Recommendations of the European Association for Palliative Care (EAPC) For the Development of Undergraduate Curricula in Palliative Medicine At European Medical Schools

Report of the EAPC Steering Group on Medical Education and Training in Palliative Care
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Foreword to the first Edition

When the EAPC was formed in 1988 many European countries were still without Palliative Care services and few medical schools taught Palliative Care. In those that did, the time allocated to it was small, students were seldom examined in it and many academics saw little reason for change.

Times have changed. Many more people have now heard of Palliative Care and come to expect it to be available and accessible when they, or a loved one, need it. Several governments are acknowledging the need for nationwide Palliative Care services, some even generously funding it. More importantly the relevance of such care for people with non-malignant conditions is being acknowledged, as are the benefits of such care earlier in the patients’ care.

Perhaps the biggest change has been the recognition that nearly 50% of deaths in Europe follow ‘chronic’ conditions, the patient becoming increasingly frail and dependent with a spectrum of suffering extending over months or years before they die. Every one of those patients needs and will benefit from Palliative Care. Their number is vast and will inexorably increase with ageing populations and medical advances capable of keeping people alive much longer than anyone expected but often with a questionable quality of life. At the same time it has come to be recognised that 90% of these people will (and should) remain under the care of their general practitioners / family doctors. Only if they have suffering of such severity, complexity or rarity that their doctors cannot be expected to look after them will specialist expertise be needed.

It follows that every undergraduate medical student will need to learn about Palliative Care - its challenges, its complexity and its immense professional and personal rewards. He / she will discover that whilst its principles are those of all good clinical care being skilled in it is not an inborn gift. It requires knowledge, compassion, sensitivity and humility. Those who teach it must have those talents in abundance but, in addition, must know how to use appropriately every teaching / tutorial technique. Preferably they should have had formal teacher - training at undergraduate or postgraduate level. Such teachers are now to be found in Palliative Care departments in several universities and it is no surprise that undergraduates are rating their teaching amongst the best in the university and the subject itself amongst the most exciting.

The experts who have produced these recommendations know how easy it is to regard Palliative Care as little more than an exercise in clinical pharmacology,
particularly in opioid prescribing and rotation. They know how central to that care is the understanding of the psychosocial and spiritual aspects of the patient’s life (so often neglected both in undergraduate years and subsequent specialist training) and the suffering and needs of caring relatives. All have a place in their proposals.

A distinguished professor of surgery, about to retire and looking back on all he had seen and learned in the previous 45 years, recently said to me that the greatest advance he had seen, the most important thing he had had to learn, was Palliative Care. “We must ensure that never again will young men and women enter our noble profession unable to care for those they cannot cure, not knowing how to listen, not being ready to learn from nurses, and not sensitive to their patients’ greatest needs”.

He, like me, would approve of these EAPC recommendations and commend them to every medical school in Europe and beyond.

Derek Doyle
Chairman of the first EAPC Education Committee
Foreword to the present Edition

The EAPC Board considers the integration of education and training in palliative medicine for all physicians as of the utmost importance. Not only is it important for the development of palliative care itself, it is crucial for the patients and families for whom we care. Accordingly, as part of the work of the Steering Group on Medical Education and Training, a number of Taskforces have engaged surveys on the state of the art of education in palliative medicine for physicians across Europe.

Palliative medicine in Europe has grown up, and already has, or is on its way to become, a specialty of its own. To assist in this development of the practice of palliative medicine, the EAPC published the “Curriculum in Palliative Care for Undergraduate Medical Education” in 2007, and in 2009 the “Recommendations for the development of postgraduate curricula” leading to certification in Palliative Medicine. Recently, a Taskforce of the Steering Group for Medical Education and Training began the process to revise and update the published recommendations. What is presented here is the revised version of the Recommendations of the European Association for Palliative Care (EAPC) for the Development of Undergraduate Curricula in Palliative Medicine at European Medical Schools, as per 2013. With the undergraduate recommendations complete, the Taskforce will now turn its attention to Recommendations for Postgraduate Education. In developing the new recommendations, the Taskforce have consulted far and wide to ensure the document is accessible, flexible and pertinent to all in the European community. In addition, the members of the taskforce themselves bring great experience of the challenges and successes in pushing the agenda of developing appropriate training in palliative medicine within existing medical curricula, and the group members are to be commended on their efforts.

As such, I believe this new publication could be extremely useful in medical schools across Europe to facilitate the development of Palliative Medicine, particularly in these countries were undergraduate education in Palliative Medicine is still in the early stages of development.

Franco De Conno
Honorary Director of the EAPC
A) Introduction

EAPC background

From its initial conception the European Association for Palliative Care (EAPC) has regarded the education and training of health care professionals as of the highest importance for the promotion and expansion of Palliative Care in Europe. In 1992 an EAPC educational committee chaired by Derek Doyle proposed specific recommendations for training in Palliative Care. Later (2000) and in recognition of ways of working collaboratively in a Palliative Care context, a multi-professional expert group revisited these recommendations for education in Palliative Care.

In order to meet the interests of specific professional curricula (medical, nursing, etc.) the project separated after an initial phase of common goal setting. In 2004 a set of recommendations for palliative nurse education were proposed (EAPC 2004). The recommendations for the development of a curriculum in Palliative Medicine is the result of an equivalent project for training in the medical field, both for the basic (undergraduate) and for the specialist (postgraduate) level in accordance with the 2003 Council of Europe recommendations on the organisation of Palliative Care (COE 144/153). The recommendations are based upon existing medical curricula formulated by the authors with a view to a pan-European perspective.

In 2012, a consultation process was undertaken to review and update the existing curriculum guidance. The curriculum was circulated to experts in Palliative Care, as well as to national associations and other key persons in the field. The comments returned were discussed within the EAPC Steering Group on Medical Education, with changes and additions integrated where appropriate.

Undergraduate training in Palliative Medicine

The number of patients with cancer and other severe diseases continues to increase. The last decades have seen a rapid growth in the concern for the severely ill and dying patients. The developments in most countries have led to the establishment of Palliative Care services in a variety of settings with diverse staff composition and disparate quality standards.

Palliative Care education at medical schools has also increased in Europe within the last few years. However, undergraduate education in Palliative Care in Europe shows wide variation and there is no standardised European core curriculum. Efforts have already been made to standardise undergraduate Palliative Care education by creating curricula for instance in Australia, Canada and the United Kingdom.

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Furthermore, efforts also have to be made to introduce Palliative Care as a compulsory subject within the medical curricula at all European medical schools. The proposed recommendations for undergraduate education in Palliative Care may help medical schools within Europe to improve Palliative Care teaching. The contents are considered essential for all medical graduates. The change in attitude towards the care of severely ill and dying patients and their families is one of the major goals. Another important aspect is to address specific skills in interprofessional teamwork as well as in the basics of symptom management. The contents of this undergraduate curriculum refer to the basic principles fixed in the definition of Palliative Care by the World Health Organization (WHO) in 2002.

EAPC definition of Palliative Care

Palliative care is the active, total care of the patients whose disease is not responsive to curative treatment. Control of pain, of other symptoms, and of social, psychological and spiritual problems is paramount.

Palliative care is interdisciplinary in its approach and encompasses the patient, the family and the community in its scope. In a sense, palliative care is to offer the most basic concept of care – that of providing for the needs of the patient wherever he or she is cared for, either at home or in the hospital.

Palliative care affirms life and regards dying as a normal process; it neither hastens nor postpones death. It sets out to preserve the best possible quality of life until death.

- Palliative care provides relief from pain and other distressing symptoms
- Affirms life and regards dying as a normal process
- Intends neither to hasten or postpone death
- Integrates the psychological and spiritual aspects of patient care
- Offers a support system to help patients live as actively as possible until death
- Offers a support system to help the family cope during the patient's illness and in their own bereavement
- Uses a team approach to address the needs of patients and their families, including bereavement counselling, if indicated
- Will enhance quality of life, and may also positively influence the course of illness
- Is applicable early in the course of illness, in conjunction with other therapies that are intended to prolong life, such as chemotherapy or radiation therapy, and includes those investigations needed to better understand and manage distressing clinical complications.
General recommendations for the curriculum development in Palliative Care

These recommendations provide only a general framework for curricula in Palliative Care as defined by the different medical schools in Europe. The integration of Palliative Care into existing undergraduate curricula is a major challenge. Instruments like the Palliative Education Assessment Tool (PEAT) help to detect already existing features of Palliative Care education which often may be found “hidden” in the curricula of medical schools.

In order to meet needs of the different health care systems and the needs of the students some general principles of curriculum planning should be followed (see (E) General Principles of Curriculum Planning).

Specific components of the undergraduate curriculum in Palliative Care

It is the advice of the authors to avoid any overload of the undergraduate curriculum in Palliative Care and to plan any education with the intention and purpose to organize the distribution of the content in the sense of “vertical integration”.

The content of the curriculum should focus on seven domains of practice, in order to achieve a basic level of competencies in Palliative Care:

- Palliative Care, Palliative Medicine (definitions, etc.)
- Pain
- Neuro-psychological symptoms (agitation, confusion etc.)
- Management of other symptoms (dyspnoea, nausea, etc.)
- Ethics and law
- Patient, family, nonclinical caregivers perspectives
- Clinical communication skills

These seven domains are integrated within the six sections of our suggested syllabus

1. Basics of Palliative Care
2. Pain and symptom management
3. Psychosocial and spiritual aspects
4. Ethical and legal issues
5. Communication

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2 www.hawaii.edu/hivandaids/Development_of_a_Palliative_Education_Assessment_Tool_for_Medical_Student_Education.pdf
The presented curriculum does not specify how and in which detail Palliative Care should be taught. However, as with PEAT, consultation of the curriculum document may help to identify relevant contents within existing undergraduate medical curricula, but also teachers who need to be contacted in order to integrate a new Palliative Care curriculum (mainly contact to teachers from oncology, family medicine, geriatrics, pharmacology, psychology, psychiatry). This approach may also keep possible resistance low.

Teamwork and recommendations for multi-professional education

The ability to work in and lead a health care team is one of the characteristics of Palliative Care and needs special attention in the planning of the curriculum and the teaching methods adopted. It is considered to be mandatory that some members of the teaching faculty have a professional background different from the medical field (nursing, pastoral services, others).

B) Goals

A learning experience has to be promoted which permits students to develop the attitude, knowledge and skills necessary to participate in effective and compassionate Palliative Care.

Goals are:

- to show that medical treatment is far beyond diagnostic investigations and healing, the patient is meant to be considered, cared for and treated holistically
- to show how to relieve symptoms (pain and others) by pharmacological and non-pharmacological means
- to show that Palliative Care of patients and their relatives is a process that does not only include crisis intervention but also includes anticipatory treatment and attention
- to show that care and treatment have to be adopted to meet the individual needs, wishes and values of individual patients and their relatives
- to show that the quality of end of life care for patients will only succeed if the attending physicians are able to reflect upon their own attitude towards disease, dying, death and mourning
- to show that the quality of medical treatment cannot only be improved by enlarging knowledge but also by the competence of team-working, communicating and the willingness to discuss ethical issues
C) Educational strategies

Educational technique and process may vary in each medical school but it is recommended that the following aspects should be noted:

- Experiential learning (including contact with inpatient units, hospital consultative service or community settings, including patient and family encounters) should be taken into account predominantly. Debriefing should be considered a priority.
- Active rather than passive techniques should be applied (problem based learning, discussion, role play).
- Multi-professional learning should be encouraged to foster cooperation.
- Horizontal integration: Specialists in Palliative Care should be aware of areas of Palliative Care taught by colleagues in other disciplines.
- Repeated occasions for self-reflection and group discussions of difficult situations, including family issues, team problems and grief, should be arranged.
- Ethical and psychosocial considerations should be integrated into all aspects of teaching.

A total of 40 hours, allocated in different years of undergraduate medical education, is recommended for achieving the goals of this curriculum. Basics should be taught as early as possible, clinical aspects later on in the medical education. Horizontal and vertical integration must be transparent for the student. Additionally a clinical experience in a Palliative Care setting is recommended.

Assessment methods should consider tools to evaluate knowledge, for instance by multiple choice questions, short questions or case studies, as well as tools to measure attitudes and skills. It is crucial to introduce Palliative Care into final examinations – in the absence of such it is unlikely that all students will attend to the core issues of the curriculum. A systematic review of undergraduate training in palliative care found a lack of consistency in what undergraduates are taught about palliative care. Teaching tends to be fragmented, ad hoc and lacks co-ordination. There are difficulties in recruiting appropriate teachers, which is linked with a lack of qualified academic leaders in palliative care within University Medical Schools. Further, Palliative Care is rarely formally assessed and this is essential if it is to have the same academic credibility and relevance as other disciplines. Teaching focused

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more on the acquisition of knowledge and skills rather than attitudes and apart from a few notable exceptions was almost universally cancer focused.

(D) Syllabus

A suggestion for the split for the percentages of the topics within the syllabus is presented below:

1. Basics of Palliative Care 5%
2. Pain and symptom management 50%
3. Psychosocial and spiritual aspects 20%
4. Ethical and legal issues 5%
5. Communication 15%
6. Teamwork and self-reflection 5%

The proposed curriculum is based upon the minimal knowledge and skills which a medical student should obtain during his/her undergraduate education. The content list below must be translated into learning objectives and the educational strategy (learning method) must be defined (see (E) General Principles of Curriculum Planning).

The teaching staff for the items below can and will vary, including faculty from other different professional background than medicine.

1. Basics of Palliative Care: 5%

<table>
<thead>
<tr>
<th>Knowledge</th>
<th>Awareness of</th>
</tr>
</thead>
<tbody>
<tr>
<td>• International development of the idea of hospice and Palliative Care</td>
<td></td>
</tr>
<tr>
<td>• Definition of Palliative Care</td>
<td>the complexity of the end-of-life</td>
</tr>
<tr>
<td></td>
<td>the physician’s task in end of life care</td>
</tr>
<tr>
<td></td>
<td>the multi-professional and interdisciplinary approach of Palliative Care</td>
</tr>
<tr>
<td></td>
<td>the necessity of an early integration of Palliative Care in disease progression</td>
</tr>
<tr>
<td>• Forms of organisation:</td>
<td>the necessity of different forms of organisation</td>
</tr>
<tr>
<td>➢ outpatient</td>
<td>the necessity of communication between services</td>
</tr>
<tr>
<td>➢ inpatient</td>
<td></td>
</tr>
<tr>
<td>➢ consulting</td>
<td></td>
</tr>
</tbody>
</table>
### 2. Pain and Symptom Management: 50%

#### a) Basic principles of symptom management

<table>
<thead>
<tr>
<th>Content</th>
<th>Awareness of</th>
</tr>
</thead>
<tbody>
<tr>
<td>• Curative therapy</td>
<td>The chance of an increase of quality of life by offering Palliative Care early</td>
</tr>
<tr>
<td>• Palliative therapy</td>
<td></td>
</tr>
<tr>
<td>• Palliative medicine</td>
<td></td>
</tr>
<tr>
<td>• Interdisciplinary options</td>
<td>The necessity of interdisciplinary and multi-professional treatment</td>
</tr>
<tr>
<td>➢ surgery</td>
<td>Balancing diagnostics and treatment with the stage of disease</td>
</tr>
<tr>
<td>➢ radiotherapy</td>
<td></td>
</tr>
<tr>
<td>➢ pharmacological</td>
<td></td>
</tr>
<tr>
<td>➢ non-pharmacological</td>
<td></td>
</tr>
<tr>
<td>• Palliative sedation-</td>
<td>Palliative sedation for intractable suffering</td>
</tr>
<tr>
<td>➢ indications</td>
<td>The conceptual difference between palliative sedation and euthanasia</td>
</tr>
<tr>
<td>➢ procedure</td>
<td></td>
</tr>
<tr>
<td>➢ care of the relatives</td>
<td></td>
</tr>
<tr>
<td>• Planning and evaluation of treatment</td>
<td>The importance of the individual and prospective treatment and care</td>
</tr>
<tr>
<td>• Symptom assessment (goals and tools)</td>
<td>The importance to define goals</td>
</tr>
<tr>
<td>• Continuous and on-demand medication</td>
<td></td>
</tr>
<tr>
<td>• Prevention and rehabilitation</td>
<td></td>
</tr>
<tr>
<td>• Documentation</td>
<td></td>
</tr>
</tbody>
</table>

#### b) Pain management:

<table>
<thead>
<tr>
<th>Knowledge</th>
<th>Comprehension of</th>
</tr>
</thead>
<tbody>
<tr>
<td>• Definition and concepts of pain</td>
<td>The multidimensional approach of pain management</td>
</tr>
<tr>
<td>• Anatomy, pathophysiology</td>
<td>The complexity of pain management in end of life care</td>
</tr>
<tr>
<td>• Mechanisms of nociceptive pain (bone pain, soft tissue pain, visceral pain)</td>
<td>The fact that there is more to pain relief than drugs.</td>
</tr>
<tr>
<td>• Mechanisms of neuropathic pain</td>
<td></td>
</tr>
<tr>
<td>• Recognition of chronic pain features</td>
<td></td>
</tr>
</tbody>
</table>
### The concept of "total pain"

#### Principles of pharmacological treatment
- Importance of achieving 'steady state'
- Using the simplest available route of administration
- Role of titration
- Necessity to prescribe rescue medication
- The role of equianalgesic doses
- The role of opioid rotation

#### Pharmacokinetics and -dynamics of opioids, non-opioids & adjuvant analgesics

#### Routes of drug administration and their indications, alternative routes when oral are not possible.

#### Further non-pharmacological and pharmacological options of pain management:
- (Oncological) Interventions (chemotherapy, radiotherapy)
- Interventional procedures (anaesthetic or neurosurgical)
- Nursing interventions
- Psychotherapy and counselling
- Social intervention
- Physiotherapy
- Complementary therapy

#### Organisational and legal problems:
- Special prescription forms
- Driving ability
- Travelling

### (c) Symptom management:

<table>
<thead>
<tr>
<th>Knowledge</th>
<th>Comprehension for</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gastrointestinal symptoms</td>
<td>The physical, psychological, social and spiritual aspects of symptom management in Palliative Care</td>
</tr>
<tr>
<td>Constipation, diarrhoea</td>
<td></td>
</tr>
<tr>
<td>Anatomy and physiology of normal defecation and bowel continence.</td>
<td></td>
</tr>
<tr>
<td>Mechanisms of constipation in end of life care (drugs, particularly opioids, altered diet)</td>
<td></td>
</tr>
<tr>
<td>Weakness</td>
<td></td>
</tr>
</tbody>
</table>

Rumours and untruths. Destroy the myth that opioid analgesics are addictive, and that if initiated too early there will be "nothing left for the end".
- Ileus

- Nausea and vomiting
  - Pathophysiology of nausea and vomiting: where are the sites, receptors?
  - Pharmacology of anti-emetics, particular sites of drug action
  - The role of the route of drug administration

- Management of bowel obstruction

- Pulmonary symptoms
  - Dyspnoea
    - Pathophysiology of respiratory symptoms
    - Relevant pharmacology (opioids, anxiolytics, steroids)
    - Principles of oxygen therapy
    - How to deal with "death rattle"
  - Cough

- Neuropsychiatric symptoms
  - Delirium, confusional states
  - Insomnia
  - Depression and other mood disorders
  - Anxiety and fear
  - Hallucinations

- Anorexia, cachexia, fatigue
  - Loss of appetite
  - Fatigue
  - Weakness, lethargy

- Thirst, dry mouth
  - Sore mouth
  - Swallowing problems

- Dermatologic symptoms
  - Wound breakdown
  - Lymphoedema
  - Itching
- Care of the dying patient
  - Emergencies in Palliative Care
  - Hypercalcemia
  - Spinal cord compression

### 3. Psychosocial and spiritual Aspects: 20%

<table>
<thead>
<tr>
<th>Knowledge</th>
<th>Comprehension for</th>
</tr>
</thead>
<tbody>
<tr>
<td>Psychological reactions to chronic illness, grief and loss</td>
<td>The patient’s autonomy</td>
</tr>
<tr>
<td>Impact on patient and family of loss of independence, role, appearance, sexuality and perceived self-worth</td>
<td>The meaning of truth</td>
</tr>
<tr>
<td>Family dynamics</td>
<td>The patient’s individuality</td>
</tr>
<tr>
<td>Ethnic, social and religious differences</td>
<td>The patient’s vulnerable self-respect</td>
</tr>
<tr>
<td>How to help patients and families to deal with practical, financial and legal issues where appropriate, in particular, to arrange for social work and legal briefing to assist with will making or revision and compensation claims, which sometimes arise as matters of urgency close to the end of life</td>
<td>The meaning of vitality and sexuality</td>
</tr>
<tr>
<td>Facilitation of work leave and travel arrangements for relatives and friends to come to visit a dying person from within the country and overseas</td>
<td>Cherishing the patient’s and relatives’ feelings</td>
</tr>
<tr>
<td>coping strategies</td>
<td>Cherishing the patient’s and relatives’ needs</td>
</tr>
<tr>
<td></td>
<td>The complexity of the patient’s social circumstances</td>
</tr>
<tr>
<td></td>
<td>The important role of the family in terms of the patient’s quality of life</td>
</tr>
<tr>
<td></td>
<td>The specific needs of children</td>
</tr>
<tr>
<td></td>
<td>The difficulties you may encounter when dealing with severe illness and imminent death</td>
</tr>
<tr>
<td></td>
<td>The importance and meaning of quality of life in Palliative Care</td>
</tr>
<tr>
<td>Grief and bereavement as a process of each concerned person</td>
<td>Identification of helpful and not helpful strategies when working with the patient’s and relatives’ mourning, including children.</td>
</tr>
<tr>
<td>Anticipatory mourning</td>
<td>at the beginning of the disease</td>
</tr>
<tr>
<td>Risk factors for difficult mourning</td>
<td>during disease</td>
</tr>
<tr>
<td></td>
<td>when patient is dying</td>
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<tr>
<td></td>
<td>after patient’s death</td>
</tr>
<tr>
<td>Spirituality</td>
<td>The differences between spirituality and religion</td>
</tr>
<tr>
<td></td>
<td>One’s own spirituality</td>
</tr>
</tbody>
</table>
4. Ethical and legal Issues: 5%

<table>
<thead>
<tr>
<th>Knowledge</th>
</tr>
</thead>
<tbody>
<tr>
<td>Discussion of decision-making at the end of life, particularly the abatement, withdrawal or withholding of a treatment</td>
</tr>
<tr>
<td>The proper ways of negotiating and placing ‘Do-not-attempt cardio-pulmonary resuscitation orders (DNACPR or DNR)</td>
</tr>
<tr>
<td>Exploration of proxy decision-making, advance directives and advanced care planning</td>
</tr>
<tr>
<td>Distinction between accepted Palliative Care practice and euthanasia</td>
</tr>
<tr>
<td>Ethical and legal differentiation in the national and international context</td>
</tr>
<tr>
<td>euthanasia</td>
</tr>
<tr>
<td>physician assisted suicide</td>
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</tbody>
</table>

<table>
<thead>
<tr>
<th>Comprehension for</th>
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</thead>
<tbody>
<tr>
<td>Ethical aspects in medical decision-making</td>
</tr>
<tr>
<td>The reflection of one’s own ethical attitude</td>
</tr>
<tr>
<td>The reflection of one’s own attitude towards death and dying</td>
</tr>
<tr>
<td>The reflection about the physician’s role in treatment of end-of-life patients</td>
</tr>
</tbody>
</table>

5. Communication: 15%

<table>
<thead>
<tr>
<th>Knowledge</th>
</tr>
</thead>
<tbody>
<tr>
<td>Models of communication</td>
</tr>
<tr>
<td>Differentiation:</td>
</tr>
<tr>
<td>verbal vs. non-verbal communication</td>
</tr>
<tr>
<td>Special situations of communication</td>
</tr>
<tr>
<td>patient’s information, prognosis</td>
</tr>
<tr>
<td>decision-making</td>
</tr>
<tr>
<td>conflict and conflict resolution</td>
</tr>
<tr>
<td>talking with relatives</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Comprehension for</th>
</tr>
</thead>
<tbody>
<tr>
<td>The perception of the patient’s attitude towards their disease</td>
</tr>
<tr>
<td>One’s own shortcomings and strong points in perception, communication and advance care planning</td>
</tr>
</tbody>
</table>
### 6. Teamwork and Self-reflection: 5%

<table>
<thead>
<tr>
<th>Knowledge</th>
<th>Comprehension for</th>
</tr>
</thead>
</table>
| - How to work in a team | - The necessity of teamwork  
- The danger of role-conflicts  
- The process of decision-making  
- The possibility of debriefing each other in a team |
| - Networking  
  ➢ support systems  
  ➢ partners | - The importance of delegation |
| - "Burn-out" -avoidance and -prophylaxis | - One's own way how to manage burdens  
- One's own way how to manage personnel concern  
- The chance of debriefing oneself by supervision |
(E) General Principles of Curriculum Planning

The educational programme in Palliative Care should be organised in each country taking into consideration the six-step approach for curriculum development for medical education formulated by Kern et al. (1998) and outlined below:

**Step 1: Identification of general needs and problems**

In Palliative Care this might include statistical information of the number of severely ill patients and causes, preference and actual place of death, deficiencies in hospital care and home care, current ethical discussion in public, health insurance issues and so on.

This kind of analysis should enable students to identify key differences between the current and an ideal approach.

**Step 2: Identification of specific needs of different target groups**

**Content**

In Palliative Care this might include orientation towards practical needs of physicians regarding Palliative Care in different settings (hospital, home care, hospice, general practice, other medical specialists in private practice etc.) Specific needs should be defined by an analysis of deficiencies.

There should be a clear distinction between the undergraduate and postgraduate level. Palliative Care is in danger of accumulating an enormous number of topics within one course or curriculum. We recommend to focus on the vertical integration of learning content relative to time in continuous medical education (exposure to concrete clinical practice will lead to other needs in postgraduate training). This approach might prevent the creation of overloaded Palliative Care curricula in the undergraduate level.

A detailed analysis of the characteristics of the learners' environment regarding barriers, enabling and reinforcing factors, will help to implement any course or curriculum more easily.

**Methods**

Preferences and experiences regarding different learning strategies might differ enormously not only among the targeted learners but also to different cultures and countries. The same is true for the resources available to learners (e.g. computers, audiovisual equipment, role models, simulated patients, teachers etc.).
Step 3: Setting of goals and objectives

Goals are of a general character and define the ideas what should be generally achieved by a course or curriculum.

Objectives are defined specifically for the measurement of outcomes taking into consideration three major fields of achievement:

- **knowledge** (cognitive)
- **skills** (psychomotor)
- **attitude** (affective)

The more precisely objectives are formulated the easier evaluation (see step 6) becomes. The formulation of a learning objective needs the definition of

- Who will perform? (e.g. "the student")
- Which level of activity? (e.g. "should be able to rank")
- What level of achievement? (e.g. "3 most frequent symptoms...")
- When should objective be reached? (e.g. "at the end of the course")

Step 4: Educational strategies

The question is what content may be presented and learnt best. Content and method must be congruent. For instance, to improve decision making skills a small group discussion (problem based learning) might be more effective than having a faculty member analyse the case for the learners. Content should include all three areas of objectives: cognitive ("**knowledge**"), psychomotor ("**skills**") and affective objectives ("**attitude**").

The use of multiple educational methods during a course or program does not only provide a more vivid atmosphere and enhanced participation but also takes into account the different learning styles of the learners. Some learn best while reading an article, others prefer role play or creating a concept for their learning (see Learning Styles Model by Kolb).

Methods for achieving cognitive objectives (Knowledge)

- Problem based learning (PBL)
- Small group work
- Lectures
- Role plays

Methods for achieving psychomotor objectives (Skills)

- Supervised clinical experience
- Simulations (e.g. simulated patients, role plays)
• Audio or visual review of skills

Methods for achieving affective objectives (Attitude)

• Exposure (experiential learning) followed by discussion
• Role models
• Role plays
• Individual and group supervision: promote openness, introspection and reflection.

These methods rely on trust, continuity and trained teachers in order to facilitate self-reflection and feedback on a personal level. In Palliative Care these types of methods need to be integrated into each course or curriculum since self-reflection is of major importance for the field.

Specific to Palliative Care are methods to promote teamwork. Teamwork can be used as a collaborative learning experience (any task or problem experienced and resolved in a group), team-teaching (teaching by two, three, etc.) and team-exercise (e.g. to solve a case together). Several models for enhancing teamwork (knowledge, skills and attitude) exist from management courses in industry and can be transferred to the field of Palliative Care.

Step 5: How to implement the course or curriculum

The following checklist facilitates the successful establishment of a curriculum in Palliative Care:

• Identify resources (personnel, time, faculties, funding)
• The program leader must be qualified to supervise and educate trainees in Palliative Care. Thus, the leader should have appropriate experience in Palliative Care and teaching
• Identify multi-professional teaching staff to meet training needs
• Develop administrative mechanisms to support the curriculum for
  o distribution of responsibilities
  o continuous communication and adaptation
  o permanent evaluation
• Find partnerships with clinical Palliative Care services, other specialities, other health care professionals.

Step 6: Evaluation and Feedback

Evaluation and feedback close the loop in the curriculum development cycle. Two questions are asked:
1. How did the participants perform? (Assessment of the individual)

Usually two different assessment strategies are used in order to evaluate the achievement of the learning objectives formulated in Step 3:

- **Formative** assessment: the change in achievement or performance over a time period, mostly monitored by a tutor or supervisor, using self-assessment tools. The exam itself is part of the educational process.

- **Summative** assessment: the level of achievement. You must pass this exam to progress. Several assessment methods are commonly used, including written or computer-interactive tests, oral examination, questionnaires (MCQ or open questions), direct observation

2. How did the curriculum perform? (Assessment of the curriculum)

In Palliative Care, as a relatively young discipline, permanent attention should be drawn to the increasing basic knowledge of the targeted groups. Therefore Step 6, 1 and 2 need to be administered carefully and repetitively. The assessment of the curriculum usually focuses on the content level as well as on the methods.

To test the impact of a curriculum on the population level (consumers' health) might be of special interest and can be administered best in clearly described environments (e.g. nursing home, community level).

The results of step 6 should be disseminated to regional, national and international groups in Palliative Care and across a multi-professional spectrum.
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