Identification of Similar Patients Through Medical Concept Embedding from Electronic Health Records: A Feasibility Study for Rare Disease Diagnosis

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Abstract. To identify patients with similar clinical profiles and derive insights from the records and outcomes of similar patients can help fast and precise diagnosis and other clinical decisions for rare diseases. Similarity methods are required to take into account the semantic relations between medical concepts and also the different relevance of all medical concepts presented in patients’ medical records. In this paper, we introduce the methods developed in the context of rare disease screening/diagnosis from clinical data warehouse using medical concept embedding and adjusted aggregations. Our methods provided better preliminary results than baseline methods, with a significant improvement of precision among the top ranked similar patients, which is encouraging for further fine-tuning and application on a large-scale dataset for new/candidate patient identification.

Keywords. Patient similarity, Electronic Health Records, word embedding, rare disease diagnosis

1. Introduction

The rapidly growing of clinical data, mainly electronic health records (EHR), presents unprecedented opportunities for precise and comprehensive patient characterization using high throughput phenotyping. In the context of rare disease, historical EHR data are inestimable sources for patient phenotyping, especially for patients with complex rare diseases. However, it is challenging for clinicians to examine and derive insight from multidimensional, large-scale EHR data [1]. Patient similarity, as a central topic

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in precision medicine to stratify patients into clinically meaningful subgroups, has also been considered in rare disease domains, aiming to identify patients with similar medical histories, diagnoses and outcomes, and derive insights from the records of similar patients to help personalized predictions [2].

The basic similarity model is vector space model (VSM), considering cosine-type similarity between patients represented as vectors of all his/her phenotypes. However, it does not take into account the dependence between phenotypes. To address this issue, the information content (IC)-based semantic similarities, such as Resnik [3] and Lin [4] similarity, are the most considered measures for concept similarity, which require topological structure of concepts in an ontological base. Another developing idea is word embedding, considering a vector representation of medical concept that encodes its meaning such that concepts with similar meaning are closer in the vector space, such as cui2vec learned from large collection of clinical notes [5], and HPO2vec learned from heterogeneous knowledge sources [6]. To obtain patient similarity, concept similarity should be aggregated, either through the average best match method as suggested in [7], or using average embeddings as suggested in [6].

In the context of C'IL-LICO program, we aim to develop transformative diagnostic, prognostic and therapeutic approaches for patients suffering from ciliopathies, a group of rare, severe and multi-systemic diseases caused by ciliary dysfunction. In this study, we describe the methods that we have developed for identifying similar patients from clinical data warehouse using medical concept embeddings and adjusted aggregations.

2. Materials and methods

2.1. NPH patients and Dr. Warehouse UMLS phenotyping

The Necker Children's Hospital is a national reference center for rare and undiagnosed diseases, hosting the Imagine research institute. The research data repository contains more than 600 patients with genetic diagnosis of nephronophthisis (NPH)-related diseases or syndromes in the NPH cohort, which is one of the major ciliopathy cohorts. The clinical data warehouse (Dr. Warehouse) [8] of Necker/Imagine Institute contains EHR data from more than 700,000 patients. The high throughput phenotyping module within Dr. Warehouse is based on the extraction of phenotypes encoded with the Unified Medical Language System (UMLS), because of its large coverage of medical concepts in French language.

As we considered the task of identifying NPH-related ciliopathies from other diseases, we selected NPH patients with sufficient information in EHRs, and re used the control patients with overlapping phenotypes described previously [9]. As renal involvement is one of the most frequent manifestations in NPH-related ciliopathies, this control cohort included patients with any kidney disease (C0022658 and all its descendants) excluding ciliopathies.

2.2. Concept similarity with clinical concept embeddings

To obtain UMLS concept embedding, first, a skip-gram fastText model [10] was trained with default parameters on a collection of 2.5 million clinical narratives from Dr. Warehouse to compute word embeddings of dimension 300 that reflect linguistic
contexts of each word. Then all labels for each UMLS concept (e.g. C0278511 with two French synonymous terms/labels “sarcome ostéogène localisé” and “ostéosarcome localisé”) were tokenized, the token embeddings were averaged to obtain label embeddings, and label embeddings were averaged to obtain the final concept embeddings. The semantic similarity between two concepts $p_1$ and $p_2$ was calculated using cosine similarity between two embeddings, denoted as $sim_{concept}(p_1, p_2)$.

2.3. Patient-patient similarity

To aggregate concept similarity into patient similarity, we first considered the average best match methods, i.e. for each concept in patient $P_1$, looking for the best matching concept in patient $P_2$, and averaging the best match similarity over all concepts in $P_1$. More precisely, for two sets of concepts (i.e., two patients), $P_1$ and $P_2$, the asymmetric set similarity is obtained with the formula:

$$sim_{set}(P_1 \rightarrow P_2) = \frac{1}{|P_1|} \sum_{P_{2j} \in P_2} \max_{P_{1i} \in P_1} sim_{concept}(p_{1i}, p_{2j}).$$ (1)

and the symmetric set similarity is the average of the two asymmetric set similarities:

$$sim_{set}(P_1, P_2) = \frac{1}{2} \left( sim_{set}(P_1 \rightarrow P_2) + sim_{set}(P_2 \rightarrow P_1) \right).$$

Here each concept in the set contributes equally to the set similarity. However, with the EHR data, both the number of patients per concept and the number of concepts per patient can be very heterogeneous. The relevant and specific concepts should be aggregated differently with higher weights. We thus developed the adjusted average best match methods:

$$sim_{set}(P_1 \rightarrow P_2) = \frac{1}{|a_1|P_1} \sum_{P_{2j} \in P_2} a_{1i} \max_{P_{1i} \in P_1} sim_{concept}(p_{1i}, p_{2j}),$$ (2)

where $a_1$ can present different weights of concepts in the set $P_1$. Three weighting methods were considered: (i) by distribution of phenotypes over patients (normalization 1), (ii) by distribution of patients over phenotypes (normalization 2), and (iii) by term frequency-inverse document frequency (tf-idf), as suggested in [9].

2.4. Performance evaluation

The EHR data of diagnosed ciliopathies and random controls were pooled. Patient-patient similarity was calculated for all patients. The patients are ranked by computing the average similarity with diagnosed ciliopathy patients, except his/herself, i.e. the diagonal similarity was removed to avoid the bias induced by the high similarity between the diagnosed ciliopathy patient and him/herself. The top k patients who are most similar with ciliopathies were predicted as highly suspected ciliopathies. The performance was assessed using metrics such as precision at k, i.e. the proportion of similar patients in the top-k set that are ciliopathies.

3. Results

3.1. Patient selection

The ciliopathy data set was composed of 79 patients with at least 6 distinct UMLS concepts extracted in EHRs. A set of 200 patients was randomly sampled from the
control cohort for this preliminary study, with the same restriction of minimum 6 concepts. The average number of distinct UMLS concepts was 64 for ciliopathy patients (max: 230, interquartile range: 15-90), and 55 for controls (max: 246, interquartile range: 17-76). The pooled dataset consisted thus of 279 patients, presenting 2593 distinct UMLS concepts.

3.2. Evaluation results

The precisions at \(k=30\), \(k=50\) and \(k=100\) are shown in Table 1 for the baseline VSM methods and our methods using embeddings and different adjusted aggregations. The area under the receiver operating characteristic (ROC) curves (AUC) is provided as well, although it is less informative for diagnosis task. The partial ROC curves were plotted in Figure 1 for \(k \leq 30\). The results showed that with a similar level of AUC, our developed methods with embeddings and adjusted aggregations could largely improve the performance for small values of \(k\).

**Table 1. Evaluation results**

| Precision@k | AUC |
|-------------|-----|
| \(k=30\)    | \(k=50\) | \(k=100\) |
| VSM norm1   | 0.43  | 0.46  | 0.36  | 0.62  |
| VSM norm2   | 0.53  | 0.48  | 0.45  | 0.70  |
| VSM tf-idf  | 0.50  | 0.48  | 0.43  | 0.69  |
| Embedding   | 0.53  | 0.42  | 0.39  | 0.64  |
| Embedding norm1 | 0.50  | 0.46  | 0.38  | 0.61  |
| Embedding norm2 | **0.70** | **0.60** | 0.43  | 0.69  |
| Embedding tf-idf | 0.67  | 0.58  | 0.43  | 0.69  |

Figure 1. Partial ROC curves for \(k \leq 30\). The precision@30 (P@30) is given at the end of partial ROC curves.

4. Discussion

In this article, we described the approaches that we have developed for identifying similar patients from clinical data warehouse using medical concepts embeddings and adjusted aggregations, which have taken into account the semantic relations between
medical concepts and the different importance of concepts. Our methods provided better preliminary results than the baseline methods, especially for small value of k, which is of particular interest as we aim to identify suspected ciliopathy patients at top ranked list who would benefit from genetic testing.

Our study had several limitations. As mentioned previously, the IC-based semantic similarity was often considered using for example Human Phenotype Ontology (HPO) as an ontological base [2] [7]. We did not use HPO for phenotyping because (i) it does not exist yet a reliable French version, and (ii) as a versatile thesaurus, the UMLS provides larger coverage of medical concepts. For the same reason as the latter, the UMLS network is less formally structured for semantic similarity. It would be interesting to compare IC-based methods and embedding-based methods. The embeddings could be improved with fine-tuning on more specific clinical records.

The next steps will be using fine-tuned medical concept embeddings to further improve the performance, and applying the patient similarity methods on a larger scale dataset for new/candidate patient identification.

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