

Wednesday, 07. June 2023

Opening Plenary

18:30 - 20:00

R6

Opening Plenary

The future of CF care

18:30 - 18:50

Speaker: Nicholas Simmonds, GB

Thursday, 08. June 2023*Meet the Experts*

07:30 - 08:20

MTE03 - Meet the Experts 03 - Management of haemoptysis in cystic fibrosis*Chair:* Andrew Jones, GB*Chair:* Malena Cohen-Cymberknoh, IL*Meet the Experts*

07:30 - 08:20

Meet the Experts*Meet the Experts*

07:30 - 08:20

MTE02 - Meet the Experts 02 - Diagnosis and management of CFTR-RD*Chair:* Carlo Castellani, IT*Chair:* Nicholas Simmonds, GB*Meet the Experts*

07:30 - 08:20

MTE01 - Meet the Experts 01 - What's new in CF-related biofilm research?*Chair:* Tom Coenye, BE*Chair:* Susanne Haussler, DE*Symposium*

08:30 - 10:00

R1

S01 - Symposium 01 - Closing the gap: treating all patients with gene-protein therapies*Chair:* Kors Van der Ent, NL*Chair:* Margarida Amaral, PT**Gene therapy in cystic fibrosis: the holy grail?**

08:30 - 08:52

Speaker: Eric Alton, GB**Rare mutations and modulators: who's left to treat?**

08:52 - 09:14

Speaker: Jeffrey Beekman, NL**CFTR modulator non-responders: pharmacogenetics and complex alleles**

09:14 - 09:36

Speaker: Nicoletta Pedemonte, IT**Treating nonsense mutations: what's left to do?**

09:36 - 10:00

Speaker: Fabrice Lejeune, FR*Symposium*

08:30 - 10:00

R2

S02 - Symposium 02 - Antibiotic therapy in cystic fibrosis - state of the art*Chair:* Jean-Luc Mainardi, FR*Chair:* Annamaria Bevivino, IT**Interspecies interactions and their effect on antibiotic efficacy**

08:30 - 08:52

Speaker: Michael Bottery, GB**Tolerance of *Pseudomonas aeruginosa* biofilms - why it**

08:52 - 09:14

matters*Speaker:* Susanne Haussler, DE**Antimicrobial susceptibility testing in cystic fibrosis: is it still relevant?**

09:14 - 09:36

Speaker: Valerie Waters, CA**New hits and validated drug targets in the fight against Mycobacterium abscessus and other nontuberculous mycobacteria**

09:36 - 10:00

Speaker: Laurent Kremer, FR*Symposium*

08:30 - 10:00

R3

S03 - Symposium 03 - Harmonising global care in cystic fibrosis*Chair:* Silvia Gartner, ES*Chair:* Ernst Eber, AT**ECFS Twinning Project - goals, progress and challenges**

08:30 - 08:52

Speaker: Pavel Drevinek, CZ**Challenges for cystic fibrosis care in developing countries**

08:52 - 09:14

Speaker: Marco Zampoli, ZA**How to overcome challenges with drug access and reimbursement**

09:14 - 09:36

Speaker: Kris De Boeck, BE**Novel programmes for enhancing drug access**

09:36 - 10:00

Speaker: Clémence Martin, FR*Symposium*

08:30 - 10:00

R4

S04 - Symposium 04 - The times are changing and we are changing with them*Chair:* Trudy Havermans, BE*Chair:* Pavla Hodkova, CZ**Change in illness perception and identity - what do we know today?**

08:30 - 08:52

Speaker: Horst Mitmansgruber, AT**Mental health and quality of life - new measures in evaluation, research and screening?**

08:52 - 09:14

Speaker: Sonia Graziano, IT**Patient education - what do patients need to know and what skills should they be taught?**

09:14 - 09:36

Speaker: Helen Chadwick, GB**Collaboration - the science of shared decision making in the light of upheaval and uncertainty**

09:36 - 10:00

Speaker: Johanna Gardecki, DE

Symposium

08:30 - 10:00

R5

S05 - Symposium 05 - Controversies in the world of screening for cystic fibrosis*Chair:* Jürg Barben, CH*Chair:* Kevin Southern, GB**Prenatal screening for cystic fibrosis remains justified in the era of CFTR modulator therapy: PRO** 08:30 - 08:48*Speaker:* Hannah Blau, IL**Prenatal screening for cystic fibrosis remains justified in the era of CFTR modulator therapy: CON** 08:48 - 09:06*Speaker:* John Massie, AU**Discussion** 09:06 - 09:16**How to make newborn screening for cystic fibrosis ethnically fairer** 09:16 - 09:38*Speaker:* Maya Desai, GB**Is it time to revisit the ECFS standards to reduce false positive results in newborn screening?** 09:38 - 10:00*Speaker:* Olaf Sommerburg, DE*Industry Sessions*

10:00 - 10:30

CF Innovation Zone - Exhibitor Presentations*Chair:* Damian Downey, GB*Symposium*

10:30 - 12:00

R1

S06 - Symposium 06 - Pulmonary challenges post access to CFTR modulators*Chair:* Jane Davies, GB*Chair:* Isabelle Durieu, FR**Strategies for infection control and surveillance in non-productive patients** 10:30 - 10:52*Speaker:* Claire Wainwright, AU**Diagnosis and management of pulmonary exacerbations in CFTR modulator responsive patients** 10:52 - 11:14*Speaker:* Barry Plant, IE**How to monitor people with preserved lung function at all ages?** 11:14 - 11:36*Speaker:* Mirjam Stahl, DE**How do we make remote monitoring work in cystic fibrosis?** 11:36 - 12:00*Speaker:* Nicholas Simmonds, GB*Symposium*

10:30 - 12:00

R2

S07 - Symposium 07 - Microbiology in the era of CFTR modulators*Chair:* Deborah Baines, GB*Chair:* Tom Coenye, BE**Effects of CFTR modulators on inflammation in cystic fibrosis** 10:30 - 10:52

Speaker: Michal Shteinberg, IL

Impact of airway inflammation and infection on the efficacy of CFTR modulators 10:52 - 11:14

Speaker: Carla Ribeiro, US

The effect of CFTR modulators on airway bacteriology in cystic fibrosis patients 11:14 - 11:36

Speaker: Helle Krogh Johansen, DK

Impact of CFTR modulators on IV antibiotic use 11:36 - 12:00

Speaker: Ruth Keogh, GB

Symposium

10:30 - 12:00

R3

S08 - Symposium 08 - Gene based-therapeutic approaches: novel delivery systems and beyond

Chair: Anna Cereseto, IT

Chair: Patrick Harrison, IE

Advances in liposome and polymer systems for cystic fibrosis gene therapy 10:30 - 10:52

Speaker: Stephen Hart, GB

Advances in AAV platforms for efficient gene therapy 10:52 - 11:14

Speaker: Hildegard Büning, DE

Inhalable RNA formulations based on lung surfactant and repurposed cationic amphiphilic drugs 11:14 - 11:36

Speaker: Koen Raemdonck, BE

Airway stem cell-based therapies for cystic fibrosis sinus disease 11:36 - 12:00

Speaker: Shafagh Waters, AU

Symposium

10:30 - 12:00

R4

S09 - Symposium 09 - Overweight and healthy lifestyle in cystic fibrosis

Chair: Chris Smith, GB

Chair: Michael Wilschanski, IL

Dietary approach to treating overweight and obesity in people with cystic fibrosis 10:30 - 10:52

Speaker: Daina Kalnins, CA

Metabolic complications in cystic fibrosis with the preventive nutrition strategy 10:52 - 11:14

Speaker: Andrea Gramegna, IT

CFTR modulators and their impact on body composition - dietitian role 11:14 - 11:36

Speaker: Dimitri Declercq, BE

Evaluating and implementing mindful eating practices in cystic fibrosis - Challenges and opportunities 11:36 - 12:00

Speaker: Helen Egan, GB

Symposium

10:30 - 12:00

R5

S10 - Symposium 10 - "Things you wish you knew" - hot topics in physiotherapy*Chair:* Gemma Stanford, GB*Chair:* Marlies Wagner, AT**Musculoskeletal issues in the modern cystic fibrosis era** 10:30 - 10:52*Speaker:* Julia Taylor, GB**Cough, spit, suck? - pros, cons and indications for microbiological sputum sampling techniques** 10:52 - 11:14*Speaker:* Carwyn Bridges, GB**Doing a lot with a little (how to do physiotherapy with limited resources/time/access to medications)** 11:14 - 11:36*Speaker:* Brenda Morrow, ZA**Fatter but fitter? - exercise requirements for the post-modulator cystic fibrosis population** 11:36 - 12:00*Speaker:* Wolfgang Gruber, DE*Industry Sessions*

12:30 - 14:00

R1

Satellite Symposium*ECFS Tomorrow Lounge Session*

12:45 - 13:45

Tomorrow Lounge

Mental Health challenges in the era of new modulators*Speaker:* Anna M. Georgiopoulos, US*Speaker:* Annette Katscher-Peitz, DE*Speaker:* Edwina Landau, IL*Speaker:* Alexandra Quittner, US*ePoster Session*

14:00 - 15:00

R2

ePoster Session 1*ePoster Session*

14:00 - 15:00

R2

EPS01 - ePoster Session 1: Triple I in cystic fibrosis: Imaging, Inflammation and Immunology*Chair:* Andrea Lakatos-Krepcik, AT*Chair:* Tim Lee, GBEPS1.01 **Using lateral decubitus computed tomography (CT) to monitor structural lung disease in young children with cystic fibrosis** 14:00 - 14:00*Oral Presenter:* Rikke Mulvad Sandvik, DKEPS1.02 **Reanalysis of N₂-lung clearance index and the comparison to SF₆-lung clearance index and magnetic resonance imaging** 14:00 - 14:00*Oral Presenter:* Eva Steinke, DEEPS1.03 **Establishing the utility of oxygen-enhanced (OE-)MRI as a structural and functional measure of cystic fibrosis lung disease: relationships with physiological and cystic fibrosis outcomes** 14:00 - 14:00

	<i>Oral Presenter:</i> Christopher Short, GB	
EPS1.04	Change in lung clearance index in adolescents with cystic fibrosis treated with elexacaftor/tezacaftor/ivacaftor: results from 12 months follow-up in the Danish cystic fibrosis cohort <i>Oral Presenter:</i> Esben Herborg Henriksen, DK	14:00 - 14:00
EPS1.05	Bronchoscopic practices in adult cystic fibrosis at Cork University Hospital 2012-2022: increased usage, new indications and the emergence of single-use flexible bronchoscopy <i>Oral Presenter:</i> Kevin Deasy, IE	14:00 - 14:00
EPS1.06	Galectin-3: potential biomarkers in children with cystic fibrosis <i>Oral Presenter:</i> Ismail Guzelkas, TR	14:00 - 14:00
EPS1.07	Impact of elexacaftor/tezacaftor/ivacaftor (ETI) therapy on <i>Aspergillus Fumigatus</i> serology in adults with cystic fibrosis <i>Oral Presenter:</i> Fraser Maxwell Curran, GB	14:00 - 14:00
EPS1.08	Immunogenic adverse events to CFTR modulators - an international survey <i>Oral Presenter:</i> Ruth M. Urbantat, DE	14:00 - 14:00
EPS1.09	Feasibility of cardiac magnetic resonance imaging in older cystic fibrosis patients <i>Oral Presenter:</i> Karuna Sapru, GB	14:00 - 14:00
EPS1.10	Prevalence, risk factors and outcomes of cardiovascular disease in cystic fibrosis: retrospective cohort study in two large patient data registries <i>Oral Presenter:</i> Freddy Frost, GB	14:00 - 14:00

ePoster Session

14:00 - 15:00

ePoster Sessions 2 - 4*ePoster Session*

14:00 - 15:00

EPS03 - ePoster Session 3 - CFTR: what is still puzzling us?*Chair:* Simon Graeber, DE*Chair:* Anabela Santo Ramalho, BE

EPS3.02	The cystic fibrosis urine test: A comprehensive analysis of all renal acid-base parameters following acute oral bicarbonate loading <i>Oral Presenter:</i> Amalie Quist Rousing, DK	14:00 - 14:00
EPS3.04	Rescuing rare CFTR mutants by a mimetic peptide targeting the AKAP function of PI3Kγ <i>Oral Presenter:</i> Angela Della Sala, IT	14:00 - 14:00
EPS3.05	Proximity Profiling of CFTR with Gating Mutations <i>Oral Presenter:</i> Melissa Iazzi, CA	14:00 - 14:00
EPS3.07	Mapping ivacaftor-induced structural changes in CFTR with computer simulations	14:00 - 14:00

Oral Presenter: David Sheppard, GB

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| EPS3.08 | Heightened mitochondrial respiration and succinate levels in cystic fibrosis cells is normalised with triple CFTR modulator therapy through mechanisms involving cell calcium flux | 14:00 - 14:00 |
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Oral Presenter: Heledd Jarosz-Griffiths, GB

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| EPS3.09 | Transcriptomic and Functional Analysis of Chloride, Bicarbonate, and Proton Secretion Along the Crypt-Villus Axis in Human Intestine: lessons for cystic fibrosis | 14:00 - 14:00 |
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Oral Presenter: Zachary Sellers, US

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| EPS3.10 | Novel CFTR modulator combinations directly address the ΔF508-CFTR NBD1 stability defect and enable full CFTR correction | 14:00 - 14:00 |
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Oral Presenter: Gregory Hurlbut, US

ePoster Session

14:00 - 15:00

EPS02 - ePoster Session 2 - Empower your patients: how to improve care right from the start

Chair: Majda Oštir, SI

Chair: Sue Braun, BE

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| EPS2.01 | Hear my voice: research <i>by</i> and <i>with</i> children with cystic fibrosis | 14:00 - 14:00 |
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Oral Presenter: Simona Caldani, FR

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| EPS2.02 | "You're the person who the decisions are going to be placed on": children with cystic fibrosis and their participation in medical care | 14:00 - 14:00 |
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Oral Presenter: Eleanor Lee Mindel, GB

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| EPS2.03 | How can we achieve self-efficacy instead of helplessness caused by frequent medical interventions in young people with cystic fibrosis? | 14:00 - 14:00 |
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Oral Presenter: Karoline Prinz, AT

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| EPS2.04 | Expanding the cystic fibrosis mental health screening guidelines: using the pediatric symptom checklist to identify and treat mental health symptoms in children 4-11 years of age | 14:00 - 14:00 |
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Oral Presenter: Emily Muther, US

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| EPS2.05 | The challenge of living with cystic fibrosis - short form - challenges experienced by parents/caregivers of children with Cystic Fibrosis in Ireland: the Irish Comparative Outcomes study (ICOS) | 14:00 - 14:00 |
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Oral Presenter: Rini Bhatnagar, IE

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| EPS2.06 | What have we been missing all these years? Empowering the patient's voice | 14:00 - 14:00 |
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Oral Presenter: Laura Bundy, GB

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| EPS2.07 | The Brief Resilience Scale (BRS) as a possible routine assessment instrument for patients with cystic fibrosis: predictive power in psychological symptoms and quality of life | 14:00 - 14:00 |
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Oral Presenter: Horst Mitmansgruber, AT

EPS2.08	Clinical effort against smoke exposure in cystic fibrosis (CEASE-CF): feasibility, acceptability, and preliminary efficacy <i>Oral Presenter:</i> Gabriela Oates, US	14:00 - 14:00
EPS2.09	Cervical cancer in cystic fibrosis: from treatment to prevention <i>Oral Presenter:</i> Aleksandra Duffy, GB	14:00 - 14:00
EPS2.10	Initial validation of the Integrated Palliative Care Outcome Scale (IPOS) in adults with cystic fibrosis (CF): data from the <i>Improving Life with CF</i> multicentre pragmatic implementation trial <i>Oral Presenter:</i> Anna M. Georgiopoulos, US	14:00 - 14:00

ePoster Session

14:00 - 15:00

EPS04 - ePoster Session 4 - What's new in CF-related diabetes?*Chair:* Stephanie Van Biervliet, BE*Chair:* Frank Bodewes, NL

EPS4.01	Using a modified glucose challenge test (GCT) as a screening tool for cystic fibrosis diabetes <i>Oral Presenter:</i> Katie Dick, GB	14:00 - 14:00
EPS4.02	Earlier identification of cystic fibrosis diabetes in children with cystic fibrosis - real-world experience on the use of continuous (flash) glucose monitoring <i>Oral Presenter:</i> Maya Chelminska, GB	14:00 - 14:00
EPS4.03	An audit of diagnosing cystic fibrosis-related diabetes using newly developed diagnostic criteria using flash glucose monitoring system (Freestyle Libreâ) at a regional adult cystic fibrosis centre <i>Oral Presenter:</i> Claire Roden, GB	14:00 - 14:00
EPS4.04	A stitch in time saves nine - abnormal glucose tolerance in patients with cystic fibrosis: systematic review and meta-analysis <i>Oral Presenter:</i> Adrienn F. Kéri, HU	14:00 - 14:00
EPS4.05	Exploring the practicalities and effectiveness of a screening tool in adult cystic fibrosis diabetes <i>Oral Presenter:</i> Hannah Burton, GB	14:00 - 14:00
EPS4.06	Alterations in incretin and somatostatin levels by glucose tolerance status in people with cystic fibrosis <i>Oral Presenter:</i> Bibi Uhre Nielsen, DK	14:00 - 14:00
EPS4.09	Role for DPP4 inhibitor therapy in cystic fibrosis-related diabetes mellitus (CFRD)- experience from Manchester Adult Cystic Fibrosis Unit <i>Oral Presenter:</i> Anjali Santhakumar, GB	14:00 - 14:00
EPS4.10	Obstetric and neonatal outcomes in women with cystic fibrosis (wwCF) and diabetes in pregnancy <i>Oral Presenter:</i> Amy Downes, GB	14:00 - 14:00

Poster Viewing

14:00 - 15:00

PS1 - Poster Viewing 1

P001	A childhood, 16 years of cystic fibrosis (CF) Newborn Screening data in East London and beyond <i>Oral Presenter:</i> Jacqui Cowlard, GB	14:00 - 14:00
P002	Results from a clinical performance study of a new neonatal PAP screening ELISA kit for cystic fibrosis-newborn screening <i>Oral Presenter:</i> Ana Marcão, PT	14:00 - 14:00
P003	Clinical variability of the CFTR variant p.Arg117Cys uncovered by newborn screening for cystic fibrosis. Does the genetic background matter? <i>Oral Presenter:</i> Emma Lundman, NO	14:00 - 14:00
P004	Clinical consequences and functional impact of the rare S737F CFTR variant <i>Oral Presenter:</i> Emanuela Pesce, IT	14:00 - 14:00
P005	An audit on adherence to guidelines on screening colonoscopies in an adult cystic fibrosis centre <i>Oral Presenter:</i> Fiona Hickey, IE	14:00 - 14:00
P008	Critical Disease Burdens of Australian Adults with cystic fibrosis: results from an online survey <i>Oral Presenter:</i> Anastasia Ward, AU	14:00 - 14:00
P009	Development of a new information video for children about cystic fibrosis <i>Oral Presenter:</i> Alison Taylor, GB	14:00 - 14:00
P010	Establishing a collaboration between the 4 university children's hospitals in Egypt and the University of Michigan to improve cystic fibrosis care in Egypt <i>Oral Presenter:</i> Samya Nasr, US	14:00 - 14:00
P011	Late diagnosis of cystic fibrosis in adulthood in Republic of North Macedonia <i>Oral Presenter:</i> Tatjana Jakovska Maretti,	14:00 - 14:00
P012	The attitude towards prenatal cystic fibrosis diagnosis in Bulgaria <i>Oral Presenter:</i> Guergana Petrova, BG	14:00 - 14:00
P013	Tuberculosis or cystic fibrosis <i>Oral Presenter:</i> Guergana Petrova, BG	14:00 - 14:00
P014	Safety and toxicity profile of SPL84, an inhaled antisense oligonucleotide cystic fibrosis therapeutic <i>Oral Presenter:</i> Gili Hart, IL	14:00 - 14:00
P015	Nebulizer selection and characterisation process with SPL84, an inhaled antisense oligonucleotide, supporting first in human clinical study in cystic fibrosis patients carrying the 3849 mutation <i>Oral Presenter:</i> Gili Hart, IL	14:00 - 14:00
P019	Complex CFTR allele [HB1] L467F-F508del is responsible	14:00 - 14:00

	for poor clinical response to elexacaftor/tezacaftor/ivacaftor CFTRm therapy in a cystic fibrosis female with the L467F-F508del/621+1 G>T genotype: a case report <i>Oral Presenter:</i> Marcela Kreslová, CZ	
P020	Therapy with elexacaftor/tezacaftor/ivacaftor in a patient with compound heterozygous CFTR mutation and the complex CFTR-allele <i>Phe508del</i>; <i>Leu467Phe</i> <i>Oral Presenter:</i> Stephanie Thee, DE	14:00 - 14:00
P021	Change in sweat chloride concentration following elexacaftor/Tezacaftor/ivacaftor in siblings with cystic fibrosis heterozygous for R334W / N1088D / R75Q mutations <i>Oral Presenter:</i> Hisham A Saumtally, GB	14:00 - 14:00
P024	Compound heterozygotes bearing the CFTRdup22 in trans with another cystic fibrosis-causing variant have a generally milder course of cystic fibrosis: analysis of 6 Czech cystic fibrosis cases <i>Oral Presenter:</i> Andrea Holubová, CZ	14:00 - 14:00
P025	Prevalence and clinical implications of the <i>p.Cys1400Ter</i> pathogenic CFTR mutation in Cyprus <i>Oral Presenter:</i> Pinelopi Anagnostopoulou, CH	14:00 - 14:00
P027	Mutational analysis of CFTR gene in Pakistani cystic fibrosis patients <i>Oral Presenter:</i> Muhammad Usman Ghani, PK	14:00 - 14:00
P028	Cystic fibrosis mutation pattern in 2019 in Albania: getting closer to the personalised therapy <i>Oral Presenter:</i> Irena Kasmi, AL	14:00 - 14:00
P029	Cystic fibrosis precision treatment - an unequal access in Brazil <i>Oral Presenter:</i> Laís Mota, BR	14:00 - 14:00
P031	Perspectives for gene therapy with the use of CFTR modulators in patients with cystic fibrosis <i>Oral Presenter:</i> Luana da Silva Baptista Arpini, BR	14:00 - 14:00
P038	Direct measurement of short-term adherence to elexacaftor/tezacaftor/ivacaftor and changes in sweat chloride levels - results from the RECOVER Study <i>Oral Presenter:</i> Roy Gavin Stone, IE	14:00 - 14:00
P039	Effect of elexacaftor/tezacaftor/ivacaftor (ETI) on sweat chloride levels in children with cystic fibrosis (CF) - a real-world experience <i>Oral Presenter:</i> Yu Ling Tan, GB	14:00 - 14:00
P040	Sweat chloride values in cystic fibrosis patients after one year on elexacaftor/tezacaftor/ivacaftor <i>Oral Presenter:</i> Natalia Cirilli, IT	14:00 - 14:00
P041	Correlation of improvements in sweat chloride and percent predicted FEV₁ across twenty studies examining different corrector therapies for F508del <i>Oral Presenter:</i> Matthew Heneghan, GB	14:00 - 14:00

P042	Nasal nitric oxide increases in patients with cystic fibrosis treated with elexacaftor/tezacaftor/ivacaftor <i>Oral Presenter:</i> Charlotte Olivia Pioch, DE	14:00 - 14:00
P043	Changes in body composition in adult cystic fibrosis patients within six months of initiation elexacaftor/tezacaftor/ivacaftor therapy in Argentina <i>Oral Presenter:</i> Lorena Mabel Tamburri, AR	14:00 - 14:00
P044	Oral and intravenous antibiotic requirements prior to and following elexacaftor/tezacaftor/ivacaftor treatment in children and adults with cystic fibrosis - analysis from the RECOVER Study <i>Oral Presenter:</i> Roy Gavin Stone, IE	14:00 - 14:00
P045	Physical fitness of paediatric cystic fibrosis patients in the era of CFTR modulators <i>Oral Presenter:</i> Laura Antonia Stöger, AT	14:00 - 14:00
P046	Early parameters to predict long-term efficacy of CFTR modulators in patients with cystic fibrosis <i>Oral Presenter:</i> Pascal Heer, CH	14:00 - 14:00
P049	Effects of elexacaftor/tezacaftor/ivacaftor after 6 months in Dutch people with cystic fibrosis with at least 1 F508del mutation <i>Oral Presenter:</i> Inez Bronsveld, NL	14:00 - 14:00
P050	Elexacaftor/tezacaftor/ivacaftor for adult cystic fibrosis patients with preserved lung function: a case series <i>Oral Presenter:</i> Zisis Balmpouzis, CH	14:00 - 14:00
P051	Real-world experience of change in lung clearance index (LCI_{2.5}) following initiation of elexacaftor/tezacaftor/ivacaftor in children with cystic fibrosis aged 6 through 11 <i>Oral Presenter:</i> Heather Dowle, GB	14:00 - 14:00
P052	Efficacy of elexacaftor/tezacaftor/ivacaftor in people with cystic fibrosis with normal/near normal FEV₁ (≥70%) <i>Oral Presenter:</i> Galit Livnat, IL	14:00 - 14:00
P053	Spirometric and anthropometric improvements in response to elexacaftor/tezacaftor/ivacaftor depend on age and lung disease severity <i>Oral Presenter:</i> Katharina Schütz, DE	14:00 - 14:00
P054	Long-term efficacy and safety of elexacaftor/tezacaftor/ivacaftor (ETI) in people with cystic Fibrosis (pwCF) ≥ 50 years of age in a real-world setting <i>Oral Presenter:</i> Matthias Welsner, DE	14:00 - 14:00
P055	No impact of elexacaftor/tezacaftor/ivacaftor (ETI) CFTRm therapy on male infertility due to CBAVD in two adult cystic fibrosis patients: case report <i>Oral Presenter:</i> Nela Stastna, CZ	14:00 - 14:00
P056	Impact of CFTR Modulator Therapy on sleep parameters in a small cohort of cystic fibrosis patients <i>Oral Presenter:</i> Joana Guimarães, PT	14:00 - 14:00

P057	Sustained Effectiveness of elexacaftor/tezacaftor/ivacaftor in lung transplant candidates with cystic fibrosis <i>Oral Presenter:</i> Filia Diamantea, GR	14:00 - 14:00
P058	New algorithm proposal to allow elexacaftor/tezacaftor/ivacaftor use for patients under 18 with liver dysfunction <i>Oral Presenter:</i> Tiphaine BIHOUEE, FR	14:00 - 14:00
P059	Therapeutic drug monitoring of elexacaftor/tezacaftor/ivacaftor over 1 year in adult patients with cystic fibrosis <i>Oral Presenter:</i> Susanne Naehrig, DE	14:00 - 14:00
P060	Measurement of the plasma concentration of elexacaftor/tezacaftor/ivacaftor (ETI) by LC/MS-MS in a patient with cystic fibrosis during pregnancy <i>Oral Presenter:</i> Mauro Leucio Mattei, IT	14:00 - 14:00
P061	FTR-Modulator therapy during pregnancy in a mother with F508del carrier status and high-grade suspicion of cystic fibrosis in the foetus <i>Oral Presenter:</i> Stephanie Thee, DE	14:00 - 14:00
P062	The impact of elexacaftor/tezacaftor/ivacaftor on adherence to inhaled medication in adults with cystic fibrosis: a 3-centre study in Greece <i>Oral Presenter:</i> Katerina Manika, GR	14:00 - 14:00
P063	Dermatological reactions to elexacaftor/tezacaftor/ivacaftor: experience within a regional adult cystic fibrosis centre <i>Oral Presenter:</i> Akhil P Sawant, GB	14:00 - 14:00
P064	Bilateral cataracts in an adolescent following the use of cystic fibrosis transmembrane conductance regulator (CFTR) modulator therapies <i>Oral Presenter:</i> Pauline Singleton, GB	14:00 - 14:00
P065	A Brazilian experience of elexacaftor/tezacaftor/ivacaftor in people with cystic fibrosis carrying at least one copy of F508del-CFTR <i>Oral Presenter:</i> Miquéias Lopes-Pacheco, PT	14:00 - 14:00
P066	Single center experience of patients using modulatory therapy <i>Oral Presenter:</i> GÖKÇEN ÜNAL, TR	14:00 - 14:00
P067	Real-life data of lumacaftor/ivacaftor therapy in children with cystic fibrosis homozygous for F508del with emphasis on small airway disease in the Netherlands <i>Oral Presenter:</i> Annelies M. Zwitterloot, NL	14:00 - 14:00
P068	Real-life experience with a generic formulation of lumacaftor-ivacaftor in patients with cystic fibrosis homozygous for the Phe508del CFTR mutation <i>Oral Presenter:</i> Alejandro Teper, AR	14:00 - 14:00
P069	Effect of Orkambi therapy on the lung microbiota in people with cystic fibrosis (PwCF) over the first 12 months of therapy (ROCK Study)	14:00 - 14:00

	<i>Oral Presenter:</i> Gisli Einarsson, GB	
P070	Lumacaftor/ivacaftor combination for cystic fibrosis patients in Bulgaria <i>Oral Presenter:</i> Guergana Petrova, BG	14:00 - 14:00
P071	Evaluation of CFTR modulator efficacy by rectal organoid morphology analysis (ROMA) indexes <i>Oral Presenter:</i> Senne Cuyx, BE	14:00 - 14:00
P073	Clinical efficacy of CFTR modulator therapy in patients carrying the I1234V mutation <i>Oral Presenter:</i> Bat El Bar Aluma, IL	14:00 - 14:00
P074	Treatment effects of CFTR modulators on people with cystic fibrosis carrying the Georgian mutation (Q359K/T360K) <i>Oral Presenter:</i> Karin Yaacoby-Bianu,	14:00 - 14:00
P075	Discrepancy between <i>in vitro</i> CFTR functional analysis and clinical response to elexacaftor/tezacaftor/ivacaftor (ETI) in a homozygous S364P (c.1090T>C) patient with cystic fibrosis <i>Oral Presenter:</i> Arthur de SEVIN, FR	14:00 - 14:00
P076	Complex CFTR allele L467F-F508del: <i>in vitro</i> and clinical response to CFTR modulators <i>Oral Presenter:</i> Eva Furstova, CZ	14:00 - 14:00
P077	Clinical and functional efficacy of elexacaftor/tezacaftor/ivacaftor in people with cystic fibrosis carrying the N1303K mutation <i>Oral Presenter:</i> Malena Cohen-Cymerknoh, IL	14:00 - 14:00
P078	Case report: elexacaftor/tezacaftor/ivacaftor as a game changer in an individual with CFTR class II mutation N1303k <i>Oral Presenter:</i> Livia Mia Gona-Hoepler, AT	14:00 - 14:00
P079	Personalized CFTR modulator therapy for G85E and N1303K homozygous patients with cystic fibrosis <i>Oral Presenter:</i> Simon Graeber, DE	14:00 - 14:00
P080	Improved clinical outcomes following ivacaftor treatment in a cystic fibrosis patient homozygous for 3272-26A>G variant <i>Oral Presenter:</i> Jasna Rodman Berlot, SI	14:00 - 14:00
P081	Therotyping - extending the success of highly effective CFTR modulators to rare mutations <i>Oral Presenter:</i> Mordechai Pollak, IL	14:00 - 14:00
P082	Is the new modulator affecting global health outcomes over time? <i>Oral Presenter:</i> Sonia Graziano, IT	14:00 - 14:00
P083	Inequal access to CFTR modulators across ECFS-CTN countries <i>Oral Presenter:</i> Fiona Dunlevy, DK	14:00 - 14:00
P084	Willingness of people treated with elexacaftor/tezacaftor/ivacaftor (ETI) to participate in	14:00 - 14:00

	randomized clinical trials of new modulators and inhaled antibiotics	
	<i>Oral Presenter:</i> Donald VanDevanter, US	
P085	Attitudes to clinical trial participation in the post Kaftrio® era: reasons to be cheerful...	14:00 - 14:00
	<i>Oral Presenter:</i> Alison Lynne Hopkins, GB	
P086	The French clinical research network in cystic fibrosis: more than 10 years of positive experience in clinical research for patients	14:00 - 14:00
	<i>Oral Presenter:</i> Alexandre Coudrat, FR	
P087	Levelling the playing field through the London Network of the UK Clinical Trials Accelerator Platform	14:00 - 14:00
	<i>Oral Presenter:</i> Sophie Pinnell, GB	
P088	A systematic review on all efficacy endpoints used in cystic fibrosis clinical trials in the past 5 years	14:00 - 14:00
	<i>Oral Presenter:</i> Marlou Bierlaagh, NL	
P089	Research priorities in cystic fibrosis: refreshing the James Lind Alliance top ten	14:00 - 14:00
	<i>Oral Presenter:</i> Alan Smyth, GB	
P090	Treatment use among SIMPLIFY trial participants through 24 weeks of follow-up	14:00 - 14:00
	<i>Oral Presenter:</i> Alex H. Gifford, US	
P091	Development and validation of revised treatment adherence and adherence barrier questionnaires for children with cystic fibrosis aged 6 through 11	14:00 - 14:00
	<i>Oral Presenter:</i> Sharon Sutton, IE	
P092	Preliminary observations of treatment and symptom reporting in the Home-Reported Outcomes in cystic fibrosis study (HERO-2)	14:00 - 14:00
	<i>Oral Presenter:</i> Cynthia Brown, US	
P093	The effect quality of life on cystic fibrosis children patients in Gaza Strip	14:00 - 14:00
	<i>Oral Presenter:</i> Asem Altorok, PS	
P094	A Phase II study to evaluate the safety, tolerability, pharmacodynamics and pharmacokinetics of BI 1291583 in patients with cystic fibrosis bronchiectasis (the Clairafly™ study)	14:00 - 14:00
	<i>Oral Presenter:</i> Marcus Mall, DE	
P095	A phase 1b/2a randomized, double-blind, placebo-controlled, multi-centre study evaluating nebulized phage therapy in cystic fibrosis subjects with chronic <i>Pseudomonas aeruginosa</i> pulmonary infection	14:00 - 14:00
	<i>Oral Presenter:</i> Urania Rappo, US	
P096	Effect of acute systemic corticosteroids on clinical outcomes in cystic fibrosis pulmonary exacerbations	14:00 - 14:00
	<i>Oral Presenter:</i> Oliver James McElvaney, US	
P097	Assessment of intrapulmonary percussive ventilation on the	14:00 - 14:00

	rheology of bronchial secretions <i>Oral Presenter:</i> Jérémy Patarin, FR	
P098	Longitudinal follow-up of exacerbated cystic fibrosis patients with sputum rheology <i>Oral Presenter:</i> Jérémy Patarin, FR	14:00 - 14:00
P099	Effect of a mucolytic agent delivered through sinus pulsating aerosol system (SPAS) device on sinonasal symptoms and patient report outcomes in people with cystic fibrosis (PwCF) <i>Oral Presenter:</i> Mari Nieves Balaguer Cartagena, ES	14:00 - 14:00
P100	Undergraduate research-based education for identifying new treatment options for cystic fibrosis <i>Oral Presenter:</i> Nikki Scheen, NL	14:00 - 14:00
P227	Real-world, single-centre evaluation of the efficacy of postal lower airway microbiology samples <i>Oral Presenter:</i> Edward Sizer, GB	14:00 - 14:00
P228	<i>In vitro</i> evolution of levofloxacin resistance in lineages of clinical <i>Pseudomonas aeruginosa</i> isolates cultured at different stages of infection from people with cystic fibrosis <i>Oral Presenter:</i> Callum Matthew Sloan, GB	14:00 - 14:00
P229	Lung and gut microbiome modifications after prolonged Kaftrio® treatment <i>Oral Presenter:</i> Carlo Castellani, IT	14:00 - 14:00
P230	The microbiome of cystic fibrosis (CF) sputum and its association with incident <i>Stenotrophomonas maltophilia</i> (SM) infections <i>Oral Presenter:</i> Lauren Bowron, CA	14:00 - 14:00
P231	Monitoring of respiratory tract infections of cystic fibrosis transplanted patients by means of a multiplex PCR assay <i>Oral Presenter:</i> Cristina Fevola*, IT	14:00 - 14:00
P232	Analysis of the lung microbiome in cystic fibrosis patients using 16s sequencing <i>Oral Presenter:</i> Manasvi Pinnaka, US	14:00 - 14:00
P233	Wanted: <i>Pseudomonas</i> and <i>Staphylococcus</i> - new diagnostic approaches in cystic fibrosis! Is 16S rRNA gene sequencing an equivalent diagnostic tool to conventional microbiology? <i>Oral Presenter:</i> Rebecca Luise Knoll, DE	14:00 - 14:00
P234	Investigating the role of hypoxia in driving the adaptation of <i>Mycobacterium abscessus</i> infection in cystic fibrosis <i>Oral Presenter:</i> Niamh Duggan, IE	14:00 - 14:00
P235	Pharmacological activation of NRF2 has protective effects during <i>Mycobacterium abscessus</i> infection by promoting host defences and reducing inflammatory damage in the context of cystic fibrosis <i>Oral Presenter:</i> Audrey Bernut, FR	14:00 - 14:00
P236	Siderophore production by the emerging cystic fibrosis pathogens of the genus <i>Achromobacter</i>	14:00 - 14:00

	Oral Presenter: Pauline Sorlin, FR	
P237	<i>Stenotrophomonas maltophilia</i> isolates from cystic fibrosis patients eliminate competitor bacteria Oral Presenter: Cristian Crisan, US	14:00 - 14:00
P238	Host-pathogen interactions in the era of antimicrobial resistance Oral Presenter: Ruggero La Rosa, DK	14:00 - 14:00
P239	Bacterial interactions in <i>Pseudomonas aeruginosa</i> and <i>Achromobacter xylosoxidans</i> co-cultures Oral Presenter: Cecilia Sahl, SE	14:00 - 14:00
P240	Lower levels of bacterial aerobic respiration in sputum from people with cystic fibrosis with chronic lung infection Oral Presenter: Peter Østrup Jensen, DK	14:00 - 14:00
P241	Using an <i>in vitro</i> model to investigate the microbiomes of people with cystic fibrosis Oral Presenter: Katrine Madsen, DK	14:00 - 14:00
P242	Small colony variants of <i>Staphylococcus aureus</i> often exhibit a mucoid phenotype in the airways of people with cystic fibrosis Oral Presenter: Christine Rumpf, DE	14:00 - 14:00
P243	The effect of hypoxic conditions on <i>Burkholderia cenocepacia</i> clinical isolates from individuals with cystic fibrosis Oral Presenter: Ciaran Carey, IE	14:00 - 14:00
P244	The impact of CFTR modulators on positive <i>Pseudomonas aeruginosa</i> (Pa) culture and antibiotic susceptibility in adults with cystic fibrosis Oral Presenter: Natalya Ellis, GB	14:00 - 14:00
P245	Real-world elexacaftor/tezacaftor/ivacaftor(ETI) changes prospective sputum collection and microbiological reporting in a single centre pilot cystic fibrosis cohort. Oral Presenter: Kevin Deasy, IE	14:00 - 14:00
P246	Being positive: Assessing the effect of CFTR modulator therapy on sputum sampling and bacterial growth at the All Wales Adult Cystic Fibrosis Centre Oral Presenter: Charlotte Addy, GB	14:00 - 14:00
P247	ETI reduces antibiotic treatment days in people with cystic fibrosis - a real-life observation.n Oral Presenter: Krystyna Poplawska, DE	14:00 - 14:00
P248	Antimicrobial activity of dry powder liposomal loaded rifampicin against <i>Mycobacterium abscessus</i> complex respiratory isolates Oral Presenter: Mona Alhamod, GB	14:00 - 14:00
P249	<i>In vitro</i> activity of liposomal loaded apramycin against <i>Pseudomonas aeruginosa</i> respiratory isolates Oral Presenter: Renlong Na, GB	14:00 - 14:00
P250	Minimum inhibitory concentration targeted antibiotic dose	14:00 - 14:00

	optimization in patients with cystic fibrosis <i>Oral Presenter:</i> David Young, US	
P251	Susceptibility to cefiderocol of an accurately identified collection of 110 <i>Achromobacter</i> strains from cystic fibrosis patients <i>Oral Presenter:</i> Vincent JEAN-PIERRE, FR	14:00 - 14:00
P252	Proposed plan for oxygen therapy as an adjuvant to antibiotics in cystic fibrosis <i>Oral Presenter:</i> Michael Tunney, GB	14:00 - 14:00
P253	Optimisation of inhaled antibiotic prescribing for <i>Pseudomonas aeruginosa</i> infections in people with cystic fibrosis: Further insights from Principal Investigators of the ECFS Clinical Trials Network <i>Oral Presenter:</i> Callum Matthew Sloan, GB	14:00 - 14:00
P254	Clinical effects of <i>Achromobacter xylosoxidans</i> in patients with cystic fibrosis <i>Oral Presenter:</i> Seyda Karabulut, TR	14:00 - 14:00
P255	Risk and time to reinfection with <i>Pseudomonas aeruginosa</i> (Pa) according to the management of Pa infection in children with cystic fibrosis <i>Oral Presenter:</i> Guillaume Thouvenin, FR	14:00 - 14:00
P256	Impact of COVID-19 pandemic and introduction of CFTR modulator therapy on sputum sampling in a large adult cystic fibrosis (CF) unit <i>Oral Presenter:</i> Katie Gaffney, GB	14:00 - 14:00
P257	Microbiological characterisation of methicillin-resistant <i>Staphylococcus aureus</i> isolates recovered from cystic fibrosis people during two Spanish multicentre studies (2013 - 2021) <i>Oral Presenter:</i> Ainhize Maruri-Aransolo, ES	14:00 - 14:00
P258	Phenotypic and gene expressional changes of <i>Pseudomonas aeruginosa</i> isolates from cystic fibrosis patient airways upon estradiol exposure <i>Oral Presenter:</i> Mareike Müller, DE	14:00 - 14:00
P259	Inevitability of treatment after culturing <i>Mycobacterium abscessus</i> complex (MABSC) in cystic fibrosis <i>Oral Presenter:</i> Nicholas Wilson, GB	14:00 - 14:00
P260	Co-culture of <i>Prevotella spp.</i> and <i>Pseudomonas aeruginosa</i> from chronic cystic fibrosis infection in artificial sputum medium <i>Oral Presenter:</i> Enna E. Gibson, GB	14:00 - 14:00
P261	Detection of viable but non culturable <i>Stenotrophomonas maltophilia</i> in cystic fibrosis sputum samples: evidence and perspectives <i>Oral Presenter:</i> Natalia Cirilli, IT	14:00 - 14:00
P262	Respiratory microbiological patterns and comparison in patients with CFTR-related disorders, cystic fibrosis and non-cystic fibrosis bronchiectasis	14:00 - 14:00

	Oral Presenter: Gregorio Basile, IT	
P263	Are cystic fibrosis patients displaying exacerbation symptoms when they culture new pathogens on respiratory samples? Oral Presenter: Paul Wilson, GB	14:00 - 14:00
P264	<i>Pseudomonas aeruginosa</i> (P. aeruginosa) - a review of seasonal patterns and symptom presentation for first and new growths Oral Presenter: Omar Lamptey, GB	14:00 - 14:00
P265	Fast-track biomarker "Anti-Exophiala IgG" in suspected <i>Exophiala</i> spp. infection Oral Presenter: Carsten Schwarz, DE	14:00 - 14:00
P266	Genomic and phenotypic comparisons of <i>Pseudomonas aeruginosa</i> ST27 strains isolated from respiratory tract and domestic environment of a cystic fibrosis patient Oral Presenter: Chloé Dupont, FR	14:00 - 14:00
P267	In vitro efficacy of inhalative antibiotics against <i>Pseudomonas aeruginosa</i> using artificial sputum medium Oral Presenter: Michael Hogardt, DE	14:00 - 14:00
P268	Outcomes after first <i>Pseudomonas aeruginosa</i> (Pa) isolate in cystic fibrosis (CF) patients 2018-2022 Oral Presenter: Ailbhe Marie Murphy, IE	14:00 - 14:00
P269	Proposed plan for oxygen therapy as an adjuvant to antibiotics in cystic fibrosis Oral Presenter: Mette Kolpen, DK	14:00 - 14:00
P270	The prevalence of <i>Candida dubliniensis</i> in cystic fibrosis - a cross sectional, one-year, single centre study Oral Presenter: Emily Krantz, SE	14:00 - 14:00
P271	Impact of <i>Achromobacter</i> spp isolation on lung function in children, a retrospective case-control study Oral Presenter: Nevine Antoun, FR	14:00 - 14:00
P272	Methicillin-resistant <i>Staphylococcus aureus</i> eradication regimens for children and adults with cystic fibrosis Oral Presenter: Rebecca Boyle, US	14:00 - 14:00
P273	Chronic airway infection and resistance pattern in children and adults with cystic fibrosis in Oman. A single centre cross sectional study Oral Presenter: Amjad Al Haddabi, OM	14:00 - 14:00
P274	Identification of <i>bla</i>_{OXA-23} in a mucoid XDR <i>Acinetobacter baumannii</i> isolated from a patient with cystic fibrosis Oral Presenter: Martina Rossitto, IT	14:00 - 14:00
P275	Prevalence of <i>Pseudomonas aeruginosa</i> infection over a five-year period in Albanian children with cystic fibrosis Oral Presenter: Irena Kasmi, AL	14:00 - 14:00
P276	Updating the consensus document: laboratory standards for processing microbiological samples from people with cystic fibrosis	14:00 - 14:00

	<i>Oral Presenter:</i> Rishi Dhillon, GB	
P277	A survey of UK cystic fibrosis centres regarding stopping anti-<i>Pseudomonas aeruginosa</i> (PA) nebulised therapy in children who become free from Pa infection <i>Oral Presenter:</i> Francis Gilchrist, GB	14:00 - 14:00
P278	Partnering for impact: the cystic fibrosis antimicrobial resistance syndicate <i>Oral Presenter:</i> CONSTANCE TAKAWIRA, GB	14:00 - 14:00
P279	Infectious status and lung clearance index <i>Oral Presenter:</i> Mihaela Dediu, RO	14:00 - 14:00
P280	Identification of infection risk areas in the domestic environment by members of the cystic fibrosis community <i>Oral Presenter:</i> Shannon Taylor Venus, GB	14:00 - 14:00
P281	Children with cystic fibrosis demonstrate high rates of asymptomatic carriage of <i>Clostridioides difficile</i> <i>Oral Presenter:</i> Keith Chee Y. Ooi, AU	14:00 - 14:00
P282	Implementing routine assessment of gastrointestinal outcome measures in the cystic fibrosis clinic <i>Oral Presenter:</i> Aleksandra Duffy, GB	14:00 - 14:00
P283	Evaluation of cystic fibrosis-related liver disease in a paediatric cohort <i>Oral Presenter:</i> Saioa Vicente Santamaría, ES	14:00 - 14:00
P284	Assessing the potential of the FIB-4 index as a screening tool for advanced liver fibrosis in an adult cystic fibrosis population <i>Oral Presenter:</i> Stephen Armstrong, GB	14:00 - 14:00
P285	Clinical and genetic risk factors for cystic fibrosis-related liver disease in Egyptian cystic fibrosis children <i>Oral Presenter:</i> Samya Nasr, US	14:00 - 14:00
P287	Characterisation of <i>CFTR</i> mutations in people with cystic fibrosis and severe liver disease who are not eligible for <i>CFTR</i> modulators <i>Oral Presenter:</i> Carla Colombo, IT	14:00 - 14:00
P288	The effects of stopping ursodeoxycholic acid in adult patients with mild cystic fibrosis-related liver disease (CFLD) <i>Oral Presenter:</i> Zaina Aloul, GB	14:00 - 14:00
P289	Intestinal obstruction syndromes in cystic fibrosis <i>Oral Presenter:</i> Alan Anderson, GB	14:00 - 14:00
P291	Screening for Coeliac Disease in adults with cystic fibrosis in Wales <i>Oral Presenter:</i> Dawn Lau, GB	14:00 - 14:00
P292	The effectiveness of pre-colonoscopy bowel preparation for colorectal cancer screening in patients with cystic fibrosis	14:00 - 14:00

in Wales*Oral Presenter:* Anna Sayers, GB

P293	An intensive protocol for bowel preparation in people with cystic fibrosis undergoing screening colonoscopy, a single centre experience <i>Oral Presenter:</i> Tal Lavi, IL	14:00 - 14:00
P294	Colorectal cancer screening in cystic fibrosis, what can iFOBT tell us? <i>Oral Presenter:</i> Nicole Taylor, AU	14:00 - 14:00
P295	The results of colorectal cancer screening in patients with cystic fibrosis in Wales <i>Oral Presenter:</i> Anna Sayers, GB	14:00 - 14:00
P296	Presentation, characteristics and management of obstructive intestinal conditions in cystic fibrosis <i>Oral Presenter:</i> Caitlin Miles, AU	14:00 - 14:00
P297	Dual delivery microbial enzymatic product as alternative PERT for cystic fibrosis patients <i>Oral Presenter:</i> Miguel Angel Poza, ES	14:00 - 14:00
P298	Gastroenterology services for patients with cystic fibrosis across Australia and New Zealand: a multi-stakeholder assessment of patients' and professionals' perspectives <i>Oral Presenter:</i> Tamarah Katz, AU	14:00 - 14:00
P299	Assessing dynamics of abdominal symptoms during a new therapy with elexacaftor/tezacaftor/ivacaftor using the new CFAbd-day2day[®] questionnaire <i>Oral Presenter:</i> Jochen Mainz, DE	14:00 - 14:00
P300	The effect of Highly Effective Modulator therapy (HEMT) on liver function tests of adult patients with cystic fibrosis-related liver disease at one year: the Welsh experience <i>Oral Presenter:</i> Zaina Aloul, GB	14:00 - 14:00
P302	Faecal calprotectin and elastase concentrations in patients before and after treatment with CFTR modulators <i>Oral Presenter:</i> Katarzyna Zybert, PL	14:00 - 14:00
P303	Real-life data for liver function in children on CFTR-modulators in Bulgaria <i>Oral Presenter:</i> Guergana Petrova, BG	14:00 - 14:00
P304	Oral glucose tolerance test in diagnosis of cystic fibrosis-related diabetes in the era of continuous glucose monitoring <i>Oral Presenter:</i> Anjali Santhakumar, GB	14:00 - 14:00
P306	Changes in glucose tolerance in people with cystic fibrosis after initiation of first-generation CFTR modulator treatment <i>Oral Presenter:</i> Rikke Spragge Ekblond, DK	14:00 - 14:00

P307	Updating the consensus document 'Management of cystic fibrosis diabetes' published by Cystic Fibrosis Trust <i>Oral Presenter:</i> Jacqueline Ali, GB	14:00 - 14:00
P308	Strategies for the screening of glucose tolerance abnormalities and diabetes in people with cystic fibrosis: a French position statement <i>Oral Presenter:</i> Laurence Weiss, FR	14:00 - 14:00
P309	UK cystic fibrosis physiotherapists' knowledge and understanding of cystic fibrosis-related diabetes in relation to physical activity and exercise-national survey <i>Oral Presenter:</i> Harbinder Sunsoa, GB	14:00 - 14:00
P310	Patient experience of a novel CFRD education clinic <i>Oral Presenter:</i> Laura Kinsey, GB	14:00 - 14:00
P311	Pregnancy in cystic fibrosis - extending the role of the cystic fibrosis pharmacist in the modulator era <i>Oral Presenter:</i> Elaine Bowman, GB	14:00 - 14:00
P312	Provider perspectives and practices related to sexual and reproductive care provision for males with cystic fibrosis <i>Oral Presenter:</i> Traci Kazmerski, US	14:00 - 14:00
P313	An overview of body composition and metabolic markers in adults with cystic fibrosis: a single centre analysis <i>Oral Presenter:</i> Clodagh Landers, IE	14:00 - 14:00
P314	Cystic fibrosis related to bone disease in children: can it be predicted? <i>Oral Presenter:</i> Ismail Guzelkas, TR	14:00 - 14:00
P315	Bone mineral density in children and adolescents with cystic fibrosis <i>Oral Presenter:</i> Gizem Tamer, NL	14:00 - 14:00
P316	Handgrip strength among children with cystic fibrosis: is there a correlation with lung function? <i>Oral Presenter:</i> Elpis Hatziagorou, GR	14:00 - 14:00
P317	Effect of high intensity interval training versus moderate intensity continuous training on appetite control in people with cystic fibrosis - a randomized controlled study <i>Oral Presenter:</i> Jana Koop, DE	14:00 - 14:00
P318	Weight and vitamin D changes in adults with cystic fibrosis in the West of Scotland post induction of triple therapy during a global pandemic <i>Oral Presenter:</i> Fiona Moore, GB	14:00 - 14:00
P319	Effects of elexacaftor/tezacaftor/ivacaftor (ETI) on nutrition parameters and resting energy expenditure (REE) in people with cystic fibrosis (pwCF) <i>Oral Presenter:</i> Lenny Sasse, DE	14:00 - 14:00
P320	Comparing faecal elastase, pancreatic enzyme doses, HbA1c and insulin doses in adult cystic fibrosis patients pre- and 6 months post-commencement of Kaftrio® <i>Oral Presenter:</i> Lidia Sheibani, GB	14:00 - 14:00

P321	Impact of short-term CFTR modulators treatment on changes in nutritional and glucose status in cystic fibrosis paediatric patients with different glucose tolerance	14:00 - 14:00
	<i>Oral Presenter:</i> Monika Mielus, PL	
P322	Lipid profiles in a Scottish adult cystic fibrosis (CF) centre - prevalence and significance in an aging population	14:00 - 14:00
	<i>Oral Presenter:</i> Lianne Robb, GB	
P323	Does elexacaftor/tezacaftor/ivacaftor triple therapy significantly raise blood lipids? A longitudinal cohort study	14:00 - 14:00
	<i>Oral Presenter:</i> Ronan Docherty, GB	
P324	Essential fatty acid deficiency in children and young adults with cystic fibrosis - micronutrient assessment is still an actual topic	14:00 - 14:00
	<i>Oral Presenter:</i> Marcela Kreslová, CZ	
P325	A paradigm shift in cystic fibrosis nutritional care: clinicians views on the management of patients with overweight and obesity	14:00 - 14:00
	<i>Oral Presenter:</i> Joanna Snowball, GB	
P326	Impact of the elexacaftor/tezacaftor/ivacaftor on the nutrition parameters and gastrointestinal symptoms in adult cystic fibrosis patients in the cystic fibrosis centre of Brno, Czech Republic	14:00 - 14:00
	<i>Oral Presenter:</i> Nela Stastna, CZ	
P327	Cross-cultural adaptation and measurement properties of the Brazilian version of the Knowledge of Disease Management-CF-Adolescent questionnaire	14:00 - 14:00
	<i>Oral Presenter:</i> Hilda Angelica Iturriaga-Jimenez, BR	
P329	Audit of change in weight and BMI centile in children on elexacaftor/tezacaftor/ivacaftor in the Liverpool paediatric cystic fibrosis population	14:00 - 14:00
	<i>Oral Presenter:</i> Clare J Woodland, GB	
P330	One-year assessment of body composition in cystic fibrosis patients on elexacaftor/tezacaftor/ivacaftor	14:00 - 14:00
	<i>Oral Presenter:</i> Veronica Zamponi, IT	
P331	What nutritional advice is being given to people with cystic fibrosis and hepatic steatosis?	14:00 - 14:00
	<i>Oral Presenter:</i> Maeve O'Driscoll, GB	
P332	Description of nutritional status in a cohort of paediatric patients with cystic fibrosis analysis of body composition changes after initiation of treatment with elexacaftor/tezacaftor/ivacaftor	14:00 - 14:00
	<i>Oral Presenter:</i> Saioa Vicente Santamaría, ES	
P333	Refresh: patient experience of an online healthy eating webinar designed specifically for adults with cystic fibrosis on CFTR modulators	14:00 - 14:00
	<i>Oral Presenter:</i> Katie Marsden, GB	

P334	Body composition changes after ellexacaftor/tezacaftor/ivacaftor treatment in adults with cystic fibrosis - single centre experience <i>Oral Presenter:</i> Andrea Vukić Dugac, HR	14:00 - 14:00
P335	Survey of oral health survey in children in a secondary paediatric cystic fibrosis service <i>Oral Presenter:</i> Naveen Rao, GB	14:00 - 14:00
P336	Evolution of the body composition of people with cystic fibrosis 6 months after the introduction of treatment with exacaftor/tezacaftor/ivacaftor (ETI) <i>Oral Presenter:</i> Camille Audousset, FR	14:00 - 14:00
P337	Bone health status and audit of DEXA scan frequency of adults with cystic fibrosis at a UK centre <i>Oral Presenter:</i> Eliza C Tassone, GB	14:00 - 14:00
P338	Body composition in children with cystic fibrosis receiving triple combination therapy <i>Oral Presenter:</i> Uros Krivec, SI	14:00 - 14:00
P339	Diet quality assessment of adults with cystic fibrosis - comparison to population dietary guidelines: a cross-sectional study <i>Oral Presenter:</i> Cian Greaney, IE	14:00 - 14:00
P340	The impact of modulators on faecal elastase in children with cystic fibrosis <i>Oral Presenter:</i> Elizabeth Sheppard, GB	14:00 - 14:00
P341	Pancreatic status and enzyme usage in the age of CFTR modulator treatment <i>Oral Presenter:</i> Eliza C Tassone, GB	14:00 - 14:00
P342	Serum sodium, chloride and potassium levels after start CFTR modulators treatment in CF pediatric patients <i>Oral Presenter:</i> Monika Mielus, PL	14:00 - 14:00
P343	Combination vitamin supplementation therapy in children with cystic fibrosis <i>Oral Presenter:</i> Anirban Maitra, GB	14:00 - 14:00
P344	How should Pancreatic Enzyme Replacement Therapy (PERT) be administered to infants? <i>Oral Presenter:</i> Eleanor McGray, GB	14:00 - 14:00
P406	Effects of the treatment with ellexacaftor/tezacaftor/ivacaftor on aerobic fitness of adolescents with cystic fibrosis <i>Oral Presenter:</i> Marcio Vinicius Fagundes Donadio, BR	14:00 - 14:00
P407	Physical activity and sedentary behavior in cystic fibrosis patients before and after triple modulator therapy: a single-centre retrospective study <i>Oral Presenter:</i> Anna Barrero, FR	14:00 - 14:00
P408	The effect of major life changes on exercise and physical activity in people with cystic fibrosis <i>Oral Presenter:</i> Owen William Tomlinson, GB	14:00 - 14:00

P409	Monitoring adherence to nebulised therapies during the first UK lockdown and school closure period of the COVID-19 pandemic in children with cystic fibrosis <i>Oral Presenter:</i> Charlotte Smith, GB	14:00 - 14:00
P410	The impact of the COVID 19 pandemic and modulator therapy on routine respiratory sampling in paediatric cystic fibrosis (CF) <i>Oral Presenter:</i> Bethany Millman, GB	14:00 - 14:00
P411	Thinking out of the box: twinning in the era of the COVID-19 pandemic <i>Oral Presenter:</i> Simona Mosescu, RO	14:00 - 14:00
P412	A patient-led review of outpatient services following the COVID-19 pandemic and widespread introduction of highly-effective modulator treatments and remote monitoring devices <i>Oral Presenter:</i> Jocelyn Choyce, GB	14:00 - 14:00
P413	How to use telemonitoring for the detection of respiratory exacerbations in cystic fibrosis: effectiveness and adherence <i>Oral Presenter:</i> Letizia Luciani, IT	14:00 - 14:00
P414	The effect of elexacaftor/tezacaftor/ivacaftor on functional capacity in children with cystic fibrosis <i>Oral Presenter:</i> Sophia Ward, GB	14:00 - 14:00
P415	A service evaluation to quantify real-world spirometry grading and frequency data following the introduction of the NuvoAir Air Next lung function device in one UK cystic fibrosis (CF) centre <i>Oral Presenter:</i> Jamie Watkins, GB	14:00 - 14:00
P416	Self-reported levels of physical activity do not give an accurate insight into the habitual levels of physical activity in children and young people with cystic fibrosis <i>Oral Presenter:</i> Helen Douglas, GB	14:00 - 14:00
P418	Patient satisfaction and clinical effectiveness of using the I-neb in children with cystic fibrosis (CF) in Wales <i>Oral Presenter:</i> Kathryn G Welsh, GB	14:00 - 14:00
P419	Lung deposition of nebulized drug using favorite nebulisation compared to mesh nebuliser in healthy and cystic fibrosis patients: a randomised cross-over trial <i>Oral Presenter:</i> Anne-Sophie Aubriot, BE	14:00 - 14:00
P420	A pilot study looking at gamification to improve the experience of airway clearance for children with cystic fibrosis <i>Oral Presenter:</i> Adam Walsh, GB	14:00 - 14:00
P421	Use of mechanical insufflation-exsufflation in children with	14:00 - 14:00

cystic fibrosis*Oral Presenter:* Kieren James Lock, GB

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|------|---|---------------|
| P422 | PEPing up your physio: how manometers improve adherence to airway clearance techniques (ACT) in children with chronic respiratory disease
<i>Oral Presenter:</i> Rachel Hawes, GB | 14:00 - 14:00 |
| P423 | A low-cost positive expiratory pressure device for people with cystic fibrosis in developing countries
<i>Oral Presenter:</i> Jamie Wood, US | 14:00 - 14:00 |
| P424 | Short-term effects of positive expiratory pressure on peripheral ventilation inhomogeneity in children with cystic fibrosis: a randomised sham-controlled crossover study
<i>Oral Presenter:</i> Simone Gambazza, IT | 14:00 - 14:00 |
| P425 | Identifying barriers to completing chest physiotherapy in the early years
<i>Oral Presenter:</i> Melissa Richmond, CA | 14:00 - 14:00 |
| P426 | The management of musculoskeletal issues in cystic fibrosis: the respiratory physiotherapists' perspective
<i>Oral Presenter:</i> Stephanie Graham, GB | 14:00 - 14:00 |
| P427 | Evaluating the provision of exercise for in-patients with cystic fibrosis across cystic fibrosis centres in the UK and identification of challenges and barriers surrounding delivery of care
<i>Oral Presenter:</i> Stephanie Graham, GB | 14:00 - 14:00 |
| P428 | An evaluation of the current Manchester Adult Cystic Fibrosis Centre (MACFC) exercise service
<i>Oral Presenter:</i> Nicole Petch, GB | 14:00 - 14:00 |
| P429 | The first UK cystic fibrosis trust exercise practitioner fellowship - an insight for the future provision of exercise services among individuals with cystic fibrosis and healthcare professionals
<i>Oral Presenter:</i> Lucia Diego-Vicente, GB | 14:00 - 14:00 |
| P430 | Are physical activity levels in children with cystic fibrosis associated to adherence to treatment?
<i>Oral Presenter:</i> Kieren James Lock, GB | 14:00 - 14:00 |
| P431 | Factors related to higher level of physical activity in Swedish cystic fibrosis-patients - a registry-based cross sectional study
<i>Oral Presenter:</i> Sofia Wilhelmsson, SE | 14:00 - 14:00 |
| P432 | Physical working capacity and health-related quality of life in adolescents and adult people with cystic fibrosis in Sweden
<i>Oral Presenter:</i> Karolina Wallin, SE | 14:00 - 14:00 |
| P433 | Association of physical activity with quadriceps force and functional performance in children with cystic fibrosis and healthy peers
<i>Oral Presenter:</i> Manon Kinaupenne, BE | 14:00 - 14:00 |

P434	Exploring cystic fibrosis patients' attitudes and beliefs regarding exercise participation during hospitalization at a regional adult cystic fibrosis unit <i>Oral Presenter:</i> Stephanie Graham, GB	14:00 - 14:00
P435	Physical activity in young people with cystic fibrosis living in Latvia <i>Oral Presenter:</i> Arta Ūdre, LV	14:00 - 14:00
P436	Feasibility and effectiveness of a low-impact, virtual exercise programme for adults with cystic fibrosis <i>Oral Presenter:</i> Ruth Watson, GB	14:00 - 14:00
P437	Some do and some don't, but who will and who won't: can patterns of physical activity maintenance be predicted in children and young people with cystic fibrosis? <i>Oral Presenter:</i> Helen Douglas, GB	14:00 - 14:00
P438	Clinimetric properties of field exercise tests in cystic fibrosis: a systematic review <i>Oral Presenter:</i> Jen Corda, AU	14:00 - 14:00
P439	Feasibility of performing the three-minute step test with remote supervision in children and adolescents with cystic fibrosis <i>Oral Presenter:</i> Marcio Vinicius Fagundes Donadio, BR	14:00 - 14:00
P440	Can the 6-minute walking test assess physical activity level among people with cystic fibrosis? <i>Oral Presenter:</i> Elpis Hatziaorou, GR	14:00 - 14:00
P441	The A-Step - an incremental exercise test defying space and infection control measures <i>Oral Presenter:</i> Natascha Remus, FR	14:00 - 14:00
P442	Sleeping issues in children, adolescents and young adults <i>Oral Presenter:</i> Lidija Skocir, SI	14:00 - 14:00
P443	A sprint not a marathon: initial outcomes from a physiotherapist-led specialist cystic fibrosis (CF) Cardio-Pulmonary Exercise Testing (CPET) service for Wales <i>Oral Presenter:</i> Rachel Young, GB	14:00 - 14:00
P444	Self-perception of fitness levels vs CPET findings in adolescents with cystic fibrosis <i>Oral Presenter:</i> Sioned Davies, GB	14:00 - 14:00
P445	A marathon not a sprint: Establishing a physiotherapist led cystic fibrosis specific Cardio Pulmonary Exercise Testing (CPET) service for Wales <i>Oral Presenter:</i> Rachel Young, GB	14:00 - 14:00
P446	Physiotherapy input to cystic fibrosis screen positive indeterminate diagnosis patients at a regional paediatric cystic fibrosis unit <i>Oral Presenter:</i> Nuala Harnett, GB	14:00 - 14:00
P447	Effects of online core stabilisation exercises on posture in cystic fibrosis children <i>Oral Presenter:</i> Kubra Kilic, TR	14:00 - 14:00

Industry Sessions

14:15 - 14:45

Tomorrow Lounge

MedTech Event*Special Symposium*

15:00 - 16:30

R1

SS01 - Best of Journal of Cystic Fibrosis, Lancet Respiratory Medicine and European Respiratory Journal*Chair:* Patrick Flume, US*Chair:* Marcus Mall, DE*Chair:* Emma Grainger, GB**Introduction**

15:00 - 15:05

Journal of Cystic Fibrosis

15:05 - 15:25

Estimating the Minimum Sample Size for Interventional and Observational Studies Using the Lung Clearance Index as an Endpoint

15:25 - 15:25

Speaker: Felix Ratjen, CA*Discussant:* Sonya Heltshe, US**Lancet Respiratory Medicine**

15:25 - 15:45

Safety and efficacy of vanzacaftor-tezacaftor-deutivacaftor in adults with cystic fibrosis: randomised, double-blind, controlled, phase 2 trials

15:45 - 15:45

Speaker: Alexander Horsley, GB*Discussant:* Felix Ratjen, CA**European Respiratory Journal**

15:45 - 16:05

The French Compassionate Program of elexacaftor-tezacaftor-ivacaftor in people with cystic fibrosis with advanced lung disease and no F508del CFTR variant

16:05 - 16:05

Speaker: Pierre-Régis Burgel, FR*Discussant:* Peter Barry, GB**Discussion - Hot Topics in Publishing**

16:05 - 16:30

Workshop

15:00 - 16:30

R2

WS01 - WS01 - Maximizing health: exploring novel strategies for exercise testing, physiotherapy, and musculoskeletal screening*Chair:* Marlies Wagner, AT*Chair:* Emma Dixon, GB

WS01.01

Cardiopulmonary exercise testing provides prognostic information in advanced cystic fibrosis lung disease

15:00 - 15:15

Oral Presenter: Thomas Radtke, CH

WS01.02

The current state of play regarding exercise testing in cystic fibrosis: co-development with the community

15:15 - 15:30

Oral Presenter: Donald Urquhart, GB

WS01.03

International Physiotherapy Group for Cystic Fibrosis (IPG/CF): international survey of physiotherapy practices in 2022 including airway clearance techniques, inhaled mucolytics and exercise

15:30 - 15:45

	<i>Oral Presenter:</i> Brenda Button, AU	
WS01.04	Longtime follow-up on exercise capacity and quality of life in people with cystic fibrosis receiving ellexacaftor/tezacaftor/ivacaftor - a Copenhagen cohort <i>Oral Presenter:</i> Lue Drasbæk Philipsen, DK	15:45 - 16:00
WS01.05	Development of a musculoskeletal screening tool for children and young people with cystic fibrosis (Addenbrooke's MST): initial findings <i>Oral Presenter:</i> Kieren James Lock, GB	16:00 - 16:15
WS01.06	Inducing sputum in the adult cystic fibrosis post-modulator era <i>Oral Presenter:</i> Alexander Williams, GB	16:15 - 16:30
<i>Workshop</i>		
15:00 - 16:30		R3
WS02 - WS02 - Fertility, pregnancy and gender-related topics		
<i>Chair:</i> Malena Cohen-Cymberknoh, IL		
<i>Chair:</i> Quitterie Reynaud, FR		
WS02.01	Decreased fertility in female cystic fibrosis patients: peering into the endometrial factor using cutting-edge organoid models <i>Oral Presenter:</i> Ellen De Pauw, BE	15:00 - 15:15
WS02.02	Ovarian reserve in women with cystic fibrosis: is this a cause of subfertility? <i>Oral Presenter:</i> Malena Cohen-Cymberknoh, IL	15:15 - 15:30
WS02.03	Correlating menstrual cycles and cystic fibrosis symptoms among women with cystic fibrosis in the era of highly effective modulators: early findings of the MENSTRUAL study <i>Oral Presenter:</i> Elinor Langfelder-Schwind, US	15:30 - 15:45
WS02.04	Triple HEMT in Pregnancy and Lactation: effects on the developing lung, gut and pancreas <i>Oral Presenter:</i> Elena Schneider-Futschik, AU	15:45 - 16:00
WS02.05	Obstetric and neonatal outcomes in cystic fibrosis (CF) show minimal variability with and without ellexacaftor/tezacaftor/ivacaftor (ETI) <i>Oral Presenter:</i> Amy Downes, GB	16:00 - 16:15
WS02.06	Prospectively evaluating maternal and fetal outcomes in the era of CFTR modulators: MAYFLOWERS study interim results <i>Oral Presenter:</i> Jennifer Taylor-Cousar, US	16:15 - 16:30
<i>Workshop</i>		
15:00 - 16:30		R4
WS03 - WS03 - Disease severity and survival: insight from registries		
<i>Chair:</i> Andreas Pflieger, AT		
<i>Chair:</i> Elpis Hatziaorou, GR		
WS03.01	Differences in disease severity among different residual function mutations: data from the ECFS Patient Registry	15:00 - 15:15

	Oral Presenter: Meir Mei-Zahav, IL	
WS03.02	Keep your enemies close: natural foes, <i>Pseudomonas aeruginosa</i> and <i>Staphylococcus aureus</i> are associated with fewer adverse clinical consequences when present in patients with chronic co-infection Oral Presenter: Micaela Mossop, GB	15:15 - 15:30
WS03.03	Clinical outcomes and long-term survival in cystic fibrosis (CF) lung transplant recipients in Belgium Oral Presenter: Muriel Thomas, BE	15:30 - 15:45
WS03.04	Survival of people with cystic fibrosis: complete overview in France and first evaluation of the impact of modulators Oral Presenter: Virginie Scotet, FR	15:45 - 16:00
WS03.05	Multicenter validation of the cystic fibrosis-ABLE score as a predictor of outcome and therapeutic response in cystic fibrosis Oral Presenter: Oliver James McElvaney, US	16:00 - 16:15
WS03.06	Effective cystic fibrosis team training in low - and medium income countries leads to significant outcome improvements Oral Presenter: Hector Gutierrez, US	16:15 - 16:30
 <i>Workshop</i> 15:00 - 16:30		
		R5
WS04 - WS04 - Understanding and correcting the CF cell: modulators and beyond		
Chair: Carlos M Farinha, PT		
Chair: Carla Ribeiro, US		
WS04.01	A PI3Ky mimetic peptide promotes F508del-CFTR plasma membrane stabilization through Protein Kinase D1 Oral Presenter: Alessandra Murabito, IT	15:00 - 15:15
WS04.02	Impact of cholesterol on the functioning of CFTR modulators Oral Presenter: Dorna Ravamehr-Lake, CA	15:15 - 15:30
WS04.03	Elexacaftor/tezacaftor/ivacaftor (ETI) treatment corrects the salt-losing phenotype in people with cystic fibrosis (pwCF) Oral Presenter: Peder Berg, DK	15:30 - 15:45
WS04.04	CFTR modulator triple combination treatment modifies exhaled breath of children with cystic fibrosis within a week Oral Presenter: Emmanuelle Bardin, FR	15:45 - 16:00
WS04.05	Alternative targets for the treatment of cystic fibrosis basic defect Oral Presenter: Floriana Guida, IT	16:00 - 16:15
WS04.06	Validation of rectal organoid morphology analysis (ROMA) as a novel physiological CFTR assay for diagnosis of cystic fibrosis Speaker: Senne Cuyx, BE	16:15 - 16:30

ECFS Tomorrow Lounge Session

15:00 - 16:00

Tomorrow Lounge

Dietetic management of the "well" person with CF*Chair:* Fiona Moore, GB*Chair:* Elizabeth Owen, GB*Workshop*

17:00 - 18:30

R1

WS05 - WS05 - Fresh results of clinical trials*Chair:* Damian Downey, GB*Chair:* George Retsch-Bogart, US

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|---------|---|---------------|
| WS05.01 | AAV mediated gene therapy for cystic fibrosis: interim results from a phase 1/2 clinical trial
<i>Oral Presenter:</i> Jennifer Taylor-Cousar, US | 17:00 - 17:15 |
| WS05.02 | A Phase-1 multiple ascending dose healthy volunteer study to evaluate the safety, tolerability, and pharmacokinetics of GDC-6988, a dry powder formulation of a selective inhaled potentiator of TMEM16A
<i>Oral Presenter:</i> Joshua Galanter, US | 17:15 - 17:30 |
| WS05.03 | A phase 3b study of the effects of elexacaftor/tezacaftor/ivacaftor (ELX/TEZ/IVA) on glucose tolerance in people with cystic fibrosis (CF) and abnormal glucose metabolism
<i>Oral Presenter:</i> Philip Robinson, AU | 17:30 - 17:45 |
| WS05.04 | Safety and efficacy of ivacaftor (IVA) in children aged 1 to <4 months with cystic fibrosis assessed with an innovative clinical trial design
<i>Oral Presenter:</i> Jane Davies, GB | 17:45 - 18:00 |
| WS05.05 | Randomised withdrawal of hypertonic saline in those with lower lung function after receiving elexacaftor/tezacaftor/ivacaftor; a sub-study of the SIMPLIFY Trial
<i>Oral Presenter:</i> Nicole Hamblett, US | 18:00 - 18:15 |
| WS05.06 | Phase 1/2a randomized, double-blind, placebo-controlled study: safety, Pk, and Efficacy outcome measures of inhaled Gallium Citrate (AR-501) in <i>P. Aeruginosa</i> infected Cystic Fibrosis patients
<i>Oral Presenter:</i> Hasan Jafri, US | 18:15 - 18:30 |

Workshop

17:00 - 18:30

R2

WS06 - WS06 - Diagnostic tools for monitoring lung disease*Chair:* Florian Singer, AT*Chair:* Felix Ratjen, CA

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|---------|--|---------------|
| WS06.01 | Outcome measures for assessment of lung function - cross-sectional analysis of relationships between impulse oscillometry, electrical impedance tomography, multiple breath washout and spirometry
<i>Oral Presenter:</i> Gemma Stanford, GB | 17:00 - 17:15 |
| WS06.02 | How accurate is home spirometry? Comparison of home to office spirometry in the PROMISE study | 17:15 - 17:30 |

	Oral Presenter: Margaret Rosenfeld, US	
WS06.03	A real-world evaluation of remote monitoring in cystic fibrosis care: a mixed-methods multicentre observational study Oral Presenter: Marc C. Oppelaar, NL	17:30 - 17:45
WS06.04	Lung volumes as possible factors explaining the heterogeneity in the response to elexacaftor/tezacaftor/ivacaftor in patients with cystic fibrosis and advanced lung disease Oral Presenter: Chiara Premuda, IT	17:45 - 18:00
WS06.05	The volume of trapped air (VTA) from multiple breath washout (MBW) substantially underestimates the extent of trapped air (TA) demonstrated with spirometry-controlled computed tomography Oral Presenter: Christopher Short, GB	18:00 - 18:15
WS06.06	Effect of elexacaftor/tezacaftor/ivacaftor on structural lung damage after one year detected by magnetic resonance imaging Oral Presenter: Samantha Conci, IT	18:15 - 18:30
Workshop 17:00 - 18:30		R3
WS07 - WS07 - Pathogenesis and adaptation of microbes in the CF airways		
Chair: Lucas Hoffman, US		
Chair: Dorothea Appelt, AT		
WS07.01	Inter-species interactions alter antimicrobial susceptibility in polymicrobial cultures Oral Presenter: Eva Bernadett Benyei, GB	17:00 - 17:15
WS07.02	Zebrafish infection model to assess persistence of <i>Pseudomonas aeruginosa</i> clinical isolates and treatment efficiency Oral Presenter: Anne Blanc-Potard, FR	17:15 - 17:30
WS07.03	A long-term study of <i>Pseudomonas aeruginosa</i> early patient isolate adaptation to the hypoxic environment Oral Presenter: Joanna Drabinska, IE	17:30 - 17:45
WS07.04	Hostile takeover factors of <i>Pseudomonas aeruginosa</i>: contact dependent secretion systems Oral Presenter: Alice Christina Zammit Collins, GB	17:45 - 18:00
WS07.05	Unraveling the pathogenicity of <i>Mycobacterium abscessus</i> clinical isolates in cystic fibrosis pulmonary epithelial cell and mouse models of respiratory infection Oral Presenter: Nicola Ivan Lorè, IT	18:00 - 18:15
WS07.06	CFTR impacts SARS-CoV-2 infection in cystic fibrosis Oral Presenter: Cristina Cigana, IT	18:15 - 18:30

Workshop

17:00 - 18:30

R4

WS08 - WS08 - Evaluating and treating gastrointestinal and liver diseases

Chair: Helmut Ellemunter, AT

Chair: Michael Wilschanski, IL

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|---------|---|---------------|
| WS08.01 | Evaluation of liver steatosis in people with cystic fibrosis using controlled attenuated parameter obtained by Fibroscan | 17:00 - 17:15 |
| | Oral Presenter: Stephanie Van Biervliet, BE | |
| WS08.02 | Change in hepatobiliary ultrasound measures, incl. shear wave elastography, after introduction of elexacaftor/tezacaftor/ivacaftor: results from 12-month follow-up in the Danish cystic fibrosis cohort | 17:15 - 17:30 |
| | Oral Presenter: Christine Højte, DK | |
| WS08.03 | Evaluating lung, liver and gut structure and function using MRI in 6 to 11 year olds - preliminary results from the GIFT-cystic fibrosis junior cohort | 17:30 - 17:45 |
| | Oral Presenter: Alexander Yule, GB | |
| WS08.04 | Faecal Immunochemical Testing - 'FIT' for purpose for bowel screening in older cystic fibrosis patients? | 17:45 - 18:00 |
| | Oral Presenter: Karuna Sapru, GB | |
| WS08.05 | Colorectal cancer-associated bacteria in adults with cystic fibrosis | 18:00 - 18:15 |
| | Oral Presenter: Laura Caley, GB | |
| WS08.06 | Pilot study evaluating the potential of <i>Akkermansia muciniphila</i> and <i>Faecalibacterium prausnitzii</i> as cystic fibrosis-specific probiotics utilising patient-derived organoid models | 18:15 - 18:30 |
| | Oral Presenter: Keith Chee Y. Ooi, AU | |

Workshop

17:00 - 18:30

R5

WS09 - WS09 - Genetics from different perspectives

Chair: Milan Jr Macek, CZ

Chair: Margarida Amaral, PT

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|---------|---|---------------|
| WS09.01 | Extending the success of Trikafta to rare mutations - insights from CFTR structure and modulator binding | 17:00 - 17:15 |
| | Oral Presenter: Batsheva Kerem, IL | |
| WS09.02 | Therotyping molecular defects of CFTR rare variants in patient derived rectal organoids | 17:15 - 17:30 |
| | Oral Presenter: Anabela S. Ramalho, BE | |
| WS09.03 | Delivery characterisation of SPL84 Inhaled Antisense Oligonucleotide | 17:30 - 17:45 |
| | Oral Presenter: Gili Hart, IL | |
| WS09.04 | Enhancement of channel activity of the CFTR protein delivered via gene therapy vectors | 17:45 - 18:00 |

Oral Presenter: Jim Hu, CA

WS09.05 **Aggregated genetic modifiers predict lung dysfunction at age 5-6 years in children with cystic fibrosis** 18:00 - 18:15

Oral Presenter: Philip Farrell, US

WS09.06 **Genetic testing donors of gametes for cystic fibrosis should be mandatory** 18:15 - 18:30

Oral Presenter: Guergana Petrova, BG

ECFS Tomorrow Lounge Session

17:00 - 18:00

Tomorrow Lounge

Psychosocial aspects of getting older with CF

Chair: Jacqueline Ali, GB

Chair: Sue Braun, BE

Friday, 09. June 2023*Industry Sessions*

07:15 - 08:15

R5

Satellite Symposium*Meet the Experts*

07:30 - 08:20

MTE04 - Meet the Experts 04 - How better characterisation of CFTR gene variants has helped our patients; will extended gene sequencing be a help or a hindrance?*Chair:* Caroline Raynal, FR*Chair:* Karen Raraigh, US*Meet the Experts*

07:30 - 08:20

Meet the Experts*Meet the Experts*

07:30 - 08:20

MTE06 - Meet the Experts 06 - Post SIMPLIFY - what pulmonary therapies to rationalise and how to do it?*Chair:* Daniel Peckham, GB*Chair:* Alex H. Gifford, US*Meet the Experts*

07:30 - 08:20

MTE05 - Meet the Experts 05 - Remote psychological counselling tools and strategies*Chair:* Marieke Verkleij, NL*Chair:* Anna M. Georgiopoulos, US*Symposium*

08:30 - 10:00

R1

S11 - Symposium 11 - Are we ready for new clinical guidelines?*Chair:* Kevin Southern, GB*Chair:* Isabelle Fajac, FR**What pathogens should we eradicate?**

08:30 - 08:52

Speaker: Tavs Qvist, DK**Antibiotic strategies for exacerbation management**

08:52 - 09:14

Speaker: Susanna Esposito, IT**How should we treat bronchopulmonary aspergillosis?**

09:14 - 09:36

Speaker: Carsten Schwarz, DE**Are referral criteria for lung transplant different for people with CFTR modulators?**

09:36 - 10:00

Speaker: Thomas Daniels, GB

Symposium

08:30 - 10:00

R2

S12 - Symposium 12 - Essential nursing skills in the CFTR modulators era

Chair: Tina D'Hondt, BE

Chair: Daniel Office, GB

Which treatments cannot be stopped and why - Supporting adherence in a healthier population 08:30 - 08:52

Speaker: Cora de Kiviet, NL

The impact of CFTR modulators on the liver, the gut and absorption 08:52 - 09:14

Speaker: Joanna Whitehouse, GB

Managing insulin and glycemic control after CFTR modulator initiation 09:14 - 09:36

Speaker: Esp rie Burnet, FR

Speaker: Deborah Grunewald, FR

Life goes on - a patient perspective of their future aspirations and what they want from their cystic fibrosis team 09:36 - 10:00

Speaker: Katrien Van Gompel, BE

Symposium

08:30 - 10:00

R3

S13 - Symposium 13 - Endocrinology and bone health in the era of CFTR modulators

Chair: Monika Mielus, PL

Chair: Dimitri Declercq, BE

The abnormalities in glucose metabolism - how and when to treat? 08:30 - 08:52

Speaker: Amir Moheet, US

Oral glucose lowering agents - do we recommend them in cystic fibrosis? 08:52 - 09:14

Speaker: Amanda Brennan, GB

The effect of CFTR modulators on the endocrine system 09:14 - 09:36

Speaker: Laurence Kessler, FR

Consequences of CFTR modulators on bone health 09:36 - 10:00

Speaker: Susannah King, AU

Symposium

08:30 - 10:00

R4

S14 - Symposium 14 - Assessment of *in vitro* biomarkers of CFTR function for prediction of clinical benefit

Chair: David Sheppard, GB

Chair: Anabela Santo Ramalho, BE

Correlating CFTR function in cell lines with clinical features to inform personalised treatment of cystic fibrosis 08:30 - 08:52

Speaker: Garry Cutting, US

Using cystic fibrosis primary airway epithelial cells to predict lung function improvements by modulator therapies 08:52 - 09:14

Speaker: Isabelle Sermet-Gaudelus, FR

Comparison of organoid swelling and *in vivo* biomarkers of CFTR function to determine modulator effects 09:14 - 09:36

Speaker: Simon Graeber, DE

Can induced pluripotent stem cells be a feasible model for theratyping? 09:36 - 10:00

Speaker: Amy Wong, CA

Symposium

08:30 - 10:00

R5

S15 - Symposium 15 - Pharmacovigilance in the real world: challenges and opportunities in an era of CFTR modulators

Chair: Gwyneth Davies, GB

Chair: Meir Mei-Zahav, IL

PRO - We capture the right outcomes on CF Registries to support post-authorisation safety studies 08:30 - 08:48

Speaker: Lutz Naehrlich, DE

CON - We capture the right outcomes on CF Registries to support post-authorisation safety studies 08:48 - 09:06

Speaker: Jamie Duckers, GB

Discussion 09:06 - 09:16

Everybody matters - inclusive approaches to population-level analyses of the effectiveness of new therapies in cystic fibrosis 09:16 - 09:38

Speaker: Elizabeth Cromwell, US

Personalised medicine vs overmedicalisation in an era of CFTR modulators: insights from CF Registries 09:38 - 10:00

Speaker: Rita Padoan, IT

Symposium

10:30 - 12:00

R1

S16 - Symposium 16 - CFTR modulators: opportunities and challenges

Chair: Nicholas Simmonds, GB

Chair: Hettie Janssens, NL

Life-long CFTR modulation - is it sustained? 10:30 - 10:52

Speaker: Pierre-Régis Burgel, FR

Extra-pulmonary benefits of CFTR modulators 10:52 - 11:14

Speaker: Bradley Quon, CA

CFTR modulators: options for those who are intolerant 11:14 - 11:36

Speaker: Daniel Peckham, GB

CFTR modulators and neonates - how low should you go! 11:36 - 12:00

Speaker: Margaret Rosenfeld, US

Symposium

10:30 - 12:00

R2

S17 - Symposium 17 - Masterclass in clinical CF - great cases

Chair: Peter Barry, GB

Chair: Andreas Hector, CH

Mycobacterium abscessus in a young child with CF - the role of IFNγ immune dysregulation	10:30 - 10:45
<i>Speaker:</i> Kushalini Hillson, GB <i>Discussant:</i> Andreas Hector, CH	
After the honeymoon with CFTR-modulators, a rude awakening with sudden and exceptionally severe infection.	10:45 - 11:00
<i>Speaker:</i> Tobias Schmidgall, DE <i>Discussant:</i> Peter Barry, GB	
Mind over modulator: A challenging case of mood disturbance in a young female following the introduction of Elexacaftor/Tezacaftor/Ivacaftor (ETI).	11:00 - 11:15
<i>Speaker:</i> Miriam Cameron, AU <i>Discussant:</i> Barry Plant, IE	
Pushing the limits in the era of CFTR modulators	11:15 - 11:30
<i>Speaker:</i> Georgia Mitropoulou, CH <i>Discussant:</i> Lieven Dupont, BE	
Altered sweat test: not only CFTR is involved.	11:30 - 11:45
<i>Speaker:</i> Joana Quaresma Vázquez, ES <i>Discussant:</i> Silvia Gartner, ES	
Increased breathless and reduced exercise tolerance in a Cystic Fibrosis patient undergoing a phase 3 randomized control trial. An unexpected and unrelated diagnosis	11:45 - 12:00
<i>Speaker:</i> Kevin Deasy, IE <i>Discussant:</i> Ian Balfour-Lynn, GB	

Symposium

10:30 - 12:00

R3

S18 - Symposium 18 - Pregnancy and early parenthood for mums and dads with cystic fibrosis - it's a team effort!*Chair:* Jennifer Taylor-Cousar, US*Chair:* David Kinoo, BE

Parenthood dilemmas - optimal medical and obstetric management strategies for pre/during and post CF pregnancy and early parenthood	10:30 - 10:52
<i>Speaker:</i> Imogen Felton, GB	
Dinner for two (how to optimise nutrition pre/during/post pregnancy and early parenthood)	10:52 - 11:14
<i>Speaker:</i> Francis Hollander-Kraaijeveld, NL	
Airway clearance and exercise - optimal physiotherapy interventions during pregnancy and early parenthood for mothers and fathers with cystic fibrosis	11:14 - 11:36
<i>Speaker:</i> Brenda Button, AU	
Keeping the "me" in pregnancy (how to cope/support CF adults pre/during/post pregnancy and early parenthood)	11:36 - 12:00
<i>Speaker:</i> Trudy Havermans, BE	

Symposium

10:30 - 12:00

R4

S19 - Symposium 19 - Is there a future for phage therapy in cystic fibrosis?*Chair:* Oana Ciofu, DK*Chair:* Gisli Einarsson, GB

Phage therapy to circumvent bacterial resistance in chronic infections 10:30 - 10:52

Speaker: Benjamin Chan, US

Combination therapy to fight bacterial infections: phages, antibiotics and immune adjuvants 10:52 - 11:14

Speaker: Anna Pistocchi, IT

Innovation in phage therapy: synergies, enzymes and engineered phages 11:14 - 11:36

Speaker: Maria del Mar Tomás Carmona, ES

Prediction of the success of phage therapy in the respiratory tract 11:36 - 12:00

Speaker: Oana Ciofu, DK*Symposium*

10:30 - 12:00

R5

S20 - Symposium 20 - Variation in the human genome is normal; get over it*Chair:* Halyna Makukh, UA*Chair:* Emmanuelle Girodon, FR

M470V and other innocent CFTR variants; let them go 10:30 - 10:52

Speaker: Karen Raraigh, US

Using populations studies to better define the impact of CFTR variants such as T5 10:52 - 11:14

Speaker: Caroline Raynal, FR

The ever-expanding CFSPID tribe, where are they going? 11:14 - 11:36

Speaker: Tanja Gonska, CA

The psychological ramification of uncertain variants 11:36 - 12:00

Speaker: Paul Weldon, GB*Industry Sessions*

12:30 - 14:00

R1

Satellite Symposium*ECFS Tomorrow Lounge Session*

12:45 - 13:45

Research topics in CF nursing : how to get started*Chair:* Espérie Burnet, FR*Chair:* Malin Heiden, DK*ECFS Tomorrow Lounge Session*

12:45 - 13:45

Tomorrow Lounge

Optimising weight management - a multidisciplinary discussion*Chair:* Gemma Stanford, GB*Chair:* Joanna Snowball, GB

ePoster Session

14:00 - 15:00

R2

EPS05 - ePoster Session 5 - Gastrointestinal and nutritional changes upon CFTR modulators*Chair:* Chris Smith, GB*Chair:* Keith Chee Y. Ooi, AU

EPS5.01	Change in gut microbiota following elexacaftor/tezacaftor/ivacaftor (ETI) therapy: preliminary analysis <i>Oral Presenter:</i> Laura Caley, GB	14:00 - 14:00
EPS5.02	Faecal microbiota changes in patients with cystic fibrosis with 6 months of elexacaftor/tezacaftor/ivacaftor: preliminary findings from the PROMISE study <i>Oral Presenter:</i> Jennifer T Duong, US	14:00 - 14:00
EPS5.03	Elexacaftor / Tezacaftor / Ivacaftor for 76 weeks is associated with a reduced oro-caecal transit time: the GIFT-CF3 extension study <i>Oral Presenter:</i> Florence Lockwood, GB	14:00 - 14:00
EPS5.04	Liver test abnormalities in adults with cystic fibrosis taking elexacaftor/tezacaftor/ivacaftor - one size doesn't fit all? <i>Oral Presenter:</i> Daniel Tewkesbury, GB	14:00 - 14:00
EPS5.05	Elexacaftor/tezacaftor/ivacaftor (ETI) - nutritional benefits are greatest for those in greatest need <i>Oral Presenter:</i> Laura Kinsey, GB	14:00 - 14:00
EPS5.06	Changes in body mass index, energy, fat and sodium intake with triple CFTR modulator therapy: preliminary analysis <i>Oral Presenter:</i> Laura Caley, GB	14:00 - 14:00
EPS5.07	Effect of elexacaftor/tezacaftor/ivacaftor on nutritional status in UK children with cystic fibrosis aged 6-11 years: a single centre service evaluation <i>Oral Presenter:</i> Elizabeth Owen, GB	14:00 - 14:00
EPS5.08	Elexacaftor/tezacafto/ ivacaftor reduces need for pancreatic enzymes, enteral feeding and omeprazole use without causing excessive weight gain: a 2-year observational study <i>Oral Presenter:</i> Naomi Scanlan, GB	14:00 - 14:00
EPS5.09	Changes in nutritional status and muscular strength in adult patients with cystic fibrosis (pwCF) treated with elexacaftor/tezacaftor/ivacaftor (ETI) <i>Oral Presenter:</i> Cecilia Brignole, IT	14:00 - 14:00
EPS5.10	Normonatremic salt depletion is not corrected in patients with cystic fibrosis treated with CFTR modulators <i>Oral Presenter:</i> Gorana Levačić, HR	14:00 - 14:00

ePoster Session

14:00 - 15:00

R2

ePoster Session 5

ePoster Session

14:00 - 15:00

R3

ePoster Session 6*ePoster Session*

14:00 - 15:00

R3

EPS06 - ePoster Session 6 - Real world studies on elexacaftor/tezacaftor/ivacaftor

Chair: Pavel Drevinec, CZ

Chair: Peter Barry, GB

EPS6.01	Effects of elexacaftor/tezacaftor/ivacaftor therapy on sputum viscoelasticity, airway infection and inflammation in patients with cystic fibrosis <i>Oral Presenter:</i> Laura Schaupp, DE	14:00 - 14:00
EPS6.02	Elexacaftor/tezacaftor/ivacaftor results in substantial improvements in chest CT scores in people with cystic fibrosis aged 12 and above over one year of treatment <i>Oral Presenter:</i> Paul McNally, IE	14:00 - 14:00
EPS6.03	Pulmonary pathogen prevalence 12 months after elexacaftor/tezacaftor/ivacaftor introduction: results from the Danish National Cystic Fibrosis Cohort <i>Oral Presenter:</i> Majbritt Jeppesen, DK	14:00 - 14:00
EPS6.04	Elexacaftor/tezacaftor/ivacaftor (ETI) reduces sputum pathogen density and lung inflammation, but infection and inflammation persist <i>Oral Presenter:</i> Samantha L Durfey, US	14:00 - 14:00
EPS6.05	A phase 3b study of the effects of elexacaftor/tezacaftor/ivacaftor (ELX/TEZ/IVA) on cough and physical activity in people with cystic fibrosis (CF) <i>Oral Presenter:</i> Eva Van Braeckel, BE	14:00 - 14:00
EPS6.06	Impact of 6 months of treatment with elexacaftor/tezacaftor/ivacaftor on clinical outcomes in children aged 6-11 with cystic fibrosis - The RECOVER study <i>Oral Presenter:</i> Paul McNally, IE	14:00 - 14:00
EPS6.07	Longitudinal impact of elexacaftor/tezacaftor/ivacaftor on abdominal symptoms assessed with the CFAbd-Score and on intestinal inflammation in people with cystic fibrosis aged \geq 12 years - The RECOVER study <i>Oral Presenter:</i> Jochen Mainz, DE	14:00 - 14:00
EPS6.08	Cystic fibrosis elexacaftor/tezacaftor/ivacaftor in liver or kidney transplanted people with cystic fibrosis using Tacrolimus, a drug-drug interaction study <i>Oral Presenter:</i> Renske van der Meer, NL	14:00 - 14:00
EPS6.09	Mid-term effect of elexacaftor/tezacaftor/ivacaftor on lung function in people with cystic fibrosis: real-world data of the German cystic fibrosis registry <i>Oral Presenter:</i> Stefanie Dillenhöfer, DE	14:00 - 14:00
EPS6.10	Evolution of psychic symptoms before and after 6 months of treatment with elexacaftor/tezacaftor/ivacaftor (ETI) in French adults patients with cystic fibrosis (pwCF)	14:00 - 14:00

Oral Presenter: Quitterie Reynaud, FR

ePoster Session

14:00 - 15:00

EPS07 - ePoster Session 7 - How much CF is it?

Chair: Philippe Reix, FR

EPS7.01	Cystic fibrosis screen positive inconclusive diagnosis (CFSPID): an Italian multicentre survey evaluating progression to definitive diagnoses <i>Oral Presenter: Cristina Fevola, IT</i>	14:00 - 14:00
EPS7.02	Cystic fibrosis Screen Positive, Inconclusive Diagnosis (CFSPID): A 15-year review	14:00 - 14:00
EPS7.03	Variability in evaluation and follow-up of newborns with CFSPID/CRMS In New York State Cystic Fibrosis Specialty Care Centres <i>Oral Presenter: Elinor Langfelder-Schwind, US</i>	14:00 - 14:00
EPS7.04	Evaluating CFSPID phenotypes and outcomes: a retrospective study from a large UK cystic fibrosis centre <i>Oral Presenter: Alison Mansfield, GB</i>	14:00 - 14:00
EPS7.05	The views of parents of children with Cystic Fibrosis Screen Positive, Inconclusive Diagnosis (CFSPID) on diagnosis, care and follow up <i>Oral Presenter: Sally Evans, GB</i>	14:00 - 14:00
EPS7.06	Hospital admissions due to pulmonary exacerbation in the first five years of life in children with cystic fibrosis in Ireland: the Irish Comparative Outcome Study of cystic fibrosis (ICOS) <i>Oral Presenter: Rini Bhatnagar, IE</i>	14:00 - 14:00
EPS7.07	Impact of newborn screening for cystic fibrosis - preliminary results from Norway <i>Oral Presenter: Magnhild Kolsgaard, NO</i>	14:00 - 14:00

ePoster Session

14:00 - 15:00

ePoster Sessions 7 - 9

ePoster Session

14:00 - 15:00

EPS08 - ePoster Session 8 - The vital role of physiotherapy in optimising airway clearance, lung function, and quality of life in the era of HEMTs

Chair: David Kinoo, BE

Chair: Lisa Morrison, GB

EPS8.01	Examining the evolution of physical activity in adults with cystic fibrosis and its relationship to lung function and maximal exercise capacity <i>Oral Presenter: Rachel Gyte, GB</i>	14:00 - 14:00
EPS8.02	Validation of the 25-level modified shuttle test in children with cystic fibrosis	14:00 - 14:00

	Oral Presenter: Jennifer Corda, AU	
EPS8.03	A pilot service conducting virtual induced sputum trials at the Blackpool Adult Cystic Fibrosis Service (BACFS) Oral Presenter: Nicole Petch, GB	14:00 - 14:00
EPS8.04	Effects of an high-intensity interval training or moderate-continuous intensity training on exercise capacity and body composition in people with cystic fibrosis - a randomized controlled study Oral Presenter: Wolfgang Gruber, DE	14:00 - 14:00
EPS8.05	One-minute sit-to-stand test improves with CFTR modulators Oral Presenter: Morgane Penelle, BE	14:00 - 14:00
EPS8.06	Prevalence of hypoglycaemia, measured by flash glucose monitoring, after moderate intensity aerobic exercise in adults with cystic fibrosis, compared to healthy controls Oral Presenter: Tiffany Dwyer, AU	14:00 - 14:00
EPS8.07	Relationship between circulating irisin levels with exercise capacity and muscle function in cystic fibrosis Oral Presenter: Kubra Kilic, TR	14:00 - 14:00
EPS8.08	Nintendo Switch™ Ring Fit Adventure™ as an alternative exercise option for people with cystic fibrosis during an inpatient admission: a pilot study of patient's perceptions Oral Presenter: Nicole Petch, GB	14:00 - 14:00
EPS8.10	Physiotherapy treatments used during pulmonary exacerbations requiring intensive therapy in Australia: data from the BEAT cystic fibrosis platform Oral Presenter: Jamie Wood, US	14:00 - 14:00

ePoster Session

14:00 - 15:00

EPS09 - ePoster Session 9 - Treatment of CF respiratory infection

Chair: Michael Hogardt, DE

Chair: Laura Sherrard, GB

EPS9.01	An <i>in vitro</i> model to predict the impact of CFTR functional restoration in the cystic fibrosis airway on <i>Pseudomonas aeruginosa</i> anti-microbial resistance, persistence and virulence Oral Presenter: John King, GB	14:00 - 14:00
EPS9.02	<i>Pseudomonas aeruginosa</i> in sputum cultures of patients with cystic fibrosis before and during one year of elexacaftor/tezacaftor/ivacaftor Oral Presenter: Dorothea Appelt, AT	14:00 - 14:00
EPS9.03	Long-term effects of Elexacaftor-Tezacaftor-Ivacaftor therapy on Cystic Fibrosis monocytes Oral Presenter: Gloria Sangiorgi, IT	14:00 - 14:00
EPS9.04	Changes in biofilm-tolerance of <i>Pseudomonas aeruginosa</i> to tobramycin following treatment with tobramycin inhalation powder (TIP)	14:00 - 14:00

	Oral Presenter: Ross P. McCleave, GB	
EPS9.05	Vantobra®, a new formulation of nebulised tobramycin: early real-world experiences at our large regional UK adult cystic fibrosis centre Oral Presenter: Catherine Brown, GB	14:00 - 14:00
EPS9.06	Development of levofloxacin resistance in clinical <i>Pseudomonas aeruginosa</i> isolates from people with cystic fibrosis in an <i>in vitro</i> variable exposure model Oral Presenter: Callum Matthew Sloan, GB	14:00 - 14:00
EPS9.07	Improving colistin activity against <i>Pseudomonas aeruginosa</i> biofilms Oral Presenter: Valentina Pastore, IT	14:00 - 14:00
EPS9.08	Antimicrobial photodynamic therapy with Ru(II)-loaded polymer nanocarriers towards treatment of bacterial lung infections: Proof-of-concept using cystic fibrosis <i>Pseudomonas aeruginosa</i> isolates Oral Presenter: Mareike Müller, DE	14:00 - 14:00
EPS9.09	Pilot study to evaluate the use of dry chlorine dioxide gas for sterilisation of virtual reality headsets Oral Presenter: Victoria Daniel, GB	14:00 - 14:00
EPS9.10	Could anti-<i>Pseudomonas aeruginosa</i> antibodies be a useful marker in monitoring the effect of ETI treatment on airways' microbiology in cystic fibrosis (CF)? Oral Presenter: Daniela Dolce, IT	14:00 - 14:00

Poster Viewing
14:00 - 15:00

PS2 - Poster Viewing 2

P033	Predictive biomarkers, and therapeutic targets, from the donor graft associated with the development of primary graft dysfunction after lung transplantation in cystic fibrosis patients Oral Presenter: Emmanuelle Brochiero, CA	14:00 - 14:00
P035	Partial correction of F508del-CFTR trafficking and stability defects by the combination of PTI-801 with ABBV-2222 or FDL-169 Oral Presenter: Miquéias Lopes-Pacheco, PT	14:00 - 14:00
P036	A comparative study of cell culture inserts for <i>in vitro</i> modeling of the cystic fibrosis airway epithelium Oral Presenter: Signe Lolle, DK	14:00 - 14:00
P037	Home spirometry engagement in a paediatric network post pandemic Oral Presenter: Philip Lawrence, GB	14:00 - 14:00
P102	Clear illustration of improved survival in cystic fibrosis using the Kaplan-Meier method: the 50-year experience of Brittany, France Oral Presenter: Virginie Scotet, FR	14:00 - 14:00
P103	Ageing with cystic fibrosis - what can we expect?	14:00 - 14:00

	<i>Oral Presenter:</i> Karuna Sapru, GB	
P105	Epidemiological characteristics of children with cystic fibrosis in Bosnia and Herzegovina <i>Oral Presenter:</i> Ganimeta Bakalovic, BA	14:00 - 14:00
P107	DeltaF508 mutations, age of diagnosis and age of death for patients in Bulgaria <i>Oral Presenter:</i> Guergana Petrova, BG	14:00 - 14:00
P108	Demographic, clinical and laboratory characteristics in cystic fibrosis population from Republic of Moldova <i>Oral Presenter:</i> Oxana Turcu, MD	14:00 - 14:00
P109	Real-life efficacy of elexacaftor/tezacaftor/ivacaftor in the Dutch cystic fibrosis population <i>Oral Presenter:</i> Dominique Zomer-van Ommen, NL	14:00 - 14:00
P110	Change in sweat chloride concentration after elexacaftor/tezacaftor/ivacaftor initiation: results from 12 months follow-up in the Danish cystic fibrosis cohort <i>Oral Presenter:</i> Thomas Bryrup, DK	14:00 - 14:00
P111	Forced oscillation techniques - is it useful for the monitoring adult patients with cystic fibrosis in the era of CFTR modulators <i>Oral Presenter:</i> Andrea Vukić Dugac, HR	14:00 - 14:00
P112	Variation in eligibility to highly effective modulator therapy between ethnic groups at our centre and how this is reflected in median lung function <i>Oral Presenter:</i> Malithi Fernando, GB	14:00 - 14:00
P113	Home reported outcomes (HERO-2) in people with cystic fibrosis taking elexacaftor/tezacaftor/ivacaftor: self-reported changes in use of chronic daily therapies at enrollment <i>Oral Presenter:</i> Cynthia Brown, US	14:00 - 14:00
P114	Predicting poor responses to elexacaftor/tezacaftor/ivacaftor: are all responses created equal? <i>Oral Presenter:</i> Matthew Craggs, GB	14:00 - 14:00
P115	The RISE study protocol: Resilience Impacted by positive Stressful Events for people with cystic fibrosis <i>Oral Presenter:</i> Sabine Elena Ineke van der Laan, NL	14:00 - 14:00
P116	Elexacaftor/tezacaftor/ivacaftor during pregnancy: the effect on maternal ppFEV₁ <i>Oral Presenter:</i> Nina Broen Jakobsen, DK	14:00 - 14:00
P117	A longitudinal study on the impact of elexacaftor/tezacaftor/ivacaftor treatment on quality of life in people with cystic fibrosis in the real world <i>Oral Presenter:</i> Sivagurunathan Sutharsan, DE	14:00 - 14:00
P118	Real-world impact of elexacaftor/tezacaftor/ivacaftor on quality of life of children with cystic fibrosis aged 6-11 years and primary caregivers in the UK: MAGNIFY, a prospective, observational, non-interventional study	14:00 - 14:00

	<i>Oral Presenter:</i> Lena P. Thia, GB	
P119	Exploring the attitudes and knowledge of the caregivers of children with cystic fibrosis and primary ciliary dyskinesia regarding COVID-19 vaccination <i>Oral Presenter:</i> Seyda Karabulut, TR	14:00 - 14:00
P120	Adherence to elexacaftor/texacaftor/ivacaftor during the first 2 years of treatment in an adolescent population <i>Oral Presenter:</i> Amanda Bevan, GB	14:00 - 14:00
P121	Vaccine coverage in people with cystic fibrosis <i>Oral Presenter:</i> Lior Tsviban, IL	14:00 - 14:00
P122	Immunization coverage and new issues of it at the age of novel CFTR modulator therapies in adults with cystic fibrosis <i>Oral Presenter:</i> Macha Tetart, FR	14:00 - 14:00
P123	Survey on medication-taking habits among patients with cystic fibrosis in a state of Argentina: a cross-sectional study <i>Oral Presenter:</i> Ezequiel Baran, AR	14:00 - 14:00
P124	Vaccination situation against COVID-19 in an adults cystic fibrosis center in Argentina <i>Oral Presenter:</i> Ezequiel Baran, AR	14:00 - 14:00
P125	Long-term economic impact using a virtual model of care in cystic fibrosis <i>Oral Presenter:</i> Paola Iacotucci, IT	14:00 - 14:00
P126	Health care resource use (HCRU) and associated costs preceding lung transplantation (LT) in Cystic Fibrosis patients <i>Oral Presenter:</i> Isabelle Durieu, FR	14:00 - 14:00
P127	Twenty years and counting: longitudinal trends in real-world primary outcomes for people with cystic fibrosis in Ireland and the evolution of a registry <i>Oral Presenter:</i> Laura Kirwan, IE	14:00 - 14:00
P128	Successful deployment of a cystic fibrosis registry solution in a low-and-middle-income country - a pilot <i>Oral Presenter:</i> Hector Gutierrez, US	14:00 - 14:00
P129	The cystic fibrosis epidemiology changing in Ukraine due to war in 2022-2023 <i>Oral Presenter:</i> Halyna Makukh, UA	14:00 - 14:00
P130	Exploring consequences of highly effective modulator therapy on patient's health pathways and cares in cystic fibrosis centers and beyond by a nationwide research programme in France (HORIZON) <i>Oral Presenter:</i> Philippe Reix, FR	14:00 - 14:00
P131	Investigation of healthcare-associated links in transmission of nontuberculous mycobacteria (HALT NTM) <i>Oral Presenter:</i> Jane E. Gross, US	14:00 - 14:00
P132	Clinical outcomes are comparable between shared care and	14:00 - 14:00

	centralised care <i>Oral Presenter:</i> Anders Lindblad, SE	
P133	Cystic fibrosis in two countries located in Northern and Eastern Europe: problems and perspectives <i>Oral Presenter:</i> Nataliya Rohovyk, SE	14:00 - 14:00
P134	Evaluation of multidisciplinary ambulatory care cost among children with cystic fibrosis: a comparative study between cystic fibrosis centers in Ireland and the United States <i>Oral Presenter:</i> Ryan Perkins, US	14:00 - 14:00
P135	Demographics profile and societal burden among persons with cystic fibrosis in the Danish population 1990 to 2018 <i>Oral Presenter:</i> Camilla Bjørn Jensen, DK	14:00 - 14:00
P136	ECFS syllabuses for the multidisciplinary team and wider field: a guide for comprehensive education <i>Oral Presenter:</i> Chris Smith, GB	14:00 - 14:00
P137	Assessment of the quality of data in the European Cystic Fibrosis Society Patient Registry <i>Oral Presenter:</i> Vibha Prasad, DE	14:00 - 14:00
P138	The impact of cystic fibrosis on periodontal disease and oral hygiene levels in adults: results from a clinical study <i>Oral Presenter:</i> Niamh Coffey, IE	14:00 - 14:00
P139	Scandinavian cystic fibrosis registry study - comparing care and clinical outcome <i>Oral Presenter:</i> Anita C. Senstad Wathne, NO	14:00 - 14:00
P140	Characterization of sleep in emerging adults with cystic fibrosis on highly effective modulator therapy <i>Oral Presenter:</i> Jane E. Gross, US	14:00 - 14:00
P141	Microbiological evaluation of an automated UV-disinfection robot on cystic fibrosis-related pathogens <i>Oral Presenter:</i> Kim Thomsen, DK	14:00 - 14:00
P142	Review of diagnostic labels for patients with unconfirmed cystic fibrosis diagnosis using registry data <i>Oral Presenter:</i> Simeon Situma Wanyama, BE	14:00 - 14:00
P143	The promise of emulating trials using registry data with an illustrative example: the effect of azithromycin on lung function <i>Oral Presenter:</i> Emily Granger, GB	14:00 - 14:00
P144	Self-medication and use of complementary and alternative medicines in cystic fibrosis - MUCAUTOMED study <i>Oral Presenter:</i> Sophie Lemagner, FR	14:00 - 14:00
P145	Cystic fibrosis in Iceland <i>Oral Presenter:</i> Helga Elíðóttir, SE	14:00 - 14:00
P146	COVID-19 pandemic in the Belgian cystic fibrosis patients compared with the international cystic fibrosis patients and the general Belgian population <i>Oral Presenter:</i> Géraldine Daneau, BE	14:00 - 14:00

P147	“What options are available for those not able to take current CFTR modulators?”: an answer hidden in plain sight? <i>Oral Presenter:</i> Tracey Daniels, GB	14:00 - 14:00
P150	Arthropathy in cystic fibrosis (protocol abstract) <i>Oral Presenter:</i> Anne Sofie Rosenborg Peretz, DK	14:00 - 14:00
P151	Pregnancies with elexacaftor/tezacaftor/ivacaftor in Argentina <i>Oral Presenter:</i> Ezequiel Baran, AR	14:00 - 14:00
P152	Development and validation of a personalised electronic patient-reported outcome measure to assess individual quality of life <i>Oral Presenter:</i> Danya Mulwijk, NL	14:00 - 14:00
P153	An Italian centre experience with elexacaftor/tezacaftor/ivacaftor therapy in 6 to 11 year olds with cystic fibrosis <i>Oral Presenter:</i> Chiara Rosazza, IT	14:00 - 14:00
P154	Cluster analysis explains heterogeneity in treatment response to elexacaftor/tezacaftor/ivacaftor in people with cystic fibrosis <i>Oral Presenter:</i> Chiara Premuda, IT	14:00 - 14:00
P155	The impact of elexacaftor/tezacaftor/ivacaftor (ETI) therapy on occurrence of bacterial lung infections in cystic fibrosis patients <i>Oral Presenter:</i> Maria Francesca Liporace, IT	14:00 - 14:00
P156	Treatment with the CFTR modulator [HB1] elexacaftor/tezacaftor/ivacaftor reduces immunological response to airway <i>Aspergillus</i> in people with cystic fibrosis <i>Oral Presenter:</i> Natalie Francis, GB	14:00 - 14:00
P157	Severe cystic fibrosis in the CFTR modulator era still exists! The implementation of a novel ‘high risk’ multidisciplinary team approach <i>Oral Presenter:</i> Thomas Tobin, GB	14:00 - 14:00
P158	Elexacaftor, tezacaftor, Ivacaftor (ETI) improves lung function but not rate of decline in a two year observational study of adolescents <i>Oral Presenter:</i> Garry Connett, GB	14:00 - 14:00
P159	Modulator`s effectiveness evaluated by lung ultrasound <i>Oral Presenter:</i> Ioana Mihaela Ciuca, RO	14:00 - 14:00
P160	Effects of elexacaftor/tezacaftor/ivacaftor triple combination therapy on glycaemic control and body composition in patients with cysticfibrosis related diabetes <i>Oral Presenter:</i> Laura Zazzeron, IT	14:00 - 14:00
P161	Desensitization to highly effective modulator therapy - international approach and call for collaboration	14:00 - 14:00

	<i>Oral Presenter:</i> Elizabeth Autry, US	
P162	Elexacaftor/tezacaftor/ivacaftor significantly improves thoracic radiological outcomes in cystic fibrosis <i>Oral Presenter:</i> Nicola Robinson, GB	14:00 - 14:00
P163	Impact of elexacaftor/tezacaftor/ivacaftor modulator therapy on cystic fibrosis chest radiographs using the Brasfield score, a single centre experience <i>Oral Presenter:</i> Mohammed Okour, GB	14:00 - 14:00
P164	Elexacaftor/tezacaftor/ivacaftor (ETI)- home spirometry during the first 14 days of treatment <i>Oral Presenter:</i> Matthieu Thimmesch, BE	14:00 - 14:00
P165	Impact of CFTR modulators on systemic inflammation	14:00 - 14:00
	<i>Oral Presenter:</i> Ivan Bambir, HR	
P166	Losses in the gains of children with cystic fibrosis who had to interrupt their modulator therapies <i>Oral Presenter:</i> Burcu Capraz, TR	14:00 - 14:00
P167	Antibiotic courses in children and adolescents with cystic fibrosis: after only a year of elexacaftor/tezacaftoriIvacaftor, there's a "before and after" <i>Oral Presenter:</i> Joana Quaresma Vázquez, ES	14:00 - 14:00
P168	Determining the relationship between vitality and C-Reactive protein in those initiating elexacaftor/tezacaftor/ivacaftor <i>Oral Presenter:</i> Jacob Gravelle, CA	14:00 - 14:00
P169	Vitamin absorption after the introduction of elexacaftor/tezacaftor/ivacaftor in children and adolescents diagnosed with cystic fibrosis <i>Oral Presenter:</i> Ana Morales Tirado, ES	14:00 - 14:00
P170	Trikafta® modulates release of extracellular vesicles in cystic fibrosis <i>Oral Presenter:</i> navya lakkappa, IE	14:00 - 14:00
P171	Clinical experience with the introduction of a programme with CFTR modulators in a paediatric cystic fibrosis centre in Poland - a preliminary report	14:00 - 14:00
	<i>Oral Presenter:</i> Katarzyna Walicka-Serzysko,	
P172	CFTR Modulators in children with cystic fibrosis: real-life evidence in Turkey <i>Oral Presenter:</i> Ismail Guzelkas, TR	14:00 - 14:00
P173	Modulatory therapy experience in patients with cystic fibrosis in Turkey: a multi-centre study <i>Oral Presenter:</i> Aynur Gulieva, TR	14:00 - 14:00
P174	Human epididymis protein 4 (HE4) plasma concentrations	14:00 - 14:00

	correlate with the improvement of ppFEV1 in response to LUM/IVA therapy in people with cystic fibrosis homozygous for p.Phe508del-CFTR	
	<i>Oral Presenter:</i> Marianna Pócsi, HU	
P175	Changes in breathprint after start of tezacaftor/ivacaftor therapy: an eNose pilot study	14:00 - 14:00
	<i>Oral Presenter:</i> Alain P. Iradukunda, NL	
P176	Implantable venous access devices in the era of Trikafta®-time for a rethink?	14:00 - 14:00
	<i>Oral Presenter:</i> Sheila Sivam, AU	
P177	Cystic fibrosis (CF)-related complications and outcomes of women during pregnancy and post-partum in elexacaftor/tezacaftor/ivacaftor (ETI)-era	14:00 - 14:00
	<i>Oral Presenter:</i> Amy Downes, GB	
P178	Impact of pre-conception lung function in women with cystic fibrosis (wwCF) on obstetric and neonatal outcomes in the elexacaftor/tezacaftor/ivacaftor (ETI)-era	14:00 - 14:00
	<i>Oral Presenter:</i> Amy Downes, GB	
P179	Maternal and foetal outcomes of multigravida cystic fibrosis patients	14:00 - 14:00
	<i>Oral Presenter:</i> Malena Cohen-Cymberknoh, IL	
P180	Evaluation of clinical features of children with cystic fibrosis and CFSPID in newborn screening programme with IRT/IRT protocol	14:00 - 14:00
	<i>Oral Presenter:</i> Tugba Sismanlar Eyuboglu, TR	
P181	Polygenic risk score and socioenvironmental factors predict early-onset lung disease in young children with cystic fibrosis	14:00 - 14:00
	<i>Oral Presenter:</i> Leslie Huang, US	
P182	Prevalence of tracheobronchomalacia is higher than previously reported in children with cystic fibrosis	14:00 - 14:00
	<i>Oral Presenter:</i> Julie Depiazzi, AU	
P184	Sex differences in annual pulmonary exacerbations in people with cystic fibrosis	14:00 - 14:00
	<i>Oral Presenter:</i> Kristina Montemayor, US	
P185	Prospective randomized observational study validating biomarkers for association with future pulmonary exacerbations in people with cystic fibrosis	14:00 - 14:00
	<i>Oral Presenter:</i> Theodore G Liou, US	
P186	Airway clearance in the STOP PEDS pilot study of oral antibiotic treatment strategies for paediatric cystic fibrosis pulmonary exacerbations	14:00 - 14:00
	<i>Oral Presenter:</i> Don Sanders, US	
P187	Small airways disease in cystic fibrosis patients	14:00 - 14:00
	<i>Oral Presenter:</i> Michela Deolmi, IT	
P188	Heterogeneity of large and small airway remodeling in human end-stage explant cystic fibrosis lungs	14:00 - 14:00

	<i>Oral Presenter:</i> Astrid Vermaut, BE	
P190	Nanostructured formulation of a novel hybrid iminosugar/steroid agent for application in cystic fibrosis lung disease <i>Oral Presenter:</i> Anna Esposito, IT	14:00 - 14:00
P191	Is abnormal glucose tolerance driving lung inflammation in cystic fibrosis? <i>Oral Presenter:</i> Stefanie Diemer, SE	14:00 - 14:00
P192	Change in lung clearance index with microbiological status in patients with cystic fibrosis <i>Oral Presenter:</i> Elpis Hatziaorou, GR	14:00 - 14:00
P193	Lung clearance index in asymptomatic CRMS/CFSPID infants progressed to a diagnosis of cystic fibrosis for pathological sweat test: a monocentric prospective experience <i>Oral Presenter:</i> Cristina Fevola, IT	14:00 - 14:00
P194	Computed cardiopulmonography (CCP) and the idealised lung clearance index (iLCI) in early-stage cystic fibrosis <i>Oral Presenter:</i> Christopher Short, GB	14:00 - 14:00
P195	Prospective longitudinal evaluation of the lung clearance index (LCI) in the clinical setting <i>Oral Presenter:</i> Jacqueline Spano, US	14:00 - 14:00
P196	Use of nose clips for routine spirometry helps maximise forced expiratory outcome measures in cystic fibrosis clinics <i>Oral Presenter:</i> Garry Connett, GB	14:00 - 14:00
P197	A pilot study to determine whether normative values of specialised pulmonary function tests differ in children from non-Caucasian, ethnic minority backgrounds <i>Oral Presenter:</i> Mary Abkir, GB	14:00 - 14:00
P198	Home spirometry as a clinical trial endpoint: qualitative needs assessment and co-production of training materials <i>Oral Presenter:</i> Margaret Rosenfeld, US	14:00 - 14:00
P199	Non-invasive monitoring of cystic fibrosis lung disease in a new era: the TERRIFIC-MILE study <i>Oral Presenter:</i> Hettie Janssens, NL	14:00 - 14:00
P207	CFHealthHub allows clinicians to identify people with long nebuliser durations and intervene to reduce duration <i>Oral Presenter:</i> Robert D Sandler, GB	14:00 - 14:00
P209	Personalised data-Linkage Understanding Treatment Optimisation (PLUTO) in the CFHealthHub Learning Health System: understanding how much is enough for normal life expectancy in the post-modulator era <i>Oral Presenter:</i> Robert D Sandler, GB	14:00 - 14:00
P210	Environmental impact of a cystic fibrosis virtual clinic <i>Oral Presenter:</i> David Green, GB	14:00 - 14:00
P211	The impact of multidisciplinary virtual cystic fibrosis clinics	14:00 - 14:00

	on patient experience, quality of life, costs and time commitments	
	<i>Oral Presenter:</i> Jody Bell, AU	
P212	Clinical and transcriptomic features of COVID-19 in cystic fibrosis: a prospective multi-centre study	14:00 - 14:00
	<i>Oral Presenter:</i> Fabiana Ciciriello, IT	
P214	Clinical course of SARS-CoV-2 infections of paediatric patients with cystic fibrosis- a single retrospective centre experience	14:00 - 14:00
	<i>Oral Presenter:</i> Justyna Sieber, AT	
P215	Intravenous antibiotic administration in children and adolescents with cystic fibrosis and catheter-associated complications: a 5-year experience	14:00 - 14:00
	<i>Oral Presenter:</i> Mónica López Rozas, ES	
P216	The impact of the COVID-19 pandemic and elexacaftor/tezacaftor/ivacaftor initiation on high-cost medication utilisation in adults living with cystic fibrosis	14:00 - 14:00
	<i>Oral Presenter:</i> Alex Chan, GB	
P217	Arthritis in cystic fibrosis - case series and review	14:00 - 14:00
	<i>Oral Presenter:</i> Anne Sofie Rosenborg Peretz, DK	
P219	Otorhinolaryngologic, audiological and genetic findings in children with cystic fibrosis: a tertiary care experience	14:00 - 14:00
	<i>Oral Presenter:</i> Raziye Atan, TR	
P220	The CFHealthHub Learning Health System - supporting a community of practice to deliver a normal life expectancy in cystic fibrosis	14:00 - 14:00
	<i>Oral Presenter:</i> Robert D Sandler, GB	
P221	Identifying knowledge gaps by using adapted cystic fibrosis R.I.S.E. in a low-resource setting	14:00 - 14:00
	<i>Oral Presenter:</i> Seyda Karabulut, TR	
P222	Multi-central profile of refugee cystic fibrosis patients in Turkey	14:00 - 14:00
P223	Transition programme as a quality improvement project: adaptation of the R.I.S.E. in a centre with limited resources - Marmara University experience	14:00 - 14:00
	<i>Oral Presenter:</i> Seyda Karabulut, TR	
P224	Eligibility of CFTR Modulator [HB1] drugs in patients registered in cystic fibrosis registry of Turkey	14:00 - 14:00
	<i>Oral Presenter:</i> Meltem Akgül Erdal, TR	
P226	Interpretation of spirometry parameters in adult patients with cystic fibrosis in North Macedonia	14:00 - 14:00
	<i>Oral Presenter:</i> Sonja Momchilovikj,	
P345	The clinical impact of reduced dose prescribing of	14:00 - 14:00

	elexacaftor/tezacaftor/ivacaftor (ETI) in children with cystic fibrosis <i>Oral Presenter:</i> Amanda Thomsen, AU	
P346	The Alfred Wellness Score (AWEScore®): measurements of quality of life before and after the introduction of elexacaftor/tezacaftor/ivacaftor (Trikafta®) in adults with cystic fibrosis <i>Oral Presenter:</i> Brenda Button, AU	14:00 - 14:00
P347	Mental health after initiating triple CFTR modulators in a Polish paediatric cystic fibrosis centre - a preliminary report <i>Oral Presenter:</i> Urszula Borawska-Kowalczyk, PL	14:00 - 14:00
P348	The impact of CFTR modulators on the quality of life of adult patients with cystic fibrosis in Croatia <i>Oral Presenter:</i> Tihana Odobasic Palkovic,	14:00 - 14:00
P349	Body image perceptions and elexacaftor/tezacaftor/ivacaftor (ETI) use in adolescents living with cystic fibrosis in the United States <i>Oral Presenter:</i> Emily Muther, US	14:00 - 14:00
P350	Impact of one year of treatment with elexacaftor/tezacaftor/ivacaftor on clinical outcomes in adult patients with cystic fibrosis - first experience from Croatia <i>Oral Presenter:</i> Ivana Lalić Čičković, HR	14:00 - 14:00
P351	Adherence to chronic treatment by adult patients with cystic fibrosis in the era of CFTR modulators <i>Oral Presenter:</i> Ivana Lalić Čičković, HR	14:00 - 14:00
P352	New therapy, new adherence idea? <i>Oral Presenter:</i> Paola Catastini, IT	14:00 - 14:00
P353	Impact of the withdrawal of CFTR modulator prescriptions on clinic attendance <i>Oral Presenter:</i> Michael Kevin Dooney, GB	14:00 - 14:00
P354	West Midlands Adult Cystic Fibrosis Centre experience of the effects of Kaftrio® on patients within the lung transplant programme <i>Oral Presenter:</i> Josephine Hussey, GB	14:00 - 14:00
P355	A qualitative study exploring parent's experience of the diagnosis of cystic fibrosis for their newborn baby <i>Oral Presenter:</i> Mark Guyers, GB	14:00 - 14:00

P356	A qualitative study on awareness, attitudes, behaviors and social adaptation of mothers of children with cystic fibrosis <i>Oral Presenter:</i> İlknur Ayvaz, TR	14:00 - 14:00
P357	Psychological study of the relationship between pain perception and fear, anxiety and quality of life in children with cystic fibrosis <i>Oral Presenter:</i> Géraldine Labouret, FR	14:00 - 14:00
P358	Anxiety and depression levels reported by parents of children with cystic fibrosis over a 6-month period <i>Oral Presenter:</i> Claire Edmondson, GB	14:00 - 14:00
P359	“No words for feelings” The factors effecting alexithymia in the patients with cystic fibrosis and their mothers <i>Oral Presenter:</i> Ayse Tana Aslan, TR	14:00 - 14:00
P360	Mental health in cystic fibrosis patients: predictive factors and psychopathology <i>Oral Presenter:</i> Carlo Castellani, IT	14:00 - 14:00
P361	Psychological and physical impact of coughing in patients with cystic fibrosis (CF) <i>Oral Presenter:</i> Ivana Arnaudova Danevska,	14:00 - 14:00
P362	Employment and life choices for PwCF in Wales - 10 years on <i>Oral Presenter:</i> Vivien S. Edwards, GB	14:00 - 14:00
P363	Socio-economic choices in the cystic fibrosis family with a child with cystic fibrosis in primary school <i>Oral Presenter:</i> Karin Risager Jakobsen, DK	14:00 - 14:00
P364	Financial and logical impacts of review appointments on parents of children with cystic fibrosis <i>Oral Presenter:</i> Anirban Maitra, GB	14:00 - 14:00
P365	‘Shouldn’t you be thinking about work now?’- employment status and influencing clinical factors in people with cystic fibrosis at a large UK adult centre <i>Oral Presenter:</i> Jacqui Wainwright, GB	14:00 - 14:00
P366	Developing an in-house and integrated sweat testing service for the routine management of adults with cystic fibrosis <i>Oral Presenter:</i> Lindsey Gillgrass, GB	14:00 - 14:00
P367	Antibiotic treatment of <i>Staphylococcus aureus</i> in small children with newly diagnosed cystic fibrosis - a nursing care experience in the Nordic countries <i>Oral Presenter:</i> Ellen Julie Hunstad, NO	14:00 - 14:00
P368	Antibiotic prescribing practice of the cystic fibrosis (CF) nursing team at a large tertiary paediatric centre <i>Oral Presenter:</i> Lauren Bartlett, GB	14:00 - 14:00
P369	Electronic prescribing: A service improvement project <i>Oral Presenter:</i> Claire Fagan, GB	14:00 - 14:00
P370	Exploring the barriers and facilitators to obtaining supplies of medicines for people with cystic fibrosis (PwCF) under	14:00 - 14:00

	the care of the All Wales Adult Cystic Fibrosis Centre (AWACFC) <i>Oral Presenter:</i> Mari Lea-Davies, GB	
P371	Art therapy to facilitate interaction among children with cystic fibrosis <i>Oral Presenter:</i> Claire Fagan, GB	14:00 - 14:00
P372	"The outcome would not have been the same" - multidisciplinary team experience of the role of occupational therapy in a Cystic Fibrosis unit <i>Oral Presenter:</i> Wendy Foo, GB	14:00 - 14:00
P373	Teamwork makes the dream work: joint working between social work and occupational therapy in adult cystic fibrosis care <i>Oral Presenter:</i> Wendy Foo, GB	14:00 - 14:00
P374	The role of a specialist youth worker in cystic fibrosis care in Wales <i>Oral Presenter:</i> Bethan Watkins, GB	14:00 - 14:00
P375	Brighter Futures with Study Buddies: an educational programme, delivered by the cystic fibrosis-MDT of tomorrow, to enable improved health literacy and essential skills for life and employability in the young PwCF <i>Oral Presenter:</i> Jacqueline Rendall, GB	14:00 - 14:00
P376	Co-developing health and lifestyle education for adolescents with cystic fibrosis <i>Oral Presenter:</i> Sally Ann Harris, GB	14:00 - 14:00
P377	Addressing the many steps to a successful transition! <i>Oral Presenter:</i> Karoline Prinz, AT	14:00 - 14:00
P378	The value of a community of practice, within the CFHealthHub learning health system <i>Oral Presenter:</i> Tracey Daniels, GB	14:00 - 14:00
P379	Impact of a mental health navigator for cystic fibrosis care in British Columbia <i>Oral Presenter:</i> Callie Waters, CA	14:00 - 14:00
P380	One stop annual review process <i>Oral Presenter:</i> Katie Baker-Wardle, GB	14:00 - 14:00
P382	Cystic fibrosis and disordered eating behaviour: a collaborative approach to improve care <i>Oral Presenter:</i> Rachel Massey-Chase, GB	14:00 - 14:00
P383	"It was all tailored around me": qualitative evaluation of Clinical Effort Against Smoke Exposure in cystic fibrosis (CEASE-CF) <i>Oral Presenter:</i> Gabriela Oates, US	14:00 - 14:00
P384	Smoking prevalence in cystic fibrosis families is higher than in the Greek general population <i>Oral Presenter:</i> Argyri Petrocheilou, GR	14:00 - 14:00
P385	Measuring what matters to patients	14:00 - 14:00

	<i>Oral Presenter:</i> Martina Kapatou, GB	
P386	Improving support for sexual and reproductive health in adult patients with cystic fibrosis <i>Oral Presenter:</i> Elizabeth C Benson, GB	14:00 - 14:00
P387	Reviews and outcomes from a new pregnancy support service at a large cystic fibrosis centre <i>Oral Presenter:</i> Beverly Govin, GB	14:00 - 14:00
P388	Three new mums, 1 big realisation: a maternity leave reflection on cystic fibrosis care postpartum <i>Oral Presenter:</i> Lucy Wadsworth, GB	14:00 - 14:00
P389	Evolving gender identification within the cystic fibrosis population and considerations on potential effects on lung function results <i>Oral Presenter:</i> Alice Darby, GB	14:00 - 14:00
P390	Ageing with cystic fibrosis: challenges for patients and team members <i>Oral Presenter:</i> Sue Braun, BE	14:00 - 14:00
P391	"I'm different": reviewing outpatient psychology support for people of colour in an inner London adult cystic fibrosis centre <i>Oral Presenter:</i> Ghiselle Green, GB	14:00 - 14:00
P392	Medical and mental status of refugees with cystic fibrosis from Ukraine <i>Oral Presenter:</i> Carsten Schwarz, DE	14:00 - 14:00
P393	Physical activity participation and mental health status of patients with cystic fibrosis during the COVID-19 pandemic: a single centre experience <i>Oral Presenter:</i> Silvia Delgado, US	14:00 - 14:00
P394	Comparisons of the personal well-being score (PWS) and health-related quality of life for people with cystic fibrosis before and during the COVID-19 pandemic <i>Oral Presenter:</i> Rana Altabee, GB	14:00 - 14:00
P395	Completeness of annual review investigations as per national recommendations after the COVID-19 pandemic - experience from a large tertiary centre in the northwest United Kingdom <i>Oral Presenter:</i> Anirban Maitra, GB	14:00 - 14:00
P396	Moral distress and burnout in healthcare workers during the COVID-19 pandemic: quantitative results from a large Australian public hospital survey <i>Oral Presenter:</i> Felicity Finlayson, AU	14:00 - 14:00
P397	Using the science of health care improvement to implement a virtual cystic fibrosis clinic alongside the traditional face-to-face model post-COVID pandemic <i>Oral Presenter:</i> Julie Knowles, GB	14:00 - 14:00
P398	Using the science of health care improvement to implement a virtual cystic fibrosis service post COVID-19 pandemic	14:00 - 14:00

	<i>Oral Presenter:</i> Julie Knowles, GB	
P399	Using a digital learning health system to reduce medicines waste in cystic fibrosis - the Easy Medicines for Burden Reduction And Care Enhancement (EMBRACE) study <i>Oral Presenter:</i> Sarah Cameron, GB	14:00 - 14:00
P400	Virtual appointments: a barrier to engagement or improving person-centred care? <i>Oral Presenter:</i> Lisa Morrison, GB	14:00 - 14:00
P401	Video call fatigue - how do patients feel? <i>Oral Presenter:</i> Fiona Moore, GB	14:00 - 14:00
P402	Initiating a home blood monitoring service for adults with cystic fibrosis <i>Oral Presenter:</i> Katie Baker-Wardle, GB	14:00 - 14:00
P403	Hospital care at home: experiences of Ipswich Hospital's Multi-disciplinary (MDT) approach to holistic and individualised cystic fibrosis care <i>Oral Presenter:</i> Karen Richards, GB	14:00 - 14:00
P404	Does knowledge of cystic fibrosis affect adherence to home monitoring? results from the CLIMB-CF study <i>Oral Presenter:</i> Claire Edmondson, GB	14:00 - 14:00
P405	Adherence in the use of digital technology for home monitoring in patients with cystic fibrosis <i>Oral Presenter:</i> Daniela Savi, IT	14:00 - 14:00
<i>Workshop</i>		
15:00 - 16:30		R1
WS10 - WS10 - Changes in the clinical landscape in the era of CFTR modulators		
<i>Chair:</i> Silke van Koningsbruggen-Rietschel, DE		
<i>Chair:</i> Lieven Dupont, BE		
WS10.01	The French Compassionate Programme of elexacaftor/tezacaftor/ivacaftor in people with cystic fibrosis with advanced lung disease and no F508del CFTR variant <i>Oral Presenter:</i> Pierre-Régis Burgel, FR	15:00 - 15:15
WS10.02	Pharmacological effects of CFTR-modulation in cystic fibrosis patients after lung transplantation: interim results of the multicenter KOALA study <i>Oral Presenter:</i> Carina M E Hansen, NL	15:15 - 15:30
WS10.03	Increasing cardiovascular risk in adults with cystic fibrosis related diabetes receiving CFTR modulator therapy (elexacaftor/tezacaftor/ivacaftor) <i>Oral Presenter:</i> Alex Chan, GB	15:30 - 15:45
WS10.04	Analyzes of worldwide inhaled antibiotic prescriptions for <i>Pseudomonas. aeruginosa</i>	15:45 - 16:00

Oral Presenter: Marianne S. Muhlebach, US

- WS10.05 **Reduction of pulmonary exacerbations in people with cystic fibrosis in Germany between 2019 and 2022** 16:00 - 16:15
Oral Presenter: Susanne Naehrig, DE

- WS10.06 **How representative are clinical trial cohorts of the general cystic fibrosis population? Implications for trial planning** 16:15 - 16:30
Oral Presenter: Rebecca Dobra, GB

Workshop

15:00 - 16:30

R2

WS11 - WS11 - Rising to the challenge: navigating mental health and adherence in the era of CFTR modulators

Chair: Edwina Landau, IL

Chair: Karoline Prinz, AT

- WS11.01 **Challenging behaviours and mood changes in a large cohort of 6-11 year old children following elexacaftor/tezacaftor/ivacaftor initiation** 15:00 - 15:15
Oral Presenter: Tim Lee, GB

- WS11.02 **Psychological wellbeing post-CFTR modulator therapy** 15:15 - 15:30
Oral Presenter: Helen Egan, GB

- WS11.03 **Positive and negative impacts of elexacaftor/tezacaftor/ivacaftor: healthcare providers' observations across US Centres** 15:30 - 15:45
Oral Presenter: Alexandra Quittner, US

- WS11.04 **Measuring barriers to adherence among people with cystic fibrosis starting treatment with elexacaftor/tezacaftor/ivacaftor: results from the RECOVER study** 15:45 - 16:00
Oral Presenter: Sharon Sutton, IE

- WS11.05 **Medication burden among adults with cystic fibrosis prescribed CFTR Modulators: a cross-sectional survey** 16:00 - 16:15

Oral Presenter: Nicola J. Shaw, GB

- WS11.06 **What to do when your patient "ghosts" you: an ethical framework for the new era of CFTR modulators** 16:15 - 16:30
Oral Presenter: Mark Chilvers, CA

Workshop

15:00 - 16:30

R3

WS12 - WS12 - Nutrition: changes in practice

Chair: Olaf Sommerburg, DE

Chair: Dee Shimmin, GB

- WS12.01 **Early growth in Danish children with cystic fibrosis since 2000** 15:00 - 15:15
Oral Presenter: Karlen Bader-Larsen, DK

- WS12.02 **Erythrocyte essential fatty acid status and its association** 15:15 - 15:30

	with breastfeeding, growth and early-onset lung disease in young children with cystic fibrosis <i>Oral Presenter:</i> HuiChuan Lai, US	
WS12.03	Impact of elexacaftor/tezacaftor/ivacaftor on fat-soluble vitamin levels in children with cystic fibrosis <i>Oral Presenter:</i> Laura Schembri, GB	15:30 - 15:45
WS12.04	An “expanding area” of interest- the effect of elexacaftor/tezacaftor/ivacaftor (ETI) therapy on body image and weight <i>Oral Presenter:</i> Alice Darby, GB	15:45 - 16:00
WS12.05	Is nutritional status still an important contributor to lung function in modern day cystic fibrosis? <i>Oral Presenter:</i> Tamarah Katz, AU	16:00 - 16:15
WS12.06	The changing landscape of tube feeding in the post modulator era <i>Oral Presenter:</i> Liz May, GB	16:15 - 16:30
<i>Workshop</i>		
15:00 - 16:30		R4
	WS13 - WS13 - Sampling and characterisation of the airway microbiome <i>Chair:</i> Valerie Waters, CA <i>Chair:</i> Michael Tunney, GB	
WS13.01	The cystic fibrosis Home Sputum-Induction Trial (CF-HomeSpIT - ISRCTN86523335) to evaluate home sputum-induction and early morning saliva sampling in children with cystic fibrosis <i>Oral Presenter:</i> Julian Forton, GB	15:00 - 15:15
WS13.02	Comparison of qPCR and 16S rRNA marker-gene Next-Generation Sequencing (NGS) for the detection and quantification of respiratory pathogens in the sputum of people with cystic fibrosis (PwCF). <i>Oral Presenter:</i> Gisli Einarsson, GB	15:15 - 15:30
WS13.03	Microbiology of upper and lower airways of cystic fibrosis (CF) patients in stable conditions and in lung transplant patients <i>Oral Presenter:</i> Daniela Dolce, IT	15:30 - 15:45
WS13.04	Shotgun metagenomic for cystic fibrosis gut-lung microbiome and antibiotic resistant genes characterisation <i>Oral Presenter:</i> Natalia Bastón-Paz, ES	15:45 - 16:00
WS13.05	The short- and long-term effects of antibiotic treatment on the oropharyngeal and fecal microbiota in infants with cystic fibrosis <i>Oral Presenter:</i> Lucas Hoffman, US	16:00 - 16:15
WS13.06	Regional bronchoscopy sampling indicates that <i>Pseudomonas aeruginosa</i> infections persist throughout the lung after elexacaftor/tezacaftor/ivacaftor (ETI) due to intra-lung bacterial migration <i>Oral Presenter:</i> Samantha L Durfey, US	16:15 - 16:30

Workshop

15:00 - 16:30

R5

WS14 - WS14 - Airway epithelial homeostasis in CF: what and how to address it

Chair: Nicoletta Pedemonte, IT

Chair: Shafagh Waters, AU

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|---------|--|---------------|
| WS14.01 | Ex vivo whole lung model of cystic fibrosis (CF) for therapeutic screening
<i>Oral Presenter:</i> Meghan Pinezich, US | 15:00 - 15:15 |
| WS14.02 | Cystic fibrosis personalised medicine; paediatric <i>in vitro</i> airway cell models to predict CFTR modulator patient outcomes
<i>Oral Presenter:</i> Laura K Fawcett, AU | 15:15 - 15:30 |
| WS14.03 | Combined therapeutic strategies for the restoration of airway epithelial integrity and function in cystic fibrosis
<i>Oral Presenter:</i> Emmanuelle Brochiero, CA | 15:30 - 15:45 |
| WS14.04 | <i>Pseudomonas aeruginosa</i> infection drives complex host responses in a cystic fibrosis-derived airway model
<i>Oral Presenter:</i> Claudia A Colque, DK | 15:45 - 16:00 |
| WS14.05 | Effect of an agro-based compound (A-bC) on remodelling and regeneration of airway epithelium in cystic fibrosis
<i>Oral Presenter:</i> Damien Adam, CA | 16:00 - 16:15 |
| WS14.06 | Single cell RNA sequencing of lung biopsies reveals altered immune-structural cell crosstalk in cystic fibrosis
<i>Oral Presenter:</i> Ralph Stadhouders, NL | 16:15 - 16:30 |

Workshop

17:00 - 18:30

R1

WS15 - WS15 - Late Breaking Science

Chair: Dorota Sands, PL

Chair: Marcus Mall, DE

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|---------|---|---------------|
| WS15.01 | SP-101 gene therapy restores CFTR function in human CF airway epithelial cultures and drives hCFTRAR transgene expression in the airways of CF and non-CF ferrets
<i>Oral Presenter:</i> Katherine Excoffon, US | 17:00 - 17:14 |
| WS15.02 | A phase 1b/2a randomized, double-blind, placebo-controlled, multicenter study evaluating nebulized phage therapy in cystic fibrosis subjects with chronic <i>Pseudomonas aeruginosa</i> pulmonary infection
<i>Oral Presenter:</i> Urania Rappo, US | 17:14 - 17:28 |
| WS15.03 | Long-term safety and efficacy of elexacaftor/tezacaftor/ivacaftor (ELX/TEZ/IVA) in people with cystic fibrosis (CF) and at least one F508del allele: an open-label, 192-week extension study
<i>Oral Presenter:</i> Deepika Polineni, US | 17:28 - 17:42 |
| WS15.04 | LONGITUDE: An observational study of the long-term effectiveness of ivacaftor/tezacaftor/elexacaftor in people with cystic fibrosis using data from the United Kingdom Cystic Fibrosis Registry
<i>Oral Presenter:</i> Gabriela Vega-Hernandez, GB | 17:42 - 17:56 |

WS15.05	Effects of elexacaftor/tezacaftor/ivacaftor (ELX/TEZ/IVA) treatment on markers of inflammation in people with cystic fibrosis (CF) <i>Oral Presenter:</i> Siobhan O'Brien, US	17:56 - 18:10
WS15.06	Mutation class dependent signatures outweigh disease associated processes in cystic fibrosis cells <i>Oral Presenter:</i> Lucia Santos, PT	18:10 - 18:30
<i>Workshop</i> 17:00 - 18:30		R2
WS16 - WS16 - Clinical effectiveness of CFTR modulators: data from registries <i>Chair:</i> Jane Davies, GB <i>Chair:</i> Andreas Jung, CH		
WS16.01	Highly-effective CFTR modulator therapy with elexacaftor/tezacaftor/ivacftor (ETI) leads to age-dependent changes of typical cystic fibrosis hallmark pathogens - data from the German cystic fibrosis registry <i>Oral Presenter:</i> Anna-Maria Dittrich, DE	17:00 - 17:15
WS16.02	Real-world (RW) clinical effectiveness of elexacaftor/tezacaftor/ivacaftor (ELX/TEZ/IVA) in children with cystic fibrosis aged 6-11 years: interim results from the HELIO study <i>Oral Presenter:</i> Lisa J. McGarry, US	17:15 - 17:30
WS16.03	Registry-based study of people with cystic fibrosis (pwCF) treated with elexacaftor/tezacaftor/ivacaftor (ELX/TEZ/IVA): Up to 2 years of real-world outcome data <i>Oral Presenter:</i> Julie K. Bower, US	17:30 - 17:45
WS16.04	Benefits of lumacaftor/ivacaftor (LUM/IVA) initiation in children with cystic fibrosis aged 2 through 5 years: Interim results from an ongoing registry-based study <i>Oral Presenter:</i> Claire Kim, US	17:45 - 18:00
WS16.05	Long-term impact of ivacaftor (IVA) in people with cystic fibrosis in Ireland <i>Oral Presenter:</i> Barry Plant, IE	18:00 - 18:15
WS16.06	International disparities in access to highly effective modulator therapies <i>Oral Presenter:</i> Jonathan Guo, GB	18:15 - 18:30
<i>Workshop</i> 17:00 - 18:30		R3
WS17 - WS17 - Strategies to replace or edit the genetic message in CF <i>Chair:</i> Anna Cereseto, IT <i>Chair:</i> Garry Cutting, US		
WS17.01	Novel approaches based on sequence-specific RNA editing by ADARs to correct CFTR nonsense mutations causing cystic fibrosis <i>Oral Presenter:</i> Viviana Barra, IT	17:00 - 17:15
WS17.02	Improved adenine base editing approach to correct W1282X-CFTR	17:15 - 17:30

	<i>Oral Presenter:</i> Carlos M Farinha, PT	
WS17.03	An alternative mutation agnostic therapy for cystic fibrosis with oligonucleotide antisense <i>Oral Presenter:</i> Christie Mitri, FR	17:30 - 17:45
WS17.04	Developing a non-viral gene therapy strategy for treating lung cystic fibrosis disease <i>Oral Presenter:</i> Bei Qiu, IE	17:45 - 18:00
WS17.05	LUNAR[®]-CF mRNA replacement therapy restores CFTR expression and function in human bronchial epithelial cells <i>Oral Presenter:</i> Javier Campos-Gomez, US	18:00 - 18:15
WS17.06	F/HN pseudotyped lentiviral vector-mediated transduction of non-human primates <i>Oral Presenter:</i> Uta Griesenbach, GB	18:15 - 18:30
 <i>Workshop</i> 17:00 - 18:30		
	WS18 - WS18 - Modifying inflammation in the CF airways	R4
	<i>Chair:</i> Isabelle Fajac, FR <i>Chair:</i> Hettie Janssens, NL	
WS18.01	Impact of elexacaftor/tezacaftor/ivacaftor therapy on sputum metabolomics in adult cystic fibrosis <i>Oral Presenter:</i> Susan Kim, US	17:00 - 17:15
WS18.02	ETI triple therapy shows sustained, progressive normalisation of airway cytokine and antiprotease balance and systemic inflammation over one year of treatment <i>Oral Presenter:</i> Mark P Murphy, IE	17:15 - 17:30
WS18.03	The effect of CFTR modulator therapy on the sputum proteome in cystic fibrosis	17:30 - 17:45
	<i>Oral Presenter:</i> Rosie Maher, GB	
WS18.04	Change in markers of systemic inflammation after elexacaftor/tezacaftor/ivacaftor initiation: results from 18 months follow-up in the Danish cystic fibrosis cohort <i>Oral Presenter:</i> Thomas Bryrup, DK	17:45 - 18:00
WS18.05	Improved diagnosis of early aspergillus lung disease in cystic fibrosis (IDEAL) study design and first results <i>Oral Presenter:</i> Federico Mollica, NL	18:00 - 18:15
WS18.06	The impact of cytomegalovirus on airway epithelial gene expression <i>Oral Presenter:</i> Julianna Svishchuk, CA	18:15 - 18:30

Workshop

17:00 - 18:30

R5

WS19 - WS19 - Complex Psychosocial/Nursing case studies*Chair:* Katrien Van Gompel, BE*Chair:* Espérie Burnet, FR

WS19.01	Non-adherence in the twilight zone: the complexity of complex medications <i>Oral Presenter:</i> Laura Moyens, BE	17:00 - 17:30
WS19.02	"Could it be the new meds?" Clinical Psychology case study reminding us to ask all the old questions in this era of all new modulator medications <i>Oral Presenter:</i> Helen Love, GB	17:30 - 18:00
WS19.03	Triple therapy for Cystic Fibrosis (Elexacaftor/Tezacaftor/Ivacaftor): desensitization after an adverse reaction with severe skin rash <i>Oral Presenter:</i> Juliana Roda, PT	18:00 - 18:30

Saturday, 10. June 2023*Symposium*

09:00 - 10:30

R1

S21 - Symposium 21 - Novel endpoints in clinical trials*Chair:* Damian Downey, GB*Chair:* Philippe Reix, FR**PK studies in children - pitfalls and challenges**

09:00 - 09:22

Speaker: Saskia De Wildt, NL**Airway clearance: can it be measured?**

09:22 - 09:44

Speaker: Gemma Stanford, GB**Radiological endpoints in pre-school children**

09:44 - 10:06

Speaker: Harm Tiddens, NL**Measuring bugs in the era of CFTR modulators**

10:06 - 10:30

Speaker: Jerry Nick, US*Symposium*

09:00 - 10:30

R2

S22 - Symposium 22 - Inflammation in CF organs*Chair:* Olaf Eickmeier, DE*Chair:* Dorota Sands, PL**Intrinsic CFTR related airway inflammation**

09:00 - 09:22

Speaker: Robert Gray, GB**Pathogen induced airway inflammation**

09:22 - 09:44

Speaker: Michael Tunney, GB**2023 Update on gut inflammation**

09:44 - 10:06

Speaker: Jochen Mainz, DE**Inflammatory joint disease in cystic fibrosis**

10:06 - 10:30

Speaker: Jobst Roehmel, DE*Symposium*

09:00 - 10:30

R3

S23 - Symposium 23 - CFTR-related pancreas disease*Chair:* Frank Bodewes, NL*Chair:* Stephanie Van Biervliet, BE**Pancreatic complications in cystic fibrosis from bench to bedside**

09:00 - 09:22

Speaker: Zachary Sellers, US**CFTR mutations in the pancreas**

09:22 - 09:44

Speaker: Grzegorz Oracz, PL**The effect of CFTR modulators on exocrine pancreatic function**

09:44 - 10:06

Speaker: Keith Chee Y. Ooi, AU**CFTR related disorders of the pancreas**

10:06 - 10:30

Speaker: Isabelle Scheers, BE

Symposium

09:00 - 10:30

R4

S24 - Symposium 24 - Understanding more about CFSPID: from diagnosis to outcomes and the role of registries

Chair: Alexander Elbert, US

Chair: Maya Desai, GB

Newborn screening programmes and CFSPID epidemiology across Europe 09:00 - 09:18

Speaker: Carlo Castellani, IT

Phenotypes, care pathways and outcomes in CFSPID 09:18 - 09:36

Speaker: Anne Munck, FR

CFSPID should be included within national CF Registries (PRO) 09:36 - 09:58

Speaker: Susanna McColley, US

CFSPID should be included within national CF Registries (CON) 09:58 - 10:20

Speaker: Andreas Jung, CH

Discussion 10:20 - 10:30

Symposium

09:00 - 10:30

R5

S25 - Symposium 25 - Exploiting novel targets for cystic fibrosis therapies

Chair: Miquéias Lopes-Pacheco, PT

Chair: Iwona Pranke, FR

Targeting SMG6-mediated pathway to rescue nonsense CFTR variants 09:00 - 09:22

Speaker: Lulu Huang, US

Global functional genomics reveals GRK5 as a therapeutic target for cystic fibrosis 09:22 - 09:44

Speaker: Hugo Botelho, PT

Esc peptides: novel therapeutic agents with dual potentiator and antimicrobial activity 09:44 - 10:06

Speaker: Maria Luisa Mangoni, IT

Potassium channels as alternative targets to modulate transepithelial fluid secretion 10:06 - 10:29

Speaker: Guy Moss, GB

Closing Plenary

11:00 - 12:30

R1

Closing Plenary

"Hurdles" on genetic therapies - Delivery, other organs.. 11:00 - 11:30

Speaker: Patrick Harrison, IE

Longitudinal data from the ECFS-Patient Registry: What happened in the last decade and what can we learn from it about the future? 11:30 - 12:00

Speaker: Eitan Kerem, IL

Closing Ceremony

12:30 - 12:50

R1

Closing Ceremony