

# OMA Palliative care advocacy plan



# Introduction

This plan was created through application of the new OMA Advocacy Toolkit to the issue of reducing hospital backlog by prioritizing palliative care. The plan is intended to serve both as an advocacy roadmap and, in part, as an example of the Toolkit's implementation to assist members in filling out the kit for their own advocacy efforts.

What needs to change? Reducing delays in patient care

What is your goal(s)? Prioritize palliative medicine.

Addressing issues with referrals, reducing wait times, increasing access to quality care through allied health supports

Who do we need to influence?	What is important to them?	How can we influence them?
Physician leaders	<ul> <li>Reducing delays in patient care</li> <li>Reducing physician burnout and moral distress</li> <li>Improving patent experience</li> </ul>	Sharing information through social media, blogs, or infographics
Ontario Health Teams	<ul><li>Improving patient experience</li><li>Improving population health outcomes</li></ul>	Building partnerships by knowledge sharing or hosting joint meetings
Patient advocate groups	<ul> <li>Higher quality care for those waiting for, or in palliative care</li> <li>Improving patient experience</li> </ul>	Supporting advocacy champions by building community networks or mentorship programs
Government	<ul> <li>Reducing healthcare cost</li> <li>Improving patient experience</li> <li>Reducing wait times</li> </ul>	Use data-driven advocacy to demonstrate cost savings and system improvements.

# What do we need to do this?

- Register for OMA advocacy network for further advocacy training
- Connect with groups via email
- Connect with people with writing/graphic design skills
- Register for the OMA Healthcare Advocates Program

# Identify your contacts

It is important to identify your contacts for your outreach and any other referrals that may come from your advocacy work to successfully engage them and develop relationships. Keep track of your meetings with them and any key points to advancing your cause.

Targeted Audience	Contact Name	Title	
OMA Physician leaders	Nicole Biordi	Sr. Director, Physician Leader	
-		Engagement, Ontario Medical	
		Association	
Palliative medicine	Joyce Cheung	Section Chair	
	Sukhjeet Dhillon	Section Vice-chair	
Emergency medicine	Angela Marocco	Section Chair	
	Tonja Stothart	Section Vice-chair	
<ul> <li>Section of General and</li> </ul>	David Barber	Section Chair	
Family Practice	Beth Perrier	Section Vice-chair	
Geriatric medicine	Rajin Mehta	Section Chair	
	Ali Ahmad	Section Vice-chair	
<ul> <li>Long term care/care of</li> </ul>	Mohamed Abu-Abed	Section Chair	
the elderly	Abhishek Narayan	Section Vice-chair	
Radiation oncology	Thomas McGowan	Section Chair	
	Peter Chung	Section Vice-chair	
Haematology &	Joanna Gotfrit	Section Chair	
medical oncology	Patricia Disperati	Section Vice-chair	
General internal	Bilal Munir	Section Chair	
medicine	Rishi Parihar	Section Vice-chair	
Hospital medicine	Stephen Chin	Section Chair	
	William Coke	Section Vice-chair	
<ul><li>Psychiatry</li></ul>	Diana Kljenak	Section Chair	
	Katherine McKay	Section Vice-chair	
Critical care medicine	David Katz	Section Chair	
	Ian Randall	Section Vice-chair	
<ul><li>Pediatrics</li></ul>	Jane Healey	Section Chair	
	Shawn Kao	Section Vice-chair	
<ul> <li>Rural Medicine Forum</li> </ul>	Christine Seidler	Fora Chair	
Patient advocate groups	Tara Walton	Manager, Ontario Palliative Care Network	
	Rick Firth	CEO, Hospice Palliative Care	
		Ontario	
		Unhoused advocacy groups	
Indigenous Physicians	Melanie Osmack	Executive Director	
Association of Canada			
Ontario Health Teams	ontariohealthteams@ontario	https://www.ontario.ca/page/o	
	.ca	ntario-health-teams	
Ministry of Health	Peter Kaftarian	Associate Deputy Minister,	
		Clinical Care and Delivery	

#### What needs to change?

To reduce delays in patient care by prioritizing palliative medicine

## What are we asking for?

#### 1. Equitable and timely access to palliative care

- 1.1. Create a heat map that shows Ontario's gaps in palliative care versus the areas of the province with the greatest number of ALC days and longest hospital wait times and use it as a guide to determine how many palliative care and hospice beds Ontario needs, and where they should be placed.
- 1.2. Address inequities and imbalances in the existing physician funding models, such as AFP and Community Palliative On-Call applications, to help reduce system fragmentation.
- 1.3. Consistent access to specialized palliative care and resources in the community to provide guidance around end-of-life care and symptom management.
- 1.4. Identify areas of the province with limited or no 24/7 support for patient, families, and caregivers, and invest in provision of on-call supports for those areas
- 1.5. Expand access to telemedicine-based palliative care consultation for patients in rural, remote, and underserviced areas.
- 1.6. Work to ensure telephone/virtual access to on-call palliative specialists for areas of the province where direct palliative care consultation is not yet available, to help provide the opportunity for all patients to die at home, should they choose.

## 2. Reduce reliance on acute care settings at end of life

- 2.1. Calculate the cost of palliative care ALC days, and reinvest that money into palliative home and community supports.
- 2.2. For the areas of the province with the greatest need, fund, create, and operate new chronic palliative care beds within existing infrastructure such as long-term care homes.
- 2.3. Amend criteria for hospice and palliative care involvement to focus on provider recommendation and patient benefit, rather than expected prognosis.
- 2.4. Address barriers to home care support—especially in rural and remote settings.
- 2.5. Increase number of skilled palliative care providers, including physicians, nurses, and other allied professionals.

#### 3. Integrated person-centered system

- 3.1. Reduce preventable admissions by identifying successful home frailty programs, investing further in them, and expanding them into other areas of the province.
- 3.2. Ensure all patients have discharge options to allow for them to die in their preferred setting, whether in the hospital or community.

	Who are you asking?			
	The OMA	The OMA	The	The public
	(corporate)	(members)	government	
What do you need from them?	(corporate) Greater emphasis on palliative medicine in next large-scale advocacy campaign	(members)  Participate in  OMA advocacy efforts	government  All Ontarians have access to quality palliative care when and where they need it	Highlight that half of Ontario may not have an option on where they choose to die. Sharing personal stories on what delays in care look like for palliative patients and families, as well as about how their lives have been affected by delays in patient care due to delayed/limited
				access to palliative care.

Stakeholder analysis					
Stakeholder	Stance	Impact	Interest	Support for OMA position	
Ontario Palliative Care Network	The OPCN is part of OH. The OPCN acts as a principal advisor to government for quality, coordinated palliative care in Ontario. The OPCN also includes regional networks, to support local palliative care delivery.  The OPCN creates delivery frameworks for the best model of care in the community, hospital and for pediatric patients.  OMA/OPCN staff had a monthly touch point for several years to align our advocacy positions and discuss regional/member specific issues. These meetings are currently on hold due negotiations.	High	High	High	
Hospice Palliative Care Ontario	Hospice Palliative Care Ontario is a provincial association for hospices and palliative care providers, professionals, and volunteers throughout Ontario.	High	High	High	

Canadian Society of	The national society for palliative care	High	High	High
Palliative Care Physicians	physicians. Their mission is to promote			
	palliative medicine in Canada through			
	advanced education, strengthening the			
	workforce, and advocacy.			
Ontario Hospital	Palliative Care is an approach to care that	High	Medium	Medium
Association	focuses on comfort and quality of life.			
	The goal is much more than comfort in			
	dying; palliative care is about living well,			
	through meticulous attention to control			
	of pain and other symptoms, supporting			
	emotional, spiritual, and cultural needs,			
	and maximizing functional status.			
Impacted OMA sections	Supports the OMA on the following core	High	High`	High
<ul> <li>Palliative medicine</li> </ul>	functions:			
Emergency	• Compensation			
medicine	Subject matter expertise			
Section of General	Engagement, leadership			
and Family Practice	Advocacy			
Geriatric medicine				
Long term				
care/care of the				
elderly				
<ul> <li>Radiation oncology</li> </ul>				
<ul> <li>Haematology &amp;</li> </ul>				
medical oncology				
<ul> <li>General internal</li> </ul>				
medicine				
<ul> <li>Hospital medicine</li> </ul>				
<ul> <li>Psychiatry</li> </ul>				
Critical care				
medicine				
<ul> <li>Pediatrics</li> </ul>				
<ul> <li>Rural Medicine</li> </ul>				
Forum				

#### **Key messages**

Why is this an issue?

Our health care system underfunds palliative/hospice beds. As such, many who would benefit from palliative care supports are instead awaiting disposition in the acute care system.

What/How do we go about fixing it?

- Review funding model to ensure equitable access to funding for palliative care in the community, especially in small, rural, and northern communities
- Improve palliative care supports for health care providers, families and communities, especially for after-hours (CPOC, AFP, FFS, virtual)
- Use PRIME to map out what palliative services are provided in small, rural, northern communities
- Increase access to palliative care for underserved populations

What would be the benefit to your audience?

Reducing delays in care/reducing the hospital backlog by providing care in more appropriate, patient-centered settings.

A shift towards increasing palliative/hospice capacity would improve measures in a number of ways:

- Investment in better palliative care flow will open up beds, which reduces the length of time that patients sit in ER waiting to be admitted, which reduces ER wait times overall
- ii. Improved patient and family satisfaction by receiving high quality palliative approach to care in teams who are trained to provide it.
- iii. Reduction in provider burnout caused by caring for end-of-life patients outside of supported environment.

#### **Tactics**

#### **Integrate into OMA Advocacy Campaigns**

• Use advocacy asks to strengthen advocacy campaigns that focus on reducing delays in patient care

# **Strengthen Collaboration with Advocacy Partners**

- When appropriate, partner with organizations such as the Ontario Palliative Care Network (OPCN), Ontario College of Family Physicians (OCFP), and Canadian Society of Palliative Care Physicians (CSPCP) to align efforts.
- Work with hospital administrators and long-term care leaders to push for increased funding and staffing for palliative care units and community hospice and palliative care beds.

#### **Engage the Public and Patient Advocacy Groups**

 Collaborate with patient advocacy organizations like the Canadian Hospice Palliative Care Association (CHPCA) and grassroots caregiver networks