

# Quality Palliative Care in Long Term Care: Tools for Change



[palliativealliance.ca](http://palliativealliance.ca)

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# The Context of LTC

- It is common for 40% to 50% of residents to die each year in LTC homes. (CIHI)
- LTC is a unique palliative care context.
  - frail older people living with progressive life limiting disease
  - A home where residents will both live and die
  - Heavily regulated and inspected (external standards)
- The majority of LTC homes in Canada lack formalized palliative care programs.

# Quality Palliative Care –Long Term Care Project Background

- Funded by Social Sciences and Humanities Research Council (SSHRC) for a five year Community-University Research Alliance called: *Quality Palliative Care in Long Term Care Alliance (QPC-LTC)*
- Knowledge Translation for this project funded by Canadian Institute for Health Research (CIHR)
- Includes 30 organizational partners and more than 20 researchers nationally and internationally
- Involves four LTC homes in Ontario as study sites

# Goals of the Project

- Improve the quality of life for residents dying in LTC
- Develop interprofessional palliative care programs
- Create partnerships between LTC homes, community organizations and researchers
- Create a toolkit for developing palliative care in LTC homes that can be shared nationally
- Promote the role of the Personal Support Worker (PSW) in palliative care

# 4 Key Messages

***“We will care for you for the rest of your life”***

***“It’s hard to watch people die for a living”***

***“Resident’s are not like paperwork”***

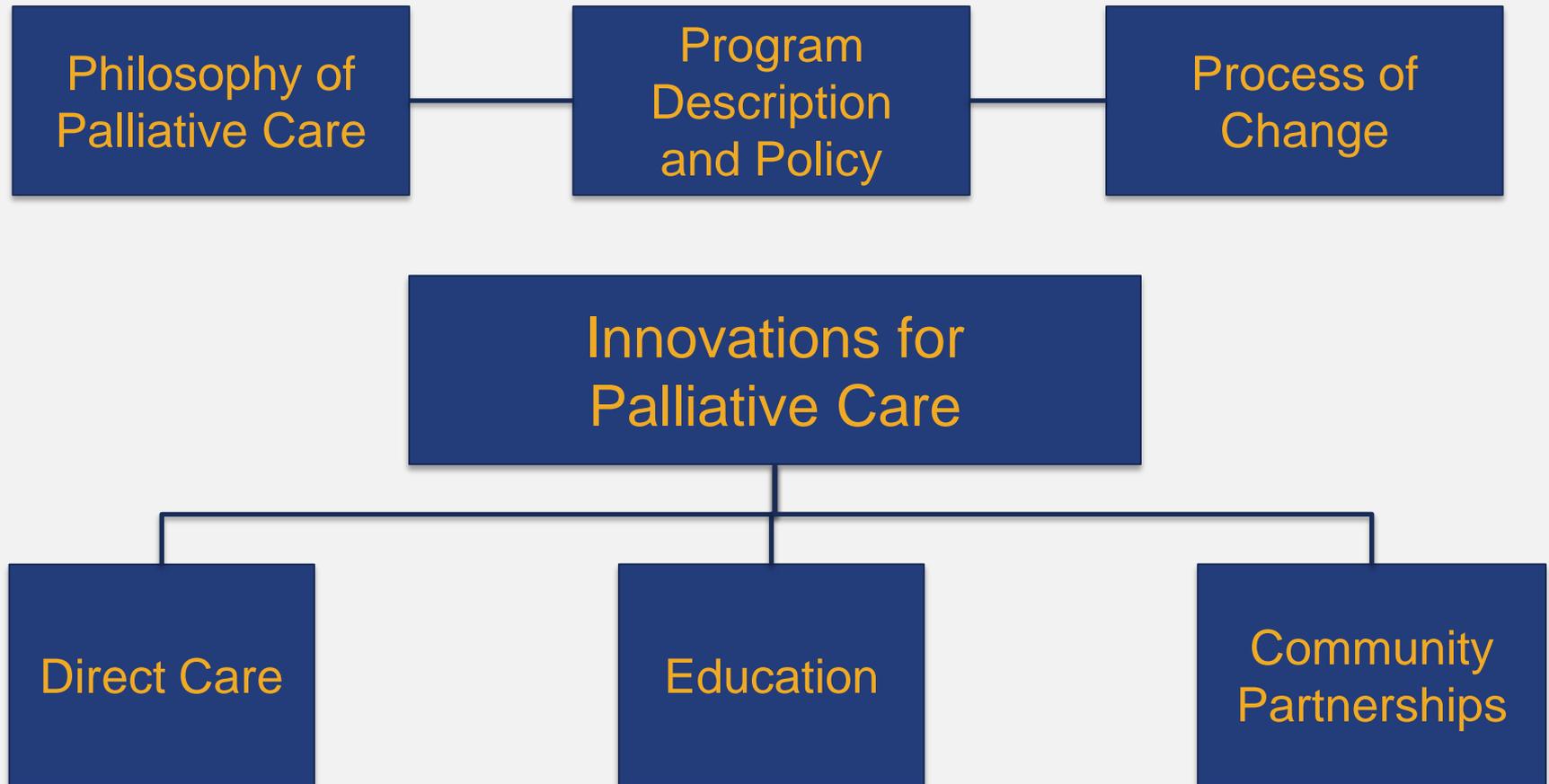
***“You can’t regulate humanistic care”***



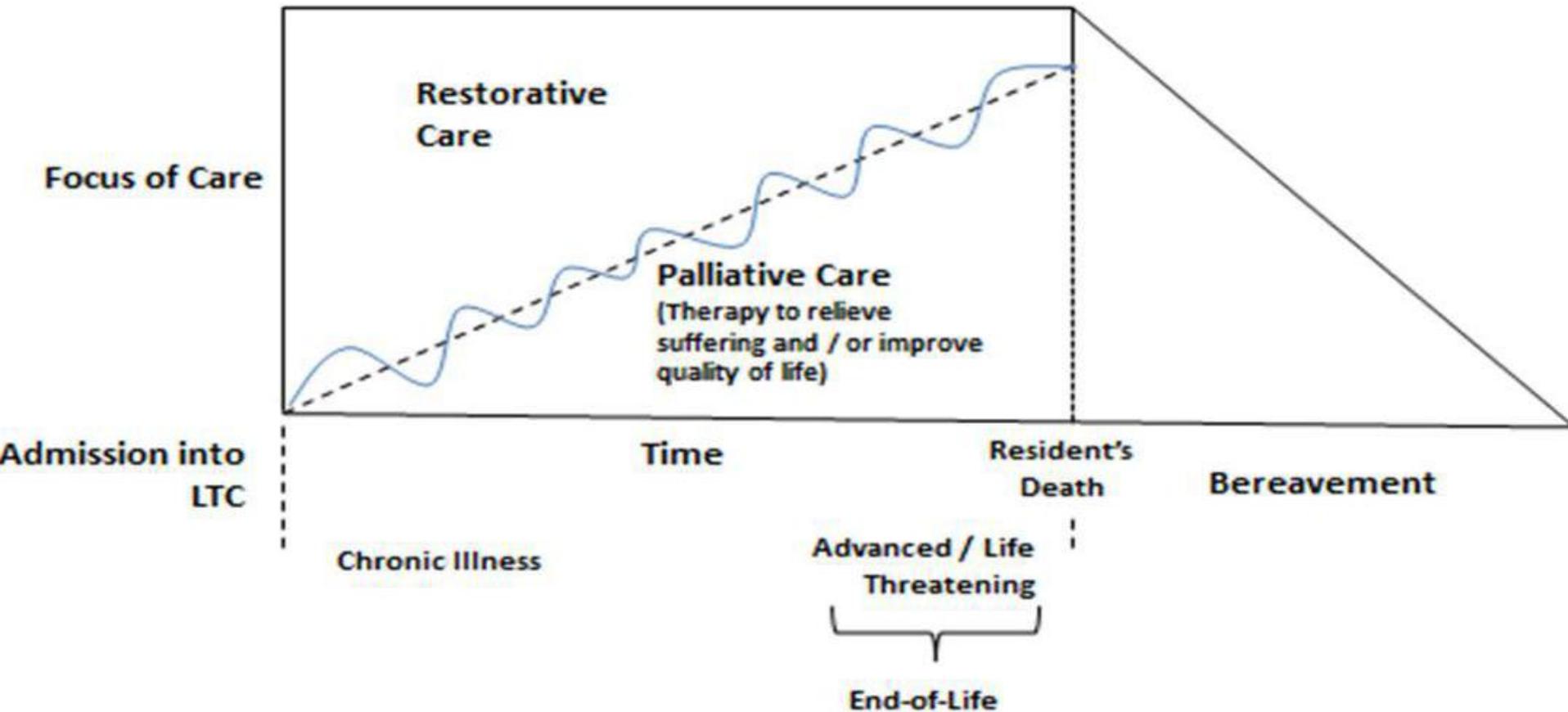
# Family Perspective

- Families want residents to die in long term care IF resources and education are available to staff
- There needs to be open communication between families and staff
- Families need to feel part of “the team”
- Families see that there is a shortage of staff
- They recognize the contribution that community partners can play in delivering palliative care

# Framework of Palliative Care in Long Term Care



# Transition from Admission to Death



CHPCA, 2002

# What is Palliative Care and End-of-Life Care?

## **Palliative Approach to Care**

- Focus is on quality of life, symptom control
- Interdisciplinary in approach
- Client centered and holistic
- Begins when death would not be “unexpected” in the next year

## **EOL Care**

- Death is inevitable
- Trajectory is short (6 months or less)
- Focus is on supporting patient and family choices
- Addresses anticipatory grief
- Supports resident with a “good death”

# When does EOL Care begin?

Palliative Performance Scale (PPSv2)

PPS Level	Ambulation	Activity & Evidence of Disease	Self-Care	Intake	Conscious Level
100%	Full	Normal activity & work No evidence of disease	Full	Normal	Full
90%	Full	Normal activity & work Some evidence of disease	Full	Normal	Full
80%	Full	Normal activity <i>with</i> effort Some evidence of disease	Full	Normal or reduced	Full
70%	Reduced	Unable Normal Job/Work Significant disease	Full	Normal or reduced	Full
60%	Reduced	Unable hobby/house work Significant disease	Occasion assistance necessary	Normal or reduced	Full or Confusion
50%	Mainly Sit/Lie	Unable to do any work Extensive disease	Considerable assistance required	Normal or reduced	Full or Confusion
40%	Mainly in Bed	Unable to do most activity Extensive disease	Mainly assistance	Normal or reduced	Full or Drowsy +/- Confusion
30%	Totally Bed Bound	Unable to do any activity Extensive Disease	Total Care	Normal or reduced	Full or Drowsy +/- Confusion
20%	Totally Bed Bound	Unable to do any activity Extensive Disease	Total Care	Minimal to sips	Full or Drowsy +/- Confusion
10%	Totally Bed Bound	Unable to do any activity Extensive Disease	Total Care	Mouth care only	Drowsy or Coma +/- Confusion
0%	Death				

Stages:  
■ Stable  
■ Transitional  
■ End of Life

Reproduced with permission from Victoria Hospice Society, 2003.

# Philosophy of Care

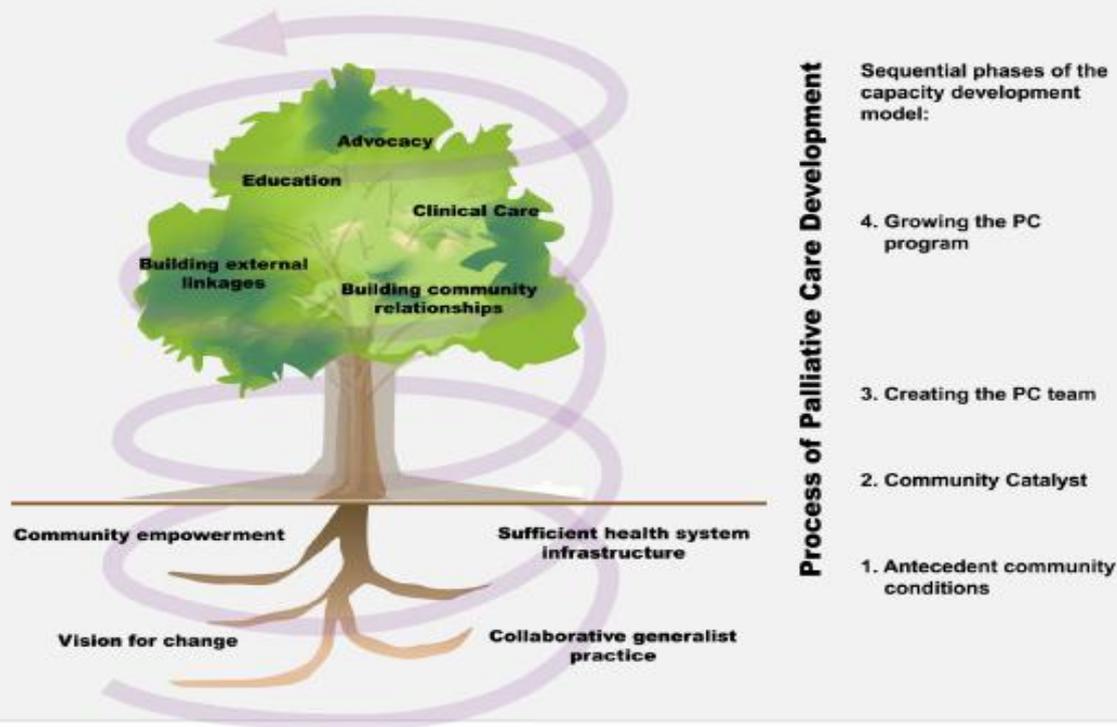
- Resident-Centred Care/Relationship-Centred Care
  - Empowers residents to be decision-makers in their own care
  - Respects residents choice, wishes, values, goals
  - Treats residents as unique, whole persons
  - Provides residents tools to care for themselves
  - Advocates for residents; acts on their concerns
  - Focuses on relationships as the core process in quality care
  - Values interdependence

# Program Description and Policy

- Goals of the program
- Program objectives
- Relevant Definitions
  - Palliative Approach
  - End-of- Life Care
  - Advance Care Planning
  - Interdisciplinary Palliative Care Resource Team
- Relevant Programs Policies and Procedures related to PC delivery

# Processes of Change

- Capacity development model for developing palliative care



# Getting Started...

- Self Assessment Audit

Description	Progress	Document or Evidence to Support
<b>Organization Context</b> <b>Formalized palliative care program Description that includes:</b>		
<ul style="list-style-type: none"> <li>• Goals and objectives of the program</li> <li>• Definition of palliative care, end-of-life care, and care planning</li> <li>• List of services that are available within the palliative care program</li> <li>• There is a process in place to identify residents who would benefit from a palliative care approach.</li> <li>• Assessments specific to palliative care and end-of-life are listed (ie. Palliative Performance Scale)</li> <li>• Staff have a formal process of communicating the palliative care needs of a resident (shift to shift report, reporting on electronic charts)</li> <li>• There is pain and symptom management built within the program</li> <li>• Residents and Families are actively contributing and participating in the palliative care program development</li> <li>• Residents and families have access to palliative care education</li> <li>• Protocols are in place to support staff around grief and loss</li> <li>• There is a quality improvement strategy in place for palliative and end-of-life care initiatives (e.g. process mapping)</li> <li>• Evaluation of resident, family and staff satisfaction</li> </ul>	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> In Development <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> In Development <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> In Development <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> In Development  <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> In Development  <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> In Development <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> In Development <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> In Development <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> In Development <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> In Development <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> In Development <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> In Development <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> In Development	

# Benefits of using Self Assessment

- Self-assessment provides an opportunity to identify gaps in your palliative care program and areas of strength/capacity
- Can guide the development of your program and policies
- When used regularly it can be an ongoing evaluation of your program
- Can help to determine organizational palliative care priorities

# Key Goals for change

- Expand Advance Care Planning
- Promote Formalized Palliative Care Programs
- Enhance human resources to provide holistic palliative care
- Support creation of palliative care teams
- Strengthen interprofessional collaboration within LTC homes and with community, and
- Integrate PC Philosophy into Resident-Centred Care

**\* Supported by Family Councils of Ontario and Concerned Friends of Ontario**

# Palliative Care Resource Team

- Interprofessional including PSWs
- Engages community partners (eg. Alzheimers Society)
- Not a clinical team
- Meets monthly
- Chaired by a staff member
- Provides leadership and mentorship within the home
- Provides a formal structure to organize education, identify needs, create strategies & support staff

# Resources for families

- Information in resident handbook
- Promote discussion of advance care planning more broadly than medical directives
- Discuss palliative care and EOL care in annual care conference
- Promote dedicated palliative care conference when appropriate
- Promote referral to hospice volunteers and grief support group when appropriate
- Promote referral to Alzheimer's Society for end of life issues workshop

# Resources for Staff

- Post death debriefing sessions
- Education about palliative and end of life care
- Pain screening and communication tools
- Palliative performance scale
- Promote use of Pain and Symptom consultants and teams for management of pain, feeding/hydration issues and delirium

# Getting Started with our toolkit

<http://www.palliativealliance.ca/news>

- Organizational Self Assessment tool of structures, process and outcomes
- Education: Palliative Care for Front Line Workers course and LEAP for Long Term Care
- Brochure on the progression of palliative care and end-of -life care to help discussions with families
- Toolkit on implementing the PPS (coming soon)
- Brochure on the role and structure of the palliative care program and team

# Innovations of Palliative Care in Long Term Care:

## Direct Care Processes

- Comfort Care Rounds
- Snoezelen
- Comfort Care Bags
- Pain Screening, Assessment and Follow-up Protocol
- PPS and Palliative Care Conferences

## Education for Staff and Volunteers

- Simulation Lab Experience for PSWs
- Palliative care for LTC workers – 10 module course
- Hospice Visits
- Spiritual Care in-services

# Innovations of Palliative Care in Long Term Care:

- Community Partnerships
  - Collaboration with community resources
  - Hospice Volunteers
  - Alzheimer's Society Education Seminars
  - Palliative Pain and Symptom Management Consultants
  - Nurse led outreach teams (nurse practitioners)

# Implementation Barriers – Require Advocacy!

- Human resources needs to be supplemented at all levels for palliative and end-of-life care
- Homes now choose between a spiritual care advisor or a social worker
- Time lacking for interprofessional teamwork
- J5 on RAI is not linked to CMI and funding
- No dedicated training dollars (to access training / backfill)

# Further Information

Visit our website:

[www.palliativealliance.ca](http://www.palliativealliance.ca)

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Special thanks to:



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