

RESEARCH ARTICLE

Design and implementation of a global site assessment survey among HIV clinics participating in the International epidemiology Databases to Evaluate AIDS (IeDEA) research consortium

Ellen Brazier^{1,2*}, Fernanda Maruri³, C. William Wester^{3,4}, Beverly Musick⁵, Aimee Freeman⁶, Angela Parcesepe⁷, Stefanie Hossmann⁸, Benedikt Christ⁸, April Kimmel⁹, John Humphrey¹⁰, Esther Freeman¹¹, Leslie A. Enane¹², Kathryn E. Lancaster¹³, Marie Ballif⁸, Jonathan E. Golub¹⁴, Denis Nash^{1,2}, Stephany N. Duda¹⁵, on behalf of the International epidemiology Databases to Evaluate AIDS (IeDEA) consortium[¶]



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1 City University of New York, Institute for Implementation Science in Population Health, New York, NY, United States of America, **2** City University of New York, Graduate School of Public Health and Health Policy, New York, NY, United States of America, **3** Department of Medicine, Division of Infectious Diseases, Vanderbilt University Medical Center, Nashville, TN, United States of America, **4** Vanderbilt Institute for Global Health, Nashville, TN, United States of America, **5** School of Medicine, Indiana University, Indianapolis, IN, United States of America, **6** Johns Hopkins Bloomberg School of Public Health, Baltimore, MD, United States of America, **7** Department of Maternal and Child Health, Gillings School of Global Public Health, University of North Carolina at Chapel Hill, Chapel Hill, NC, United States of America, **8** Institute of Social and Preventive Medicine, University of Berne, Berne, Switzerland, **9** Department of Health Behavior and Policy, Virginia Commonwealth University School of Medicine/VCU Health, Richmond, VA, United States of America, **10** Division of Infectious Diseases, Department of Medicine, Indiana University, Indianapolis, IN, United States of America, **11** Department of Dermatology, Massachusetts General Hospital, Harvard Medical School, Boston, MA, United States of America, **12** The Ryan White Center for Pediatric Infectious Disease and Global Health, Department of Pediatrics, Indiana University School of Medicine, Indianapolis, IN, United States of America, **13** Division of Epidemiology, College of Public Health, The Ohio State University, Columbus, OH, United States of America, **14** Center for Tuberculosis Research, Johns Hopkins University School of Medicine, Baltimore, MD, United States of America, **15** Department of Biomedical Informatics, Vanderbilt University Medical Center, Nashville, TN, United States of America

[¶] The Complete Membership of the Study Group can be found in IeDEA Study Membership in [S3 Text](#).
* ellen.brazier@sph.cuny.edu, ekbrazier@gmail.com

Abstract

Introduction

Timely descriptions of HIV service characteristics and their evolution over time across diverse settings are important for monitoring the scale-up of evidence-based program strategies, understanding the implementation landscape, and examining service delivery factors that influence HIV care outcomes.

Methods

The International epidemiology Databases to Evaluate AIDS (IeDEA) consortium undertakes periodic cross-sectional surveys on service availability and care at participating HIV treatment sites to characterize trends and inform the scientific agenda for HIV care and implementation science communities. IeDEA's 2020 general site assessment survey was

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Abbreviations: IeDEA, International epidemiology Databases to Evaluate AIDS; VUMC, Vanderbilt University Medical Center; REDCap, Research Electronic Data Capture; PMTCT, Prevention of maternal to child transmission.

developed through a consultative, 18-month process that engaged diverse researchers in identifying content from previous surveys that should be retained for longitudinal analyses and in developing expanded and new content to address gaps in the literature. An iterative review process was undertaken to standardize the format of new survey questions and align them with best practices in survey design and measurement and lessons learned through prior IeDEA site assessment surveys.

Results

The survey questionnaire developed through this process included eight content domains covered in prior surveys (patient population, staffing and community linkages, HIV testing and diagnosis, new patient care, treatment monitoring and retention, routine HIV care and screening, pharmacy, record-keeping and patient tracing), along with expanded content related to antiretroviral therapy (differentiated service delivery and roll-out of dolutegravir-based regimens); mental health and substance use disorders; care for pregnant/postpartum women and HIV-exposed infants; tuberculosis preventive therapy; and pediatric/adolescent tuberculosis care; and new content related to Kaposi's sarcoma diagnostics, the impact of COVID-19 on service delivery, and structural barriers to HIV care. The survey was distributed to 238 HIV treatment sites in late 2020, with a 95% response rate.

Conclusion

IeDEA's approach for site survey development has broad relevance for HIV research networks and other priority health conditions.

1. Introduction

HIV care and treatment programs around the world continue to evolve and expand. However, the changing characteristics of HIV services are not well described across countries and global regions. Timely descriptions of the evolution of and variations in the characteristics of HIV services over time are important for documenting the comprehensiveness of care provided and progress in the scale-up of services and evidence-based service delivery strategies, as well as advancing the global HIV community's implementation science agenda [1]. Such data can enhance understanding of the implementation landscape for HIV care across the globe, while being essential for efforts to identify service delivery factors that are associated with care outcomes of interest among those newly enrolling in HIV care and those engaged in long-term HIV care.

The International epidemiology Database to Evaluate AIDS (IeDEA) consortium (<http://www.iedea.org>) is a global research consortium of HIV care and treatment sites in seven geographic regions: the Asia-Pacific; the Caribbean, Central, and South America; North America; Central Africa; East Africa; Southern Africa; and West Africa. Established in 2006, the IeDEA research consortium currently comprises close to 400 HIV care and treatment sites in 44 countries that contribute longitudinal data for more than 2 million patients, including adults and children, ever enrolled in HIV care—data that are used in epidemiology research on HIV care and treatment outcomes.

IeDEA undertakes periodic surveys among sites that are active in the consortium in order to document HIV service availability across sites and over time—information that is helpful for understanding patient-level data and examining how patient outcomes are associated with

health facility and service delivery characteristics. We herein describe the process and methodology used in developing IeDEA's 2020 site assessment survey and fielding it across treatment sites within the research consortium.

2. Materials and methods

Study aim and objectives

IeDEA comprises a heterogeneous mix of HIV care and treatment clinics at academic and community-based hospitals and health centers that have joined the consortium at different timepoints. In view of the dynamic and changing composition of the research consortium, along with the evolving global and national HIV response, IeDEA has conducted periodic surveys of participating clinics since 2009 in an effort to document service availability and practices. Because the consortium does not dictate clinical practices or policies to participating sites, these periodic site surveys allow for the characterization of clinics and services across the consortium, which is particularly important as IeDEA sites may not be representative of all HIV clinics in a given country or region because of the specialized HIV care they offer and/or the large volume of patients they serve. Conducted every three to five years, these surveys have targeted all active sites in most IeDEA regions, with convenience sampling generally used in the Southern Africa region, where patient data are contributed by both stand-alone HIV care and treatment centers and by large HIV programs that compile data from numerous small clinics that they are supporting (hereafter referred to as programmatic cohorts).

Designed as a cross-sectional facility-level survey of HIV treatment sites, the objectives of IeDEA's 2020 consortium-wide survey were to: (1) systematically document facility, program, and service characteristics of active sites, including selected clinic and service delivery attributes that change over time; (2) collect data on specialized topics related to HIV care that constitute important gaps in the scientific literature; (3) compile preliminary data for future research studies; and (4) collect data to inform the development and implementation of future in-depth topic-specific site assessment surveys, as well as intervention studies.

Survey development process

The content of IeDEA's 2020 site assessment survey was developed through a consultative 18-month process (Fig 1) that built on lessons learned through prior IeDEA-wide surveys [2–4], while harnessing the expertise of researchers across IeDEA. As a first step in this process, members of IeDEA's technical working groups [5] were invited to review the content of the previous general site assessment survey, conducted in 2017, and make recommendations for

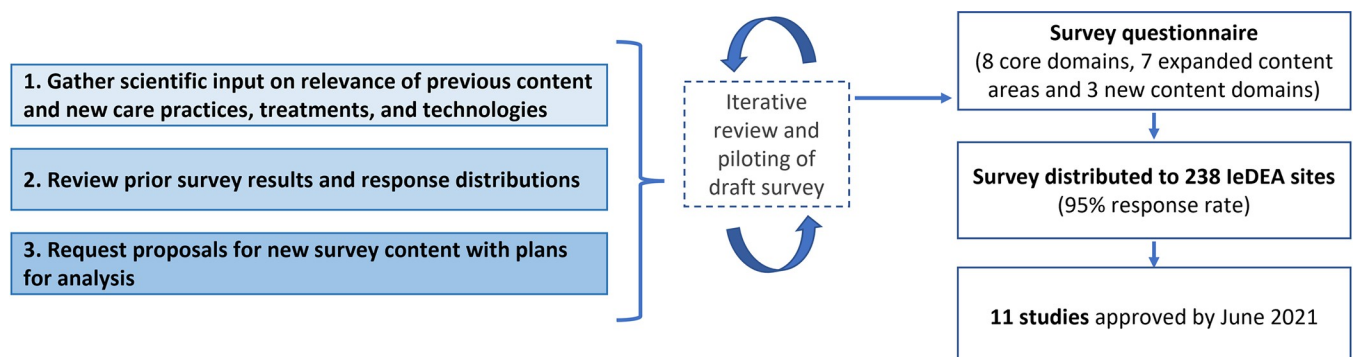


Fig 1. Survey development process (2019–2021).

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content that should be retained to facilitate longitudinal analyses of service delivery and practices across IeDEA sites. IeDEA's technical working groups were also asked to identify questions that could be retired because they were no longer relevant or because of concerns with the quality of data generated during the previous survey. In addition, data from the 2017 survey were analyzed to identify response options that should be collapsed or combined to avoid counts too small for meaningful analysis. Through this review process, core domains of the survey questionnaire were defined.

In addition to reviewing historical content, a complementary proposal-driven process was undertaken for the development of expanded and new survey content. Each of IeDEA's technical working groups were invited to prepare brief research proposals to support the addition of new survey content, articulating focused objectives and proposing specific survey questions to address key gaps in the literature. Investigators proposing new survey content were requested to specify plans for using the data (e.g., as the basis for an analysis that would result in a manuscript and/or as pilot data to inform a more comprehensive research study). An iterative review process was undertaken to standardize the format of new survey content, ensuring that questions conformed with best practices in site-level survey design and measurement developed through prior IeDEA site assessment surveys and used uniform response options, branching logic and skip patterns to help minimize respondent burden. Through this process, expanded content was developed for several domains covered by prior surveys, and several new survey domains were proposed.

While it was originally intended that the survey would explore the status of service delivery at the time of survey implementation, calendar year 2019 was selected as the primary reference period for most survey questions because of disruptions related to the SARS-CoV-2 pandemic in 2020. This pre-COVID-19 reference period was also selected because several of the associated research proposals planned to link site survey data with patient-level data for the pre-pandemic period.

After compiling the survey content as a draft paper questionnaire, the questionnaire was programmed in REDCap version 10.3.2 (Research Electronic Data Capture)—a web-based software platform for secure data capture in research studies. REDCap provides a user-friendly interface for data capture; audit capabilities for tracking data manipulation; automated export routines for common statistical packages (e.g., R, SAS, Stata, SPSS); and procedures for data integration with external sources [6, 7].

Housed in a local data center hosted at Vanderbilt University Medical Center (VUMC), the REDCap database for the 2020 survey was extensively tested and refined through an iterative review process to ensure that the online survey was streamlined for data entry and easy to navigate. Once the online data capture instruments were completed in mid-2020, the survey was piloted in English with HIV clinic staff at eight sites across four IeDEA regions. The pilot served to identify questions and wording that were not consistently understood, as well as various modifications to improve the survey instructions, response options, skip patterns, and technical content of the questionnaire.

After addressing issues emerging through the pilot, the survey content and REDCap database were finalized in English, and the questionnaire was then translated into French for use in Francophone settings. The translation was completed by a professional translator with specialized expertise in health care, pharmaceuticals, and HIV. The draft translation was then reviewed by IeDEA colleagues in the West Africa region to ensure that it accurately reflected the idiomatic and technical terms predominant in the region. The final text in the French questionnaire was used in a second deployment of the REDCap database for IeDEA-participating sites in Francophone countries.

3. Results

Survey content

The survey developed through IeDEA's consultative 18-month process covered eight core domains included in prior surveys, including patient populations served, clinic staffing and community linkages, HIV testing and diagnostic capacity, care for newly-enrolling patients, antiretroviral therapy (ART) adherence and retention strategies, routine care and screening for HIV patients, pharmacy services, and record-keeping (S1 Text). In addition, it included expanded content in seven thematic areas (differentiated delivery of antiretroviral therapy; roll-out of dolutegravir-based regimens; screening and management of mental health and substance use disorders; tuberculosis preventive therapy; pediatric and adolescent tuberculosis care; and pregnancy and postpartum care for women and HIV exposed infants) and three new content domains (structural barriers to HIV-related care, Kaposi's sarcoma diagnostics, and the impact of COVID-19 on routine HIV care and treatment).

Sampling procedures

All sites that were actively contributing longitudinal patient-level data to IeDEA in 2020 were eligible for inclusion in the survey. Sites that were exiting IeDEA in 2020 ($n = 3$) were ineligible for inclusion, along with sites that were part of an interval cohort ($n = 14$) (i.e., sites where patient data reflect systematic assessments at pre-specified and regular intervals, rather routine patient care extracted from medical records). In the Southern Africa region of IeDEA, where participating sites include 12 stand-alone clinics and 201 clinics that are part of six large programmatic cohorts, a systematic, hybrid sampling strategy was used. First, to facilitate longitudinal analyses of HIV care and service attributes across time, we included all stand-alone sites, along with five sites that had been selected through convenience sampling in prior surveys to represent five of the region's large programmatic cohorts. Further, to ensure the representativeness of sites selected from within large programmatic cohorts, stratified random sampling was used to select an additional 18 cohort-representative sites, with the number of urban vs. rural sites determined based on the distribution of clinics within each cohort.

The final sample eligible for the survey comprised 238 sites (Fig 2), including all active sites in the Asia-Pacific region ($n = 52$), Caribbean, Central, and South America ($n = 9$), Central Africa ($n = 21$), East Africa ($n = 74$), West Africa ($n = 14$), along with all sites in North America that are not part of an interval cohort ($n = 36$), and 32 sites from the Southern Africa region (15% of all active sites in the region).

Ethical review

The conduct of the 2020 IeDEA-wide site assessment survey was reviewed and designated a non-human subjects operational/quality improvement project by the VUMC Institutional Review Board (#200013).

Survey implementation

The survey was self-administered in either English or French, using paper forms and online REDCap electronic versions of the questionnaire, depending on country context. In Spanish and Portuguese-speaking contexts, the English-language questionnaire was used. In each IeDEA region, data managers distributed the survey instructions, a PDF of the questionnaire, and unique site-specific survey links to investigators and/or research staff at eligible sites. Site-level partners selected staff with in-depth knowledge about the care and services provided to adult and pediatric HIV patients to complete the questionnaire. These survey respondents

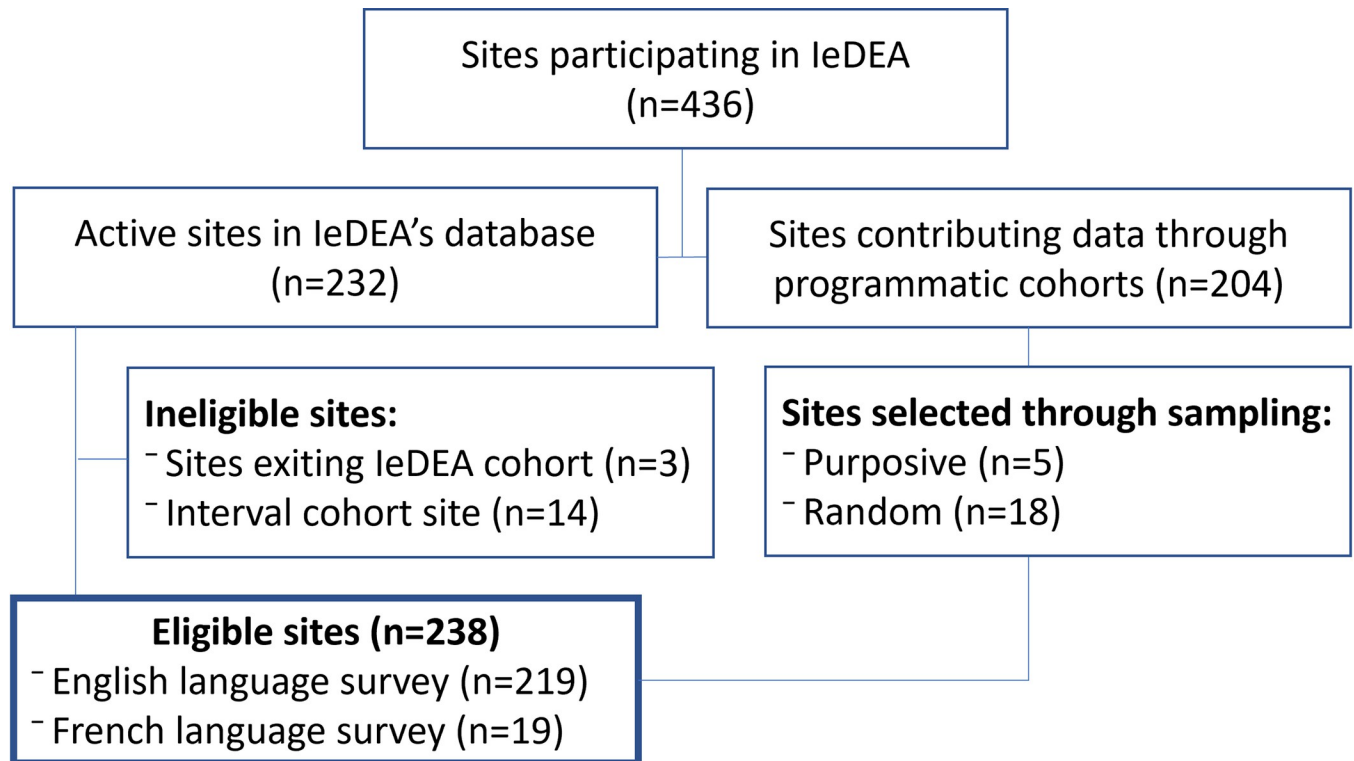


Fig 2. Site inclusion flowchart.

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were encouraged to consult with staff from other units of the health facility, such as pharmacy and laboratory departments, where necessary. Sites that had participated in the pilot were asked only to respond to new and revised questions.

The English-language version of the survey launched on September 11, 2020, and the French-language version launched one month later, on October 16, 2020. Throughout the data collection period, data quality checks were performed centrally by regional data managers to identify incomplete surveys and surveys with inconsistent data so that in-country coordinators could follow-up with specific sites, as needed, with reminders and queries to maximize the survey response rate and resolve data quality issues. Surveys completed on paper were entered into the REDCap electronic database by local or regional data managers.

When the survey was closed on March 1, 2021, responses had been received from 227 sites across leDEA's seven regions (Fig 3). The overall response rate was 95% (11 sites did not begin the survey), and the urban vs. rural distribution of responding sites (69% and 31%, respectively) matched the distribution of eligible sites (Table 1). Of 227 submitted surveys, 98% were complete. Most survey respondents (88%) identified themselves as clinical staff or managers (e.g., clinical officer in charge, clinic manager, or other clinical officer), with 9% of surveys completed by site-level data managers or affiliated research staff and 4% not specified. Almost half (48%) of the surveys were completed by more than one staff (range: 2 to 18), meaning that the survey responses reflected the perspectives of multiple staff and providers involved in HIV-related care.

Data management

After the survey closed, additional data quality checks were performed to review data completeness and consistency, and site-level respondents were queried, as needed, to resolve data

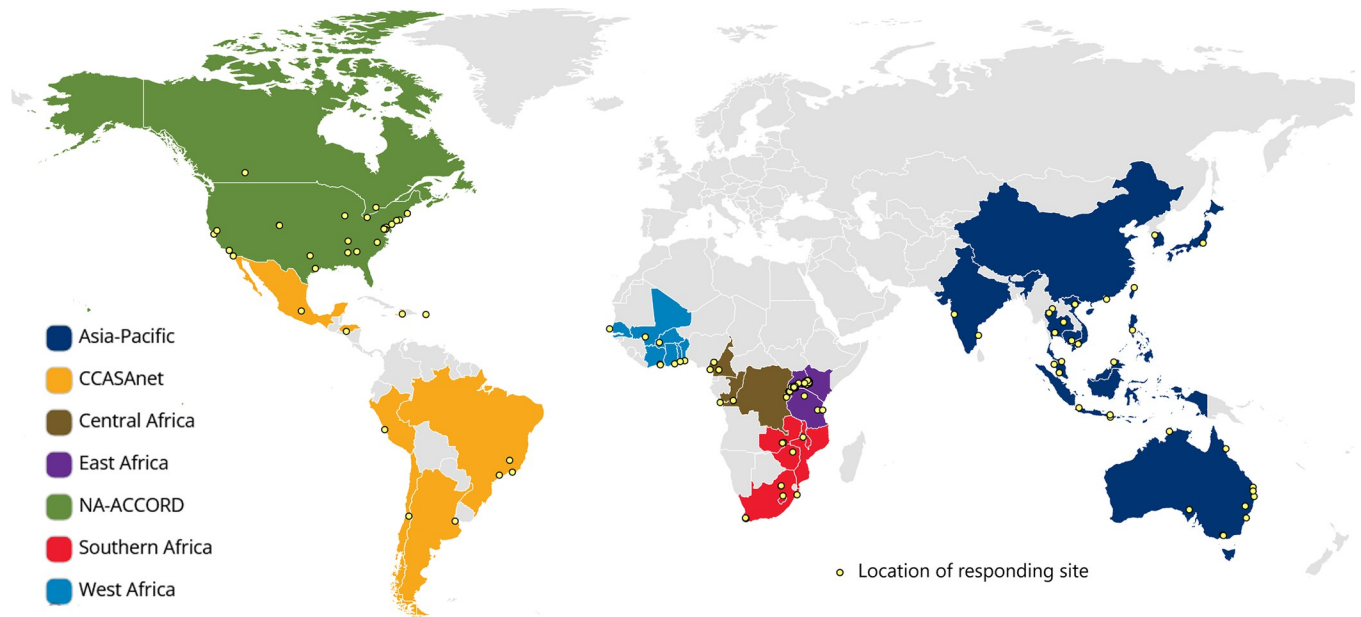


Fig 3. IeDEA regions and locations of responding sites.

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quality issues. Following the completion of data quality checks in April 2021, data from the English and French databases were merged, and REDCap’s automated export tools were used to extract the data for use in analyses. All survey information, including survey instructions, English and French questionnaires, linked research proposals, and data exports were stored in a secure drive at VUMC for dissemination to IeDEA investigators. By June 2021, data had been released for all analyses planned through the survey development process, along with an additional longitudinal analysis (Table 2).

Table 1. 2020 IeDEA-wide site assessment survey response and completion rates.

Region	Eligible sites			Responding sites ^c			Survey Completion
	N	Urban	Rural	N (%) ^d	Urban	Rural	n (%) ^e
		n (%) ^d	n (%) ^d		n (%) ^e	n (%) ^e	
Asia-Pacific	52	45 (88%)	6 (12%)	51 (98%)	45 (88%)	6 (12%)	51 (100%)
Caribbean, Central, and South America	9	19 (90%)	2 (10%)	9 (100%)	19 (90%)	2 (10%)	8 (89%)
Central Africa ^a	21	9 (100%)	0 (0%)	21 (100%)	9 (100%)	0 (0%)	21 (100%)
East Africa	74	19 (26%)	55 (74%)	74 (100%)	19 (26%)	55 (74%)	74 (100%)
North America	36	36 (100%)	0 (0%)	30 (83%)	30 (100%)	0 (0%)	29 (97%)
Southern Africa	32	22 (69%)	10 (31%)	28 (88%)	20 (71%)	8 (29%)	26 (93%)
West Africa ^b	14	14 (100%)	0 (0%)	14 (100%)	14 (100%)	0 (0%)	14 (100%)
Total	238	164 (69%)	73 (31%)	227 (95%)	156 (69%)	71 (31%)	223 (98%)

^a English version survey was completed by 15 sites, French version survey was completed by 6 sites

^b English version survey was completed by one site, French version survey was completed by 13 sites

^c Survey responses included 75 questionnaires initially completed on paper and 152 completed in REDCap.

^d Denominator: All eligible sites

^e Denominator: All responding sites

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Table 2. Analyses using the IeDEA 2020 site assessment survey data.

Data collected through IeDEA's 2020 site assessment survey are being used in studies addressing the following aims:

1. Describing the availability of differentiated HIV services including antiretroviral therapy delivery models and the patient populations eligible for differentiated antiretroviral therapy delivery.
2. Characterizing the introduction of Dolutegravir-based regimens, including its use as first, second, and/or third-line therapy across HIV clinics in low- and middle-income countries, patient eligibility criteria used (e.g., age, sex, and clinical characteristics), and practices related to transitioning patients to Dolutegravir-based regimens, including practices related to viral load testing, resistance testing, and monitoring of side effects among patients transitioned to Dolutegravir-based regimens.
3. Describing facility-level practices surrounding user fees for adult HIV diagnosis and laboratory management and describing the accessibility of these services (e.g., availability on- vs. off-site).
4. Describing the availability of prevention of maternal to child transmission (PMTCT) services across IeDEA clinics, including the proportion of programs offering integrated PMTCT services and differences in the characteristics of PMTCT service delivery across HIV clinics.
5. Describing the availability and approaches used in screening and management of mental health disorders in HIV clinic settings.
6. Describing clinic practices for substance use disorder screening and management across IeDEA and the integration of these services in HIV care.
7. Describing routine practices for tuberculosis disease screening, diagnosis, and treatment for children and adolescents who are living with HIV.
8. Describing clinical practices for screening for tuberculosis disease, testing for latent tuberculosis infection, and practices related to prescribing and monitoring tuberculosis preventive therapy among PLWH.
9. Characterizing the capacity, clinical practices, and existence of user fees for diagnosis of Kaposi sarcoma, including the types of Kaposi sarcoma diagnostics available in low- and middle-income countries.
10. Describing the availability of Pre-Exposure Prophylaxis across adult HIV clinics, along with trends in the availability of Pre-Exposure Prophylaxis since 2014.
11. Characterizing service delivery challenges and constraints posed by the COVID-19 pandemic at HIV clinics, as well as site-level mitigation strategies introduced to ensure the continuity of HIV care.

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4. Discussion

IeDEA's consortium-wide site assessment surveys provide an opportunity to document the evolution of HIV care and treatment for adults and children among a globally diverse cohort of clinics at all levels of the health system, in urban and rural areas and in low-, middle-, and high-income settings. IeDEA's established relationships with participating sites allow us to collect timely data on the adoption and implementation of new guidance and best practices among real-world service delivery programs in order to track progress and identify implementation gaps. The ability to link this programmatic information with data for patients enrolled in HIV care at participating sites, further allows for investigation of associations between program and policy implementation and patient outcomes of interest.

Important strengths of IeDEA's 2020 site assessment survey development process include data-driven and proposal-driven approach, which engaged diverse methodological and scientific expertise within the consortium. By starting with an analysis of prior IeDEA-wide survey data, we were able to identify questions and response options that have not provided reliable or useful information previously. In addition, ensuring that new survey content was developed by investigators involved in IeDEA's technical working groups and supported by detailed, aim-driven proposals with specific plans for using the data helped ensure that time and effort expended on data collection led to analyses and manuscripts that address salient and timely questions for the global HIV response. With work in progress for 11 papers addressing gaps in the literature, IeDEA's 2020 site assessment survey promises to generate more scientific productivity than IeDEA's prior consortium-wide general site assessment surveys [2–4] and specialized surveys focused on specific patient populations (e.g., pediatrics, pregnant and postpartum women) [8] and areas of service delivery (e.g., tuberculosis, cancer, mental health disorders, etc.) [9–15].

Several limitations of IeDEA's site assessment survey implementation process are worth noting. First, these surveys rely on voluntary self-report by survey respondents. While the presence or absence of specific services or care components can be captured rather reliably with minimal misclassification, it is also likely that knowledge about HIV care and service delivery varies across respondent types and across sites. Moreover, social desirability bias may result in some respondents reporting practices outlined in relevant norms and guidelines, as opposed to their day-to-day practices. Recall bias is another potential concern, particularly as the survey primarily focused on practices prior to the COVID-19 pandemic and because 2020 was a year with extraordinary challenges for health service delivery around the globe, with many health care facilities introducing changes in practice to mitigate risks associated with the pandemic. It is possible that recall of services and practices prior to the pandemic were inaccurate or incomplete, potentially resulting in either over- or under-reporting of services. While such biases cannot be completely avoided in voluntary self-administered surveys, our high response and survey completion rates, as well as the fact that almost half of the sites involved multiple staff in completing the survey may partially mitigate the impact of these biases.

In addition, while the high overall response rate for the survey means that data are broadly representative for sites participating in IeDEA, these sites may not be representative of HIV service delivery within some countries and regions, particularly in contexts where IeDEA sites were initially identified because they served a sizable patient population and/or offered advanced levels of care. This limitation notwithstanding, as IeDEA's 2020 site assessment survey included a large number of sites and settings that have participated in previous IeDEA-wide surveys, it is expected that temporal trends in HIV service delivery among IeDEA sites may be broadly reflective of prevailing trends across other HIV sites that are not part of the consortium [16].

5. Conclusion

IeDEA's consortium-wide site assessments position it to address timely knowledge gaps related to HIV care and treatment service delivery, as well as to substantially contribute to the field of HIV implementation science globally. The collaborative approach used in developing IeDEA's 2020 global site assessment survey has broad applicability for other large HIV service provider networks and programs, along with research and service provision networks addressing other health conditions, such as tuberculosis and other non-communicable conditions.

Supporting information

S1 Text. IeDEA 2020 site assessment survey.

(PDF)

S2 Text. Acknowledgments.

(DOCX)

S3 Text. IeDEA study membership.

(DOCX)

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creating a map with the locations of sites that responded to the 2020–2021 IeDEA-wide site survey.

Author Contributions

Conceptualization: Ellen Brazier, Fernanda Maruri, C. William Wester, Angela Parcesepe, Stefanie Hossmann, Benedikt Christ, April Kimmel, John Humphrey, Esther Freeman, Leslie A. Enane, Kathryn E. Lancaster, Marie Ballif, Jonathan E. Golub, Denis Nash, Stephany N. Duda.

Funding acquisition: Denis Nash, Stephany N. Duda.

Methodology: Ellen Brazier, Fernanda Maruri, C. William Wester, Beverly Musick, Aimee Freeman, Denis Nash, Stephany N. Duda.

Writing – original draft: Ellen Brazier.

Writing – review & editing: Fernanda Maruri, C. William Wester, Beverly Musick, Aimee Freeman, Angela Parcesepe, Stefanie Hossmann, Benedikt Christ, April Kimmel, John Humphrey, Esther Freeman, Leslie A. Enane, Kathryn E. Lancaster, Marie Ballif, Jonathan E. Golub, Denis Nash, Stephany N. Duda.

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