developing small molecule ENaC inhibitors to prevent dehydration of airway epithelia in CF
- Analyse the potential of targeting ATP12A to restore host defence mechanisms to CF airway

Chair: David Sheppard, GB Chair: Marcel Bijvelds, NL

voido, ita	
Modulation of TMEM16A to substitute for CFTR dysfunction in CF epithelia	09:00 - 09:22
Speaker: Raimund Dutzler, CH	
SLC26A9 and SLC6A14 as gene modifiers of disease severity and drug response in cystic fibrosis Speaker: Harriet Corvol, FR	09:22 - 09:44
Innovative ENaC inhibitors to suppress dehydration of CF	09:44 - 10:06
epithelia	03.44 - 10.00
Speaker: Henry Danahay, GB	
ATP12A as a therapeutic target in CF lung disease	10:06 - 10:30
Speaker: Gilles Crambert, FR	

Closing Plenary		
11:00 - 12:30		R1
<b>Closing Plenar</b>	y	
Chair: Harm Tid	dens, NL	
Chair: Isabelle F	Fajac, FR	
	ECFS standards for the provision of CFTR modulators	11:00 - 11:30
	Speaker: Kevin Southern, GB	
	Cystic fibrosis research and treatment, what is next?	11:30 - 12:00
	Speaker: Jeffrey Beekman, NL	
	ECFS President Address	12:00 - 12:30
	Speaker: Isabelle Fajac, FR	
Closing Ceremo	ny	
12:30 - 13:00		R1
<b>Closing Cerem</b>	ony	