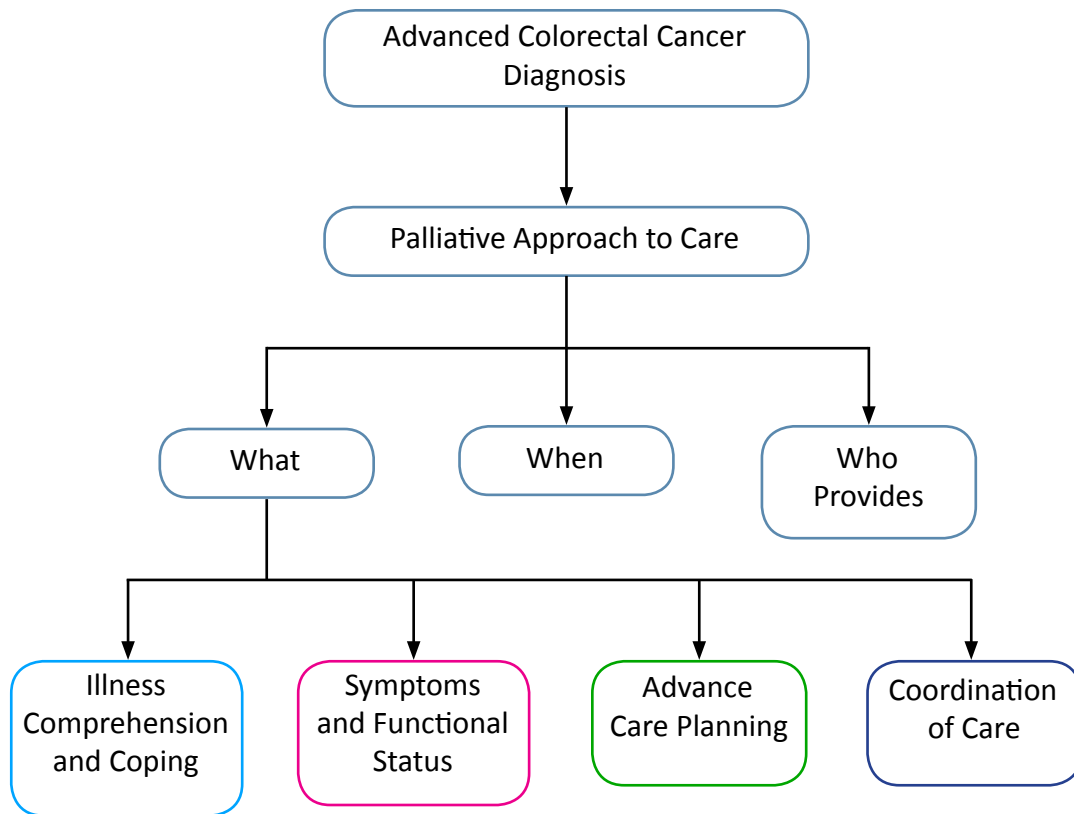


# Integrating an Early Palliative Approach into Advanced Colorectal Cancer Care - Interactive Care Pathway

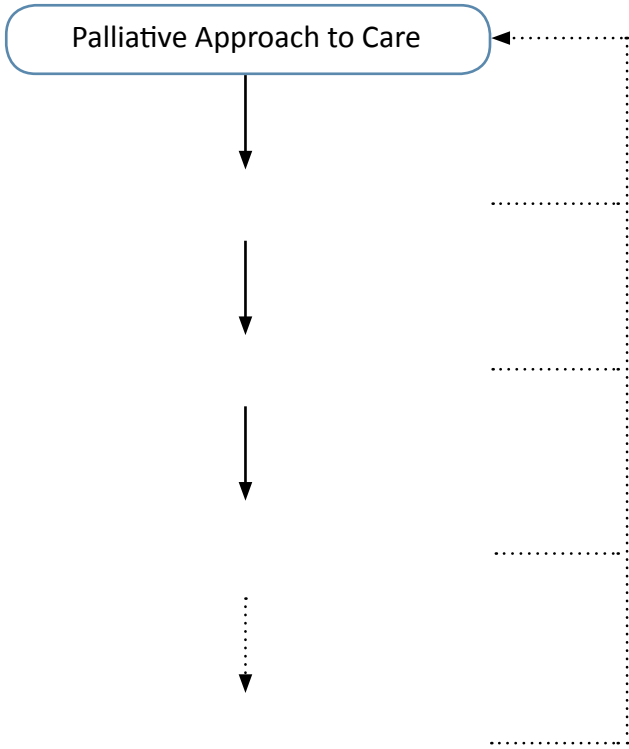
Effective Date: January, 2019



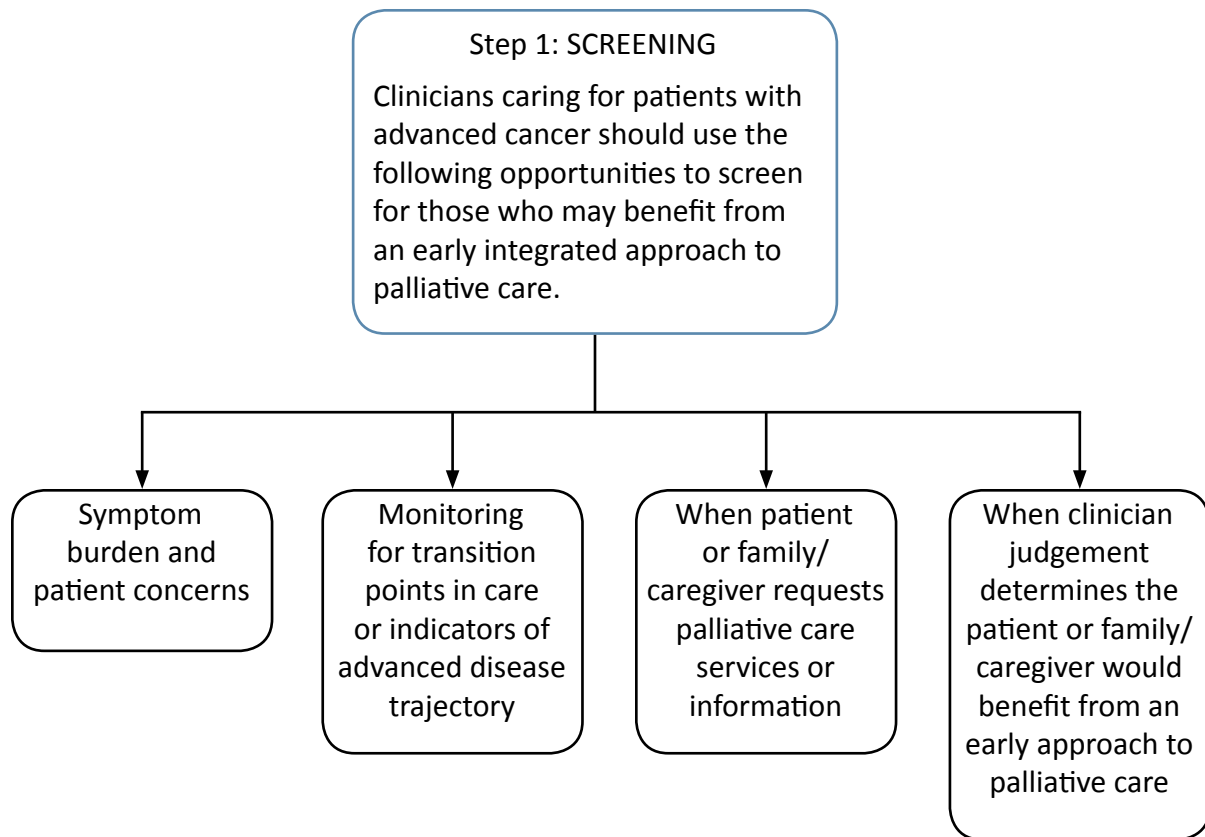
Please refer to the [Advanced Colorectal Cancer Care Clinical Practice Guideline](#) for additional information



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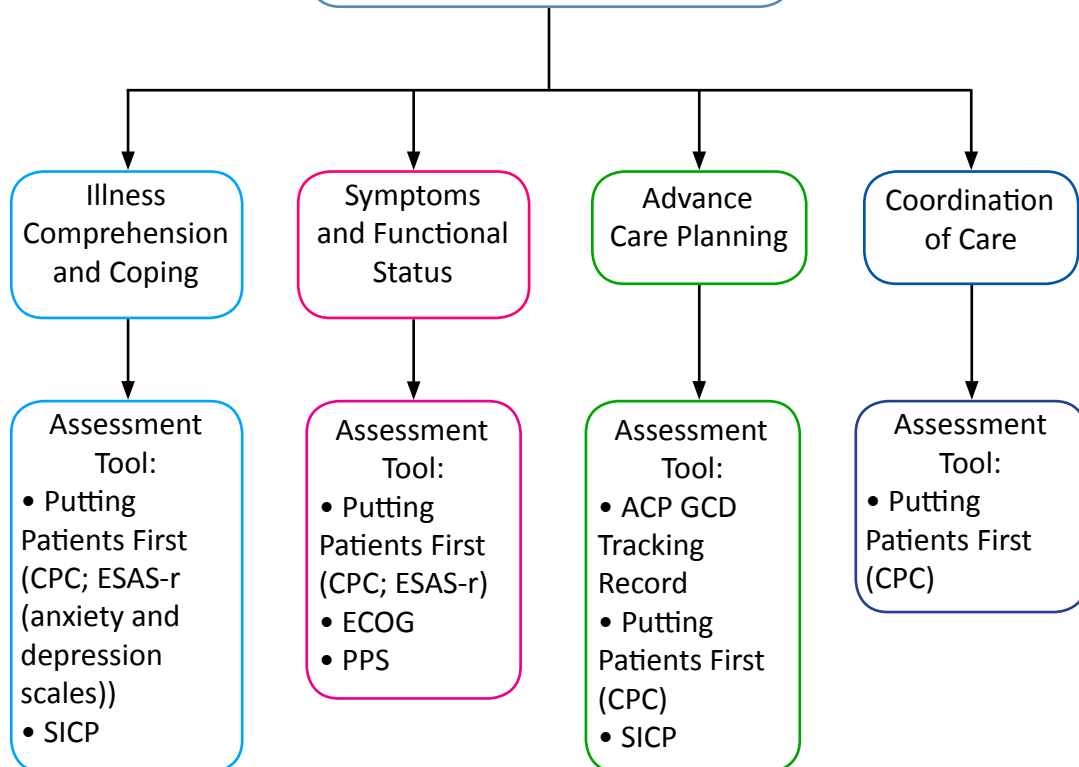
Patients who screen positive for one or more of these screening methods meet criteria for use of an early integrated palliative approach to care.

Patients who screen negative should be rescreened regularly.

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## Step 2: IDENTIFY PATIENT NEEDS

Once patients have been screened, clinicians should identify unmet patient needs. This can be guided by the four essential components of an early palliative care approach and corresponding assessment tools. Patient needs should be re-assessed at regular intervals.

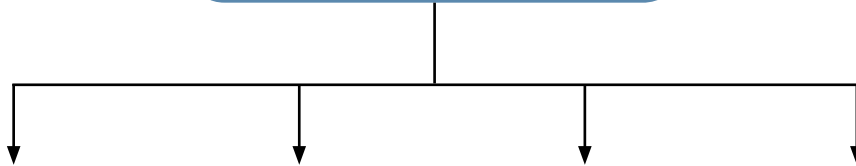


PPF= Putting Patients First, CPC= Canadian Problem Checklist, ESAS-r= Edmonton Symptom Assessment System- Revised, SICIP= Serious Illness Care Program, ECOG= Eastern Cooperative Oncology Group, PPS= Palliative Performance Scale, ACP/GCD Tracking Record= Advance Care Planning/Goals of Care Designation

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**Step 3: PRIMARY PROVIDER  
MANAGEMENT OF UNMET NEEDS**

Primary palliative care management of unmet patient needs is strongly recommended using the four essential components of an early palliative care approach as a guide.



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**Step 3: Illness  
Comprehension and Coping**



Evidence has shown that addressing a patient’s understanding of their illness, how they are coping and how those close to them are coping, is associated with improved patient outcomes. In addition to consulting secondary palliative care providers, below are some other methods that can be used to support patients and family/caregivers.

	<b>Method</b>	<b>Description</b>
Specialist provided	Spiritual care	Existential and faith based supports
	Art therapy, music therapy	Facilitated creative art therapies
	Psychosocial counselling	Including: psychotherapy, Cognitive Behavioral Therapy, family therapy, individual and group therapy, CALM (Managing Cancer and Living Meaningfully)
Provided by family physicians, oncologists and nurses (with training)	Vital talk	Strategies and clinician resources for responding to difficult conversations
	SICP	Program of resources, education workshop and systems change processes to increase meaningful conversations between people living with serious illness and their clinicians about their values and priorities
	Dignity conserving therapy (including life review)	Question guides and approaches e.g. asking “what do I need to know about you as a person to give you the best care possible”

Patient preference, available time and clinician skills may be factors in the extent to which exploring patient illness understanding and coping occurs during clinical encounter, and clinician judgment is required about whether and when to refer for additional formal coping supports. This guideline encourages primary providers to explore and identify techniques to use in their practice.

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Step 3: Management of symptoms by providers and patient self management

Providers

Patient

Management of symptoms is a cornerstone of palliative care. Symptom management tips sheets are available at [www.ahs.ca/guru](http://www.ahs.ca/guru) under Palliative & Supportive Care:

- Anxiety
- Constipation
- Decreased appetite and cachexia
- Delirium
- Depression
- Diarrhea
- Dyspnea
- Fatigue
- Malignant bowel obstruction
- Nausea and Vomiting
- Oral care
- Sleep disturbance
- Pain
- Tenesmus (a feeling, sometimes painful, of constantly needing to pass stool despite an empty rectum)

Patient self-management strategies can enhance symptom management, ability to cope and quality of life. Providers should encourage use of the following resources:

- Telephone triage or nurse navigator available at each Alberta Cancer Centre
- [Healthlink \(811\)](#) for health questions, or [Inform Alberta \(211\)](#) for questions about services available within the community
- Patients on Palliative Home Care can call for support 24/7
- [CancerControl Alberta Patient Education Resources](#)

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Step 3: Advance care  
planning patient's  
preferred method of  
decision making

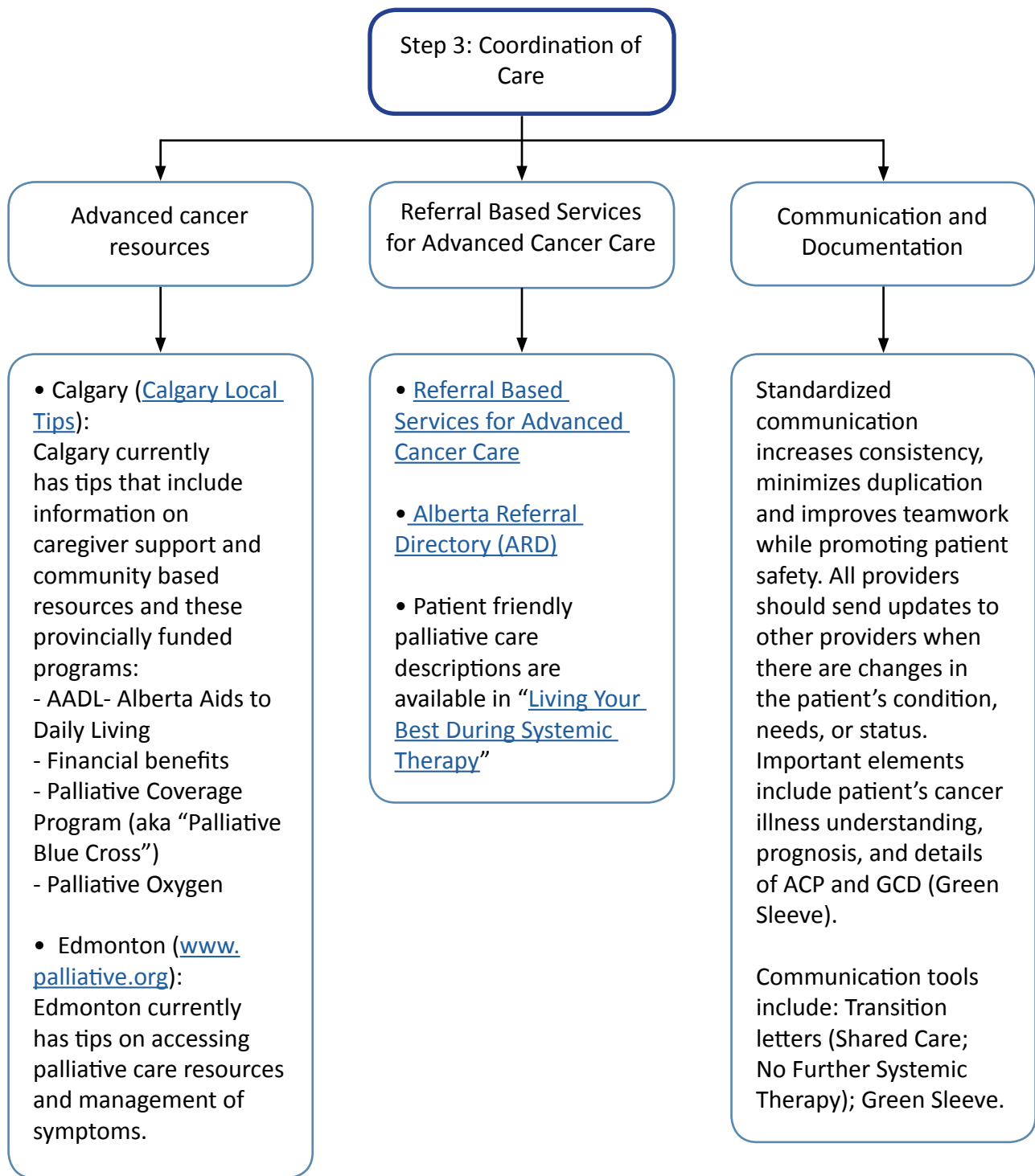
Encourage patients to have conversations about ACP throughout cancer treatment. Clinicians, within their scope of practice, can be instrumental in:

- Encouraging reflection on and expression of personal values, beliefs and preferences related to personal goals, health care and their preferred method of decision making (e.g. shared decision making, supported decision making with family/caregiver, self-determined, physician-directed)
- Sharing prognostic information, tailored to the patient's readiness and preferences around illness understanding.
- Supporting the patient in selecting an appropriate ADM.
- Encouraging the patient to communicate with their ADM, family/caregivers, other health care team members about their values, beliefs and preferences and who they have selected as ADM.
- Encouraging the patient to create a Personal Directive.
- Documenting ACP conversations on the ACP GCD Tracking Record.

**Resources:**

- Patient and health care provider resources are available at [www.conversationsmatter.ca](http://www.conversationsmatter.ca)
- [Serious Illness Care Program](#) (SICP)

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#### Step 4: EXPLORING END OF LIFE TOPICS

Clinicians are recommended to support patients who are exploring end of life topics.

##### End of Life Planning

- Review ACP discussions and patient preferences for care at end of life (e.g. switch to focus on comfort care near end of life).
- Revisit caregiver supports.
- Ensure patient is on Palliative Coverage Program.
- Develop a plan for expected death in the community if patient would like to remain at home:
  - AHS Guideline on [Expected death in the home](#)

##### Estate and Funeral Planning

- Remind patient & their families to begin this planning early on, including wills and financial enduring power of attorney. Connect to appropriate resources (e.g. social work) as needed.

##### Hospice Access

- Hospice care is a specialized service available in certain locations that provides 24/7 facility based care to those approaching end of life and whose needs cannot be met at home.
- There are highly specific criteria for referral to hospice and access is often determined by secondary palliative care clinicians. Refer to "Service Description" or talk to your local palliative care clinician about whether hospice may be appropriate.

##### Medical Assistance in Dying

- AHS has developed the [Medical Assistance in Dying Policy](#) to guide medical assistance in dying services. A [physician FAQ](#) sheet is available.

##### Grief and Bereavement

- Ask patients about their emotional supports.
- Let patients and caregivers know that grief and bereavement resources are available both before and after death.
- Most services are accessed by self-referral and clinicians can help by providing the access information. Refer to "Service Description" document.

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