Web-Based Module for the Collection of Electronic Patient-Reported Outcomes in People Living With HIV in Nouvelle Aquitaine, France: Usability Evaluation

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Abstract

Background: Patient-reported outcomes (PROs) can be of great value for both research and chronic disease management. We developed a new module of the ANRS CO3 Aquitaine cohort study’s Web-based data capture and visualization solution (APPEGE 2.0) for the collection of electronic PROs among people living with HIV cared for in Nouvelle Aquitaine, France.

Objective: This study aimed to evaluate the usability of 2 successively developed prototypes of ARPEGE 2.0’s electronic PROs module before launching a pilot study, owing to the novelty of the proposed data collection method for our setting and specific characteristics of the target population.

Methods: A total of 2 sequential rounds of empirical, task-based usability evaluations were conducted, involving 8 research staff and then 7 people living with HIV. Evaluators provided written feedback during round 1 and oral feedback during round 2. Evaluators who completed the full set of tasks responded to the System Usability Scale (SUS). We assessed changes in SUS scores between rounds and concluded usability testing when SUS scores reached a ceiling effect, defining good usability a priori as a usability score of 70.

Results: Insights were generated regarding the visibility of system status and the match between the system and the real world that improved the module’s usability. Research staff evaluators reported mean SUS scores of 65 (SD 18.87) and patient evaluators reported mean SUS scores of 85 (SD 5.4; P=.032).

Conclusions: Software modifications, informed by successive rounds of usability testing, resulted in sufficient gains in usability to undertake piloting. Insights generated during evaluations prompted us to find the appropriate balance between optimal security and ease of use.

Trial Registration: ClinicalTrials.gov NCT03296202; https://clinicaltrials.gov/ct2/show/NCT03296202

International Registered Report Identifier (IRRID): RR2-10.2196/10.2196/resprot.9439

(JMIR Form Res 2019;3(4):e15013) doi: 10.2196/15013
KEYWORDS
patient reported outcome measures; patient generated health data; quality of life

Introduction
HIV, once fatal, is now a manageable chronic illness [1]. In Western Europe, the majority of individuals who received a diagnosis of HIV are in care and on potent antiretroviral therapy, which prevents serious diseases both related and unrelated to AIDS [2]. The improved prognosis and the increased life expectancy of people living with HIV (PLWH) makes preserving health and ensuring good quality of life the cornerstone of their care [3-5]. One strategy to help providers respond to PLWH’s evolving needs and improve the quality and efficiency of their overall care is collecting and using patient-reported outcomes (PROs) [6].

PROs or “any report of the status of the patient’s health condition that comes directly from the patient, without interpretation of the patient’s response by a clinician or anyone else” [7] have been used extensively in clinical research [6]. PROs can be used at the population level for research and to improve the quality of care or at the individual level to support clinical decision making [8]. Their use may allow for more accurate symptom detection, better patient-provider communication, and improved outcomes [9]. Logistical, technical, and ideological barriers have nevertheless limited their use in routine care [10]. The adoption of electronic medical records coupled with the adaptation of paper questionnaires to computerized and internet-based formats may help overcome these barriers [10,11].

With evidence from the United States suggesting that the collection of PROs by using touchscreen-based information technology was both feasible and of value for both research and clinical HIV care [12-14], a prototype of an electronic PRO module linked to the ANRS CO3 Aquitaine cohort’s data capture and visualization system (ARPEGE 2.0) was developed in 2017 [15]. As the overall usefulness of interactive health care applications or their usability is likely to affect their acceptability and adoption, usability evaluations of 2 successively developed prototypes of the ARPEGE 2.0 solution were conducted in preparation for a pilot study [15].

Methods
This formative research study took place in Bordeaux, France, at the Inserm UMR 1219-Bordeaux Population Health Research Centre and the St André Bordeaux University Hospital. It was designed as part of the ANRS CO3 Aquitaine study, an open, prospective hospital-based cohort of PLWH in care in 13 clinics in southwestern France. A local institutional review board approved the study’s protocol (Comité de Protection de Personnes Sud-Ouest et Outre-Mer III) on September 18, 2017.

Description of the Electronic Patient-Reported Outcome Module Powered by ARPEGE 2.0
ARPEGE 1.0 is a proprietary, secure, electronic case report form developed in Microsoft ASP.NET (WebForm). Data are stored within a Microsoft SQL Server 2014–based data management system. The ANRS CO3 Aquitaine cohort relies on ARPEGE 1.0 for data capture. Clinical data, extracted from both medical records and laboratory data, derived from the hospital’s laboratory information management systems, have been collected systematically since 1987 and electronically via ARPEGE 1.0 since 2013 with the support of Clinical Research Associates. ARPEGE 2.0 is a generic Web-based data capture and visualization system also developed in Microsoft ASP.NET (WebForm). ARPEGE 2.0 has enabled the creation of the module for the collection of electronic PROs in routine care for observational research and, ultimately, clinical care.

The content of ARPEGE 2.0’s initial electronic PRO module is based on current treatment guidelines for people being treated for HIV and associated comorbidities [16]. Prototyping was carried out over 2017 with the support and regular feedback from a working group comprising research staff, local stakeholders, and end users (clinicians and patient representatives). The questionnaires were evaluated individually according to their psychometric properties, administration method, and length. The following areas are covered by the electronic PRO module: socioeconomic status and individual social and material deprivation [17], multidimensional quality of life (WHOQOL-HIV BREF) [18], treatment burden (Treatment Burden Questionnaire) [19], physical activity (the Short Version of the International Physical Activity Questionnaire), alcohol use and screening for at-risk drinking behavior (Alcohol Use Disorders Identification Test Consumption, Fast Alcohol Consumption Evaluation) [20], tobacco and nicotine use and screening for tobacco dependency (Fagerström), cannabis (Cannabis Abuse Screening Test) and drug use, and, finally, depression (Patient Health Questionnaire) [21].

Conditional branching was used where appropriate. The module also allows patients to report any other treatment-related issues in a free text field. Where applicable, the International Society for Pharmacoeconomics and Outcomes Research ePRO Task Force’s recommendations on adapting paper-based instruments were followed, ensuring that data produced are equivalent or superior to those generated from paper-based administration methods [22].

Recruitment
Nielsen’s recommendations that favor conducting several iterative studies, each with a small number of participants, were adopted [23]. In round 1 (May 2018), evaluators were employees of the Inserm UMR 1219 Bordeaux Population Health Research Center or affiliated with the project, referred to herein as research staff. In round 2 (June 2018), a convenience sample of PLWH being cared for at the St André Bordeaux University Hospital was identified by clinical staff either before or during their routine visit.

Procedure
The evaluation procedure differed between round 1 and round 2. However, for both rounds, oral consent was obtained. It was
then explained that each study participant (evaluator) would be provided with a unique identifier, which would allow him/her to create a personal account and access the questionnaires. Evaluators were shown the study-specific brochure where the number would be written on a detachable coupon (Multimedia Appendix 1). They were asked to complete 5 tasks: (1) navigate between pages on the publicly available website and locate key information, (2) create a user account, (3) confirm their account, (4) initiate the electronic PRO assessment, and (5) complete the electronic PRO assessment. Whether or not each task was completed with ease, assistance or not was monitored, and a score of 2 to 0 was attributed (2=the task was completed with ease and 0=it was not completed). The highest possible score was therefore 10, and the lowest score was 0. Neither round 1 nor round 2 evaluators were compensated.

In round 1, research staff were provided with instructions detailing the background of the study and how it would be implemented in a clinical setting. Evaluators were given a link to a staging version of the electronic PRO module. They were asked to complete the previously described tasks. They then responded to an Web-based questionnaire that included the System Usability Scale (SUS), a widely used robust tool for measuring usability. It consists of 10 items with 5 response options, from strongly agree to strongly disagree [24,25]. Evaluators provided written feedback in an open text field and by email.

In round 2, patients participated in one-on-one testing sessions, lasting between 1 and 2 hours, with a researcher in a dedicated, private space at the hospital (June 2018). The researcher based each session on a standardized qualitative interview guide. A personal computer (Mac Book Air) with access to the staging site was provided to complete the study tasks. Patient evaluators were also allowed to complete the questionnaire on their personal smartphones, matching how the electronic PRO module might be accessed in routine care. Evaluators were instructed to use the think aloud method, in which users are asked to verbalize all thoughts as they interact with the system while carrying out tasks. Subsequently, those who completed all tasks responded orally to the SUS and provided open-ended feedback [24]. All sessions were audio recorded, and field notes were taken.

**Analysis**

Task completion and SUS scores were calculated for each evaluator, and means and standard deviations were calculated for each round. We performed a t test assuming unequal variance to determine if each round of testing produced significant difference in the mean SUS scores. A priori, we defined success in usability when the SUS score reached a ceiling effect, with a minimum score of 70—generally accepted as a cut-off for good usability [26].

Qualitative analysis included review of written feedback, audio recording–enhanced field notes, and responses to open-ended questions. We performed thematic content analysis on written feedback and audio recording–enhanced field notes, abstracting and compiling emerging themes from each round of testing. These are reported according to Nielsen’s usability heuristic categories [27].

**Results**

**Overview**

Table 1 presents evaluators’ characteristics and mean task completion scores for rounds 1 and 2. The majority of round 1 evaluators were women (7/8). They reported using computers either regularly (5/8) or often (3/8). In all, 5 out of 7 round 2 evaluators were men. A total of 3 reported using a computer regularly, 3 often, and 1 never. Overall, mean task completion scores were 7.8 (out of 10) in round 1 and 7.1 in round 2. In round 1, 7 evaluators completed all tasks compared with 4 out of 7 in round 2. Task completion was hampered owing to 2 evaluators being locked out of their accounts and 1 evaluator being unable to complete tasks owing to poor eyesight. This evaluator was attributed 0 on all tasks.

The usability insights uncovered during the 2 rounds of usability evaluations together with the solutions adopted are presented in Table 2.
Table 1. Evaluator characteristics and task scores.

| Evaluator characteristics | Task 1—information found | Task 2—account created | Task 3—account confirmed | Task 4—PRO\(^a\) assessment initiated | Task 5—PRO assessment completed | Total |
|---------------------------|--------------------------|------------------------|--------------------------|--------------------------------------|---------------------------------|-------|
| **Round 1 (N=8)**         |                          |                        |                          |                                      |                                 |       |
| Male (n=1)                | 2.0                      | 1.1                    | 1.4                      | 1.5                                  | 1.8                             | 7.8   |
| 30-40                     | 2.0                      | 0.0                    | 0.0                      | 0.0                                  | 0.0                             | 2.0   |
| Female (n=7)              | 2.0                      | 1.3                    | 1.6                      | 1.7                                  | 2.0                             | 8.6   |
| <30                       | 2.0                      | 1.5                    | 1.5                      | 1.5                                  | 2.0                             | 8.5   |
| 30-40                     | 2.0                      | 1.3                    | 1.3                      | 1.7                                  | 2.0                             | 8.3   |
| 41-50                     | 2.0                      | 1.0                    | 2.0                      | 2.0                                  | 2.0                             | 9.0   |
| >50                       | 2.0                      | 1.0                    | 2.0                      | 2.0                                  | 2.0                             | 9.0   |
| **Round 2 (N=7)**         | 1.7                      | 1.1                    | 1.4                      | 1.7                                  | 1.1                             | 7.1   |
| Male (n=5)                | 1.6                      | 1.2                    | 1.6                      | 1.6                                  | 1.6                             | 7.6   |
| <30                       | 2.0                      | 1.5                    | 2.0                      | 2.0                                  | 2.0                             | 9.5   |
| 30-40                     | 2.0                      | 2.0                    | 2.0                      | 2.0                                  | 2.0                             | 10.0  |
| >50                       | 1.0                      | 0.5                    | 1.0                      | 1.0                                  | 1.0                             | 4.5   |
| Female (n=2)              | 2.0                      | 1.0                    | 1.0                      | 2.0                                  | 0.0                             | 6.0   |
| >50                       | 2.0                      | 1.0                    | 1.0                      | 2.0                                  | 0.0                             | 6.0   |

\(^a\)PRO: patient-reported outcome.
Table 2. Usability insights per round and solution adopted according to Nielsen’s usability heuristics.

| Usability categories | Round 1—research staff | Round 2—patients | Solution |
|----------------------|-------------------------|------------------|----------|
| Visibility of system status | Login procedure was confusing owing to the complexity of password, requiring 2 symbols | Challenges adhering to password requirements for certain patients | Password requirements were spelled out for users in bold. A password visualization button was also added to the password field to allow users to ensure that passwords created matched before registering their account |
| —^a | Unclear whether the QuAliV number (required for creating the account) is case sensitive | Information incorporated into the presentation of the study to participants |
| Validation of questionnaire unclear | — | Information buttons added to the home page of the electronic PRO module instructing users on how the questionnaires functioned and reminding them to submit their completed questionnaires. The button was also relabeled to make its functionality clearer |
| Match between system and the real world | Date picker was in English and began in 2018, requiring users to click to go back in time | — | The date picker was replaced with a French version. It allowed users to type in their birth dates without using the calendar |
| — | Nonmutually exclusive modalities or response missing | Minor modifications made to question modalities to ensure clarity |
| — | Issues stemming from the translation of questionnaire from English to French | Further cognitive debriefing with native speakers to identify the best translation of the item in question |
| — | Difficulties understanding the meaning of certain questions | Less formal language substituted where possible and examples given to facilitate the comprehension of certain questions |
| — | Confirmation of account on one’s smartphone (email) resulted in being locked out of one’s account on another device | Automatic connection to the site after creating one’s account deleted (temporarily) to avoid users locking themselves out of their account. Users must reenter their username and password |
| User control and freedom | Need for returning back to last page completed in the questionnaire | — | The user is now redirected back to the most recent page completed within each questionnaire. Scrolling from one page of a questionnaire to another automatically saves entered data |
| — | Radio button could not be unclicked or erased | A refresh button was added to each item to allow users to erase their responses and therefore leave items unanswered |
| — | Questionnaire opens in a pop-up window whose size cannot be modified | Double checked to ensure that text could be easily read in each window |
| — | Unclear whether users had to provide first and last name | We added text indicating that typing one’s first and last name was optional |
| Consistency and standards | Format of certain questions was noted as being inconsistent between questionnaires, Yes/No questions appeared in a table format as soon as they used the same response thesaurus | — | Minor improvements in formatting were made where possible. Further development required to accommodate this change in the longer run |
| — | Typos in certain questions were identified | — |
| Error prevention | Aberrant response possible for certain free text fields | — | Stricter constraints added |
| — | The password required was complex. Instructions on password requirements were missing from the account creation page | Instructions on password requirements added |
| Need to clarify units in free text fields | — | Units added in gray in each text field |
An asterisk was added to indicate which questions were mandatory. The user is sent back to mandatory questions before being allowed to progress in the questionnaire. These questions were marked in red to indicate that they were mandatory.

Error message added to the module explaining that users would be able to reaccess their accounts after 20 min.

Questionnaires are programmed to open successively.

Further trouble shooting using full array of browsers and devices.

An 11-point radio button scale was proposed as a temporary solution.

Alternative formatting used to improve readability.

Progression bar for each questionnaire goes from orange to green as soon as all nonconditional questions are answered. Users are directed to unanswered obligatory questions upon attempting to go on to the next page of the questionnaire.

What Did Not Work

The account creation task was the most challenging for users. One of the issues identified was the complexity of the password requirements. The password had to be entered twice and contain at least 8 alphanumeric characters, including 2 special characters and a capital letter (Figure 1). Many evaluators, both research staff and patients, attempted this step more than once. We clarified the password requirements and ensured that error messages were informative regarding the system status, and we made it possible to visualize the password after round 1 (Multimedia Appendix 2). As errors still occurred, we added additional error prevention features. The password is validated as the user types as opposed to the user receiving an error message upon clicking register.

What Worked

The first task involved navigating the external website that patients would access from home, unassisted, to create their account. Users found the information provided on the external website quickly and found its structure clear. All users quickly understood how the attributed unique identifier would be used to create their personal account. Once users had created their account, efforts to guide him/her through electronic PROs by having each questionnaire open one after the other appeared to work well. The use of stoplight-style color coding and a progress bar allowed users to see if they had missed a question and helped them recognize, diagnose, and recover from errors seamlessly. The order of the PROs was received positively by users and therefore remained unchanged between prototype versions.
Figure 1. Initial log-in page (round 1).

![Initial log-in page](image1)

Fields (top to bottom): Name, First Name, E-mail address, Confirm e-mail address, Password, Confirm password, QuAliV Number.

Password instructions (in bold): The password must contain at least 8 characters (letters and numbers) of which one must be uppercase and two special characters among the following: @ # $ % & * + ! ?

Figure 2. Revised log-in page (round 2).

![Revised log-in page](image2)

Fields (top to bottom): Name (optional), First Name (optional), E-mail address, Confirm e-mail address, Password, Confirm password, QuAliV Number.

Password instructions (in bold): The password must contain at least 8 characters: 1 uppercase, 1 number and 2 special characters among the following: @ # $ % & * + ! ? /
A login problem, also detected during the second round of usability testing, was being locked out of one’s account accidentally. This issue arose from a security measure included in the electronic PRO module’s design. Users were logged out automatically after a period of 20 min of inactivity. If users accidently left the page without logging out of their accounts, they could no longer log back in owing to the Bordeaux University servers’ restrictions. If the user attempted to return to their account, they received an error message indicating that they were already connected. This issue could not be resolved without completely releasing the automatic logout timeframe (shortening it). We therefore modified the error message indicating that the user could access their account again in 20 min.

**System Usability Scale Scores**

In round 1, experts reported mean SUS scores of 65 (SD 18.87), and patients, in round 2, reported mean SUS scores of 85 (SD 5.4) ($P = .032$).

**Discussion**

**Principal Findings**

Iterative usability evaluations of 2 successively developed prototypes allowed us to see how easy our electronic PRO module was to use and identify when and where users encountered problems or experienced confusion. We were able to improve the module’s usability markedly, specifically the visibility of system status and the match between the system and the real world, and take into account the specific needs of our patient population (their level of computer literacy and age) and the specificities of our clinical setting. Finally, we were pushed to find the appropriate balance between optimal security and ease of use.

Unlike PRO collection methods employed in clinics in the United States [12–14], where patients complete an electronic PRO assessment by using touchscreen information technology with the assistance of a research assistant/administrator at clinics, we aimed to design a Web-based Bring Your Own Device solution. We therefore assumed that the majority of users would have access to a smartphone or personal computer with a reliable internet connection. The proposed solution, developed in-house, had to work well enough to allow a group of users, with varying levels of computer familiarity, to use it with little to no assistance.

**Strengths and Limitations**

Some caveats should be considered in the interpretation of our results. We conducted the first round of usability testing in a sample of research staff who may not fully represent end users. This strategy, recognized as an easy way of catching obvious usability issues, resulted in high-quality, detail-oriented, and exhaustive feedback, allowing for a number of basic usability problems to be resolved before evaluations with patients. Most evaluators were comfortable using computers and the internet. They may not fully reflect the diversity of the cohort of PLWH in the region. More purposeful sampling of evaluators with lower computer literacy may have resulted in the detection of additional usability insights.

In round 2, we used the think aloud method. This method has been known to slow the thought process and increase mindfulness, which might prevent errors that might have normally occurred [28]. However, when evaluators are asked to perform simple tasks, the method has been shown to have no effect on user performance [29]. We opted for this method as the tasks were not considered complex.

**Conclusions**

Nevertheless, software modifications, informed by successive rounds of usability testing, resulted in sufficient gains in usability to undertake piloting.

**Acknowledgments**

Alain Volny-Anne (European AIDS Treatment Group) and Eugenie Destandau provided valuable feedback on the design and content and the QuAliV website and electronic PRO module. The authors would like to thank all of the evaluators, both research staff and patients, who contributed to this study. They thank the scientific committee, clinical sites and their investigators, and clinical research associates, further cited, who have been involved in the successful implementation of the subsequent phase of the study.

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The Aquitaine ANRS Cohort is sponsored by the Bordeaux University Hospital and funded by the ANRS (France REcherche Nord& Sud Sida-hiv Hépatites) and the Bordeaux University Hospital. The cohort is coordinated from within the Inserm UMR 1219—Bordeaux Population Health Research Centre. Seed funding was granted by the ANRS in 2017 via the CSS-5 call to develop the electronic PRO module. Diana Barger was awarded a 36-month young researcher grant from Sidaction to design and conduct a study on quality of life in PLWH within the ANRS CO3 Aquitaine cohort as part of her doctoral research.

Conflicts of Interest
DB has received a speaking fee from Gilead. FB declares to have received reimbursement for attending a symposium from ViiV Healthcare, Gilead, Bristol-Myers Squib, Merck and Janssen, speaking fee and consultancy fee from ViiV Healthcare, Gilead, Bristol-Mys-Squib, Merck and Janssen, and funds for research from Gilead and ViiV Healthcare. The other authors do not have any conflict of interest to declare.

Multimedia Appendix 1
Patient brochure with unique identifier. [PNG File, 478 KB-Multimedia Appendix 1]

Multimedia Appendix 2
Login page with password verification. [PNG File, 234 KB-Multimedia Appendix 2]

References
1. Deeks SG, Lewin SR, Havlir DV. The end of AIDS: HIV infection as a chronic disease. Lancet 2013 Nov 2;382(9903):1525-1533 [FREE Full text] [doi: 10.1016/S0140-6736(13)61809-7] [Medline: 24152939]
2. INSIGHT START Study Group, Lundgren JD, Babiker AG, Gordin F, Emery S, Grund B, et al. Initiation of antiretroviral therapy in early asymptomatic HIV infection. N Engl J Med 2015 Aug 27;373(9):795-807 [FREE Full text] [doi: 10.1056/NEJMoa1506816] [Medline: 26192873]
3. Rasmussen LD, Obel N. How do we preserve health among adults living with HIV? Lancet HIV 2019 Feb;6(2):e69-e70. [doi: 10.1016/S2352-3018(18)30336-9] [Medline: 30683626]
4. Antiretroviral Therapy Cohort Collaboration. Survival of HIV-positive patients starting antiretroviral therapy between 1996 and 2013: a collaborative analysis of cohort studies. Lancet HIV 2017 Aug;4(8):e349-e356 [FREE Full text] [doi: 10.1016/S2352-3018(17)30066-8] [Medline: 28501492]
5. Lazarus JV, Safreed-Harmon K, Barton SE, Costagliola D, Dedes N, Valero J, et al. Beyond viral suppression of HIV - the new quality of life frontier. BMC Med 2016 Jun 22;14(1):94 [FREE Full text] [doi: 10.1186/s12916-016-0640-4] [Medline: 27334606]
6. Engler K, Lessard D, Lebouché B. A review of HIV-specific patient-reported outcome measures. Patient 2017 Apr;10(2):187-202. [doi: 10.1007/s40271-016-0195-7] [Medline: 27637488]
7. Food and Drug Administration. 2009 Oct. Patient-Reported Outcome Measures: Use in Medical Product Development to Support Labeling Claims URL: https://www.fda.gov/regulatory-information/search-fda-guidance-documents/patient-reported-outcome-measures-use-medical-product-development-support-labeling-claims [accessed 2019-11-12]
8. Snyder CF, Aaronson NK, Choucair AK, Elliott TE, Greenhalgh J, Halyard MY, et al. Implementing patient-reported outcomes assessment in clinical practice: a review of the options and considerations. Qual Life Res 2012 Oct;21(8):1305-1314. [doi: 10.1007/s11136-011-0054-x] [Medline: 22048932]
9. Greenhalgh J, Long AF, Flynn R. The use of patient reported outcome measures in routine clinical practice: lack of impact or lack of theory? Soc Sci Med 2005 Feb;60(4):833-843. [doi: 10.1016/j.socscimed.2004.06.022] [Medline: 15571900]
10. Jones JB, Snyder CF, Wu AW. Issues in the design of internet-based systems for collecting patient-reported outcomes. Qual Life Res 2007 Oct;16(8):1407-1417. [doi: 10.1007/s11136-007-9235-z] [Medline: 17668293]
11. Gensheimer SG, Wu AW, Snyder CF, PRO-EHR Users’ Guide Steering Group, PRO-EHR Users’ Guide Working Group. Oh, the places we’ll go: patient-reported outcomes and electronic health records. Patient 2018 Dec;11(6):591-598. [doi: 10.1007/s40271-018-0321-9] [Medline: 29968179]
12. Crane HM, Lober W, Webster E, Harrington RD, Crane PK, Davis TE, et al. Routine collection of patient-reported outcomes in an HIV clinic setting: the first 100 patients. Curr HIV Res 2007 Jan;5(1):109-118. [doi: 10.2174/157016207779316369] [Medline: 17266562]

13. Kozak MS, Mugavero MJ, Ye J, Aban I, Lawrence ST, Nevin CR, et al. Patient reported outcomes in routine care: advancing data capture for HIV cohort research. Clin Infect Dis 2012 Jan 1;54(1):141-147 [FREE Full text] [doi: 10.1093/cid/cir727] [Medline: 22042879]

14. Schumacher JE, McCullumsmith C, Mugavero MJ, Ingle-Pang PE, Raper JL, Willig JH, et al. Routine depression screening in an HIV clinic cohort identifies patients with complex psychiatric co-morbidities who show significant response to treatment. AIDS Behav 2013 Oct;17(8):2781-2791 [FREE Full text] [doi: 10.1007/s10461-012-0342-7] [Medline: 23086427]

15. Barger D, Leleux O, Conte V, Sapparrart V, Gapillout M, Crespel I, et al. Integrating electronic patient-reported outcome measures into routine HIV care and the ANRS CO3 Aquitaine Cohort's Data Capture and Visualization System (QuAliIV): protocol for a formative research study. JMIR Res Protoc 2018 Jun 7;7(6):e147 [FREE Full text] [doi: 10.2196/resprot.9439] [Medline: 29880467]

16. Ministry of Solidarity and Health. 2013. Medical management of people living with HIV, recommendations of the expert group, 2013 report URL: http://solidarites-sante.gouv.fr/IMG/pdf/Rapport_Morlat_2013_Mise_en_ligne.pdf [accessed 2017-11-16] [WebCite Cache ID 6v1OhYDd1]

17. Labbe E, Blanquet M, Gerbaud L, Poirier G, Sass C, Vendittelli F, et al. A new reliable index to measure individual deprivation: the EPICES score. Eur J Public Health 2015 Aug;25(4):604-609. [doi: 10.1093/eurpub/cku231] [Medline: 25624273]

18. O'Connell KA, Skevington SM. An international quality of life instrument to assess wellbeing in adults who are HIV-positive: a short form of the WHOQOL-HIV (31 items). AIDS Behav 2012 Feb;16(2):452-460. [doi: 10.1007/s10461-010-9863-0] [Medline: 21181253]

19. Tran V, Montori VM, Eton DT, Baruch D, Falissard B, Ravaud P. Development and description of measurement properties of an instrument to assess treatment burden among patients with multiple chronic conditions. BMC Med 2012 Jul 4;10:68 [FREE Full text] [doi: 10.1186/1741-7015-10-68] [Medline: 22762272]

20. Dawson DA, Grant BF, Stinson FS, Zhou Y. Effectiveness of the derived Alcohol Use Disorders Identification Test (AUDIT-C) in screening for alcohol use disorders and risk drinking in the US general population. Alcohol Clin Exp Res 2005 May;29(5):844-854. [doi: 10.1097/01.alc.0000164374.32229.a2] [Medline: 15897730]

21. Kroenke K, Spitzer RL, Williams JB. The PHQ-9: validity of a brief depression severity measure. J Gen Intern Med 2001 Sep;16(9):606-613 [FREE Full text] [doi: 10.1046/j.1525-1497.2001.016009606.x] [Medline: 11556941]

22. Coons SJ, Gwaltney CJ, Hays RD, Zhou Y, Revicki DA, ISPOR ePRO Task Force. Recommendations on evidence needed to support measurement equivalence between electronic and paper-based patient-reported outcome (PRO) measures: ISPOR ePRO Good Research Practices Task Force report. Value Health 2009 Jun;12(4):419-429 [FREE Full text] [doi: 10.1111/j.1524-4733.2008.00470.x] [Medline: 19900250]

23. Nielsen J, Landauer TK. A Mathematical Model of the Finding of Usability Problems. In: Proceedings of the INTERACT '93 and CHI '93 Conference on Human Factors in Computing Systems. 1993 Presented at: CHI'93; April 24-29, 1993; Amsterdam, The Netherlands p. 206-213.

24. Brooke J. Hell - Jens Oliver Meiert. 1996. SUS - A Quick and Dirty Usability Scale URL: https://hell.meiert.org/core/pdf/sus.pdf [accessed 2019-11-12]

25. Bangor A, Kortum PT, Miller JT. An empirical evaluation of the system usability scale. Int J Human-Comput Interact 2008;24(6):574-594. [doi: 10.1080/10447310802055776]

26. Sauro J. User Experience Magazine. 2011 Aug. SUStisfied? Little-Known System Usability Scale Facts URL: https://uxpamagazine.org/sustisfied/ [accessed 2019-06-15]

27. Nielsen J, Nielsen Norman Group: UX Training, Consulting, & Research. 2012 Jan 3. Nielsen Norman Group: UX Training, Consulting, & Research. 2012 Jan 3. Usability 101: Introduction to Usability URL: https://www.nngroup.com/articles/usability-101-introduction-to-usability/ [accessed 2019-04-12]

28. Dumas JS, Redish JC. A Practical Guide To Usability Testing. Bristol: Intellect Ltd; 1994.

29. Ericsson AK, Herbert SA. Protocol Analysis: Verbal Reports As Data. Boston: The Mit Press; 1993.

Abbreviations

PLWH: people living with HIV
PRO: patient-reported outcome
SUS: System Usability Scale
Please cite as:
Barger D, Leleux O, Conte V, Sapparrart V, Gapillout M, Crespel I, Erramouspe M, Delveaux S, Wittkop L, Dabis F, Bonnet F
Web-Based Module for the Collection of Electronic Patient-Reported Outcomes in People Living With HIV in Nouvelle Aquitaine, France: Usability Evaluation
JMIR Form Res 2019;3(4):e15013
URL: http://formative.jmir.org/2019/4/e15013/
doi: 10.2196/15013
PMID: 31850847

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