Piloting a Hospital Pharmacy Performance Model in the Face of Province-Wide Implementation of Activity-Based Funding in Quebec Health Care Centres

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ABSTRACT

Background: In the face of province-wide implementation of activitybased hospital funding in Quebec, a need arose to effectively measure pharmacists' contributions along the patient care trajectory and to enable pharmacy benchmarking using valid performance indicators.

Objectives: A 3-phase project was initiated to measure the performance and impact of pharmacists and pharmacy departments. Phases 2 and 3, described here, focused on gradually implementing, in various health care centres, the priority indicators selected in phase 1.

Methods: The project involved multiple committees overseeing the implementation, data collection, analysis, and documentation of 18 performance indicators. Specific tools were developed to facilitate data collection and encourage pharmacists' participation. A feedback survey was used to document pharmacists' experiences.

Results: Substantial data were gathered over 3 years (2017 to 2020), involving 358 pharmacists from 6 health care centres. The overall contribution rate to the daily data collection from front-line pharmacists was 55%. The feedback survey revealed that, of the various communication tools used to promote the project, in-person events were better perceived by the front-line pharmacists than online tools. Of the 183 respondents to the survey, most (94%, n = 172) believed it was important to collect data to document pharmacists' activities, and 82% (n = 150) saw the project as relevant to the upcoming activity-based funding system.

Conclusions: Despite challenges, progress was made in defining relevant indicators, adjusting the list generated during phase 1, and reaching a consensus on 16 indicators. Stakeholders expressed interest, emphasizing the importance of documenting pharmacists' activities. The project has laid the foundation for demonstrating the value of pharmacists along the patient care trajectory and measuring pharmacy departments' performance. However, more integrated technological solutions are needed for province-wide implementation.

Keywords: performance indicators, hospital pharmacy benchmarking, quality improvement, pharmacy performance measurement

RÉSUMÉ

Contexte : Dans le contexte de la mise en œuvre à l'échelle provinciale du financement à l'activité des hôpitaux au Québec, le besoin s'est fait sentir de mesurer efficacement la contribution des pharmaciens tout au long du parcours de soins aux patients et de permettre l'analyse comparative des départements de pharmacie à l'aide d'indicateurs de performance valables.

Objectifs : Un projet en trois phases a été lancé pour mesurer la performance et l'impact des pharmaciens et des départements de pharmacie. Les phases 2 et 3 décrites dans cet article portaient sur la mise en œuvre progressive dans différents établissements de santé des indicateurs prioritaires retenus lors de la phase 1.

Méthodologie : Le projet impliquait plusieurs comités chargés de superviser la mise en œuvre, la collecte de données, l'analyse et la documentation des 18 indicateurs de performance. Des outils spécifiques ont été préparés pour faciliter la collecte des données et encourager la participation des pharmaciens. Un sondage de rétroaction a été utilisé pour recueillir les impressions des pharmaciens.

Résultats: Une quantité considérable de données a été recueillie sur une période de 3 ans (2017 à 2020) auprès de 358 pharmaciens de 6 établissements de santé. Le taux global de participation à la collecte quotidienne de données auprès des pharmaciens était de 55 %. Le sondage de rétroaction auprès des pharmaciens a révélé que, parmi les divers outils de communication utilisés pour promouvoir le projet, les événements en personne étaient jugés plus utiles que les outils en ligne. Des 183 répondants au sondage, la plupart (94 %, n = 172) ont estimé qu'il était important de recueillir des données pour recenser les activités des pharmaciens, et 82 % (n = 150) ont estimé que le projet était pertinent pour le futur système de financement à l'activité.

Conclusions: Malgré les défis, des progrès ont été réalisés quant à la définition des indicateurs pertinents en ajustant la liste générée au cours de la phase 1 pour établir un consensus concernant 16 indicateurs. Les parties prenantes ont manifesté leur intérêt, soulignant l'importance de recenser les activités des pharmaciens. Le projet a jeté les bases pour démontrer la valeur des pharmaciens tout au long du parcours de soins des patients et mesurer la performance des départements de pharmacie. Cependant, des solutions technologiques plus intégrées sont nécessaires pour permettre la mise en œuvre à l'échelle provinciale. **Mots-clés**: indicateurs de performance, analyse comparative de la pharmacie hospitalière, amélioration de la qualité, mesure de la performance en pharmacie

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INTRODUCTION

Performance measurement in health care is the subject of ongoing research.¹ Large-scale performance measurement helps identify areas for improvement, streamline processes, and ultimately enhance patient outcomes.^{2,3} It also helps in comparing practices across different health care centres and against national and international standards, fostering the sharing of knowledge to drive best practices.⁴ The surge in activity-based funding for hospitals underscores the significance of comprehensive activity measurement, challenging administrators and various caregivers to justify their value in contributing to patient outcomes.5-7 To tackle this issue, many authors in the United States and Europe have proposed various sets of indicators.⁸⁻¹⁴ These initiatives reflect differing priorities and methods across countries. In Canada, no nationwide system exists for benchmarking hospital pharmacy practices. About a decade ago, the Canadian Society of Hospital Pharmacy introduced clinical pharmacy key performance indicators (cpKPIs) to evaluate pharmaceutical care, although these do not cover all aspects of pharmacy practice.³

Continuous, large-scale measurement of hospital pharmacy practice has proven difficult due to the heterogeneity of data sources, the lack of specific, automated collection tools, and the intensity of resources required.¹⁵⁻¹⁷ As such, most attempts at performing measurements have focused on a narrower scope of activities, such as medication error rates, adherence to clinical guidelines, or the efficient management of pharmaceutical resources.¹⁸⁻²⁴ Many efforts have centred on aspects of workload or technical efficiency, whereas a patient-centric approach might work better to justify the added value of pharmacists.²⁵⁻²⁸

In the face of province-wide implementation of activitybased hospital funding in Quebec, a need emerged to better quantify hospital pharmacy activity, impact, and performance. With valid and relevant performance indicators representing all areas of professional practice, pharmacy departments would be better equipped to measure and benchmark their performance, demonstrate their value throughout the patient care trajectory, and provide input for an activity-based funding system. In a recent study,²⁹ our team used a consultative approach to develop a performance framework and associated indicators, emphasizing the need for balanced, easily documented indicators. The pilot project described here was undertaken to validate these indicators, which are proposed for use in hospital activity-based funding systems.

METHODS

The initiative was led by a steering committee of 3 senior pharmacists (F.B., F.P., and L.V.) and 3 health care data analysis consultants (including S.S.), with support from an external advisory committee, consisting of a former hospital Chief Executive Officer (CEO), 5 Chief Pharmacy Officers, a quality assurance coordinator, and a regulatory authority observer, all selected for their expertise and strategic roles in the health care system. The central steering committee met monthly to review the project's progress and to direct the work according to the issues and problems raised. At the beginning and the end of each of the 3 phases, the advisory committee gave its opinion on the direction to be taken.

Phase 1, described in our previous article,²⁹ ended with the steering committee's selection of a limited set of indicators that can be used to demonstrate the benefits of pharmacy activities along the patient care continuum. The development of a performance framework, a literature review, and an extensive consultation process supported their choice of indicators. Selection rounds were used to ensure representation of 5 framework dimensions (appropriateness, quality and safety, efficiency, innovation and continuous improvement, and organizational structure) and 5 professional roles (pharmaceutical care, drug distribution, education of trainees and colleagues, research, and management and professional matters). Of the 150 indicators initially identified during phase 1, 24 were selected. With the experimental phases in mind, 13 indicators were prioritized for their low measurement complexity and immediate relevance to managers, concretely improving the performance assessment tools in use, as well as allowing for benchmarking between institutions (Table 1; see column for indicators at start of the pilot project). The prioritized indicators covered all 5 professional roles, but only 4 of the 5 dimensions of the framework. Indeed, indicators lying in the organizational structure dimension were given a high complexity score, leading to a long-term priority level. As such, they could not be used in the pilot project.

Quebec's health care centres coordinate and deliver services in designated regions, encompassing facilities with specific mandates such as acute or long-term care. Each heath care centre has one integrated pharmacy department, which serves all of the centre's facilities, from one or multiple sites. From among all health care centres, those suitable to serve as pilot sites were identified through interest expressed by department heads and anticipated resource availability, with representation of academic and non-academic centres, diverse clienteles (short-term, long-term, and ambulatory), and a variety of pharmacy information systems. The participating health care centres treated this project as a quality improvement initiative and therefore did not seek exemptions from their respective research ethics boards. Each health care centre's CEO signed a commitment and confidentiality agreement. Results from the pilot sites were anonymized to comply with confidentiality requirements.

Phases 2 and 3 are referred to jointly as the "pilot project", which was set up according to the following principles:

TABLE 1	(part 1 of 2). Pe	rformance Indicators	TABLE 1 (part 1 of 2). Performance Indicators Tested in the Pilot Study				
	Fra	Framework	India	Indicator Definition	Sour	Source of Data	
Code ^a	Dimension	Professional Role	Start of Pilot Project	End of Pilot Project	Manual by Pharmacists	Manual by Information Pharmacists Systems	Ad hoc
BUNDLE 1							
Q1	Quality and safety	Pharmaceutical care	Number of medication reconciliations on admission carried out or validated by pharmacists by volume of clientele	Number of medication reconciliations during general admissions and hemato-oncology outpatient visits carried out or validated by pharmacists by volume of clientele ^b	>	>	
Q2	Quality and safety	Pharmaceutical care	Proportion of patients for whom there is a documented medication reconciliation at discharge	Withdrawn	>	>	
A3	Appropriateness	Pharmaceutical care	Number of individual counselling sessions offered by pharmacists by volume of clientele	Same	>	>	
Q3	Quality and safety	Drug distribution	Number of medication errors by volume of clientele	Same		>	
E2	Efficiency	Drug distribution	Ratio of pharmacy technicians' worked hours to pharmacists' worked hours	Same		>	
12	Innovation and continuous improvement	Management and professional matters	Proportion of the hospital's clinical governance bodies on which a pharmacist sits ^c	Same			>
BUNDLE 2							
A2	Appropriateness	All 5 roles ^d	Share of each of the professional role categories in the total worked hours of all pharmacists	Same	>		
A1	Appropriateness	Pharmaceutical care	Hours worked in pharmaceutical care by volume of clientele	Same	>	>	
A4	Appropriateness	Research	Proportion of pharmacists' worked hours devoted to drug utilization reviews	Withdrawn		>	>
Q4	Quality and safety	Management and professional matters	Proportion of pharmacists with an advanced pharmacotherapy master's degree	Same			>

TABLE 1	TABLE 1 (part 2 of 2). Performance Indicators Tested in the Pilot Study						
	Frai	Framework	Indic	Indicator Definition	Sour	Source of Data	
Code ^a	Dimension	Professional Role	Start of Pilot Project	End of Pilot Project	Manual by Pharmacists	Information Systems	Ad hoc
BUNDLE 3	e						
A7	Appropriateness	Pharmaceutical care	New indicator ^e	Share of each of the care sectors in the total worked hours of all pharmacists	>		
E7	Efficiency	Pharmaceutical care	New indicator ^e	Share of drug therapy problems resolved by pharmacists assigned to either pharmaceutical care or drug distribution	>		
E8	Efficiency	Pharmaceutical care	New indicator ^e	Number of care activities per day for pharmacists assigned to pharmaceutical care	(🗸 †)		
ES	Efficiency	Pharmaceutical care	Indicator imported from the lower-priority scores in phase $1^{\rm e}$	Number of drug therapy problems resolved by pharmacists assigned to direct patient care by volume of clientele	>	>	
Q10	Quality and safety	Drug distribution	New indicator ^e	Share of incident reports for medication errors that are attributed to pharmacy		>	
E1	Efficiency	Drug distribution	Ratio of the number of sterile drug preparations to pharmacy technician assistants' worked hours	Same			>
Ξ	Efficiency	Drug distribution	Average processing time for urgent or priority prescriptions	Same			>
Ξ	Innovation and continuous improvement	Research and education of trainees and colleagues	Proportion of pharmacists who participated in outreach activities ^g	Same			>
^a The codes improveme ^b Medication medication ^c Board of C Committee ^d All 5 roles ^e For phase ^f Indicator E therapy pro	^a The codes were assigned using the fi improvement, Q = quality and safety. ^b Medication reconciliation performed medication management is a major co Gboard of Directors; Council of Physici Committee; Infection Prevention and ^d All 5 roles: pharmaceutical care, dru ^e For phase 3, indicators A7, E7, E8, ar ^f indicator E8 was computed using dat therapy problems resolved).	⁴ The codes were assigned using the first letter of the framework dimension improvement, Q = quality and safety. ^b Medication reconciliation performed for hemato-oncology outpatient visit medication management is a major component of care. ⁹ Doard of Directors, Council of Physicians, Dentists and Pharmacists Execut Committee; Infection Prevention and Control Committee; and Risk and Que ⁴ All 5 roles: pharmaceutical care, drug distribution, education of trainees a ^e For phase 3, indicators A7, E7, E8, and Q10 were created, and indicator E5 ^f Indicator E8 was computed using data already collected for other indicato therapy problems resolved).		⁴ The codes were assigned using the first letter of the framework dimension to which the indicator belonged, followed by a consecutive number. A = appropriateness, E = efficiency, I = innovation and continuous improvement, Q = quality and safety. ^b Medication reconciliation performed for hemato-oncology outpatient visits was added to test the measurement of an Accreditation Canada Required Organizational Practice for ambulatory care visits when medication management is a major component of care. ^b Medication management is a major component of care. ^c Borand of Directors; Council of Physicians, Dentists and Pharmacists Executive Committee; Pharmacy and Therapeutics Committee; Credential Review Committee; Medical, Dental and Pharmaceutical Evaluation Committee; Infection Prevention and Control Committee; and Risk and Quality Management Committee. ^c All 5 roles: pharmaceutical care, drug distribution, education of trainees and colleagues, research, and management and professional matters. ^c For phase 3, indicators A7, E7, E8, and Q10 were created, and indicator E6 was imported from the lower-priority scores in phase 1 to better reflect front-line pharmacists' practice. ^c For phase 3, indicators A7, E7, E8, and Q10 were created, and indicators (e.g., individual counselling sessions, medication reconciliation on general admission or hemato-oncology outpatient visits, and drug therapy problems resolved). ^c For phase accounted using data already collected for other indicators (e.g., individual counselling sessions, medication reconciliation on general admission or hemato-oncology outpatient visits, and drug therapy problems resolved).	icy, I = innovatio r ambulatory can al and Pharmace al and outpatient ce. another hospital	n and continuo e visits when utical Evaluatio visits, and dru	

- Ensure the support and commitment of the CEO from each pilot site.
- Deploy priority indicators in 3 successive bundles for progressive implementation in the pilot sites, maintaining a steady pace.
- Implement the simplest and easiest-to-document indicators first, to generate enthusiasm and mobilization.
- Favour documentation solutions that minimize pharmacist data entry time and manual data entry.
- Ensure data quality control.

On-site working committees were created to deploy the indicators at their respective sites. Each working committee comprised the head of the pharmacy department, a project manager, and a clinical-administrative information systems manager.

With the support of the on-site working committees, the steering committee carried out the following activities sequentially and repeatedly throughout the deployment, by indicator bundle and pilot site:

- 1. Organization of deployment
 - a. Adjustment of indicator variables
 - b. Update of data collection tools
 - c. Communication and promotion activities at pilot sites
- 2. Data collection and extraction
 - a. Daily data collection from pharmacists
 - b. Quarterly data extraction from facilities' information systems
 - c. Quarterly ad hoc data collection for specific indicators
- 3. Analysis
 - a. Processing and analysis of indicator results
 - b. Monitoring of pharmacists' participation in data collection
- 4. Documentation and feedback to pilot sites

Figure 1 displays the timeline of the major deployment steps of the pilot project, which ran from June 2017 to October 2020. Phase 2 marked the start of experimentation, with the first bundle of indicators (Q1, Q2, A3, Q3, E2, and I2; see Table 1) being deployed in certain facilities at the first pilot site. The first bundle was then deployed in a few facilities at 2 other pilot sites. Subsequently, bundles 1 and 2 (A2, A1, A4, and Q4) were deployed in all short- and long-term facilities at the 3 sites. This second phase of the overall project lasted 16 months. Phase 3 saw the addition of indicators from bundle 3 (A7, E7, E8, E5, Q10, E1, E3, and I1) and, more importantly, aimed to validate or question the previously adopted solutions and to experiment on a larger scale, over 2 years and across 6 pilot sites.

Master indicator fact sheets included the definition of each indicator, data sources, calculation and interpretation methods, and update frequency. For 9 of the 18 indicators tested, manual data collection by pharmacists was required. A data collection tool was built using a stand-alone web survey form (SurveyGizmo) with embedded definitions and instructions, allowing pharmacists to report their activities on a daily basis. Pharmacist participation remained voluntary throughout the whole pilot project. With the support of the advisory committee, the steering committee set an ambitious, arbitrarily chosen target of 70% for the daily participation rate at each pilot site, calculated as the average number of forms submitted divided by the average number of expected pharmacist shifts during weekdays. During phase 3, individual participation was disclosed to the on-site working committees, so that they could follow up with less assiduous pharmacists. The actual data collected by each pharmacist remained confidential, including for pharmacy administration, in accordance with the partnership agreements.

Other specific tools were developed and updated as the pilot project progressed. For example, paper checklists

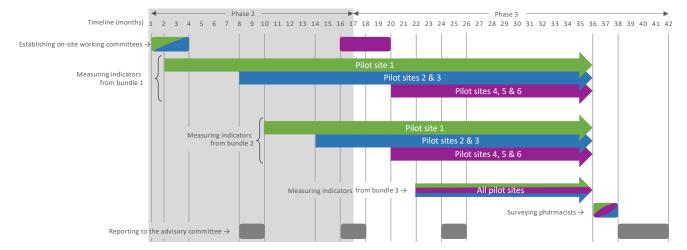


FIGURE 1. Timeline of the major deployment steps of the pilot project. The measurement of indicators was deployed in 3 bundles at the pilot sites. Green refers to activities at pilot site 1; blue to activities at pilot sites 2 and 3; and purple to activities at pilot sites 4, 5, and 6.

enabled pharmacists to collate requested data over the course of each day. The on-site working committees received monthly updates, with tables and graphs summarizing pharmacists' participation. In response to the participation rate observed during phase 2, the steering committee implemented various communication tools to engage pharmacists in data collection during phase 3. These included a dedicated webpage, an animated video, an interactive instructional document, a quarterly newsletter, systematic weekly email reminders, access to the history of selfsubmitted forms and personal statistics, and in-person meetings to promote the project and obtain feedback.

Quarterly data extraction and compilation were completed by the on-site working committees using pharmacy and financial information systems, along with accidentincident registries and some ad hoc data collection based on forms provided by the steering committee. All results were integrated into a master reporting template. Upon receiving the data, the steering committee supplemented them with information from the web-based daily data collection tool. The data were also validated against the health care centres' financial and statistical reports to ensure quality. Consistency checks were performed by comparing results across the 6 pilot sites, and corrections were applied with support from the on-site working committees when necessary. Standardized tables and graphs facilitated clear presentation and analysis. During phase 2, each pilot site had access only to its own results, to comply with confidentiality agreements. Data were analyzed by facility and care sector where this could be done without compromising respondents' anonymity. Only the steering committee could perform benchmarking. Following requests from the pilot sites and amendment of project agreements, consolidated data were disclosed to the on-site working committees in phase 3, which allowed the pilot sites to benchmark their respective results.

After daily data collection ended, all pharmacists at the pilot sites, regardless of their participation in data collection, were invited to complete an online survey concerning their experience of the project. The purpose of this survey was to gather relevant information to complete the analysis and support recommendations at the conclusion of the project.

RESULTS

The pilot project encompassed 358 hospital pharmacists across 6 pilot sites, representing different types of hospitals (with various missions, including academic and non-academic) and different health regions (Table 2). Data collection lasted 1 to 3 years, depending on the pilot site. By the end of the project, the overall participation rate by pharmacists in daily data collection stood at 55%, falling short of the preset target of 70%. Notably, only a small percentage (7%, n = 25) of pharmacists failed to submit at least 1 form. Analysis of the participation rate over time showed a downward trend at several pilot sites, reflecting a gradual loss of momentum.

Data extraction from health care centres' information systems posed several difficulties owing to the number of systems involved and their lack of integration. In most cases, the extracted files had to be manipulated before the data could be integrated into the indicator calculation tool. Additionally, substantial modifications to the definitions of indicators and their component variables were required. For instance, the indicator reporting on admission medication reconciliation (indicator Q1) was initially measured only for inpatients, including those in long-term care. However, feedback from the pilot sites highlighted the need to extend the scope of this indicator to encompass medication reconciliations conducted during hemato-oncology outpatient visits, to align with Required Organizational Practices issued by Accreditation Canada for ambulatory care visits when medication management is a major component of care.30

Two indicators were withdrawn at the end of phase 2. Despite its significance for care continuity, the indicator measuring medication reconciliation at discharge (indicator Q2) was withdrawn after nearly a year of the pilot due to challenges with data collection. According to the

				No. of	Facilities
Pilot Site	No. of Pharmacists	Academic	Region	Acute Care	Long-Term Care
1	97	Yes	Urban	5	1
2	57	No	Suburban	3	15
3	44	No	Suburban	7	11
4	85	Yes	Urban	5	12
5	45	No	Rural	6	13
6	30	Yes	Urban	1	0
Total	358			27	52

TABLE 2. Characteristics of Pilot Sites

project definitions established in phase 1, this indicator was intended to account only for medication reconciliation directly involving a pharmacist at some point in the process. However, the tools and definitions used at the pilot sites did not enable differentiation between medication reconciliation exercises that did and did not involve a pharmacist. The second indicator that was withdrawn related to hours worked on drug utilization reviews (indicator A4). It was deemed to concern too few hospitals and too few worked hours to justify manual data collection by pharmacists.

Feedback during phase 2 revealed that several frontline pharmacists felt that the measured activities did not reflect a substantial portion of their practice. To address this issue and increase participation, 5 new indicators measuring pharmaceutical care and drug distribution were added in phase 3. During phase 1, indicator E5, which reports on drug therapy problems resolved by pharmacists, was initially assigned a lower priority due to the anticipated intensity of data collection and thus was not included at the start of the pilot project. However, it was reinstated to better highlight the clinical activities of frontline pharmacists. Among the new indicators, 4 required manual data collection by pharmacists (indicators A7, E5, E7, and E8). One of these (indicator E8) was calculated using data already recorded by pharmacists for other indicators (Table 1).

Table 3 shows a few selected results from the 16 final indicators. Results must be interpreted with caution, as they are subject to many contextual factors across the various pilot sites, such as staff shortages and level of participation

TABLE	3 (part 1 of 2). Selected Results from Testing	Indicators during the Pilot Project
Code ^a	Indicator Definition	Selected Results ^b
All 5 pr	rofessional roles ^c	
A2	Share of each of the professional role categories in the total worked hours of all pharmacists	Pharmacists spent 38% of their time offering pharmaceutical care, followed by 36% performing drug distribution activities and 19% handling management and professional matters.
Profess	ional role: pharmaceutical care	
A1	Hours worked in pharmaceutical care by volume of clientele	Acute care inpatients received an average of 1 hour of care during their hospital stay. Emergency department clients received only 4 minutes per visit on average. Long- term care residents received an average of 3 hours of pharmaceutical care per year. Hemato-oncology patients received 21 minutes of care per visit. ^d
A3	Number of individual counselling sessions offered by pharmacists by volume of clientele	Very few individual counselling sessions were offered. Hemato-oncology patients received on average 9 counselling sessions over 100 chemotherapy visits. Only 7% of acute care patients received a counselling session during their stay, and 3% of long-term care residents received one such session over a year.
A7	Share of each of the care sectors in the total worked hours of all pharmacists	62% of pharmaceutical care hours were offered to acute care inpatients, of which general adult medicine and surgery patients represented 63% (i.e., 39% of all pharmaceutical care hours), followed by intensive care (19% [i.e., 12% of all pharmaceutical care hours]). Long-term care residents accounted for only 8% of all pharmaceutical care hours.
Q1	Number of medication reconciliations during general admissions and hemato-oncology outpatient visits carried out or validated by pharmacists by volume of clientele	On average, only 17 medication reconciliations were done per 100 acute care admissions. This rate increased to 24 per 100 admissions for patients 75 years of age and older. Variability among long-term care residents was substantial, ranging from 0 to 91 medication reconciliations per 100 admissions.
E5	Number of drug therapy problems resolved by pharmacists assigned to direct patient care by volume of clientele	On average, 63 drug therapy problems were resolved per 100 admissions to acute care, and 139 per 100 beds over a year in long-term care. On average, 11 problems were resolved per 100 outpatient chemotherapy visits, and 3 problems per 100 emergency department visits.
E7	Share of drug therapy problems resolved by pharmacists assigned to either pharmaceutical care or drug distribution	The majority (72%) of drug therapy problems were resolved by pharmacists assigned to pharmaceutical care. The remaining proportion of such problems (28%) were resolved by pharmacists assigned to drug distribution.
E8	Number of care activities per day for pharmacists assigned to pharmaceutical care	On average, a pharmacist carried out 12.7 care interventions per working day, most of which (9.6) involved resolving drug therapy problems.

TABLE 3 (part 2 of 2). Selected Results from Testing Indicators during the Pilot Project

Code ^a	Indicator Definition	Selected Results ^b		
Profess	ional role: drug distribution			
Q3	Number of medication errors by volume of clientele	The average error rate was higher for acute care (7.7 errors per 1000 patient-days) than for long-term care (2.0 errors per 1000 patient-days). The error rate was also higher for patients seen in emergency departments (5.1 errors per 1000 visits) than for those seen in hemato-oncology visits (1.8 errors per 1000 visits).		
Q10	Share of incident reports for medication errors that are attributed to pharmacy	The average proportion of errors attributed to pharmacy was 3.8%.		
E1	Ratio of the number of sterile drug preparations to pharmacy technician assistants' worked hours	On average, pharmacy technician assistants made 3.5 sterile hazardous drug preparations per hour and 8.3 sterile nonhazardous drug preparations per hour.		
E2	Ratio of pharmacy technicians' worked hours to pharmacists' worked hours	The ratio of pharmacy technician assistants' worked hours to pharmacists' worked hours was 1.46. The ratio varied from 1.0 to 2.0 across the pilot sites.		
E3	Average processing time for urgent or priority prescriptions	The average processing time for urgent prescriptions was 36 minutes, whereas the average processing time for priority prescriptions was 59 minutes.		
Professional roles: research and education of trainees and colleagues				
11	Proportion of pharmacists who participated in outreach activities	Over the course of a year, 29% of pharmacists wrote an article (16%), presented a poster (13%), and/or gave a talk at a conference (18%). Pharmacists from academic pilot sites reported a substantially higher rate of research activities.		
Professional role: management and professional matters				
Q4	Proportion of pharmacists with an advanced pharmacotherapy master's degree	On average (across all pilot sites), 83% of pharmacists had an advanced pharmacotherapy master's degree. One pilot site reached 100%.		
12	Proportion of the hospital's clinical governance bodies on which a pharmacist sits ^e	Only 2 of the 6 pilot sites reached the preset target of 100%. The average was 89% across all the pilot sites.		

^aThe codes were assigned using the first letter of the framework dimension to which the indicator belonged, followed by a consecutive number. A = appropriateness, E = efficiency, I = innovation and continuous improvement, Q = quality and safety.

^bUnderestimation is assumed for indicators requiring manual data collection, as nonrespondents performed activities that were not counted.

^cAll 5 roles: pharmaceutical care, drug distribution, education of trainees and colleagues, research, and management and professional matters. ^dFor this indicator, an adjustment was made to account for under-reporting, identifying discrepancies between the hours documented by pharmacists via the web form and the hours allocated by the pharmacy departments, and adjusting the result proportionally.

^eBoard of Directors; Council of Physicians, Dentists and Pharmacists Executive Committee; Pharmacy and Therapeutics Committee; Credential Review Committee; Medical, Dental and Pharmaceutical Evaluation Committee; Infection Prevention and Control Committee; and Risk and Quality Management Committee.

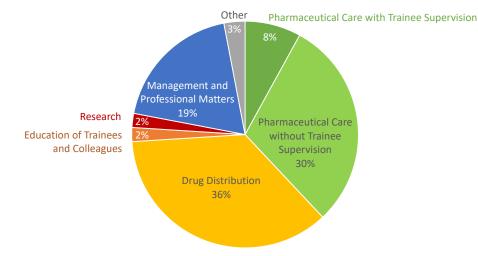


FIGURE 2. Average distribution of hours worked by pharmacists according to professional roles. These results cover a 10-month period across all 6 pilot sites. The breakdown of hours of care with and without supervision of trainees was added during phase 3 to obtain a broader picture of teaching activities, beyond the hours worked in association with the education role and to reflect the additional workload generated by hosting students.

from front-line pharmacists. Figure 2 shows an example of results for the indicator reporting the distribution of hours worked by pharmacists in each professional role (indicator A2). Time devoted to pharmaceutical care ranged from 30% to 55% across pilot sites, which could reflect available human resources rather than a focus on certain activities by administrators.

Indicators calculated as a ratio of manually collated data to data from facilities' information systems are subject to under-reporting bias. To test and illustrate a method for adjusting the calculation to reduce the impact of this bias, a corrective calculation was applied to the indicator reporting the amount of time devoted to pharmaceutical care per volume of clientele (indicator A1). This correction relied on identifying discrepancies between the hours documented by pharmacists via the web-based form and the hours allocated by the respective pharmacy departments. For instance, if, in a specific facility and care sector, the recorded hours for pharmaceutical care within the indicators project represented only 60% of the allocated hours, the indicator result was adjusted proportionally to account for the under-reporting.

The feedback survey was sent to 337 pharmacists, of whom 183 submitted responses, yielding a 54% response rate. Most respondents (94%, n = 172) stated their belief that it is important to collect data to document pharmacists' activities, and 82% (n = 150) saw the project as relevant to the upcoming activity-based funding system. Almost all (97%, n = 177) took part in daily data collection, but some (28%, n = 51) admitted to dropping out during the project. Many (65%, n = 119) mentioned that nonparticipation on certain days was due to omission or oversight. For those who dropped out or did not participate (n = 51), three-quarters (n = 38) said the data collected did not represent their clinical practice, and one-third (n = 17) cited excessive workload.

The primary barriers to participation in data collection were the effort and time required, along with difficulties related to definitions and data compilation, as reported by 65% (n = 119) and 59% (n = 108) of respondents, respectively. Among pharmacists who undertook daily data collection, the most challenging aspect was reported to be tracking the number of pharmacotherapeutic problems resolved (68%, n = 86/126). Respondents noted it was difficult to remember all of the day's activities for proper reporting.

The majority of survey participants reported that they knew about the available tools (84% [n = 154] to 98% [n = 179], depending on the tool), and a good number used them (56% [n = 102] to 80% [n = 146], depending on the tool). The most popular tool was the weekly reminder to access the form (80%, n = 146). The most helpful communication methods were meetings and face-to-face interactions with facility managers (64%, n = 117) and the steering committee (57%, n = 104). Webcasts, promotional emails (excluding

reminders), and an interactive slideshow explaining data collection were useful for 45% (n = 82) of respondents. A promotional video on the pilot project was the least useful (17%, n = 31).

In general, 55% (n = 101) of the 183 survey respondents were satisfied with the pilot project, while 39% (n = 71) were dissatisfied. Better communication of results and more frequent project updates were the top suggestions for increased engagement (35% [n = 64] and 26% [n = 47], respectively). Only 7% (n = 13) would have increased their participation if their department head had made it mandatory.

DISCUSSION

Despite its challenges, the pilot project allowed for significant progress toward a consensus on the most relevant indicators and their definitions for pharmacy departments in Quebec health care systems. For many of the selected indicators, data on their frequency of occurrence or extent of application across all facilities had never previously been collected. The phased deployment of multiple indicators across various pilot sites proved effective. The project's large scale allowed the collection of substantial data over 3 years, involving approximately 20% of the provincial workforce. Also, the project generated interest among province-level stakeholders and hospital pharmacists, reinforcing the importance of documenting the added value associated with pharmacists' activities.

Although many studies have detailed the careful selection of performance indicators for hospital pharmacy activities, few have demonstrated the practicality of implementing such a model on a large scale. Lo and others¹⁵ reported on 5 different experiences in Canadian hospitals where the renowned cpKPIs were implemented to various extents. These authors proposed solutions to the barriers identified, focusing mainly on simplifying processes, increasing the use of automation, and enhancing transparency for stakeholders. Our project's developers believed in the usefulness of cpKPIs, and half of the 8 Canadian cpKPIs were tested (indicators A3, Q1, Q2, and E5). However, these required manual data collection from front-line pharmacists, which contributed to the perceived significant added workload and undue burden.

One of the main challenges of this pilot project was getting front-line pharmacists to participate diligently in daily data collection. Compulsory participation, had it been declared, could have improved the validity and acceptability of adopting the proposed performance indicators. However, due to the external nature of the steering committee, it could not impose data collection, so pharmacists' participation remained voluntary. Nonetheless, it is noteworthy that the majority of pharmacists persisted in their participation, despite the long data collection period. However, encouraging participation beyond a certain level proved difficult. The feedback survey revealed the challenge of changing organizational culture, with the main reason for nonparticipation being the difficulty of integrating the new habit of completing the data collection form into pharmacists' daily activities.

The culture of performance measurement is a challenge in itself, which may be rooted in negative perceptions. Frontline pharmacists who are mandated to collect data routinely may perceive that their time would be better invested in direct patient care. In their conclusions following focus group discussions to explore pharmacists' perceptions of the barriers to and facilitators of cpKPI implementation, Minard and others¹⁶ found that, despite facing challenges, frontline pharmacists generally supported cpKPI measurement. Another group surveying Canadian hospital pharmacists found that involvement in cpKPI activities was positively correlated with overall job satisfaction.³¹ Similarly, our project achieved some success in changing negative perceptions: 94% of pilot site pharmacists surveyed at the project's end recognized the importance of collecting data to document their activities.

In a foundational paper published in 1978, Donabedian divided quality indicators into 3 categories: structure, process, and outcome.³² All but one of the Canadian cpKPIs are process indicators. These are most valuable when there is strong evidence associating processes with clinically meaningful outcomes.¹⁵ Although most of the indicators selected in our exercise were of a structure or process nature, many belonged to the outcome category (specifically indicators E3, E5, E7, Q3, and Q10). Outcome indicators are essential because concrete results such as costs or medication errors help the public and stakeholders understand the impacts of clinical pharmacists. Conversely, the analysis of outcome indicators is inherently flawed, because outcomes are most often influenced by multiple factors that may not depend entirely on pharmacists' activities.

Amid a movement toward privatization of the Saudi Arabian health care system, Al-Jazairi and Alnakhli³³ tested a series of 18 indicators similar to ours over 1 year in a single tertiary care hospital. Instead of daily data collection, they asked pharmacists to collect data on a monthly basis, which may be more acceptable in the long run. Indeed, their participation rate reached 95%, and they were able to show the value of clinical pharmacists and associated cost savings. As they note, a health care institution must then benchmark the collected indicators against national or international indicators to determine where their services stand when compared with other institutions. Attempting to do so, our project highlighted another significant challenge of the benchmarking exercise, that is, the need to validate the data and account for contextual factors across various sites. This requirement presents a potential hurdle to large-scale deployment due to the meticulous analysis and understanding of the field's reality that is required.

CONCLUSION

This project has laid the foundation for demonstrating the value of pharmacy activities along the patient care trajectory and measuring the performance of individual pharmacy departments. Although the tools developed for the pilot project worked well within the project's resource limits, they may not be suitable for province-wide use. This caveat is especially true of the requirement for daily data collection by pharmacists and the technologies used for data collection and processing. Scaling up and deploying certain indicators at the provincial level will require more advanced technological solutions than those used in the pilot project, minimizing or even eliminating manual data collection, along with centralized measurements for system-wide consistency. Achieving large-scale deployment that aligns with government requirements for system development and financing will require support from public authorities.

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