# AN ONTOLOGY FOR SUPPORTING CLINICAL RESEARCH ON CERVICAL CANCER

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Abstract: This work presents an ontology for cervical cancer that is positioned in the center of a research system for conducting association studies. The ontology aims at providing a unified "language" for various heterogeneous medical repositories. To this end, it contains both generic patient-management and domain-specific concepts, as well as proper unification rules. The inference scheme adopted is coupled with a procedural programming layer in order to comply with the design requirements.

## **1 INTRODUCTION**

Medical Knowledge, being the compilation of many years of research, has grown vast both in volume and in complexity. Recently, the need for employing semantically-aware models of medical knowledge has become evident. Since then, ontologies have been successfully used in various medical domains, disciplines and even aspects of medical practice and research. Examples of this include the Gene Ontology (Ashburner et al., 2000), the ReMINE ontology for adverse events (http://www.reminethe Ontology of Clinical Reproject.eu/), (http://rctbank.ucsf.edu/home/ocre.html), search the Ontology for Clinical Investigators (http://www.bioontology.org) etc.

This paper presents an ontology designed to facilitate research in the domain of cervical cancer (CxCa). This is, to the best of our knowledge, the first ontology to deal with concepts of the CxCa domain. It's main targets are (i) to provide a means for unification of various result coding formats and (ii) to extract implicit knowledge in order to produce potential query terms as inferred types of individuals. Through this process, medical researchers are given the ability to form patient groups, large enough to provide statistically significant results in association studies.

The paper is structured as follows: In section 2 we describe the problem, while in section 3 we try

to provide some intuition about the CxCa domain. Section 4 is devoted to the description of the basic structure of the proposed ontology, section 5 outlines the medical unification rules and section 6 the inferencing procedure followed. Section 7 contains the technologies utilized for implementation and some ontology statistics and section 8 concludes the paper.

## 2 PROBLEM STATEMENT

In an effort to gain a more comprehensive and holistic insight on the origin of complex diseases, genetic association studies (Hirschhorn and Daly, 2005) constitute a significant approach for clinical researchers. In order to perform statistically meaningful association studies, large amounts of clinical data are required - especially when performing studies among many study factors. However, the current clinical practice of constructing disposable and isolated clinical research repositories hinders the construction of collections big enough to facilitate the execution of complex association studies. To tackle this problem, comes the unification of existing medical repositories that contain heterogeneous cervical cancer related information.

The need to resolve this heterogeneity, makes the ontological representation of the CxCa knowledge the perfect candidate solution to this type of problems.

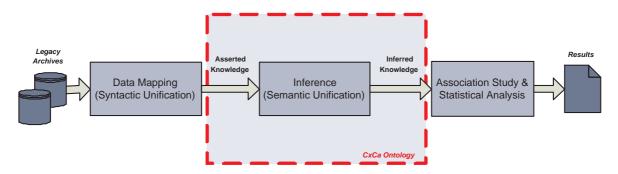


Figure 1: The Context of use of CxCa Ontology.

Ontologies have been widely used as unifying models to deal with heterogeneity issues of legacy archives. Furthermore, reasoning on top of ontologies is a very active field of ongoing research that has already produced fruitful and concrete results (Baader et al., 2006), (Tsarkov and Horrocks, 2006), (Hustadt et al., 2004), (Dyckhoff, 2000), (Sirin et al., 2007). Finally, there are plenty of semantic querying languages suitable for querying on ontologies (Jeen et al., 2004).

The ontology presented here has been developed within the context of ASSIST (Mitkas et al., 2008), a research project funded by the European Commission. The main objective of this project is the semantic unification of physically isolated and heterogeneous medical repositories that include cervical cancer related data into one semantic repository in order to facilitate the execution of association studies. In this context, the CxCa ontology was built to serve as the schema of a 'container' repository that enables the above unification.

The resulting data are employed by the project to perform association studies in an automatic way. The context of use of the CxCa Ontology is depicted in Figure 1.

## **3 DOMAIN BACKGROUND**

In this section we present some basic notions regarding the domain and try to provide the reader an insight about the ontology design requirements.

Cervical cancer is the second leading cause of cancer-related deaths after breast cancer for women between 20 and 39 years old (Landis et al., 1999) and one of the leading types of cancer affecting women worldwide. Despite a significant progress in early diagnosis and treatment of cervical cancer, there are more than 60,000 new cases and 30,000 deaths each year in Europe alone. Recently, it has been proved that infection by the human papilloma virus (HPV) is necessary condition for the disease (Walboomers et al., 1999). However, since HPV infection is highly unlikely to be the sole cause for developing cancer, recent trends in medical research combine genetic with clinical data and attempt to discover underlying associations of the disease with environmental agents, virus characteristics and genetic attributes, in order to identify new markers of risk, diagnosis and prognosis.

Diagnosis of CxCa is based mainly on three types of examinations, namely *cytology*, *colposcopy*, and *histology*. A problem that arises here is that examinations results are often encoded in Hospital Information Systems (HIS) using different coding standards, and/or custom formats.

On these grounds, a unification procedure is essential. For medical research purposes (ie, for conducting association studies), four levels of discretization have been considered adequate for each examination (Agorastos et al., 2009). These are presented in table 1.

Furthermore, in order to combine the findings of these types of examinations, the severity of the disease is again quantized into four discrete levels, a number that was considered adequate to capture it's different stages (Agorastos et al., 2009). When the aggregate result needs to be based on more than one types of examinations, the following practice is assumed. If a histology has been conducted, then it's result is considered regardless of the existence of other results. In other words, histology is regarded the "golden standard" examination having the highest reliability. Then follow colposcopy, and, finally, cytology.

Another important aspect of the clinical research procedure is that when more than one examinations of the same type have been conducted, the "worst" (ie, most severe) is considered valid.

The same thing holds when a patient is associated with more than one cases (ie, various series of examination and treatment procedures at different moments in the course of time). Again, for this person, only

Table 1: Summary	of the adopted	classification scher	ne for CxCa stages.

Normal (+Within Normal Limits)		
Low Grade Cervical Intraepithelial Neoplasia (LCIN)	stage 1	
High Grade Cervical Intraepithelial Neoplasia (HCIN)		
Invasive Cervical Cancer	stage 3	

it's "worst" case is taken under consideration for the execution of an association study.

## **4 ONTOLOGY STRUCTURE**

The conceptual schema of CxCa Ontology basically consists of two types of entities (concepts).

1. *Patient Management Entities*. These are generic entities capturing information about patient records as they are stored in the corresponding Hospital Information Systems (HIS). Their organization is based on the concepts of *Case* and *Visit*.

A Case can be defined as "a collection of data referring to a patient for a certain period of time, within which a diagnosis on the disease is meaningful, i.e., makes medical sense" and is related to a *Person*. Each Case comprises of one or more Visits. Every Visit is essentially a medical record and may contain one or more *Medical Interventions*. Each Medical Intervention is associated with a single *Result*. However, since within the context of a Case multiple Medical Interventions of the same type with conflicting Results may be conducted, each Case is associated with one and only one *Collective Result* for each type of Medical Intervention.

When a diagnosis is made (i.e. the *Severity Index* of the disease is calculated) it refers to a single Case, taking into consideration all the Visits and interventions the latter may be associated with. This means that each Person is possibly related with more than one Cases with different Diagnoses through the *hasCase* object property. However, since for the purpose of performing an association study only one Case per Person has to be considered, the *hasWorstCase* object property associates each Person with its Case with the worst Diagnosis. A new Case is instantiated should the patient return after a long period of time, which would yield any previous examinations obsolete.

This type of entities also includes notions such as *Clinic*, *Medical Intervention*, *Result*, *Lifestyle Choice* etc.

2. Domain Specific Entities. These model terms that

are strongly associated with the CxCa disease, it's stages and the related interventions and genotypic and phenotypic factors. This type of entities also contains concepts that do not directly correspond to the information stored in the EHR and are essentially inferred through a reasoning process according to appropriate definition rules.

Examples of such entities include *Colposcopy*, *HPV Vaccination*, *MTHFR C667T Polymorphism*, *Stage1 CxCa Severity Index* etc.

The scheme adopted is designed to be generic enough to be potentially able to incorporate information about other types of disease as well. The key-concepts of it, along with their interconnecting properties, are represented in Figure 2.

Furthermore, the notion that the stages of the disease and the risk factors under investigation are not affected by interrelations between different patients, motivated us to design the ontology so that the ABox forms independent individual groups. This means that two Person entities are neither directly nor indirectly connected to each other. In this way, reasoning can be performed independently within each subgraph concerning a person, making the reasoning process an explicitly parallel task.

## 5 MEDICAL UNIFICATION RULES

The unification tasks of the ontology can be roughly divided into two categories. The first category translates results from various different coding schemes to a common format, while the second reveals implicit classification criteria.

#### 5.1 Result Unification Rules

The ontology contains all major standards for each diagnostic intervention. Their results are translated to a 4-category classification scheme for each type of examination. For instance, in cytology, the Munich, Bethesda and Pap coding schemes are considered and characterization of a Cytology Result as a Stage0 Cytology Result is based on satisfaction of the restriction posed in formula (1).

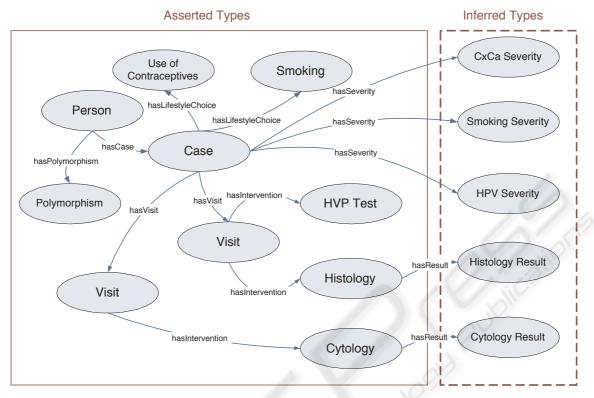


Figure 2: Basic Entities and their Interconnecting Properties.



5.2 Diagnostic Unification Rules

This type of rules aims at producing aggregate selection criteria for patient record retrieval.

An indispensable selection criterion when performing association studies is the Severity Index (i.e. the stage) of the disease. As there exist three types of diagnostic interventions that may be associated with a Case in our domain, calculation of the severity of the specific Case is based upon them, giving priority to existence of a Histology, in absence of which, considering Colposcopy and if both of the former are absent, Cytology is considered.

As Description Logics adhere to a strict open world assumption, it is impossible to deduce the absence of an examination result, as knowledge is considered incomplete. We overcome this problem by instantiating "triplets" of results (one for each examination) and explicitly storing a "NoResult" property for the examinations that have not actually been conducted. An example of such a rule is given in formula (2).

## **6** INFERENCE

Medical inference is essentially based on incomplete knowledge and is thus non-monotonic. Doctors suggest a treatment conjecturing about the most probable cause for some observed symptoms or examination results, disregarding theoretically possible but unlikely alternative causes. Moreover, these results may contradict one another.

stage1_Severity $\equiv$ Cervical_Cancer_Severity $\sqcap$	
(∃isDiagnosedBy	
(Case ⊓	
$((\exists \texttt{caseHasCollectiveResult} \{\texttt{stage1\_Collective\_Histology\_Result}\}) \sqcup$	
$((\exists \texttt{caseHasCollectiveResult} \{\texttt{NO\_Collective\_Histology\_Result}\}) \sqcap$	
$(\exists \texttt{caseHasCollectiveResult} \{\texttt{stage1\_Collective\_Colposcopy\_Result}\})) \sqcup$	
$((\exists \texttt{caseHasCollectiveResult} \{\texttt{NO\_Collective\_Histology\_Result}\}) \sqcap$	
$(\exists \texttt{caseHasCollectiveResult} \{\texttt{NO\_Collective\_Colposcopy\_Result}\}) \sqcap$	
$(\exists \texttt{caseHasCollectiveResult} \{\texttt{stage1\_Collective\_Cytology\_Result}\}))$	
)	
)	
) (2	2)

This is also the case in our setting. Before concluding to a triplet of results associated with a case in order to apply the rules of section 5.2, one has to produce a *Collective Result* for each examination type, taking into account all examinations of this type.

Because there may be an undefined number of each one of them, setting an upper limit of instances, creating all of them and applying the trick of "no result" as in section 5.2 is not an option here. In order to overcome this problem we choose to add an intermediate external inference step and consider closed world semantics by using appropriate queries in an RDF query language. This procedural layer executes sequentially a number of queries, implementing the set difference operator, in order of decreasing severity.

The process for doing this for cytology is as follows:

- First we query for cases containing cytologies associated with a stage3 Cytology Result. Obviously, since this is the worst outcome, these are considered the Collective Cytology Results for the corresponding cases.
- Then we query for examinations associated with a stage2 Cytology Result. This query also returns the Cytologies of the previous query, which are subtracted with a proper query, and the remaining ones are considered as the Collective Cytology Results for the corresponding cases.
- The steps are continued for the other two stages and the process is repeated for all examinations.

We employ the same methodology in order to identify the worst Case for each Person. Again, consecutive queries are employed, this time retrieving Persons w.r.t. their Collective Severity (as calculated with the rules of section 5.2), in order of decreasing severity.

In summary, the unification procedure consists of the steps depicted in Figure 3.

We have to point out that, because of it's nonmonotonic nature, this process must be repeated each time the ontology ABox is populated with new data.

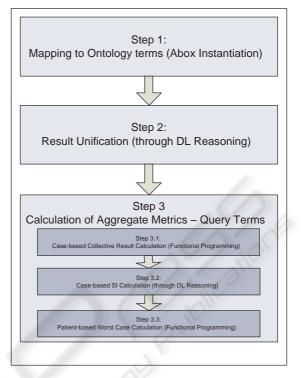


Figure 3: The Unification Procedure.

However, this does not yield problems for our purpose, as the system is research-oriented and does not need to be concurrent w.r.t. the EHR records of the associated hospitals.

## **7 IMPLEMENTATION DETAILS**

The Ontology was developed in Protege knowledge modeling tool (Noy et al., 2001), while the knowledge representation language employed was OWL-DL. Sesame (Broekstra et al., 2002) and OWLIM (Kiryakov et al., 2005) were chosen for storage and reasoning, respectively. DL reasoning is performed by the IRRE component of Sesame and the queries are expressed in SeRQL (Broekstra and Kampman, 2003).

The ontology TBox currently consists of 174 classes, 22 object and 96 datatype properties, 26 equivalent class axioms, while current instantiation of the ontology ABox includes about 680,000 individual 620,000 role assertions, containing data about 3,200 patients and 8,400 cases.

## 8 CONCLUSIONS

We have described the basic structure and inference mechanisms of a medical ontology in the domain of cervical cancer. The ontology is the central component of a system that aims at the unification of various virtual medical repositories and acts as a common language designed as a means for conducting association studies. Limitations and requirements of this cause have made the use of an ad-hoc reasoning scheme inevitable.

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