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Authors

Sim, Ida
Carini, Simona
Tu, Samson
[et al.](#)

Publication Date

2010-03-01

Peer reviewed

The Human Studies Database Project: Federating Human Studies Design Data Using the Ontology of Clinical Research

Ida Sim, MD, PhD¹, Simona Carini, MA¹, Samson Tu, MS², Rob Wynden¹, Brad H. Pollock, PhD³, Shamim A. Mollah, MA⁴, Davera Gabriel, RN⁵, Herbert K. Hagler, PhD⁶, Richard H. Scheuermann, PhD⁶, Harold P. Lehmann, MD, PhD⁷, Knut M. Wittkowski, PhD⁴, Meredith Nahm, MS⁸, Suzanne Bakken, RN, DNSc⁹

¹Univ. of California San Francisco, CA; ²Stanford University, Stanford, CA; ³Univ. of Texas Health Science Center, San Antonio, TX; ⁴The Rockefeller University, New York, NY; ⁵Univ. of California Davis, CA; ⁶UT Southwestern Medical Center, TX; ⁷Johns Hopkins University, Baltimore, MD; ⁸Duke University, Durham, NC; ⁹Columbia University, New York, NY

Abstract

Human studies, encompassing interventional and observational studies, are the most important source of evidence for advancing our understanding of health, disease, and treatment options. To promote discovery, the design and results of these studies should be made machine-readable for large-scale data mining, synthesis, and re-analysis. The Human Studies Database Project aims to define and implement an informatics infrastructure for institutions to share the design of their human studies. We have developed the Ontology of Clinical Research (OCRe) to model study features such as design type, interventions, and outcomes to support scientific query and analysis. We are using OCRe as the reference semantics for federated data sharing of human studies over caGrid, and are piloting this implementation with several Clinical and Translational Science Award (CTSA) institutions.

Introduction

Human studies are one of the most central and valuable activities in biomedical research. Study designs and results should be made machine-readable to facilitate large-scale data mining and synthesis.

The Human Studies Database (HSDB) Project is a consortium of research institutions that is developing semantic and data sharing technologies to federate descriptions of human studies design over caGrid. In this paper, we describe 1) our use cases; 2) the Ontology of Clinical Research (OCRe), a rich model of human study designs; 3) our HSDBgrid data sharing architecture incorporating i2b2 and caGrid technologies; and 4) early results on using OCRe as the semantic standard for sharing over HSDBgrid.

Overview

There is a growing interest in sharing raw clinical research data to facilitate science and to promote

transparency and accountability (1, 2). Because of competing regulatory and intellectual property concerns, it is unlikely that such sharing will be accomplished by aggregating all data into a single database. Instead, the most feasible data sharing approach is to “federate” queries over locally controlled databases that are standardized to a common model of clinical research.

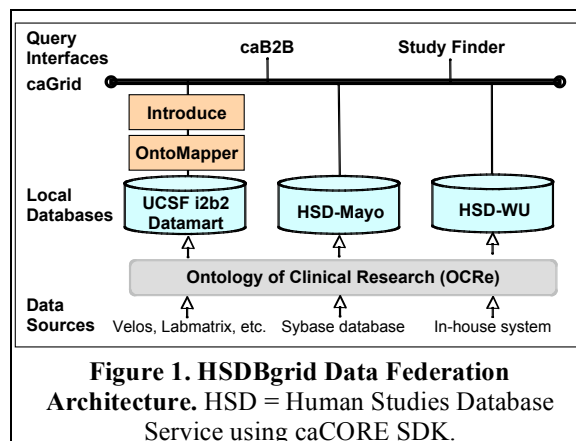


Figure 1 illustrates the HSDBgrid architecture for federating human studies databases. OCRe serves as the common semantic model, which defines the concepts that can be queried over the individual databases. The content and richness of HSDBgrid queries is therefore critically dependent on OCRe. For example, if OCRe does not include the concept of primary outcome, then HSDBgrid cannot support queries about primary outcomes.

It is especially important that OCRe be a rich model of human study designs, because the use and interpretation of study data depends critically on the context in which those data were collected. For example, data from a trial enrolling only patients with advanced breast cancer will not be representative of breast cancer patients in general. Similarly, a diabetes study that excludes patients with heart disease is non-representative of diabetes patients in general. Studies

may even include subjects who do not have the condition of interest: e.g., a study with a non-specific case definition, or a study with healthy volunteers.

Effective sharing of clinical research data therefore requires sharing study design metadata as well as results data. OCRE is a model of human studies design and results data that can serve as a common semantic for data sharing. The HSDB Project's initial goal is to share study design information (e.g., study design type, eligibility criteria, outcomes) among CTSA and other institutions as a prelude to the more complicated task of sharing results data.

Use Cases for Sharing Human Studies Data

Shared human studies data have two broad uses: 1) for researchers, to inform the design of new studies and to aggregate and analyze existing data for new findings; and 2) for research administrators, to inform the optimization of research oversight and processes. We canvassed researchers and administrators from six CTSA institutions to describe and prioritize their needs for shared human studies data (3). The top three priority needs were 1) research characterization (e.g., of population characteristics, outcome variables); 2) registration of studies into ClinicalTrials.gov; and 3) facilitating research collaborations. We broadened research characterization to cover scientific query and analysis in general, and adopted that as our target need for HSDB. For example, Dr. A, a researcher, seeks data on the prevalence of asthma in school-aged children to inform the design of a new study. These needs would not be met by searching PubMed or ClinicalTrials.gov, because studies may have collected relevant data without that collection being mentioned. In contrast, HSDB would meet Dr. A's needs by supporting queries of key study features standardized across large numbers of human studies of varied design. But what kind of queries would Dr. A submit to HSDB? And what modeling is needed in OCRE to support those queries?

To determine whether a study's asthma prevalence data is relevant to her needs, Dr. A must first identify studies enrolling school-aged children. Next, she needs to select studies whose design types are suitable for assessing prevalence (e.g., observational cohort and cross-sectional studies). She then needs to examine how individual studies specified the phenomenon of asthma (e.g., extrinsic asthma, status asthmaticus), how the phenomenon was represented as study variables (e.g., peak flow, billing code), how and when these variables were measured, and whether any study interventions might have increased or decreased the reported prevalence of asthma. She

will also want to adjust for clinically relevant covariates (e.g., air quality) and will want to know if they were measured. This use case illustrates the depth to which OCRE must model eligibility criteria, study design types, study outcomes and variables, and study exposures. This scientific depth of modeling is not present in existing clinical research models (e.g., BRIDG, CDISC SDTM) that serve primarily operational and administrative needs.

Ontology of Clinical Research

OCRe is an OWL 1.1 ontology that focuses on the design and analysis of human studies. Its scope includes human investigations of any design type (e.g., interventional, observational) for any intent (e.g., therapeutic, diagnostic, preventive) in any clinical domain on any type of data (e.g., clinical, imaging, genomics). OCRE includes 1) a representation of the structure of human studies and associated entities, 2) informational entities (e.g., study protocols), 3) terms for describing study characteristics, and 4) bindings to standard terminologies (e.g., SNOMED CT).

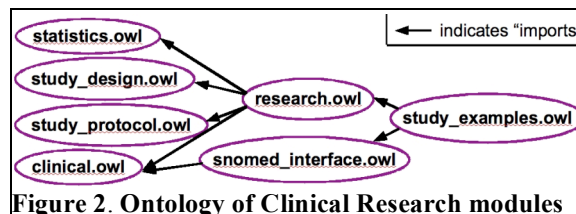


Figure 2. Ontology of Clinical Research modules

OCRe is organized as a set of modular components related by their import relationship (Figure 2). The *research* module imports the *clinical*, *study_design*, *statistics*, and *study_protocol* modules to describe a study. The *study_protocol* module imports from the BRIDG model (4) terms that specify temporal aggregates (e.g., epochs and arms) and sequencing relationships among protocol-driven activities.

OCRe modules are independent of any clinical domain because the clinical content is expressed through external ontologies and terminologies such as NCI Thesaurus or SNOMED-CT. OCRE interfaces to these terminologies by relating OCRE entities (e.g., outcome phenomenon) to these external concepts (e.g., acute myocardial infarction) and their associated terminology codes (e.g., SNOMED-CT code for acute myocardial infarction).

In the next sections, we discuss OCRE's modeling of several key domains of clinical research.

Study Design Typology

We postulated that there exist a small number of high-level study design types that represent distinct

approaches to human investigations, and that we could reliably classify all human studies into these design types. Since each study type is subject to a distinct set of biases and interpretive pitfalls, a study's design type would strongly inform the interpretation and reuse of its data and biosamples.

Through iterative consultation with statisticians and epidemiologists, we defined a typology of study designs based on discriminating factors that define mutually exclusive and exhaustive study types (hybrid studies can be of more than one type). We use these factors as questions in a web-based classification tool (5). Our tool first classifies studies into human and non-human studies (Does the study use or collect measurements, assessments or observations about individual humans?). It then classifies human studies into qualitative or quantitative studies, and subsequently classifies quantitative studies into four interventional or four observational high-level design types (in red in Figures 3 and 4).

For interventional studies (Figure 3), discriminating factors include whether the investigator has a choice of interventions to which s/he can assign participants, whether the main comparison is within or across participants, and whether intervention assignment and data analysis are only within a single participant. Additional descriptors elaborate on secondary design features (e.g., randomization, blinding) that introduce or mitigate additional interpretive concerns.

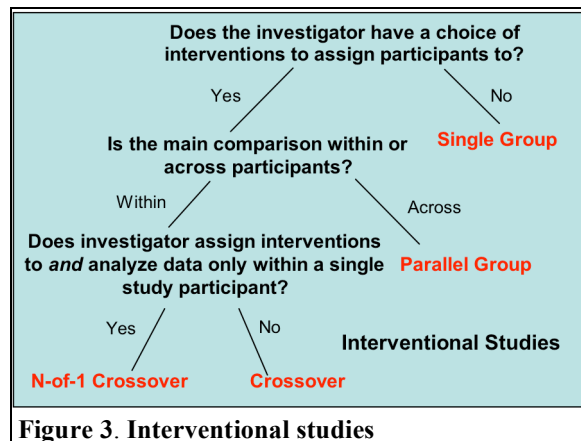


Figure 3. Interventional studies

For observational studies (Figure 4), the four design types are based on whether the main control group is defined by case (outcome) or exposure (predictor) status, whether the case and control are in the same person, and whether outcomes are measured at the same time as predictors or after. Additional descriptors other than the ones for interventional studies apply to these observational study types (e.g., retrospective or prospective). The design typology is formalized in OCRE as an OWL hierarchy.

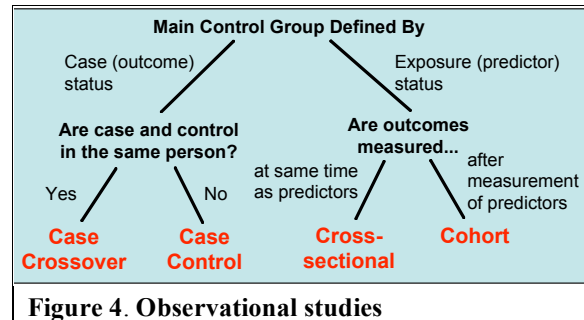


Figure 4. Observational studies

Eligibility Criteria

OCRe uses Eligibility Rule Grammar and Ontology (ERGO) Annotation (6) to capture the clinical content of eligibility criteria in machine-readable form. ERGO Annotation is a declarative representation of eligibility criteria that is informed by both the complexity of natural language and the requirements for computability. ERGO Annotation models three statement types: 1) simple statements making single assertions, 2) statements about quantitative comparisons, and 3) complex statements, which are simple and/or comparison statements joined by Boolean connectives or semantic connectors (e.g., evidenced_by).

Study Outcomes and Analyses

In OCRe, the study protocol specifies the study activities to achieve the study's scientific objectives, such as the collection and analysis of study data. Figure 5 shows our conceptualization of the entities related to outcomes and analyses in human research. We first define a study phenomenon as "a fact or event of interest susceptible to description and explanation." Study phenomena are represented by one or more specific study variables that may be derived from other variables. For example, the study phenomenon of cardiovascular morbidity may be represented as a composite variable derived from cardiovascular death, myocardial infarction (MI), and stroke variables. Each variable can be further described by its type (e.g., dichotomous), coding (e.g., death or not), timepoints of assessment (e.g., 6 months after index MI), and assessment method (e.g., death certificate). All variables are associated with participant-level and study-level observations (observations aggregated across subjects).

A study protocol may specify several analyses, each having dependent and independent variables that represent various study phenomena. Variables may play the role of dependent or independent variables in different analyses. If the study protocol designates a primary analysis, the dependent variable of that analysis represents what is conventionally known as the primary outcome of the study. To our knowledge,

first end-to-end use of ontologies to share semantically standardized data over caGrid.

Discussion

The HSDB project is a multi-institutional collaboration that has made substantial technical progress towards integrating human studies design data to address high priority scientific query and analysis needs. Our approach uses OCRE, a semantic model of human studies design and analysis, as the common semantics for interoperating local caBIG and i2b2 human studies databases over caGrid.

There are several challenges facing this project and caGrid data sharing in general. One challenge is to more fully understand the respective roles of OWL ontologies and UML models in federated data sharing, and to use or develop appropriate caGrid and other technologies consonant with these roles. We are exploring using SPARQL views of OCRE to generate reproducible mappings between OCRE and HSDB UML models. This mechanism will be important for propagating OCRE updates to UML models.

A second challenge is data acquisition. How will disparate human studies design data be gathered and aggregated from study protocol documents, ethics board applications, and clinical research management systems (CRMSs) into HSDB repositories? Presently, the process is entirely manual, but increasing automation will be possible if ethics boards and CRMSs begin to adopt OCRE's conceptualization and definitions of human research. We also need to continue our early efforts at harmonizing OCRE with BRIDG (and thus to HL7's Clinical Trials Registry and Results Project), to Open Biomedical Ontologies, and to other data sharing and reporting initiatives (e.g., FDA Amendments Act of 2007, NCI's Clinical Trials Reporting Program, NIH's Data Sharing Initiative). A multi-institutional international human studies database will be an incomparably rich resource for clinical and translational research.

Acknowledgements

Funded in part by R01-LM-06780 and by UL1RR024156, UL1RR024128, UL1RR02500, UL1RR024143, UL1RR024146, UL1RR024131, UL1RR025767, and UL1RR024982 from the National Center for Research Resources (NCRR), a component of the National Institutes of Health (NIH) and NIH Roadmap for Medical Research. Its contents are solely the responsibility of the authors and do not necessarily represent the official view of NCRR or NIH. We thank Shannon Hastings and Rakesh Dhaval from OSU for their technical contributions.

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