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Scalable Collaborative Infrastructure for a Learning Healthcare System (SCILHS): Architecture

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ABSTRACT
We describe the architecture of the Patient Centered Outcomes Research Institute (PCORI) funded Scalable Collaborative Infrastructure for a Learning Healthcare System (SCILHS, http://www.SCILHS.org) clinical data research network, which leverages the $48 billion dollar federal investment in health information technology (IT) to enable a queriable semantic data model across 10 health systems covering more than 8 million patients, plugging universally into the point of care, generating evidence and discovery, and thereby enabling clinician and patient participation in research during the patient encounter. Central to the success of SCILHS is development of innovative ‘apps’ to improve PCOR research methods and capacitate point of care functions such as consent, enrollment, randomization, and outreach for patient-reported outcomes. SCILHS adapts and extends an existing national research network formed on an advanced IT infrastructure built with open source, free, modular components.

INTRODUCTION
The Scalable Collaborative Infrastructure for a Learning Healthcare System (SCILHS, pronounced ‘skills’) is one of 11 clinical data research networks (CDRNs) funded by the Patient Centered Outcomes Research Institute (PCORI) in 2014. PCORI, a non-governmental organization created under the Patient Affordable Care Act seeks to build an information technology (IT) backbone to support comparative effectiveness research at a national scale across both CDRNs and also patient powered research networks (PPRNs).

SCILHS engages patients, clinicians, health systems leadership, and key healthcare stakeholders as collaborators to build on an existing network of hospitals and health systems that have already adopted a common clinical and translational research IT and regulatory framework. SCILHS, comprising 10 health systems (box 1), is a step toward answering the Institute of Medicine’s call for a learning healthcare system (LHS)1–3 to ‘generate and apply the best evidence for the collaborative healthcare choices of each patient and provider; to drive the process of discovery as a natural outgrowth of patient care; and to ensure innovation, quality, safety, and value in health care’.

Fifteen years ago, SCILHS informatics leaders began a quest to develop informatics infrastructure and regulatory innovation that would convert the emerging electronic health record (EHR) into a research tool for improving patient outcomes. All of our work and open source toolkits have been supported by grants from the National Institutes of Health, Centers for Disease Control and Prevention, and Office of the National Coordinator of Health Information Technology (ONC). First, we built Indivo3–4 the first personally controlled health record, which gave patients their data, and apps to make those data useful. Then, i2b2 (Informatics for Integrating Biology and the Bedside)5–7 created an open source analytic platform to the EHR, to fuse and analyze data produced by the delivery system, and identify research cohorts. i2b2’s flexible common semantic data model readily accommodates a variety of clinical data. Our next advance was SHRINE (Shared Health Research Information Network),8–10 a tool enabling investigators to query i2b2 nodes in real time across multiple sites for collaborative population research. i2b2 has been successfully implemented at more than 100 sites across the USA, thereby enabling investigators to use delivery system data to identify patients with specific illnesses and clinical characteristics. A recent PCORI survey of all PCORnet sites revealed that 37% of the existing CDRN nodes and 31% of the PPRN nodes already used i2b2. Finally, we built SMART (Substitutable Medical Applications, Reusable Technologies)—a platform to enable any developer to contribute to an ‘App Store for Health and Research’ compatible with i2b2-SHRINE instances or compliant EHRs.11–13

These informatics tools and associated research policy advances have already contributed to transformation in the clinical research enterprise—real-time, collaborative population health research is now enabled across SHRINE member sites distributed nationally—but they have yet to yield substantial improvements in the health of our patients. Now, in establishing PCORnet, PCORI has catalyzed a new national research dialog to answer patient-oriented questions and improve human health. We directly address this challenge via a strategy intended to avoid prior mistakes of large-scale, top-down, costly software infrastructure efforts that
failed to scale (e.g., caBIG), instead building SCILHS with open source, free, modular components with vibrant user and software developer communities that have already spread virally to scale across heterogeneous health systems.

Here, we detail the informatics approaches taken by SCILHS to identify large cohorts of patients and engage them for research. Our technology strategy links lockstep to processes for regulatory innovation, development of robust governance constructs and policies, and local adoption by hospital leadership and institutional review boards.

**THE SIDECAR APPROACH**

SCILHS adopts and extends a strategy of establishing a freely accessible health data ‘sidecar’ warehouse to the EHR, effectively leveraging existing data collected by EHRs during routine care while avoiding costly, time-consuming EHR integrations (Figure 1). Developed intensively over the past 5 years at Harvard Medical School, this approach employs vendor agnostic, free, open source, scalable, and interoperable technologies to produce the only research-based, shared repository of EHR data that can be queried in real-time. Of already proven value in the research ecosystem, these components support a cost-effective and sustainable research network of >8 million patients.

We consider the heterogeneity of collaborating institutions to be a key measure of success; via adoption of the sidecar approach, we enable any institution to join our SCILHS network. Specifically, a primary goal is inclusion of diverse populations within our CDRN network, thereby enabling capture of the genetic, genomic, and socioeconomic variation that exists beyond insured populations in managed care settings alone. Further, by freely sharing the processes and software that have been developed and supported by Harvard, we hope to catalyze the formation of many other new networks across heterogeneous health systems and institutions, and involve new

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**Box 1 Alphabetical list of Scalable Collaborative Infrastructure for a Learning Healthcare System (SCILHS) sites**

- Beth Israel Deaconess Medical Center
- Boston Children’s Hospital
- Boston Health Net (Boston Medical Center and Community Health Centers)
- Cincinnati Children’s Hospital Medical Center
- Columbia University Medical Center and New York Presbyterian Hospital
- Morehouse School of Medicine/Grady Memorial Hospital (Research Centers in Minority Institutions)
- Partners HealthCare System (includes Massachusetts General and Brigham & Women’s Hospital)
- University Mississippi Medical Center
- The University of Texas Health Science Center at Houston
- Wake Forest Baptist Medical Center

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**Figure 1** Each site will install the Scalable Collaborative Infrastructure for a Learning Healthcare System (SCILHS) Sidecar, for identifying and reviewing cohorts and the mySCILHS suite to: (a) manage linkage of contact data to the de-identified Patient Cohort list produced by the multisite Shared Health Research Information Network (SHRINE) query; (b) administer and store consent documents; (c) outreach to patients through web-based survey and telephony; and (d) promote ongoing patient engagement through outgoing messaging, including (in the future) return of research results to patients. The web-based survey will be administered using REDCap and Indivo technologies, and will be accessible either by patients at home, or at the point of care, through tablet/kiosk-based interaction. Once completed, patient-reported data will have subject identifiers encoded; its standardized survey metadata will then be loaded into the corresponding SCILHS sidecar (i2b2 node), enabling semantic data linkage with electronic health record data via SHRINE/i2b2, while preserving subject confidentiality. These software platforms will be provided to sites as self-contained, pre-configured virtual machines, enabling rapid dissemination of these technologies while minimizing administrative and software development overhead at each site.
partners in improving our core components, common data models, and ontologies.

The sidecar infrastructure is composed of the following:

- **i2b2** (Informatics for Integrating Biology and the Bedside). Data analytic platform employed for EHR data analytics and clinical research at >100 academic medical centers worldwide (NIH funded).

- **SHRINE** (Shared Health Research Informatics Network). Federated query and response system that enables investigators to discover EHR data housed in i2b2 nodes across multiple independent institutions (NIH CTSA funded).

- **SMART Platforms**. First described in the *New England Journal of Medicine*, SMART has programmatic interfaces and applications that transform both EHRs and their sidecars into platforms that run substitutable iPhone-like apps. SMART enables a national scale ‘App Store’ for PCOR for rapid cycle innovation of PCOR methods (ONC funded).

- **Indivo**. The original personally controlled health record links patients to clinical and research settings. Used by hundreds of thousands of individuals, Indivo was also the initial software codebase for Microsoft’s HealthVault platform (NIH, CDC, and ONC funded).

- **REDCap** (Research Electronic Data Capture). Electronic data capture tool with 757 institutional partners, used to survey patients online (NIH CTSA funded).

### DATA MODELS AND ONTOLOGIES

SCILHS will combine EHR data with payer claims to facilitate longitudinal tracking of patients over time and across sites of care. The sidecar approach provides the capability to implement new data models without transforming all of the stored source data—a key element in the scalability and interoperability of our platform (table 1). By enabling well-designed, cross-mapped ontologies that support a PCORnet common data model, this approach incorporates otherwise disparate clinical data sources into an easily-queried system that stores data in a flexible format. Data are stored in i2b2 using an entity–attribute–value model, employing a central ‘fact’ table based upon Kimball’s *Star Schema* wherein each row stores a flexibly defined, atomic ‘fact’ or observation for a patient. Much of i2b2’s versatility arises from its focus on a semantic definition of patient observations that can represent various existing and newly defined data elements: claims, EHR, genetic and imaging data, as well as patient reported outcomes and demographics. Analogous to a capacious warehouse with adjustable shelves and bins, i2b2 accommodates various nomenclatures for data elements, and supports robust tags of associated modifiers and values. This approach enables database indexing of facts and observations to support high performance execution of expressive queries and filters.

**Table 1** Approaches to scalability and interoperability

<table>
<thead>
<tr>
<th>Sidecar approach</th>
<th>‘Community-extensible ontologies’</th>
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<tbody>
<tr>
<td>EHR data are managed in a sidecar, readily established at any institution, regardless of EHR vendor product (Epic, etc)</td>
<td>All schemas and ontologies we produce are open source, free, and already widely adopted</td>
</tr>
<tr>
<td>i2b2 uses a simple data model (Star Schema) greatly simplifying the Extract, Transform, and Load procedure. These ETL procedures are established for all major EHR products</td>
<td>Ontologies can be imposed on the data after the fact, enabling a hospital in our network to readily adapt to any ratified PCORI Common Data Model</td>
</tr>
<tr>
<td>i2b2 employs an ontology-based approach that supports flexible, on-the-fly incorporation of new data elements and coding systems. Terminologies such as ICD, NDC, and LOINC may be pre-loaded as hierarchical concept trees; new or ad-hoc terminologies including patient-reported outcome measures or locally defined data dictionaries readily coexist and may be cross-mapped in i2b2. Concepts may be grouped using simple hierarchies and then optionally re-mapped into other reference coding systems and data models (eg, Observational Medical Outcomes Partnership (OMOP) data model). In this way, i2b2 accommodates diverse real-world coding systems while maintaining a straightforward query interface for its users. The SHRINE Adaptor Cell maps local i2b2 terminologies into a common, standards-based SHRINE ontology. This enables a common shared ontology for federated queries while allowing individual i2b2 instances within institutions to retain local hierarchies and terminologies. The Adaptor transforms a federated SHRINE query into a query that runs on the local i2b2 database. The Adaptor then converts the result of that query back into the common SHRINE message format, using well-maintained standards including RxNorm, ICD9, and LOINC. In addition, SHRINE includes tools for ontology mapping and ontology-based data mining. Simple SHRINE customizations enable use of other query systems, for example the QueryHealth distributed query system (ONC) uses PopMedNet to query i2b2.</td>
<td></td>
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**SUCCESS TO DATE**

SHRINE and i2b2-based research includes characterization of rare morbidities of common diseases, very rare diseases such as peri-partum cardiomyopathy (discovered in SHRINE and published in *Nature*), detections of drug–drug interactions, and measures of quality and clinical efficacy across self-organized SHRINE networks in Europe, the University of California healthcare systems, and a just-in-time network to study the prevalence of complication rates of type 1 and type 2 diabetes in hospitals across this country. Others have used SHRINE to characterize and track the rising incidence of colorectal cancer and further characterize it, and to identify and optimize practice variation in inflammatory bowel disease and intervene to change that practice. i2b2 and SHRINE have been implemented as the base infrastructure for a variety of enhanced chronic disease registry-based research efforts. The Childhood Arthritis and Rheumatology Research Alliance uses the SHRINE/i2b2 registry framework to federate clinical care data and patient-reported data from 62 academic medical centers in the USA and Canada and is currently piloting consensus treatment protocol trials. The Harvard Inflammatory Bowel Disease (IBD) Longitudinal Data Repository employs the same infrastructure and utilizes i2b2 as its centralized data warehouse for IBD-related quality improvement development at 50 centers.
**PATIENT ENGAGEMENT**

The health systems that have joined SCILHS reflect the American demographic—an essential requirement for reaching statistically valid, clinically meaningful, and patient-centric conclusions about therapies across the diverse spectrum of all healthcare consumers. In order to achieve the comprehensive, patient-centered outcomes infrastructure called for by PCORI, we introduce a new, patient-centric platform (mySCILHS) based on the Indivo system and incorporating the REDCap electronic data capture tool.

mySCILHS will support the Blue Button REST API for standards-based interactions with PPRNs and other patient-selected tools. This API exposes up-to-date, structured clinical summary data for each participating patient. Via a consumer-friendly workflow based on web standards including OAuth2, patients can authorize third-party apps and services, including PPRNs, to access their clinical data.

**ANTICIPATED WORKFLOW**

Figure 2 shows the workflow from an initial query through the analytic phase in a comparative effectiveness study. Each node in the network maintains an instance of i2b2 containing claims and de-identified electronic medical record data. SCILHS is a true peer-to-peer network, meaning that any SHRINE-based node can initiate a query, using a common ontology, that aggregates results from all participating sites. After the initial query, the investigator can automatically pass the query to each site where duly authorized local site investigators may review individual subject data for study eligibility using i2b2 SMART apps (figure 3). The final patient list is transmitted to the mySCILHS.
IMPLEMENTING AND SCALING
SCILHS includes 10 legally and financially independent institutions whose CEO or equivalent senior institutional official has committed to active participation in governance, policy development, data sharing, and sustainability planning. Each member has pledged to invest additional personnel and resources to ensure the network meets local patient and clinical stakeholder needs. By harmonizing informatics infrastructure, data models, regulatory processes and policies, and patient participation within and across member institutions, we anticipate that SCILHS will achieve and remain a successful model for inter-institutional PCOR. Utilizing the innovative SCILHS sidecar IT approach to EHR access, we minimize local informatics burden, further enabling a sustainable and adaptable PCOR infrastructure.

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Collaborators SCILHS Network.

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accuracy or integrity of any part of the work are appropriately investigated and resolved.

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