



PALLIATIVE CARE AND ADVANCED CANCER

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GYNECOLOGIC CANCER SOCIETY

BACKGROUND

- Care as a continuum.
- Impact of incurable diseases, role of a multidisciplinary approach.
- QOL & Patient – Centered Medicine.
- Family Members & Caregivers.
- Disparity/Inequity.

DEFINITION

- PALLIATIVE CARE IS AN APPROACH THAT IMPROVES THE QUALITY OF LIFE OF PATIENTS AND THEIR FAMILIES FACING THE PROBLEM ASSOCIATED WITH LIFE-THREATENING ILLNESS, THROUGH THE PREVENTION AND RELIEF OF SUFFERING BY MEANS OF EARLY IDENTIFICATION AND IMPECCABLE ASSESSMENT AND TREATMENT OF PAIN AND OTHER PROBLEMS, PHYSICAL, PSYCHOSOCIAL AND SPIRITUAL.

WHO DEFINITION

- Holistic.
- Not only pain.
- All dimensions.
- Multidisciplinary team.
- Relatives/Caregivers.
- Loss of hope.

IAHPC DEFINITION 2018

PALLIATIVE CARE IS THE ACTIVE HOLISTIC CARE OF INDIVIDUALS ACROSS ALL AGES WITH SERIOUS HEALTH-RELATED SUFFERING DUE TO SEVERE ILLNESS AND ESPECIALLY THOSE NEAR THE END OF LIFE. IT AIMS TO IMPROVE THE QUALITY OF LIFE OF PATIENTS, THEIR FAMILIES AND THEIR CAREGIVERS.

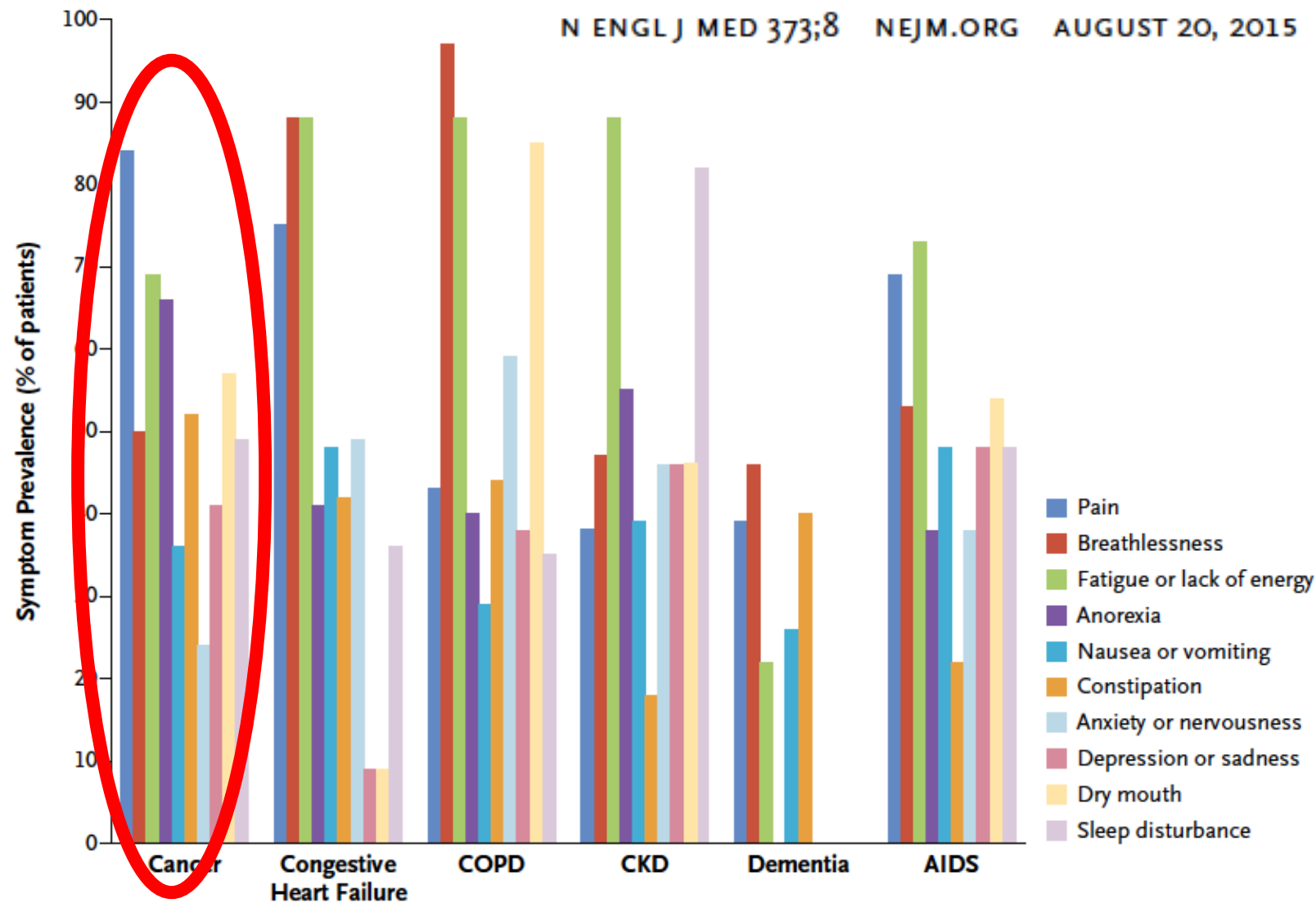


Figure 1. Symptom Prevalence in Advanced Illness.

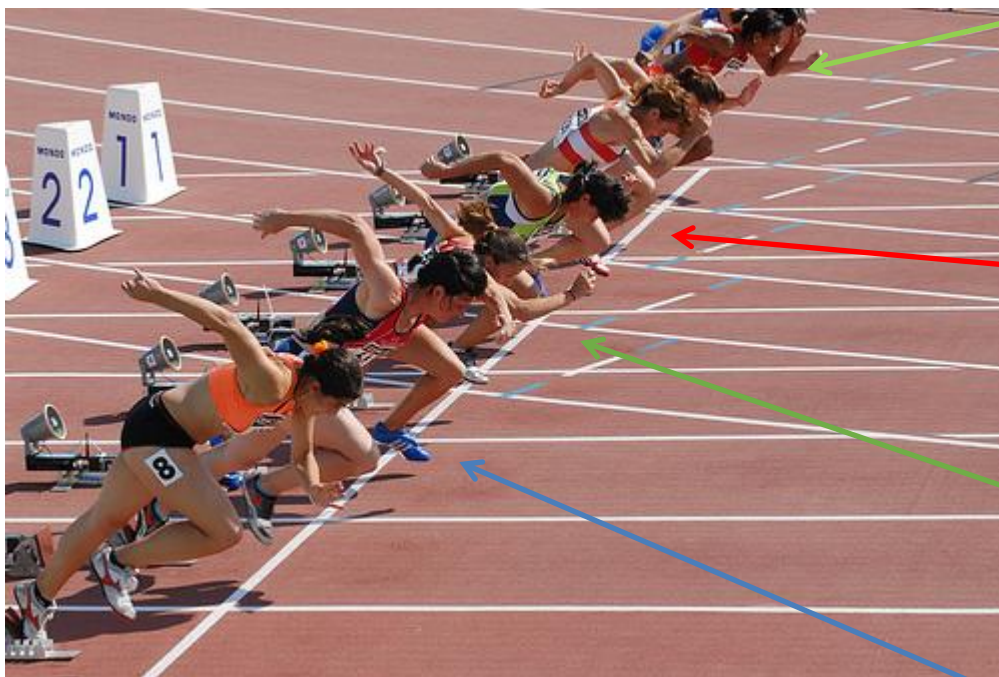
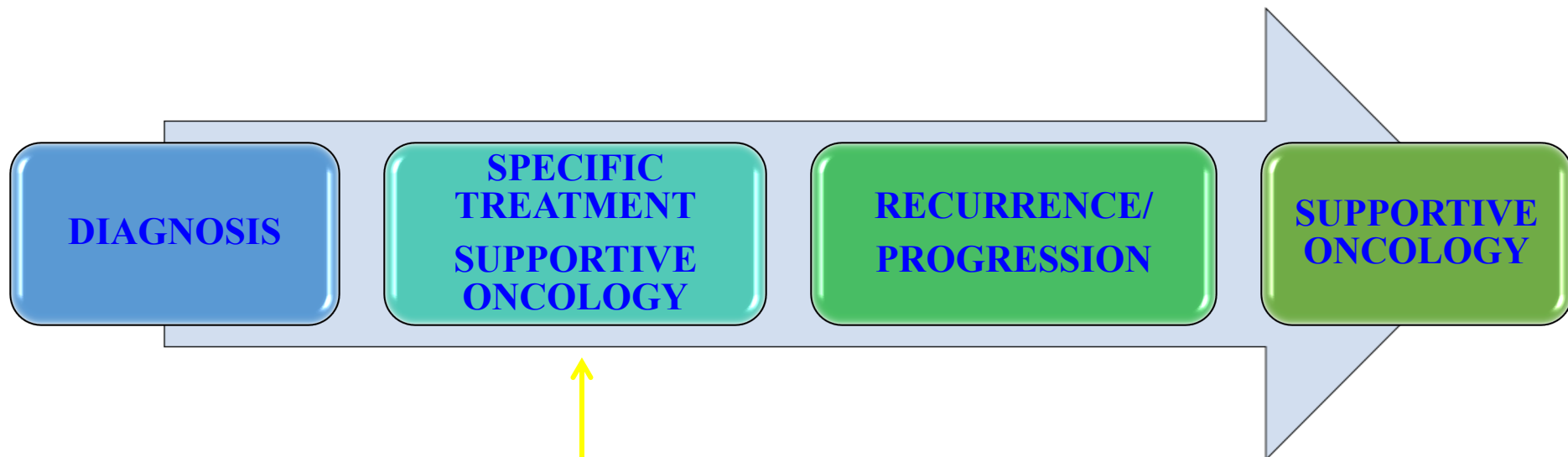
Data are from representative studies of symptom prevalence among patients with cancer,⁸⁻¹² congestive heart failure,^{13,14} chronic obstructive pulmonary disease (COPD),¹⁵ chronic kidney disease (CKD),^{13,14} or dementia^{16,17} and among patients who received highly active antiretroviral therapy for the acquired immunodeficiency syndrome (AIDS).¹⁸ Self-reported data regarding some symptoms were unavailable for patients with dementia.

Traditional palliative care



Early palliative care





SUPPORTIVE ONCOLOGY

RADIOTHERAPY

MEDICAL ONCOLOGY

SURG/GYN ONCOLOGY



Cochrane
Library

Cochrane Database of Systematic Reviews

Early palliative care for adults with advanced cancer (Review)

Haun MW, Estel S, Rücker G, Friederich HC, Villalobos M, Thomas M, Hartmann M

Cochrane Database of Systematic Reviews 2017, Issue 6. Art. No.: CD011129.

DOI: [10.1002/14651858.CD011129.pub2](https://doi.org/10.1002/14651858.CD011129.pub2).

Figure 4. Forest plot of comparison: I Health-related quality of life, outcome: I.1 Health-related quality of life.

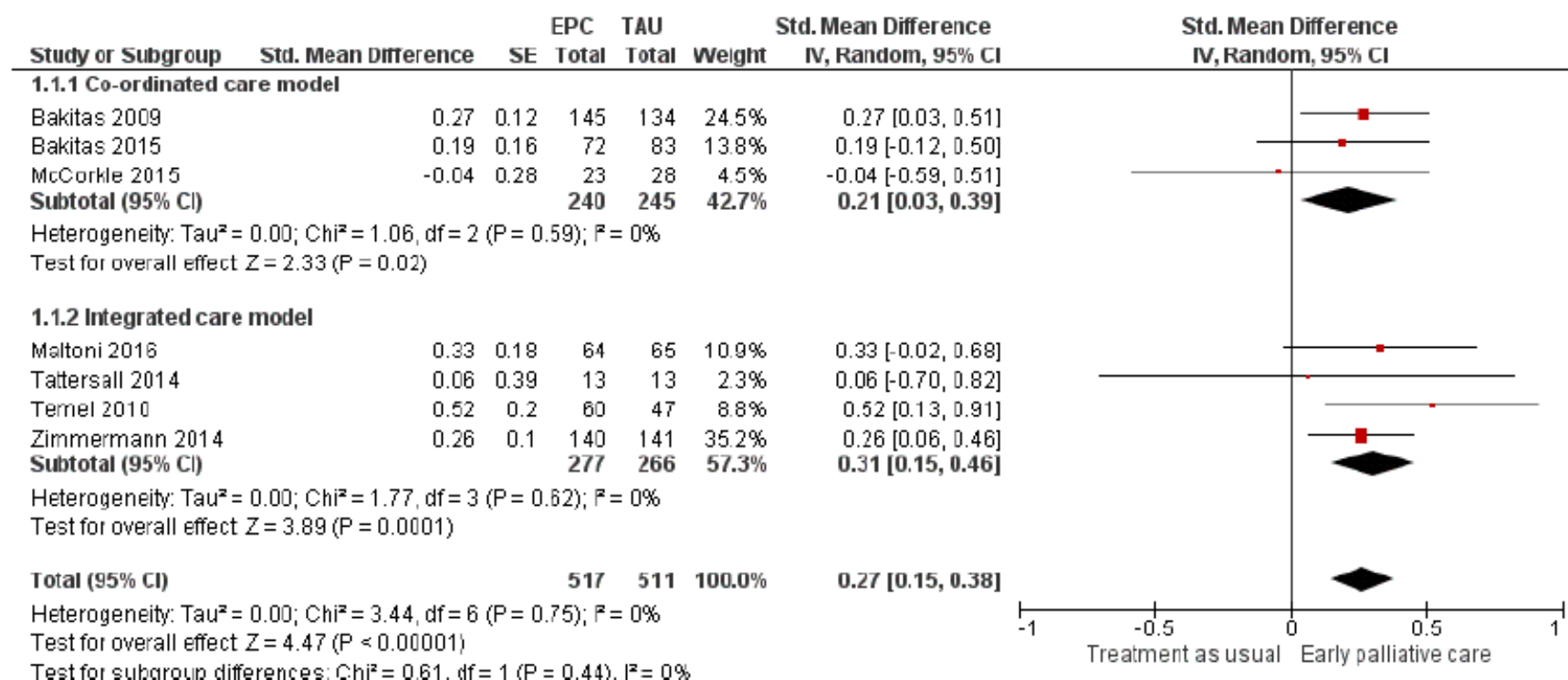


Figure 7. Forest plot of comparison: I Early palliative care vs standard oncological care, outcome: I.4 Symptom intensity.

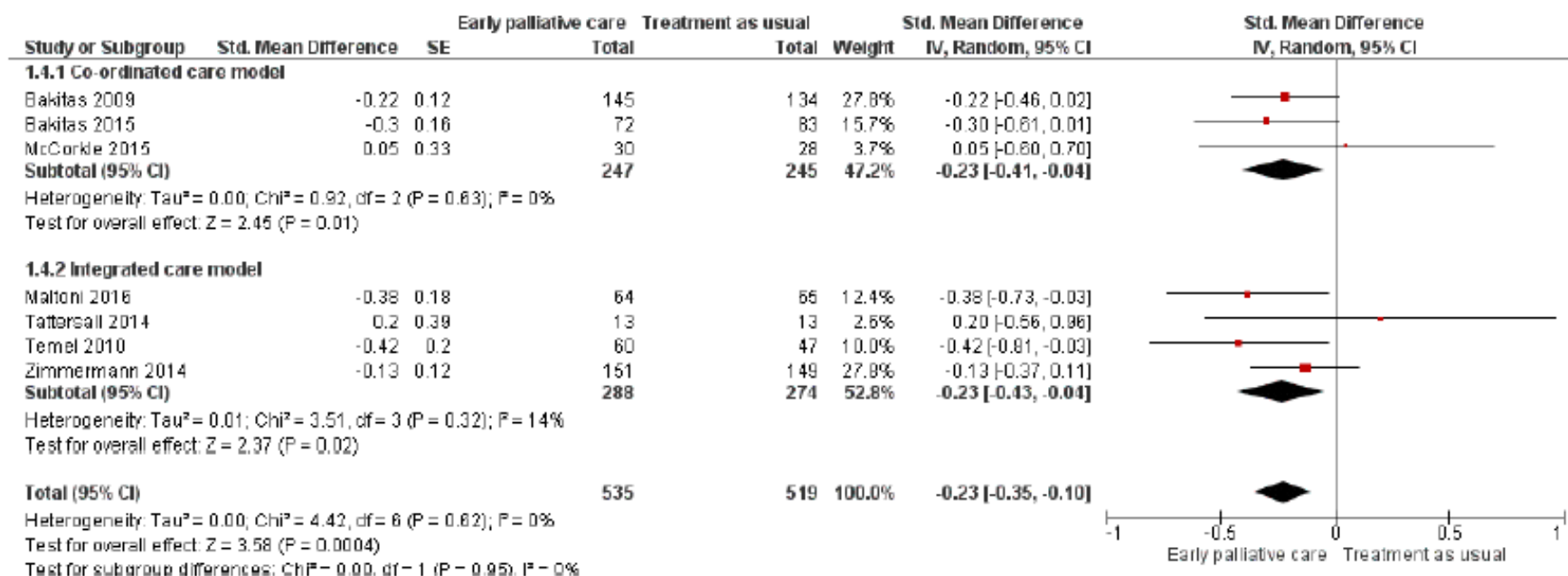


Figure 5. Forest plot of comparison: I Early palliative care vs TAU, outcome: I.2 Survival.

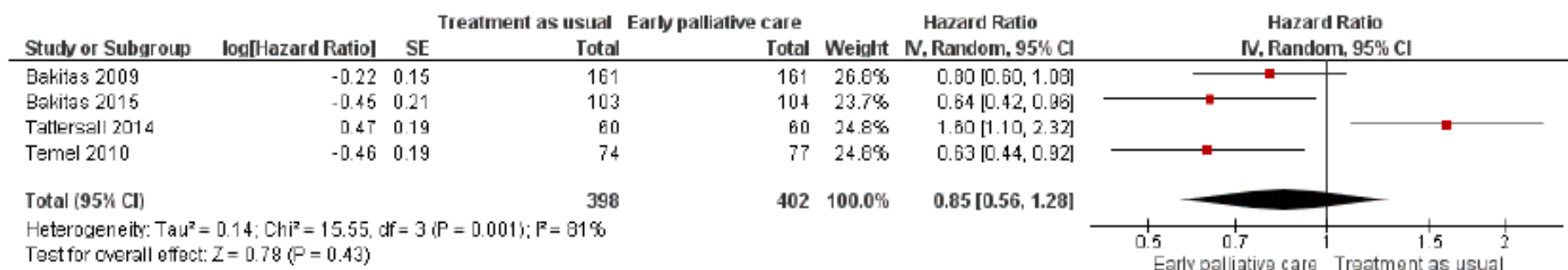
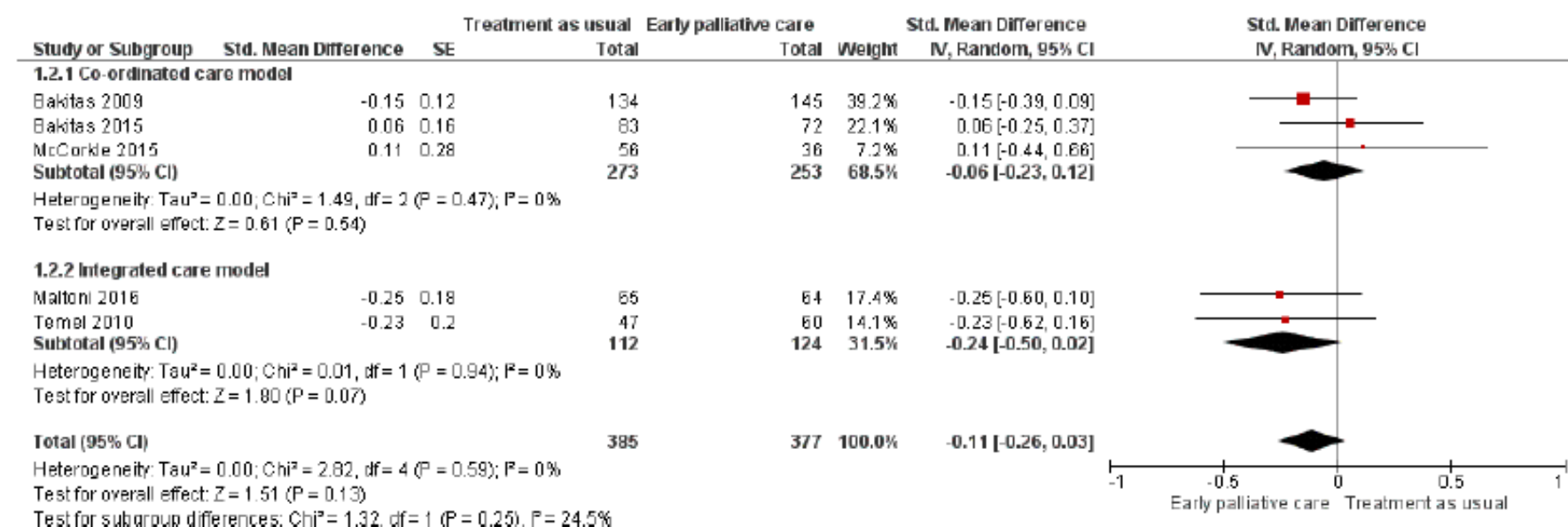


Figure 6. Forest plot of comparison: I Early palliative care vs standard oncological care, outcome: I.2 Depression.



Strengthening of palliative care as a component of comprehensive care throughout the life course

The Sixty-seventh World Health Assembly,

Having considered the report on strengthening of palliative care as a component of integrated treatment throughout the life course;¹

Recalling resolution WHA58.22 on cancer prevention and control, especially as it relates to palliative care;

Integration of Palliative Care Into Standard Oncology Care: American Society of Clinical Oncology Clinical Practice Guideline Update

Betty R. Ferrell, City of Hope Medical Center, Duarte, CA; Jennifer S. Temel and Jeffrey M. Peppercorn, Massachusetts General Hospital; Tracy A. Balboni,

Betty R. Ferrell, Jennifer S. Temel, Sarah Temin, Erin R. Alesi, Tracy A. Balboni, Ethan M. Basch, Janice I. Firn, Judith A. Paice, Jeffrey M. Peppercorn, Tanyanika Phillips, Ellen L. Stovall,† Camilla Zimmermann, and Thomas J. Smith

J Clin Oncol 34. © 2016 by American Society of Clinical Oncology



Definition of Palliative Care^{a,c}

Palliative care is an approach to patient/family/caregiver-centered health care that focuses on optimal management of distressing symptoms, while incorporating psychosocial and spiritual care according to patient/family/caregiver needs, values, beliefs, and cultures. The goal of palliative care is to anticipate, prevent, and reduce suffering and to support the best possible quality of life for patients/families/caregivers, regardless of the stage of the disease or the need for other therapies. Palliative care can begin at diagnosis and should be delivered concurrently with disease-directed, life-prolonging therapies and should facilitate patient autonomy, access to information, and choice.

Palliative care becomes the main focus of care when disease-directed, life-prolonging therapies are no longer effective, appropriate, or desired. Palliative care should be provided by the primary oncology team and augmented as needed by collaboration with an interdisciplinary team of palliative care experts.

Standards of Palliative Care^{b,c}

- Institutions should develop processes for integrating palliative care into cancer care, both as part of usual oncology care and for patients with specialty palliative care needs.
- All cancer patients should be screened for palliative care needs at their initial visit, at appropriate intervals, and as clinically indicated.
- Patients/families/caregivers should be informed that palliative care is an integral part of their comprehensive cancer care.
- Educational programs should be provided to all health care professionals and trainees so that they can develop effective palliative care knowledge, skills, and attitudes.
- Palliative care specialists and interdisciplinary palliative care teams, including board-certified palliative care physicians, advanced practice nurses, physician assistants, social workers, chaplains, and pharmacists, should be readily available to provide consultative or direct care to patients/families/caregivers and/or health care professionals who request or require their expertise.
- Quality of palliative care should be monitored by institutional quality improvement programs.



INDICATIONS^{d,e}

One or more of the following:

- Uncontrolled symptoms
- Moderate-to-severe distress related to cancer diagnosis and cancer therapy
- Serious comorbid physical and psychosocial conditions
- Complex psychosocial needs
- Poor prognostic awareness
- Potentially life-limiting disease
- Metastatic solid tumors
- Patient/family/caregiver concerns about course of disease and decision-making
- Patient/family/caregiver requests for palliative care
- Patient request of hastened death

- Inform patients/families/caregivers about palliative care services
 - Anticipate symptoms and discuss preventative measures

ASSESSMENT

- Benefits/burdens of anticancer therapy
- Personal goals/values/expectations
- Symptoms
- Psychosocial or spiritual distress
- Educational and informational needs
- Cultural factors affecting care
- Criteria for consultation with palliative care specialist ([See PAL-7](#))

PALLIATIVE CARE INTERVENTIONS^e

- Anticancer therapy
- Appropriate treatment of comorbid physical and psychosocial conditions
- Coordination of care with other health care providers
- Symptom management
- Advance care planning
- Psychosocial and spiritual support
- Culturally appropriate care
- Resource management/social support
- Consultation with palliative care specialist
- Hospice referral
- Response to request to withdraw or withhold life-sustaining treatment
- Response to requests for hastened death (physician-assisted dying)
- Care of imminently dying hospitalized patient
- Palliative sedation

REASSESSMENT

Acceptable outcomes:

- Patient satisfied with response to anticancer therapy
- Adequate symptom management
- Reduction of patient/family/caregiver distress
- Acceptable sense of control
- Decreased caregiver burden
- Strengthened relationships
- Optimized quality of life
- Personal growth and enhanced meaning

If unacceptable

- Re-evaluate intervention options and intensify as possible

AFTER-DEATH INTERVENTIONS

- For family and caregiver(s):
 - Immediate after-death care
 - Bereavement support
 - Cancer risk assessment and modification
- For health care team:
 - General support (ie, debriefing)

→ Death →

Ongoing reassessment

Ongoing reassessment

Lancet Oncology Commission



Integration of oncology and palliative care: a *Lancet Oncology* Commission



PATIENT - CENTRED CARE

ELEMENTS

- Respect for patients' values, preferences and expressed needs.
- Coordination & integration of care.
- Information, communication & education.
- Physical comfort (bothersome symptoms).
- Emotional support (fear & anxiety).
- Involvement of family & friends.

RECOMMENDATIONS

- Integrate the concept “patient – centred care”.
- Implement routine use of patient – reported outcomes (PROMs).
- Integrate shared – decision making and Advanced Care Planning.
- Involve and assess the family as part of early integration of cancer palliative care.
- Develop the content and basic method of standardised care pathway for use as a tool for early integration of pall care into oncology.
- Mandatory training of oncology & pall care specialists in patient – centred care.

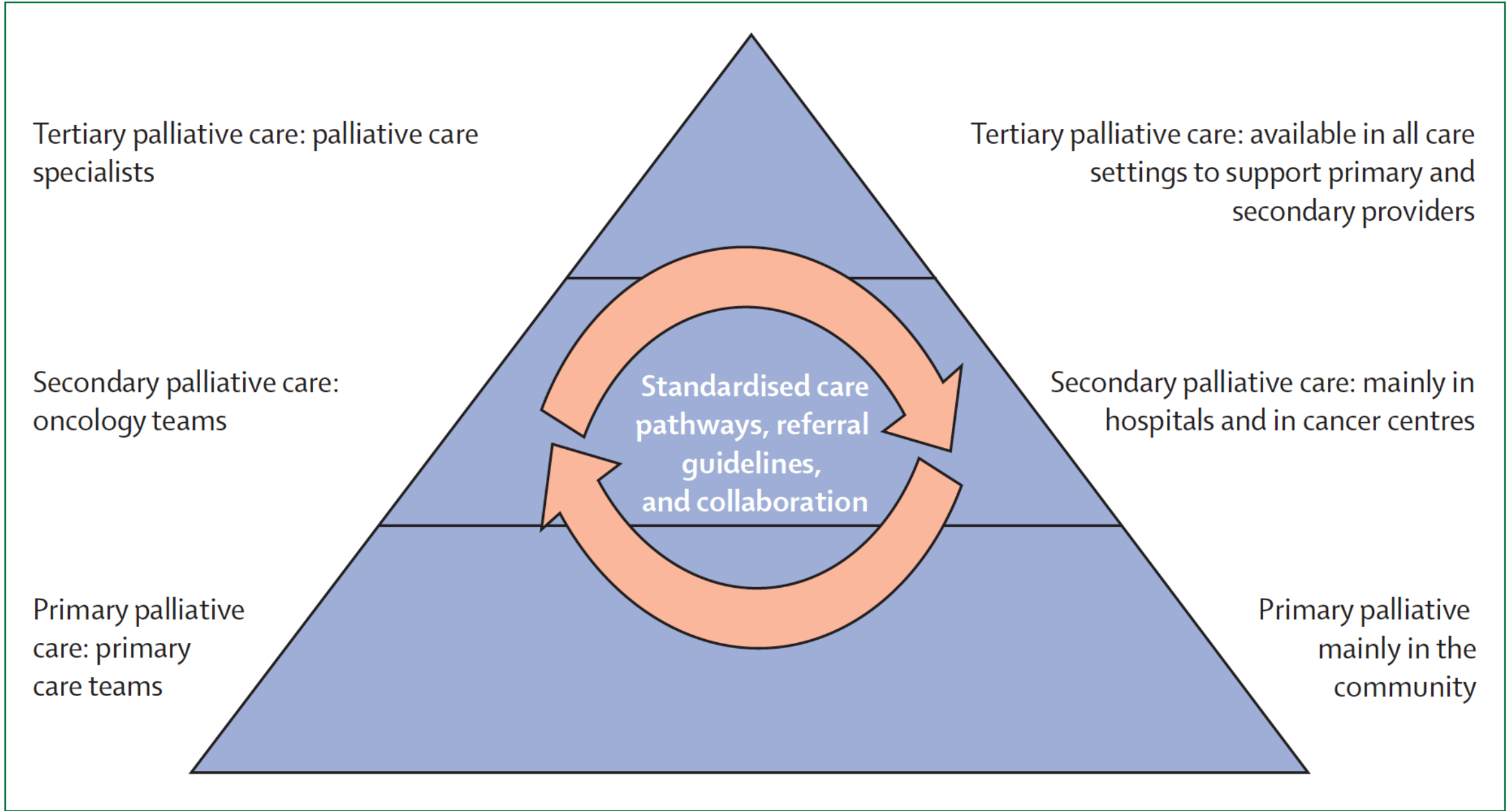


Figure 5: Proposed model of optimal oncology palliative care provision, including integration across providers and settings

PATIENT'S MANAGEMENT

DISEASE-DIRECTED THERAPIES

- Disease-directed therapies. .
- Treatment options: balance risks & benefits.
- Personalized treatments.

SUPPORTIVE ONCOLOGY

- Symptom management
- Psychosocial Oncology Services.
- Patient-Physician Communication (prognosis, end-of life.
- Advance care planning.
- End of Life.

PALL CARE/SUPPORTIVE ONCOLOGY AND GYNECOLOGICAL MALIGNANCIES

TREATMENT – RELATED

A. Surgery.

B. Chemotherapy:
-Neuropathic pain.

C. Radiotherapy:
-Fistulae.
-Actinic Disease.
-Lymphedema.

ADVANCED DISEASE

A. General Symptom
Burden.

B. Specific Symptoms:
-Central Pelvic Recurrence.
-Pelvic Sidewall
Recurrence.
-Peritoneal Carcinomatosis.



**Malignant Ureteric Obstruction.
Malignant Bowel Obstruction.
Fistulae/Vaginal Discharge.**



PALL CARE/SUPPORTIVE ONCOLOGY AND BREAST CANCER

TREATMENT – RELATED

A. Surgery.

B. Chemotherapy:
-Neuropathic pain.

C. Radiotherapy:
-Actinic Disease.
-Lymphedema.

ADVANCED DISEASE

A. General Symptom
Burden.

B. Specific Symptoms:
-Bone/Brain Metastases.
-Breast Lump:

-Ulcerated.

-Malodorous.

-Hemorrhagic.

-Nodal Metastases:

-Superior Cava Vein
Syndrome.



Global Atlas of Palliative Care at the End of Life



Rates for people in need of palliative care

All ages and both sexes per 100,000 (including adults and children)

178.36 - 272.85

272.86 - 281.84

281.85 - 307.16

307.17 - 467.52

Data not available

Not applicable

0 2500 5000 7500 10000 Kilometers



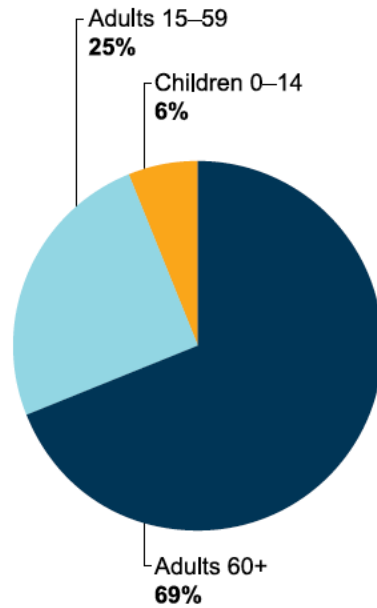
Rates for adults in need of palliative care

Both sexes per 100,000

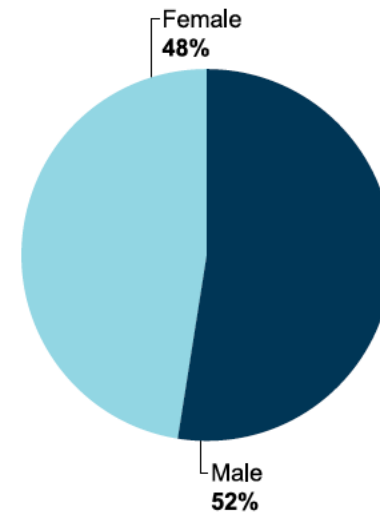
- 234.12 - 353.39
- 353.40 - 385.78
- 385.79 - 378.45
- 378.46 - 561.52
- Data not available
- Not applicable

0 2'500 5'000 7'500 10'000 Kilometers

DISTRIBUTION PER AGE AND SEX

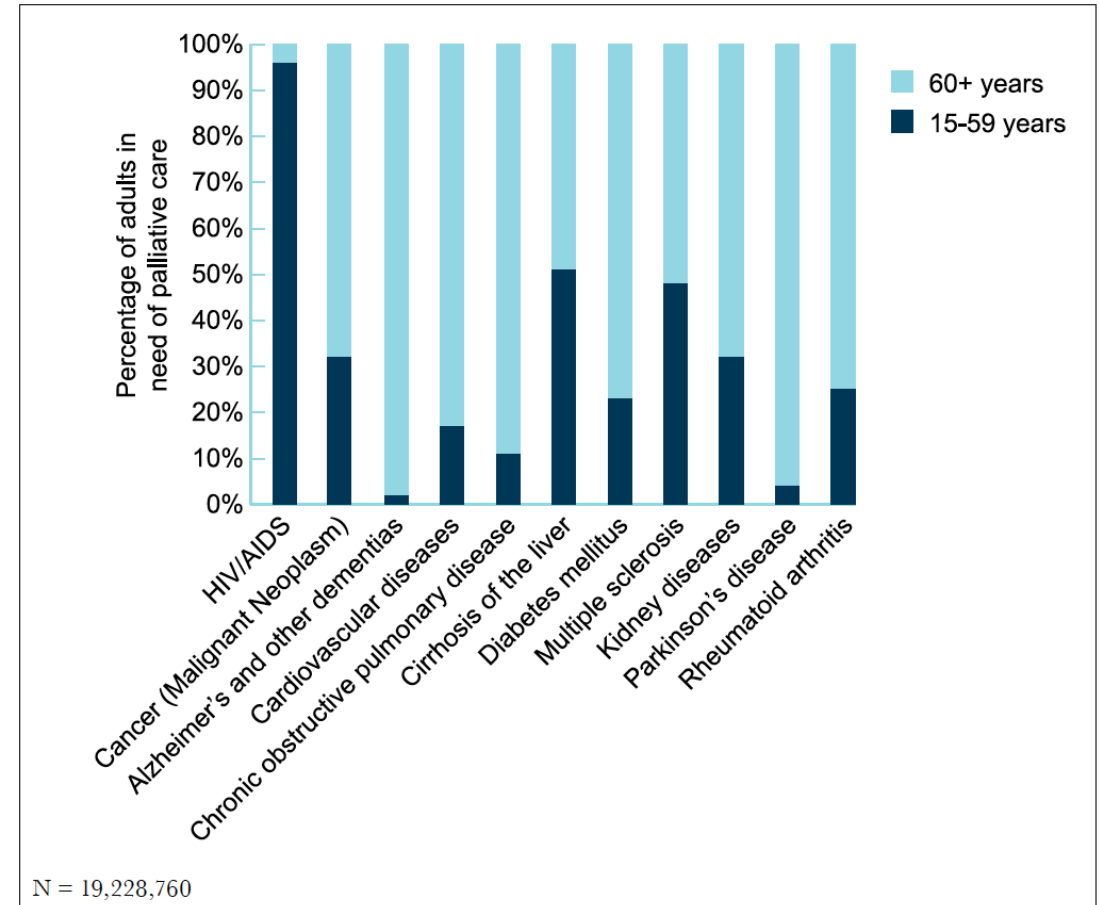
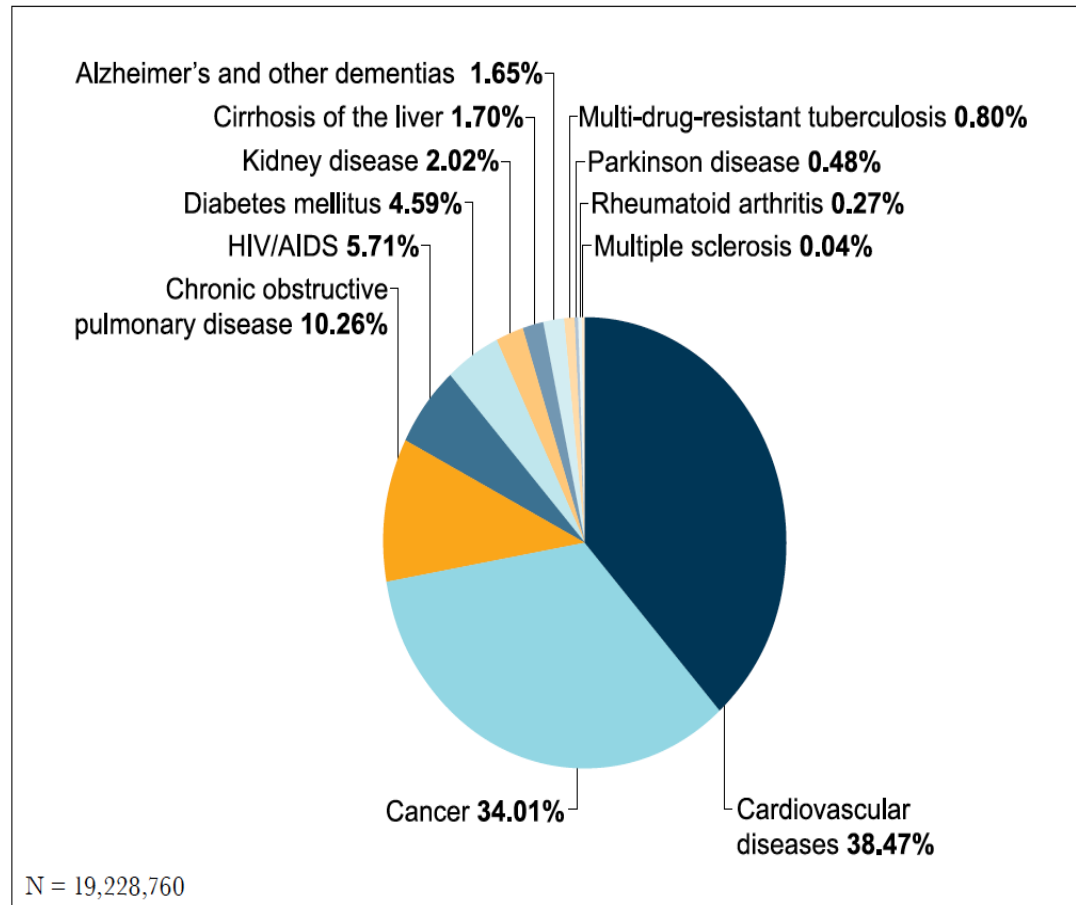


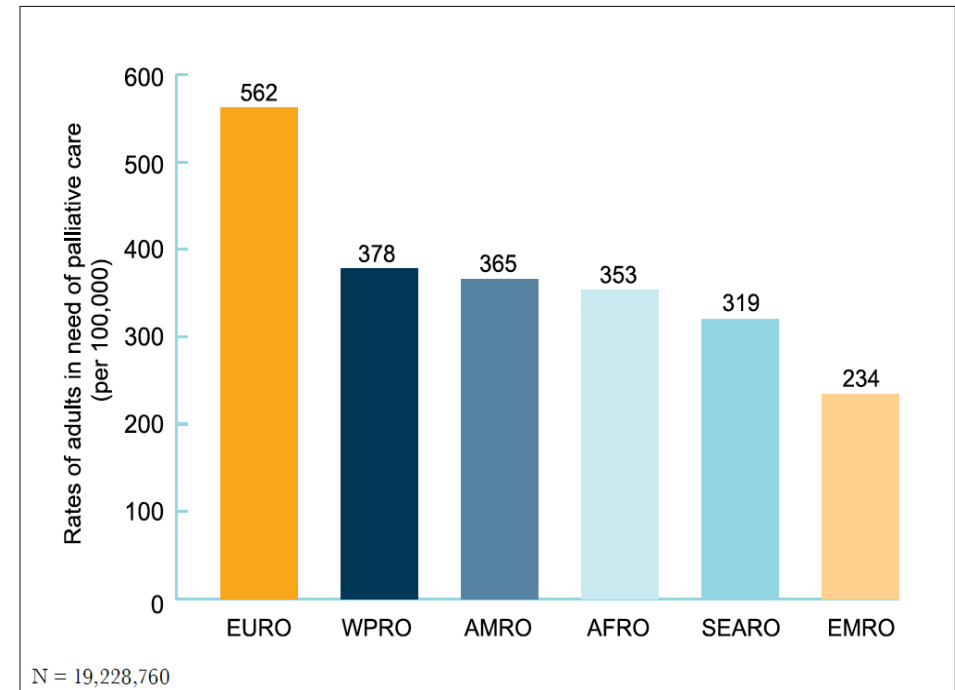
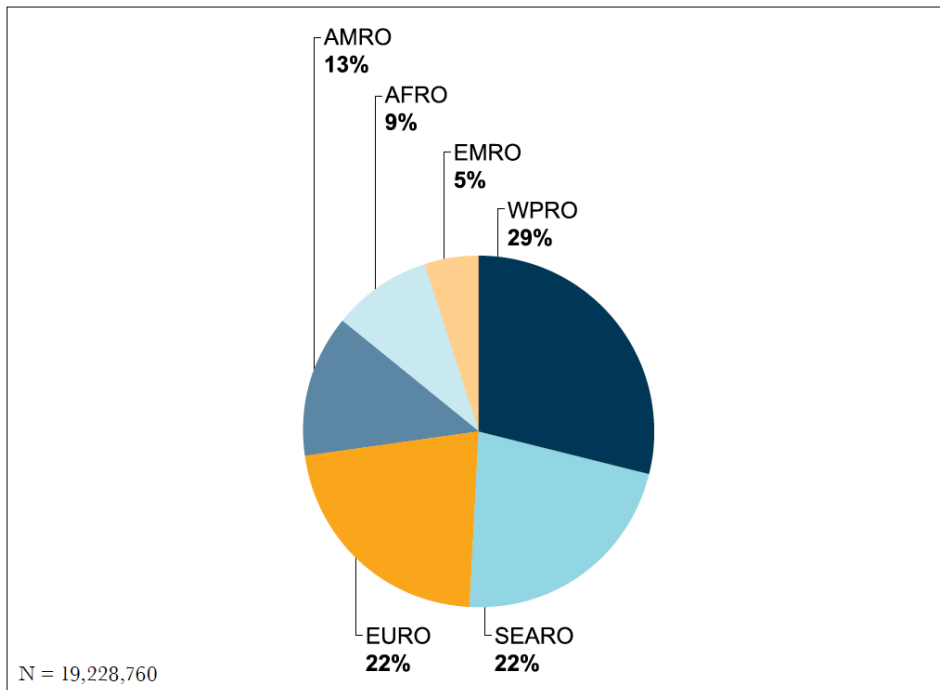
N = 20,398,772



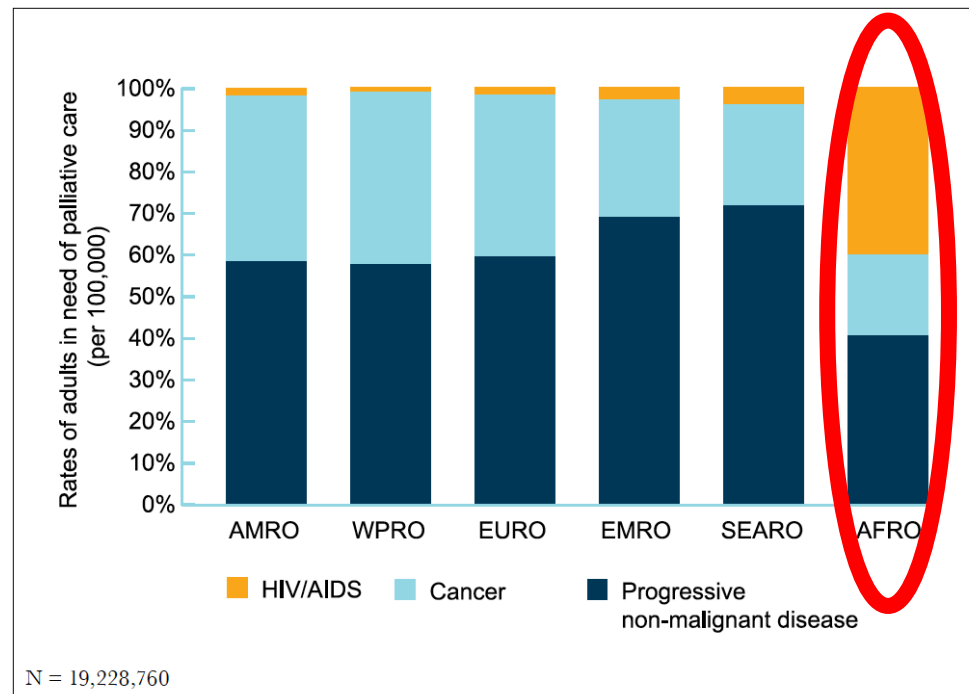
N = 19,228,760

TYPE OF INCURABLE DISEASES IN ADULTS

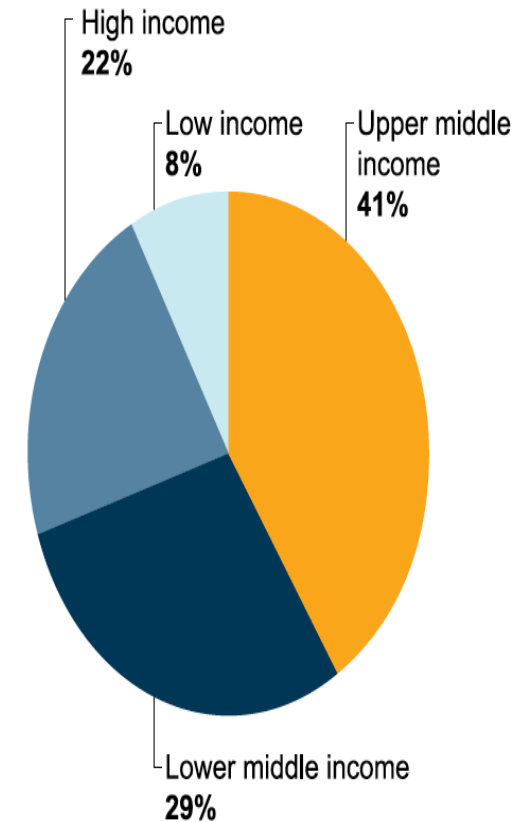
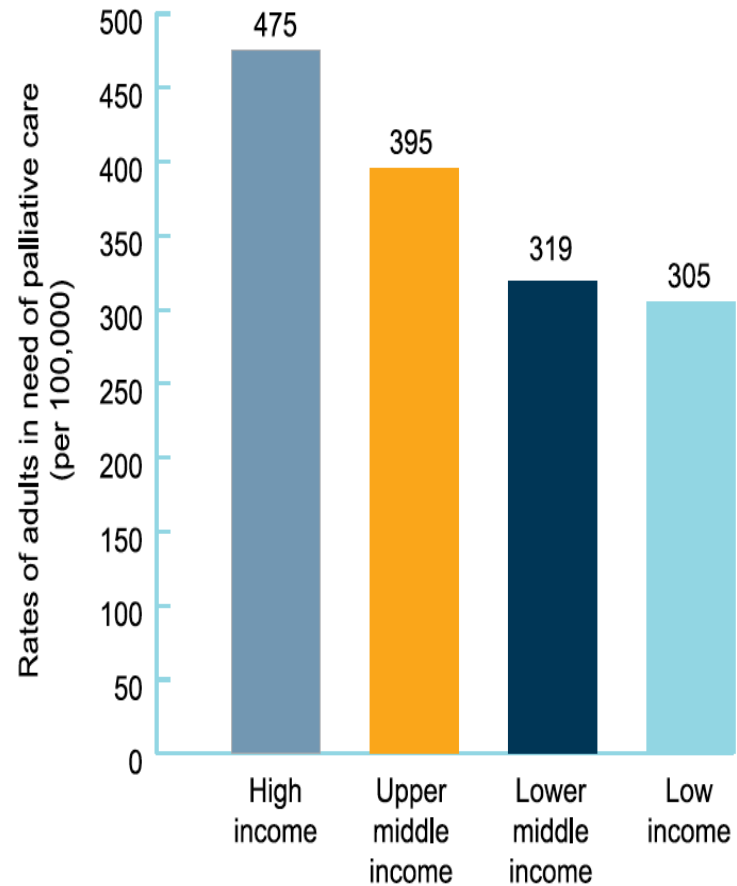




WPRO: Western Pacific.
 EURO: Europe.
 SEARO: South East.
 AMRO: Americas.
 AFRO: Africa.
 EMRO: Eastern Mediterranean.



DISTRIBUTION ACCORDING TO WORLD BANK CLASSIFICATION



N = 19,228,760

CANCER AND HIV NEED FOR PALLIATIVE CARE



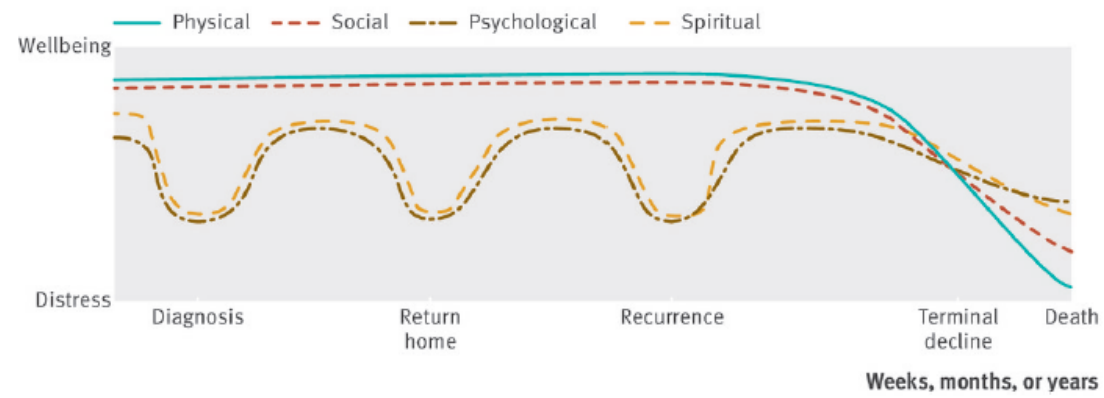


Fig 1 Wellbeing trajectories in patients with conditions such as cancer causing rapid functional decline

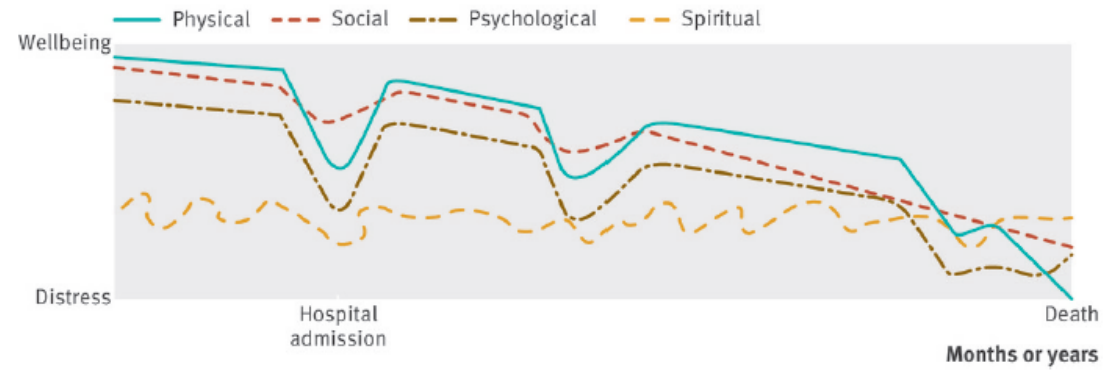


Fig 2 Wellbeing trajectories in patients with intermittent decline (typically organ failure or multimorbidity)

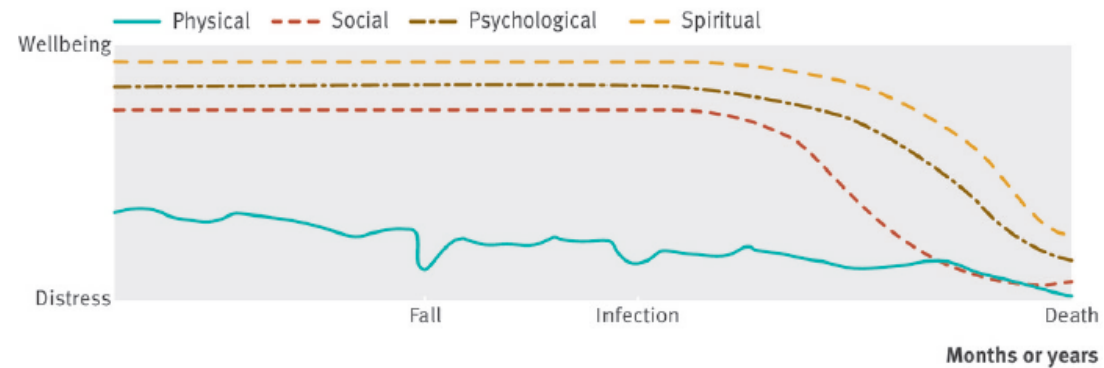


Fig 3 Wellbeing trajectories in patients with gradual decline (typically frailty or cognitive decline)

BARRIERS

- Policy.
- Education.
- Medications.
- Implementation.

Policy

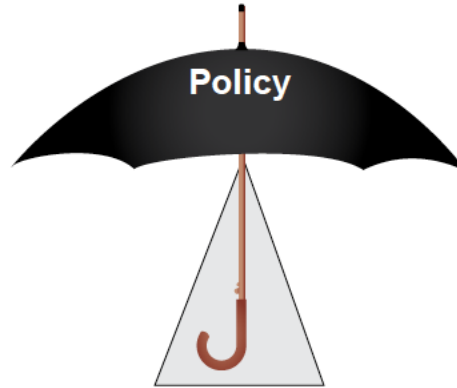
- Palliative care part of national health plan, policies, related regulations
- Funding/service delivery models support palliative care delivery
- Essential medicines

(Policy makers, regulators, WHO, NGOs)

Medicine availability

- Opioids, essential medicines
- Importation quota
- Cost
- Prescribing
- Distribution
- Dispensing
- Administration

(Pharmacists, drug regulators, law enforcement agents)



Education

- Media and public advocacy
- Curricula, courses – professionals, trainees
- Expert training
- Family caregiver training and support

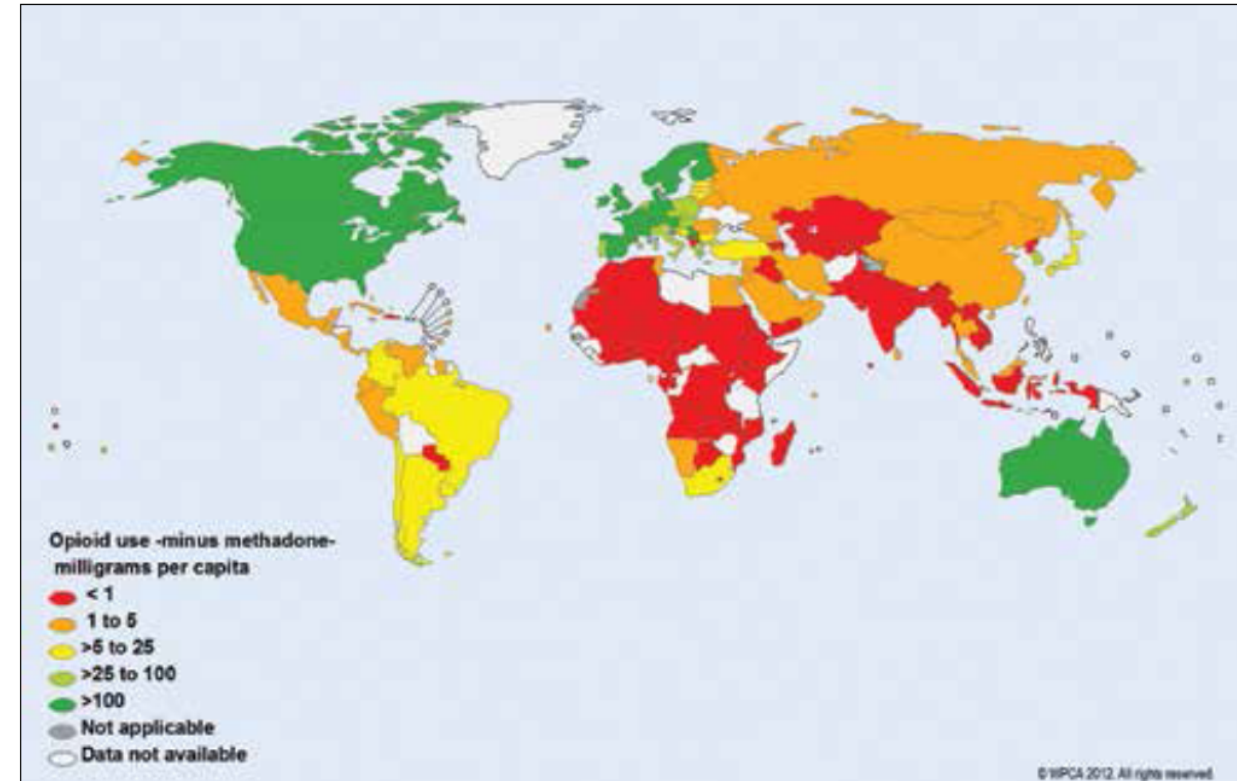
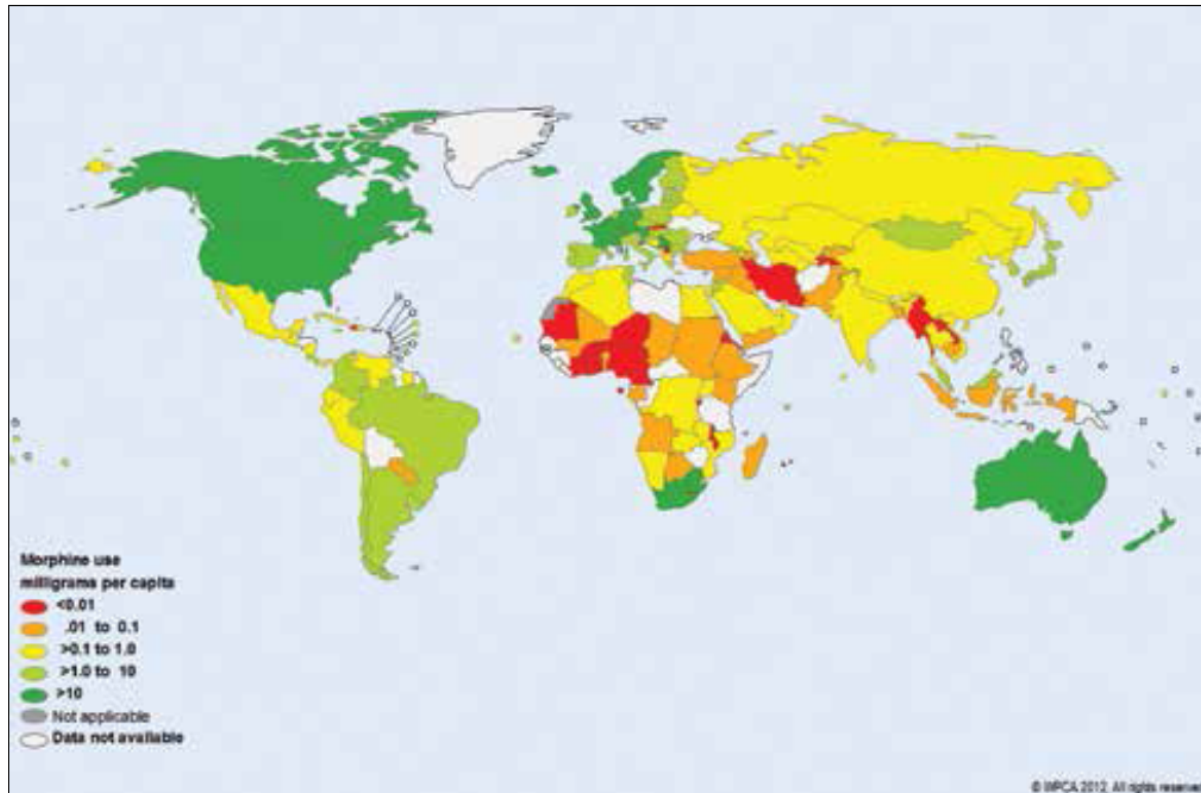
(Media and public, healthcare providers and trainees, palliative care experts, family caregivers)

Implementation

- Opinion leaders
- Trainer manpower
- Strategic and business plans – resources, infrastructure
- Standards, guidelines measures

(Community and clinical leaders, administrators)

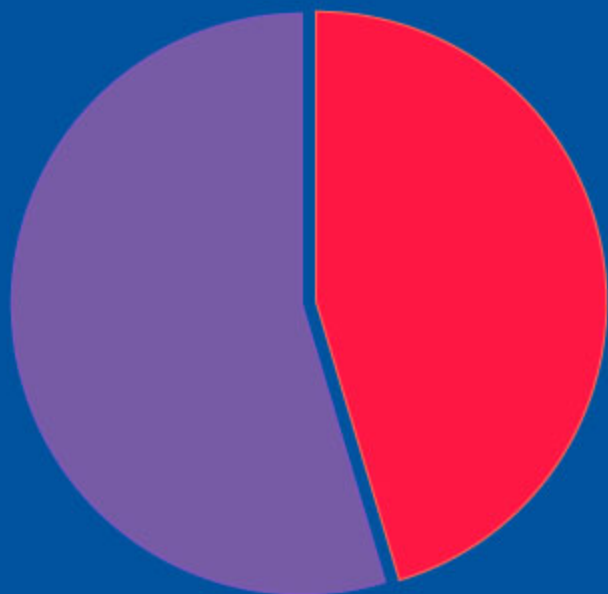
MORPHINE AND OPIOID CONSUMPTION WORLDWIDE



Alleviating the access abyss in palliative care and pain relief— an imperative of universal health coverage: the *Lancet* Commission report



Felicia Marie Knaul, Paul E Farmer, Eric L Krakauer*, Liliana De Lima, Afsan Bhadelia, Xiaoxiao Jiang Kwete, Héctor Arreola-Ornelas, Octavio Gómez-Dantés, Natalia M Rodriguez, George A O Alleyne, Stephen R Connor, David J Hunter, Diederik Lohman, Lukas Radbruch, María del Rocío Sáenz Madrigal, Rifat Atun†, Kathleen M Foley†, Julio Frenk†, Dean T Jamison†, M R Rajagopal†, on behalf of the Lancet Commission on Palliative Care and Pain Relief Study Group‡*



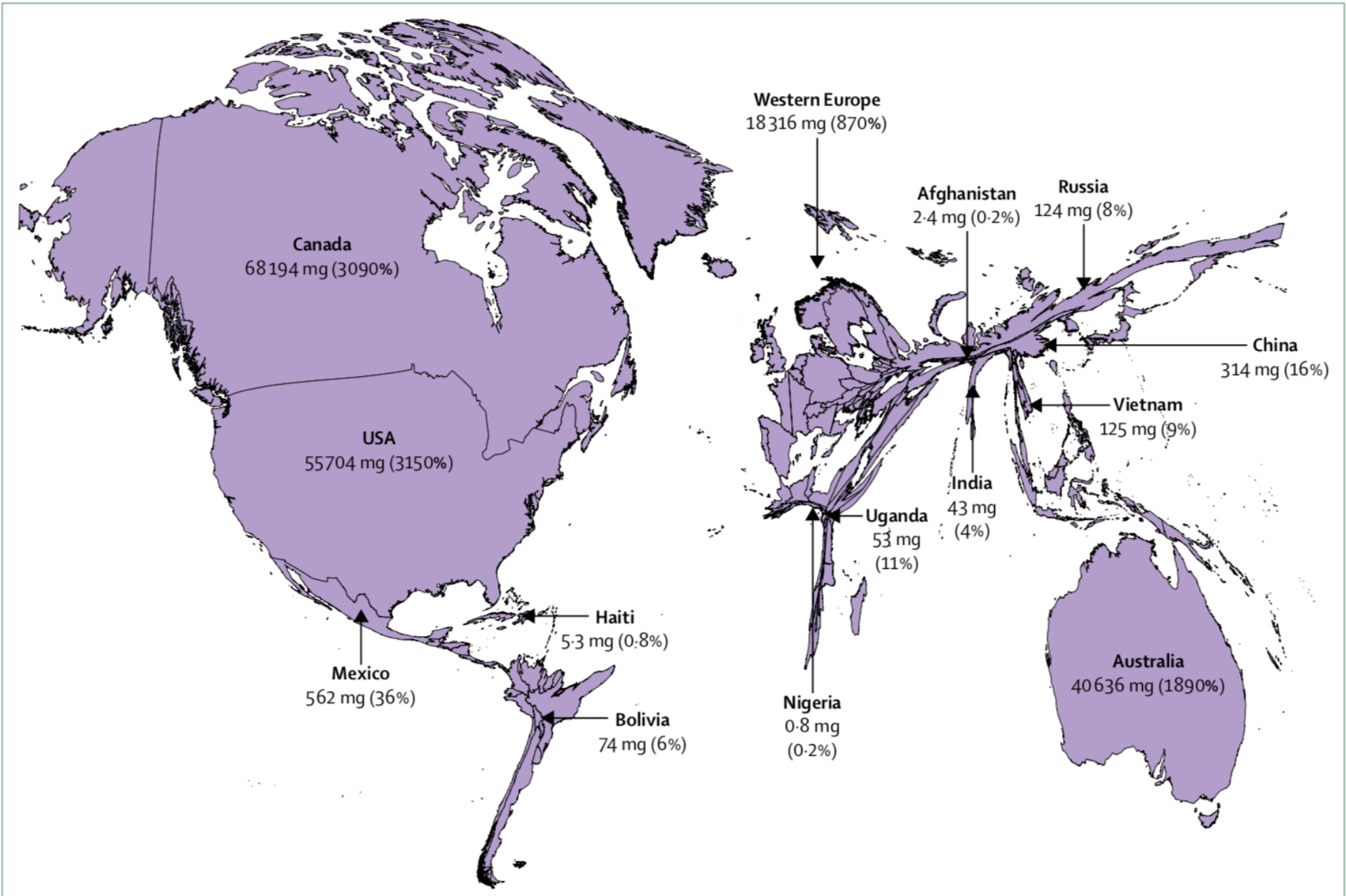
Every year, 25.5 million people die with serious health-related suffering that requires palliative care. That is nearly half of all deaths.

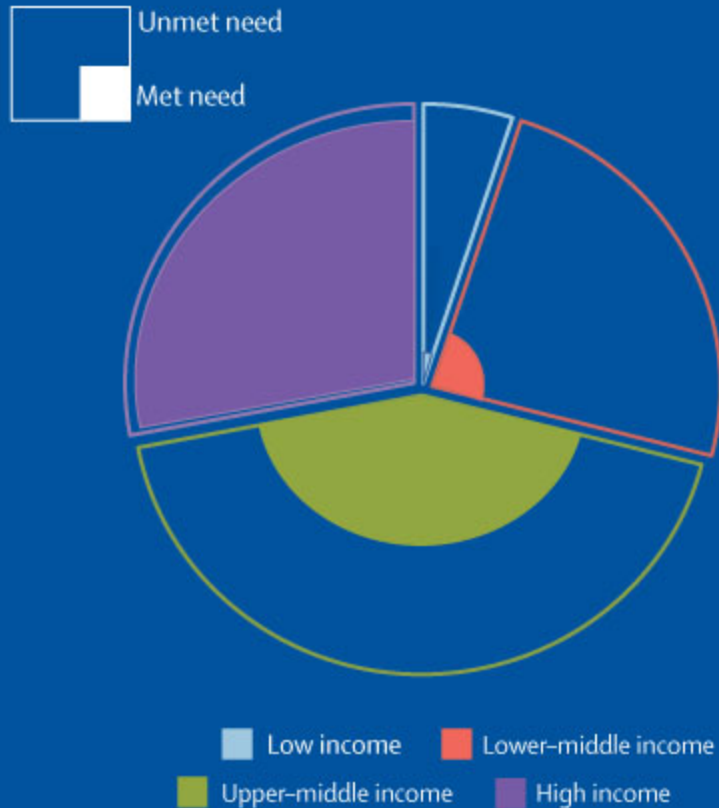
Over 80% live in low-income and middle-income countries where access is severely lacking.

The Lancet Commission on Global Access to Palliative Care and Pain Relief

THE LANCET

The best science for better lives





Immediate-release morphine is an essential component of palliative care, yet over 97% of need in low-income and middle-income countries goes unmet.

Providing pain relief for all those with unmet medical need could cost as little as US\$145 million.

The Lancet Commission on Global Access to Palliative Care and Pain Relief

THE LANCET

The best science for better lives

Panel 2: An Essential Package Of Palliative Care And Pain Relief Health Services

The Essential Package contains the inputs for safe and effective provision of essential palliative care and pain relief interventions to alleviate physical and psychological symptoms, including the medicines and equipment that can be safely prescribed or administered in a primary care setting. The list of essential medicines in the Essential Package is based on WHO's list of essential medicines,³⁵ and considers the medicines, doses, and administration routes for palliative care for both adults and children.

The Essential Package is designed to be lowest cost by including only off-patent formulations, frugal innovation for needed equipment, and a staffing model based on competencies rather than professions. Tasks often undertaken by specialised medical personnel in high-income countries can be performed by other specialised and general practitioners and nurses or by community health workers empowered with the necessary training and medical supervision to participate effectively in the delivery of palliative care and pain treatment at all levels of care, from the hospital to the home.^{36,37}

With the key exception of morphine, the medicines in the Essential Package are available in most countries even if supply is limited. For morphine, an essential palliative care medicine, assuring safety and accessibility is complex. Ensuring a balance between appropriate medical access to controlled medicines and the prevention of their diversion and non-medical use is crucial, and the Commission not only designed appropriate human resource models but also the strategies to provide the complementary policy and stewardship to expand access to an Essential Package that includes morphine.³²

The health services of the Essential Package must be complemented by interventions for the relief of social and spiritual suffering to preserve the dignity of patients, facilitate access to health interventions, and prevent financial hardship and impoverishment. Yet, these social supports are neither part of the remit of health ministries nor should they be financed from a health budget.

Antipoverty and social development policies, publicly funded safety nets, programmes, and ministries must give special attention to ensure that families do not sacrifice their basic needs in desperate attempts to care for loved ones. These persons with life-limiting or life-threatening health conditions and their families should be mainstreamed into existing social support and social welfare programmes, yet they are often ignored, excluded, or marginalised, preventing them from being effectively integrated into these programmes.

Medicines

- Amitriptyline
- Bisacodyl (Senna)
- Dexamethasone
- Diazepam
- Diphenhydramine (chlorpheniramine, cyclizine, or dimenhydrinate)
- Fluconazole
- Fluoxetine or other selective serotonin-reuptake inhibitors (sertraline and citalopram)
- Furosemide
- Hyoscine butylbromide
- Haloperidol
- Ibuprofen (naproxen, diclofenac, or meloxicam)
- Lactulose (sorbitol or polyethylene glycol)
- Loperamide
- Metaclopramide
- Metronidazole
- Morphine (oral immediate-release and injectable)
- Naloxone parenteral
- Omeprazole
- Ondansetron
- Paracetamol
- Petroleum jelly

Medical equipment

- Pressure-reducing mattress
- Nasogastric drainage or feeding tube
- Urinary catheters
- Opioid lock box
- Flashlight with rechargeable battery (if no access to electricity)
- Adult diapers (or cotton and plastic, if in extreme poverty)
- Oxygen

Human resources (varies by referral, provincial or district hospital, community health center, or home)

- Doctors (specialty and general, depending on level of care)
- Nurses (specialty and general)
- Social workers and counsellors
- Psychiatrist, psychologist, or counsellor (depending on level of care)
- Physical therapist
- Pharmacist
- Community health workers
- Clinical support staff (diagnostic imaging, laboratory technician, nutritionist)
- Non-clinical support staff (administration, cleaning)

Additional detail is provided in the additional online material.



special article

Palliative Care in the Global Setting: ASCO Resource-Stratified Practice Guideline

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Nahla Gafer
Harmala K. Gupta
Susan Horton
Graciela Jacob
Ruiuo Jia
Frank L. Lu
Daniela Mosoiu
Christina Puchalski
Carole Seigel
Olaitan Soyannwo
James F. Cleary

executive summary

Purpose The purpose of this new resource-stratified guideline is to provide expert guidance to clinicians and policymakers on implementing palliative care of patients with cancer and their caregivers in resource-constrained settings and is intended to complement the *Integration of Palliative Care Into Standard Oncology Care: American Society of Clinical Oncology Clinical Practice Guideline Update* of 2016.

Methods ASCO convened a multidisciplinary, multinational panel of experts in medical oncology, family medicine, radiation oncology, hematology/oncology, palliative and/or hospice care, pain and/or symptom management, patient advocacy, public health, and health economics. Guideline development involved a systematic literature review, a modified ADAPTE process, and a formal consensus-based process with the Expert Panel and additional experts (consensus ratings group).

Results The systematic review included 48 full-text publications regarding palliative care in resource-constrained settings, along with cost-effectiveness analyses; the evidence for many clinical questions was limited. These provided indirect evidence to inform the formal consensus process, which resulted in agreement of $\geq 75\%$ (by consensus ratings group including Expert Panel).

Recommendations The recommendations help define the models of care, staffing requirements, and roles and training needs of team members in a variety of resource settings for palliative care. Recommendations also outline the standards for provision of psychosocial support, spiritual care, and opioid analgesics, which can be particularly challenging and often overlooked in resource-constrained settings. Additional information is available at www.asco.org/resource-stratified-guidelines.

Recommendations It is the view of ASCO that health care providers and health care system decision makers should be guided by the recommendations for the highest stratum of resources available. The guideline is intended to complement but not replace local guidelines.

- CHILEAN EXPERIENCE.

- IGCS ENVISION.

TAKE HOME MESSAGES

- Real need for palliative care.
- Importance of multidisciplinary approach and not only pain management.
- Huge differences among countries and areas.
- Promissory future, but still a lot of work ahead of us!!

KEY MESSAGE: EARLY INTEGRATION NOT REFERRAL



THANKS

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