Case Report

Each patient is a research biorepository: informatics-enabled research on surplus clinical specimens via the living BioBank

Alexander V. Alekseyenko, Bashir Hamidi, Trevor D. Faith, Keith A. Crandall, Jennifer G. Powers, Christopher L. Metts, James E. Madory, Steven L. Carroll, Jihad S. Obeid, and Leslie A. Lenert

1Biomedical Informatics Center, Medical University of South Carolina, Charleston, South Carolina, USA, 2Department of Public Health Sciences, Medical University of South Carolina, Charleston, South Carolina, USA, 3Department of Oral Health Sciences, Medical University of South Carolina, Charleston, South Carolina, USA, 4Department of Healthcare Leadership and Management, Medical University of South Carolina, Charleston, South Carolina, USA, 5Department of Biostatistics & Bioinformatics, Computational Biology Institute, Milken Institute School of Public Health, George Washington University, Washington DC, USA, 6Department of Dermatology, University of Iowa, Iowa City, Iowa, USA, 7Division of Pathology Informatics, Department of Pathology and Laboratory Medicine, Medical University of South Carolina, Charleston, South Carolina, USA, 8Department of Pathology and Laboratory Medicine, Medical University of South Carolina, Charleston, South Carolina, USA, and 9Department of Medicine, Medical University of South Carolina, Charleston, South Carolina, USA

Corresponding Author: Alexander V. Alekseyenko, PhD, FAMIA, Biomedical Informatics Center, Medical University of South Carolina, 22 WestEdge St, MSC 200, WG213C, Charleston, SC 29403, USA (alekseye@musc.edu)

Received 18 May 2020; Revised 11 August 2020; Editorial decision 6 September 2020; Accepted 9 September 2020

ABSTRACT

The ability to analyze human specimens is the pillar of modern-day translational research. To enhance the research availability of relevant clinical specimens, we developed the Living BioBank (LBB) solution, which allows for just-in-time capture and delivery of phenotyped surplus laboratory medicine specimens. The LBB is a system-of-systems integrating research feasibility databases in i2b2, a real-time clinical data warehouse, and an informatics system for institutional research services management (SPARC). LBB delivers deidentified clinical data and laboratory specimens. We further present an extension to our solution, the Living mBiome Bank, that allows the user to request and receive phenotyped specimen microbiome data. We discuss the details of the implementation of the LBB system and the necessary regulatory oversight for this solution. The conducted institutional focus group of translational investigators indicates an overall positive sentiment towards potential scientific results generated with the use of LBB. Reference implementation of LBB is available at https://LivingBioBank.musc.edu.

Key words: just-in-time biobanking, phenotyped specimens, phenotyped specimen data, translational research services

STATEMENT OF THE PROBLEM

The need for molecular and other data from precisely phenotyped human specimens is paramount in translational research. Although a part of the general solution, expansive biobanking of such specimens is not always viable due to the high burden and expense of biobank establishment and maintenance. Although not a complete replacement, just-in-time capture of specimens from standard-of-
care surplus sources can support many use cases of traditional biobanking, if this can be accomplished with minimal disruptions to the clinical laboratory’s mission. Examples of existing platforms that provide just-in-time specimen capture functionality include (i) Harvard’s Crimson system, which combines its Informatics for Integrating Biology and the Bedside (i2b2) with a link to its laboratory specimen processing system, and (ii) the Vanderbilt BioVU system, which operates more like a traditional biobank with storage of a broad range of specimens.

Living BioBank (LBB) seeks to extend the model developed in Crimson from its proprietary platform to a more generally applicable solution and to extend the approach across research networks. In addition, a limitation of the Crimson system is the necessity for the specimens and patient data to be identified to the end user. The specimen recruitment process involves identified chart review by investigators to accept or reject the available specimen. This typically requires a higher degree of regulatory oversight than if the specimens were to be deidentified and clinical abstraction limited to just a predefined phenotype. As a result, documentation and time requirements for such studies increase. A more streamlined solution would allow for a precise electronic phenotype to guide the surplus specimen capture without the need to validate every individual patient’s record.

In this case report, we describe the Medical University of South Carolina Biomedical Informatics Center LBB solution for phenotyped surplus specimen capture that integrates a number of institutionally adopted informatics systems to deliver some functionality of just-in-time biobanking in a reduced regulatory burden environment. We demonstrate the use of LBB in an institutional review board (IRB)-approved Living μBiome Bank study protocol that further seeks to augment the phenotype with phenotyped specimen microbiome data and limited deidentified clinical data to support microbiome data interpretation. We assess the feasibility of the LBB solution in terms of the accuracy of e-phenotypes and the time required for delivery of a set of specimens. We also report on evaluation of user acceptance of the solution.

TECHNICAL DESCRIPTION OF THE LIVING BIOBANK

Figure 1 outlines the LBB solution players, systems, processes, and deliverables. Although many electronic phenotyping systems may suit the purpose, the current implementation of the LBB (https://LivingBioBank.musc.edu/) uses i2b2 as an entry point for specimen requests. A request is specified by an investigator seeking surplus laboratory materials. An honest data broker retrieves and reviews the i2b2 phenotype with the investigator to ensure the best match of the specified inclusion/exclusion characteristics to the intended phenotype. The query is then used to match patients who have laboratory testing ordered and to deliver just-in-time surplus specimen

availability reports to laboratory honest brokers. The primary specimen source for the LBB is the MUSC Pathology and Laboratory Medicine phlebotomy operations. These specimens are available for capture after clinical testing is completed and just prior to when the specimen is normally discarded. At this point all identifiers are stripped from the physical specimen, and the specimen is no longer considered human subject-derived material. The laboratory honest brokers fulfill the requests by delivering a collection of phenotyped specimens to investigators. Thus, the final LBB deliverables include the reviewed i2b2 phenotype (query) and phenotyped specimens. Specimens are only collected when matching to a specified phenotype request and become surplus and no longer needed for provision of care.

A walk-through of the LBB user and honest broker interfaces is available as Supplementary Data.

Regulatory status of the LBB

The development of LBB proceeded under regulatory consultation with the MUSC IRB. LBB was determined to be a process and thus exempt from review. From the MUSC IRB standpoint, the phenotyped specimens do not constitute human subject-derived materials and therefore are exempt from regulations regarding informed consent. Further, the phenotypes used for specimen capture are deemed generic enough to not constitute identifiable data. LBB users are required to abide by a Data Use Agreement, which explicitly prohibits reidentification of the subjects. In conjunction with the LBB process to capture phenotyped specimens, the users may use institutional data request services to obtain linked clinical variables. Such requests require proper oversight, including IRB-approved protocols, which are reviewed by data honest brokers to determine compliance and allowable clinical data elements to be delivered. For our study, the MUSC IRB has determined that the surplus specimens delivered via LBB process comprise deidentified nonhuman subject material. However, each study would still have its own IRB review, and if the requested data or samples were deemed potentially identifiable, HIPAA would have to be addressed either via a limited data set agreement or a HIPAA waiver. This may happen, for example, if the phenotyped specimen capture occurs in such a short time span that, even if the specimen date is not explicitly provided, it may be viewed as implicitly available, thus potentially being interpreted as a limited identifier in the data.

LIVING BIOME BANK FOR PHENOTYPED MICROBIOME DATA REQUESTS

As an illustration of a practical use case of the LBB solution, we present an NIH National Center for Advancing Translational Sciences (NCATS)-funded research study protocol, the Living μBiome Bank (LμBB), developed to deliver phenotyped specimen microbiome data using LBB solution (Figure 2A). The LμBB protocol was approved
by IRB (Pro00079660) to deliver to the users processed phenotyped specimen microbiome data and an additional limited set of non-PHI clinical variables from the medical records to facilitate data interpretation. The institutional research services necessary for L\textsuperscript{μ}BB request fulfillment are managed by integration with the Service, Pricing, and Applications for Research Centers, or SPARCRequest (SPARC) system, which is an interoperable, searchable research–resource electronic storefront that supports research services operations. L\textsuperscript{μ}BB uses the LBB solution to remove the user from physical access to the specimen by capturing the phenotyped specimens and processing them to derive deliverable microbiome assay data. L\textsuperscript{μ}BB finds suitable specimens for many possible microbiome studies at MUSC Infection Control active surveillance programs. At MUSC, surveillance for methicillin-resistant \textit{Staphylococcus aureus} and vancomycin-resistant \textit{Enterococcus} generate over 15,000 tests for each pathogen annually. Recently, COVID-19 testing nasopharyngeal swabs have become available as a specimen source. The associated clinical data is available to aid in interpretation of the microbiome data and is delivered deidentified by the data honest broker.

Multi-institutional access to the L\textsuperscript{μ}BB

We implemented a model for interinstitutional requests within the L\textsuperscript{μ}BB. The working solution presented in Figure 2B uses NIH Clinical and Translational Science Awards (CTSA) Accrual to Clinical Trials (ACT) Network Shared Health Resource Information Network (SHRINE)\textsuperscript{8} to share the phenotype specification from the requesting institution (eg, George Washington University) to MUSC L\textsuperscript{μ}BB. Authentication to the LBB application is enabled by InCommon, a widely accepted authentication infrastructure across academic institutions in the USA. Non-ACT SHRINE member sites (eg, University of Iowa) can be credentialed at the LBB host site using institutional affiliate-sponsored access processes. Under the network model only the data, not the specimens, cross the institutional boundaries, which only requires a Data Use Agreement to operate.

Additional regulatory considerations for L\textsuperscript{μ}BB

L\textsuperscript{μ}BB study is granted a waiver of consent by the IRB because it is impracticable to obtain consent given that there is no way to know who would have leftover samples, and most patients would already be gone by the time the samples were pulled for research. To further mitigate ethical research concerns, L\textsuperscript{μ}BB considers (when present) the patient’s response to Research Permissions Questionnaire (RPQ) implemented by MUSC. The RPQ allows patients to indicate if they “agree” or “do not agree” to research on their surplus materials that would otherwise be discarded.\textsuperscript{9–12} By default, opt-out preferences are honored; however, since RPQ responses are available in research data warehouse (RDW), the i2b2 users may request that recruitment only happens for subjects indicating opt-in preferences. Our data indicate that among the patients who have been presented with the RPQ, 70%–75% tend to respond in the affirmative.\textsuperscript{11,13} Because of the adoption of the Common Rule, the function of the RPQ is not required for LBB implementation but is an optional additional feature.

VALIDATION, USER EXPECTATIONS, AND EXPERIENCES

We identified 21 individual investigators across MUSC, including College of Medicine, College of Dental Medicine, and College of Nursing, to serve as clinical area experts to conduct a validation focus group study, deemed by the IRB to be quality improvement and exempt. Each expert was offered an opportunity to specify their own phenotype of interest using MUSC i2b2 and asked to abstract 20 retrospective charts from each phenotype to determine if those were a “Match,” “Mismatch,” or if they were “Unsure” for any rea-
son. Nineteen investigators completed chart abstractions and an exit questionnaire.

Many investigator-defined phenotypes result in reasonable patient matches upon review

The chart match rate to the phenotype varied dramatically across experts (Figure 3). Within each clinical area, certain e-phenotypes captured the desired cohort better than others. On average, infectious, rheumatic, neonatal, and cancer e-phenotypes performed better, whereas psychiatric, gastrointestinal, skin, and pulmonary performed worse at capturing the intended cohort of the clinical experts. We speculate that the domains that did well are inpatient-focused clinical areas which naturally collect more data in the electronic health record, while the domains that did not perform as well are more outpatient-focused. Further study of the factors that contribute to successful e-phenotype specification is thus warranted and will be forthcoming from our group.

Expected specimen accrual times are reasonable and practical for pilot data collection

Using historical data, we have estimated average accrual rates for each phenotype. Most phenotypes yielded accrual rates reasonable for small-scale pilot studies. For instance, as shown in Table 1, the majority of specified phenotypes would be able to accrue a sample of 20 specimens in under 4 weeks. Only a quarter of the specified phenotypes would need more than 6 months to accrue this sample size; these represent the cases where multi-institutional collaborations are needed. Specific clinical areas where collaborations are needed include pediatric and neonatal phenotypes, rheumatic diseases, and opioid use disorders. Two e-phenotype queries (GI 11 and skin 15) specified pediatric populations, which are excluded from LyBB requests due to limitations of our current implementation of research preferences capture.

Clinical experts expect to trust scientific findings from LBB and are likely to recommend

A REDCap questionnaire has been offered to our investigator focus group to reflect on their experience using i2b2 system and their recommendations for the specimens obtained from LBB (Table 2). Responses to the questionnaire were assessed for association with chart-to-phenotype match rates in R (version 3.6.1) using logistic regression models with ordered Likert-scale factors as trends. With the study coordinator and honest broker support provided, most describe i2b2 as easy to use and would be willing to use it again to define a phenotype. Clinical experts who would be willing to use the system again had, on average, a higher match rate, with a positive linear trend (coefficient = 0.95, P value = .006). Experts generally tended to have confidence in the scientific data from LBB, while 6 of

---

Table 1. Number of e-phenotypes by expected waiting time (weeks) for accrual of 20 specimens

| Specimen | < 4 weeks | 4–26 weeks | 26–52 weeks | > 52 weeks |
|----------|-----------|------------|-------------|-----------|
| Blood    | 14        | 5          | 0           | 2         |
| Nasal    | 6         | 8          | 2           | 3         |
| Gut      | 5         | 8          | 2           | 4         |
the experts had some reservations. We also identified a positive linear trend between match rate and the level of confidence in scientific findings from LBB (coefficient = 0.55, P value = .016). Experts generally reported a willingness to recommend the system to others, with little to no reservation (n = 17). The willingness to recommend came with a positive linear relationship with chart match rate (coefficient = 0.73, P value = .005). More than half of the experts were willing to pay to obtain specimens from the system. Phenotypes specified by this group of experts had an average match rate of 77% (CI 60.7–93.7).

CONCLUSIONS AND FUTURE DIRECTIONS

LBB provides a solution for phenotyped surplus specimen capture that is open source (Ruby source code available at https://github.com/living-biobank/living-biobank) and eliminates the need for identified chart review to capture each individual specimen, which is necessary in the Harvard Crimson model. Further the LBB implementation of just-in-time capture of user-requested phenotypes does not require comprehensive biobanking of large volumes of specimens without a specific user needing them as in the BioVu model, potentially cutting the costs of human-derived research material provision. Our solution integrates processes across 2 widely adopted infrastructure systems within the CTSA community, namely SPARC and i2b2. LBB is not a replacement for a proper biorepository, which allows for full access to clinical data for consenting subjects but can be a useful auxiliary service to accelerate research, as our LmBB study demonstrates. The LBB solution access to phenotypes and phenotyped specimens simplifies regulatory considerations for the users. LmBB further extends LBB to allow for surplus clinical specimens to be processed using high-throughput molecular laboratory techniques (e.g., sequencing) to deliver deidentified phenotyped microbiome and clinical data. This allows the user to deal only with the data rather than the physical specimens. The focus groups study conducted indicated high phenotype match rates and acceptance of scientific potential of the data generated by the LBB process. Further work is needed to scale the system to allow for integrated interinstitutional specimen requests and to understand the factors and best practices in phenotyping to ensure highest possible match rates.

FUNDING

AVA, BH, LAL, JSO, KC, and JGP are supported by NIH/NCATS R21 TR002513. AVA and BH are supported by NIH/NLM R01 LM012517. SLC is supported by NIH/NCI 5P30CA138313, 1UM1CA239752, and 1U54CA210962 and by NIH/NIA 1RO1AG055132. The project described was supported by the NIH/NCATS UL1 TR001450.

AUTHOR CONTRIBUTIONS

LAL and AVA conceived the essential concepts of the manuscript and directed the research; BH and TDF contributed creatively to the implementation of the concept; KAC and JGP contributed to evaluation and network solution; and CLM, JEM, and SLC contributed to feasibility of the project from the laboratory medicine perspective. JSO oversaw implementation of essential informatics infrastructure for the project. AVA drafted the manuscript. All coauthors have revised and approved the manuscript.

### Table 2. Questionnaire captures expert reflections on just-in-time biobanking with LBB

| Question and response | N | Validation match rate (%) | Trend test | Coefficient | P value |
|-----------------------|---|---------------------------|------------|-------------|---------|
|                       |   | Mean (95% CI)             | Contrast*  |             |         |
| In the future, would you use the i2b2 system again to define a cohort? | | | | | |
| Definitely will not use again | 0 | | | | |
| Probably will not use again | 1 | 35 | | | |
| Probably will use again | 9 | 57.8 (42.1–73.5) | linear | 0.95 | 0.006 |
| Definitely will use again | 9 | 67.2 (45.1–89.4) | quadratic | −0.22 | 0.362 |
| How difficult was the use of i2b2 to define a cohort? | | | | | |
| Not at all difficult | 10 | 64.5 (44.4–84.6) | | | |
| Somewhat difficult | 5 | 71 (52–90) | | | |
| Moderately difficult | 3 | 30 (24.3–35.7) | linear | −0.22 | 0.528 |
| Very difficult | 1 | 70 | quadratic | 0.7 | 0.025 |
| Would you trust scientific results that derive from a system like this? | | | | | |
| Definitely will not trust | 0 | | | | |
| Probably will not trust | 6 | 44.2 (30–58.3) | | | |
| Probably will trust | 10 | 70.5 (19.4–107) | linear | 0.55 | 0.016 |
| Definitely will trust | 3 | 63.3 (30–58.3) | quadratic | −0.59 | 0.001 |
| Would you recommend this system to your colleagues? | | | | | |
| Definitely will not recommend | 0 | | | | |
| Probably will not recommend | 2 | 52.5 (18.2–86.8) | | | |
| Probably will recommend | 10 | 52.5 (35.4–69.6) | linear | 0.73 | 0.005 |
| Definitely will recommend | 7 | 75.7 (54.2–97.3) | quadratic | 0.42 | 0.027 |
| Would you pay to obtain specimens from such system? | | | | | |
| Definitely will not pay | 1 | 70 | | | |
| Probably will not pay | 8 | 46.2 (29–63.5) | | | |
| Probably will pay | 9 | 77.2 (60.7–93.7) | linear | −1 | 0.037 |
| Definitely will pay | 1 | 25 | quadratic | −0.66 | 0.078 |

*Only linear and quadratic trends are presented.*
SUPPLEMENTARY MATERIAL

Supplementary material is available at Journal of the American Medical Informatics Association online.

ACKNOWLEDGMENTS

The Biomedical Informatics Center personnel worked tirelessly on the LBB: John Clark, Andrew Cates, Katie Kirchoff, Ito Eta, Kayla Glick, Wenjun He, and Tami Crawford. AVA and JGP were delegates to 2017 Microbiome Data Science Innovation Labs sponsored by BD2K Training Coordinating Center. We thank Kimberly Snow for providing valuable feedback on clarity and wordmanship.

CONFLICT OF INTEREST STATEMENT

AVA is a scientific advisory board member for Second Genome, Inc., which has not contributed to this research.

REFERENCES

1. Paoli PD. Biobanking in microbiology: from sample collection to epidemiology, diagnosis and research. FEMS Microbiology Reviews 2005; 29: 897–910.
2. Murphy SN, Weber G, Mendis M, et al. Serving the enterprise and beyond with informatics for integrating biology and the bedside (i2b2). J Am Med Inform Assoc 2010; 17 (2): 124–30.
3. Murphy S, Churchill S, Bry L, et al. Instrumenting the health care enterprise for discovery research in the genomic era. Genome Res 2009; 19 (9): 1675–81.
4. Kurreeman F, Liao K, Chibnik L, et al. Genetic basis of autoantibody positive and negative rheumatoid arthritis risk in a multi-ethnic cohort derived from electronic health records. Am J Hum Genet 2011; 88 (1): 57–69.
5. Bowton E, Collier S, Wang X, et al. Phenotype-driven plasma Biobanking strategies and methods. J Pers Med 2015; 5 (2): 140–52.
6. Bowton E, Field JR, Wang S, et al. Biobanks and electronic medical records: enabling cost-effective research. Sci Transl Med 2014; 6 (234): 234cm3.
7. Sampson RR, Glenn JL, Cates AM, et al. SPARC: A multi-institutional integrated web based research management system. AMIA Jt Summits Transl Sci Proc 2013; 2013: 230.
8. Weber GM, Murphy SN, McMurry AJ, et al. The Shared Health Research Information Network (SHRINE): a prototype federated query tool for clinical data repositories. J Am Med Inform Assoc 2009; 16 (5): 624–30.
9. Obeid JS, Gerken K, Madathil KC, et al. The development of an electronic research permissions management system to enhance informed consents and capture research authorizations data. AMIA Jt Summits Transl Sci Proc 2013; 2013: 189–93.
10. Sanderson IC, Obeid JS, Madathil KC, et al. Managing clinical research permissions electronically: a novel approach to enhancing recruitment and managing consents. Clin Trials 2013; 10 (4): 604–11.
11. Marshall EA, Oates JC, Shoaibi A, et al. A population-based approach for implementing change from opt-out to opt-in research permissions. PLoS One 2017; 12 (4): e0168223.
12. Obeid JS, Shoaibi A, Oates JC, et al. Research participation preferences as expressed through a patient portal: implications of demographic characteristics. JAMIA Open 2018; 1 (2): 202–9.
13. Shoaibi A, Obeid JS, Oates JC, et al. The association between method of solicitation and patient permissions for use of surplus tissues and contact for future research. JAMIA Open 2018; 1 (2): 195–201.
14. Harris PA, Taylor R, Thielke R, et al. Research electronic data capture (REDCap)—a metadata-driven methodology and workflow process for providing translational research informatics support. J Biomed Inform 2009; 42 (2): 377–81.
15. R Core Team. R: A Language and Environment for Statistical Computing. Vienna, Austria: R Foundation for Statistical Computing, 2019.