Feasibility of Large-Scale Implementation of an Electronic Patient-Reported Outcome Remote Monitoring System for Patients on Active Treatment at a Community Cancer Center

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QUESTION ASKED: Can a community oncology practice successfully enroll, engage, retain, and manage patients with cancer undergoing intravenous cancer therapy using electronic patient-reported outcome (ePRO)-based digital symptom monitoring? Would patients agree to participate and continue to report using the tool? Why did patients discontinue use? What proportion of reports generated alerts and contained severe symptoms? How were these alerts managed by nursing, ie, telephone consultations or urgent office visits?

SUMMARY ANSWER: This single-center experience indicates that an ePRO-based digital symptom monitoring platform can be effectively implemented at a large scale with a high level of long-term patient enrollment, engagement, and retention. Most reports could be effectively resolved by nurse specialists, and physician intervention was infrequently required.

WHAT WE DID: Clinical implementation of a proprietary digital symptom monitoring ePRO platform was undertaken by the Highlands Oncology Group, a 22-physician community oncology practice in the state of Arkansas. Rollout was initiated in June 2020. Patients receiving parenteral therapy were invited to enroll in the digital symptom monitoring system using either the application or an interactive voice response interface. All patients received in-person training using the reporting modality of their choice. Patients submitted reports on a schedule defined by the practice on the basis of disease, treatment regimen, and clinical factors. Symptoms reported that exceeded a severity threshold were graded by the practice as standard or severe. These reports were monitored in a dashboard managed by nursing staff who determined best management for the patient including telephone intervention or urgent office visit. Patients in this study were enrolled any time before December 1, 2021, and follow-up was through February 28, 2022.

WHAT WE FOUND: Over an approximately 17-month period, 923 patients were successfully enrolled. Retention rates at 3, 6, 9, and 12 months were 94%, 88%, 73%, and 67%, respectively. Few patients discontinued use for reasons related to the platform (n = 47; 5%). Of the 25,311 ePRO reports submitted, 49% (n = 12,334) exceeded the predefined alert thresholds and 8% (n = 1,920) included severe symptoms. The nursing team responded within 24 hours by telephone to 15.5% (n = 3,910) of all reports. All phone calls were in response to reports that exceeded the severity threshold (31.7% of reports), and 72.7% (n = 1,395) of reports that had severe symptoms received a call. Only 6.4% (n = 249) of phone calls required an office evaluation within 72 hours of the report.

BIAS, CONFOUNDING FACTORS: The experience in a single site limits the generalizability of these results, and this warrants reproduction in other centers with different clinical workflows as well as patient populations differing in age, ethnicity, case distributions, and treatments. Specific implementation strategies were not tested. We acknowledge the potential for selection bias between users and nonusers of the ePRO system. Finally, it is difficult to speculate how the pandemic influenced enrollment, engagement, and management of patient reports.

REAL-LIFE IMPLICATIONS: The use of digital symptom monitoring with patient-reported outcomes has been shown to improve patient outcomes. The evidence of benefit has been largely derived from research studies in the academic setting recruiting only a small number of patients. We report that it is feasible to adopt this technology in the community oncology setting at a large scale. Consideration of inclusion of ePROs in alternative payment models is reasonable. The impact on health outcomes remains to be determined.

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PURPOSE The use of digital symptom monitoring with patient-reported outcomes (PROs) has been shown to improve patient outcomes. The evidence of benefit has been largely derived from research studies. The feasibility of adopting this technology in the real-world setting is unknown.

METHODS We report on the clinical implementation of a proprietary electronic patient-reported outcome (ePRO)-based digital symptom monitoring platform at the Highlands Oncology Group practice, a large community oncology practice. We present here our experience with patient enrollment, engagement, and retention; reasons for discontinued use; proportion of reports generating alerts and containing severe symptoms; and the responses to alerts including nursing telephone consultations and urgent office visits.

RESULTS Over an approximately 17-month period, 923 patients were successfully enrolled. Patients enrolled from June 20, 2020, through November 30, 2021, with follow-up through February 28, 2022. Retention rates at 3, 6, 9, and 12 months were 94%, 88%, 73%, and 67%, respectively, with greater retention at 12 months in patients age 65 years or older. Few patients discontinued use for reasons related to the platform (n = 47; 5%). Of the 25,311 ePRO reports submitted, 49% (n = 12,334) exceeded the predefined alert thresholds and 8% (n = 1,920) included severe symptoms. The nursing team responded within 24 hours by telephone to 31.2% (n = 3,910) of all reports with alerts. Of reports with severe symptoms, 72.7% (n = 1,395) received a call. Only 6.4% (n = 249) of phone calls required an office evaluation within 72 hours of the report.

CONCLUSION This single-center experience indicates that an ePRO-based digital symptom monitoring platform can be effectively implemented at a large scale with a high level of long-term patient engagement. Most reports could be effectively resolved by nurses, and physician intervention was infrequently required.

INTRODUCTION Quality care for patients with cancer requires open lines of communication between patients and their care teams. Real-time patient reporting of issues and concerns and timely clinician responses are necessary to improve patient outcomes. Delivering this real-time patient engagement presents a major organizational challenge, and lends itself to a technology-based solution.

The use of digital symptom monitoring with patient-reported outcomes (PROs), often referred to as electronic PROs (ePROs), has been shown to improve patient satisfaction and quality of life, increase time on treatment, reduce ER/hospital utilization, and prolong overall survival.1-8

Despite these many advantages, and multiple platforms that have been developed and modeled,9-16 the evidence of benefit has been largely derived from research studies that recruited only a small number of patients3,5,8 or larger studies with only short follow-up.15

The seminal publication in this area by Basch et al3 randomly assigned 766 patients at Memorial Sloan Kettering to symptom monitoring intervention or usual care. These patients were enrolled over 4 years. Patient diagnoses were limited to genitourinary (32%), gynecologic (22%), breast (20%), and lung (25%) cancers. The extent of patient reporting was not described, although median time on study was only 3.7 months. Although the primary end points of this landmark study were met, clearly establishing the value of remote
monitoring, the ability to scale this approach more widely and rapidly cannot be assessed by this publication.

Basch has subsequently launched a multicenter trial of a digital monitoring system in the community setting, the PRO-TECT trial. The trial was opened in 2017 in 50 sites with an accrual goal over 4 years of 1,100 patients. The trial randomized 52 community practices by site to active patient engagement or usual care with each single site limited to 50 patients. As reported in an ASCO plenary abstract, 1,191 patients were accrued in these 50 sites, and 593 patients were randomly assigned to the active patient management arm. The study reported patient and clinician satisfaction, physical function, symptom burden, and health-related quality of life, all of which were improved by the active intervention. Patient engagement remained high during the study period. These encouraging preliminary results confirm the findings Basch reported previously. However, the generalizability of this experience is diminished by the extremely modest accrual per site over the time period of the study.

The largest experience in the community has been reported by Patt et al with Texas Oncology. In the report, 73% (1,841) of enrolled patients (there were 4,375 patients approached and 2,522 patients enrolled) submitted an ePRO report over a 6-month period. Although patient engagement was initially high, it fell from 72% to 54%, with the overall follow-up time very limited; at the 3-month point, only 10% of enrolled patients had been on the platform long enough to be assessed for continued engagement. The clinical implications of these patient reports were not described. Although this report does establish the feasibility of a large-scale launch in the community, the extremely short follow-up and the lack of details regarding the impact of the data reporting on patient outcomes make it unclear whether the deployment was in fact successful.

The adoption of digital monitoring systems into routine clinical practice remains limited. The feasibility of more widespread adaptation of this approach must start with high patient enrollment, patient engagement in submitting scheduled and as-needed reports, and a high level of patient retention over time. We hypothesized that with workflow and electronic medical record integration, appropriate training, and a user-friendly patient interface, a digital symptom monitoring ePRO platform could be effectively executed at a large scale in the community oncology care setting. We report here the preliminary experience of Highlands Oncology Group, a large community oncology practice with a proprietary ePRO-based digital symptom monitoring platform.

**METHODS**

**Design**

This study reports the preliminary experience of implementing a proprietary ePRO-based digital symptom monitoring platform in a large community practice. The primary aim was to evaluate feasibility using criteria of patient enrollment, engagement, and retention, and the reasons for discontinued use. The secondary aim was to evaluate the proportion of reports generating alerts and the subgroup of alerts including reports of severe symptoms; and the responses to these reports including nursing telephone consultations and urgent office visits.

**Setting**

Highlands Oncology Group is a community oncology practice in the state of Arkansas consisting of 22 physicians and 28 advanced practice providers. The practice has four sites of service. They participated in the Oncology Care Model and use a centralized nurse triage model.

**Patient Population**

All adult patients with cancer in each participating treatment site starting parenteral therapy were invited to use the ePRO system at the time of their pretreatment education program.

**Intervention**

The Canopy Care ePRO-based digital symptom monitoring platform is a proprietary modular, cloud-based communication system that incorporates integrated interfaces for patients and clinical staff.

Patients can submit scheduled and unscheduled reports of their distress and specific symptoms using either a symptom reporting application, which can be used on a smartphone or tablet, or an interactive voice response (IVR) interface (telephone report). The symptom monitoring application incorporates a 10-point linear analog patient distress scale and a comprehensive problem list that includes physical symptoms as well as psychosocial issues reported using a verbal rating scale. The symptoms list in the application was modeled after several validated symptom evaluation scales including the National Comprehensive Cancer Network distress thermometer, the Memorial Symptom Assessment Scale, and the Edmonton Symptom Assessment Score. Patients rated each problem on a four-point rating scale (mild, moderate, severe, and worst possible). The IVR interface facilitated reporting of overall patient well-being and a more focused list of relevant symptoms.

At the time of enrollment, a reporting schedule was defined for each individual patient on the basis of the diagnosis, treatment, and comorbidities. Most frequently, the schedule was weekly. In addition patients were instructed to submit unscheduled reports on an as-needed basis in the event of worsening symptoms or deterioration in general well-being.

A clinician-facing web-based interface incorporates a continuously updated work queue of patients’ reports generated by the digital symptom monitoring. Symptom-specific thresholds were defined by the practice for each
TABLE 1. Patient Characteristics

| Characteristic | Total (N = 923), No. (%) | Female (n = 562), No. (%) | Male (n = 360), No. (%) |
|----------------|--------------------------|---------------------------|-------------------------|
| Tumor type     |                          |                           |                         |
| Breast         | 184 (19.9)               | 182 (32.4)                | 1 (0.3)                 |
| Hematologic    | 182 (19.7)               | 87 (15.5)                 | 95 (26.4)               |
| GI             | 161 (17.4)               | 70 (12.5)                 | 91 (25.3)               |
| Thoracic       | 157 (17.0)               | 79 (14.1)                 | 78 (21.7)               |
| Genitourinary  | 145 (15.7)               | 95 (16.9)                 | 50 (13.9)               |
| Other          | 30 (3.3)                 | 14 (2.5)                  | 16 (4.4)                |
| Melanoma       | 26 (2.8)                 | 14 (2.5)                  | 12 (3.3)                |
| Unknown        | 20 (2.2)                 | 16 (2.8)                  | 4 (1.1)                 |
| Head and neck  | 15 (1.6)                 | 4 (0.7)                   | 11 (3.1)                |
| Brain          | 3 (0.3)                  | 1 (0.2)                   | 2 (0.6)                 |
| Age, years     |                          |                           |                         |
| < 20           | 2 (0.2)                  | 0 (0.0)                   | 2 (0.6)                 |
| 20-29          | 14 (1.5)                 | 9 (1.6)                   | 5 (1.4)                 |
| 30-39          | 49 (5.3)                 | 40 (7.1)                  | 9 (2.5)                 |
| 40-49          | 100 (10.8)               | 71 (12.6)                 | 29 (8.1)                |
| 50-59          | 192 (20.8)               | 124 (21.1)                | 68 (18.9)               |
| 60-69          | 289 (31.3)               | 168 (29.9)                | 121 (33.6)              |
| 70-79          | 211 (22.9)               | 119 (21.2)                | 92 (25.6)               |
| ≥ 80           | 64 (6.9)                 | 31 (5.5)                  | 33 (9.2)                |
| Missing age    | 2 (0.2)                  | 0 (0.0)                   | 1 (0.3)                 |
| Race/ethnicity |                          |                           |                         |
| Non-Hispanic White | 822 (89.1)          | 499 (88.8)                | 323 (89.7)              |
| Non-Hispanic other | 50 (5.4)            | 36 (6.4)                  | 14 (3.9)                |
| Unknown/missing | 26 (2.8)               | 10 (1.8)                  | 15 (4.2)                |
| Hispanic or Latino | 14 (1.5)            | 7 (1.2)                   | 7 (1.9)                 |
| Non-Hispanic Black | 11 (1.2)            | 10 (1.8)                  | 1 (0.3)                 |
| Metastatic status |                          |                           |                         |
| Metastatic     | 520 (56.3)               | 299 (53.2)                | 221 (61.4)              |
| Not metastatic | 403 (43.7)               | 263 (46.8)                | 139 (38.6)              |

NOTE. One patient was missing documentation of sex in the electronic medical record; hence, the number of male and female patients does not add up to the overall number.

Demographic variables included age at the time of enrollment (analyzed as a continuous variable, as 10-year categories, and as a binary variable [<65 v ≥65 years]), sex (male v female), and race/ethnicity (non-Hispanic White, non-Hispanic Black, Hispanic or Latino, and other). Patients were categorized by primary tumor type using the International Classification of Disease, Ninth and Tenth Revisions, Clinical Modification (ICD-9-CM and ICD-10-CM) codes. Tumor types were placed into the following broad categories: GI, thoracic, genitourinary, breast, hematologic, melanoma, head and neck, brain, other, and unknown. A patient was identified as having metastatic disease if there was a record of any ICD code containing secondary and neoplasm that occurred on or before the first relevant structured administration.

Computer-guided triage pathways for structured telephone evaluation by oncology nurses were developed to triage patients who can be managed with a telephone consultation alone, and those who require expedited or urgent in-person evaluation. On the basis of individual patient circumstances and nursing judgment, not all alerts exceeding the threshold resulted in a telephone consultation or an urgent visit.

The practice rollout was initiated in June 2020. Rollout occurred sequentially across practice sites. Patients in this study were enrolled any time before December 1, 2021, and follow-up was through February 28, 2022.

Data Collection

Demographic and clinical variables were obtained from structured electronic health record (EHR) data. ePRO-specific data (eg, date of enrollment, date and reason for report, alerts/symptoms, and reason for opt-out) were collected through the ePRO platform and linked to patient-level EHR data.

Data Evaluation and Statistics

In patients enrolled on the ePRO, descriptive statistics (absolute and relative frequencies) were reported for categorical demographic variables overall and by sex. Crude reporting frequency was calculated for each patient as the total number of reports per time (in months) from enrollment to last report; and was categorized as <1, 1 to <2, 2 to <3, 3 to <4, or >4 reports per month. The distribution...
of reporting frequency was reported overall and by age (< 65 v ≥ 65 years).

Time to last report was calculated as the time from date of enrollment to date of last report. A Kaplan-Meier approach was used to calculate retention. Analyses were indexed to the date of enrollment. An event was defined as the date of last report in patients who did not opt out of Canopy and who had ≥ 45 days between their last report and last structured visit date; or who had an opt-out reason that was related to the platform. Patients were censored at the last report date if they had not opted out but had < 45 days between the last report and last structured visit date; or if they opted out for reasons not related to the Canopy platform. Patients with follow-up after the study period end date were administratively censored on February 28, 2022. Kaplan-Meier curves were generated for all patients enrolled on Canopy, overall and stratified by age (< 65 v ≥ 65 years). The log-rank test was used to compare the retention distribution by age. Since the median retention was not reached, retention probabilities were calculated at 3, 6, 9, and 12 months after enrollment.

The number and proportion of reports that generated any alert (standard or severe) or an alert because of at least one severe symptom was calculated. Reports that resulted in at least one telephone evaluation within 24 hours of the report date/time were identified; and among those reports that generated an alert and resulted in a telephone evaluation, the distribution of alert-generating symptoms was reported. Additionally, the number and proportion of reports followed by at least one office evaluation within 72 hours of the report date was calculated.

Analyses were conducted using R statistical computing software, version 4.0.2.19

### RESULTS

#### Patient Population and Enrollment

Between June 22, 2020, and November 30, 2021, 3,072 patients were treated with parenteral anticancer therapies at Highlands. Since the program was rolled out sequentially across the different practice sites, not all treated patients were eligible to enroll at the start of the measurement period. Overall, 1,173 eligible patients were invited to enroll into the digital symptom monitoring system and 923 (79%) were successfully enrolled with multiple reports. The most common reason patients chose not to enroll was lack of interest (60%). Of these 923 patients, 296 discontinued at some point after enrollment. The most common reasons for discontinuation were unrelated to the ePRO platform: death and discharge to hospice (47% and 18% of opt-out reasons, respectively).

**TABLE 2.** Reports per Month Among Successfully Enrolled Patients

| Reporting Frequency | Total (N = 923), No. (%) | < 65 Years (n = 499), No. (%) | ≥ 65 Years (n = 422), No. (%) |
|---------------------|--------------------------|-----------------------------|-----------------------------|
| < 1 report per month| 168 (18.2)               | 91 (18.2)                   | 77 (18.2)                   |
| 1 to < 2 reports per month | 158 (17.1) | 88 (17.6) | 70 (16.6) |
| 2 to < 3 reports per month | 155 (16.8) | 84 (16.8) | 69 (16.4) |
| 3 to < 4 reports per month | 223 (24.2) | 140 (28.1) | 83 (19.7) |
| ≥ 4 reports per month | 219 (23.7) | 96 (19.2) | 123 (29.1) |

NOTE. Two patients were missing documentation of date of birth/age in the electronic medical record; hence, the number of patients in the age categories does not add up to the overall number.

**FIG 1.** Kaplan-Meier estimates of time to last Canopy report: (A) retention probability of all patients and (B) retention probability stratified by age (< 65 v ≥ 65 years). Two patients had a missing value for age and were not included in the analysis stratified by age.
Very few patients (approximately 5%) discontinued use of the monitoring system for reasons related to the platform: not interested (n = 31), bothersome (n = 10), too complicated (n = 4), too stressful (n = 1), and does not own a phone (n = 1).

Enrolled patients varied across tumor type (solid and hematologic), age, and metastatic status (Table 1), and included patients on active treatment at the time of implementation as well as new treatment starts over the approximately 17-month period. Most patients elected to use the application (794; 86%), whereas 71 (7.7%) elected to use the IVR interface and for 58 (6.3%), the preferred interface was not documented.

### Reporting and Retention

The median duration of follow-up was 9 months. The average number of reports submitted by patients per month is presented in Table 2. Most patients (64.7%) submitted two or more reports per month. The patient retention at 3, 6, 9, and 12 months was 94%, 88%, 73%, and 67%, respectively (Fig 1). We observed a statistically significant difference in retention between patients age older or younger than 65 years, with patients age 65 years or older more likely to be continuously reporting at 12 months (73% v 62%, P = .0011).

### Management of ePRO Reports

During the study period, 25,311 ePRO reports were submitted. Of the 25,311 ePRO reports submitted, 49% (n = 12,334) exceeded the predefined alert thresholds and 8% (n = 1,920) included severe symptoms. The nursing team responded within 24 hours by telephone to 31.2% (n = 3,910) of all reports with alerts. Of reports with severe symptoms, 72.7% (n = 1,395) received a call. Overall, 15.5% of all reports generated a phone call. In reports that triggered nursing team/staff telephone calls within 24 hours, the most common alert-generating symptoms were high levels of distress, pain, nausea, weakness, diarrhea, dyspnea, and fatigue (Table 3).

| Alert-Generating Symptom | No. | % of Alerts | Alert-Generating Symptom | No. | % of Alerts |
|--------------------------|-----|-------------|--------------------------|-----|-------------|
| High distress level      | 2,930 | 74.9        | Indigestion              | 220 | 5.6         |
| Pain                     | 1,968 | 50.3        | Swelling                 | 209 | 5.3         |
| Weakness/fatigue         | 1,416 | 36.2        | Cough                    | 208 | 5.3         |
| Nausea                   | 1,326 | 33.9        | Loss of interest in usual activities | 197 | 5.0         |
| Difficulty breathing     | 955   | 24.4        | Sadness                  | 174 | 4.5         |
| Diarrhea                 | 922   | 23.6        | Fever under 100.4°F      | 159 | 4.1         |
| Chest pain               | 640   | 16.4        | Tingling in hands or feet | 157 | 4.0         |
| Fatigue                  | 632   | 16.2        | Nervousness              | 156 | 4.0         |
| Eating/appetite          | 525   | 13.4        | Bathing/dressing         | 148 | 3.8         |
| Sleep                    | 505   | 12.9        | Nausea/vomiting          | 143 | 3.7         |
| Breathing                | 498   | 12.7        | Fever over 100.4°F       | 129 | 3.3         |
| Getting around           | 495   | 12.7        | Fears                    | 123 | 3.1         |
| Anxiety                  | 363   | 9.3         | Insurance/financial      | 120 | 3.1         |
| Numrness/tingling         | 347   | 8.9         | Urination problems       | 118 | 3.0         |
| Memory/concentration     | 346   | 8.8         | Mouth/throat sores       | 116 | 3.0         |
| Vomiting                 | 335   | 8.6         | Other                    | 108 | 2.8         |
| Constipation             | 301   | 7.7         | Feeling swollen           | 106 | 2.7         |
| Depression               | 250   | 6.4         | Hot flashes              | 105 | 2.7         |
| Worry                    | 239   | 6.1         | Rash                     | 94  | 2.4         |
| Nose dry or congested    | 238   | 6.1         | Skin dry or itchy        | 94  | 2.4         |
| Dizziness/lightheadedness| 237   | 6.1         | Cold sensitivity          | 85  | 2.2         |
| Falls/balance            | 236   | 6.0         | Skin/nail problems       | 83  | 2.1         |
| Headache                 | 231   | 5.9         | Visual                   | 82  | 2.1         |
| Dry mouth                | 220   | 5.6         | Physical appearance      | 80  | 2.0         |

NOTE. In the above table, we have only included symptoms that were reported in 2% or more of alerts. May be more than one symptom per call.

3,661 (93.6%) of symptom reports leading to a phone call were managed over the telephone by the nurse. The remaining 249 (6.4%) required an office evaluation within 72 hours of the report.
TABLE 4. Organizational, Clinician, Patient, and Implementation-Related Barriers to ePRO Uptake

| Organization barriers                                                                 |    |
|---------------------------------------------------------------------------------------|--|
| 1. Integrating the system into the clinic workflow                                      |    |
| a. Minimize disruption                                                                 |    |
| b. Optimal use of administrative, nursing, and physician resources                    |    |
| 2. Acceptability                                                                       |    |
| a. Clinicians                                                                         |    |
| b. Patients                                                                           |    |
| 3. Up-front investment                                                                 |    |
| 4. Technical maintenance                                                               |    |
| Implementation barriers                                                                |    |
| 1. Necessity for comprehensive and role-specific training for all parties (clinicians, staff, and patients) using the ePRO system |    |
| 2. Inadequate clinician response to patient reports, eg, to address critical PRO results |    |
| Clinician’s barriers                                                                   |    |
| 1. Information overload                                                                |    |
| a. Resistance to generation of extra workload, potential to lengthen clinical visits   |    |
| b. Fear of identification of problems that clinicians may feel unqualified to address |    |
| 2. Added time-sensitive clinical responsibility                                        |    |
| Patient barriers                                                                       |    |
| 1. Excessively burdensome ePRO collection software                                      |    |
| a. Too many questions                                                                  |    |
| b. Too much complexity                                                                 |    |
| c. Difficult log-in requirements                                                       |    |
| 2. Difficulty with technology                                                          |    |
| a. Literacy and language                                                                |    |
| b. Familiarity with touch-screen computers                                             |    |
| c. Availability of home computers and internet connectivity                            |    |
| d. Impaired manual dexterity                                                           |    |

Abbreviations: ePRO, electronic patient-reported outcome; PRO, patient-reported outcome.

DISCUSSION

Despite the established evidence base that digital symptom monitoring with PROs can improve patient outcomes and the multiplicity of platforms that have been developed and modeled, adoption of this approach in community oncology practices has been limited. Barriers to the more widespread diffusion of this intervention have been described and can be classified as organizational, clinician, patient, and implementation-related barriers (Table 4). The design of the specific proprietary ePRO digital symptom monitoring platform used in this study has been informed by these considerations, with emphasis on ease on the patient use experience, and the development of a clinician interface that is easily integrated into practice workflows and the electronic medical record. This descriptive report does not purport to describe an optimal implementation strategy.

The success reported here compares favorably with published experience. The long-term patient retention in this real-world study was comparable with that reported in the recently presented US national cluster randomized trial. The generalizability of this experience is supported by the observation that the enrolled population of patients had similar demographics in relation to age, sex, and cancer site distribution to patients seen in routine clinical practice. Age and lack of familiarity with technology have been reported as potential barriers to ePRO implementation. In our experience, advanced age was not a barrier to retention. Indeed, we observed a statistically significant difference in patient reporting frequency and retention by age, with patients age 65 years and older more likely to be continuously reporting at 12 months, contrary to expectations. In Basch’s MSKCC trial, one third of patients were computer-inexperienced, and this group benefited at least as much from the symptom monitoring intervention. In our cohort, fewer than 10% chose interactive voice response. Of those using the app-based response, very few patients discontinued the use of the system because of difficulty or burden.

Patients submitted a significant number of reports each month: close to half exceeded a predetermined symptom threshold and 10 percent were classified by the clinical team as severe symptoms. As expected, this resulted in a higher rate of telephone calls for patients with severe alerts (72.7%) than those with standard alerts (30%). Most issues could be effectively resolved by the telephone consultation and only approximately 5% of patients required an urgent office visit. The primary symptoms responsible for these interventions mirror the most common complaints in this treated population, and also mirror the most common reasons for emergency room visits in patients with cancer. Not all alerts generated a telephone call. Clinical judgment was exercised by nurses during the evaluation of patient reports including all reports defined as severe.

Our study has strengths and limitations. The cohort described is large and was enrolled over a short period of time. There was a very low rate of patient drop-out. The patients enrolled are similar in age, sex, and cancer type to those seen in routine clinical care. In addition, the symptoms they report are also qualitatively similar to those seen in routine clinical practice. The experience in a single site allowed for standardization of processes; however, it diminishes the generalizability of these results, and this warrants reproduction of the experience in other centers with different ethnicity and case distributions. We acknowledge the potential for selection bias between users and nonusers of the ePRO system. Finally, it is difficult to evaluate the impact of the COVID-19 pandemic. Like many community oncology practices, Highlands Oncology Group successfully continued treating...
patients during the pandemic. It is difficult to speculate how the pandemic influenced enrollment, engagement, and management of patient reports.

On the basis of these preliminary data, we are pursuing several projects. Further work is needed to determine impact on patient outcomes, and the generalizability of this experience across different treatment settings and different disease and treatment groups.

In conclusion, this single-center experience supports the hypothesis that implementing an ePRO-based digital symptom monitoring platform is feasible in routine clinical practice at a large scale. We provide evidence that patients can remain engaged over a prolonged period of time with regular reporting and low attrition. These reports result in telephonic interventions in a sizable percentage of patients, with only a small proportion requiring office evaluation.
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AUTHORS' DISCLOSURES OF POTENTIAL CONFLICTS OF INTEREST

Feasibility of Large-Scale Implementation of an Electronic Patient-Reported Outcome Remote Monitoring System for Patients on Active Treatment at a Community Cancer Center

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