Biomedical Named Entity Recognition

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Introduction

- Electronic health records (EHRs) are the databases used by general practitioners (GPs) to store the medical history of patients.
- Most of the information held in these EHRs is in the form of natural language.
- Therefore it is largely inaccessible for statistical analysis.
- Unlocking this information can bring a significant advancement to biomedical research.

Literature Survey

- Rule-based systems can extract medical information with good accuracy in simpler situations, such as information that follows regular speech patterns (Karystianis et al., 2018).
- Neural networks are being especially successful in complex NLP tasks (Young et al., 2017), where more traditional rule-based and other ML-based methods fail (Cambria & White, 2014).

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- For instance, a combination of the long short-term memory (LSTM) type of recurrent neural networks (RNNs) and a convolutional neural network (CNN) has been successfully applied to set new state-of the-art performance for NER tasks based on CoNLL-2003 and OntoNotes 5.0 data (J. Chiu & Nichols, 2016).
- Amounts of annotated text, which is especially troublesome for their application to EHRs. Namely, only a few publicly available datasets exist for medical NLP, and even fewer exist with annotations for NLP tasks (e.g. document classification; or slot filling). The best known of these datasets are MIMIC-III (A. E. W. Johnson et al., 2016) and i2b2 (i2b2, 2018, p. 2).

Problem Statement

To apply the concept of NER natural language processing on the Electronic Health Records in order to make the process of extracting the required information from the unstructured, huge, unorganized medial history of a patient in a more faster and efficient way to reduce the effort of the medical official doing the work.

Dataset

```
1 -DOCSTART- -X- -X- O
   -DOCSTART- -X- -X- O
                                                                             3 Clustering NN 0 0
 3 EU NNP B-NP B-ORG
                                                                             4 of NN 0 0
 4 rejects VBZ B-VP 0
                                                                             5 missense NN O O
 5 German JJ B-NP B-MISC
                                                                             6 mutations NN 0 0
 6 call NN I-NP O
                                                                             7 in NN 0 0
 7 to TO B-VP O
                                                                             8 the NN 0 0
 8 boycott VB I-VP 0
                                                                             9 ataxia NN O B-Disease
 9 British JJ B-NP B-MISC
                                                                            10 - NN O I-Disease
10 lamb NN I-NP O
                                                                            11 telangiectasia NN O I-Disease
11 . . 0 0
                                                                            12 gene NN 0 0
12
                                                                            13 in NN 0 0
13 Peter NNP B-NP B-PER
                                                                            14 a NN 0 0
   Blackburn NNP I-NP I-PER
                                                                            15 sporadic NN O B-Disease
15
                                                                            16 T NN O I-Disease
16 BRUSSELS NNP B-NP B-LOC
                                                                            17 - NN O I-Disease
   1996-08-22 CD I-NP 0
                                                                            18 cell NN O I-Disease
18
                                                                            19 leukaemia NN O I-Disease
19 The DT B-NP O
                                                                            20 . NN 0 0
20 European NNP I-NP B-ORG
                                                                            21
21 Commission NNP I-NP I-ORG
                                                                            22 Ataxia NN O B-Disease
22 said VBD B-VP O
                                                                            23 - NN O I-Disease
23 on IN B-PP O
                                                                            24 telangiectasia NN O I-Disease
24 Thursday NNP B-NP O
                                                                            25 ( NN 0 0
25 it PRP B-NP O
                                                                            26 A NN O B-Disease
26 disagreed VBD B-VP O
                                                                            27 - NN O I-Disease
27 with IN B-PP O
                                                                            28 T NN O I-Disease
28 German JJ B-NP B-MISC
                                                                            29 ) NN 0 0
29 advice NN I-NP O
                                                                            30 is NN 0 0
30 to TO B-PP O
                                                                            31 a NN 0 0
31 consumers NNS B-NP O
                                                                            22 nococcive NM O P Dicease
```

Methodology

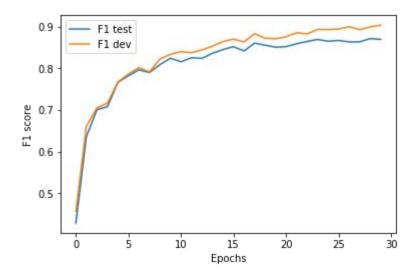
- The model has three inputs character-level, word-level and casing input.
- The architecture starts processing these three inputs independently and then merges them to process further.
- We create a word-level and character-level embedding and the set of unique words and labels from the three datasets are stored.
- These word and character embeddings are mapped to a 50 dimensional and 30 dimensional vector embeddings using the Global Vectors for Word representation.
- All these mapped tokens are undergone the process of casing where we consider 6
 different types of cases before they are mapped using the GLoVE model.
- We make use of the CNN-BLSTM model, a joint learning of shape, appearance and dynamics done by a deep learning technique. This includes a convolutional neural networks and bidirectional long short term memory(CNN-BLSTM).
- We use the word embeddings from the GLoVE model and use them to generate a dictionary of all possible characters from the available set of words.

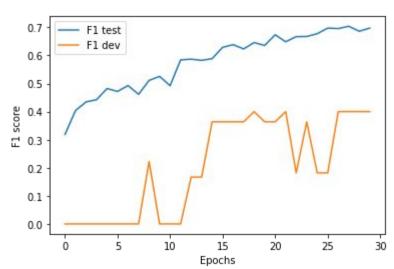
Contd.

- We tag the datasets with the numerical values and generate a class predictor.
- Build Model defines all the models that are called and we concatenate the processed character-level vector, word-level vector and casing data vector into a single vector.
- Dropout has been used with a rate of 0.5 to prevent the risks of overfitting in the model.
- The parameter we considered for the evaluation purpose if F1 score which is compared against the values as mentioned in the results sections.

Results

We are using F1-score as a metric for our model and the graph plotted over 30 epochs is pasted below. The first figure is the graph over medical data and the second one over non- medical data





References

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Thank You