

Palliative Care

History of Palliative Care (1)

1967	St Christopher's Hospice founded (UK)
1975	Royal Victoria Hospital (Canada) opens palliative care ward
1981	Seirei Mikatahara General Hospital (Shizuoka Prefecture) establishes hospice
1984	Yodogawa Christian Hospital (Osaka Prefecture) establishes hospice ward
1990	Palliative care ward admissions established as medical fee item
1994	Home medical management established as medical fee item
2002	Palliative care surcharge established as medical fee item
2006	Cancer Control Act passed
2007	Basic Plan to Promote Cancer Control Programs (1 st term) starts; palliative care from early treatment
2008	Palliative care training for physicians involved in cancer care starts (basic palliative care, raising public awareness)
2012	Basic Plan to Promote Cancer Control Programs (2 nd term) starts; earlier start of palliative care, from cancer diagnosis Outpatient palliative care management established as medical fee item Local at-home palliative care coordination initiative starts (Training for at-home care support physicians)

Partially revised from documents submitted to Conference on Palliative Care Promotion (16 Mar 2016)

History of Palliative Care (2)

Apr 2012	Conference on palliative care promotion launched (- Mar 2016)
Sep 2012	Conference on palliative care promotion: Interim report
2013	Palliative care advancement initiative launched (Improvement of palliative care centers)
Jan 2014	Guidelines for Advancement of Designated Cancer Care Hospitals revised for Palliative Care WHO 'Global Atlas of Palliative Care at the End of Life' issued
Dec 2015	Accelerated Cancer Control Plan starts
Apr 2016	New medical fee items established: At-home palliative care support surcharge for clinics/hospitals Cancer outpatient at-home coordination guidance Palliative care ward emergency admission initial surcharge
May 2016	Conference on cancer palliative care Advancement launched (-May 2018)
Nov 2017	Palliative Care Provision for Cardiovascular Disorders Patients working group established (-Apr 2018)
Dec 2017	Palliative care training workshop guideline revised, including cancer
Apr 2018	Palliative care surcharge for advanced heart failure patients added to medical fee table
Mar 2019	Conference on living well with and beyond cancer
Jul 2021	Conference on living well with and beyond cancer
Mar 2023	Basic Plan to Promote Cancer Control Programs (4 th term) starts Promotion of palliative care from the time of cancer diagnosis Palliative care is shifted from the symbiotic field of the basic plan to the medical field.

Partially revised from documents submitted to Conference on Palliative Care Promotion (16 Mar 2016)

Palliative Care Workshops

- For medical professionals, including cancer care -

1. Context

In Dec 2016, the Cancer Control Act (Law No. 98, 2006) was amended to incorporate palliative care; the Conference on Cancer Palliative Care Advancement advised that palliative care should be provided beyond cancer patients, and that training should extend to other medical professionals, not only physicians and dentists. Thus, workshops were established.

2. Objective

Developing an accurate understanding of basic palliative care, with knowledge of, techniques and ways of approaching palliative care, to ensure participants may provide appropriate palliative care from diagnosis.

3. Participants

- All cancer care physicians/dentists
 - Staff at designated cancer care hospitals
 - Staff at-home care support clinics/hospitals working with designated cancer care hospitals and at hospitals with palliative care wards
- Other palliative care healthcare workers

4. Workshops mode

- E-learning
- In-person sessions



5. Syllabus

i) Required

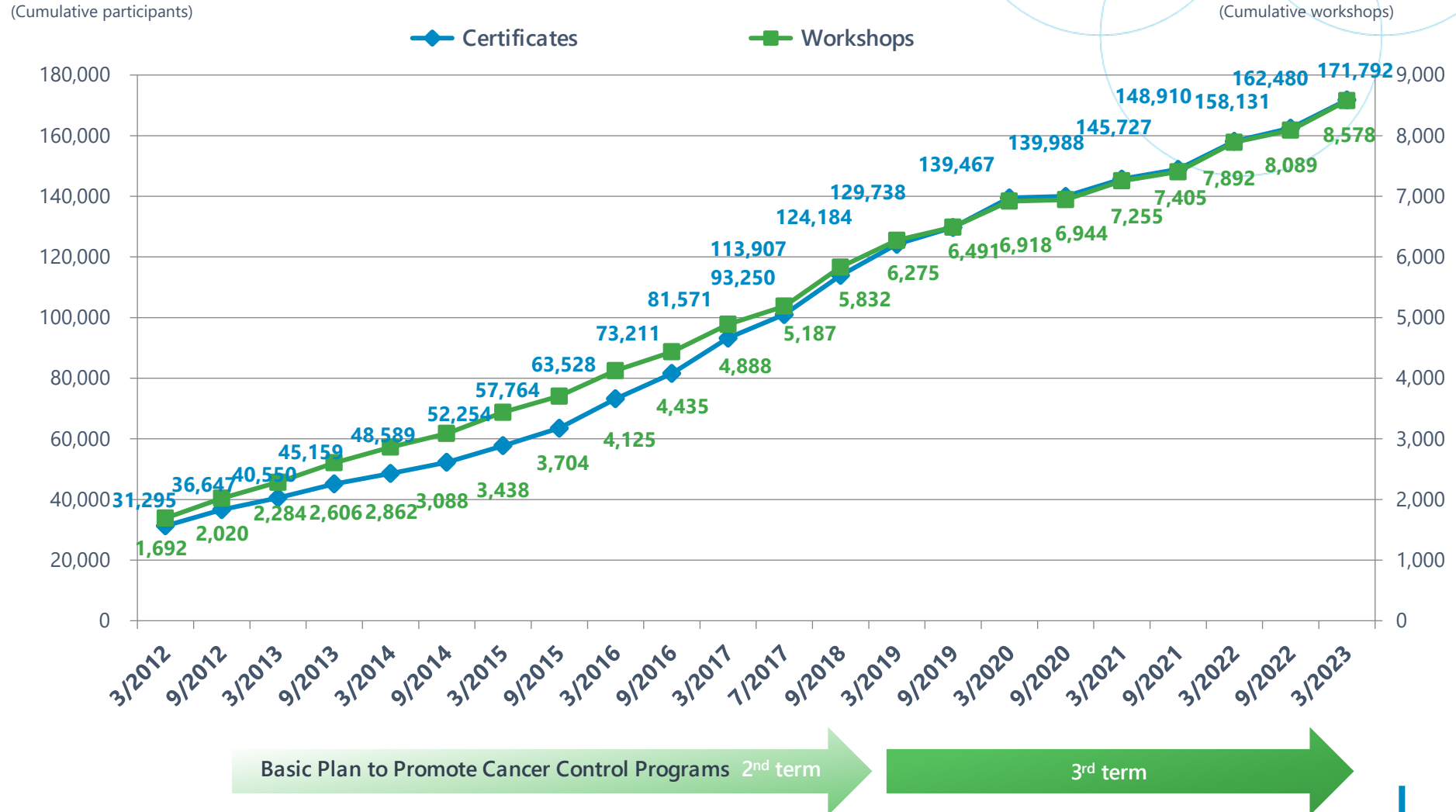
Holistic palliative care with the patient's perspective/ pain screening followed by symptoms relief, referrals to specialists/ cancer pain evaluation and management/ care of dyspnea, digestive tract symptoms, anxiety, depression, delirium/ communication/ care facility decisions, regional care network, at-home palliative care/ advance care planning, care for families and the bereaved

ii) Optional

Non-cancer palliative care, physical palliative care for discomfort other than pain, dyspnea, digestive tract symptoms/ mental palliative care for discomfort other than anxiety, depression, and delirium/ alleviation of symptoms through palliative radiotherapy, nerve blocks/ palliative care for social suffering



Palliative Care Workshops/Participation



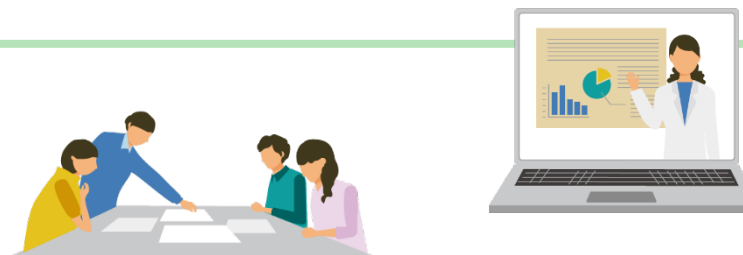
Palliative Cancer Care Training Initiative

Project Overview

- Article 17 of the Cancer Control Act decrees “palliative care suited to patient conditions provided from diagnosis,” and “training for medical professionals towards improving quality of cancer patients’ lives.” The Conference on Cancer Palliative Care Advancement advice lead to expand palliative care to non-cancer patients.
- To build capacity, palliative care e-learning platforms and workshops are organized to raise the level and quality of palliative care for cancer, and to raise public awareness of palliative care for cancer.

Palliative Care Training

- Basic training for all medical professionals
- Logistical management of workshops
- Operational support for of e-learning lecture platforms



Training of Instructors

Focus on lecturers/instructors capable of planning/running workshops



To the Public

Disseminating reliable information and raising awareness of its importance



(Commissioned to the Japanese Society for Palliative Medicine)

Cancer Patients in the Final Stage of Receiving Care - A survey




Current Status and Issues

- Palliative care at the final stage of cancer care can only be comprehended by surveying those directly involved in the patients' care. To raise the Quality of Life (QOL) for both patients and their families, surveying the bereaved is required.
- The Cancer Control Promotion Council notes that earlier surveys of bereaved families are biased as they are conducted through Designated Cancer Care Hospitals.
- In some countries, surveys are conducted on samples extracted from death registries.
- Acceleration Plan for Cancer Control (dec 2015) dictates to conduct a survey of bereaved families to comprehend a clearer picture of patients in final stages receiving medical and nursing care, in order to improve their quality of life, working with related organizations.

Inconsistent results due to survey methods

	Research A			Research B		
Method	Survey bereaved, via medical institutions			Survey bereaved identified from general public		
Feature	Mostly designated hospitals and palliative care wards			Mostly general hospitals		
Doctors promptly dealt with patient's difficult symptoms	Designated Hospital	Palliative care ward	Hospice at Home	Hospital	Palliative care ward	At-home care
	56%	78%	77%	39%	56%	52%

Surveys on bereaved family members (Based on death registrations)

	 United Kingdom	 America	 Italy
Number of subjects surveyed	22,292 Persons	1,578 Persons	1,289 Persons
method	Post	Phone interview	Interview
survey item	Quality of care pain/symptoms/ treatments communication use of decision-making support services	Physical pain emotional support decision-making dignity support for family	Quality of care pain/symptoms/ treatments communication overall satisfaction socio-economic issues

Overview

To be conducted with support from patient advocacy groups, by sampling cases from death questionnaires of [Vital Statistics](#). A [survey of bereaved families to be carried out, towards improving quality of life of cancer patients.](#)

Palliative Care Promotion Initiative (Advancement of Palliative Care Centers)

