Electronic Patient Reporting of Adverse Events and Quality of Life: A Prospective Feasibility Study in General Oncology

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QUESTION ASKED: Is it feasible and acceptable, and what level of compliance is achieved when asking oncology patients to regularly self-report both adverse events (Patient-Reported Outcomes version of the Common Terminology Criteria for Adverse Events [PRO-CTCAE]) and health-related quality of life (HRQOL) at home on an electronic system (Internet/telephone option)?

SUMMARY ANSWER: High compliance was observed for weekly patient self-reported PRO-CTCAE plus HRQOL completions every 4 weeks. Patient feedback was positive, highlighting a willingness to complete questionnaires, with the system being easy to use and problem free.

WHAT WE DID: Two hundred forty-nine oncology patients receiving a variety of treatments from 2 United Kingdom hospitals were sent weekly auto-reminders to complete PRO-CTCAE and every 4 weeks, the HRQOL was also available. These self-reported questionnaires were completed online or via a telephone-based interactive voice response system (IVRS) for a 12-week period.

WHAT WE FOUND: Overall compliance was more than 60% at each individual timepoint and the cumulative compliance was 75% to 80% for each questionnaire. Some differences were observed across the treatment groups (chemotherapy higher compliance), patient age (older patients completed the PRO-CTCAE more often), and choice of system (greater HRQOL completion with IVRS). The Internet option was more popular (82.3%) and quicker, but some older patients (16.9%) preferred the IVRS. Most participants remained on the study throughout the 12-week period, with only 13 active withdrawals, 6 deaths, and 18 passive withdrawals (not completing any electronic questionnaires postbaseline).

BIAS, CONFOUNDING FACTORS: This feasibility study was carried out in a general oncology setting in a hypothetical trial scenario, rather than among active clinical trial participants. This influenced how the study was presented and some of the practicalities (baseline completion was defined flexibly rather than stringently as per clinical trial protocols), and potentially affected uptake and compliance. The recruitment rate was relatively low; therefore, there may be biases as a result of self-selection.

REAL-LIFE IMPLICATIONS: The value of routine PRO collection is increasingly recognized in cancer clinical trials, and the need for innovative methods to achieve this remotely have been heightened further during and after the coronavirus disease 2019 pandemic. Our findings highlight the willingness of patients to self-report from home and the acceptability of the electronic system. Additional exploration of this system within active clinical trials is warranted to explore whether implementation can improve the precision and accuracy of adverse event reporting, leading to improved data quality and patient safety.

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abstract

PURPOSE Adverse event (AE) reporting is essential in clinical trials. Clinician interpretation can result in under-reporting; therefore, the value of patient self-reporting has been recognized. The National Cancer Institute has developed a Patient-Reported Outcomes version of the Common Terminology Criteria for Adverse Events (PRO-CTCAE) for direct patient AE reporting. A nonrandomized prospective cohort feasibility study aimed to explore the compliance and acceptability of an electronic (Internet or telephone) system for collecting patient self-reported AEs and quality of life (QOL).

METHODS Oncology patients undergoing treatment (chemotherapy, targeted agents, hormone therapy, radiotherapy, and/or surgery) at 2 hospitals were sent automated weekly reminders to complete PRO-CTCAE once a week and QOL (for a maximum of 12 weeks). Patients had to speak/understand English and have access to the Internet or a touch-tone telephone. Primary outcome was compliance (proportion of expected questionnaires), and recruitment rate, attrition, and patient/staff feedback were also explored.

RESULTS Of 520 patients, 249 consented (47.9%)—mean age was 62 years, 51% were male, and 70% were married—and 230 remained on the study at week 12. PRO-CTCAE was completed at 2,301 (74.9%) of 3,074 timepoints and QOL at 749 (79.1%) of 947 timepoints. Individual weekly/once every 4 weeks compliance reduced over time but was more than 60% throughout. Of 230 patients, 106 (46.1%) completed 13 or more PRO-CTCAE, and 136 (59.1%) of 230 patients completed QOL questionnaires. Most were completed on the Internet (82.3%; mean age, 60.8 years), which was quicker, but older patients preferred the telephone option (mean age, 70.0 years). Positive feedback was received from patients and staff.

CONCLUSION Self-reporting of AEs and QOL using an electronic home-based system is feasible and acceptable. Implementation of this approach in cancer clinical trials may improve the precision and accuracy of AE reporting.

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INTRODUCTION

As more cancer treatments with innovative modes of action are evaluated via clinical trials, the need to monitor and record adverse events (AEs) increases to ensure safety and guide prescribing and patient information. The Common Terminology Criteria for Adverse Events (CTCAE) is the standard method of reporting AEs, allowing clinicians to uniformly classify AEs and their severity, but its focus is on safety rather than exploring patient experiences. Clinician interpretation can result in under-reporting of lower-grade morbidity and severity downgrading; therefore, patient self-reporting is a complementary alternative. The CTCAE system is suitable for acute AEs when patients receive regular oncologist review; however, many new treatments are outpatient-based oral therapies that require less frequent reviews. Although these treatments do not result in the acute severe AEs observed with chemotherapy, they often result in low- to moderate-grade prolonged symptoms that significantly impact patients’ lives and treatment adherence. CTCAE is less appropriate for monitoring AEs during these treatments.

Patient-reported outcomes (PROs) enable systematic, subjective measurement of symptoms and experiences. Health-Related Quality of Life (HRQOL)
questionnaires are well-established PRO measures used in phase II and III randomized clinical trials. Implementing PROs to routinely monitor individual patients’ AEs is relatively uncommon, despite accumulating evidence demonstrating improvement in communication, patient well-being, symptom control, and survival when clinicians receive these data.

The National Cancer Institute developed the Patient-Reported Outcome version of the CTCAE item bank (PRO-CTCAE) to obtain direct patient AEs in trials. Technology can help PRO-CTCAE to become a standardized trial approach, and Web-based PRO efficiency and patient acceptability has been demonstrated. Of the United Kingdom population, 90% are regular Internet users; however, 24% of those age 65 years and older reported no Internet use in the past 3 months. Therefore, a telephone interactive voice response system (IVRS) widens eligibility.

Oncology clinicians, trialists, the pharmaceutical industry, and regulatory authorities recognize the importance of collecting PRO data in drug development and evaluation. Concerns still exist about potential patient burden and data collection frequency. There is a pressing need to develop robust systems to collect PRO-CTCAE alongside clinician-reported CTCAE, as well as explore the feasibility of collecting frequent PRO-CTCAE in addition to HRQOL at prespecified trial timepoints where QOL is an important primary or secondary outcome.

This feasibility study evaluated a self-reported electronic system—Internet/IVRS—to collect PRO-CTCAE and HRQOL in a mixed group of oncology patients and to combine the data with a standard clinical trial database. Primary outcome was compliance and secondary outcomes included recruitment rates, Internet/IVRS uptake, attrition, patient and staff feedback, and system testing.

**METHODS**

**Patient Population and Study Design**

This proof-of-principle, nonrandomized, prospective, cohort feasibility study was conducted in 2 United Kingdom hospitals between August 19, 2014, and October 7, 2015, with 12-week follow-up. The study design modeled a hypothetical clinical trial scenario in which PRO-CTCAE data are collected for safety (real time, as frequently as needed) and HRQOL at predefined timepoints to assess treatment outcomes.

Our aim when defining the study population was to capture a range of patients across the main cancer sites and treatments. We anticipated that this would increase the generalizability of the findings. Consecutive eligible patients were approached. Therefore, the sample target was 210 participants with early or metastatic cancer receiving either chemotherapy, targeted agents, hormone therapy, radiotherapy, or surgery (42 patients per group, allowing for 30% attrition) plus 42 patients with Eastern Cooperative Oncology Group (ECOG) performance status score of 2 to 3 receiving any treatment. Eligibility criteria included speaking/understanding English and access to a touch-tone telephone and/or Internet. Patients on clinical trials with regular HRQOL assessments were excluded. Written informed consent was obtained from study participants. Ethical approval was granted from the National Health Service Leeds-East Research Ethics Committee (14.YH.0181).

**Study Outcomes**

- Compliance was defined as the proportion of completed questionnaires out of expected completions. This was calculated cumulatively for each questionnaire—PRO-CTCAE and HRQOL—and at each expected timepoint weekly (7-day window)/every 4 weeks (28-day window) across gender, system, study group, and age. Multiple completions per week were discounted from compliance calculations. The number of completions per participant was also calculated, including multiple completions.
- Recruitment rate was defined as the proportion of patients who consented to the study relative to eligible patients approached. Demographics—age, gender, diagnosis, and treatment—were compared between consenting patients and declining patients.
- Internet/IVRS uptake was calculated in 2 ways: the proportion who preferred each system at baseline and the proportion who used each system most often during the study.
- Active withdrawal was the predominant attrition outcome, defined as the proportion of participants who actively requested withdrawal from the study out of participants who consented. Any withdrawal reasons were recorded where provided. In addition, passive withdrawal was defined as the proportion of participants who did not complete any electronic questionnaires postbaseline but did not request withdrawal. Passive withdrawals are included in the compliance calculations, but active withdrawals were excluded after the withdrawal date.
- Patient and staff feedback was captured on a paper end-of-study (EOS) questionnaire that explored experiences of the system(s) and staff views.
- System testing was assessed through the documentation of system issues and phone calls and comments from patients and staff. Call and online session time logs were also recorded.

**Electronic System and Questionnaires**

Participant registration via the clinical trials research unit (CTRU) registration system provided the trial number and username and password for accessing the electronic system to complete the 2 self-reported questionnaires.

First, 21 PRO-CTCAE items covering 11 common symptoms (pain and/or discomfort, nausea, vomiting, diarrhea,
constipation, headache, mouth/throat sores, fatigue, skin rash, flu-like symptoms, and muscle ache) were available at any time. Second, the European Organization for the Research and Treatment of Cancer Quality of Life Questionnaire C30 (EORTC-QLQ-C30) 30-item cancer-specific measure exploring HRQOL24 was available once every 4 weeks.

Holch et al23 provide a detailed overview of the Internet (QTool) system. Internet system users were presented with 1 (PRO-CTCAE) or 2 questionnaires (if EORTC-QLQ-C30 was due). The IVRS that was developed specifically for this study by CTRU technicians was a freephone number. To ease the log-in process on the IVRS, all usernames were a 5-digit numerical code, and passwords were a 4-digit numerical code. The same format was used on the Internet system to enable participants to switch between systems if required. Once logged in, IVRS users heard a welcome message, then heard the EORTC-QLQ-C30 questionnaire (if due, every 4 weeks) and/or the PRO-CTCAE. Each question was read out, followed by the response options (eg, “Question number 13. In the last 7 days, what was the SEVERITY of your fatigue, tiredness, or lack of energy at its worst? Press 1 then hash for none, 2 then hash for mild, 3 then hash for moderate…..”) Patients could interrupt the responses with their selected response and could take a break—hang up and re-ring—after the first questionnaire if needed.

A specific application pulled the data from a separate server that held the Internet/QTool data, which was then stored in a specific PRO database on CTRU servers. A program was created to generate/send automated reports to the local research team detailing completions. The PRO database was combined with the standard trial database (Elsevier’s MACRO Data Capture System) that contained all the participant clinical record forms for statistical analysis.

**Procedure**

Eligible patients were initially approached by clinical/research staff with the study information sheet. Recruitment was undertaken sequentially across the treatment subgroups and stopped once each prespecified target number was reached.

Demographic details and computer use (frequency and perceived confidence) were self-reported at baseline. Clinical data captured from medical records included diagnosis, treatment, ECOG performance status, and treatment changes (discontinuation/new treatments). Baseline electronic questionnaires were completed in clinic, or a brief demonstration was provided and the full questionnaires completed at home on the preferred system. All participants received step-by-step instructions covering both systems. Participants were reminded (by automated e-mail, text, or both) weekly—every 7 days from the registration date—for 12 weeks to complete the electronic questionnaires. EOS paper questionnaires were sent/given once each participant had completed the 12 weeks.

Any deaths or withdrawal requests were actioned promptly (reminders stopped). All participants were reminded that the system was not intended to replace usual care or methods of reporting any symptoms or concerns to their clinical team. However, researchers endeavored to attach a paper report summarizing any recent AEs to participants’ medical notes at any scheduled outpatient visits.

**Analysis**

Data analysis was performed using STATA between August 2016 and February 2017. Descriptive analysis of the study outcomes was conducted. Missing data were not imputed. System issues were coded thematically into generic and patient issues.

**RESULTS**

**Study Population**

Of 888 patients who were assessed for eligibility during the study period, 368 were ineligible, and of 520 approached, 249 consented to the study (recruitment rate 47.9%; Fig 1). Consent rates were higher in targeted agents (57 of 93; 61.3%) and chemotherapy groups (49 of 97; 50.5%), and lower in surgery (42 of 91; 46.2%), radiotherapy (47 of 107; 43.9%), and hormone therapy (54 of 132; 40.9%; $x^2 = 10.3; P = .04$).

Participants (mean age, 62.1 years; standard deviation [SD], 13.3 years) were significantly younger than those who declined (mean age, 69.4 years; SD = 12.3 years; $t = −6.5; df = 518; P < .001$). No differences were found by gender ($P = .46$) or disease site ($P = .08$). Of 271 who declined, reasons included being too busy (n = 56; 20.7%), not interested (n = 26; 9.6%), and being too worried/stressed (n = 15; 5.5%). Access to the required technology prevented participation for 21 non-Internet users (who did not want to use IVRS; 7.7%), 15 patients who were hard of hearing (5.5%), and 5 who were unskilled and/or disliked both options (1.8%).

**Table 1** lists the demographic and clinical characteristics of the sample. This shows almost similar numbers of male and female patients (n = 128, 51.4%; and n = 121, 48.6%, respectively), and most were married (n = 173; 69.5%) and 124 were retired (49.8%). Breast (n = 57; 22.9%), colorectal (n = 39; 13.7%), and prostate (n = 33; 13.3%) cancer were the most common cancer types. IVRS participants were older (mean age, 70.0 years; SD, 9.9 years; Internet mean age, 60.8 years; SD, 13.7 years), had lower education levels, and more had prostate cancer (12 of 43; 27.9%) compared with Internet users.

**Internet/IVRS Uptake**

At baseline, 82.3% of participants (205 of 249) indicated they preferred the Internet, whereas 16.9% (42 of 249) preferred the IVRS (1 had no preference and 1 withdrew before registration). Most participants used their preferred system throughout the study period, but a few used both
**FIG 1.** Study recruitment CONSORT diagram. (*) Passive withdrawals are those who did not complete the online questionnaires postbaseline but who did not formally request to be withdrawn. (NOTE. These were included as expected in the compliance calculations.) (†) Two patients who withdrew were sent and completed the (end-of-study (EOS) questionnaire as they had used the systems for some time. ECOG PS, Eastern Cooperative Oncology Group performance status; EORTC-QLQ-C30, European Organization for the Research and Treatment of Cancer Quality of Life Questionnaire C30; PIS, Patient Information Sheet; PRO-CTCAE, Patient Reported Outcomes version of the Common Terminology Criteria for Adverse Events.

**Assessed for eligibility (N = 888)**

- Ineligible (n = 368)
  - Did not receive eligible treatment (n = 77)
  - Eligible treatment group already filled (n = 65)
  - Discharged (n = 53)
  - Advised not appropriate to approach (n = 40)
  - Histology not eligible (n = 36)
  - No Internet and no landline phone (n = 19)
  - Timing/identified too late (n = 19)
  - Patient died (n = 20)
    (after being given PIS [n = 4])
  - Enrolled in another study (n = 15)
  - Performance status/not ECOG PS 2/3 when only recruiting this group (n = 11)
  - Language barrier (n = 8)
  - Out of area (n = 5)

- Eligible (n = 520)

  - Declined (n = 271)
    - No reason given (n = 57)
    - Too worried (n = 15)
    - Taken part in other research (n = 12)
    - Too ill (n = 11)
    - Other (n < 10)

  - Not interested (n = 26)

  - No Internet and said no to using phone (n = 21)

  - Did not want to do research (n = 19)

  - Hard of hearing (n = 15)

**Consented to REPORT-UK study (n = 249)**

- Eligible participants, 47.9%

**Completed baseline questionnaires, including both electronic questionnaires (n = 229)**

- Did not complete either electronic questionnaires at baseline (n = 7)
- Did not complete both electronic questionnaires at baseline
  - PRO-CTCAE (n = 17)
  - EORTC-QLQ-C30 (n = 10)

**Active withdrawal requests (n = 13)**

- Week 0 (n = 6)
- Week 1 (n = 2)
- Week 2 (n = 2)
- Week 3 (n = 1)
- Week 4 (n = 2)
- Week 10 (n = 1)
- Patient died (n = 6)

**Passive withdrawals* (n = 35)**

- Did not complete any electronic questionnaires postbaseline (n = 18)
- Did not complete any PRO-CTCAE questionnaires postbaseline (n = 19)
- Did not complete any EORTC-QLQ-C30 questionnaires postbaseline

**No. formally enrolled in the study at 12-week point excluding active withdrawals and deaths (n = 230)**

- Completed week 12 PRO-CTCAE electronic questionnaire (n = 155)
- Completed EOS questionnaire feedback (n = 192 of 232)†
### TABLE 1. Demographic Characteristics of the Study Population

| Characteristic                  | Overall (N = 249) | Internet (n = 196) | IVRS (n = 43) | Missing* (n = 10) |
|--------------------------------|-------------------|-------------------|--------------|------------------|
| **Mean age, years (SD)**       | 62.6 (13.4)       | 60.8 (13.7)       | 70.0 (9.9)   | 66.6 (8.4)       |
| **Gender**                     |                   |                   |              |                  |
| Male                           | 128 (51.4)        | 99 (50.5)         | 23 (53.5)    | 6 (60.0)         |
| Female                         | 121 (48.6)        | 97 (49.5)         | 20 (46.5)    | 4 (40.0)         |
| **Marital Status**             |                   |                   |              |                  |
| Married/civil partnership      | 173 (69.5)        | 138 (70.4)        | 30 (69.8)    | 5 (50.0)         |
| Cohabiting                     | 17 (6.8)          | 16 (8.2)          | 1 (2.3)      | 0 (—)            |
| Separated/divorced             | 16 (6.4)          | 11 (5.6)          | 2 (4.7)      | 3 (30.0)         |
| Widowed                        | 14 (5.6)          | 6 (3.1)           | 6 (14.0)     | 2 (20.0)         |
| Single                         | 29 (11.6)         | 25 (12.8)         | 4 (9.3)      | 0 (—)            |
| **Employment**                 |                   |                   |              |                  |
| Full time                      | 47 (18.9)         | 43 (21.9)         | 3 (7.0)      | 1 (10.0)         |
| Part time                      | 25 (10.0)         | 23 (11.7)         | 1 (2.3)      | 1 (10.0)         |
| Unable                         | 42 (16.9)         | 34 (17.3)         | 7 (16.3)     | 1 (10.0)         |
| Retired                        | 124 (49.8)        | 86 (43.9)         | 31 (72.1)    | 7 (70.0)         |
| Unemployed, not looking        | 5 (2.0)           | 5 (2.6)           | 0 (—)        | 0 (—)            |
| Unemployed, looking            | 1 (0.4)           | 0 (—)             | 1 (2.4)      | 0 (—)            |
| Other                          | 5 (2.0)           | 5 (2.6)           | 0 (—)        | 0 (—)            |
| **Education**                  |                   |                   |              |                  |
| Basic school level             | 93 (37.3)         | 64 (32.7)         | 23 (53.5)    | 6 (60.0)         |
| Higher education (eg, A level) | 61 (24.5)         | 51 (26.0)         | 8 (18.6)     | 2 (20.0)         |
| Degree level                   | 95 (38.2)         | 81 (41.3)         | 12 (27.9)    | 2 (20.0)         |
| **Study group**                |                   |                   |              |                  |
| Chemotherapy                   | 43 (17.3)         | 37 (18.9)         | 6 (14.0)     | 0 (—)            |
| Targeted agents                | 46 (18.5)         | 40 (20.4)         | 5 (11.6)     | 1 (10.0)         |
| Hormone therapy                | 43 (17.3)         | 26 (13.3)         | 13 (30.2)    | 4 (40.0)         |
| Radiotherapy                   | 43 (17.3)         | 34 (17.3)         | 8 (18.6)     | 1 (10.0)         |
| Surgery                        | 42 (16.9)         | 36 (18.4)         | 4 (9.3)      | 2 (20.0)         |
| ECOG PS 2-3                    | 32 (12.9)         | 23 (11.7)         | 7 (16.3)     | 2 (20.0)         |
| **Disease group**              |                   |                   |              |                  |
| Breast                         | 57 (22.9)         | 45 (23.0)         | 8 (18.6)     | 4 (40.0)         |
| Colorectal                     | 39 (15.7)         | 35 (17.9)         | 3 (7.0)      | 1 (10.0)         |
| Prostate                       | 33 (13.3)         | 19 (9.7)          | 12 (27.9)    | 2 (20.0)         |
| Leukemia                       | 15 (6.0)          | 13 (6.6)          | 2 (4.7)      | 0 (—)            |
| Myeloma                        | 13 (5.2)          | 10 (5.1)          | 3 (7.0)      | 0 (—)            |
| Lung                           | 12 (4.8)          | 6 (3.1)           | 5 (11.6)     | 1 (10.0)         |
| Renal                          | 12 (4.8)          | 11 (5.6)          | 1 (2.3)      | 0 (—)            |
| Head and neck                  | 11 (4.4)          | 10 (5.1)          | 1 (2.3)      | 0 (—)            |
| Rectal                         | 10 (4.0)          | 9 (4.6)           | 1 (2.3)      | 0 (—)            |
| Ovarian                        | 9 (3.6)           | 6 (3.1)           | 3 (7.0)      | 0 (—)            |
| Lymphoma                       | 9 (3.6)           | 8 (4.1)           | 0 (—)        | 1 (10.0)         |
| Pancreas                       | 7 (2.8)           | 7 (3.6)           | 0 (—)        | 0 (—)            |
| Bladder                        | 6 (2.4)           | 5 (2.6)           | 1 (2.3)      | 0 (—)            |

(continued on following page)
(10 patients when completing the PRO-CTCAE and 6 when completing the EORTC), and 1 patient who preferred IVRS and 3 who preferred the Internet used the opposite system more frequently. Questionnaire completion was quicker on the Internet (PRO-CTCAE median time, 3.0 minutes; interquartile range [IQR], 2.0 to 4.0 minutes; EORTC median time, 6.0 minutes; IQR, 4.0 to 8.0 minutes) compared with IVRS (PRO-CTCAE median time, 7.0 minutes; IQR, 6.1 to 13.6 minutes; EORTC median time, 14.5 minutes; IQR, 13.5 to 15.9 minutes).

Attrition and Compliance
At week 12, 92.4% of participants (230 of 249) remained on study, with 6 deaths and 13 active withdrawals (5.2%; 5 in week 1, 7 by week 5, and 1 in week 10). Five patients who withdrew were receiving hormone therapy and 8 were across the other treatment groups (ECOG/targeted/radiotherapy, n = 2; chemotherapy/surgery, n = 1). Withdrawal reasons included the time required to participate (n = 2), being unwell (n = 2), stopped treatment (n = 1), and computer problems (n = 2). Eighteen participants were classified as passive withdrawals (7.2%), having not completed any electronic questionnaires postbaseline.

Overall combined compliance for both systems was 74.9% for PRO-CTCAE (completed at 2,301 of 3,074 timepoints) and 79.1% for EORTC (completed at 749 of 947 timepoints). Table 2 lists compliance at each timepoint split by gender, system, study group, and age. Individual timepoint compliance (Figs 2A and 2B) shows more than 60% compliance throughout. Completions per participant illustrates that 46.1% (106 of 230) completed 13 or more PRO-CTCAE questionnaires, more than 80% (185 of 230) completed 8 or more, and 59.1% (136 of 230) completed the maximum 4 EORTC questionnaires (Figs 2C and 2D).

Compliance was higher in chemotherapy patients, with week 12 cumulative rates of 85.3% and 90.4% for PRO-CTCAE and EORTC, respectively (Table 2). Surgical participants had the lowest PRO-CTCAE compliance (baseline, 88.1%; week 12 cumulative, 67.3%). ECOG participants had lower EORTC compliance (baseline, 93.8%; month 3 cumulative, 68.7%).

Men and women had fairly similar compliance at each timepoint (Table 2). PRO-CTCAE compliance was slightly higher in the age group ≥ 60 years (week 12 cumulative, 78.5%) compared with the younger age group ≤ 59 years (week 12 cumulative, 68.9%). In contrast, EORTC compliance was relatively similar at each timepoint. PRO-CTCAE compliance fluctuated on both systems, but cumulative compliance was similar (IVRS, 74.5%; Internet, 76.7%). In contrast, EORTC compliance was higher on IVRS (cumulative month 3 IVRS, 85.6%; Internet, 79.8%); however, the number of IVRS users was small.

Patient and Staff Feedback
EOS questionnaires were completed by 192 (82.8%) of 232 patients (Internet, n = 157; IVRS, n = 35). Among nonresponders (n = 40), 20 had not completed any electronic PRO-CTCAE questionnaires postbaseline (ie, passive withdrawals; IVRS, n = 3; Internet, n = 17) and 20 had engaged postbaseline (IVRS, n = 4; Internet, n = 16). Almost all patients found the system(s) easy to use (IVRS, n = 35 [100%]; Internet, n = 156 [98.1%]) and access (IVRS, n = 32 [94.1%]; Internet, n = 148 [93.1%]); felt prepared post-training (IVRS, n = 35 [100%]; Internet, n = 153 [96.8%]), and more than 80% of all patients (151 of 188) were happy completing weekly questionnaires. Qualitative comments echoed this usability. Time to complete was “about right” (IVRS, n = 29 [82.9%]; Internet, n = 157 [99.4%]), although 6 IVRS users felt it was slow or took “too long”. Approximately one third (62 of 184; 33.7%) reported missing a week mainly because of illness (n = 21) or forgetting (n = 24). Fourteen hormone therapy patients (14 of 29; 48.3%) found few of the symptom questions to be relevant to them.

Staff surveys (32 of 50; 64%) indicated positive views about regular/weekly symptom reporting. Despite some concerns over resources and the time needed to review data, the
benefit for auditing symptoms and directing consultations and in clinical trials was emphasized.

**System Testing**

During the 17-month study, 3,170 questionnaires were completed (2,413 PRO-CTCAE questionnaires and 757 EORTC questionnaires) and proportionately few issues were recorded—38 patient-related and 69 system-related issues. Notable system issues included reminder problems (n = 20), information technology/server issues (n = 7) affecting access or auto-emails, completion report issues (n = 12), and some technical difficulties (Internet, n = 11; IVRS, n = 8). Overall downtime was not systematically recorded, but isolated incidences were experienced. Patient issues included initial access problems (n = 19), completion queries (n = 8), and lost/forgotten username and/or password (n = 6). The PRO database was combined successfully with the standard clinical trial database with only a single issue during the data retrieval/combining process.

**DISCUSSION**

To our knowledge, this is the first study illustrating the feasibility and acceptability of an electronic system to collect both self-reported PRO-CTAE and HRQOL data from a diverse sample of United Kingdom patients with cancer. The results demonstrate patient willingness, low attrition (5.2%; 13 of 249 active withdrawals), and high compliance (approximately 60% to 70% week 12; cumulative, approximately 75% to 80%). Most participants preferred the Internet, but approximately 17% of those who were older used IVRS, indicating an alternative mode was desirable. Combining PRO data with the clinical trial database worked effectively with few system issues recorded. This approach is therefore recommended and a comparative evaluation is now required.

The 47.9% recruitment rate is lower than the 70% uptake rate routinely observed in PRO studies, but is encouraging compared with the 31% to 64% uptake rate in mailed cancer surveys. Target participants included various diagnoses, which made robust interpretation of any differences between cancer groups challenging as a result of small frequencies. Disease site was not associated with the rate of consent for the study; however, uptake was specifically low in some groups (eg, prostate and hormone therapy). Therefore, the study findings cannot be generalized to all cancer populations and may not reflect

### TABLE 2. Cumulative Compliance Over Time

| Questionnaire | Gender | Overall | Male | Female | System | Targeted | Hormone | Radiotherapy | Surgery | ECOG PS 2-3 | Age, Years |
|---------------|--------|---------|------|--------|--------|----------|---------|--------------|---------|-------------|------------|
| **PRO-CTCAE** |        |         |      |        |        |          |         |              |         |             |            |
| Baseline      | 93.2   | 93.0    | 93.4 | 95.3   | 96.9   | 100.0    | 93.5    | 86.0         | 97.7    | 88.1        | 93.8       |
| Week 1        | 86.6   | 85.7    | 87.4 | 85.7   | 90.0   | 95.3     | 85.9    | 87.8         | 90.7    | 74.7        | 83.9       |
| Week 2        | 84.0   | 83.4    | 84.6 | 80.8   | 87.5   | 93.0     | 81.2    | 85.1         | 92.1    | 70.2        | 81.5       |
| Week 3        | 82.2   | 81.9    | 82.5 | 79.5   | 85.3   | 90.6     | 77.7    | 85.6         | 90.5    | 70.3        | 77.0       |
| Week 4        | 80.3   | 80.4    | 80.2 | 79.6   | 82.8   | 88.3     | 73.8    | 82.8         | 89.0    | 72.3        | 74.3       |
| Week 5        | 79.3   | 79.6    | 79.1 | 78.9   | 81.7   | 87.5     | 73.4    | 82.3         | 87.6    | 71.1        | 73.1       |
| Week 6        | 78.9   | 79.3    | 78.4 | 76.9   | 81.3   | 88.2     | 72.7    | 81.4         | 85.9    | 71.3        | 72.2       |
| Week 7        | 78.3   | 78.5    | 78.0 | 77.0   | 80.5   | 88.2     | 72.5    | 80.4         | 84.3    | 70.9        | 71.8       |
| Week 8        | 77.0   | 77.1    | 76.9 | 77.3   | 78.9   | 86.4     | 70.9    | 79.7         | 82.8    | 69.9        | 71.1       |
| Week 9        | 76.4   | 76.8    | 76.0 | 77.1   | 78.2   | 85.8     | 70.7    | 79.6         | 81.4    | 69.5        | 70.1       |
| Week 10       | 76.0   | 76.4    | 75.6 | 76.5   | 77.8   | 86.4     | 70.5    | 79.6         | 80.0    | 68.6        | 69.5       |
| Week 11       | 75.5   | 76.0    | 74.9 | 75.3   | 77.3   | 85.9     | 69.9    | 79.1         | 79.2    | 68.1        | 69.7       |
| Week 12       | 74.9   | 75.4    | 74.3 | 74.5   | 76.7   | 85.3     | 69.5    | 78.7         | 78.0    | 67.3        | 69.0       |
| **EORTC**     |        |         |      |        |        |          |         |              |         |             |            |
| Baseline      | 96.0   | 95.3    | 96.7 | 100.0  | 99.0   | 100.0    | 95.7    | 93.0         | 100.0   | 92.9        | 93.8       |
| Month 1       | 88.5   | 88.0    | 89.0 | 92.7   | 90.3   | 95.3     | 85.7    | 89.0         | 94.0    | 86.6        | 77.4       |
| Month 2       | 84.5   | 84.5    | 84.6 | 90.1   | 85.7   | 95.2     | 80.7    | 85.0         | 88.8    | 81.1        | 73.0       |
| Month 3       | 79.1   | 79.6    | 78.6 | 85.6   | 79.8   | 90.4     | 74.9    | 77.8         | 83.1    | 76.5        | 68.7       |

**NOTE.** Data are given as percentages, unless otherwise noted.

**Abbreviations:** ECOG PS, Eastern Cooperative Oncology Group performance status; EORTC, European Organization for Research and Treatment of Cancer Quality of Life Questionnaire-C30; IVRS, interactive voice response system; PRO-CTCAE, Patient Reported Outcomes version of the Common Terminology Criteria for Adverse Events.
the compliance/views of specific cancer groups who were under-represented in this study. Patients required Internet access or a touch-tone telephone, which excluded some patients or those not confident with technology. In a clinical trial setting, offering paper-based PROs should continue, although this may diminish as Internet use increases. Those who declined tended to be older, but reasons focused on being too busy/overwhelmed with treatment. This study provided limited personal benefit beyond altruism; therefore, if electronic PRO reporting was integrated into treatment trials, uptake could be higher. Basch et al report higher uptake, although their approach included clinic-based computer completions. Future trials, combining remote/home-based completions with clinic-based completions may optimize compliance.

Compliance overall was high (approximately 60% to 70% at week 12; cumulative, approximately, 75% to 80%) and higher still in chemotherapy patients regularly going to the hospital for treatment reviews. PRO-CTCAE items chosen were particularly applicable to chemotherapy-related symptoms. Lower weekly compliance for surgical patients may be a result of the intense recovery period immediately postsurgery and less frequent review post-surgery. Some clinical trials that included paper-based QOL measures have observed higher compliance (75% at 6 months), but the current study had more intense frequency (weekly/every 4 weeks). In an active clinical trial, compliance may be higher because of patient selection, trial-related incentives, and item relevance. Basch et al reported 92% cumulative compliance for home PRO-CTCAE reporting in trial patients, but 14.7% of questionnaires were collected via a coordinator directly ringing patients.

Despite most participants choosing to use the Internet, some older patients preferred the IVRS. An alternative real-time method is desirable if the purpose of collection is tracking symptoms/AEs, whereas if the aim is obtaining HRQOL outcomes, paper-based alternatives may suffice.

Overall compliance did not differ across systems, but different patterns were observed, potentially influenced by the order of presentation: EORTC was presented first on the IVRS at the once every 4 weeks intervals (explaining higher EORTC IVRS compliance), whereas Internet users had to proactively choose both questionnaires. Furthermore, completions on the Internet were quicker, which may be a result of the systems themselves (ie, IVRS required the questions/answers to be read out at a comprehensible speed), but it may also reflect the younger, more educated Internet users.
Patient feedback was positive, finding the systems acceptable and problem free. Most issues were related to initial queries or access problems and were quickly resolved. Reasons for noncompletion were a result of being unwell or forgetting, rather than negativity about the systems.

The systems were only available in English, which is a limitation that should be addressed in future work. Furthermore, the general oncology setting influenced how the study was presented (focusing on system testing rather than on the value of collecting PROs in trials) and potentially affected uptake and compliance. The definition of baseline differed to what would be done in a formal clinical trial setup: A lenient baseline PRO-CTCAE/HRQOL completion (up to 3 days postregistration) was used, rather than prerandomization, as in many clinical trials.

The study highlights implementation considerations for future actual clinical trial use, including the need for a serious AE (or suspected unexpected serious adverse reaction) alert system, clinicians directly receiving symptom summary reports or alerts, and a function for individual symptom reporting or adding other free-text symptoms. Some of the considerations also relate to concerns voiced regarding the ethics of PRO data in clinical trials and the timescale of data review.30 Using an electronic PRO system with automated alerts helps with the timely management of these issues compared with paper-based methods, but creates new training needs for trial staff.31 Furthermore, electronic data collection requires continuous assessment to ensure that information technology systems and infrastructure meet clinical trial regulation and Medicines and Healthcare products Regulatory Agency and European Medicines Agency requirements.32

In conclusion, routine collection of PRO data in cancer clinical trials is increasingly important. This study illustrates the feasibility and patient acceptance of electronic PROs during treatment and incorporating PRO data into clinical trials. Additional exploration of this system is warranted where PROs are combined with other trial data—alongside traditional approaches—to improve data quality and patient safety.

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AUTHORS' DISCLOSURES OF POTENTIAL CONFLICTS OF INTEREST

Electronic Patient Reporting of Adverse Events and Quality of Life: A Prospective Feasibility Study in General Oncology

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