

A Clinically Derived Terminology: Qualification to Reduction

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Mayo Foundation is developing synonym rich entry points for the recording of patient problems by clinicians, which will map to the KP-Mayo Convergent Medical Terminology. We describe the empirical sources for these terminology components, and how the number and complexity of the terms could be substantially reduced by the introduction of a Qualifier axis. The expressive power of these entry points is dramatically enhanced by this axis. This work is being integrated into terminology navigation modules being jointly developed with Lexical Technology, which leverages UMLS content. It will form the basis for structured problem entry into Mayo's Computer-based Electronic Record.

INTRODUCTION

Computer-based Patient Records are penetrating the American healthcare environment with an accelerating rapidity. Less rapid in appearance are standard terminologies that can capture the clinical detail of patient problems, diagnoses, events, or interventions with sufficient detail to support evidence based medicine. While some terminologies approach this functional behavior, most do not¹ and arguably none presently seem natural to clinicians using them for the routine documentation of healthcare.

The Mayo Foundation has indexed patient problems, diagnoses, and procedures for a century. These indices form the basis of outcomes research, disease natural history studies, treatment response evaluations, continuous quality improvement efforts, and clinical epidemiology. During more recent decades, reimbursement and capitation have been computed from their derivative mappings against statistical classifications such as the ICD-9-CM. The process of indexing has become increasingly computerized. It is approaching the logical final step of being integrated with the Computer-based Patient Record as a structured entry using controlled medical terminology.

This report characterizes some preliminary work at Mayo by our Clinical Terminology Committee

to make the presentation of such a controlled terminology familiar and relevant to Mayo practitioners. This terminology development is a module within several larger projects, including Mayo's Electronic Medical Record initiative, the Kaiser-Permanente and Mayo Foundation partnership to evolve toward a Convergent Medical Terminology², and the Lexical Terminology Inc. initiatives to produce a multi-functional terminology navigation and browsing resource.

METHODS

Clinical Sources

Two clinical applications served as the empirical source of problem statements entered by clinicians, the Mayo Master Sheet and Assessment statements within electronic Clinical Notes.

The Mayo Master Sheet has captured summary diagnoses, problems, and dismissal impressions since its introduction at the Clinic in 1907. The full text of these entries has been electronically available since 1993. This evaluation sorted the 2,548,119 strings captured over a 15 month period into frequency order, yielding 811,060 unique strings. The most frequent string appeared 30,092 times, the 5000th rank ordered string was present 37 times. From these, the 5000 most frequent strings formed the basis of the Master Sheet input.

The Clinical Notes application was introduced at Mayo in 1994, and is being used by an increasing number of Mayo's 1,200 staff clinicians in the outpatient setting. The 1,389,781 strings which appeared in the Assessment or Impression part of this repository during 1995 were sorted into 547,979 unique terms. The most frequent string was recorded 38,550 times, and the 5000th rank ordered string appeared 18 times. These strings were then merged with the Master Sheet input.

Clinical Review

The unique reduction of the merged Master Sheet and Clinical Notes terms were reviewed subjectively. Two conclusions were immediately evident:

1. Logistic and operational terms were embedded in these lists, which have little clinical relevance.
2. Many terms were combinations of diagnoses and operational qualifiers.

We carefully pruned the list of logistical elements, and recorded the reason for reductions. In the course of this process, a preliminary pattern of embedded qualifiers was created based on their appearance in the clinical sources.

At this point, we established the utility and feasibility of distinguishing operational qualifiers from clinical modifiers. Qualifiers we defined as words or phrases that operationally or administratively qualify the meaning of a diagnosis or problem, e.g. *history of* a condition, *status post* a procedure, or *rule out* a condition (Mnemonic: Q for Quality Assurance Studies). These are opposed to a different but equally important group of variations which we class as Modifiers. Modifiers are words or terms which modify the severity, location, acuity, or other intrinsic clinical detail of a diagnosis or condition, e.g. *Stage I*, *acute*, *antero-lateral*, etc. (Mnemonic: M for Medical).

These distinctions are practical in part because we deemed Qualifier enumeration and recognition to be straightforward. Modifiers are

a vastly more complex problem³, which invoke semantic and linguistic issues we chose to defer.

Qualifiers

Having distinguished the role of Qualifiers, and establishing an empirical starting point for a Qualifier ontology, we completed a preliminary thesaurus and ordering for Qualifiers. These concepts and their synonyms could then be identified within our merged clinical terms corpus, and removed. This process was facilitated by invoking the lexical normalization tools⁴ developed by Alexa McCray at the NLM and distributed with the UMLS.

The underlying Mayo Terminology was reinforced by the re-addition of terms that had been present only in combination with a Qualifier. Similarly, some phrases had Qualifiers intrinsically embedded within a composite term, where the meaning would be distorted by the excision of the Qualifier; these were retained in the terminology.

RESULTS

Terminology edits which resulted from combination, review and reduction of the Master Sheet and Clinical Notes sources appear in the table. Overlap between these sources approximated only 20%. An additional 20% were deleted due to superfluous combinations of terms already present with our Qualifier lexicon.

Process or Code	Process Step or Code Meaning	Count of Terms for Each Step		
		Added	Deleted	Remaining
Master Sheet	Most Frequent MS Terms over 3 years	5,000		5,000
Clin. Notes	Most Frequent from Clinical Notes	5,000		10,000
Unique Merge	Drop Terms Redundant Across Sources		2,047	7,953
D	Qualifier marked for Deletion		2,032	5,921
E	Type of Examination		691	5,230
A	Abbreviation		461	4,769
S	Site of Diagnoses or Examination Including anatomy.		4	4,765
L	Location of Visit		105	4,660
Pr	Prescription drugs or therapy		12	4,648
Q	A new term added for a Qualifier	129		4,777
T	A new Term added for an Abbreviation	387		5,164

Table: "Balance register" of term volume, by review "transaction."

Measurable deletions took place for administrative or provincial reasons, accumulating another 10% reduction in aggregate. Thus, from our original source of 10,000 terms, just 5,164 unique terms remained after review and reduction.

The Appendix enumerates the 269 strings that form our preliminary ontology of Qualifiers. These include 104 Preferred Terms, or unique concepts; the remainder are variants, synonyms, or abbreviations. The two character prefix to the term identifiers, borrowed from the UMLS, indicates:

PT	Preferred Term
SY	Synonym
LV	Lexical Variant
AB	Abbreviation

The three digit term code conveys a superficial hierarchy and notation of synonymy. Terms with identical numbers are functionally synonymous (even if not exactly synonymous – they are so interpreted for purposes of Problem List entry).

DISCUSSION

This report begins a suite of presentations that address multiple steps in the empirical creation of a clinical terminology. While the desirable attributes of clinical vocabularies have become relatively well understood^{5,6}, their actualization into widely used or useful terminologies for computer-based patient records has lagged.

The practical consequences of introducing a Qualifier axis are almost self-evident. Whereas our raw lexicon had an ability to recognize about 8,000 terms as potential problem statements, the simple combinatorial of 270 qualifiers across some 5,000 disease specific terms parleys this by three orders of magnitude – a considerable advantage. Invoking the lexical mapping utilities associated with the Specialist Lexicon⁴ would further multiply the number of “strings” which could match our preliminary terminology.

The recognition that an independent axis can geometrically expand the expressive power of a terminology is hardly novel. Multi-axial coding dates at least to the development of the SND⁷ by the New York Academy of Medicine in 1928; this of course is the ancient forerunner to

SNOMED⁸. Nevertheless, the development of a well-structured “General” axis, capturing Qualifiers and Modifiers lags behind in the evolution of most terminologies. This has practical manifestations, since reimbursement codes presently cannot discriminate *Rule-Out* conditions from final diagnoses, nor have they any reasonable mechanism to recognize disease severity.

Our empirical modality has prompted a series of Usability Laboratory evaluations⁹ which focus on the practical ability of clinicians to navigate or use our terminology in prototypes of computer-based patient records. The ability to qualify existing terms to fit the clinical scenario is warmly received in these subjective studies. Indeed, a moderate clamor to implement parallel functionality for clinical modifiers affirms a known limitation.

The recognized limitations of this present work are well understood. Future research must address an ontology of Modifiers, analogous to that created for Qualifiers. Whether these should be inheritable within classes of diagnoses to guard against non-sense term coordination and to inform term composition, bears scientific cost-benefit consideration. This work also does not provide an ontology for the residual 5,164 Mayo Clinical Terms, although we are approximating that in our pre-alpha evaluations by mapping to UMLS hierarchies. Finally, while an empirically derived starting point affords familiarity and relevance, it suffers incompleteness and non-comprehensiveness. This is being addressed by Mayo specialist reviews of our terminology content for completeness.

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No	Qualifier	Family History	AB542	NEED	PT804	Preliminary	AB874	by path
PT100	Indeterminate	Family Hx	PT542	Ruled out	AB804	Prelim	AB874	by path
SY100	Unknown	FH	PT543	Rd/O	PT810	RULE Out	PT875	Cytologic
PT101	Indeterminate etiology	Family history of	AB543	Rd/O	AB810	RO	SY875	Cytology
LV101	of indeterminate etiology	Family Hx of	AB544	without	AB810	RO	AB875	by cytology
SY101	of undetermined etiology	Positive family history	PT544	W/O	PT820	Differential	PT890	Positive
SY101	Unknown cause	Positive family Hx	AB544	w/o	AB820	Diff	AB890	Pos
SY101	Unknown etiology	Positive family history of	SY544	Free from	PT821	Working	AB890	+
LV101	of unknown etiology	Positive family Hx of	LV544	Free of	PT830	Include	LV890	Positive for
PT110	Uncertain	Strong family history	SY544	stn	SY830	Includes	PT891	Present
PT111	Uncertain etiology	Strong family history of	AB544	s/	PT832	Cannot exclude	PT892	Known
LV111	of uncertain etiology	Strong family Hx	AB544	s	AB832	Cannot rule out	PT893	Specific
PT120	Unspecified	First degree relative with	PT550	Resolved	AB832	Cannot RO	PT893	Specific for
SY120	Not Otherwise Specified	No family history	AB550	Resolv	AB832	Cannot R/O	LV893	Specific for
AB120	NOS	No family Hx	PT560	Remission	PT840	High Risk	PT894	Identified
SY120	Non-specific	No family history of	AB560	Remiss	PT841	Likely	SY894	Identifiable
AB120	NS	No family Hx of	LV560	in remission	PT850	Probable	PT895	With
LV120	Non-specific	Negative family history	SY560	Remitted	SY850	Most likely	AB895	W/
PT121	Unidentified	Negative family Hx	PT561	In continued remission	PT851	Suspicious for	AB895	W
SY121	Not identified	Negative family history of	PT562	No evidence of recurrent disease	AB851	Susp	AB895	w/
PT200	History	Negative family Hx of	PT570	No Additional Disease	PT852	Evidence of	AB895	w
AB200	Hx of	Follow Up	AB570	NAD	LV852	Evidence for	SY895	cum
AB200	HO	FU	PT600	Treated	SY852	Evidence	AB895	c
AB200	H/O	F/U	AB600	Fixed	AB852	Evid	AB895	c/
LV200	History of	Follow Up Care	PT610	on therapy	PT853	Compatible	PT896	Absolute
AB200	Hx	F/U Care	LV610	under therapy	LV853	Compatible with	PT899	Certain
PT210	by history	FU Care	AB610	on treatment	SY853	Consistent	PT900	Principal
AB210	by Hx	Check	SY610	on Rx	LV853	Consistent with	PT901	Primary Diagnosis
PT220	Previous	Recheck	LV610	under treatment	AB853	CW	PT910	Secondary Diagnosis
PT230	Status Post	ReCK	PT611	on medical therapy	AB853	CW	PT911	Incidental
AB230	SP	Normal	PT612	on replacement therapy	AB853	cW	PT912	Innocuous
AB230	SP	NI	PT620	on replacement therapy	PT860	Unconfirmed	PT920	Administrative
SY231	Prior history of	Within normal limits	SY620	Controlled	PT861	Unconfirmed	PT921	Reimbursement
PT231	Prior Hx of	WNL	SY620	under control	SY861	Seen	PT921	Billing
AB231	Prior Hx of	Negative examination	SY620	under adequate control	SY861	Witnessed	PT923	Management
PT240	Past	Neq exam	PT621	under good control	SY861	Watched	PT924	Accounting
SY240	Past history	Neq examination	SY621	well controlled	PT862	Reported	PT925	Accreditation
LV240	Past history of	Normal examination	PT630	Controlled on therapy	PT870	Clinical	PT926	Review
PT241	Remote	Normal exam	SY630	Controlled on treatment	PT870	Clinical	PT930	Research
SY241	Remote History of	No Dx	PT631	Controlled on medication	PT871	Surgical	PT931	Protocol
LV241	Remote History of	Satisfactory	PT640	Stable	SY871	Surgically	PT932	Study
AB241	Remote Hx of	Satis	PT690	Uncontrolled	SY871	by surgery	PT933	Statistical
PT250	Recent	Negative	SY690	Not controlled	PT872	Radiologic	PT940	Reportable
PT260	Recurent	Neq	SY690	Out of control	AB872	Radiol	PT942	Mandatory Public Health
SY261	Relapsing	-	PT691	Unstable	SY872	X-Ray	PT950	Screening
SY261	Re-occurring	Negative for	SY691	Not stable	SY872	Radiologically		
PT290	No history	Neq for	PT692	Labile	SY872	Radiological		
AB290	No Hx	No	PT800	Possible	PT873	Laboratory		
SY290	Negative history	Not	PT801	Suspected	AB873	Lab		
AB290	Neq history	No evidence	SY801	Suspect	SY873	by laboratory		
AB290	Negative Hx	No evidence of	AB801	Susp	AB873	by lab		
SY290	Neq Hx	No evidence for	PT803	Questionable	PT874	Pathologic		
AB290	No past history	Not seen	SY803	Question	AB874	Path		
AB290	No past Hx	Not observed	AB803	Quest	SY874	Pathologically		
PT291	Never	No evidence of disease	LV803	Question of	SY874	by pathology		

Appendix: Tabular list of Qualifiers derived from clinical sources and used in reduction.