



EAPC 17th World Congress Online

“EXPLORING NEW DIMENSIONS”

Interactive Online Sessions

6 - 8 October 2021

EAPC 17th World Congress Online Programme

06.10.2021

09:00 – 9:50

PL 1 Opening Ceremony

Is Palliative Care having an existential crisis?

Seamus O’Mahony, Ireland

EAPC Award: Inequalities and Injustice in Palliative Care

Irene Higginson, UK

10:00 – 10:50

PS 1 (E) Massive Open Online Courses (MOOCs) to enhance education in palliative care

Using MOOCs to develop community discussion around death and dying

Jennifer Tieman, Australia

Using MOOCs to achieve international dissemination of research

Nancy Preston, United Kingdom

What’s online line learning like? Experiences of a user from Lebanon

Silva Dakessian, Lebanon

10:00 – 10:50

PS 2 (A) Innovative models of home palliative care

Updated Cochrane review results on the effectiveness and cost-effectiveness of home palliative care services for adults

Barbara Gomes, United Kingdom

Community-based short-term integrated palliative care for older people with chronic noncancer conditions: a randomised controlled mixed method trial

Catherine Evans, United Kingdom

Accelerated transition to palliative care at home in advanced cancer: A randomized clinical trial

Per Sjogren, Denmark

11:00 – 11:50

PS 3 (B) Improving care for people unable to self-report: the role of person-centred proxy-reported assessment measures

Using structured, person-centred measures for people with dementia unable to self-report to identify and meet palliative care needs

Susanne de Wolf-Linder, Switzerland

Nuriye Kupeli, United Kingdom

Person-centred proxy measures: clinical and methodological challenges, and recommendations to maximise valid and reliable assessment.

Clare Ellis-Smith, United Kingdom

	<p>Assessing palliative care needs in people with dementia using proxy measurement: validation of the Integrated Palliative care Outcome Scale for Dementia (IPOS-Dem) in German nursing homes Michaela Schunk, Germany</p>
12:00 – 12:50	<p>PS 4 (B) Management of fatigue in advanced cancer - psychological, physical and pharmacological</p> <p>Psycho-educational interventions for fatigue in palliative care Asanga Fernando, United Kingdom</p> <p>Exercise interventions for fatigue in palliative care María del Rocío Rojí Buqueras, Spain</p> <p>Drug treatments for fatigue in palliative care Paddy Stone, United Kingdom</p>
10:00 – 10:50	<p>FC 01 (B) Measuring and managing symptoms</p> <p>B-05 Cost-effectiveness of sustained-release morphine for refractory breathlessness in COPD: a randomized clinical trial Cornelia Van den Berg - Verberkt, The Netherlands</p> <p>B-06 Healthcare Resource Utilisation by Patients with Advanced Cancer Living with Pain: An Analysis from the STOP Cancer Pain Trial Nikki McCaffrey, Australia</p> <p>B-08 Longitudinal symptom profile of palliative care patients receiving a nurse-led end of life (PEACH) Program to support preference to die at home Meera Agar, Australia</p> <p>L-08 Identifying the consequences of restrictive visitor policies on palliative care providers, their patients, and caregivers. Kirsten Wentlandt, Canada</p> <p>L-13 Factors Influencing Acceptance or Refusal of Palliative Care among Patients with Moderate to Severe Symptoms: A Qualitative Study Rachel Sue-A-Quan, Canada</p>
11:00 – 11:50	<p>FC 02 (D) ACP & CPR</p> <p>D-01 Do physicians discuss Cardio-Pulmonary Resuscitation (CPR) prognosis with hospitalized older patients? An analysis of admission interviews Clara Castillo, Switzerland</p> <p>D-13 Advanced Care Planning in Nursing Home Patients with and without Cognitive Impairment who Are at the End of Life Rafael Montoya Juárez, Spain</p> <p>N-01 When to initiate advance care planning conversations in patients with advanced cancer? Results of the ACTION trial Judith Rietjens, The Netherlands</p>

<p>11:00 – 11:50</p>	<p>M-12 Advance care planning in primary care for people with gastrointestinal cancers: a feasibility trial Kirsty Boyd, UK</p> <p>FC 03 (A) Innovation and new technologies</p> <p>A-02 Measurement equivalence of the paper-based and electronic version of the Integrated Palliative care Outcome Scale: a randomised crossover trial Christina Ramsenthaler, Germany</p> <p>A-06 Development and Evaluation of a Web-based Program for Advance Care Planning Doris van der Smissen, The Netherlands</p> <p>A-04 Developing a Digital Game to Improve Public Perception of Dementia Gillian Carter, United Kingdom</p> <p>A-07 'Educating RITTA': Evaluation of an Artificial Intelligence Programme in Opioid Prescribing - A Pilot Project and Needs Assessment Mark Taubert, United Kingdom</p> <p>A-09 Implementation of eHealth Interventions to Support Assessment and Decision Making for Residents with Dementia in Care Homes: A Systematic Review Juliet Gillam, UK</p>
<p>15:00 – 15:050</p>	<p>PL 2</p> <p>Inclusivity in palliative care – are we as good as we think? Richard Harding, UK,</p> <p>Prize: The IJPN Palliative Care Nurse of the Year</p> <p>Best Abstract: B-01 The SILENCE (Scopolamine butylbromide given prophylactically for death rattle) study: a double-blind randomized, placebo-controlled multicenter trial. Jet van Esch, The Netherlands</p>
<p>16:00 – 16:50</p>	<p>PS 5 (F) Economics of palliative care: new approaches to understand and demonstrate effects</p> <p>Can palliative care mitigate inequities at end of life? Joachim Cohen, Belgium</p> <p>Trajectories of health care expenditures after the onset of functional disability: new insights from latent class analysis Claire Ankuda, United States</p> <p>Projecting future needs, costs and outcomes in palliative care: an economic policy evaluation for the Department of Health (Ireland) Peter May, Ireland</p>

<p>16:00 – 16:50</p>	<p>PS 6 (A) "There is an app for that" – leveraging technology to advance palliative care to patients and families across care settings</p> <p>When digitalization goes viral, in the pandemic and in palliative care David Blum, Switzerland</p> <p>What do people need from technology in the provision of palliative care? Findings from user engagement and technology implementation across low, middle and high-income settings Matthew Allsop, United Kingdom</p> <p>"I went to the beach without leaving my bed" -- virtual reality for symptom management in advanced illness: current evidence and applications Hunter Groninger, United States</p>
<p>18:00 – 18:50</p>	<p>PS 7 (A) Peer support in palliative care: how to leverage its potential?</p> <p>An introduction to peer support: what do we know, what questions remain, and what is the future of peer support? Kate Lorig, United States</p> <p>Face-to-face peer support groups in palliative care: findings from the first large-scale SMART study. Erica Witkamp, The Netherlands</p> <p>Viral- International online peer support & learning in palliative care Mark Taubert, United Kingdom</p>
<p>16:00 – 16.50</p>	<p>FC 04 (B) Symptoms and Sedation</p> <p>B-07 Clinical Aspects of Palliative Sedation: A Systematic Review of Prospective Studies Alazne Belar, Spain</p> <p>B-10 Prophylactic Drugs for Irradiation-induced Neurocognitive Decline: A Systematic Review on Efficacy and Outcomes Filipa Tavares, Portugal</p> <p>B-14 Understanding palliative care needs of people with frontotemporal lobar degeneration (FTD); systematic review Ali-Rose Sisk, United Kingdom</p> <p>F-02 Continuous Deep Sedation until Death (CDS) in France: What Is at Stake for Patients, Proxies and Physicians in the Context of the 2016 Law? Sandrine Bretonniere, France</p> <p>Q-02 Guidelines to Reduce, Handle and Report Missing Data in Palliative Care Trials Co-produced Using a Multi-stakeholder Nominal Group Technique Jamilla Hussain, United Kingdom</p>

18:00 – 18:50	<p>FC 05 (R) COVID 1</p> <p>R-02 Initiating Advance Care Planning Discussions during Covid-19: A Mixed Methods Study of Healthcare Professionals' Experiences Mary Turner, UK</p> <p>R-03 Barriers and Facilitators to Specialized Palliative Care Integration for Patients with COVID-19 Kirsten Wentlandt, Canada</p> <p>R-06 Care for dying patients under the Covid-19 pandemic in Norway: a survey of bereaved relatives Dagny Haugen, Norway</p> <p>R-08 Experiences, challenges and potential solutions of generalist palliative care in inpatient setting during the SARS-CoV2 pandemic Manuela Schallenburger, Germany</p> <p>R-09 Changes in patterns of mortality and place of death during the COVID-19 pandemic: a retrospective analysis of data from the four nations of the UK Sean O'Donnell, UK</p>
07.10.2021	
09:00 – 09:50	<p>PL 3</p> <p>Meeting the challenge of dementia care in the future Deborah Parker, Australia</p> <p>Leadership Education in Palliative Care Award</p> <p>Best Abstract: L-01 Palliative day care in Belgium: health economic analysis using population-level administrative data Kristof Faes, Belgium</p>
10:00 – 10:50	<p>PS 8 (M) Expanding palliative care beyond health care services: lessons learned from international research</p> <p>How do specialized palliative care services engage with the general public? A survey from Belgium, Sweden and the UK Sally Paul, United Kingdom</p> <p>Creating Collaborative Community Culture at the End of Life: Insights from an Asset Based Community Development project in Australia Andrea Grindrod, Australia</p> <p>Creating palliative care capacity across society: evaluating compassionate city programs Joachim Cohen, Belgium</p>
10:00 – 10:50	<p>PS 9 (G) Revealing the current state of palliative care for persons with persistent mental illness in three high income countries</p> <p>The views and experiences of people with mental illness, and their carers, on their palliative care needs Jed Jerwood, United Kingdom</p>

<p>10:00 – 10:50</p>	<p>Palliative care for people diagnosed with mental illness : The New Zealand perspective Helen Butler, New Zealand</p> <p>Palliative care in the gaps: Experiences of palliative care for persons with persistent mental illness in the community Erin E. Donald, Canada</p> <p>FC 06 (J) Compassionate Communities and Civic responsibilities</p> <p>E-02 Implementing a Compassionate Communities Framework to Raise Public Awareness of Palliative Care in Alberta, Canada Konrad Fassbender, Canada</p> <p>G-02 The preferred place of burial in the context of migration: The example of Turkish migrants in Germany Marco Hajart, Germany</p> <p>J-04 Civic Engagement Concerning Serious Illness, Dying, Loss and Bereavement: A Systematic Integrative Review Louise D'Eer, Belgium</p> <p>J-05 Compassionate Cities: A Systematic Integrative Review of Existing Initiatives Worldwide Bert Quintiens, Belgium</p> <p>L-16 Key Areas of Clinical Practice that Enable Optimal Palliative Care in Acute Hospitals: A Mixed Methods Study Claudia Virdun, Australia</p>
<p>11:00 – 11:50</p>	<p>PS 10 (B) (Potential) Pitfalls in palliative care pharmacotherapy</p> <p>Darwin's Legacy - Drug Interactions and Dietary/Herbal Supplements Andrew Dickman, United Kingdom</p> <p>Is any jerk a seizure? Constanze Rémi, Germany</p> <p>The Cannabis Conundrum Jennifer Schneider, Australia</p>
<p>12:00 – 12:50</p>	<p>PS 11 (D) Developing an evidence base to support advance care planning in dementia</p> <p>Developing guidance in addressing the challenges of advance care planning in dementia: an EAPC Delphi study Jenny T van der Steen, The Netherlands</p> <p>Implementing advance care planning with family caregivers of nursing home residents in Italy Paola Di Giulio, Italy</p> <p>Engaging in advance care planning conversations: lessons learned from people with dementia and their family Lieve van den Block, Belgium</p>

11:00 – 11:50	<p>FC 07 (L) Models of Care</p> <p>L-04 Enhanced Supportive Care Impact for Patients and Healthcare System Ann Griffiths, United Kingdom</p> <p>L-06 Thank goodness, you are here: a pre-post mixed methods study to assess the impact of 7-day specialist palliative care on clinical outcomes and experiences of patients, carers and staff. Sarah Brearley, United Kingdom</p> <p>L-07 Evaluation of an enhanced 7-day Specialist Palliative Care Service: Findings from a quantitative analysis Ceu Mateus, United Kingdom</p> <p>L-12 Specialist palliative care teams and characteristics related to referral rate: a national cross-sectional survey among hospitals in the Netherlands Manon Boddaert, The Netherlands</p> <p>M-08 The role of emergency medical services in the palliative care – scoping review David Peran, Czech Republic</p>
12:00 – 12:50	<p>FC 08 (H) Psychological symptoms and communication</p> <p>H-05 Let It Out (LIO): a mixed-methods study to optimise the design of an online, emotional disclosure intervention for adults with terminal illness receiving hospice care Daisy McInnerney, United Kingdom</p> <p>H-09 The Correlation between Suffering and Spiritual Distress in Elderly Cancer Patients: A Cross-sectional Study Helga Martins, Portugal</p> <p>M-10 Perceived Quality of Care and Emotional Functioning of Patients with Advanced Cancer and their Relatives: Results of a Multicenter Observational Cohort Study (eQuiPe) Janneke van Roij, The Netherlands</p> <p>Q-01 Using a behavioural theory to gain insight into critical determinants of health promoting behaviour around serious illness - A case example applied to starting a conversation about palliative care Anne-Lore Scherrens, Belgium</p>
12:00 – 12:50	<p>FC 09 (D) Older & frail people</p> <p>D-02 Older People with Severe Frailty talking about their Palliative Care Needs: An Interview and Survey Study during the Covid-19 Pandemic Richard Green, United Kingdom</p> <p>C-03 Predictors of pre-death and post death grief in family carers of people with dementia. A systematic review. Sophie Crawley, United Kingdom</p>

15:00-15:50	<p>D-03 Symptom Control and Palliative Care Outcomes in Patients at the End of Life in Nursing Homes Rafael Montoya Juárez, Spain</p> <p>D-11 Perspectives of Older People Living with Mild Dementia about Eating and Drinking Problems at the Later Stages of Dementia: A Qualitative Study Kanthee Anantapong, United Kingdom</p> <p>M-13 Experiences of a Novel Integrated Care Service for Older Adults at Risk of Severe Frailty: An Analysis of Survey and Interview Data Mabel Okoeki, United Kingdom</p> <p>PL 4</p> <p>Technology, Innovation and Artificial Intelligence – a vision for Palliative Care Björn Eskofier, Germany</p> <p>Prize: “Palliative Medicine” - Paper of the Year</p> <p>Best Abstract: L-03 Demonstrating the impact of palliative care: a secondary analysis of routinely-collected person-centred outcomes data among hospice community patients Fliss Murtagh, UK</p>
16:00 – 16:50	<p>PS 12 (B) Treatment decision-making: Communicating, informing and supporting patient choice</p> <p>Clinician-patient communication and treatment decision-making support in advanced kidney disease -Protocol for the OSCAR Study Lucy Selman, United Kingdom</p> <p>Information-provision in advanced cancer care Liesbeth van Vliet, The Netherlands</p> <p>Supporting treatment decision-making in practice Martin Loucka, Czech Republic</p>
16:00 – 16:50	<p>PS 13 (C) Children and Young People: The Hidden Carers</p> <p>Children and young people caring for dying parents in the UK Steve Marshall, United Kingdom</p> <p>U.S. Caregiving interventions for children and youth caregivers Melinda Kavanaugh, United States</p> <p>Protecting mental health by identifying children and their needs when an adult is ill Jeffrey Hanna, United Kingdom</p>
16:00 – 16:50	<p>FC 10 (E) Education and Workforce</p> <p>C-08 Adaptation and continuous learning to start and persevere in Palliative Care. Revisiting professional coping strategies through an integrative review Maria Arantzamendi, Spain</p>

	<p>E-03 Objective structured clinical examination (OSCE) in assessment of Palliative Care competence of advanced nurse practitioner students Virpi Sulosaari, Finland</p> <p>E-04 Building Capacity for Early Palliative Care within Primary Care Teams: Implication and Evaluation of the CAPACITI Program Hsien Seow, Canada</p> <p>G-05 Navigating tensions between ideal and realistic palliative care in socially deprived areas: a multiple case study analysis of healthcare professionals' experiences Maddy French, United Kingdom</p> <p>R-01 Reflections and experiences of physicians working during the COVID-19 pandemic: a qualitative study. Lindsay Hurlburt, Canada</p> <p>PS 14 (E) Rebalancing death and dying: the Lancet Commission on the Value of Death</p> <p>Death, medicine, immortality, and climate change Richard Smith, United Kingdom</p> <p>The complex death system: mapping people, pathways, and structural determinants of end of life care Afsan Bhadelia, United States</p> <p>The Lancet Commission on the Value of Death: recommendations and next steps Libby Sallnow, United Kingdom</p> <p>PS 15 (Q) Researcher Award Session</p> <p>Clinical Impact Harvey Chochinov, Canada</p> <p>Post Doc</p> <p>English Palliative and End of Life Care: Several Observations from an Anthropologist Erica Borgstrom, United Kingdom</p> <p>Early Researcher</p> <p>Improving patient and family care using patient centred outcome measures - from development to implementation into practice Barbara Antunes, Portugal</p>
08.10.2021	
09:00 – 09:50	<p>PL 5</p> <p>Using the lived experience for re-designing care pathways Ingeborg Griffioen, The Netherlands</p> <p>Award Photo competition</p>

	<p>Best Abstract: D-06 Is Palliative Care Provision Associated with Prevalence of Death in Hospital at the Population Level? Evidence from 30 European Countries in the Years 2005-2017 Jingjing Jian, China</p>
10:00 – 10:50	<p>PS 16 (B) Opioids for chronic breathlessness - practical evidence-based prescribing</p> <p>Does morphine help? Evidence for effectiveness, dose and preparations David Currow, Australia</p> <p>What about other opioids: what do we know, what don't we know? Miriam Johnson, United Kingdom</p> <p>Are opioids safe in chronic breathlessness? Daisy Janssen, The Netherlands</p>
10:00 – 10:50	<p>FC 11 (J) Dementia and Care homes</p> <p>J-03 Factors Associated with Emergency Department Attendance by People with Dementia Approaching the End-of-Life: A Systematic Review Lesley Williamson, United Kingdom</p> <p>D-09 Development of an advance care planning by proxy intervention for residential aged care residents without decision making capacity Laura Jones, Switzerland</p> <p>D-14 What influences Quality of Death and Dying in Dutch nursing homes? Judith Meijers, The Netherlands</p> <p>J-09 Patterns of unplanned hospital admissions among people with dementia: From diagnosis to the end of life Emel Yorganci, United Kingdom</p> <p>L-10 Developing a Benchmarking Tool for Evaluating the Quality of End-of-life care in Care Homes Helen Chan, Hong Kong, China</p>
11:00 – 11:50	<p>PS 17 (D) Exploring new dimensions in dementia palliative care: making research work in practice</p> <p>Providing a firm base: A system-based logic model of integrated palliative dementia (the EMBED-Care programme) Catherine Evans, United Kingdom</p> <p>Learning from trials: LIVE@Home.path, an RCT of care coordination for people with dementia Bettina Husebo, Norway</p> <p>The reality: Understanding why in practice, interventions don't work as expected Jenny T van der Steen, The Netherlands</p>

11:00 – 11:50	<p>FC 12 (M) PC identification and impact</p> <p>J-01 Population-based Projections of Place of Death for Northern Ireland by 2040 Tracey McConnell, United Kingdom</p> <p>K-01 Palliative care, COVID-19 and Universal Health Coverage: Results of a Global Survey Stephen Connor, United Kingdom</p> <p>M-03 Implementation of the Supportive and Palliative Care Indicators Tool (SPICTTM) in General Practice in Germany – An Interventional Study Kambiz Afshar, Germany</p> <p>M-05 Concurrent validity and prognostic utility of the Needs Assessment Tool: Progressive Disease - Heart Failure Christina Ramsenthaler, Germany</p>
12:00 – 12:50	<p>PS 18 (F) Ethical decision making at the end-of-life</p> <p>Nudging and Palliative paternalism Carlo Peruselli, Italy</p> <p>Autonomy: a perspective from the ethics of care Ludovica De Panfilis, Italy</p> <p>Personal autonomy and end-of-life decision making Kevin Brazil, United Kingdom</p>
12:00 – 12:50	<p>FC 14 (C) Families and Carers</p> <p>C-02 Dying at ‘Home’: Bereaved Family Caregivers’ Experiences of Just-in-Case Medications at End of Life Christina Faull, United Kingdom</p> <p>C-04 Family Carers’ Support from Healthcare Professionals in Conducting ACP Conversations with their Seriously Ill Relatives: A Cross-sectional Survey of Bereaved Family Carers Isabel Vandenbogaerde, Belgium</p> <p>C-06 A Systematic Review with Narrative Synthesis on Mutual Support between Patients and Family Caregivers in Palliative Care Rachel McCauley, Ireland</p> <p>C-07 Caregiving experiences of informal carers of people living with MND; a qualitative evidence synthesis Kate Flemming, United Kingdom</p> <p>L-05 How to provide grief support to caregivers systematically? The evaluation of a grief support toolkit for generalist healthcare professionals. Marcella Tam, The Netherlands</p>
15:00 – 15:50	<p>PS 19 (B) Sedation at the end of life outside specialist palliative care – international perspectives</p>

	<p>Use of sedatives and sedation at the end of life outside specialist palliative care in Germany Eva Schildmann, Germany</p> <p>Challenges in the practice of palliative sedation outside specialist palliative care: data from an international questionnaire survey and in-depth interviews in the Netherlands Agnes van der Heide, The Netherlands</p> <p>Palliative sedation in Flemish nursing homes: reported complexities and the development of a setting-specific practice protocol Kenneth Chambaere, Belgium</p>
16:00 – 16:50	<p>PS 20 (C) Strategies to improve the psychosocial wellbeing of family carers: how international collaborations make a difference</p> <p>Evidence synthesis of factors affecting family carers' psychological wellbeing" Tracey Shield, United Kingdom</p> <p>Supporting family carer decision-making in palliative dementia care Kevin Brazil, United Kingdom</p> <p>Key considerations when selecting family carer outcome measures James Dionne-Odom, United States</p>
16:00 – 16:50	<p>FC 16 (R) COVID 2</p> <p>O-02 The support needs and experiences of people bereaved in the UK during the Covid-19 pandemic: baseline results from a mixed-methods longitudinal survey Emily Harrop, United Kingdom</p> <p>R-10 Bereaved Relatives' Quality of Life Pre- and post-COVID-19 Pandemic: Results of the Prospective, Multicenter, Observational eQuiPe Study Laurien Ham, The Netherlands</p> <p>R-12 How the COVID-19 pandemic affects palliative care inpatients and outpatients without a COVID-19 diagnosis and their families - a qualitative interview study Natalie Berges, Germany</p> <p>R-14 Death and bereavement due to COVID-19: A discourse analysis of online newspapers with implications for end of life care Lucy Selman, United Kingdom</p> <p>R-15 "I promised my husband I wouldn't leave him alone": Burdens due to visit restrictions for dying patients during the COVID-19 pandemic in Germany: a mixed-methods study of bereaved relatives. Karlotta Schlösser, Germany</p>
18:00 – 18:50	<p>PL 6 Closing Ceremony</p> <p>Floriani Lecture: How can palliative care navigate the opioid crisis? Jim Cleary, United States</p> <p>Prize: "Journal of Palliative Medicine" European Paper of the Year</p>

	<p>Best Abstract: B-04 Analgesic prescribing patterns for people dying with and without dementia in the UK: general practice cohort study Liz Sampson, United Kingdom</p> <p>EAPC World Research Congress 2022 Camilla Zimmermann, Canada</p>
08.10.2021	Paediatric Palliative Care Day
08:00 – 08:50	<p>Meet the experts Richard Hain, United Kingdom Joana Mendes, Portugal</p>
10:00 – 10:50	<p>PL 1 Interventions at the end of life in children: where evidence meets experience Richard Hain, United Kingdom</p>
11:00 – 11:50	<p>FC 13 (I) Advancing ACP I-01 Evaluating attitude, self-efficacy, behaviour, and intention to engage in advance care planning communication in paediatric oncology: development and face validation of a new measurement instrument Anne Van Driessche, Belgium I-02 Parental decision-making behaviours in paediatric ACP: A qualitative study Kathrin Knochel, Switzerland I-03 Patterns in discussing End-of-Life in paediatric Advance Care Planning conversations Julie Brunetta, The Netherlands I-04 Impact of Advance Care Planning in Paediatric Oncology Miguel Vieira Martins, Portugal I-10 Conceptualising paediatric advance care planning; is it more than just 'do not attempt resuscitation'? Sidharth Vemuri, Australia</p>
12:00 – 12:50	<p>PS 1 Transitioning children with palliative care needs between settings and phases From hospital to home or hospice care Mette Asbjörn Neergaard, Denmark From NICU to home care Finella Craig, United Kingdom From chronic phase to EoL Laura Madanat, Finland</p>

13:00 – 13:50	<p>Lunch time with Paediatrics</p> <p>Ana Lacerda, Portugal Lizzie Chambers, United Kingdom</p>
15:00 – 15:50	<p>FC 15 (I) Sharing learning from across the globe</p> <p>I-07 What matters to children and families facing advanced cancer? Identification of priority outcomes to improve paediatric palliative care in Turkey and Jordan Sabah Boufkhed, UK</p> <p>I-08 End of Life Care for Newborns with Major Congenital Malformations in a Tertiary Neonatal Referral Center in Brazil Maria Augusta Gibelli, Brazil</p> <p>I-09 The Butterfly Pathway: Supporting Families Following the Antenatal Diagnosis of a Life-limiting Condition in a Neonatal Intensive Care Unit Setting Charlotte Burleigh, United Kingdom</p> <p>I-12 Existential and spiritual care needs of parents of children with a life-limiting condition, a qualitative interview study Marije Brouwer, The Netherlands</p> <p>I-15 Existential Concerns of Children and Young People with Life-limiting or Life-threatening Conditions Hannah Scott, United Kingdom</p>
16:00 – 16:50	<p>PL 2</p> <p>Balancing aggressive treatment and palliative care – is it achievable? Joana Mendes, Portugal</p>