

7TH PUBLIC HEALTH PALLIATIVE CARE INTERNATIONAL CONFERENCE

Democratizing caring, dying and grieving participation, action, understanding and evaluation

20-23 SEPTEMBER 2022 - BRUGES, BELGIUM

Programme Poster exhibition

Conference Venue: Bruges Meeting & Convention Centre (BMCC), Beursplein 1, 8000 Bruges

Poster exhibition: Foyer, level 3

Group 1: Poster exhibition on 20-21 September 2022

ACP & End of Life communication

140	Tessa	Bergman	<i>Information meetings on end-of-life care organized by GPs and other organizations; do they differ in audience and results?</i>
101	Marcelo	Caballero	<i>Idiolectic Conversation and Symptom Management in Palliative Care Patients</i>
211	Ludovica	De Panfilis	<i>Development and evaluation of an in-hospital service to inform people about end-of-life rights: the SCEGLIERE service</i>
36	Nicholas	Jennings	<i>Medical end-of-life decision-making in a small resource-poor Caribbean country: a mortality follow-back study of home deaths</i>
147	Maria Fidelis	Manalo	<i>Talking about death to advanced cancer patients: not if, but how</i>
171	Fanny	Monnet	<i>Developing an advance care planning website for dementia: Integrating a user-centered approach and patient and public involvement</i>
235	Marta	Perin	<i>Advance Care Planning in Italian palliative care networks: a grounded theory study</i>
173	Galmiche	Perrine	<i>Mirror methodology: using marketing codes for better talking about end-of-life</i>

195	Ramona	Schenell	<i>To make and execute decisions throughout life- a model to facilitate self-determination in residential care.</i>
184	Steffen	Eychmüller	<i>Sense of security in palliative homecare: A qualitative descriptive study on the experiences of patients, family caregivers and outpatient healthcare professionals.</i>
120	Willemijn	Tros	<i>General practitioners' evaluations of optimal timing to initiate ACP for patients with cancer, organ failure or multimorbidity</i>
218	Isabel	Vandenbogaerde	<i>Support from health care professionals in empowering family carers to discuss Advance Care Planning: A population-based survey</i>
256	Sarah	Vilpert	<i>Role of end-of-life care planning in knowledge of partner's preferences for end-of-life aspects</i>

Community

293	Gabriela	Hidalgo	<i>Implementation of community-based palliative care in Primary Health Care in integration with Specialized Outpatient Care</i>
178	Briony	Hudson	<i>Exploring the support needs of the British Muslim Community with palliative care needs – a collaborative qualitative project (TBC)</i>
152	Helen	MacGregor	<i>Supporting compassionate communities in dying, death & bereavement by providing space for orgs to gather, discuss & learn"</i>
259	Steven	Vanderstichelen	<i>Charters are not enough: supporting volunteering at the volunteer level</i>
117	Julia	Verne	<i>A workplace compassionate community approach to support bereaved staff in the two national public health organizations in England (TBC)</i>

COVID

97	Shirley H.	Bush	<i>Experiencing and making sense of illness in an inpatient specialist palliative care unit during a pandemic: an interview study</i>
272	Barbara	Gale	<i>Hospice care - where do we go now?</i>
136	Kathy	Kortes-Miller	<i>Not Just A Visitor; Caregiving experiences in Long Term Care during COVID-19</i>
26	Venetia	Lawless	<i>Home and Community Based Palliative Care: Shaping the Future from Lessons Learned During the COVID-19 Pandemic</i>
77	Matteo	Moroni	<i>Elderly Covid -19 patients in nursing home cared by young doctors together with Palliative Care teams. A model beyond pandemic?</i>

208	Stina	Nyblom	<i>Experiences among Swedish patients in specialized palliative home care and their significant others during the Covid-19 pandemic</i>
25	Vilma	Oliveros	<i>Are we now more empathetic, compassionate, and understanding towards difference?</i>
298	Gabriela	Rezende	<i>Impacts of the COVID-19 pandemic on palliative care: a qualitative and multicenter study</i>
90	Ahmed	Sharaf	<i>Opportunities and challenges to family carers end of life care research in the COVID pandemic: A researcher's reflections</i>
82	Katherine	Sleeman	<i>Variation in palliative care service response and association with Covid-19 mortality rates: an ecological study in UK regions</i>
232	Stavroula Myrto	Trianti	<i>Impact of Covid Pandemic on ambulatory palliative care</i>
119	Julia	Verne	<i>Learning lessons from COVID-19 experience for palliative care for people with dementia in England (TBC)</i>
267	Masha	Zee	<i>The relation between distress and feeling appreciated when providing end-of-life care during the COVID-19 pandemic (CO-LIVE).</i>

Various Topics

174	Agathe	CANT-DIOT	<i>How is end-of-life care organized around the world?</i>
93	Marius	Čiurlionis	<i>Home-based palliative care and re-hospitalizations for palliative care patients</i>
49	Ascensión	Doñate-Martínez	<i>Needs for timely palliative care in older patients with non-malignant diseases: a multicenter qualitative study</i>
261	Rodeyns	Julie	<i>Fostering empathy and compassion through art observation</i>
59	Anne	van Driessche	<i>Recruitment and retention in palliative care trials: challenges and strategies</i>
72	Sarah	Yardley	<i>Getting palliative prescription medications right at home, in hospital & hospice: Activity Theory analysis</i>
74	Sarah	Yardley	<i>Reclaiming relationships as a legitimate tool for systems change: A Change Laboratory approach to palliative care</i>
71	Sarah	Yardley	<i>Patient safety in palliative care: improving system design for safe, responsive medication management</i>

Group 2: Poster exhibition on 22-23 September 2022

Cancer

57	Torbjørn	Paulsen	<i>Early referral to a palliative team improves end- of- life care among gynecological cancer patients</i>
194	Germans Natuhwera	Germans	<i>Cultural and religious beliefs and traditions affecting attending the clinic for cancer and palliative care in Uganda. findings from a mixed-methods study</i>
196	Germans Natuhwera	Germans	<i>Lived experiences of health professional cancer patients and survivors. a phenomenological evidence</i>
192	Tara	Joyce McDonnell	<i>A Controlled Clinical Trial of the effects of Meditation on patient's Perceptions of their Quality of Life in an acute hospital</i>
297	Marysia	M R do Prado De Carlo	<i>Total Pain in The Context of Palliative Care and its Assessment in People with Advanced Lung Cancer</i>
143	Lisa	Shirt	<i>"Patient and Caregiver-reported Acceptability of an "Automatic" Supportive and Palliative Care Referral for Advanced Lung Cancer Patients</i>
44	Lidija	Veterovska Miljkovic	<i>Experience from the use of oxycodone / naloxone versus morphine in the treatment of severe cancer pain in palliative care in the Republic of North Macedonia</i>

Equity and Access

160	Clare	Howie	<i>Improving equity of palliative care provision within advanced heart failure via a realist synthesis: Palliative Heart Synthesis</i>
285	Jamilla	Hussain	<i>Equity, diversity and inclusion in palliative and end of life care practice and research: personal reflections</i>
213	Terhi Kukkonen	Kukkonen	<i>Developing the competence of palliative care –project, multidisciplinary cooperation in long distance region</i>
287	Rudina	Rama	<i>Barriers and Challenges to provide palliative care in developing countries- Case of Albania</i>

Euthanasia & MAID

156	Charlotte	Boven	<i>Relatives of cancer patients' experiences regarding the interaction with healthcare providers during a euthanasia process</i>
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157	Charlotte	Boven	<i>Healthcare providers' experiences regarding the interaction with family during a euthanasia process: A qualitative study</i>
288	Lavoie	David	<i>Creating narratives to support experiences of medical assistance in dying in Canada: qualitative investigation of nurses' views</i>
193	Gordon	Gubitz	<i>The Canadian Medical Assistance in Dying (MAiD) Curriculum Development Project</i>
132	Kathy	Kortes-Miller	<i>Untold Stories; Accompanying someone who used medical assistance in dying at the end of life</i>
138	Rikmenspoel	Marjolein	<i>Information and support regarding euthanasia: a qualitative study on the needs of nursing home residents in the Netherlands</i>

Family caregiving

83	Kristen	Abbott-Anderson	<i>Conversations and care at end-of-life for persons with dementia: Family care partners share their perspectives</i>
216	Dr Kabindra	Bhagabati	<i>Concerns of the patients' relatives about Breaking Bad News to patients – An experience in a tertiary cancer center in NE India</i>
46	Tineke	De Keyser	<i>Empowerment of carer in outpatient home care through small chunks of accessible intel"</i>
233	Nina	Malmström	<i>Being a child of a parent diagnosed with ALS - from the child's perspective</i>
229	Anneli	Ozanne	<i>Experiences of parenthood when a parent has been diagnosed with ALS</i>
202	F H D SHEHAN	SILVA	<i>Cultural adaptation and validation of Sinhala version of Modified Caregiver Strain Index</i>

Health Promotion

43	Madeleine	Juhrmann	<i>Exploring a Public Health Palliative Care Approach to Emergency Medical Services</i>
205	Tracey	McConnell	<i>Loneliness at end of life: A public health approach to promoting the health and wellbeing of terminally ill patients and their carers</i>
248	Katarzyna	Patynowska	<i>Promoting the health and wellbeing of family caregivers providing end-of-life care – the role of community Healthcare Assistant</i>

95	Marianne	Tinkler	<i>A framework for developing co-designed integrated service model of palliative care for people living with severe mental illness.</i>
52	Vincent	Van Goethem	<i>Development of a self-management eHealth program to support and empower people with advanced cancer and their family caregivers</i>

Literacy & Education

144	Ashley	Chastain	<i>Development and Testing of the Palliative Care-Related Knowledge, Attitudes & Confidence in Home Health Care Questionnaires</i>
37	Zara , Linda	Höök, Frode	<i>Education for palliative representatives increases the quality of palliative care</i>
258	Clément	Meier	<i>Perceptions and knowledge of end-of-life medical situations among older adults in Switzerland</i>
155	Danny	Vereecke	<i>Knowing, being proficient, and being in palliative care starts in education, but does it?</i>
88	Ruth Si Man	Wong	<i>A Systematic Scoping Review on Moral Distress in Undergraduate Medical Students</i>

Loss & grief

27	Mo	Blishen	<i>The Wind Telephone - How a disconnected telephone can connect to grief</i>
159	Charlotte	Boven	<i>Walking the walk in connection: Experiences of healthcare providers with bereavement care in oncology</i>

Paediatric Palliative Care

126	Soham	Bandyopadhyay	<i>A 12-month international, observational study of palliative care in childhood cancer patients</i>
227	Leigh	Donovan	<i>Beyond bricks and mortar: enabling equity of access to pediatric hospice care for Queensland children and families</i>
84	Leana	Spies	<i>A Study to understand the experiences of adolescents and young adults with cancer in a Northern Cape public health setting</i>

Poster Lightning Sessions

Wednesday 21 September 2022

During Lunch break: level 4, catering hall

49	Ascensión	Doñate-Martínez	<i>Needs for timely palliative care in older patients with non-malignant diseases: a multicenter qualitative study</i>
132	Kathy	Kortes-Miller	<i>Untold Stories; Accompanying someone who used medical assistance in dying at the end of life</i>
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233	Nina	Malmström	<i>Being a child of a parent diagnosed with ALS - from the child's perspective</i>
77	Matteo	Moroni	<i>Elderly Covid -19 patients in nursing home cared by young doctors together with Palliative Care teams. A model beyond pandemic?</i>

Thursday 22 September 2022

During Lunch: level 4, catering hall

227	Leigh	Donovan	<i>Beyond bricks and mortar: enabling equity of access to paediatric hospice care for Queensland children and families</i>
193	Gordon	Gubitz	<i>The Canadian Medical Assistance in Dying (MAiD) Curriculum Development Project</i>
25	Vilma	Oliveros	<i>Are we now more empathetic, compassionate, and understanding towards difference?</i>
202	F H D SHEHAN	SILVA	<i>Cultural adaptation and validation of Sinhala version of Modified Caregiver Strain Index</i>
59	Anne	van Driessche	<i>Recruitment and retention in palliative care trials: challenges and strategies</i>

Friday 23 September 2022

During Lunch: level 4, catering hall

285	Jamilla	Hussain	<i>Equity, diversity and inclusion in palliative and end of life care practice and research: personal reflections</i>
43	Madeleine	Juhrmann	<i>Exploring a Public Health Palliative Care Approach to Emergency Medical Services</i>
52	Vincent	Van Goethem	<i>Development of a self-management eHealth program to support and empower people with advanced cancer and their family caregivers</i>
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