

Poster presentations

Set 1

Setup: Thursday, 08.00 – 09.00

Removal: Friday, 14.15 – 14.30

Assessment and measurement tools

148	P1	Integration of the ESAS into Daily Cancer Symptom Management for Cancer Inpatients	Erin	Mutterback	Ottawa, CA
149	P2	Identifying Suffering - A Study to Improve Understanding of Care	Heather	Davies	Bristol, UK
150	P3	Translation and Validation of the Dutch Version of the 'Multidimensional Fear of Death Scale'	Peter	Pype	Antwerp, BE
151	P4	Care of the Dying Evaluation (CODE) – Developing an 'Outcome Measure' to Assess Quality of Care in the Last Days of Life	John	Ellershaw	Liverpool, UK
152	P5	A Theory-driven Evaluation on the Integration of a Palliative Care Network (PCN)	Daryl	Bainbridge	Hamilton, Ca
153	P6	Validation of the Minimal Documentation System (MIDOS) in Palliative Care	L	Bertram	Aachen, DE
154	P7	A Comparison of Different Data Collection Tools when Researching Spirituality in Palliative Care	Gudlaug Helga	Asgeirsdottir	Kopavogur, IS
155	P8	A Totally Computerized Home-care Medical Record in Palliative Care: Our Experience	Furio	Zucco	Garbagnate Milanese, IT
156	P9	Physical Activity Level as an Outcome Measure for Use in Cancer Cachexia Trials: A Feasibility Study	Matthew	Maddocks	Nottingham, UK
157	P10	Acceptability of the SPARC© Questionnaire for Identifying Supportive and Palliative Care Needs in People with a Recent Diagnosis of Thoracic Cancer	Matthew	Maddocks	Nottingham, UK
159	P12	Palliative and Supportive Care Needs of Thoracic Cancer Patients at Diagnosis	Ruth	England	Nottingham, UK
160	P13	What Does the International Classification of Functioning Disability and Health (ICF) Add to the Traditional Assessment Tools in Palliative Care Patients at End of Life?	Anna	Giardini	Montescano, IT
161	P14	Predicting Weight Loss in People with Cancer: Validation of a Screening Instrument	Vanessa	Halliday	Nottingham, UK
163	P16	Functional Assessment Predicts Ability to Complete Trial Follow Up in Cancer-induced Bone Pain	Angela	Scott	Edinburgh, UK
165	P18	A Comparison of Numerical Rating Scale (NRS) and Borg Scales in the Assessment of Breathlessness Severity in People with Chronic Heart Failure (CHF)	Stephen	Oxberry	Leeds, UK
166	P19	The Use of Two Common Palliative Outcome Measures in Clinical Care and Research: A Systematic Review about POS and STAS	Charles	Le Grice	London, UK
167	P20	Prognostic Factors for Survival in Cancer Patients in Iceland	Sigrídur	Helgadóttir	Reykjavík, IS
168	P21	Symptom Relief by Erythrocyte Transfusions - An Evaluation Using the ESAS Tool in an Advanced Home Care Unit	Ulla	Martinsson	Uppsala, SE
169	P22	The Team Observed Structured Clinical Encounter (TOSCE): An Assessment Tool for Team and Interprofessional Competencies in Palliative Care and Primary Care	Daryl	Bainbridge	Hamilton, CA

171	P24	New Results on Prognosis at the End of Life by Home Care Supportive Palliative Care Teams	Maria	Nabal	Lleida, ES
172	P25	Cross Cultural Adaptation of the Spanish Version of the Memorial Delirium Assessment Scale (MDAS)	Antonio	Noguera	Madrid, ES
173	P26	Physical Activity: A Responsive and Meaningful Patient-centred Outcome Measure in Palliative Oncology?	Richard	Skipworth	Edinburgh, UK
174	P27	Do We Really Use Pain Assessment Tools?	Sophie	Pautex	Collonge-Bellerive, CH
175	P28	Domains of Quality-of-Life Instruments Appropriate for Use in Palliative Care: A Review	Michael	Echteld	Amsterdam, NL
176	P29	Depression in Palliative Care Patients: Comparing the Use of Screening Tools (BDI-II, HADS) with a Psychiatric Diagnostic Instrument (SCAN 2.1)	Franca	Warmenhoven	Nijmegen, NL
177	P30	A Tool for Early Identification of Palliative Care Patients in General Practice, the Radboud Indicators for Palliative Care Needs (RADPAC)	Bregje	Thoonsen	Nijmegen, NL
178	P31	Psychometric Development of a Single-item End-of-Life Patient-reported Outcome (EOLPRO): The First Piece of the Puzzle	Nikki	Mc Caffrey	Adelaide, AU
179	P32	The Use of Questionnaires in Longitudinal Research with Patients with Advanced Cancer	Laurie	Dunn	Liverpool, UK
180	P33	Assessing Physical, Psychosocial and Spiritual Distress of Palliative Cancer Patients: Development of a Multidimensional Screening Instrument	Martin	Weber	Mainz, DE
181	P34	Carer Support Needs Assessment in End of Life Home Care, Phase 2: Design, Development and Testing of a Tool for Routine Practice	Gail	Ewing	Cambridge, UK
182	P35	The Palliative Sedation Checklist and Clinical Flowsheet (PASEF): A Pilot Study	Shirley	Bush	Ottawa, CA
183	P36	Technology-based Data Collection Allows Longitudinal Assessment across Palliative Care Settings	Amy	Abernethy	Durham, US
184	P37	Development and Initial Validation of an Instrument to Identify Children Requiring Palliative Care: The Paediatric Palliative Screening Scale (PaPaS Scale)	Eva	Bergsträsser	Zurich, CH
185	P38	Views and Preferences of People from Ethnic Minority Backgrounds in the UK in Improving Participation in a Palliative Care Stroke Questionnaire	Tariq	Saleem	London, UK
186	P39	The Most Common Symptoms and Needs of Patients Receiving Palliative Care in Iceland, a Longitudinal Study	Bryndis	Gestsdottir	Reykjavik, IS
187	P40	The Development and Testing of a New Outcome Measure for Hospices and Specialist Palliative Care Services: The SC-OM	Julia	Addington-Hall	Southampton, UK
188	P41	Acceptability and Validity of a Computerised Body Map for Pain Assessment in Cancer Patients	Ellen	Jaatun	Trondheim, NO
189	P42	Use of the Consciousness Scale for Palliative Care as a Prognostic Factor	José António	Ferraz Gonçalves	Porto, PT
192	P45	Validation of the Utrecht Symptom Diary	Ellen	de Nijs	Utrecht, NL
193	P46	Patient-focused Endpoints for Clinical Trials: Criterion-based Validation of Accelerometer-based Activity Monitoring in Advanced Cancer	Richard	Skipworth	Edinburgh, UK
194	P47	Being Depressed in the Palliative Phase – The Patients' Experiences	Elisabeth	Brenne	Trondheim, NO

195	P48	An Evaluation of the Usefulness of the Distress Thermometer for the Elderly (DTE) for Older Adults (60+) in Care Homes in Identifying Distress and Need	Jan	Oyebode	Birmingham, UK
196	P49	Symptom Assessment and Identification of Distress in the Oncology Day Unit	Sharon	Beatty	Drogheda, IE
197	P50	From MNUPAL to PERFORM: Improving Research and Education on Cancer Cachexia	Robert	Kilgour	Montreal, CA
198	P51	Malnutrition, Depression, Anxiety: A Comorbid Condition	Wadih	Rhondali	Lyon, FR

Audit and quality control

199	P52	A Prospective Audit of the Current Management of Dyspnoea among Day Hospice and Inpatients at a Hospice with Reference to the Local Guidelines	Emma	Dymond	Glasgow, UK
200	P53	Radiotherapy (RT) Treatment of Brain Metastases: Survival and Factors Influencing Outcome	Elaine	Wallace	Dublin, IE
201	P54	The Use of Melthylalntrexone in Clinical Practice in the West of Scotland	Fiona	Mcmunnigall	Denny, UK
202	P55	An Audit of In-patient Steroid Use in a Hospice Setting	Mohd Shazrul	M. Ramly	Limerick, IE
203	P56	Prognostic Characteristics of a Medical Emergency Team Population with a Known Life-limiting Illness	Fiona	Stafford-Bell	Sydney, AU
204	P57	Evaluation of a Specialist Pharmacist's Interventions in an Inpatient Hospice Setting	Andrew	Dickman	Liverpool, UK
205	P58	Does Daily Hospice Treatment Improve the Physical and Mental Health of Breast Cancer Patients Treated with Mastectomy?	Samir	Husić	Tuzla, BO
206	P59	A 1-year Quality Project to Develop a Generic Model for Improving Palliative Care in a Hospital Ward	Lene	Joergensen	Vejle, DK
207	P60	Quality Management Stages in a Palliative Care Program	Beatrice	Pop	Navalmoral de la Mata, ES
208	P61	Audit of Phlebotomy Practices in a Specialist Palliative Care Unit	Lucy	Balding	Cork, IE
209	P62	A Review of the Use of Anti-epileptic Drugs (AEDs) in High Grade Central Nervous System (CNS) Tumours	Elaine	Wallace	Dublin, IE
210	P63	Patients in the Last Year of Life Referred to the Acute Receiving Unit	Jennifer	Gray	Edinburgh, UK
212	P65	Staff Meetings in a Specialist Palliative Care Team: A Qualitative Observational Study	Claudia	Cugno	Torino, IT
213	P66	Agitation and Restlessness in the Last 24 Hours of Life on the Liverpool Care Pathway for the Dying Patient (LCP)	Tamsin	McGlinchey	Liverpool, UK
214	P67	Sedative use in End-of-Life Care: An Audit of Patterns of Potentially Sedative Medication Use in a Specialist Palliative Care Inpatient Unit	Marlise	Poolman	Wrexham, UK
215	P68	Back to Basics: Development of a Bowel Assessment Chart	Geraldine	Tracey	Dublin, IE
216	P69	Great Expectations? Do Expectations of Admission Outcome of Patients, Their Relatives and Referrers on Admission to a Specialist Palliative Care Unit Correlate with Each Other and the Final Outcome?	Damien	McMullan	Belfast, UK
217	P70	Audit of Discharge Communication for Inpatients Discharged on Corticosteroids from a Specialist Palliative Care Unit	Damien	McMullan	Belfast, UK

219	P72	A Review of Non Malignant Referrals to a Homecare Service from January 2008 to December 2009	Denise Hayes	Hayes	Dublin, IE
220	P73	Steroid Prescribing in Oncology Patients in a Regional Cancer Centre	Carolyn	Datta	Glasgow, UK
221	P74	Audit of Laxative Use in Patients on Regular Strong Opioids	Brenda	O'Connor	Cork, IE
222	P75	The Economic Balance of a Palliative Care Unit in Garbagnate Milanese, Italy: Home Care Costs	Furio	Zucco	Garbagnate Milanese, IT
223	P76	An Audit of Care of Patients in the Last Days/Hours of Life Using the Liverpool Care Pathway (LCP) to Measure Care Practice	Melanie	Rosúa Rodriguez	Malaga, ES
224	P77	Monthly Costs of Supportive Cancer Care	Sharon	Beatty	Drogheda, IE
225	P78	Survey of the Development of Hospice and Palliative Care in Austria from 2006 to 2008	Leena	Pelttari	Vienna, AT
226	P79	Development of Nine Quality Indicators for the National Danish Palliative Care Database (DPD): Literature Review and Consensus Process	Mogens	Groenvold	Copenhagen, DK
227	P80	An Audit of the Management of Malignant Bowel Obstruction in a Tertiary Referral Cancer Centre	Nick	Gough	London, UK
228	P81	ESAS symptom Profile of Advanced Cancer Patients Referred to a Palliative Care Consult Team in a University Hospital and Evolution in the First Week of Attention	Ana	de Santiago	Madrid, ES
229	P82	Results from a Facility-led Full Clinical Audit Cycle Conducted in 5 Palliative Care Centres in South Africa and Uganda: Evidence of Improved Patient and Family Outcomes	Richard	Harding	London, UK
230	P83	Admissions to the Specialist Palliative Care Unit: What Does the Documentation Tell Us?	Aoife	Gleeson	Dublin, IE
231	P84	Doctor and Nurse Perspectives on End-of-Life Care in Irish Hospitals	David	Clark	Dumfries, UK

End of life care and quality of death

232	P85	Current Practice in the Management of Terminal Haemorrhage by Palliative Care Teams in the UK	Simon	Noble	Swansea, UK
233	P86	Unsuitable for ICU – What Happens Next?	Alistair	Mckeown	Glasgow, UK
234	P87	Relationship between End-of-Life Decisions and Care Provided in the Final Three Months of Life: The Nationwide SENTI-MELC Study in Belgium	Lieve	Van den Block	Brussels, BE
235	P88	Measuring Patients' Experiences with Palliative Care: The CQ-index Palliative Care	Susanne Jaquelin Johannes	Claessen	Amsterdam, NL
236	P89	How and Why Use a Subcutaneous Needle in End of Life Care?	Brita	Jørgensen	Aarhus, DK
237	P90	Proportion of Incurable Cancer Patients who Receive Palliative Chemotherapy during the Last Month of Life	Bertil	Axelsson	Östersund, SE
238	P91	Non-pharmacological Care-giving Activities: A Multi-national and Multi-professional Inventory of what Is Done in the Last Days of a Patient's Life	Carol Carl Johan	Tishelman Fürst	Stockholm, SE
239	P92	Strong Opioids' Consumption before and after the Establishment of Palliative Care Service in an Egyptian Cancer Center	Samy	Alsirafy	Cairo, EG

240	P93	"To Cherish Each Day as It Comes" - Spirituality and Palliative Care in a Cultural Context	Gudlaug Helga	Asgeirsdottir	Kopavogur, IS
241	P94	Radiotherapy (RT) in the Last Month of Life: Factors Predictive of Poor Outcome	Elaine	Wallace	Dublin, IE
242	P95	Causes of Admission to a Palliative Care Inpatient Unit in an Egyptian Cancer Center	Khaled	Galal	Cairo, EG
244	P97	Factors Associated with Home Return in Cancer Patients Hospitalised in an Acute Care Hospital and Referred to a Palliative Care Team	Monica	Escher	Geneva, CH
245	P98	Artificial Nutrition and Hydration in Last Week of Life of Cancer Patients: Practices, Attitudes and Effects. A Literature Review	Natasja	Raijmakers	Rotterdam, NL
248	P101	Decision-making in Euthanasia and Physician Assisted Suicide, a Qualitative Research	Marianne	Dees	Nijmegen, NL
250	P103	Evaluation of Factors Interfering with the Wishes of Terminally-ill Patients to Receive Home-care rather than In-patient Treatment	Toshihiko	Nakatani	Izumo, JP
251	P104	The Dying Phase: How Is Psychological and Psychosocial Support Understood by Palliative Care Experts across Europe?	Maren	Galushko	Cologne, DE
252	P105	The Dying Phase: Who Is in Need of and who Can Deliver Psychological and Psychosocial Support - The Perspective of Palliative Care Experts in Europe	Maren	Galushko	Cologne, DE
253	P106	Patient Related Factors Considered by Hospice Staff when Recognising the Dying Phase	Richard	Latten	Liverpool, UK
254	P107	"He's Deteriorating": Language Used by Palliative Care Staff Providing Care of Dying Patients	Richard	Latten	Liverpool, UK
255	P108	Transitions to Palliative Care in Acute Hospitals: Findings from a Qualitative Study with Medical and Nursing Staff in the UK	Clare	Gardiner	Sheffield, UK
256	P109	An Audit on the Use of Phenobarbitone for Sedation in Intractable Distress in the Dying	Pauline	Kane	Dublin, IE
257	P110	Exploring the Extent of Communication Surrounding the Transition to Palliative Care in Patients with Heart Failure	Emilie	Green	Sheffield, UK
258	P111	Fear of Death and Dying in Palliative Day Care Patients	Denise	Traue	Cambridge, UK
259	P112	Psico-social Variables in Patients with Psychological Discomfort at Life'End	Rita	Marson	Aviano, IT
262	P115	Evaluation of the Inpatient and Outpatient End-of-Life Care in Germany	Luis Carlos	Escobar Pinzón	Mainz, DE
263	P116	Meeting the Needs of Dying Patients – Reality or Ideal Idea	Iris	Schaefer	Bern, CH
264	P117	Capturing the Invisible: A Survey of Deathbed Phenomena (DBP) in an Irish Hospice	Regina	Mc Quillan	Dublin, IE
266	P119	Maintaining Integrity in the Face of Death	Gillian	Horne	Nottingham, UK
267	P120	Identification and Communication: An Electronic End of Life Pilot Register	Clare	Smith	London, UK
268	P121	Dying behind Bars: A Regional Evaluation of Palliative Care in Prisons in the North West of England	Mary	Turner	Lancaster, UK
269	P122	Dignity for Nursing Home Patients: Design of the Study	Mariska	Vlug	Amsterdam, NL

270	P123	Prospective Study of Medical Emergency Team (MET) Calls to Define Issues of End of Life Decision-making, Symptoms and Transition in Goals of Care	Meera	Agar	Sydney, AU
271	P124	Palliative Sedation for Patients Dying at Home in Belgium: A Descriptive Study	Livia	Anquinet	Jette, BE
272	P125	Implementation in General Practice of a Safety Box for Treating Last Days of Life Symptoms for Patients Dying at Home	Jacob Borgaard	Thellesen	Vejle, DK
273	P126	"I'm Afraid! What Happens?" EPR and Decision Making	Emanuela	Porta	Milan, IT
274	P127	Use of Palliative Care Services and End-of-Life GP Visits in the Netherlands and Belgium	Ebun	Abarshi	Amsterdam, NL
275	P128	Implementation of the Gold Standards Framework and the Liverpool Care Pathway in Nursing Care Homes in Scotland: Qualitative Interviews with Bereaved Relatives and Nursing Care Home Managers	Julie	Watson	Edinburgh, UK
276	P129	Advanced Dementia: What Care Do Patients and Carers Need?	Jacqueline	Crowther	Liverpool, UK
277	P130	Understanding of and Attitudes towards Organ and Tissue Donation amongst Hospice Daycare Patients and Nursing Staff	Dee	Traue	Cambridge, UK
279	P132	Cancer Patients' Care at the End-of-Life in a Critical Care Environment: Perspectives of Families, Patients and Practitioners	Natalie	Pattison	London, UK
280	P133	Evaluation of Needs of Patients and Their Caregivers in a Stationary Hospice in Relation to the Provided Care	Karolina	Pietruk	Warsaw, PL
281	P134	The Development of a Dignity Care Pathway	Bridget	Johnston	Stirling, UK
282	P135	A Comparison of Health Care Expenditure of Cancer Terminal Stage Care between Hospice and Nonhospice Patients	Chia-Chin	Lin	Taipei, TW
283	P136	Dying in Hospital	Anne	MacLennan	Wellington, NZ
284	P137	Is It Feasible to Compare Religion, Spirituality and Spiritual Well-being with Quality of Death?	Rebecca	Evans	Edinburgh, UK
285	P138	How Can Quality of Death (QOD) in a Hospice Be Measured in a Simple Manner?	Rebecca	Evans	Edinburgh, UK
286	P139	The Liverpool Care Pathway for the Dying Patient (LCP): An Audit of its Use in a Hospice Environment and Review of Locality Prescribing Guidelines	Catherine	Rolls	Wirral, UK
287	P140	Does Information about the Imminent Death to a Cancer Patient Influence the Content of Care during the Last Week of Life?	Gunilla	Lundquist	Ludvika, SE
289	P142	Advance Directives for Euthanasia in Dementia: What is Their Feasibility in Practice?	Marika E.	de Boer	Amsterdam, NL
290	P143	Improving Methods for Surveying End of Life Care and the Quality of Death through Cognitive Interviewing	Barbara	Gomes	London, UK
291	P144	Management of End-of-Life Care of Terminally Ill Cancer Patients in Two Different Units within the Same Institution: Palliativists vs Oncologists Attitude	Anna Maria	Cuomo	Pavia, IT
292	P145	Advance Directives – Directive Indeed? An Empirical Study of Advance Care Directives in Medical Decision Making at the End of Life	Pauline	Kouwenhoven	Utrecht, NL

294	P147	The Young Ones: Discharging Young Dying Patients Home to Die	Sue	Haig	Southampton, UK
295	P148	Developments in End of Life Care in Patients with Non Small Cell Lung Cancer; 2002 Compared to 2007	Eline L	Aarden	Groningen, NL
296	P149	Palliative Sedation Therapy: A Limited Systematic Review on Content and Quality of Guidelines	Eva	Schildmann	Berlin, DE
297	P150	Client Biography and a Peaceful Death: Supporting Intra-, Inter-personal and Community Relationships through Volunteer Facilitated Life Stories	Francis	Icasiano	Melbourne, AU
298	P151	PCOC: Introducing X-CAS Casemix Adjusted Score for Benchmarking Pain and Symptoms	David	Currow	Wollongong, AU
299	P152	PCOC: Setting the First Benchmarks for Palliative Care in Australia	David	Currow	Wollongong, AU
300	P153	Communication about Dying in a Dutch Hospital	Erica	Witkamp	Rotterdam, NL
301	P154	Evidence for Dignity Conserving Care at End of Life: A Narrative Review	Ulrika	Östlund	Stirling, UK
302	P155	How Does Access to, and Interaction with, Health Services Support or Undermine 'Living with Distress' in People with Advanced Cancer?	Thomas	Lynch	Lancaster, UK
304	P157	Physicians' Perceptions on Euthanasia in Finland	Maria	Silvoniemi	Turku, FI
305	P158	Dying Women - Gender and Palliative Care	Sigrid	Beyer	Vienna, AT
306	P159	Circumstances at the End of Life in the Brussels Metropolitan Region and the Influence of Specialist Palliative Care Involvement. Results from an Epidemiological Population-based Study	Joachim	Cohen	Brussels, BE
307	P160	Patient Led Case Conferencing at the Interface between Acute and Community Care to Facilitate Advanced Care Planning the End of Life	Clare	Smith	London, UK
308	P161	Do Oncology and Palliative Care Institutions Respond Differently to the Needs of Patients with Lung Cancer in Their Last Month of Life? Preliminary Results from a Multicenter Survey	Bernd	Alt-Epping	Göttingen, DE
309	P162	Advance Care Planning in UK Primary Care: A Feasibility Study	Bruce	Mason	Edinburgh, UK
311	P164	A European Co-ordinating Action to Identify and Drive forward Best Clinical Research Practice in End of Life Cancer Care: PRISMA	Richard	Harding	London, UK
312	P165	Patient Participation in End-of-Life Decision-making: Design of a Study	Linda	Brom	Amsterdam, NL
313	P166	Inequalities in the Prevalence of End-of-Life Decisions in the Past Decade: A Systematic Review	Steven	Gorlé	Jette, BE
314	P167	When Euthanasia Requests Are Refused. Qualitative Interviews with Patients and Physicians about Reasons and Consequences	H. Roeline	Pasman	Amsterdam, NL
315	P168	Medical End-of-Life Decisions: Who Decides?	Natalie	Evans	Barcelona, ES
316	P169	Communication Strategies to Initiate Conversations Planning for End of Life Care (EOLC)	Jane	Seymour	Nottingham, UK
318	P171	Autonomy and Actual Request for Euthanasia	Marianne	Dees	Nijmegen, NL

319	P172	Experiences of End of Life Care in Older People in Care Homes	Katherine	Froggatt	Hatfield, UK
320	P173	Symptoms and Variance Documentation in the Last Days of Life Using the Liverpool Care Pathway in Three Hospital Units in Iceland	Svandis	Halfdanardottir	Kopavogur, IS
321	P174	Dying behind Bars: Scoping the Literature on Palliative Care in Prisons	Sheila	Payne	Lancaster, UK
322	P175	Quality of Life, Emotional Well Being and Functional Status of Patients with Cancer, CHF, and COPD: Does Diagnosis Matter or Is Sick, Sick?	Karen	Steinhauser	Durham, US
323	P176	Unmet Needs in End-of-Life Care Skills among Finnish Physicians	Maria	Silvoniemi	Turku, FI
324	P177	Changes in Patients' End-of-Life Preferences in Course of Time: Design of the Study	Matthijs	van Wijmen	Amsterdam, FI
325	P178	Community Nursing Staff and the Advance Care Planning Tool, 'Preferred Priorities for Care': An Evaluation of Perceptions, Perspectives and Process	Iris	Fineberg	Lancaster, UK
326	P179	Attitude towards Death and Dying in Health Care Professions	Feliks	Blaszczyk	Wroclaw, PL
328	P181	Research at the End of Life - Palliative Care Patients Perspectives	Hermann	Ewald	Kiel, DE
329	P182	Hospice Physician Home Visits	Declan	Walsh	Cleveland, US
330	P183	Advance Care Planning: A Serial Interview Study with Palliative Clinical Nurse Specialists	Deirdra	Sives	Stirlingshire, UK
331	P184	Health Care Assistants and Care of the Dying: An Exploration of Their Role and Experiences	Daniel	Munday	Coventry, UK
332	P185	Advance Care Planning: A Serial Interview Study with Patients and Their Carers	Deirdra	Sives	Denny, UK

Pain

333	P186	Opioid-induced Hyperalgesia - Fiction or Reality?	Uwe	Junker	Remscheid, DE
334	P187	Aspect of the Impact of a New Palliative Care Unit: Opioids	Antimo	De Salve	Mantova, IT
335	P188	Effectiveness of Micro Doses Ketamine in Controlling Neuropathic Pain in Cancer Patients	Nikolay	Yordanov	Vratsa, BG
336	P189	Classification of Neuropathic Cancer Pain in Randomized Controlled Drug Trials: A Systematic Review	Geana	Kurita	Copenhagen, DK
337	P190	The Social Workers Role in Pain Management: A US Study	Debra	Parker Oliver	Columbia, US
338	P191	The Use of Opioids Other than Morphine as First Line to Treat Cancer Pain	Margaret	Clifford	Limerick, IE
339	P192	Opioid Prescription Practices, Patient Education, Educational Needs, and Satisfaction with Pain Management in Icelandic Cancer Patients on Opioids	Valgerdur	Sigurdardottir	Kopavogur, IS
340	P193	Study of Pain in Sarcoma Patients	Cinzia	Brunelli	Milano, IT
341	P194	The Use of Systemic Lidocaine in a Hospice Setting to Manage Complex Pain by Utilising a Titration Protocol	Fiona	McFatter	Dundee, UK
343	P196	Mind-body Medicine: Meditation Used in Cancer and Non Cancer Pain Patients	Paolo	Poli	Pisa, IOT
344	P197	Pharmacokinetic Reliability of a Fentanyl Buccal Soluble Film in Normal Healthy Volunteers	Ignacio	Tagarro	San Fernando de Henares, ES
345	P198	A Pilot Study to Evaluate the Safety and Effectiveness of Co-administration of Teprenone and Captopril for Mucositis in Patients Receiving Concurrent Chemoradiation Therapy for Head and Neck Cancer	Eiko	Yashiro	Chiba, JP

346	P199	Don't Have a Title Yet	John	Smith	Springfield, US
347	P200	Patient-perceived Barriers for Cancer Pain Management between Western and Eastern Cultures - A Systematic Review by Meta-analysis	Chen Hsiu	Chen	Tao-Yuan, TW
348	P201	Pain, Depression and Spirituality in Advanced Cancer Patients	Filipo	Valentini	Torino, JO
349	P202	The Psychological Benefits of Radiotherapy for Cancer-induced Bone Pain	Angela	Scott	Edinburgh, UK
350	P203	Chronic Pain in Cancer Patients in Catalunya: Epidemiology, Appropriateness and Satisfaction with Pain-relief Treatment: Preliminary Data	Josep	Porta-Sales	L'Hospitalet de Llobregat, ES
351	P204	Do Different Opioids for Cancer Pain Differ in Incidences of Nausea and Somnolence during the Commencing Time?	Yoshiaki	Okamoto	Suita, JP
352	P205	Should Palliative Patients be Allowed to Drive?	Thomas	Sitte	Bochum, DE
353	P206	Pilot Study to Test the Feasibility and Pre-test the Efficacy of the German Language Adapted PRO-SELF© Plus Pain Control Program, an Educational Intervention Directed at Patients and Their Family Caregivers to Reduce Cancer Pain and Related Symptoms	Antje	Koller	Basel, CH
354	P207	PCOC: Demonstrating Outcomes in Pain Management in Australia	Deb	Rawlings	Adelaide, AU
355	P208	Fentanyl Buccal Tablet for the Treatment of Breakthrough Cancer Pain – Interim Analysis of a Non-interventional Study Evaluating its Routine Use in Germany and Austria	Hendrik	Mittendorf	Munich, DE
356	P209	Review of Current Morphine Prescribing Guidelines- 'Could Do Better'	Iloro	Finlay	Newport, UK
357	P210	Use of Hydromorphone in Combination with Lorazepam and Their Effect on Ventilation in Palliative Care Patients with Cancer Pain	Katri Elina	Clemens	Bonn, DE
358	P211	The Use of Transdermal Buprenorphine to Alleviate Radiotherapy-related Pain in Head and Neck Cancer Patients	Johan	Menten	Leuven, BE
359	P212	Population Pharmacokinetic Analysis of Transdermal Fentanyl in Japanese Patients with Cancer Pain	Hideya	Kokubun	Kanagawa, JP
360	P213	Epidemiology of Chronic Pain, Treatment Appropriateness and Satisfaction in Ambulant Cancer Patients: Preliminary Results	Josep	Porta-Sales	L'Hospitalet de Llobregat, ES
361	P214	Systematic Review on Alternative Routes for Opioid Application: An EPCRC Opioid Guidelines Project	Lukas	Radbruch	Aachen, DE
362	P215	Investigating the Neural Basis of Pain Processing in Diabetic Peripheral Neuropathy (DPN) and Chemotherapy-induced Peripheral Neuropathy (CIPN)	Elaine	Cachia	Sheffield, UK
363	P216	Differential Contribution of Opioid and Noradrenergic Mechanisms to the Anti-nociceptive and Anti-allodynic Efficacy of Tapentadol in Rat Models of Nociceptive and Neuropathic Pain	Wolfgang	Schroder	Aachen, DE
364	P217	Systematic Review on the Role of Hydromorphone for Moderate to Severe Cancer Pain: An EPCRC Opioid Guidelines Project	Cinzia	Brunelli	Milan, IT
365	P218	Concerning the further Development of Adequate Pain Management in Patients Eligible for Palliative Care in Georgia	Tamari	Rukhadze	Tbilisi, GE
366	P219	A Revision of EAPC and WHO Three Step Analgesic Ladder May Be Needed	Olav Magnus	Fredheim	Trondheim, NO
367	P220	Fentanyl Pectin Nasal Spray Delivers Consistent and Reliable Effects in the Treatment of Breakthrough Cancer Pain: Results from a Long-term Study	Luis	Torres	Cadiz, ES
368	P221	Systematic Review on the Role of Transdermal BUPRENORPHINE (TB) for Moderate to Severe Cancer Pain: An EPCRC Opioid Guidelines Project	Marco	Maltoni	Forli, IT

368B	P221	Systematic Review on the Role of Transdermal Fentanyl (TF) for Moderate to Severe Cancer Pain: An EPCRC Opioid Guidelines Project	Marco	Maltoni	Forli, IT
369	P222	Cognitive Functioning of Pain Patients at the Beginning of Opioid Therapy	Paulina	Andryszak	Bydgoszcz, PL
370	P223	Using the Edmonton Classification System for Cancer Pain - Is there a Pattern amongst the Patients in Our Service who Receive Methadone? Are the Features of Complex Pain Appropriately Assessed and Documented at First Review?	Faith	Cranfield	Brisbane, AU
371	P224	Safety and Tolerability of Fentanyl Pectin Nasal Spray in the Treatment of Breakthrough Cancer Pain: Results from a Long-term Study	Lukas	Radbruch	Aachen, DE
372	P225	Patient Acceptability and Tolerability of Fentanyl Pectin Nasal Spray (FPNS) in the Treatment of Breakthrough Cancer Pain (BTCP): Results from a Double-blind, Double-dummy, Multiple-crossover Study of FPNS versus Immediate-release Morphine Sulphate (IRMS)	Andrew	Davies	Surrey, UK
373	P226	Effect of Morphine and Fentanyl on Innate Immune Function	Jason	Boland	Sheffield, UK
374	P227	Impact of Adverse Effects of Pain Medication on Quality of Life in Myeloma Patients	Jason	Boland	Sheffield, UK
374B	P227	Genetic Variability of the Response to Opioids in Pain Treatment	Silvia	Librada	Merida, ES

Symptoms other than pain

375	P228	Symptomatic Degree of Control in Terminal Ill Patients Based on Specific Diagnoses Skills: ESAS and HADS	Miguel Ángel	Cuervo Pinna	Badajoz, ES
376	P229	Symptom Experience, Palliative Care, and Spiritual Well-being in Patients with Advanced Cancer	Suchira	Get-Kong	Bangkok, TL
377	P230	Suffering: A Study to Explore Palliative Care Nurse Specialists Understanding	Heather	Davies	Bristol, UK
378	P231	Relief of Breathlessness with Nasal Fentanyl in Palliative Home Care - Relation of Single Dosage Compared with the Background Opioid	Thomas	Sitte	Fulda, DE
379	P232	An Evaluation of Supportive Care Needs in Newly Diagnosed Lung Cancer Patients and Lung Cancer Patients within the Last 3 Months of Life	Deans	Buchanan	Dundee, UK
380	P233	Platelet Transfusion in Palliative Care - A Postal Survey of Current Practice	Viv	Lucas	Letchworth Garden City, UK
381	P234	Randomised Phase III Clinical Trial of 5 Different Arms of Treatment for Patients with Cancer-related Anorexia/Cachexia Syndrome (CACS)	Franco	De Conno	Cagliari, IT
382	P235	Professionals Perceptions of the Role of the Dietitian in Fatigue Management within Specialist Palliative Care	Wilhelm	von Hornstein	Co Louth, IE
383	P236	Malignancy Related Bowel Obstruction: Challenging the Convention	Edith	Ubogagu	London, UK
384	P237	Can "Steroid Switching" Improve Steroid-induced Musical Hallucinations in a Terminal Cancer Patient?	Seitetsu	Kanemura	Mino, JP
385	P238	Dispnoea-etiology Factors and Treatment in Patients with Terminal Diseases	Lidija	Veterovska Miljkovic	Skopje, MA
386	P239	A Systematic Review of Drug Management for the Treatment of Cancer Related Fatigue	Ollie	Minton	London, UK
387	P240	Symptomatology Like Possible Determining Factor of the Performance of a Team of Palliative Home Care	Alfredo	Zamora Mur	Huesca, ES
388	P241	Depression Symptoms among Patients with Metastatic Ovarian Cancer in a Programme of Palliative Cancer Care: A Prospective and Cross-sectional Study	Peter	Priester	Hradec Kralove, CH
389	P242	Prognostic Factors in Advanced Cancer Patients: Refinement of Survival Prediction Using Nutritional Variables	Vickie	Baracos	Edmonton, Ca

390	P243	Morphine Treatment of Advanced Breast Cancer Stage in Association with MAP-HD Plus ACTH-Depot: Clinical Evaluation of the Association on Body Weight, Depression, Intensity of Pain and Consumption of Morphine Day	Francesco	Buda	Udine, IT
391	P244	Fatigue in Breast Cancer Patients: Prevalence and Associated Factors	Cibele	Mattos Pimenta	Sao Paulo, BR
392	P245	Is Fatigue Similar among Patients with Colo-rectal, Breast or Prostate Cancer?	Cibele	Mattos Pimenta	Sao Paulo, BR
393	P246	Laxative Use in Patients with Advanced Illness and Opioid-induced Constipation Treated with Subcutaneous Methylnaltrexone	Robert	Israel	Orem, US
394	P247	The Management of Diabetes in Cancer Patients Receiving Palliative Care in a Hospital Setting	Lucy	Balding	Waterford, IE
395	P248	Analysis of Response to Methylnaltrexone by Response to Previous Dose in Patients with Advanced Illness and Opioid-induced Constipation	Robert	Israel	San Diego, US
396	P249	Fatigue during Chemotherapy - Patients' Experiences with the Symptom in Daily Life, with Fatigue Related Self Care Activities and Communication with Health Care Professionals: A Qualitative Study	Elisabeth	Spichiger	Basel, CH
397	P250	Constipation in Palliative Care Patients. Incidence, Risk Factors and the Diagnostic Tools	Tomasz	Dzierzanowski	Warsaw, PL
398	P251	Evaluation of Central Serotonin Sensitivity in Breast Cancer Survivors with Cancer-related Fatigue Syndrome (CRFS)	Susanna	Alexander	London, UK
399	P252	Differences in Fatigue Experiences between Advanced Cancer Patients and Cancer Survivors	Johan	De Raaf	Rotterdam, NL
400	P253	Opioids for Dyspnoea in Chronic Heart Failure	Stephen	Oxberry	Leeds, UK
401	P254	Dyspnoea Descriptors in Chronic Heart Failure (CHF): Changes with Opioid Therapy	Stephen	Oxberry	Leeds, UK
402	P255	Making Exercise Count - The Management of Fatigue in Palliative Care Patients	Graham	Whyte	Liverpool, UK
403	P256	Do Co-morbidity and Symptoms' Intensity Influence Survival Prognosis?	Maria	Nabal	Lleida, ES
404	P257	Randomized, Double-blind, Placebo-controlled Trial of Docusate in the Management of Constipation in Patients in Palliative Care Units	Yoko	Tarumi	Edmonton, CA
405	P258	A Pilot Randomised Controlled Trial of a Customised Rehabilitation Intervention for Managing Fatigue in Patients with Advanced Cancer	Gail	Eva	Oxford, UK
406	P259	The Delirium in a Service of Palliative Care	Alfredo	Zamora	Huesca, ES
407	P260	Multicenter Randomized Double-blinded Placebo-controlled Trial of Methylphenidate for Depressive Symptoms in Advanced Cancer Patients	Carlos	Centeno	Pamplona, ES
408	P261	Mental Symptoms in Patients with Cancer from General Practice	Florien	van Heest	Odoornerveen, NL
410	P263	Palliative Care Emergency Kit: Our First Results	Antonio	Ortega Morell	Antequera, ES
411	P264	Palliative Care Patients' Perspectives of Factors with Positive Impact on Mood in Facing Terminal Illness	Franca	Warmenhoven	Nijmegen, NL
413	P266	A Systematic Scoping Review of the Range and Content of Rehabilitation Interventions to Optimise Physical Function in Adults with Metastatic Spinal Cord Compression	Gail	Eva	Oxford, UK
416	P269	OPCARE9. Signs and Symptoms of Approaching Death	Susanne	Linder	St.Gallen, CH
417	P270	Indicating Anticoagulant Therapy in Palliative Care - A Multicenter Survey	Bernd	Alt-Epping	Göttingen, DE
418	P271	The Mediation Effect of Anxiety on the Relationship between Posttraumatic Stress Symptoms and Preparatory Grief in Advanced Cancer Patients	Kyriaki	Mystakidou	Athens, GR

419	P272	Prevalence and Severity of Opioid Induced Bowel Dysfunction Symptoms and Differences among Strong Opioids. DIO Study	Jaume	Canal	Lérida, ES
421	P274	Factors Associated with the Perception of Constipation in Advanced Cancer Patients: Preliminary Results	Anna	Libran	Terrassa, ES
422	P275	Decision-support for Diagnosing Depression through Case-based Reasoning	Odd Erik	Gundersen	Trondheim, NO
423	P276	Methylnaltrexone for Opioid-induced Constipation: Clinical Experience at a Tertiary Cancer Centre	Jennifer	Dutka	Edmonton, CA
424	P277	Breast Cancer Patients Receiving Postoperative Radiotherapy: Distress, Depressive Symptoms and Unmet Needs of Psychosocial Support	Eeva	Salminen	Turku, FI
425	P278	Pharmacological Interventions for Pruritus in Adult Palliative Care Patients	Carola	Xander	Freiburg, DE
426	P279	Prevalence and Determinants of Loss of Dignity among Danish Patients Receiving Palliative Care	Lise	Jul Houmann	Copenhagen , DK
427	P280	Fatigue and its Relationship with Other Quality of Life Markers in Terminally Ill Cancer Patients in Kenya	Zzipporah	Ali	Nairobi
428	P281	Cancer Cachexia (CC) in Dependence of Tumour Behaviour under Anticancer Treatment: A Literature Review of Advanced Pancreatic, Non-small Cell Lung and Prostate Cancer	Florian	Strasser	St. Gallen, CH
429	P282	Prospective Study about Depression in Oncology and Assessment of the Medical Decision Making's Strategy	Wadih	Rhondali	Lyon, FR

Set 2

Setup: Friday, 14.30 – 15.00

Removal: Saturday, 15.30 – 15.45

Basic science and translational research

430	P1	Chemical Compatibility/Stability of Common Drug Combinations Administered by Continuous Subcutaneous Infusions for End of Life Care	Andrew	Dickman	Liverpool, UK
431	P2	1st Contact: Video Interview Method in Advance Care Planning Research	Sarah	Russell	Berkhamsted, UK
432	P3	The Effect of Complementary Therapies on Multidisciplinary Teamwork in Palliative Care	Claudia	Wenzel	Vienna, AU

Bereavement

433	P4	Bereavement's Myths in Students at the End of Master Course in Palliative Care versus Other Masters Courses	Manuel	Capelas	Lisboa, PT
434	P5	Supporting Parentally Bereaved Children and Their Families: Findings from a Narrative Study	Jacqueline	Ellis	Liverpool, UK
435	P6	Did You Talk about what Was Important with Your Dying Parent? Feelings of Guilt in Bereaved Teenage Daughters	Tove	Bylund Grenklo	Stockholm, SE

Education

436	P7	Influences on GP Decision Making for Choice of CME Sessions in Palliative Care	Peter	Pype	Antwerp, BE
-----	----	--	-------	------	-------------

437	P8	Lessons to Learn: Research about Teaching in Palliative Medicine	Stephanie	Stiel	Aachen, DE
438	P9	Physicians' Knowledge of Pain Management and the Use of Opioids at the End of Life	C.A.	Rhodius	Amsterdam, NL
439	P10	The Information, Myth, Believes that the Patients and the Community Has on Cervical Cancer	Stella	Rithara	Nairobi, KE
441	P12	How Do Newly Qualified Doctors Learn to Care for Patients who Are Dying?	Jane	Gibbins	Bristol, UK
442	P13	Why Do Newly Qualified Doctors Feel Unprepared to Care for Patients at the End of Life?	Jane	Gibbins	Bristol, UK
443	P14	The Other Side of the Gate: An Evaluation of 'Opening the Spiritual Gate' Programme	Kathryn	Gaunt	Southport, Merseyside, UK
444	P15	A Palliative Care Education Course for Professionals Coming from Developing Countries Engaged in a New Italian Hospice	Filippo	Valentini	Torino, IT
445	P16	Working with Older People to Develop a Training Programme for Volunteer Peer Educators in End of Life Care Education	Jane	Seymour	Nottingham, UK
446	P17	Evaluation of the Acceptability of an Electronic Information Programme (DVD) Concerning Cancer Associated Weight Loss and Fatigue to Patients with Lung Cancer	Max	Watson	Belfast, UK
447	P18	Skill Development: Involving Patients in Decisions about Their Care	Margaret	O'Connor	Frankston, AU
448	P19	How to Teach and Learn Core Attitudes in Palliative Care?	Christina	Ramsenthaler	Oldenburg, DE
449	P20	Inventory of Education in Palliative Care; A Signal Report!	Marieke	Groot	Nijmegen, NL
450	P21	Education Changed My Life: Experiences of Learners from a Foundation Degree in Palliative Care	Jane	Wale	Leicester, UK
451	P22	"Ways of Seeing" - Views on the Most Important Elements in the Patient and Support Worker Relationship in Palliative Care	Jane	Wale	Leicester, UK
452	P23	Translating Results of Research to Practice. The Activities of the Scientific Committee of the V&VN Dutch Nurses' Association; Department Palliative Nursing in the Netherlands	Madeleen	Uitdehaag	Utrecht, NL
453	P24	What Factors Affect the Likelihood of "Success" in Online Study?	Deb David	Rawlings Currow	Adelaide, AU
454	P25	An Evaluation of a Pilot Introductory End of Life Education Programme for Staff Working in Residential Care Settings for Older People in the Republic of Ireland	Orla	Keegan	Dublin, IE
455	P26	A Palliative Care Approach: Education and Training for Health Care Assistants	Orla	Keegan	Dublin, IE
456	P27	The Patient as a Teacher – The Meaning of Death and Dying for Medical Students	Martina	Pestinger	Aachen, DE
457	P28	Symptoms Control in Palliative Care: Assessment of Two Different Teaching Methodologies	António	Barbosa	Lisboa, PT
458	P29	Evaluation of Palliative Care Teaching in the Undergraduate Medical Curriculum	Joanna	Bowden	Edinburgh, UK
460	P31	Development of a Competency-based Curriculum in Neurologic Palliative Care	Michel	Beauverd	Lausanne, CH
461	P32	Public Attitudes to Death, Dying and Bereavement: A Synthesis of the Research Literature	Jane	Seymour	Nottingham, UK

462	P33	The EPOS Study: Experiences in the Statistical Analysis of Genetic Association	Peter	Fayers	Aberdeen, UK
463	P34	What Makes Informed Consent Documents User-friendly?	Kari	Sand	Trondheim, NO

Epidemiology

464	P35	A Prospective Comparative Survey of Patient Characteristics and Treatment Outcomes in a Palliative Care and in an Oncology Unit	Bernd Oliver	Maier	Wiesbaden, DE
465	P36	Description of a Tertiary Swiss University Hospital Palliative Population Based on the International Classification of Disease (ICD): A Retrospective Pilot Study	Michel	Beauverd	Lausanne, CH
466	P37	Survey on In-hospital Palliative Patients: Prevalence and Characteristics (Part I)	Paul	Vanden Bergh	Bruxelles, BE
467	P38	Survey on In-hospital Palliative Patients: Treatment Strategies (Part II)	Paul	Vanden Bergh	Bruxelles, BE
468	P39	Survey on In-hospital Palliative Patients: Hospital Resources Use and Discharge Management (Part III)	Paul	Vanden Bergh	Bruxelles, BE
469	P40	Description of a Patients' Transverse Cut Attended by Barbastro'S Home Care Support Team	Alfredo	Zamora	Huesca, ES
470	P41	Amyotrophic Lateral Sclerosis Gender Differences	M. Teresa	Salas	Madrid, ES
471	P42	Anti-cholinergic Load, Health Care Utilisation and Survival in People with Advanced Cancer. A Pilot Study	Meera	Agar	Adelaide, AU
472	P43	Death and the City. A Population Based Study of End-of-Life Decision-making in the Brussels Metropolitan Region and Non-metropolitan Flanders	Joachim	Cohen	Brussels, BE
473	P44	Trends in Medical End-of-Life Practices in Belgium, 1998-2001-2007	Johan	Bilsen	Brussel, BE
474	P45	Characteristics of Patients with Prostate Cancer Enrolled in Swedish Palliative Care	Staffan	Lundström	Stockholm, SE
475	P46	Vitamin D Insufficiency in Cancer: The Tip of the Iceberg?	Declan	Walsh	Cleveland, US
476	P47	Caregiving Trajectories Mirror Different Patterns of Functional Decline by Diagnosis	David	Currow	Adelaide, AU
477	P48	Prescription of Antidepressants to Cancer Patients in Their Last Year of Life	Siri	Brelin	Oslo, NO
478	P49	Illness Attribution in Advanced Disease	L Willi	Rayner Lee	London, UK
479	P50	Epidemiology of Depression in Palliative Care	Willi	Lee	London, UK
480	P51	The Association between Physical Activity and Health Related Quality of Life. A Population Based Study	Gro F.	Bertheussen	Trondheim, NO
481	P52	Cancer Patients Account for Only 3% of Patients Receiving Opioids in Norway	Olav Magnus	Fredheim	Trondheim, NO

Ethics

483	P54	Ethical and Methodological Issues in End of Life Care Research with People from Minority Cultures	Margaret	O'Connor	Melbourne, AU
-----	-----	---	----------	----------	---------------

484	P55	Characteristics of Granted and Rejected Euthanasia Requests in Belgium	Yanna	Van Wesemael	Jette, BE
486	P57	(Bio)Ethical Aspects of Palliative Care	Tomislav	Cengic	Rijeka, HR
487	P58	Implementation of Ethics Support in Nursing Homes and Primary Care – What Does Health Personnel Want?	Georg	Bollig	Bergen, NO
488	P59	A Study on Moral Distress among Health Care Workers Providing Home-based Palliative Care	Kevin	Brazil	Hamilton, CA
489	P60	Deep Continuous Sedation as a Preferable Alternative to Physician-assisted Death? Some Ethical Thoughts	Kasper	Raus	Ghent, BE
490	P61	Euthanasia and Physician Assisted Suicide in the Polish Sixth Year Medical Students Views	Aleksandra	Kotlinska-Lemieszek	Poznan, PL
491	P62	Research among Bereaved Relatives of Deceased Cancer Patients: Is it Feasible and Acceptable?	Jonathan	Koffman	London, UK
492	P63	I Don't Want to Die Like Daddy. Impact of Motivations on the Concept of an Individual Advance Directive – A Qualitative Analysis	Birgit	Jaspers	Goettingen, DE
493	P64	Review of Oregon's Official Annual Reports on its Death with Dignity Act	Ilorá	Finlay	Cardiff, UK

Family and care givers

494	P65	Capturing Caregivers' Views: Challenges and Strategies Used in Undertaking Research with Caregivers of Patients with Advanced Disease	Farida	Malik	London, UK
495	P66	Family Experiences of Having a Person with Dementia Nearing the End of Life in a Care Home: A Systematic Review	Jean	Hennings	Lancaster, UK
496	P67	How Many Dying Patients Can a Team Cope with? A Survey of Hospices in Germany Compared with Data from Palliative Care Units	David	Pfister	Aachen, DE
497	P68	Predictors of Caregiver Burden in a Palliative Care Randomized Trial to Improve Patients' Quality of Life	Marie	Bakitas	Lebanon, US
498	P69	Communication between Hospice Teams and Informal Caregivers during Team Meetings	Elaine	Wittenberg-Lyles	Denton, US
499	P70	Caregiver Participation in Hospice Team Meetings: A Pilot Study of the ACTIVE Intervention	Debra	Parker Oliver	Seattle, US
500	P71	The Impact of Breast Cancer in a Patient and the Community at Large	Stella	Rithara	Nairobi, KE
501	P72	Family Meetings in Specialist Palliative Care Units: Are they Effective?	Breffni	Hannon	Dublin, IE
502	P73	The Carers' Drama in Home Palliative Care	Jenny	Newbury	Bradford-on-Avon, UK
504	P75	The Significance of Fatigue in Relatives of Palliative Patients	Maria	Carlsson	Uppsala, SE
505	P76	Main Fears of Caregivers of Patients Treated by Palliative Care Teams in Extremadura	Victoria	Romero	Cáceres, ES
506	P77	Indicators of Ability to Care, in Families of Cancer Patients Receiving Palliative Care	Carla	Reigada	Porto, PT
507	P78	The Experience of the Experimental Process of Education of the Care Giver in Palliative Home Care	Antimo	De Salve	Mantova, IT

508	P79	Reconciling Employment with Caring for a Husband with an Advanced Illness	Marjolein	Gysels	London, UK
509	P80	Informal Caregivers Needs of People at the End of Their Life at Home: Supervision Strategies	Maria João	Teixeira	Aveiro, PT
510	P81	Quality of Palliative Care through the Relatives' Eyes: Findings of the CQ-index Palliative Care	Anneke	Francke	Utrecht, NL
511	P82	Including Carer Effects in Economic Evaluations: Willingness to Care, a Missing Link?	Nikki	Mc Caffrey	Adelaide, AU
512	P83	Making Words Count: Readability Issues for Palliative Care Consumers	Deb	Rawlings	Adelaide, AU
513	P84	The Ultimate Measure of the Quality of Health Care Is the Outcomes that Patients and Carers Achieve	Deb	Rawlings	Adelaide, AU
514	P85	Modelling the Carer Support Component of a Complex Intervention for Breathlessness in Advanced Disease	Morag	Farquhar	Cambridge, UK
514B	P85B	Support, Loneliness and Well-being amongst Very Old People in Their Last Year of Life	Morag	Farquhar	Cambridge, UK
515	P86	Mental Health in Significant Others of Patients Dying from Lung Cancer	Ulrika	Östlund	Stockholm, SE
517	P88	Caregivers' Anxiety and Self Efficacy in Palliative Care	Kyriaki	Mystakidou	Athens, GR
518	P89	Delivering Feasible and Acceptable Palliative Care Interventions Targeted at Informal Home Caregivers: How Should We Proceed? Findings from an Intervention Development Study	Richard	Harding	London, UK
519	P90	Which Themes Are Relevant when Assessing the Relatives of Cancer Patients' Situation and Their Perception of Health Care?	Line	Lund	Copenhagen, DK
520	P91	Family Conferences – How Do We Use It?	Ana	Querido	Leiria, PT
521	P92	The Context of Respite Care	Virginia	Dunn	Oxford, UK
523	P94	Quality of Life in Hospice Caregivers Involved in a Telephone Counseling Program (Support TECH)	Jean	Kutner	Aurora, US
524	P95	Spiritual Pain as an Expression of Suffering in Advanced Cancer Patients' Caregivers in the Palliative Care Setting	Marvin	Delgado Guay	Houston, US

Medical sociology

525	P96	Patient Barriers to Cancer Pain Management in Denmark and Lithuania	Ramune	Jacobsen	Copenhagen, DK
526	P97	Continuous Deep Sedation at the End of Life: The 'Natural' Hypothesis	Kasper	Raus	Ghent, BE
528	P99	Deprivation Scores and Access to Specialist Palliative Care Services in Cancer Patients in Dundee	Claire	Douglas	Dundee, UK
529	P100	A National Wide Survey of Staffing Power of Hospice/Palliative Care Unit in Japan	Paul	Abe	Chiba, JP
531	P102	Participant Observation of Decision Making over the Last Six Months of Life for Patients with Advanced Cancer in Greece	Despina	Anagnostou	London, UK
532	103	A Missing Link? The Process of Recognizing Dying and its Nature in Patient Records	Sabine	Pleschberger	Vienna, AT

Non-cancer

533	P104	Symptom Prevalence of Patients with Chronic Obstructive Pulmonary Disease (COPD) who Access Palliative Care in Salford	Catherine	Hayle	Salford, UK
534	P105	Different Patterns of Death in a Palliative Care Unit among End-stage Non-oncological and Advanced Cancer Patients	Jose	Garcia-Garcia	Alcalá Guadaira, ES
535	P106	Use of the Liverpool Care Pathway in Patients Dying with Advanced Chronic Kidney Disease: A Survey of Hospital Renal Services in England	Sarah	Wenham	Salford, UK
536	P107	Developing Effective Patient-centred Palliative Care for Patients with Advanced Progressive Non-malignant Disease and Their Carers: A Qualitative Study Exploring the Views of Patients, Carers and Health Care Professionals	Fiona	Paterson	Dundee, UK
537	P108	Managing Medical Crises: The Experiences of People Affected by Motor Neurone Disease (MND) and Their Carers	Jacqueline	Hill	Clay Cross, UK
538	P109	Does Regular Palliative Care Consultation Improve Symptom Burden in Patients with Advanced Chronic Kidney Disease (CKD) Attending a Renal Conservative Management Service?	Claire	Douglas	Dundee, UK
539	P110	Palliative Care for People Living with HIV/AIDS in Uganda: An Investigation of Patients & Caregivers' Outcomes and Professional Perspectives	Wesley	Too	Nottingham, UK
540	P111	Increased Non-cancer Admissions to Hospice Palliative Care Units – Will this Hurt Cancer Care?	Rebekah	Gilbert	Edmonton, CA
541	P112	“Something Bubbling in my Blood”: The Experience of Advanced Chronic Kidney Disease as a Hidden Illness	Fliss	Murtagh	London, UK
542	P113	End of Life Care in Patients on Renal Replacement Therapy: A Retrospective Review of Deaths over a 5 Year Period	Bernadette	Brady	Dublin, IE
543	P114	Palliative Care for Renal Patients - The First Three Years	Jeena	Ackroyd	Leeds, West Yorkshire, UK
544	P115	Between Chronic and Advanced: Can We Find the Difference?	Deb David	Rawlings Currow	Adelaide, AU
545	P116	Shoulder Pain in Motor Neuron Disease: An Under Appreciated Problem	David	Currow	Adelaide, AU
546	P117	Dialysis Patients Are a Palliative Care Population: Results of the DIA-PAIN Study	Claudia	Gamondi	Bellinzona, CH
547	P118	The Experience of Withdrawing from Dialysis: Perceptions of Bereaved Carers	Fliss	Murtagh	London, UK
548	P119	Quality of Care for Patients with Idiopathic Pulmonary Fibrosis – Perspectives from Patients and Carers	Catriona	Killin	Cardiff, UK
549	P120	Unknown Group of Movement Disorders with Palliative Care Needs – Parkinson's Plus Syndromes: Multiple System Atrophy (MSA) and Progressive Supranuclear Palsy (PSP)	Tariq	Saleem	London, UK
550	P121	Quality of Life in Patients with Beta-thalassemia. A 2001-2009 Repeated Cross Sectional Survey	Massimo	Costantini	Genoa, IT

551	P122	Measuring Non Malignant Chronic Pain Interference in the Elderly	Ana	Querido	Leiria, PT
552	P123	Defining the Role of Specialist Palliative Care in the Management of Motor Neurone Disease: A Retrospective Review of Hospice Involvement in Motor Neurone Disease over a Ten Year Period	Richella	Ryan	Dublin, IE
553	P124	Pattern of Non-cancer Referral to Palliative Care in a Saudi Tertiary Care Hospital	Samy	Alsirafy	Cairo, EG

Organisation of services

555	P126	Palliative Care in Canada: The Economic Perspective for Families and Health Care System	Serge	Dumont	Quebec City, CA
556	P127	A Ten Years Experience Following Cancer Patients in Home-care: Organisation of the Service	Mario	Ranuzzi	Monterotondo, IT
557	P128	Ambulance Based Palliative Care in Northern Thuringia - 2 Years of Experiences	Heike	Buentzel	Nordhausen, DE
558	P129	Palliative Home Care Teams in Portugal: How Many and where We Need	Manuel	Capelas	Lisboa, PT
559	P130	Transitions in the Last Year of Life to Access Health Care	Donna	Wilson	Edmonton, CA
560	P131	What is the Scope of Information Available on Hospice Websites?	Grace	Yang	Ipswich, UK
561	P132	An Exploration of the Experience of Patients who Are Discharged from Community Specialist Palliative Care (CSPC)	Miranda	Stacpoole	London, UK
562	P133	A Retrospective Review of the Outcome of Patients Discharged Following an Inpatient Hospice Stay	Niamh	O'Connor	Dublin, IE
563	P134	Extending Palliative Care to Non Cancer Patients: Challenges and Resources	Simone	Veronese	Torino, IT
564	P135	Possible Determining Factors of the Action of an Equipment of Support for Domiciliary Attention	Alfredo	Zamora Mur	Huesca, ES
565	P136	“This Huge Feeling as if Arms Are Going Round You to Protect You and Comfort You” – A Sense of Sanctuary Deriving from the Ethos and Atmosphere of a Cancer Support Service	Zoe	Cockshott	Lancaster, UK
566	P137	Physiotherapist in a Team of Palliative Care: Multidisciplinary	Antimo	De Salve	Mantova, IT
567	P138	Pain and Symptom Control and Palliative Radiotherapy Consultation by Telehealth for Rural Cancer Patients: Symptom, Cost and Satisfaction Outcomes	Sharon	Watanabe	Edmonton, CA
569	P140	Palliative Care Consults and Outcomes for Patients with Cerebral Infarct or Intracranial Hemorrhage	Hallie	Mason	Akron, US
570	P141	Palliative Care Consults and Outcomes for Hospital Patients with Renal Failure	Charina	Gayomali	Akron, US
571	P142	A Novel Program of Mental Health Service Delivery for Palliative Patients in a Hospice Setting: The Continuing Care Psychiatric Consulting Service (CCPCS)	William	Chimich	Edmonton, CA
572	P143	The Project of the Palliative Care in the Public Health Department of the Primorsko-Goranska County	Morana	Brkljacic Zagrovic	Rijeka, HR

573	P144	Expert Consultation in Palliative Care in South-west Netherlands	Agnes	van der Heide	Rotterdam, NL
574	P145	A Descriptive Analysis of Referrals to a Cancer Unit Specialist Palliative Care Team from December 2001- March 2009 during the Establishment on the N Ireland Cancer Network	Claire	Daly	Londonderry, UK
575	P146	Clinical and Organisational Outcomes of Palliative Care Hospital Support Teams, in Spain	Josep	Porta-Sales	L'Hospitalet, ES
576	P147	Negotiating the Boundary between Paid and Unpaid Hospice Workers: A Qualitative Study of how Hospice Volunteers Understand Their Work	Sarah	Field-Richards	Nottingham, UK
577	P148	A Cross Sectional Audit of Palliative Care Needs in the Acute Hospital Setting	Timothy	To	Daw Park, AU
578	P149	How Stable Are Preferences for Place of Death? Findings from a Pilot Study	Irene	Higginson	London, UK
579	P150	Development of Palliative Medicine in Russia in XXI a Century - The Organisation of the Palliative Help on Federal and Regional Levels	Tofik	Biktimirov	Ulyanovsk, RU
580	P151	Hospital Use Near Death in Alberta, Canada	Donna	Wilson	Edmonton, CA
581	P152	A Corporate Assessment of the Palliative Care Needs of People with Intellectual Disability	Karen	Ryan	Dublin, IE
582	P153	Telephone Consultations: Reasons for Consultation and Strategies Developed for Problem Solving	Cátia	Ferreira	Porto, PT
584	P155	A Randomised Pilot of Academic Detailing in General Practitioners for Palliative Dyspnoea Management	David	Currow	Adelaide, AU
585	P156	Transitions between Care Settings in the Last Year of Life for People Living with Heart Failure, Stroke and Lung Cancer: A Qualitative Study	Louise	Holmes	Liverpool, UK
586	P157	Living Alone with Cancer at the End of Life: An Exploration of the Experiences of Older Adults	Lucy	Cooper	Liverpool, UK
587	P158	Hospices Network in Italy: An Unpredictable Growth	Furio	Zucco	Garbagnate Milanese, IT
588	P159	The Development of Quality Indicators in Palliative Care: European Panel on Indicators of Best Practice in the Organisation of Palliative Care in Europe: A 2-round Delphi	Karen	Van Beek	Leuven, BE
589	P160	A Descriptive Study of the Role of Pastoral Care/Chaplaincy in Specialist Palliative Care Services in Ireland	Ursula Michael	Bates Bramwell	Drogheda, IE
590	P161	Organizing Methodology to Improve the Accessibility of Palliative Care Unit, Hospice and Home Care of Mantova: The Case Management	Antimo	De Salve	Mantova, IT
591	P162	Establishing a National Framework for Coordination of Palliative Care	Mai-Britt	Guldin	Aarhus, DK
592	P163	Hospital Palliative Care Supportive Team and Palliative Care Unit See Different Patient Profile	Joaquim	Julià	Badalona, ES
593	P164	Support Staff in British Hospices: Invisible Workers?	Mary	Turner	Lancaster, UK
594	P165	Relaxation Outpatient Clinic – Prospective Evaluation	Jenny	Gilbert	Stirling, UK
595	P166	Supportive and Palliative Care in England: Differences between the Ten Strategic Health Authorities	Bill	Noble	Sheffield, UK

596	P167	Different Approaches to Care for the Terminally Ill – A Four Country Comparison	Christopher	Klinger	Toronto, CA
597	P168	The Impact of Variations in out of Hours End of Life Care Provision on Patient Experience, Staff and Health Systems: A New UK National Study	Julia	Addington-Hall	Southampton, UK
598	P169	Dynamic Evaluation of All Services to Inform Policy Development	Ilorá	Finlay	Cardiff, UK
599	P170	Impact of a Palliative Day Care Unit on Advanced Cancer Patient Life's Journey	Georges	Czapiuk	Paris, FR
600	P171	Palliativnetz Berlin - A Regional Network for Palliative Care in Berlin	H. Christof	Müller-Busch	Berlin, DE
601	P172	Preferences and Choices for Place of Death among Patients with Advanced Cancer in Greece: Findings from a Longitudinal Qualitative Study	Despina	Anagnostou	London, UK
602	P173	Hospice Care Facilities in the East of Netherland; Organization, Supply and Use of Care	Marieke	Groot	Nijmegen, NL
604	P175	Implementing 7 Day Working by Clinical Nurse Specialists across Rural and Urban Areas	Ilorá	Finlay	Cardiff, UK

Palliative care in children and adolescents

605	P176	Evaluating Clinical Competency Following Multinational Association of Supportive Care in Cancer [MASCC] Involvement Nairobi-Hospice	Stella	Rithara	Nairobi, KE
606	P177	Physicians' Attitudes and Actual Practices Concerning Physician-assisted Dying in Minors: A Nationwide Survey	Geert	Pousset	Ghent, BE
607	P178	Novel Use of Pregabalin for Neuropathic Pain in Paediatrics	Sinead	Kelly	Dublin, IE
608	P179	Pain in Children with Cancer at the End of Life: A Review	Anne-Sophie	Darlington	Rotterdam, NL
609	P180	e-KernPäP – An Innovative Web Based Documentation System for the Specialized Ambulatory Pediatric Palliative Care in Germany	Margit	Baumann-Köhler	Münster, DE
610	P181	Pain in Children at the End of Life and Parents' Hope	Anne-Sophie	Darlington	Rotterdam, NL
612	P183	My Space or Social Space – Navigating the Rapid Discharge of Teenagers and Young People Home to Die	Sue	Duke	Southampton, UK

Palliative care in elderly

613	P184	How Does Palliative Care Training Influence Attitude, Stability and Fear of Nursing Staff in Homes for the Elderly?	Hans-Ulrich	Giesen	Duisburg, DE
614	P185	Improving End of Life Care in Long Term Care Facilities: Perspectives of Healthcare Providers and Bereaved Family Caregivers	Christine	Beck	Halifax, CA
615	P186	Illuminating Carer's Understanding of End of Life Care in Care of the Elderly Community Units	Una	Molloy	Dublin, IE
616	P187	PRISMA WP6: Best Practice in Long Term Care Facilities in Europe	Gwenda	Albers	Amsterdam, NL

618	P189	Communication Skills Training for Health Care Professionals and Family Carers in Dementia Care. A Systematic Review	Eva	Eggenberger	Vienna, AT
619	P190	First Steps towards Validation of a Symptom Scale for Palliative Elderly	Ruth	Piers	Ghent, BE
620	P191	How Can Do Not Resuscitate Orders and Advance Directives Be Implemented in Nursing Homes?	Georg	Bollig	Bergen, NO
621	P192	A Validated Risk Score to Estimate Mortality Risk in Patients with Dementia and Pneumonia: Barriers to Clinical Impact	Jenny	van der Steen	Amsterdam, NL
622	P193	Dying at Home: Connection between Primary Health Care and Palliative Care: A Necessity	Rosa Maria	Jauma Pou	Banyoles, ES
623	P194	Palliative Care in the Oldest of Old in the Dutch Community: A Nationwide Study	Ebun	Abarshi	Amsterdam, NL
625	P196	Acupuncture is a Useful Adjunct to Medical Treatment for Painful Elderly. A Pilot Study	Marie	Couilliot	Bobigny, FR
626	P197	Gender-sensitive End of Life Care in Long Term Care Facilities for Frail Elderly	Elisabeth	Reitingner	Vienna, AT
627	P198	Guidelines for a Palliative Approach for Aged Care in the Community Setting	David Deb	Currow Rawlings	Adelaide, AU
628	P199	Standards of Quality for Hospice and Palliative Care in Austrian Nursing Homes	Sigrid	Beyer	Vienna, AU
629	P200	Do Opioids Have an Effect on Survival in the Elderly Oncologic Hospice Population?	Johan	Menten	Leuven, BE
631	P202	The Economic Background of Dementia	Médea	Kui	Budapest, HU
632	P203	Aging People and Palliative Care in São Paulo-Brazil: Preliminary Report	Angélica M	Yamaguchi	São Paulo, BR
633	P204	First Interviews: Aging People and Palliative Care in São Paulo-Brazil	Angélica	Yamaguchi	São Paulo, BR

Policy

634	P205	Public Health Initiatives to Improve Palliative Care in Germany	Nils	Schneider	Hannover, DE
635	P206	Policies to Improve End-of-Life Decisions in Flemish Residential Care Facilities Accommodating Persons with Intellectual Disabilities: Prevalence, Content and Implementation Strategies	Luc	Deliens	Brussels, BE
636	P207	The Relation between Hospice and Health Care System. A Comparative Contemporary History of the Developments in England and the Netherlands	Aalbertha	van Reuler	Manchester, UK
637	P208	Palliative and Hospice Care in Russia: A New Approach to Improve Public Health Policy	Olga	Usenko	Lithia, UK
638	P209	Grappling with Economic Evaluation in Palliative Care - A Promising Foothold	Nikki	Mc Caffrey	Adelaide, AU
639	P210	The Old Dog & the Poor Relation: Informing Funding Decisions in Palliative Care with Cost-consequence Analysis in the Cost-disutility Plane	Nikki	Mc Caffrey	Adelaide, AU
640	P211	Current Perspectives on the Pediatric Palliative Care in Russia	Sergey Georgiy	Rudoy Novikov	Moscow, RU

641	P212	Palliative Care Resource Nurses as a Strategy to Improve Palliative Care Health Network: A Qualitative Study to Investigate the Factors of Success through the Nurses' Role Perception and the Institutional Point of View	Thierry	Currat	Lausanne, CH
642	P213	A Survey of End-of-Life Cancer Care Research in Europe	Katrin	Sigurdardottir	Trondheim, NO
643	P214	Translating the WHO Definition of Palliative Care into Scientific Practice: The Multiprofessional Academic Network at the Interdisciplinary Center for Palliative Medicine in Munich	Gian Domenico	Borasio	Munich, DE

Psychology and communication

644	P215	Patients' Psychosocial Experiences of Attending Specialist Palliative Day Care (SPDC): A Systematic Review	Miriam	Johnson	Hull, UK
646	P217	End-of-Life Care Communication Barriers for Doctors and Patients with Advanced HIV Disease	Elena	Vvedenskaya	Nizhny Novgorod, RU
647	P218	Spirituality and Depressed Mood in Patients with Advanced Cancer	Mari	Lloyd-Williams	Liverpool, UK
648	P219	Expert Opinion on Unresolved Issues Relating to the Detection and Treatment of Depression in Palliative Care: A Delphi Exercise	Lauren	Rayner	London, UK
649	P220	Teaching Communication Skills Using a 'Toolkit'. Is it Effective?	Helen	Mitchell	Cardiff, UK
650	P221	Communication Issues in Patients with Breakthrough Cancer Pain	Katherine	Webber	London, UK
651	P222	Exercise as a Supportive Therapy in Incurable Cancer: Exploring Patient Preferences	Matthew	Maddocks	Nottingham, UK
652	P223	An Investigation of Terminally Ill Taiwanese Patients' and Palliative Care Professionals' Perspectives on Dignity: A Preliminary Study for Conducting Dignity Therapy in Taiwan	Hui-Ching	Li	London, UK
654	P225	Assessing Feasibility and Potential Effectiveness of a Variant of Dignity Therapy for Terminally Ill Cancer Patients in Taiwan: A Pilot Study	Hui-Ching	Li	London, UK
655	P226	Psychologists in Palliative Care in Europe: A Discipline "Under Construction"	Saskia	Jünger	Aachen, DE
656	P227	Patients' Perceptions of Depression: Implications for Health Professionals	Mari	Lloyd-Williams	Liverpool, UK
657	P228	A Comparative Study by Literature Review and Interview of the Roles in Palliative Care of the Clinical Psychologist in France and the Social Worker in the United Kingdom	Anella	Gallen	Lyon, FR
658	P229	Improving Continuity of Care for Palliative Patients: What Discharge Information Do General Practitioners Want?	Helen	Mitchell	Cardiff, UK
659	P230	Explorative-linguistic Analysis of Face-to-Face Interviews on Euthanasia with Terminal Patients, Their Physicians and Their Relatives	Marianne	Dees	Antwerpen, BE
660	P231	Talking about Advance Care Planning: How Do Terminally Ill Patients and Their Closest Carers Approach these Conversations?	Philomena	Swarbrick	Lancaster, UK

661	P232	Diaries and Narratives as a Coping Mechanism in Palliative Care	Naomi	Mor	Ramat Hasharon, IL
662	P233	Communication with People with Dementia in End of Life Care: The Relevance of Gender-related Issues	Elisabeth	Reitinger	Vienna, AT
663	P234	Palliative Medicine: Ethnocultural Mechanisms of Psychological, Psychopathological and Psychosocial Factors in Formation Professional Desadaptations, Relations to Psychopathology and Psychopharmacotherapy	Tofik	Biktimirov	Ulyanovsk, RU
664	P235	Oncologists' Disclosure of Prognostic Information – Reflections upon Own Practice	Lotte	Rogg	Oslo, NO
665	P236	Acute Distress Even before the Diagnosis is Confirmed: A Qualitative Longitudinal Study of People with Malignant Glioma and Their Relatives	Scott	Murray	Edinburgh, UK
666	P237	Getting 'Connected': Investigating Doctors' and Nurses' Attitudes to Undertaking Advanced Communication Skills Training	Mary	Turner	Lancaster, UK
667	P238	Differences and Similarities in the Disclosure of a Cancer Diagnosis and Prognosis between Three Mediterranean Countries: A Comparative Overview	Arantza	Meñaca	Barcelona, ES
668	P239	Validation and Psychometric Evaluation of the German Spouse Response Inventory in Patients Suffering from Cancer Pain and Headache	Frank	Elsner	Dresden, DE
669	P240	Correlation between Change in Targeted Psychosocial Resources and Quality of Life-related Outcomes in Advanced Breast Cancer Patients Participating in a Psychosocial Intervention	Amy	Abernethy	Durham, US
671	P242	The Perception of Time as a Means of Detecting Suffering	Alberto	Alonso-Babarro	Madrid, ES
672	P243	A Taxonomy of End-of-Life Communication and Decision Making	Reginald	Deschepper	Brussels, BE
673	P244	Do Patients' Preferences for Information and Participation Change between Diagnosis and Death? A Longitudinal Study of Lung Cancer Patients in Flanders (Belgium)	Reginald	Deschepper	Brussels, BE
110	P244B	Suffering: Threats and Resources	Alberto	Alonso Babarro	Madrid, ES

Research methodology

674	P245	Behind the Scenes of an EAPC Taskforce Survey on Education for Psychologists in Palliative Care	Eva	Eggenberger	Vienna, AT
675	P246	Contribution of Ethnography to Palliative Care Research	Tania	Pastrana	Lancaster, UK
676	P247	Attitudes of Health Care Professionals toward Euthanasia and Physician-assisted Suicide: A Review of the Literature: 2002-2008	Miguel Angel	Cuervo Pinna	Badajoz, ES
677	P248	Interpretive Description in Palliative Care Research: An Example from South Africa	Liz	Gwyther	Winnipeg, CA
679	P250	A Multi-method Approach to Questionnaire Development for a Cross-national Survey of Public Views on End of Life Care	Barbara	Gomes	London, UK

680	P251	Conducting Research in Palliative Care. Regional Observatory of Palliative Care in a Regional Palliative Care Program	Maria	Solano	Mérida, ES
681	P252	Searching for Best Scientific Evidence on Palliative Care: A Methodology Scheme by a Palliative Care Work Group	Miguel Angel	Cuervo Pinna	Badajoz, ES
683	P254	Analysis of the Barriers to Recruitment to the Prognosis in Palliative Care Study (PiPS)	Bridget	Gwilliam	London, UK
684	P255	Involving 'Frail' Older People in Palliative Care Research Studies: Methodological Challenges	Julie	Skilbeck	Nottingham, UK
685	P256	Validating the Findings of Qualitative Research: An Innovative Approach for Palliative Care	Lucy	Selman	London, UK
686	P257	Using Qualitative Research to Examine Decision Making in Continuous Deep Sedation until Death: Design Issues in a Newly Funded Study	Jane	Seymour	Nottingham, UK
688	P259	"Mind the Gatekeepers" – Reflections on Recruitment in an Ethnographic Study in Palliative Home Care	Sabine	Pleschberger	Vienna, AT
690	P261	Investigation of the Criteria Applied by Palliative Care Journals to Evaluate Qualitative Research	Nikki	Mc Caffrey	Adelaide, AU
691	P262	Considerations in Searching and Retrieving the Palliative Care Literature	David Deb	Currow Rawlings	Adelaide, AU
692	P263	Valuing Palliative and End of Life Care: An Economic Perspective	Konrad	Fassbender	Edmonton, CA
693	P264	User Involvement in Palliative Care Research: Impact and Challenges	Katherine	Froggatt	Lancaster, UK
694	P265	Bereavement Research: Methodological Issues when Studying Vulnerable People	Virginia	Dunn	Oxford, UK
695	P266	InForm: A Step forwards in Clinical Trial Data Management	Ruth	Branford	London, UK
696	P267	Palliative Care Making a Difference in Africa: A Rapid Evaluation Methods Study	Scorr	Murray	Edinburgh, UK
697	P268	End of Life Care Research in Intensive Care Units: Challenges and Solutions	Caroline	Rumble	London, UK
698	P269	In which Impact Journals Publish Contributions the Most Active Researchers of Palliative Care?	Ana	Carvajal	Pamplona, ES
699	P270	Are Studies in Palliative Care Biased? Characteristics of Patients Recruited to the EPOS Study	Peter	Fayers	Aberdeen, UK
700	P271	An Information Technology (IT) Infrastructure that Supports Longitudinal Patient-reported Outcome (PRO) Assessment and Correlation of Symptoms with Patient Parameters	Amy	Abernethy	Durham, US
702	P273	Researching Day-to-Day Ordinary Practice	Erna	Haraldsdottir	Stirlingshire, UK
703	P274	Review Methods in Palliative Care: Learning from an Innovative Realist Review Approach	Catherine	Walshe	Manchester, UK
704	P275	Translation and Cross Cultural Adaptation of Instruments to Measure Comfort, Hope and Quality of Life in Palliative Care Research	Ana	Querido	Leiria, PT
705	P276	Two Roles in the Same Soul: Being Both Clinical Researcher and Specialist Palliative Care (PC) Professional (Nurse/Doctor)	Susanne	Linder	St. Gallen, CH

