

Benchmarking Cancer Centers: From Care Pathways to Integrated Practice Units

Anke Wind, PhD^{a,b}; Francisco Rocha Gonçalves, PhD^c; Edit Marosi, MSc^d; Lucia da Pieve, MSc^e; Monica Groza, MSc^f; Marco Asioli, MSc^g; Marco Albini, MSc^h; and Wim van Harten, MD, PhD^{a,b,i}

Abstract

Background: Structuring cancer care into pathways can reduce variability in clinical practice and improve patient outcomes. International benchmarking can help centers with regard to development, implementation, and evaluation. A further step in the development of multidisciplinary care is to organize care in integrated practice units (IPUs), encompassing the whole pathway and relevant organizational aspects. However, research on this topic is limited. This article describes the development and results of a benchmark tool for cancer care pathways and explores IPU development in cancer centers. **Methods:** The benchmark tool was developed according to a 13-step benchmarking method and piloted in 7 European cancer centers. Centers provided data and site visits were performed to understand the context in which the cancer center operates and to clarify additional questions. Benchmark data were structured into pathway development and evaluation and assessed against key IPU features. **Results:** Benchmark results showed that most centers have formalized multidisciplinary pathways and that care teams differed in composition, and found almost 2-fold differences in mammography use efficiency. Suggestions for improvement included positioning pathways formally and structurally evaluating outcomes at a sufficiently high frequency. Based on the benchmark, 3 centers indicating that they had a breast cancer IPU were scored differently on implementation. Overall, we found that centers in Europe are in various stages of development of pathways and IPUs, ranging from an informal pathway structure to a full IPU-type of organization. **Conclusions:** A benchmark tool for care pathways was successfully developed and tested, and is available in an open format. Our tool allows for the assessment of pathway organization and can be used to assess the status of IPU development. Opportunities for improvement were identified regarding the organization of care pathways and the development toward IPUs. Three centers are in varying degrees of implementation and can be characterized as breast cancer IPUs. Organizing cancer care in an IPU could yield multiple performance improvements.

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Healthcare systems struggle with the rising cost of cancer care¹ and are increasingly under pressure to deliver high-quality services.² Cancer care is often provided across different settings and involves healthcare professionals from multiple disciplines,³ and therefore high-quality multidisciplinary care can be compromised by inadequate coordination.⁴ The implementation of care pathways has been shown to reduce variability in clinical practice and to improve

outcomes.⁵ “Clinical/Care pathways,” with varying nomenclature such as critical pathways, integrated care pathways, case management plans, and care maps, are used to systematically manage a patient-focused care program.⁶ Although clinician views on purpose, content, and implementation diverge,⁷ consensus exists on pathway characteristics: it should have strong multidisciplinary character aimed at improving quality and efficiency, and strong emphasis

^aDepartment of Psychosocial Research and Epidemiology, Netherlands Cancer Institute, Amsterdam, the Netherlands; ^bDepartment of Health Technology and Services Research, University of Twente, Enschede, the Netherlands; ^cInstituto Português de Oncologia do Porto, Porto, Portugal; ^dInternational Office, National Institute of Oncology, Budapest, Hungary; ^eClinical Risk Management and Accreditation Unit, Centro di Riferimento Oncologico (CRO), IRCCS, National Cancer Institute, Aviano (Pordenone), Italy; ^fThe Oncology Institute “Prof. Dr Ion Chiricuta” Cluj-Napoca, Cluj-Napoca, Romania; ^gScientific Directorate, Fondazione IRCCS Istituto Nazionale dei Tumori, Milan, Italy; ^hQuality Monitoring Office, Humanitas Clinical and Research Center, Milan Italy; and ⁱRijnstate Hospital, Arnhem, the Netherlands.

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Correspondence: Wim van Harten, MD, PhD, Department of Health Technology and Services Research, University of Twente, PO Box 217, 7500 AE, Enschede, the Netherlands. Email: w.v.harten@nki.nl

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on alignment issues throughout the care chain for a homogeneous patient group. Vanhaecht et al⁸ showed that improvement in care pathway concepts and methodology demand international knowledge-sharing, which can be facilitated by international benchmarking (eg, on specific topics such as information technology integration). According to Polite et al,⁹ oncology pathways offer many potential advantages; nevertheless, several issues must be addressed, such as who should control the development of pathways.

A recent development in healthcare is the transformation from volume-based to value-based care.¹⁰ In terms of value (ie, patient health benefits per healthcare dollar spent),¹¹ most current models lack (1) an ability to measure outcomes that matter to patients, (2) transparency around measured clinical and financial outcomes, and (3) care coordination across all providers in the care pathway. According to Porter and Teisberg,¹⁰ transformation to value-based healthcare requires a shift from silos organized by specialty to care organized around a patient's medical condition, including the whole pathway. They propose integrated practice units (IPUs), in which a team of clinical and nonclinical personnel provides the full care pathway.¹⁰ Wherever IPUs exist, consistent results are found, including faster treatment, better outcomes, and lower costs.¹ An example of IPU development in cancer care is the multidisciplinary care centers at MD Anderson Cancer Center.¹² However, research regarding this subject is limited.

The primary goal of the present study was to describe the development and outcomes of a benchmark

tool for oncology care pathways. Benchmarking is defined as the continual and collaborative measuring and comparing of results of key work processes with those of the best performers.¹³ Learning how to adapt these best practices can help achieve breakthrough process improvements and build healthier communities.¹³ The secondary goal was to explore the degree of IPU development in cancer centers based on the benchmark data.

Methods

Study Design

This international benchmarking study—part of the BENCH-CAN project,¹⁴ a European project aimed at benchmarking cancer care to contribute to improving the quality of interdisciplinary patient treatment—involved 7 European cancer centers (4 in South Europe and 3 in Central/East Europe). Participating cancer centers were members of the Organisation of European Cancer Institutes (OECI), and 5 were designated Comprehensive Cancer Centers by the OECI.¹⁵ The benchmark tool used to collect data was developed and executed according to the 13 steps developed by van Lent et al¹⁶ (Table 1). Steps 7 through 12 are further elaborated in the following section.

Indicator Development and Collection

Indicators (step 7) were derived from the literature and expert opinion. Experts included a representative from the Netherlands Comprehensive Cancer Organisation (IKNL), who was researching pathways in oncology,

Table 1. Thirteen-Step Benchmarking Method

Step	Action	Application in This Study
1	Determine what to benchmark	Organizational aspects of pathways and tumor services
2	Form a benchmarking team	BENCH-CAN consortium
3	Choose benchmarking partners	7 cancer centers in South and Central/East Europe
4	Define and verify the main characteristics of the partners	Mapping exercise of the external environment of the cancer centers and the influence this could have on the pathway and tumor service development
5	Identify stakeholders	Patients, clinicians, administrators, and researchers
6	Construct a framework to structure the indicators	This was not specifically done for this study; framework of the BENCH-CAN project was used ¹⁴
7	Develop relevant and comparable indicators	Based on literature and expert opinion (from cancer centers and an oncology care pathway expert from the Netherlands Comprehensive Cancer Organisation)
8	Stakeholders select indicators	Stakeholders from the BENCH-CAN project and other experts from cancer centers provided feedback on the indicators
9	Measure the set of performance indicators	Data collection phase was 3 months A team (consisting of the first author, other members of the BENCH-CAN consortium, and a member of European Cancer Patient Coalition) performed a site visit to each pilot center to verify the data, understand the context, and clarify any questions arising from the data
10	Analyze performance differences	Deductive qualitative content analysis ²¹ performed
11	Take action: present results in a report and provide recommendations	For each participating cancer center, a report was prepared containing the anonymized outcomes of the benchmark for all centers Improvement recommendations were sent in a separate document
12	Develop improvement plans	If in agreement, pilot sites developed improvement plans
13	Implement the improvement plans	Outside the scope of this study

Adapted from van Lent W, de Beer R, van Harten W. International benchmarking of specialty hospitals. A series of case studies on comprehensive cancer centres. *BMC Health Serv Res* 2010;10:253, with permission.

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and those from the cancer centers.^{17,18} Stakeholders of the BENCH-CAN project (eg, clinicians and [quality] managers) and experts from other cancer centers (OEI members, n=71) were then asked for feedback and a consensus was reached on the indicator set (step 8), containing 51 qualitative indicators and 193 quantitative indicators. Indicators assessed multiple topics concerning the organization of cancer care pathways, specifically those for breast and colorectal cancers, with a focus on organizational aspects of care pathways, not the clinical interpretation. Pilot sites appointed a team responsible for data collection covering multiple departments, including different stakeholder groups (ie, patients, clinicians, researchers, and management). Data were collected for the year 2012. After a quick data scan, a 1-day visit to each center was performed to verify the data, understand the context, and clarify questions arising from the data (step 9). Each visit consisted of semistructured interviews and a tour of the cancer center, with a specific focus on the breast unit (if available). The visits were also used to collect additional information and acquire feedback on the benchmark tool. The validity of the indicators was checked using feedback from pilot sites based on 3 criteria^{19,20}: (1) clear definition, (2) data availability and reliability, and (3) discriminatory features.

Analyses

After the completion of all site visits, data for each indicator were compared. A deductive form of qualitative content analysis²¹ was used to report on the collected data (step 10). This method, which contains 9 separate steps from the benchmarking method tool, are described in Table 2.

Data in our study were anonymized, and pathway description was structured based on the criteria by Kinsman et al.²² Data analysis focused mainly on the pathway for breast cancer because all centers could present sufficient data, which allowed for comparison. Indicators developed for the pathway tool were based on the IPU criteria developed by Harvard Business School,²³ such as organization of multidisciplinary teams (MDTs) and description of the steps taken in the development of the care pathway. To explore the degree of IPU development, every center was scored against these criteria (eg “organized around the patient medical condition or set of closely related conditions” and “co-located in dedicated facilities”)

Table 2. Deductive Qualitative Content Analysis Steps

Step	Action
1	Read through the (benchmark) data transcript, making notes in the margins
2	Assess the notes made and list the different types of information found
3	Read through this list and categorize each item
4	Repeat the first 3 stages again for each data transcript
5	Collect all of the categories or themes and examine each in detail and consider the fit and relevance
6	Categorize all data (all transcripts together) into minor and major categories/themes
7	Review all categories and ascertain whether some can be merged or some need to be subcategorized
8	Return to the original transcripts and ensure that all the information has been categorized
9	To ensure the validity, send a report containing all data to the pilot sites for verification

Adapted from Zhang Y, Wildemuth BM. Qualitative Analysis of Content. Available at: https://www.ischool.utexas.edu/~yanz/Content_analysis.pdf. Accessed September 15, 2016.

(available at: <https://www.isc.hbs.edu/health-care/vb-hcd/pages/integrated-practice-units.aspx>).

Results

Indicators

After data collection, definition clarity, data availability, data reliability, and discriminative value of the indicators were evaluated with the pilot centers. Based on this evaluation a total of 7 qualitative and 52 quantitative indicators were deemed irrelevant and removed, and 1 indicator regarding minimal volume of surgeries was added. This evaluation resulted in a final set of 45 qualitative indicators and 141 quantitative indicators that were considered suitable for wider use in benchmarking care pathways and exploring IPUs ([supplemental eAppendix 1, available with this article at JNCCN.org](#)).

Pathway Benchmark

An overview of the status of breast cancer pathways is provided in Table 3. Most cancer centers only started using official pathways recently; centers D and E had not implemented and formalized all pathways and were recommended to do so. Pathways are based on guidelines, either national or international, and have a clear “director” to guide development. All centers perform at least mammography, ultrasound, and physical examination before breast surgery, and all perform annual mammographies in the first 5 years of follow-up for all patients. However, center E indicated that they perform the follow-up for approximately 60% of their patients. Because mammography plays an important role in both

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Table 3. Overview of Breast Cancer Pathways

Cancer Center	Date of Pathway Introduction	Basis	Pathway Director	Diagnosis ^a and Follow-Up ^b	Other Pathways
A	2007	Pathways were developed using national and international guidelines and based on experience of professionals in cancer treatment	(Breast) Clinic coordinator	Diagnosis: 100% of patients Follow-up: 100% of patients Efficiency mammography (scans/device/year): 4,341	Endocrine tumors, skin/melanoma, pediatric oncology, onco-hematology, and nervous system, lung, soft tissue/bone, head and neck, breast, gynecologic, gastrointestinal, and urologic cancers
B	2011	Pathways are part of quality management system documentation Treatment guidelines are reviewed and updated by a commission at least once a year Guidelines are based on EUSOMA guidelines ³² and SIS accreditation ³³	Head of the breast surgery department	Diagnosis: 90%–95% of patients Follow-up: Unknown Efficiency mammography (scans/device/year): 7,617	Lung, breast, gynecologic, urologic, head and neck, and gastrointestinal cancers
C	2010	Pathways were developed by center leadership and members of the relevant MDT based on international and national guidelines, relevant legislation, and clinical practices	Director of the breast cancer pathway is the head of the breast cancer MDT Director of each pathway is the head of the relevant MDT	Diagnosis: 100% of patients Follow-up: 100% of patients Efficiency mammography (scans/device/year): 6,490	Lung, breast, gastrointestinal, and urologic cancers, and melanoma
D	2014	Institutional protocols and working procedures were developed based on international and national guidelines Periodically revised by MDTs and management Some pathways are not formalized, but they have existed for years because there have been MDTs for years	Director of the cancer care pathway is the chief of the department	Diagnosis: 100% of patients Follow-up: 100% of patients Efficiency mammography (scans/device/year): 6,427	Hematology, pediatric oncology, and digestive, urologic, lung, colorectal, breast, head and neck, and gynecologic cancers
E	2014	Clinical pathways were developed in accordance with the regional and national legislation, as well as clinical practice Currently, the pathways are partially implemented	There are clear directors within the development of pathways For breast cancer it is the head of the breast surgical oncology division	Diagnosis: 100% of patients Follow-up: \pm 60% receive follow-up at this center; of these, 100% receive mammography Efficiency mammography (scans/device/year): 10,444	Melanoma, lymphoma, sarcoma, and ovarian, gastric, head and neck, breast, and colon cancers
F	2005	Clinical pathways are based on international and national guidelines and conferences to continually stay up to date with the clinical literature and keep them current	Cancer care pathways are developed by the clinical and surgery departments with the supervision of the Healthcare Directorate and in close collaboration with the Medical Directorate	Diagnosis: 100% of patients Follow-up: 100% of patients Efficiency mammography (scans/device/year): 5,852	Biliary tumors, sarcomas, and thyroid/parathyroid, prostate, neuroendocrine, pancreatic, hepatic, head and neck, and breast cancers
G	2011	PDSA cycles are used for developing, implementing, and reviewing clinical pathways, which are based on national and international guidelines	Disease units (integrated practice units) and MDTs are responsible for defining and implementing evidence-based pathways and algorithms as a support for their activities The cancer center managing director and the clinical and scientific director are responsible for facilitating, overseeing, and approving clinical pathways	Diagnosis: 100% of patients Follow-up: 75% of patients Efficiency mammography (scans/device/year): 4,125	Chronic lymphatic leukemia, chronic myeloid leukemia, melanoma, mesothelioma, multiple myeloma, sarcoma, thymoma, onco-thrombosis, supportive treatments, female fertility and sexuality preservation, and urologic, cervical, colorectal, endometrial, liver, lung, ovarian, pancreatic, pituitary, prostate, and upper gastrointestinal cancers

Abbreviations: EUSOMA, European Society of Breast Cancer Specialists; MDT, multidisciplinary team; PDSA, plan-do-study-act; SIS, Senologic International Society.

^aPercent of patients undergoing presurgical mammography, ultrasound, and physical examination.

^bPercent of patients undergoing routine mammographic screening.

diagnosis and follow-up, efficient use of the machines is essential. The number of scans performed per device per year varied from 4,125 to 10,444.

Pathway Characteristics

An overview of breast cancer pathway characteristics is provided in Table 4. The pathways were developed for MDTs, with different roles for the various healthcare professionals. Center D was the least structured and was recommended to include discussion by an MDT in the

pathway for all patients. Centers E and F were recommended to evaluate their protocol for patient transition to other healthcare facilities, because this was lacking in the formal pathway. In most countries, maximum waiting and throughput times for the different steps in the pathway were set by the government.

Evaluation

Table 5 shows that evaluation methods vary from an informal evaluation by MDTs to an extensive internal

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Table 4. Breast Cancer Pathway Characteristics					
Cancer Center	Multidisciplinary	Guidelines or Evidence Base	Content of Pathway Description	Criteria for Waiting and Throughput Times	
A	Care pathway describes the roles of the different disciplines involved in the MDT Formalized position	Breast MDT core members: surgical and medical oncology, radiotherapy, imaging, pathology, and nursing Non-core members: plastic surgery, psycho-oncology, medical physics and rehabilitation, genetics, neurosurgery, orthopedics, and palliative care	Pathways were developed using national and international guidelines	Pathway describes all steps, from patient admission process and referral requirements through follow-up	Pathway contains maximum waiting times between the various steps
B	Care pathway describes the roles of the different disciplines involved in the MDT	Breast MDT members: breast surgeons, radiologists, medical oncologists, radiotherapist, pathologist, geneticist, nurse, and administrator	Pathways are based on well-known international guidelines for breast cancer (EUSOMA ³² and SIS ³³ guidelines)	Pathway contains information about epidemiologic, scientific, genetic, and clinical steps	Government sets maximum waiting times for certain steps of the pathway
C	Care pathway describes the roles of the different disciplines involved in the MDT	Breast MDT members: breast surgeon, clinical oncologist, plastic surgeon, psychologist, imaging diagnostician, and pathologist	Pathways are based on international guidelines, such as NCCN ³⁴ and ESMO ³⁵ guidelines, and national guidelines	Pathway describes all steps, from patient admission through follow-up, including all clinical treatment algorithms	Government does not limit maximum waiting times in general
D	Care pathway describes the roles of the different disciplines involved and which cases should be presented in the MDT meeting	Breast MDT members: medical oncologist, radiotherapist, radiologist, surgeon, and pathologist	Pathways are based on international and national guidelines	Pathway describes all steps, from patient registration through follow-up, including all clinical diagnosis and treatment algorithms	Government only limits the time from referral by the general practitioner to the time the specialized physician sees the patient There are currently no standard means of measuring waiting and throughput times between the steps in the pathway
E	MDT for each pathway	Breast MDT members, preoperative setting: radiologist, surgeon, pathologist, and breast nurse Breast MDT members, postoperative setting: surgeon, medical oncologist, radiotherapist, pathologist, nuclear medicine doctor, and breast nurse	Pathways are based on national and international guidelines	Pathway describes all steps, from patient admission process and referral requirements through follow-up	Government decides on maximum waiting times between the steps within the pathway
F	Care pathway describes the roles of the different disciplines involved	Breast MDT members: breast and plastic surgeons, medical oncologists, radiotherapists, geneticists, and pathologists	Pathways are based on international and national guidelines, clinical study protocols, and conferences	Pathway describes the different steps taken in the diagnostic and treatment phases, and the available clinical diagnosis and treatment algorithms for each of these steps; follow-up is also described	Government sets maximum waiting times between the various steps in the pathway Patient access to the hospital is described and monitored in the pathway
G	All participants in MDTs were involved in developing and reviewing the clinical pathways Therefore, clinical pathways are integrated in the activities of MDTs and, vice versa, clinical pathways include MDTs as a crucial step	Breast MDT members: surgeons, oncologists, radiotherapists, pathologists, and nurses On call: radiologists, plastic surgeons, and psychologists	Pathways are based on international and national guidelines and conferences	Pathway describes all steps, from patient registration through follow-up, including all clinical diagnosis and treatment algorithms	There are set waiting times within the pathway; these are monitored inside the regional portal

Abbreviations: EUSOMA, European Society of Breast Cancer Specialists; MDT, multidisciplinary team; SIS, Senologic International Society.

and external evaluation. The clinical governance department at center A performs a patient pathway audit for every pathology clinic (IPU). The breast cancer pathway at center B was evaluated through external accreditation. At center D, evaluation is performed internally and externally. At center E, the pathways were not systematically evaluated; an indicator matrix containing indicators from various sources was being developed for this purpose. At center F, pathways were collectively discussed through MDT meetings in close collaboration with the Healthcare Directorate and the Medical Directorate, who are in charge of supervising care pathways. Center G organized an extensive

internal and external evaluation (by a regional and national agency). Overall pathway evaluation seems to focus mainly on waiting and throughput times and less on quality performance.

IPU Development

By definition, cancer centers are organized around a medical condition (ie, cancer). However, when differentiating and considering specific types of cancer, Table 6 shows that for breast cancer this can only be seen at center A (other than the breast IPU, this center has an IPU for 10 other cancer types), center B (breast cancer only IPU), and center G (IPU for 5 types of cancer, including breast). Table 6 shows the degree of implemen-

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Table 5. General Pathway Evaluation Policies

Cancer Center	Evaluation Method	Examples of Type of Measurements	Frequency	Outcomes Evaluation
A	Clinical governance conducts a patient pathway audit for every pathology clinic, in which times between referral/appointment/diagnosing/treatment are evaluated, including reasons for delay Compliance with therapeutic guidance documents is also checked	Efficiency, clinical patient outcomes	Every trimester	Quarterly reports and an annual report Another annual report with clinical indicators also compares the results of the past 2 years
B	Evaluation of the breast pathway is performed by an external evaluation body by means of accreditation	Efficiency; each year the pathway leadership conducts or participates in ≥2 studies (eg, CP3R) that measure quality and/or outcomes	Once annually	Outcome of the evaluation was a structured pathway
C	MDT members evaluate the applicable pathways	Evaluation of patient flow efficiency	Every 3 years	Outcomes of the pathway evaluation are discussed by members of the relevant MDT and included in the annual report
D	Pathways are evaluated through review of patient files There are no goals set beforehand Internal evaluation is performed by 2 means: quality-of-care control team and internal audit External evaluation is performed by the national healthcare insurance	Monitoring of patient complaints, suggestions, and diagnosis and treatment guidelines adherence, and evaluation of quality of services	Internally every 3 months, externally quarterly	The quality-of-care team and the external evaluation reports are presented to the center's medical council and the general manager As a result of the evaluation, the responsible parties are informed of their individual performance and provided with suggestions for improvement
E	Pathways were not systematically monitored; they will be monitored through specific indicators To date, proposed indicators are EUSOMA's indicators for breast unit accreditation, integrated with internal performance indicators and indicators proposed by BENCH-CAN ¹⁴	EUSOMA's indicators for breast unit; BENCH-CAN indicators ¹⁴	Not applicable	Not applicable
F	Medical Directorate coordinates the activities for the evaluation of diagnostic and therapeutic pathways (PDTA) The evaluation process is performed inside the center in close collaboration with the clinical and surgery departments and under supervision of the Healthcare Directorate	Efficacy and efficiency of pathways measured through evaluation of clinical outcomes (eg, mortality rates, readmissions, external audits) and quality outcomes (eg, adherence to the PDTA, customer satisfaction)	Every 6 months	Findings from the evaluation process are implemented by the clinical and surgery departments in order to translate results into practice Quality, Training, and Privacy Office coordinates the quality improvement process of the pathways in close collaboration with the departments
G	<u>External evaluation:</u> External evaluation and benchmark are provided by 2 different centers (regional and national) Regional benchmarks for single hospital performance against other regional hospitals 2 types of evaluation are provided: <ul style="list-style-type: none"> • General evaluation at the department level • Evaluation on the pathway level The national evaluation is performed by a national agency that once a year displays on a Web portal the outcome measures (eg mortality), benchmarked against all national hospitals <u>Internal evaluation:</u> Internal system calculates and provides all performance indicators to the hospital Review of clinical pathways is based on a periodical review of patient records Planning is in process to include the pathways inside the EPR system in order to facilitate clinical decisions and improve standardization	30-day mortality, readmissions, surgical reinterventions, transfers to other acute hospitals, efficiency (eg, percentage of patients operated on within 60 days from mammography)	Continuous improvement Small/relevant every year as evidence changes Formally/big every 3 years	Several actions are implemented to improve outcomes and quality provided to patients as a result of the pathway evaluation In breast cancer, for example, starting from the group of evaluation indicators, there is continuous improvement with an annual evaluation visit by EUSOMA

Abbreviations: CP3R, Cancer Program Practice Profile Reports; EPR, electronic patient record; EUSOMA, European Society of Breast Cancer Specialists; MDT, multidisciplinary team; PDTA, Diagnostic-Therapeutic-Healthcare Protocol; SIS, Senologic International Society.

tation of the IPU features, scored with a 3-point system. This is linked to the common organization unit where, in all centers, providers involved are members of the cancer center, but in center A, B, and G they are members of a specific organizational unit.

All centers work with tumor-specific MDTs, hence dedicating time to the specific condition. Members of the MDT are however not always solely or clearly dedicated to one tumor type or unit. A

physician heads the MDT, although only centers E and G have a dedicated case manager.

Regarding responsibility for the full care cycle, this was the case for MDTs in all centers. Nevertheless, only centers A and G indicated that the unit encompassed rehabilitative care and supporting services (eg, psychosocial services). For other centers, these were part of services for the whole center, as were patient education, engagement, and follow-up, which are seen as integral to care but not specifically arranged for a unit (except

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Table 6. Degree of Development of Breast Cancer IPUs Based on Criteria

Criterion	Cancer Center						
	A	B	C	D	E	F	G
Organized around a medical condition	+++	+++	++	++	++	++	+++
MDT with dedicated time to the condition	+++	+++	++	++	++	++	+++
Providers involved are members of or affiliated with a common organizational unit	+++	++	++	++	++	++	+++
Physician team captain or clinical care manager oversees care process	++	++	+	+	++	+	+++
Providers function as a team, meeting formally and informally on a regular basis	+++	+++	+++	+++	+++	+++	+++
Responsibility for the full cycle of care for the condition, encompassing outpatient, inpatient, and rehabilitative care, as well as supporting services (eg, nutrition, social work, behavioral health)	+++	++	++	++	++	++	+++
Incorporates patient education, engagement, and follow-up as integral to care	+++	+	+	+	+	+	+++
Measures outcomes, costs, and processes for each patient using a common information platform	+++	+	+	+	+	+	+++
Accepts joint accountability for outcomes and costs	+++						+++
Single administrative and scheduling structure	+++	+	++	+	+	+	+++
Co-located in dedicated facilities	+++	+++	+++	+++	+++	+++	+++
Total degree of implementation out of 33	32	21	19	18	19	18	33

A triple plus symbol (+++) indicates that the feature is fully developed, a double plus symbol (++) indicates that the feature is mostly developed, and a single plus symbol (+) indicates that the feature is partially implemented. Blank cells indicate that the criterion could not be assessed. Abbreviations: IPUs, integrated practice units; MDT, multidisciplinary team.

for centers A and G). However, center G mentioned that many patients admitted to the unit already have a diagnosis and many complete follow-up elsewhere. This compromises the measurement of outcomes of the entire care pathway for each patient.

All centers measure outcomes, costs, and processes; however, most (n=5) do this for the whole center and not for specific units. Because the IPU-related definition of “outcomes that most matter to patients” is not specified, and consensus-based International Consortium for Health Outcomes Measurement sets²⁴ were not yet available for breast cancer at the time of measurement, we could not establish whether this was actually the case. Two centers, however, reported that they have patient-based focus groups and measure breast cancer-specific process indicators and outcomes and use this feedback to improve care provision.

Centers are either lacking data platforms that allow for the collection of specific inputs (in terms of human resources and finances) and outcomes per unit, or do not currently collect these data. Similarly, specific administrative and scheduling structures were lacking or not fully implemented, because centers C, D, E, and F do not have a formalized unit. Although center B indicated they have a breast IPU, whether they have an accompanying dedicated scheduling structure was unclear.

Overall, 3 types of centers can be identified: those that have the IPU structure implemented (centers A and G), those that have the IPU partially implemented (center B), and those that have certain features of the IPU but did not develop or implement the IPU (centers C, D, E, F). IPU imple-

mentation outcomes for the breast unit in center A are described in [supplemental eAppendix 2](#).

Discussion

This study developed a benchmark tool to assess development, implementation, and evaluation of cancer care pathways. The tool was successfully tested in 7 cancer centers to assess its suitability for yielding improvement suggestions regarding pathway organization and providing data for exploring the status of IPU organization.

The director concerned with pathways is usually a medical specialist, which is in accordance with the assertion of Polite et al⁹ that the responsibility for oncology pathway development must always lie primarily with clinicians. Our data suggest that centers have no clear strategy when developing pathways, which is consistent with the findings of Vanhaecht et al,⁸ who showed that a minority of sampled countries (43%) used a systematic approach to develop, implement, and evaluate care pathways. None of the centers work with previously established goals for pathway evaluation. In-built continuous evaluation and follow-up should guarantee the effectiveness of care pathways,²⁵ and was therefore recommended as an improvement opportunity for cancer centers.

Porter²⁶ stated that care pathways are beneficial but not sufficient for delivering value-based, high-quality care and recommended the establishment of IPUs. In this explorative study assessing the centers' benchmark data against the criteria of an IPU,²⁰ we identified 3 groups. Group 1, consisting of centers fully meeting all

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criteria (centers A and G), and group 2, consisting of centers partially meeting the criteria (center B), both reported having a breast IPU. Based on our data, we agree that centers A and G have an IPU for breast cancer. However, for center B, improvements in 4 criteria are required to have a fully implemented IPU. Group 3, consisting of centers meeting only some of the criteria, did not report having an IPU. The fact they meet some of the criteria is likely due to the pathway organization and the organization of MDTs (which are key features of an IPU) that has been developed for years.²⁷ Dedicating specific resources, such as staff to measure outcomes specific to the IPU and IPU-specific administrative and scheduling systems, were lacking or not fully implemented. Centers were recommended to improve registration of patient outcomes using permanent patient-based input, both for clinical (eg, securing state-of-the-art treatment innovations, introducing cognitive testing for side effects on brain functioning as a consequence of chemotherapy) and organizational aspects (eg, determining the impact of waiting on a diagnosis or treatment on a patient's quality of life). Measuring outcomes "that matter most to patients" requires improvement in all centers; however, this is a continuously developing objective and requires a permanent or periodic update involving patient input.

Sarai et al²⁸ identified IPUs as units in which providers commit a substantial portion of their time to treating a focused set of care pathways, implying that pathways are IPU building blocks. In our study, we used pathways as tools to map current organizational processes and identified areas to improve these processes where necessary (value and quality improvement on the process level). IPUs are structured organizational units in which the process identified through the pathway occurs. IPU organization seems to require an organizational change from a focus on discipline-based departments (eg, radiotherapy) to those departments facilitating the IPUs, which are pathway-based. In this study, IPUs are therefore seen as tools for quality and value improvement on a strategic level.

Evaluation of the breast IPU in center A showed that improved efficiency led to more time available for patients, though higher volumes, economies of scale, and improvements in quality. Although based on a small number, this corresponds to findings by Low et al,²⁹ who found that IPUs resulted in reduced readmissions among patients at highest risk of readmission. Enthoven et al,³⁰ however, argue that for patients with multiple

morbidities, the IPU concept is unfavorable. This stems from the original ideas of Porter and Teisberg¹⁰ that IPUs should be entities with sufficient degrees of "self-organization." No published information exists on how patients feel about this development, which needs to be known in order to facilitate true patient-centered care.

Developing and implementing an IPU has barriers, which will vary based on the condition being treated, provider organization, and health system characteristics. Keswani et al³¹ divided these barriers into 3 subcategories: operational, technology, and payment/contracting. Supplemental eAppendix 3 provides an overview of the barriers.

This study has several limitations. First, we did not emphasize the detailed clinical content of the pathway. Future research focusing on the exact pathway content as part of the benchmark will further enable international knowledge-sharing. Furthermore, because the initial focus of the indicators was pathways, we had to deduct the IPU criteria from the material. This assessment was aided by focusing the site visits on breast cancer departments, which showed whether centers had the IPU organization in place. Indicators that specifically examine IPU organization should be refined and could help provide a more thorough assessment based on IPU criteria. This study only focused on breast cancer; future studies should include patient cohorts that have shared medical (and social) needs. Additionally, because few scientific publications are available, some degree of subjectivity was inevitable in the explorative assessment of the benchmark data against the IPU criteria. Further studies of scale and cutoff points that distinguish between different levels of IPU development are recommended.

This study's evaluation of an IPU for breast cancer care showed performance improvements in terms of efficiency and finances. Future research focusing on more extensive (patient) outcome evaluation over multiple years will allow comparison between IPU-organized and non-IPU-organized cancer centers to determine whether IPUs add value and actually decrease costs for the health system.

Finally, this study focused on a limited number of cancer centers (n=7) and collected data for 1 year, with all of the centers located in Europe, although the original theory stems from the United States. However, although health and financing systems differ, the findings are likely applicable in the United States. Future research should include larger series

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(including general hospitals with an oncology department) in multiple countries (including different continents) and over multiple years to assess actual discriminative capabilities of the tool and establish the sensitivity for changes over time.

Conclusions

We successfully developed and piloted a benchmark tool to compare and elaborate on organizational performance, and provided recommendations for improving the organization of cancer care pathways

(both implementation and evaluation). The data generated through the benchmark enabled exploration of the status of IPU development. Pathways can be seen as the process that occurs in the strategic unit of the IPU. Our assessments found that centers varied in fulfilling IPU criteria, usually lacking specific resources such as staff; in measuring IPU-specific outcomes; and in their IPU-specific administrative and scheduling systems. Development of an IPU requires a strategic organization change with several implications for operations, technology, and payment and contracting.

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