Seminar on Standardization of Laboratory Data with LOINC
Hong Kong Hospital Authority



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Regenstrief Institute

Medical Informatics

Overview

- 1. Origins and Evolution of LOINC
- 2. LOINC Introduction
- 3. Mapping Tools and Resources
- 4. Closing Thoughts

Regenstrief Institute

established 1969

Biomedical Informatics
Health Services Research
Aging Research

Biomedical Informatics at Regenstrief Institute

Building data-rich systems for clinicians



McDonald et al. Canopy Computing: using the Web in Clinical Practice. JAMA. 1998;280(15):1325-1329.

Individual healthcare computer systems are rich with patient data, but rather than a canopy linking all the trees in the forest, the data "fruit" come from a diverse forest of individual computer "trees".



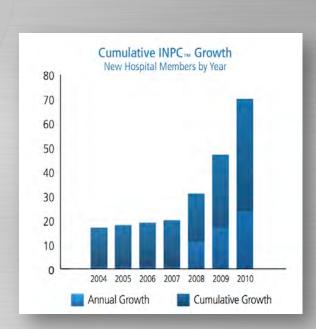
Regenstrief Biomedical Informatics

40 years of EMR work Indiana Network for Patient Care

Nation's most comprehensive and longest tenured HIE

Regenstrief

- the neutral 3rd party convener



A "Humongous" Database

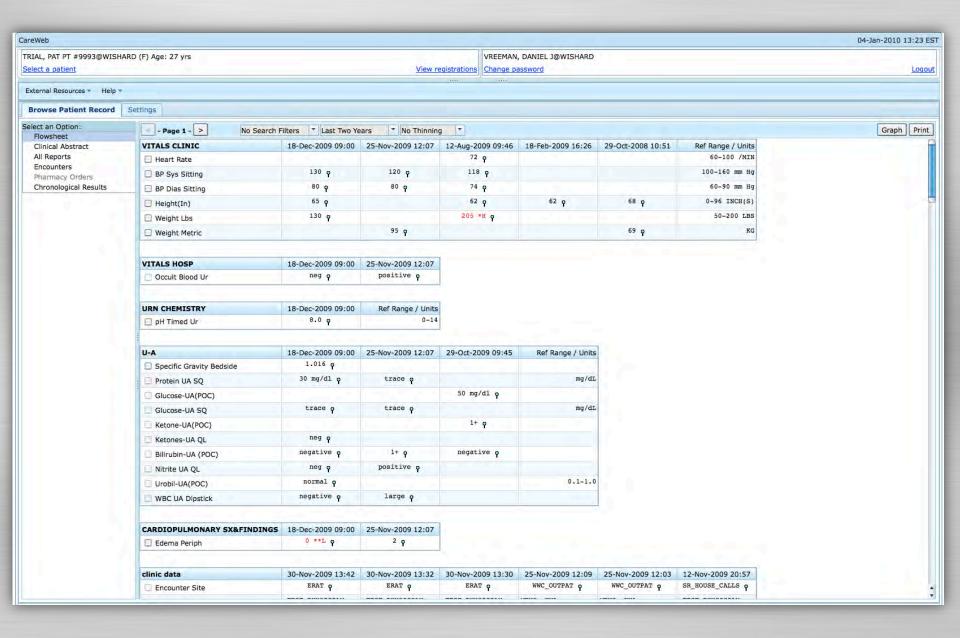
- 200+ source systems
- 12.1 million patients
- 4.1 billion results
- 79.4 million text reports

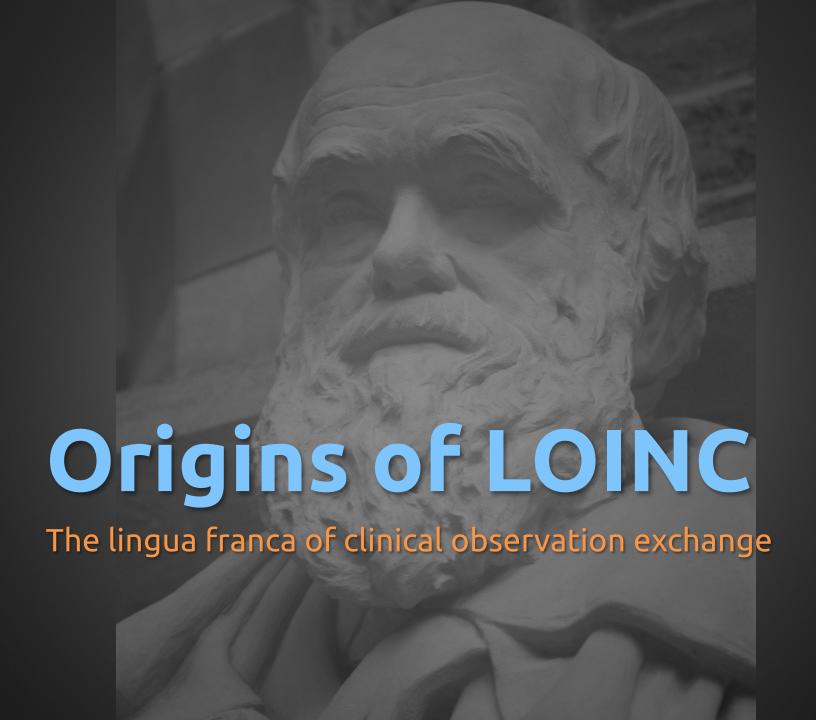
1 million transactions per day

Indiana Network for Patient Care

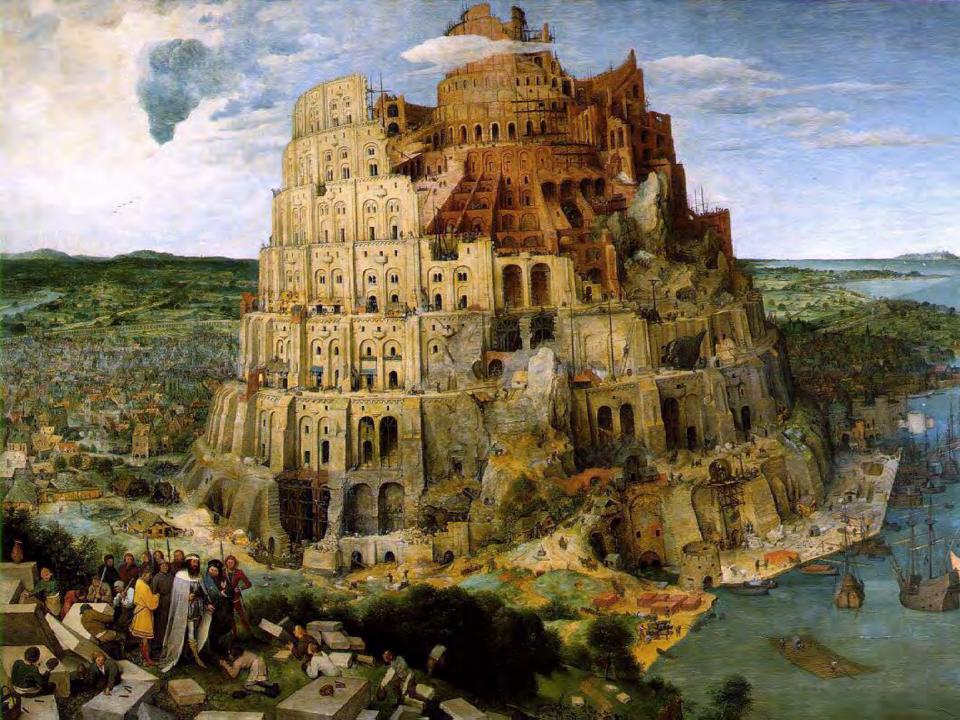


A Unified Patient Record









Fundamental challenge: local systems have different ways of identifying the same concept

A vocabulary standard would serve as the *Rosetta Stone*.

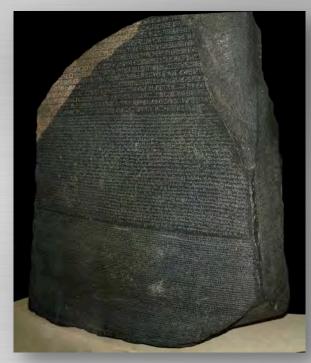


Image © Hans Hillewaert / CC-BY-SA-3.0

Logical Observation dentifiers lames and Codes

A universal code system that facilitates exchange, pooling, and processing of results

Established in 1994 by Regenstrief Institute.

Vocabulary standard for observation identifiers.



Same or Different?

what you see in the order list

Lab A

Test Name: Lyme Disease Serology

Measures: B. burgdorferi Ab IgG

Method: ELISA

Scale: quantitative

e.g.: Titer 1:40

LOINC Code = 5062-5

Lab B

Test Name: Lyme Disease Antibody

Measures: B. burgdorferi Ab IgM

Method: Immune blot

Scale: qualitative

e.g.: Positive

LOINC Code = 6321-4

Similar name, different meaning...



photo via Caption Time

meerkat meerkat mere cat meerkat

If an observation is a question and the observation value is an answer...

LOINC provides codes for questions

Other vocabularies provide codes for the *answers*

What is my patient's hemoglobin level?



718-7:Hemoglobin:MCnc:Pt:Bld:Qn

How fast does my patient usually walk?

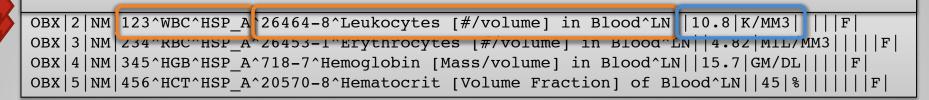


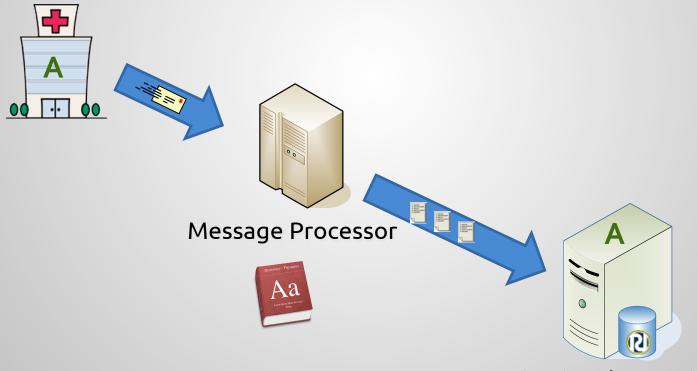
41959-8: Walking speed: Vel: 1W^mean: Patient: Qn: Calculated

Indiana Network for Patient Care



LocalCode^LocalName^CodeSystem LOINCcode^LOINCname^CodeSystem





Institutional Repository

Result with a Coded Value

Data type of result (OBX-5) is a coded element

This code is from LOINC

This code is from SNOMED

OBX||CE|6609-2^Listeria ID^LN||36094007^L. monocytogenes^SCT

Code identifying this observation (what are these results? Listeria culture)

Code identifying the result (L. monocytogenes)

Laboratory LOINC

Microbiology Tests

Challenge chemistry tests

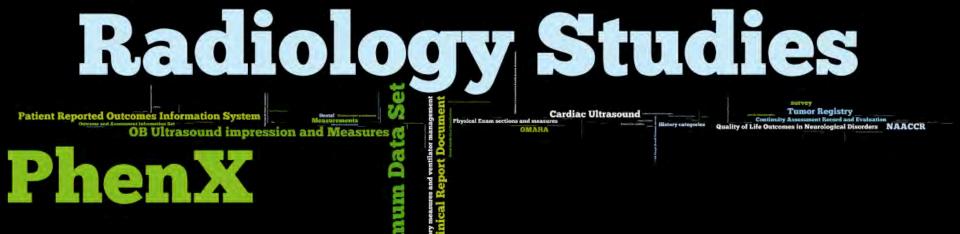
Chemistry non challenge tests

Allergy Testing

Ceel Markers Mutations

Drug toxicology tests

Clinical LOINC



Laboratory LOINC Committee Chair: Clem McDonald, MD



Clinical LOINC Committee Chair:

Stan Huff, MD







International Classification of Diseases (ICD)

and the rest of the family

RXNorm

normalized names for clinical drugs

The LOINC Community

Open. Nimble. Pragmatic.

Open Development

Worldwide distribution at no cost End-user content additions
Welcome all comers
Volunteers



Funding Support

Principal current sources are:
U.S. National Library of Medicine
Regenstrief Foundation

Prior support from

Several other U.S. federal agencies, John A. Hartford Foundation

Regenstrief Institute

Steward

Developer of content

Developer of tools

Developer of community

Distributor

Voice



log in new user?

you are here: home

Logical Observation Identifiers Names and Codes (LOINC®)

A universal code system for identifying laboratory and clinical observations.

From serum levels of hepatitis B surface antigen to diastolic blood pressure, LOINC has standardized terms for all kinds of observations and measurements that enable exchange and aggregation of electronic health data from many independent systems.

More than 14,000 people in 145 countries use LOINC to help make bridges across their islands of health data.

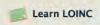
It's free, but invaluable. Both LOINC and the RELMA mapping program that helps link your local codes to LOINC terms are distributed at no cost by the Regenstrief Institute. LOINC is your key to interoperable data exchange.

How do **you** say glucose?



Ready to get started?

Or, search LOINC with our online app



Background FAO Users Guide Presentations/Tutorials



Use LOINC

Download LOINC Map to LOINC Search LOINC "Best of" LOINC



Get Involved

Forum Meetings Mailing Lists Directory of Adopters



Develop LOINC

Submit Term Requests What's Coming Translate LOINC



LOINC 2.38 Released: 2011-12-30

RELMA 5.5

Released: 2011-12-30

Download Now ----

Recent Forum Posts

Microbiology | Re: Microbiology Text results and LOINC

Microbiology | Re: Organism Names and LOINC?

More posts...



LOINC

LOINC New paper published in IBI about helping map to #LOINC with RELMA by augmenting local test names http://t.co /plwkalfM about 1 hour ago - reply - retweet favorite

Join the conversation



Copyright @ 1994-2012 by the Regenstrief Institute, Inc. Read the full LOINC® and RELMA® Terms of Use

Content

oinc.org stats: 14697 users from 145 countries

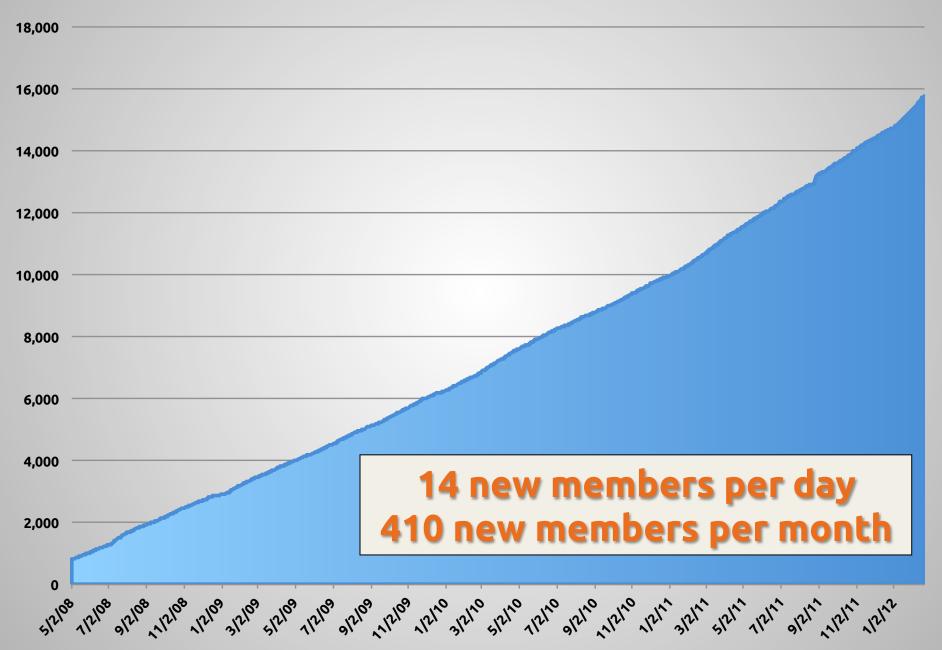
Downloads LOINC RELMA Accessory Files

LOINC Usage Value Sets Newborn Screening Document Ontology

Documentation

LOINC Users' Guide Recommended Readings Presentations/Tutorials

loinc.org members



15,800+ users in 145 countries



The LOINC Distribution

LOINC and RELMA Complete Package



DINC and RELMA Complete Download File (All Formats Included) (216.1 MB)

File Version: LOINC 2.34 and RELMA 5.1 | Release Date: 2011-02-17 | File type: application/zip

This file contains the RELMA® program installer (which also includes the LOINC® and RELMA® Users' Guides), and the LOINC® database in both MS Access™ and tab-delimited formats.

To download this file you need to log in with your user name and password. If you do not have an account here, head over to the registration form.

LOINC Table



LOINC 2.34 Released: 2010-12-29

The LOINC table with fields for LOINC codes, each of the six parts of the formal name of the LOINC, synonyms, comments, and other information. Distributed as a tab-delimmited file and Access database, and a release to release change file. Documentation includes LOINC Release Notes and a the comprehensive LOINC User's Guide.

Download Now Ifree -

Accessory Files



LOINC 2.34

Released: 2010-12-29

Additional files available in the LOINC distribution.

- LOINC Panels and Forms File
- LOINC Multiaxial Hierarchy File
- LOINC Context-specific Hierarchy Template File

Download Now Ifree |

RELMA



RELMA 5.1

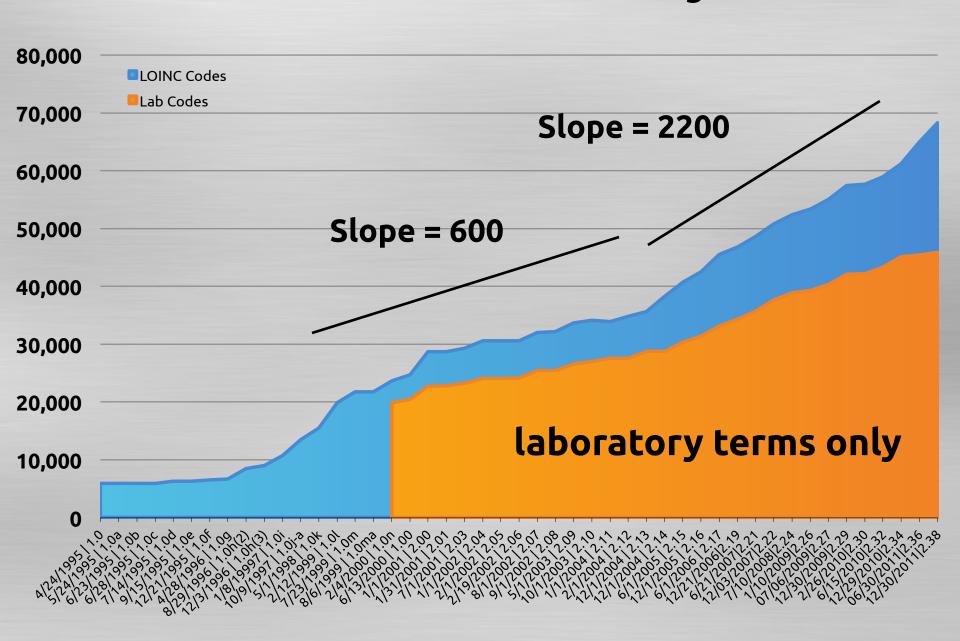
Released: 2011-02-17

Regenstrief LOINC Mapping Assistant (RELMA®) is a Windows program for searching the LOINC database and helping you map local codes to LOINC codes. Documentation includes <u>RELMA Release Notes</u> and a comprehensive RELMA User's Manual.

Download Now Irree

Major releases twice per year (June and December)

LOINC Codes Over Time By Release





loinc.org/terms-of-use

No cost Worldwide In perpetuity





Cannot use any Licensed Material to develop or promulgate a different standard for orders or observations.

That would defeat the purpose of having a standard!



LOINC Submitters



Since 2009
79 organizations from 14 countries

LOINC Translators



19 organizations

How do you say glucose?



the *lingua franca* of clinical observation exchange















Γλυκόζη Glucosio 포도당 Glicose

Glucosa

Glucosa

Glucosa

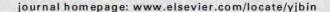
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Enabling international adoption of LOINC through translation

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Computerized/standards

ABSTRACT

Interoperable health information exchange depends on adoption of terminology standards, but international use of such standards can be challenging because of language differences between local concept names and the standard terminology. To address this important barrier, we describe the evolution of an efficient process for constructing translations of LOINC terms names, the foreign language functions in RELMA, and the current state of translations in LOINC. We also present the development of the Italian translation to illustrate how translation is enabling adoption in international contexts. We built a tool that finds the unique list of LOINC Parts that make up a given set of LOINC terms. This list enables translation of smaller pieces like the core component "hepatitis c virus" separately from all the suffixes that could appear with it, such "Ab.IgG", "DNA", and "RNA". We built another tool that generates a translation of a full LOINC name from all of these atomic pieces. As of version 2.36 (June 2011), LOINC terms have been translated into nine languages from 15 linguistic variants other than its native English. The five largest linguistic variants have all used the Part-based translation mechanism. However, even with efficient tools and processes, translation of standard terminology is a complex undertaking. Two of the prominent linguistic challenges that translators have faced include: the approach to handling acronyms and abbreviations, and the differences in linguistic syntax (e.g. word order) between languages. LOINC's open and customizable approach has enabled many different groups to create translations that met their needs and matched their resources. Distributing the standard and its many language translations at no cost worldwide accelerates LOINC adoption globally, and is an important enabler of interoperable health information exchange.

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In-progress

Catalan

Dutch

French

Russian

Turkish











Adopted as National Standard

Brazil

Canada

France

Germany

The Netherlands

Mexico

Rwanda



Large Implementations

SIGA Saúde project
Canada Health Infoway
ePSOS
Assistance publique - Hôpitaux de Paris
Red Agrolab

BITAC

Many more...

HL7 and Regenstrief Institute Sign Statement of Understanding

November 14, 2011

INDIANAPOLIS — Health Level Seven® (HL7®) International, the global authority on standards for interoperability of health information technology with members in 55 countries, and the Regenstrief Institute, Inc., an internationally respected healthcare and informatics research organization, today announced an agreement to create a complementary process to develop and extend comprehensive standards in the healthcare industry.

"This agreement further solidifies and extends the wonderful relationship HL7 has enjoyed with Regenstrief for many years," said Bob Dolin, chair of HL7 Board of Directors. "HL7 is committed to working with Regenstrief and other standards bodies to advance the delivery of safe and effective patient care."

Logical Observation Identifiers Names and Codes (LOINC®) is a universal code system developed by the Regenstrief Institute for identifying laboratory and clinical observations. When used in conjunction with the data exchange standards developed by HL7, LOINC's universal observation identifiers make it possible to combine test results, measurements, and other observations from many independent sources. Together, they facilitate exchange and pooling of health data for clinical care, research, outcomes management, and other purposes.

"Regenstrief has been a long-standing contributor to the standards developed by HL7, and likewise, LOINC has been enhanced by its adoption in HL7's standards," said Daniel Vreeman, DPT, M.Sc., associate director of terminology services at the Regenstrief Institute. "With this agreement, we look forward to an even closer collaboration with HL7 that improves the semantic interoperability of health data exchange worldwide."

LOINC began in the mid 1990's when Regenstrief investigators, using their decades of experience with electronic medical records, began the Indiana Network for Patient Care, the nation's first citywide health information exchange. The researcher clinicians found they could receive data from various INPC member institutions but that the clinical content was difficult to interpret because each used a different code for the same test or observation so it was like receiving messages in French, Spanish and Italian when all they could understand was English.

LOINC was born from the desire to develop a lingua franca. From the beginning it has been a free and open system, encouraging additions, comments and feedback. Two new versions of LOINC are issued annually, with more than 2,000 new terms for tests or clinical observations per release. These new additions are based on requests from end users.

Tagged with: electronic medical records, medical informatics, Regenstrief



Media Contacts

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Sherold Barr HL7 503-595-5546 sherold@sheroldbarr.com

US Adoption

A few key highlights



US Federal Agencies

NLM

VA

DoD

IHS

NCI

CDC

EHR Incentive Program

a.k.a. "Meaningful Use"

Documents Using HL7 CCD Component HITSP/C32 (incorporated by reference

§ 170.207 Vocabulary standards for

time, patient identification, and user identification must be recorded when

(2) Stando Specification Record and

in § 170.299

(incorporate (b) Electro Standard, T Prescription Prescriber/P standard. In Version 8, R

in § 170.299 (2) Stando Standard, In Version 10.6 in § 170.299

October 200

(c) Electro results to pu Standard. H

reference in § 170.299). Implementation specifications. HL7 Version 2.5.1 Implementation Guide: Electronic Laboratory Reporting to Public Health, Release 1 (US Realm) (incorporated by reference in § 170.299).

representing electronic health information.

(2) Standard. The code set specified at 45 CFR 162.1002(a)(5).

(c) Laboratory test results. Standard. Logical Observation Identifiers Names and Codes (LOINC®) version 2.27, when such codes were received within an electronic transaction from a laboratory

(incorporated by reference in § 170.299).

45 CFK 162.1002(a)(5).

(c) Laboratory test results. Standard. Logical Observation Identifiers Names and Codes (LOINC®) version 2.27, when such codes were received within an electronic transaction from a laboratory (incorporated by reference in § 170.299). disclosures for treatment, payment, and health care operations, as these terms are defined at 45 CFR 164.501.

§ 170.299 Incorporation by reference.

(a) Certain material is incorporated by reference into this subpart with the

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TABLE 5–12. OBSERVATION/RESULT SEGMENT (OBX)										
Seq	Len	DT	Cardinality	Lab Result Sender Usage	ELR Receiver Usage	NHSN Receiver Usage	Lab to EHR Receiver Usage	Value Set	HL7 Element Name	Description/Comments
3		CWE	[11]	R	R	R	R	Laboratory Observation Identifier Value Set	Observation Identifier	Unique identifier for the type of observation. This field provides a code for the type of observation. OBX-3 in conjunction with OBX-4 Observation Sub-ID should uniquely identify this OBX from all other OBXs associated with this OBR.
										LOINC is used as the coding system for this field except where the test being reported has no equivalent LOINC code. In this case, use of local codes is allowed. This should only occur for new tests that have yet been coded by LOINC.
										When populating this field with values, this guide does not give preference to the triplet in which the standard (LOINC) code should appear.
										Lab to EHR - LOINC® is an HL7 approved code system and shall be used for the Observation Identifier as described in the appropriate HITSP Interoperability Specification. Use of LOINC codes for additional tests is strongly encouraged.
4	120=	ST	[01]	CE	CE	С	С		Observation Sub-ID	Harmonized condition predicate: Required if there is more than one OBX with the same OBX-3 Observation Identifier associated with the same OBR. Normally, this field is populated with a number, but text values may be used also.



September 9, 2011

Farzad Mostashari, MD, ScM National Coordinator for Health Information Technology Department of Health and Human Services 200 Independence Avenue, SW Washington, DC 20201

Dear Dr. Mostashari:

The HIT Standards Committee's (HITSC) Clinical Quality Measures Workgroup (CQMWG) and Vocabulary Task Force (VTF) jointly developed recommendations on the assignment of code sets to clinical concepts [data elements] for use in quality measures.

The CQMWG and VTF held a series of joint meetings to develop the set of recommendations. This letter transmits the recommendations to the Department of Health and Human Services (HHS) on the assignment of code sets to clinical concepts for use in quality measures. On August 17, 2011, the CQMWG and VTF reported on and discussed their findings with the HITSC, which were subsequently approved as outlined below.

LOINC Names

Introduction to LOINC Naming Conventions

Anatomy of a LOINC Term

5193-8:Hepatitis B virus surface Ab:ACnc:Pt:Ser:Qn:EIA

5193-8 LOINC Code

Hepatitis B virus surface Ab

ACnc

Pt

Ser

Qn

EIA

Component

Property Measured

Timing

System

Scale

Method

There are six major LOINC axes

NOT part of a LOINC Name

Reason for the test (disease it diagnoses)

Testing instrument

Specific details about the specimen

Priority (e.g. STAT)

Where testing was done

Who did the test

Test interpretation

Anything not part of naming the test

Stuff carried in other parts of HL7 message

Component

The substance or entity that is measured, evaluated, or observed

Sodium
Glucose
Brucella sp. <u>organism</u>
Influenza A Virus <u>antigen</u>
Cytomegalovirus <u>antibody</u>
Lipids.total



5193-8:Hepatitis B virus surface Ab:ACnc:Pt:Ser:Qn:EIA

Component Structure

Analyte Name^Challenge^Adjustments

Formal analyte name

Specify "subanalytes" May have subclasses Calcium Coronavirus Ag Calcium.ionized

Challenge

Two parts separated by "post"

1H post 100 g Glucose PO <time delay>post<challenge type>

Adjustments

Adjusted to pH 7.4

Property

The characteristic or attribute of the analyte that is measured, evaluated, or observed.

Major Categories:

mass substance catalytic activity arbitrary number



5193-8:Hepatitis B virus surface Ab: ACnc: Pt: Ser: Qn: EIA

Property

* the most difficult LOINC axis

Fully Named Properties

MCnc mass concentration

SCnc substance concentration

MCnt mass content

CCnc catalytic concentration

Prid presence or identity

Imp impression

Type "kind of"





Property is related to units of measure

MAPPING CAUTION



Common Property Issues

Fraction versus Ratio

Fraction = Part/Whole

NFr: % Eosinophils / leukocytes

SFr: % HGB which is A2

Ratio = multiple analytes from *same* system

MCrto: BUN/Creatinine in urine

Relative Ratio = measures from *different* system

RelRto: actual to normal control

Timing

The interval of time over which the observation or measurement was made

Pt point in time

12H *12 hour collection*

24H 24 hour collection





Non-Pt timings are often

5193-8:Hepatitis B virus surface Ab:ACnc:Pt:Ser:Qn:EIA

round with kate Property

System

The system (context) or specimen type upon which the observation was made.

Ser serum

Ser/Plas serum or plasma

Bld whole blood

Ur *urine*

Flu body fluid

Tiss tissue

XXX specified elsewhere



photo via AlishaV

System Structure

System[^]Super System

Super System

Patient is the default

Used to indicate:

blood product unit, bone marrow donor, fetus

818-5:A Ag:ACnc:Pt:RBC^BPU:Ord:

11670-7:Blood flow.mean:Vel:Pt:Aortic arch^fetus:Qn:US.doppler

Scale

Qn Quantitative

continuous numeric can have operators

Ord Ordinal

Ranked set (1+, 2+, 3+)

Nom Nominal

unranked collection

Taxonomy (e.g. bacteria)

Nar Narrative



photo via puuikibeach

5193-8:Hepatitis B virus surface Ab:ACnc:Pt:Ser:Qn:EIA

Method

Only needed if interpretation affected

Different normal ranges
Test sensitivity

Listed at the generic level

Agglutination

Enzyme Immunoassay

Probe with target amplification



5193-8: Hepatitis B virus surface Ab: ACnc: Pt: Ser: Qn: EIA



Standardized Assessments and Collections

Representing Patient Assessments in LOINC® Daniel J. Vreeman, PT, DPT, MSc^a, Clement J. McDonald, MD^b, Stanley M. Huff, MD^c aRegenstrief Institute, Inc and Indiana University School of Medicine, Indianapolis, IN; Lister Hill Center, National Library of Medicine, Washington DC; University of Utah and Intermountain Healthcare, Salt Lake City, UT ABSTRACT

Without being included in accepted vocabulary standards, the results of completed patient assessment instruments cannot be easily shared in health information exchanges. To address this important barrier, we have developed a robust model to represent assessments in LOINC through iterative refinement and collaborative development. To capture the essential aspects of the assessment, the LOINC model represents the hierarchical panel structure, global item attributes, panel-specific item attributes, and structured answer lists. All assessments are available in a uniform format within the freely available LOINC distribution. We have successfully added many assessments to LOINC in this model, including several federally required assessments that contain functioning and disability content. We continue adding to this "master question file" to further enable interoperable exchange, storage, and processing of assessment data.

INTRODUCTION

Despite progress on many fronts, interoperable health information exchange continues to be hampered by the plethora of idiosyncratic conventions for representing clinical concepts in different electronic systems. Many times, the lack of interoperable connections between systems means that valuable results are unavailable to clinicians when they need it.1 LOINC® (Logical Observation Identifiers Names and Codes) is a universal code system for identifying

and Codes) is a universal code system for identifying it.1 LOINC® (Logical Observation Identifiers Names results are unavailable to clinicians when they need connections between systems means that valuable

representation of assessments since its early development when it included codes for standardized scales such as the Glasgow Coma Score and the Apgar Score. Prior work^{5,6} has demonstrated the capability of LOINC's semantic model to represent many assessments with only modest extensions.

Over time, we have both significantly refined LOINC's model for patient assessments and added much new content. Here we present a summary of this progress. Specifically, the purpose of this paper is to describe LOINC's model for assessments, the methods and rationale by which this model was developed, the current assessment content, and some of the lessons learned in the process.

BACKGROUND

Fully specified LOINC names are constructed on six main axes (Component, Property, Timing, System, Scale, and Method) containing sufficient information to distinguish among similar observations. 2 Different LOINC codes are assigned to observations that measure the same attribute but have different clinical meanings. The LOINC codes, names, and other attributes are distributed in the main LOINC database made available at no cost in regular releases on the LOINC website (http://loinc.org). In addition to the LOINC database, Regenstrief develops and distributes at no cost a software program called RELMA that provides tools for searching the LOINC database, viewing detailed accessory content, and for

mapping local terminology to LOINC terms. mapping local terminology to LOINC terms. database, viewing detailed accessory content, and for RELMA that provides tools for searching the LOINC Int. J. Functional Informatics and Personalised Medicine, Vol. x, No. x, xxxx

LOINC®: a universal catalogue of individual clinical observations and uniform representation of enumerated collections

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Stanley M. Huff

University of Utah and Intermountain Healthcare, 4646 W. Lake Park Blvd. Salt Lake City, UT 84120,USA E-mail: Stan.Huff@imail.org

Abstract: In many areas of practice and research, clinical observations are recorded on data collection forms by asking and answering questions, yet without being represented in accepted terminology standards these results cannot be easily shared among clinical care and research systems. LOINC contains a well-developed model for representing variables, answer lists and the collections that contain them. We have successfully added many assessments and other collections of variables to LOINC in this model. By creating a uniform representation and distributing it worldwide at no cost, LOINC aims to lower the barriers to interoperability among systems and make this valuable data available across settings when and where it is needed.

Keywords: clinical observations; framework; health information technology; patient data; patient assessments; data sets; public health; research; standards;

patient data, patient assessments, data sets, pubble health, research, standards, Keywords: clinical observations, framework, health information technology,

Vreeman DJ, McDonald CJ, Huff SM. Representing patient assessments in LOINC®. AMIA Annu Symp Proc. 2010;832-836. PMID: 21347095.

Vreeman DJ, McDonald CJ, Huff SM. LOINC® - A Universal Catalog of Individual Clinical Observations and Uniform Representation of Enumerated Collections. Int J Funct Inform Personal Med. 2010;3(4):273-291.

Iteratively expanded the base lab panel model to accommodate more complex attributes



Hierarchy of a Panel in LOINC

57021-8 CBC W Auto Differential panel in Blood

PANEL HIERARCHY

0 1	LOINC#	LOINC Name
	57021-8	CBC W Auto Differential panel in Blood
	58410-2	Complete blood count (hemogram) panel in Blood by Automated count
	6690-2	Leukocytes [#/volume] in Blood by Automated count
	789-8	Erythrocytes [#/volume] in Blood by Automated count
	718-7	Hemoglobin [Mass/volume] in Blood
	4544-3	Hematocrit [Volume Fraction] of Blood by Automated count
	787-2	Erythrocyte mean corpuscular volume [Entitic volume] by Automated count
	785-6	Erythrocyte mean corpuscular hemoglobin [Entitic mass] by Automated count
	786-4	Erythrocyte mean corpuscular hemoglobin concentration [Mass/volume] by Automated count
	21000-5	Erythrocyte distribution width [Entitic volume] by Automated count
	788-0	Erythrocyte distribution width [Ratio] by Automated count
	777-3	Platelets [#/volume] in Blood by Automated count
	32207-3	Platelet distribution width [Entitic volume] in Blood by Automated count
	32623-1	Platelet mean volume [Entitic volume] in Blood by Automated count
	57023-4	Auto Differential panel in Blood
	770-8	Neutrophils/100 leukocytes in Blood by Automated count
_	25222 6	Neutraphile hand form/100 laukacutes in Blood by Autamated count



Purpose

Assessments are widely used...





. How do you rate	your confid	en ce that you	can getand ke	ep an erection								
	Very low Low Moderate High Very											
	1	2	8	4	6							
. With sexual stim or penetration (ent			ou relections be	en sufficient to	aflow							
fmnd stall sexually active	Amos rever iNever	(Much less than had the sime)	Occasionally (About helf the fine)	Most of the time (Much more than half the time)	Amos: dways /Aways							
0	1	2	3	4	- 6							
active	never (Never	(Much less than had the time)	(About half the sine)	(Much more than half the time)	alvays /Aways							
0	- 1	2	3	4	- 6							
During sexual intuition of it intil completion of it inaven't tried to have sexual intercourse.			oman	inta in you'r erec Signiy diffcut	Not officult							
0	1	2	3	4	- 6							
5. When you attem Imnoratal sexually active	Amost never /Never	Rarely	Occasionally	Most of the time	you?: Amos alvays /Away:							
		(Much less	(About helf the	fluch more than half the								
	1											

...and not unlike other clinical observations

LOINC could be a

master question file and uniform representation

Look back period for all items is 7 days unless another time frame is indicated.

E1100. Change in Behavior or Other Symptoms

Consider all of the symptoms assessed in items E0100 through E1000

Enter Code

How does resident's current behavior status, care rejection, or wandering compare to prior assessment (OBRA or PPS)?

- 0. Same
- 1. Improved
- 2. Worse
- 3. N/A because no prior MDS assessment

54696-0 Change in behavioral or other symptoms in last 7D: Find: Pt: ^Patient: Ord: MDSv3

NAME

Fully-Specified Name:	Component	Property	Time Aspect	System	Scale	Method
	Change in behavioral or other symptoms in last 7D	Find	Pt	^Patient	Ord	MDSv3

Long Common Name: Change in behavioral or other symptoms in last 7 days MDSv3

OBSERVATION ID IN FORM

E1100

FORM CODING INSTRUCTIONS

Consider all of the symptoms assessed in items E0100 through E1000.

NORMATIVE ANSWER LIST:

Source: Change in behavioral or other symptoms

Code System OID: 1.3.6.1.4.1.12009.10.1.34

SEQ#	Answer	Code	Answer ID
1	Same	0	LA11009-0
2	Improved	1	LA65-8
3	Worse	2	LA11011-6
4	N/A because no prior MDS assessment	3	LA11012-4

SURVEY QUESTION:

Text: Change in behavioral or other symptoms -

How does resident's current behavior status, care rejection, or wandering compare to prior assessment (OBRA or PPS)?

Source: MDSv3.E1100



MM MM

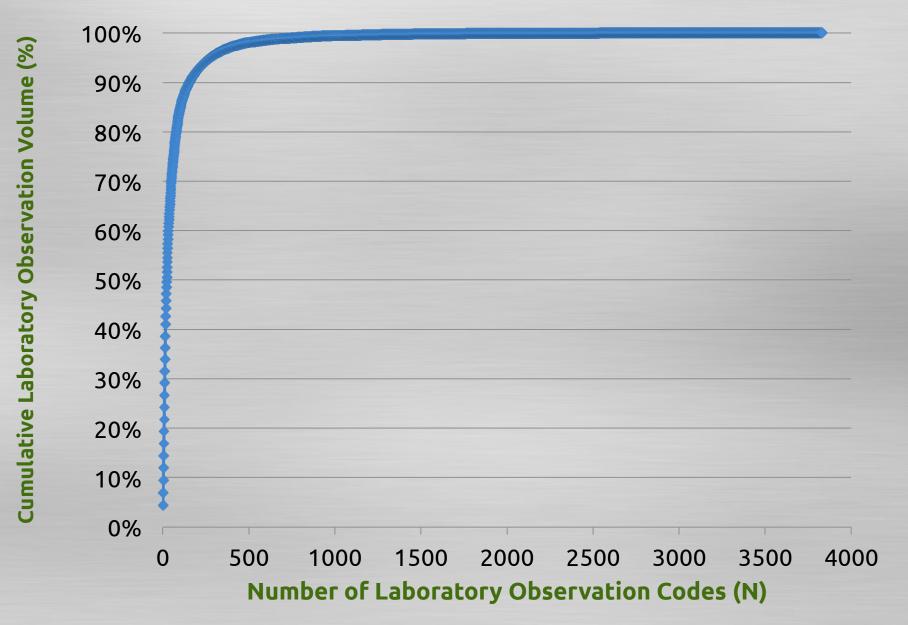
FORMS LOINCS ANSWERS +

Panels/Forms Available as Separate Download

+	@http://lo	inc.org/downlo	oads											¢
						Sheets	Charts	Smar	tArt Graph	ics	WordArt			
0	A B	C	D	EFF	G H	-	K		M	N	0	P	Q	R
1	PARENT_ID PARENT_L	C PARENT_NAME I	D 5	EQUENCE LOINC_NU	I LOINC_NAM DISPLAY_	N. OBSERVATI SUBM	NITTEF SKIP_LOGI	CSKIP_LOG	IC SKIP_LOGI	CANSWER	R MAXIMUN	DEFAULT_V	TYPE_OF_E	DATA_TYPE DA
982	28138 54580-6	Minimum Data Set 3.0:	28138	589 54580-6	Minimum Data Set 3.	0:-:Pt:^Patient:-::								
983	28138 54580-6	Minimum Data Set 3.0:	28139	1 54501-2	Identification informa	ation:-:Pt:^Patient:-:	:							
984	28138 54580-6	Minimum Data Set 3.0:	28187	2 54508-7	Hearing, speech, and	vision:-:Pt:^Patient	1421		If No, Cont	inue to BO	200, Hearing.	If Yes, Skip	to G0100, Ac	tivities of Daily
985	28138 54580-6	Minimum Data Set 3.0:	28198	3 54509-5	Cognitive patterns:-:P	t:^Patient:-::						200		
986	28138 54580-6	Minimum Data Set 3.0:	28231	4 54633-3	Mood:-:Pt:^Patient:-:	MDSv3:								
987	28138 54580-6	Minimum Data Set 3.0:	28282	5 54511-1	Behavior:-:Pt:^Patien	to::								
988	28138 54580-6	Minimum Data Set 3.0:	28309	6 54518-6	Preferences for custo	mary routine and a	tivities:-:Pt:^Pat	ent:-::		11-				
989	28138 54580-6	Minimum Data Set 3.0:	28354	7 54522-8	Functional status:-:Pt	:^Patient:-::								
990	28138 54580-6	Minimum Data Set 3.0:	28387	8 54528-5	Bladder and bowel:-:	Pt:^Patient:-::				-				
991	28138 54580-6	Minimum Data Set 3.0:	28404	9 54531-9	Active disease diagno	sis:-:Pt:^Patient:-::								
992	28138 54580-6	Minimum Data Set 3.0:	28483	10 54556-6	Health conditions:-:P	t:^Patient:-::								
993	28138 54580-6	Minimum Data Set 3.0:	28530	11 54565-7	Swallowing Swallowin	ng/Nutritional Statu	s							
994	28138 54580-6	Minimum Data Set 3.0:	28551	12 54570-7	Oral ∨ de Oral/Deni									
995	28138 54580-6	Minimum Data Set 3.0:	28561	13 54572-3	Skin conditions:-:Pt:^	Patient:-::								
996	28138 54580-6	Minimum Data Set 3.0:	28637	14 55094-7	Medications:-:Pt:^Pat	tient:-:MDSv3:							- T-1	
997	28138 54580-6	Minimum Data Set 3.0:	28647	15 54990-7	Special treatments an	nd procedures:-:Pt:^	Patient:-::							
998	28138 54580-6	Minimum Data Set 3.0:	28733	16 55042-6	Restraints:-:Pt:^Patier	11 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1								NE II
999	28138 54580-6	Minimum Data Set 3.0:	28751	17 55059-0	Therapy supplement	for Medicare PPS:-:	et:^Patient:-::						-	
000	28138 54580-6	Minimum Data Set 3.0:	28757	18 55063-2	Assessment administ									
001	28139 54501-2	Identification informati	28140	1 54581-4	Facility provider num	bers:-:Pt:Facility:-:N	IDSv3:							
002	28139 54501-2	Identification informati	28143	2 54582-2	Provider ty Type of P									
003	28139 54501-2	Identification informati	28144	3 54502-0	Assessmen Type of A									
004	28139 54501-2	Identification informati	28150	4 54896-6	Submission requirem	ent:Find:Pt:^Patient	:Ord:MDSv3:							
005	28139 54501-2	Identification informati	28151	5 54503-8	Legal name of resider									
006	28139 54501-2	Identification informati	28157	6 45966-9	Social security & med		^Patient:Set::						1 - 1	
007	28139 54501-2	Identification informati	28161	7 54505-3	Language: -: Pt: ^Patier									
800	28139 54501-2	Identification informati	28165	8 54506-1	Optional resident iter									
009	28139 54501-2	Identification informati	28169	9 54589-7	Preadmission Screeni		view (PASRR):Fin	d:Pt:^Patier	nt:Nom:MDS	v3:				
010	28139 54501-2	Identification informati	28170	10 45973-5	Conditions Condition									
011	28139 54501-2	Identification informati	28178	11 54590-5	Entry type: Type of E	The second second second								
012	28139 54501-2	Identification informati	28179	12 54591-3	Admitted fr Entered fr									
013	28139 54501-2	Identification informati	28180	14 54592-1	Previous assessment		ignificant correct	ion:TmStp:	Pt:^Patient:C	n:MDSv3:				
014	28139 54501-2	Identification informati	28181	15 54593-9	Assessment reference					1				
015	28139 54501-2	Identification informati	28182	16 54507-9	Medicare stay:-:Pt:^P			-		to 80100.	Comatose. If	Yes, Continu	ue to A24008	3, Start date of
016	28139 54501-2	Identification informati	28859	13 55128-3	Discharge status:Find		IDSv3:			1				
017	28144 54502-0	Assessment ∨ tracking	28145	1 54583-0	Federal OB Federal O			g						

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Current Versions

LOINC 2.38

Released: 2011-12-30

RELMA 5.5

Released: 2011-12-30





Common LOINC Laboratory Observation Codes

We created an empirically-based list of the most common LOINC result codes for laboratories, practices, researchers, and others who wish to map their int for much of the result volume, we think that Top 2000+ list will be an

"Top 2000 Results"

represent about 98% of the test volume carried by three large

ab Tests with a wealth of advice and guidance about which codes to

LOINC Top 2000+ Lab Observations and Mapper's Guide

- LOINC Mapper's Guide to the Top 2000+ Lab Observations Introduction to Version 1.0 (pdf)
 - File Version: Top 2000+ Lab Observations Introduction 1.0 | File type: application/pdf | Release Date: 2011-05-31

File Description: Background document describing the origins, development, and organization of the LONIC Mapper's Guide to the Top 2000+ Lab Observations.

- LOINC Mapper's Guide to the Top 2000+ Lab Observations Version 1.0a (pdf)
 - File Version: LOINC Mapper's Guide to the Top 2000+ Lab Observations 1.0 | File type: application/pdf | Release Date: 2011-06-01

File Description: Document containing mapping guidance for the LOINC Top 2000+ Lab Observations.

OID: 1.3.6.1.4.1.12009.10.2.3

- LOINC Mapper's Guide to the Top 2000+ Lab Observations Version 1.0a (xlsx)
 - File Version: LOINC Mapper's Guide to the Top 2000+ Lab Observations 1.0 | File type: application/vnd.openxmlformats-officedocument.spreadsheetml.sheet | Release Date: 2010-06-01 File Description: Document containing mapping guidance for the LOINC Top 2000+ Lab Observations. Requires a loinc.org account.

OID: 1.3.6.1.4.1.12009.10.2.3

LOINC Top 2000+ Lab Observations - Version 1.0 (csv)

File Version: Top 2000+ Lab Observations 1.0 | File type: text/comma-separated-values | Release Date: 2010-05-31

File Description: The list of LOINC Top 2000+ Lab Observations in a simple CSV file. Does NOT contain mapping guidance.

OID: 1.3.6.1.4.1.12009.10.2.3

Notes about Version 1.0a: This version contains a couple of corrections for UCUM units. The rest of the content is unchanged from version 1.0 (2010-05-31)

Recent Forum Posts

Microbiology | Re: Microbiology Text results and LOINC

Microbiology | Re: Organism Names and LOINC?

More posts.



LOINC New paper published in JBI about helping map to #LOINC with RELMA by augmenting local test names loinc.org/articles /Kim20... about 1 hour ago - reply - retweet -

Join the conversation

Common Result Codes in LOINC and RELMA

The Top 2000+ LOINC terms have their relative rank stored in the COMMON_TEST_RANK field of the main LOINC table. Starting with RELMA version 5.3, the program's "common lab tests only" filter will limits searches to only return codes in this common test list.

Through LOINC version 2.34, the INPC_PERCENTAGE field of the main LOINC table contained the relative percent volume by test result based on an analysis of 3 years (2006–2008) of laboratory result data from the Indiana Network for Patient Care (INPC), one of the oldest and largest health information exchanges in the US. Within the RELMA mapping program, the "common lab tests only" filter limits searches to only return codes in this common test list.

Background Reading

■ Vreeman DJ, Finnell JT, Overhage JM. A Rationale for Parsimonious Laboratory Term Mapping by Frequency. Proc AMIA Symp. 2007;771-775.



Value set of universal laboratory order codes from LOINC

This value set defines a collection of universal laboratory order codes for use by developers of order entry systems that deliver them in HL7 messages to laboratories, where they could be understood and fulfilled. This value set is designed to cover greater than 95% of the test ordering volume in the US, and was by the HITSP C80 Clinical Document and Messaging Terminology

sidered a minimum 'starter' set" and "does not attempt to include all

a.k.a. "Top 300 Orders"

Value Set

- Common Lab Orders Value Set Preface to Version 1.2 (pdf)
 File Version: Common Lab Orders Preface 1.2 | File type: application/pdf | Release Date: 2011-06-03
- Common Lab Orders Value Set Version 1.2 (pdf)
 File Version: Common Lab Orders 1.2 with Preface | File type: application/pdf | Release Date: 2011-06-03
 O(D: 1.3.6.1.4.1.12009.10.2.2
- Common Lab Orders Value Set Version 1.2 (xlsx) or if you prefer (xlsx) (xlsx)

 File Version: Common Lab Orders 1.2 | File type:application/vnd.openxmlformats-officedocument.spreadsheetml.sheet | Release Date: 2011-06-03

 OID: 1.3.6.1.4.1.12009.10.2.2

HITSP C80

HITSP C80 - Clinical Document and Message Terminology Component
 Table 2-96 "Laboratory Order Value Set" reference to the LOINC Common Lab Orders value set



Example UCUM codes for units of measure in electronic messages

This is a draft enumeration of The Unified Code for Units of Measure (UCUM), designed to make it clear what the UCUM syntax would produce for specific unit patterns in electronic communication. This early version, composed in a relatively short time frame, is based on content provided by Intermountain Healthcare, from a National Library of Medicine and Regenstrief Institute project that is analyzing raw units from more than 23 laboratory sources and their translation to UCUM and from the HL7 table of units. In this version we have not included all of the content from all of these sources. Specifically for this version, we excluded units for which we could not quickly find definitions or clear patterns of usage, units of measure that we believed would only be used in pharmacy dispensing, and units used for purely clinical reporting (e.g. cigarette pack-years). We have included most of the pure metric units that were in the source table whether they apply



Example UCUM codes for units of measure in electronic messages

This is a draft enumeration of The Unified Code for Units of Measure (UCUM), designed to make it clear what the UCUM syntax would produce for specific unit patterns in electronic communication. This early version, composed in a relatively short time frame, is based on content provided by Intermountain Healthcare, from a National Library of Medicine and Regenstrief Institute project that is analyzing raw units from more than 23 laboratory sources and their translation to

a.k.a. "Common Units of Measure"

. Specifically for this version, we excluded d only be used in pharmacy dispensing, and were in the source table whether they apply ould be excluded.

Table of Example UCUM Codes for Electronic Messaging

- X Table of Example UCUM Codes for Electronic Messaging Preface to Version 1.1 (pdf)

 File Version: Table of Example UCUM Codes for Electronic Messaging Preface 1.1 | File type: application/pdf | Release Date: 2011–10-04
- Xable of Example UCUM Codes for Electronic Messaging Version 1.1 (pdf)

 File Version: Table of Example UCUM Codes for Electronic Messaging 1.1 | File type: application/pdf | Release Date: 2011–10-04
- Table of Example UCUM Codes for Electronic Messaging Version 1.1 (xlsx)

 File Version: Table of Example UCUM Codes for Electronic Messaging 1.1 | File type:application/vnd.openxmiformats-officedocument.spreadsheetml.sheet | Release Date: 2011-10-04

by Daniel Vreeman - last modified 2011-10-12 03:00



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Content

LOINC Usage Value Sets Newborn Screening Document Ontology Documentation

LOINC Users' Guide Recommended Readings Presentations/Tutorials

LOINC MAPPER'S GUIDE TO TOP 2000+ LAB TESTS v1.0a

SORT BY: Class Override >System Adjusted > Long Common Name

В	С	E	F	G	Н		P
LOINC # Long Common	Name	Class Override	Rank	Example UCUM	UCUM Display	Comment	System Adjusted

126 Antibacterial susceptibility

The statistics for antibiotic susceptibility tests in the Top 2000 List are not as broadly based as most of the other test categories, because antibiotic susceptibilities were available from only one of our 3 sources.

LOINC provides codes for antibiotic suceptibility testing based on method used. The four major categories are as follows:

- 1) A general flavor that does not specify the method of testing used
- 2) Minimum Inhibitory Concentrations (MIC)
- 3) Kirby Bauer disc testing (KB) and
- 4) Gradient strip (E-test)

The general flavor can be used to report results for any of the three more specific approaches (Kirby Bauer, MIC or E-test susceptibilities) assuming that the details regarding the method of testing is provided elsewhere in the messages or in other OBX segments.

The majority of the antibiotic susceptibility tests that made it into the top 2000 list are of this general flavor type, but a few MIC tests and gradient strip LOINC codes also appear. In case your laboratory prefers the more specific codes for the antibiotics listed here, you can find them under the Antibiotic susceptibility class in the full LOINC database.

Some of the antibiotics used to treat tuberculosis are also used to treat more common bacterial infections. LOINC provides specific codes for reporting antibiotic susceptibilities to slow growing Mycobacteria – such as M.tuberculosis, M.avium and M.intracellular, and these codes should be used for reporting antibiotic susceptibilities for such bacteria. These codes can be identified by the phrase "slow growing mycobacteria" in the method part of the LOINC name. Antibiotic susceptibilities to a fast growing mycobacteria can be reported under the same codes as any other bacteria.

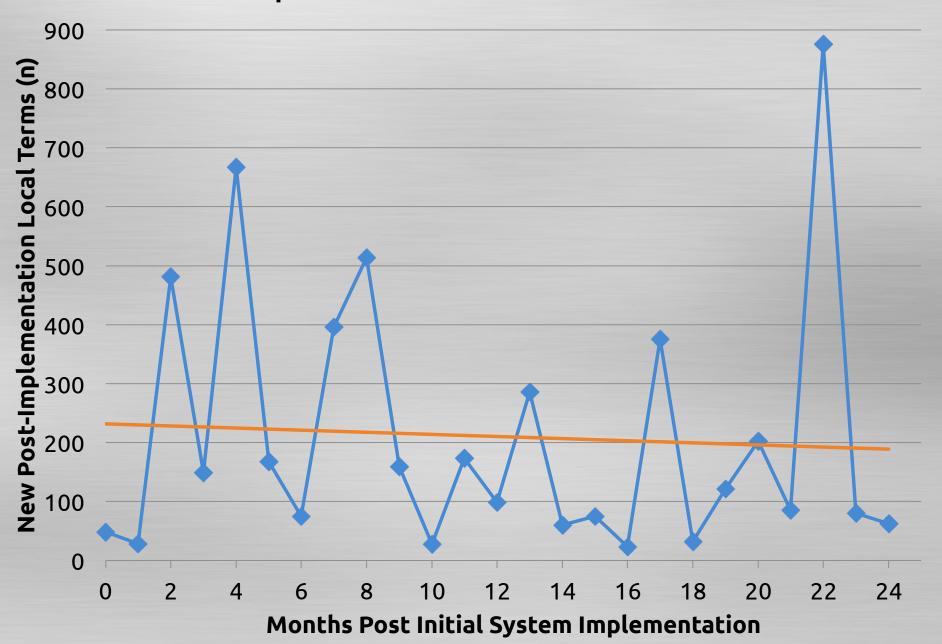
4	-	-
п	,	_

128	17-3 Methicillin resistant Staphylococcus aureus [Presence] in Unspecified specimen by Organism specific culture	Antibacterial susceptibility	146	Methicillin Resistant Staphlocuss via culture	Any
129	50-7 Amikacin [Susceptibility]	Antibacterial susceptibility	414		Isolate
130	62-3 Amoxicillin+Clavulanate [Susceptibility]	Antibacterial susceptibility	549		Isolate
131	54-9 Ampicillin [Susceptibility]	Antibacterial susceptibility	331		Isolate
132	55-6 Ampicillin+Sulbactam [Susceptibility]	Antibacterial susceptibility	330		Isolate
188	58-0 Aztreonam [Susceptibility]	Antibacterial	454		Isolate





Post-Implementation Terms Since Go-live



Add half as many terms after go-live.

Most new local terms could be mapped to existing LOINCs.

Jump In!





ivdconnectivity.org







Next Generation IVD Connectivity

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CREATING SOLUTIONS

GENERATING MOMENTUM Countdown to Connectathon 2012



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"Today's clinical labs and their vendors—the IVD industry—spend a great deal of time and money connecting analyzers and IT systems to one another. IICC unites the stakeholders around a common goal-to solve our shared problem through modern connectivity technology."

Eric Olson, President, IICC, and Vice President, Automation &IT, Siemens Healthcare Diagnostics MORE>

"The current explosion of healthcare information technology (HIT) will continue to heavily impact healthcare in the foreseeable future. Modernizing instrument interface and interoperability standards will integrate the clinical lab into the mainstream of this health information flow."

Jay B. Jones, PhD, DABCC, Board Member and Chair of the Provider Review Committee, IICC, and Director, Chemistry & Regional Labs, Geisinger Health System MORE >

Founding Members













Ortho Clinical Diagnostics



SIEMENS



The Race is On!



Happy LOINCing!

photo via ryarwood

Acknowledgements

LOINC Development Team

Clem McDonald, Kathy Mercer, Jaci Phillips, Jami Deckard, David Baorto, Kelly Malott

RELMA Development Team

John Hook, Mark Fisher, Karen Ahmed, Anandhi Sowmyan, James Dennis

LOINC Committee

Supporters: NLM, Regenstrief Institute, Regenstrief Foundation

