



European Framework for the certification of CCCNs in the course of iPAAC

WP 10

Author(s): Lead author: Simone Wesselmann

Co-authors: Ellen Griesshammer

Version: 1.0

Date: 27. 09. 2019





Contents

Pr	Prologue	3
Di	Division of Authority	3
Αι	Auditors	5
Αι	Audit	6
5.1	Audit Plan	6
5.2	On-site Audit	6
5.3	Audit Report	7
C	Certificate	7
E	xample	8
E	Excerpt from the Auditreport	11
	5.1 5.2 5.8 6 6 6 7 7 8	Prologue

This report arises from the Innovative Partnership for Action Against Cancer Joint Action, which has received funding from the European Union through the Consumers, Health, Agriculture and Food Executive Agency of the European Commission, in the framework of the Health Programme 2014-2020. The European Commission is not responsible for the content of this report. The sole responsibility for the report lies with the authors, and the Consumers, Health, Agriculture and Food Executive Agency is not responsible for any use that may be made of the information contained herein. The authors are not responsible for any further and future use of the report by third parties and third-party translations.





1 Prologue

The goal of Work Package 10 is to further develop practical instruments ensuring a standardised integrated and comprehensive oncological care in all European Member States that is tumour-specific and delivers all-encompassing high-quality care to all patients. Based on the results of the previous Joint Action CanCon WP 10 has defined two Sets of Standards (SoS) for Comprehensive Cancer Care Networks (CCCNs). One Set is tumour-specific for colorectal and pancreatic cancer and includes guideline-based requirements, structural requirements, e.g., staffing and technical infrastructure and key performance figures. The second SoS outlines non-tumour-specific, generic standards for the basic organization of oncological care within a CCCN.

With the goal of setting up CCCNs the contents of both sets will be implemented in two pilot centres (Lower Silesian Oncology Center, Wroclaw, Poland and Charité, Berlin, Germany). Building on these two Sets of Standards for Comprehensive Cancer Care Networks (see chapter 3). The purpose of this document is to define the conditions necessary to verify the successful and sustainable implementation of the defined standards.

The implementation of CCCNs is an ongoing process that requires a continuous reflection and assessment of the treatment outcomes and of the underlying processes and structures. The prerequisites, which are defined in this framework are therefore an important element in order to monitor and evaluate the status quo of implementation and with this the overall quality of oncological care within CCCNs.

The described process and outlined standards in this document are only to be used in the context of the iPAAC pilot centres certification. Any further use requires that the described criteria remain unchanged and should only take place in close co-ordination with the WP 10 task 5 leader.

2 Division of Authority

The trustworthiness and value of a certification system is reflected by the quality of the stated requirements and moreover by the underlying principles of the monitoring and evaluation processes that form the basis of the framework.

It must be ensured that each section of the framework works independent from another and that potential conflicts of interest are avoided.

The following sections of the framework should therefore be separated from each other:

1. Definition of the Sets of Standards

Working group iPAAC WP 10, task 5 defined the requirements for Comprehensive Cancer Care Networks (CCCNs) and Colorectal and Pancreatic Cancer Care Networks. The requirements consist of a set of standards including key performance indicators.

2. Review/audit of the implementation of the set of standards





Oncology experts who are not affiliated with the pilot centres will review the implementation of the Standards in the two pilot CCCNs.

3. Awarding of the certificate

On the basis of the results of the audit the certificate is awarded on behalf of iPAAC WP 10. The awarding of the certificate will take place after agreement by a certificate awarding committee consisting of at least five oncology experts.

4. Evaluation of the overall process

Since this is a pilot, an independent evaluation group assesses the process by conducting structured interviews with the pilot sites, the audit teams, and the awarding committee. The evaluators produce a Report about how effective the process in the pilots was and whether any modifications to the process are needed going forward.

3 Documents needed for certification

The following documents must be sent to the WP 10 leader (Deutsche Krebsgesellschaft e.V.) in advance of the audit:

- a) Master data sheet with overview of all partners of the CCCN (Link)
- b) Standard for Comprehensive Cancer Care Networks (Link)
- c) Standard for Colorectal and Pancreatic Cancer Networks (Link)
 - a. including Self-Assessment with scoring system

The Master data sheet (a) contains the information (discipline, name, department, address, mail, etc.) of all partners of the CCCN and the Colorectal and Pancreatic Cancer Networks.

In the Sets of Standards (b) and (c), the pilot centres describe their individual compliance with the Standards. The SoS includes a Self-Assessment with a scoring system. The completed Set of Standards including the key performance indicators are the basis for the on-site audit.

The scoring system for the Self-Assessment lists one score for each chapter of the Set of Standards. The score is an indicator for the implementation degree of each chapter in the Set of Standards.

The completed Set of Standards will be assessed by the auditors prior to the audit. In the case of implausible or incomplete information, the CCCN will be consulted and may need to provide further information or explanations in preparation for the on-site audit. The auditors will provide a list with the names of documents which should be translated in English for the onsite audit.





4 Auditors

The Set of Standards summarises the requirements that, when implemented in CCCN's, should provide patients with comprehensive, high-quality care at all stages and in all areas of an oncological disease. The requirements are tumour specific and generic and pursue the goal of a continuous improvement process within the CCCN.

The auditors have the task of checking the degree of implementation of the Standards and identifying the need for quality improvements. In order to address these improvement opportunities and increase the quality of care, appropriate actions with the CCCNs have to be discussed and agreed upon by the time of the finalisation of the Report.

In order to fulfil the outlined tasks, the auditors must be fully knowledgeable of the processes, procedures and day-to-day clinical practice in oncological care structures. In addition, the auditor must be familiar with the current diagnostic and therapeutic concepts of the oncological disease which is being audited and be able to assess the effects and side effects of all treatment steps of an oncological therapy.

Only when the auditor can fulfil the above described prerequisites, an assessment of the implementation of the requirements and the detection of quality improvements required with simultaneous development of meaningful quality improvement measures can be achieved.

Further it is highly desirable that at least one person of the audit team is fluent in the language of the audited CCCN to ensure that all members of the reviewed network can benefit from the audit and the discussions.

Based on these guiding principles, at least two auditors must meet the following requirement: Medical doctor with a board qualified specialist training; preferably in the relevant discipline of the tumour entity being audited.

It is suggested that the size of overall audit team for each pilot CCCN be based on the size of the CCCN and the number of sites being visited, but be at least 4 persons. These should include a senior oncology nurse, and an oncologist who has experience of conducting clinical trials. All members of the audit team should meet the following requirements:





- at least 7 years of experience in working in the field of oncology (the experience cannot date back more than 3 years)
- Successful participation in the CCCN-Audit-Training Course* (*to be developed)
- Cannot be a member of the iPAAC WP 10
- should declare the absent of potential conflict of interest to the centres being audited
- must sign a non-disclosure statement

Guests and the extent of their involvement, should be agreed by the audited CCCN. They can participate in the on-site audit but have no authority to independently assess or verify the implementation of the Set of Standards.

5 Audit

5.1 Audit Plan

An audit plan will be prepared for the audit.

The contents of the audit plan are based on the chapters of the Set of Standards.

Basically, all specialist disciplines and cooperation partners of the CCCN are visited on site. The responsible clinical personnel of the respective area will present their work and will be available for the exchange/discussion with the audit team.

The overall structure of the CCCN including the relevant key performance figures and quality of results are presented in the beginning with an introductory presentation by the Directors of the CCCN.

The audit plan is prepared by the WP 10 leader in consultation with the WP 10 task 5 members and in co-ordination with the pilot sites.

The duration of the audit is two full days on site.

The audit plan lists:

- Departments/Facilities to be visited on-site (e.g., operating room)
- Processes being discussed (e.g., preparation of chemotherapy protocols)
- People responsible for the individual areas / departments / processes within the CCCN
- Times and location
- List of documents that have to be available in English

5.2 On-site Audit

During the on-site audit, the CCCN substantiates the information provided in the SoS with the presentation of the accompanying documents (e.g.: SOPs, cooperation agreements, standard chemotherapy protocols, training calendars, etc.).





During the on-site audit, the auditors verify the compliance with the set of standards. They determine the degree of implementation of the CCCN and deduce an overall impression of the maturity stage of the CCCN. Areas with improvement potential will be identified and, together with the CCCN, improvement measures discussed and agreed upon. Through randomly selected patient records, the adherence of the guideline-based care of the patients and the implementation of the requirements laid out in the SoS will be verified and checked by the auditors.

5.3 Audit Report

The auditors have three options to evaluate the performance of the CCCN if the requirements are not fully implemented or if quality deficiencies are identified during the audit: (a) observations, (b) recommendations and/or (c) deviations. Through this, the CCCN receives a feedback and a reference point of the importance of the pointed-out short-comings which need to be addressed and remedied.

After the audit, the auditors prepare an Audit Report in which they either recommend or do not recommend that the certificate should be issued. In the report, the auditors write a summary of the actual situation for each chapter of the SoS and assess the chapters via the listed comment options. In addition, the auditors use the self-assessment tool to score each chapter/subchapters.

Comment options:

Observation(s)	Observations describe general impressions from the audit process that are neither recommendations nor deviations.
Recommendation(s)	Recommendations for the further development of the Centre. If the "must" formulation has been used, failure to comply with the recommendation may lead to a deviation being formulated in the next re-audit.
Deviation(s)	Deviations document non-compliance with the technical and medical requirements. The deviations must be remedied within 3 months (Exceptions must be explained) by the Network and this must be proven to the working group of WP 10 within the period specified in the deviation report.

6 Certificate

On the basis of the audit report, the scoring instrument and the documents provided by the CCCN (a-d), a committee (Certificate Awarding Committee) will issue the certificate on behalf of iPAAC, WP 10.

The certificate awarding committee consists of at least five members of Task 5, WP 10 including the WP 10 leader.





The certificate is valid for 5 years. If the improvement potential of the CCCN is very high the period of validity of the certificate can be reduced. In the years 1 and 3 after the initial audit, the CCCN must submit the annual results of the quality indicators and submit a written statement on the state of play in the areas where recommendations were made. The information will be evaluated by the audit team and the Certificate Awarding Committee will decide on the continuation of the certificate.

Continuation of the certificate beyond the defined term requires a re-audit. During the validity of the certificate, the CCCN is allowed and encouraged to use the certificate for public communication (e.g., website, letter heads, etc.).

In case there have been significant changes to the Set of Standards over the course of 5 years the auditors should receive an update/refresher training

7 Example

Assumption:

The aim of WP 10 is to improve oncological care. The structures and processes summarized in the set of standards should be implemented and with this good oncological care is made possible. There are different ways in which the individual standards of a chapter can be implemented in a CCCN. However, it is crucial that the overall process is implemented. For the example of Chapter 1.8, this means that nursing care is fully integrated in a CCCN with the right qualified professionals and sufficient human resources and the described tasks in the sense of the CCCN and the patients are fulfilled. However, it is not sufficient to show only the current state of the CCCN. Recommendations for further development must be listed to enable continuous quality improvement within the CCCN.

Excerpt of the Set of standards provided by the CCCN before the audit:

(see Chapter 3 Documents needed for certification)

1.8 Nursing care

Chapter	Requirements	Explanatory remarks of the CCCN
1.8.1		The CCCN describes here how it has implemented the standards described in the right column





1.8 Nursing care

	 At least 2 full-time active specialist oncology nurses must be employed on day duty in the CCCN to facilitate care coordination and provide specialized care. Specialist oncology nurses are identified by name. Active care by a specialist oncology nurse must be proven in the units in which patients receive inpatient oncology care. 	
	Pre-condition for the recognition as oncology nursing staff are	
	 Further training as oncology nursing staff according to the country specific regulations If there are no country specific regulations the CCCN must demonstrate how oncology nurses are educated (recommendation: European Oncology Nursing Society Educational Framework http://www.cancernurse.eu/education/cancernursingeducationframework.html) 	
1.8.2	Responsibilities / tasks	
	 Patient related tasks include: Specialist assessment of symptoms, side effects and stress/strain Conduct and evaluation of nursing measures Pain and symptom identification Identification of individual patient-based counselling needs. The specialist counselling needs are already to be defined in the nursing concept of the individual 	





1.8 Nursing care

	A specialised nurse should be	
	present in the consultation hours	
	where the diagnosis and further	
	diagnostic/treatment steps are	
	planned	
	Ongoing information and	
	counselling of patients (and their	
	family members) during the entire	
	course of the disease	
	 Conduct, coordination and 	
	documentation of structured	
	counselling sessions and	
	guidance of patients and family	
	members. Depending on the	
	concept this can also be done by	
	specialist nurses with many years'	
	experience and specialist	
	expertise.	
	Participation in the tumour board	
	is desirable.	
	Initiation of and participation in	
	multi-professional case	
	discussions/nursing visits; the aim	
	is to find a solution in complex	
	nursing situations.; Criteria for the	
	selection of patients are to be laid	
	down; per year and centre at least	
	12 case discussions/nursing visits	
	are to be documented	
Self	Yes – the chapter has been implemented as a wild as a larger and a second as a wild as a larger and a second as a la	Yes
assessment	implemented on a wide scale, and	
	the Deming cycle has been	
	completed twice.	Mostly
	Mostly – the chapter has been implemented in critical places, the	
	implemented in critical places, the	
	Deming cycle completed once.	Partially
	 Partially – the chapter has been only partly implemented, or only 	
	' ' ' ' ' ' ' ' ' ' ' ' ' ' ' ' ' ' '	
	recently introduced and not evaluated.	No
	 No – the chapter has not been implemented 	
	Not Applicable (rare).	Not Applicable
	140t Applicable (Tale).	110t Applicable





8 Excerpt from the Auditreport

(see Chapter 5c On-site-audit, comment options)

1.8 Nursin g care	Observation(s)		n describes here how the standards of the plemented in the CCCN
	Recommendatiom ns	agreed impr	describes here recommendations and the overnent measures for the further of the integration of nursing care within the
		-	implementation of a further training oviding more full-time employees
	Deviation(s)		am describes here deviations (= non- th the technical and medical requirements) remedied
	Overall assessment	Yes	The audit team states the current state of play with the overall assessment
	Yes – the chapter has been implemented on a wide scale, and the Deming cycle has	Mostly	
	been completed twice.	Partially	
	Mostly – the chapter has been		
	implemented in critical places, the Deming cycle completed once.	No	
	Partially – the chapter has been only partly implemented, or only recently introduced and not evaluated.	Not Applicable	
	No – the chapter has not been implemented		



