

Seminário

Vida com dignidade e qualidade até ao fim

14 de outubro de 2017

Fundação Calouste Gulbenkian

Programa

Seminário com
tradução em simultâneo

Imagem: Ad Médic

COM O ALTO PATROCÍNIO
DE SUA EXCELÊNCIA
UNDER THE HIGH PATRONE OF THE
PRESIDENT OF THE PORTUGUESE REPUBLIC



O Presidente da República

Dia mundial
dos cuidados paliativos



World hospice &
palliative care day
Voices for hospices

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Auditório 2 | Fundação Calouste Gulbenkian

13.00h Abertura do Secretariado

14.00–14.20h

Cuidados Paliativos – Um direito humano: O que os portugueses sabem deste direito?

Palliative Care – A human right: What the portuguese know about this right?

Prof. Doutor Manuel Luís Capelas

14.20–15.05h

Abordagem de Saúde Pública nos Cuidados em Fim de Vida: Construção de Comunidades e Cidades Compassivas

A Public Health Approach to End-of-Life Care: Building Compassionate Communities and Cities



Allan Kellehear, PhD

Professor de Cuidados em Fim de Vida na Universidade de Bradford, no Reino Unido. Sociólogo clínico e de saúde pública. Fundador da primeira unidade académica do mundo de cuidados paliativos em saúde pública (Austrália, 1990). Autor das primeiras publicações no âmbito dos cuidados paliativos numa perspetiva de saúde pública. Cofundador com Sallnow e Kumar da Série de Conferências de Cuidados Paliativos em Saúde Pública. Membro da Academia de Ciências Sociais e Presidente da Association for the Study of Death and Society (ASDS) and Public Health Palliative Care International (PHPCI).

50th Anniversary Professor End-of-Life Care) at the University of Bradford, UK. He is a medical and public health sociologist. Founder of the world's first academic public health palliative care unit in Australia in the 1990s, he also wrote some of the first academic literature expounding the public health approach to palliative care, and co-founded (with Sallnow and Kumar) the Public Health Palliative Care Conference Series. He is a Fellow of the Academy of Social Sciences, and President of both the Association for the Study of Death and Society (ASDS) and Public Health Palliative Care International (PHPCI).

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15.05–15.50h

Os Cuidados Paliativos como um Direito Humano: Perspetiva de justiça social

Palliative Care as a human right: A social justice perspective



Philip J. Larkin, PhD

Professor Responsável de Enfermagem Clínica (Cuidados Paliativos) na Escola de Enfermagem Obstétrica e de Sistemas de Saúde do University College Dublin; Coordenador do Our Lady's Hospice & Care Services em Dublin, Irlanda. Presidente da Direção da Associação Europeia de Cuidados Paliativos.

Joint Chair, Professor of Clinical Nursing (Palliative Care), University College Dublin School of Nursing Midwifery and Health Systems, and Our Lady's Hospice & Care Services, Harold's Cross, Dublin, Ireland. President of the Board of Directors of the European Association for Palliative Care.

15.50–16.15h *Coffee-break*

16.15–17.45h

Debate Público

Moderadores: Jornalista Ana Filipa Nunes e Prof. Doutor Manuel Luís Capelas

Comentadores: Prof. Doutor Alexandre Castro Caldas e Dra. Sofia Silva

17.45h

Receção a Sua Excelência o Presidente da República

Professor Doutor Marcelo Rebelo de Sousa

18.00h

Encerramento do Seminário

Com a participação e comunicação de Sua Excelência

o Presidente da República Professor Doutor Marcelo Rebelo de Sousa

Resumo da comunicação

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Abordagem de Saúde Pública nos Cuidados em Fim de Vida: Construção de Comunidades e Cidades Compassivas

A Public Health Approach to End-of-Life Care: Building Compassionate Communities and Cities

Allan Kellehear, PhD., FAcSS
University of Bradford, UK

This lecture will commence with some preliminary remarks about the globally widespread clinical, acute-care culture of palliative care and the common myths and distortions in the idea of 'community' as traditionally used by this style of palliative care before then introducing an outline of public health/health promotion approaches to palliative care. I will then introduce a public health approach to palliative care as one that addresses the social, psychological and spiritual troubles that people living with dying, living with bereavement, and living with long term care. These 'co-morbidities' are all amenable to prevention, harm reduction, and early intervention by public health methods. These interventions embrace community development, health promotion, death education, and ecological strategies. I will argue that the future of palliative care must embrace this approach if it is to successfully achieve greater access for all and provide a meaningful continuity of care at the end of life. Current global examples of compassionate communities or compassionate cities will serve as illustrations of this approach .



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Philip J. Larkin, PhD

University College Dublin, UK

There has been an ongoing dialogue led by world palliative care leaders for palliative care to be defined as a human right. This dialogue has suggested that to deny people access to palliative care when they need it, particularly in the context of effective pain relief, is both cruel and inhuman. There are many international examples where poor access, poverty, lack of services and limited political will to improve and develop palliative care mean that many citizens are unable to access palliative care when they most need it. The World Health Authority resolution 69.14 (2014) calls on nations to promote and embed palliative care in their health systems. Yet, three years on, impact seems limited.

An alternative argument is that palliative care as a human right is a worthy aspiration but hard to implement. Given the challenges of disadvantage and marginalization which impacts many people, a call for a model based on social justice which looks at care and choice within existing care provision may be a stronger framework in which to promote optimal palliative care.

This presentation will look at the Human Rights debate in palliative care and offer an alternative proposal, a model of Affective Equality, based on love, care and solidarity (Baker & Lynch 2009) to understand how we can improve the place of palliative care within our health systems. Using examples from recent research in palliative care using this framework, the application of the model to care of patients in our Society who often fail to access palliative care will be discussed and debated.





VERSÃO DIGITAL
DO PROGRAMA



Organização



Associação Portuguesa
de Cuidados Paliativos

Secretariado

admedic⁺

ORGANIZAÇÃO E SECRETARIADO
DE EVENTOS

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