# **Medical Knowledge-enriched Textual Entailment Framework**

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### Abstract

One of the cardinal tasks in achieving robust medical question answering systems is textual entailment. The existing approaches make use of an ensemble of pre-trained language models or data augmentation, often to clock higher numbers on the validation metrics. However, two major shortcomings impede higher success in identifying entailment: (1) understanding the focus/intent of the question and (2) ability to utilize the real-world background knowledge to capture the context beyond the sentence. In this paper, we present a novel Medical Knowledge-Enriched Textual Entailment framework that allows the model to acquire a semantic and global representation of the input medical text with the help of a relevant domain-specific knowledge graph. We evaluate our framework on the benchmark MEDIQA-RQE dataset and manifest that the use of knowledgeenriched dual-encoding mechanism help in achieving an absolute improvement of 8.27% over SOTA language models. We have made the source code available here.<sup>1</sup>

### 1 Introduction

The entailment task is similar with natural language inference (NLI), involves identifying the semantic similarity between two natural language texts, premise (P) and hypothesis (H). The NLI task's effectiveness is crucial for developing a robust natural language understanding system that functions at a human level (Ben Abacha et al., 2019; Abacha and Demner-Fushman, 2016; Romanov and Shivade, 2018). Recent literature suggests the use of contemporary language models (LMs) (Devlin et al., 2018; Beltagy et al., 2019), often ensembled, to achieve better performance (Zhu et al., 2019; Bhaskar et al., 2019; Xu et al., 2019) on the NLI task. However, our qualitative interpretation of the dataset and results suggests that LMs fails when it comes to textual entailment (TE), despite being the on-demand language model. The limitations belong to two major categories:

- Multiple word form: Medical text offers high degree of variability in the form of synonym and abbreviated words. The same can be witnessed in Table-1, where BERT (Devlin et al., 2018) is unable to predict the entailment between P and H as they have different terms 'Kartagener's Syndrome' and 'Primary Ciliary Dyskinesia', while both are synonym.
- Focus/Intent understanding: Given P and H, LMs often fails to capture the focus/intent of both the sentences. Table-1 shows an example, where the focus of P and H are misunderstood. It can be seen that P emphasizes the possibility of 'atypical pneumonia' occurring within a month after treatment, whereas H talks about the possible treatments for the disease.

These findings indicate that existing LMs lack semantic interpretation of the input, which is crucial in the inferencing tasks. In this paper, we deal with the question:

Does the medical textual entailment task benefit from the external domain knowledge to distinguish semantically identical medical sentences to recognize entailment?

To address this question and above-mentioned limitations, this paper presents a novel framework for

<sup>1</sup>https://github.com/VishalPallagani/Medical-Knowledge-enriched-Textual-Entailment

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Table 1: Examples from the MedQA-RQE dataset, where the text highlighted in blue are semantically similar words and red represents the lexically similar but semantically dissimilar words.

recognizing textual entailment, Sem-KGN: **Sem**antic Knowledge-enriched Graph Network that explores the domain-specific knowledge to enhance the semantic interpretability in the LMs. The proposed method devise a dual-encoding mechanism to enrich the classical document encoding (obtained from BERT) with the knowledge-enriched graph encoder. Specifically, our method builds a heterogeneous dictionary graph of the given P and H to encode the global context, while BERT's proficiency lies in capturing the local contextual information. Rather than constructing graphs solely based on the triples (subject, object, predicate), more *semantic units* are introduced into the graph as additional nodes to enrich the relations between the entities. These additional nodes add medical-entities centered factual information which are generated by expanding the medical knowledge graphs (KGs) such as UMLS (Bodenreider, 2004), SNOMED-CT (Donnelly, 2006) and ICD10 (Quan et al., 2005). The medical entities present in P and H are used to query the related information, 'diseases/syndromes', dosage, 'side-effects', and 'drug-interaction' from the mentioned KGs. Finally, Graph Convolutional Network (GCN) (Kipf and Welling, 2017) is employed to generate the graph encoding, augmented with the regular document encoder, and later fused through a multi-headed attention layer to support entailment.

**Contributions:** (i) Proposed medical-TE framework, by utilizing domain-specific medical KGs to encode the global and semantic information of the premise and hypothesis, (ii) Exploited the capabilities of semantic units in a graph network to encode medical-entities centered factual information for recognizing textual entailment, and (iii) Evaluated the effectiveness of the proposed method over state-of-the-art language models and knowledge-infused baseline methods on the benchmark MEDIQA-RQE dataset.

**Related Work :** The development of annotated TE and NLI medical datasets (Abacha et al., 2015; Ben Abacha et al., 2019; Abacha and Demner-Fushman, 2016; Romanov and Shivade, 2018) and a variety of pre-trained language models has led to a rise of extensive ongoing research in this field. Majority of the systems developed for the TE task adopts the multi-task learning (MTL) framework (Zhu et al., 2019; Bhaskar et al., 2019; Kumar et al., 2019; Zhou et al., 2019; Xu et al., 2019), ensemble method (Sharma and Roychowdhury, 2019), and transfer learning (Bhaskar et al., 2019) for achieving better accuracy. Xu et al. (2019) employed the MTL approach (Liu et al., 2019; Yadav et al., 2018; Yadav et al., 2019; Yadav et al., 2020) in TE task to learn from the auxiliary tasks of question answering (QA) and NLI. The best performing system at MedQA 2019-RQE shared task (Zhu et al., 2019) utilized the MTL approach to learn from intermediate NLI task. Further they used knowledge distillation approach to condense the information obtained from various models and transfer it into an single model. Few of the works (Wang et al., 2019; Bhaskar et al., 2019) in extracting information for the entailment task. However, the consideration of building a vocabulary graph from the textual and later enriching them with information from the medical KGs is still an unexplored territory.

## 2 Proposed Approach

The overall architecture of the proposed Sem-KGN is illustrated in Fig.-1. The rest of the section elaborates Sem-KGN in detail.

## 2.1 Document Encoder

Given a  $n_P$  words premise  $P = \{w_1^P, w_2^P, \dots, w_{n_P}^P\}$  and  $n_H$  words hypothesis  $H = \{w_1^H, w_2^H, \dots, w_{n_H}^H\}$ . The document encoder is responsible to capture basic lexical and syntactic in-



Figure 1: Architecture of proposed methodology

formation from the premise and hypothesis in the form of local context information. We employed the BERT model to serve as the document encoder in our proposed Sem-KGN framework. Formally, the local context features are computed as  $d_1, d_2, \ldots, d_n = \text{Document-Encoder}(w_1, w_2, \ldots, w_n)$ .

### 2.2 Knowledge-enriched Graph Encoder

**Vocabulary Graph Construction:** We first construct a dictionary based on all the unique words in the training dataset. Thereafter, we build a graph G = (V, E) based on the word co-occurrence information in the dictionary. Instead of building graph based on a given P and H, we were motivated by the work of (Lu et al., 2020) to build graph by considering all the lexicon in the dataset, which aims at encoding the global information of the particular domain (in our case medical domain). The nodes of the graph G are words in the dictionary, the edge between two nodes  $w_i$  and  $w_j$  is determine by the normalized point-wise mutual information (NPMI) (Bouma, 2009).

$$NPMI(w_i, w_j) = -ln \frac{p(w_i, w_j)}{p(w_i)p(w_j)} \frac{1}{ln \ p(w_i, w_j)}$$
(1)

where  $p(w_i, w_j) = \frac{\#s(w_i, w_j)}{\#W}$ ,  $p(i) = \frac{\#s(w_i)}{\#W}$ , #s(.) is the number of sliding windows containing a word or a pair of words, and #W is the total number of sliding windows. We make an edge between the nodes if the value of NPMI( $w_i, w_j$ ) exceed a particular threshold value.

**Graph-expansion with Medical Knowledge:** We expand the existing graph G with the additional nodes and corresponding edges to form a graph  $G^* = (V \cup \overline{V}, E \cup \overline{E})$ . Towards this, first we extracted the medical entities by exploiting the entity recognition model<sup>2</sup> trained on MedMentions dataset (Mohan and Li, 2019), from a given pair P and H. Once the medical entities are identified, KGs such as SNOMED-CT, ICD-10, UMLS and Clinical Trials are exploited to extract the information of two-hop connected 'diseases/syndromes', 'dosage', 'side effects', and 'drug-interaction' type medical-concepts. The final expanded entities are the additional nodes  $\overline{V}$  which act as the semantic unit to capture the domain-specific relationship (e.g., treats, caused by) and hierarchical relations (e.g., is a) between medical-concepts.

**Graph-expanded Knowledge Encoding:** The Graph-expanded Knowledge Encoder (GxK Encoder) computes the representation of each node from graph  $G^*$ . We are interested to compute the representation for each extracted entities  $(e_1, e_2 \dots e_m)$  for given pair of P and H. Formally, we get  $h_1, h_2, \dots, h_m = GxK-Encoder(G^*, e_1, e_2, \dots, e_m)$ 

We model it using the 2-layer GCN architecture. For the entities, we first make a input matrix  $M \in \mathbb{R}^{m \times |V|}$ , where row of the matrix M is the one-hot vector of length of |V| (size of dictionary). Given the adjacency matrix D of expanded graph  $G^*$  a and input matrix M, a single layer of graph convolution is computed as follows:

$$H^{(1)} = relu(MDW^{(1)}), \text{ where } D \in \mathbb{R}^{|V| \times |V|} \text{ and } W^{(1)} \in \mathbb{R}^{|V| \times d}$$

$$\tag{2}$$

<sup>&</sup>lt;sup>2</sup>https://go.aws/37SD7ae

where  $H^{(1)} \in \mathbb{R}^{m \times d}$  is matrix which rows are the node features in. For a given input M, we are interested to captures the part of the graph from D by multiplying them together as MD. The feature at a given node  $w_i$  computed by the interaction between all the neighbouring nodes as  $h_{w_i}^{(l+1)} = relu\left(\sum_j \frac{1}{c_{ij}}h_{w_j}^{(l)}W^{(l)}\right)$ where  $c_{ij}$  is the normalization constant (Kipf and Welling, 2017). The second layer convolution is obtained as  $H^{(2)} = relu(H^{(1)}W^{(2)})$ , where  $W^{(2)} \in \mathbb{R}^{d \times d}$ 

### 2.3 Multi-headed Aggregator

We fuse the information from Document-Encoder and GxK-Encoder using the multi-headed selfattention (Vaswani et al., 2017). Our aims to utilize the best of both the worlds. We form a single feature sequence by augmenting both the encoder representations obtained from both the encoders, with additional [CLS] token representation. By applying the self-attention on the augmented encoding sequence, we facilitate the network to attend the useful information across the individual encoding. We model our Multi-Headed Aggregator (MH-Aggregator) using the 12 layers of Transformer-block with 16 heads. We use the last layer output of the aggregator and consider the [CLS] token representation as the final representation  $F \in \mathbb{R}^d$  of given P and H pair. We employ a feed-forward layer to classify a pair of premise P and hypothesis H into the corresponding *'entail'* or *'non-entail'* classes.

$$f_1, f_2, \dots, f_{n+m} = \text{MH-Aggregator}(d_1, d_2, \dots, d_n, h_1, h_2, \dots, h_m)$$

$$prob(class = entail|P, H, \theta) = exp(W_{entail}^T F + b) / \sum_{i} exp(W_j^T F + b)$$
(3)

Models		Accuracy	Precision	Recall	F1-Score
Baseline 1	BERT	47.90	46.16	48.26	47.18
	BioBERT	50.14	46.28	48.97	47.58
	ClinicalBERT	49.60	48.79	49.56	49.17
Baseline 2	BERT + KI	49.56	46.06	48.69	47.33
	BioBERT + KI	51.15	48.56	49.56	49.05
	ClinicalBERT + KI	50.14	50.00	50.00	50.00
(Zhu et al., 2019)	BERT + linear projection	51.30	51.53	51.30	49.45
Proposed Model	Sem-KGN	56.17	63.18	56.18	59.47

Models Components	Accuracy	Precision	Recall	F1-Score
Sem-KGN	56.17	63.18	56.18	59.47
(-) Knowledge-enriched Graph Encoder	47.90 (8.27 ↓)	$46.16~(17.02\downarrow)$	$48.26~(7.90\downarrow)$	$47.18~(12.29\downarrow)$
(-) Medical Knowledge-graph	$50.65~(5.52\downarrow)$	$62.27~(0.91\downarrow)$	$50.66~(5.52\downarrow)$	55.86 (3.61 $\downarrow$ )

 Table 3: Ablation study showing the role of Table 2: Experimental results of our proposed model each component in the model. The values (Sem-KGN) and the baseline methods on the official within the bracket show the absolute decretest set.

## **3** Experimental Results and Analysis

**Dataset and Metrics:** We used widely adopted benchmark entailment dataset, MEDIQA-RQE created by (Abacha and Demner-Fushman, 2016), released in the BioNLP 2019 shared task. The dataset is derived from consumer health questions (CHQs) and frequently asked questions (FAQs) from the U.S. National Library of Medicine and National Institute of Health respectively. The training and validation set consists of total 8890 CHQ and FAQ pairs with the entail label of 4784 instances. The test set consist of 230 pairs with 115 entail labels. We used official evaluation metrics (Accuracy) to evaluate our model. Additionally, we also provided the Precision, Recall, and F1-Scores for the evaluation.

**Implementation Details:** We have chosen models' hyper-parameters empirically on the validation set. The base-uncased version of BERT<sup>3</sup> of hidden size 768 with a max sequence length of 200 (160 for P and 40 for H) is used in all experiments reported in the paper. The size of dictionary to create the dictionary matrix was 30000. The dimension of The threshold of NPMI is set to 0.3 to obtain meaningful relation between words. The last layer's hidden size of the graph-expanded knowledge encoder is set to 16. We use the Adam optimiser (Kingma and Ba, 2014) for parameters update after every epoch of training. We set the standard batch size of 16, and trained for 5 epochs with a dropout rate of 0.2 and 2e - 5 learning rate in all the experimental results reported in this work.

**Baseline Models:** To show the effectiveness of Sem-KGN, we adopted following competitive baseline models:

1. Language Models (LMs): We utilized the SOTA LMs (BERT) as well as the LMs adapted for the

<sup>&</sup>lt;sup>3</sup>https://bit.ly/2ZajZR1

Sample Examples and Error Type	Premise	Hypothesis	True Label	BioBERT+ KI	Sem-KGN
Example 1	Can you mail me patient information about Glaucoma, I was recently diagnosed and want to learn all I can about the disease.	How is glaucoma diagnosed ?	Non-entailed	Non-entailed	Non-entailed
Example 2	I was writing to inquire about more information regarding the diagnosis of OI. We have family members who are in the process of waiting for genetic testing to come back but are under allegations of child abuse. Is there any information that may be helpful to us?	How to diagnose Osteogenesis Imperfecta ?	Entailed	Non-entailed	Entailed
Error Type 1: Complex and Long Question	Hello, my dad, 68 years old, has gastritis, it did ache occasionally over the last several years. The other day, he went to hospital to have medical check-up with endoscopic ultrasonography, and found GIST with about 1cm in size. Dr. told him that he may consider surgery or not, it is up to him. What are we supposed to do?	How is an endoscopic ultrasound performed ?	Non-entailed	Entailed	Non-entailed
Error Type 2: Ambiguous Question	Can you please send me as much information as possible on "hypo- thyroidism". I was recently diagnosed with the disease and I am struggling to figure out what it is and how I got it.	How is Hypothyroidism diagnosed ?	Non-entailed	Entailed	Non-entailed

Table 4: Qualitative and error analysis of our proposed model (Sem-KGN) with the best baseline model.

#### clinical (ClinicalBERT) and medical domain (BioBERT), fine-tuned for MEDIQA-RQE task.

**2. Knowledge-Infused Language Models (+KI):** These baselines model works on the principle of *shallow knowledge infused learning*, where we provided the knowledge about the medical entities at the token level to the LMs. The intuition behind using this baseline was to understand at what layer if knowledge is integrated into the LMs, it is going to be beneficial.

**Results:** Table-2 provides an overview of the results, which demonstrates that Sem-KGN, equipped with KGs enriched graph encoding performs the best over all the baselines model. A considerable increase in the accuracy of 8.27% can be observed over vanilla BERT model in comparison to proposed Sem-KGN. The similar set of improvement (over 6%) can be observed with BioBERT and Clinical-BERT. We also observed the power of basic shallow (+KI) over the vanilla LMs showing the absolute improvement of 1.5%. Further, in comparison to baseline 2, Sem-KGN achieved the average increment of 6% over all the knowledge-infused LMs. Finally, from our ablation study (*c.f.* Table-3), it can be noticed that enriching the graph encoder with the domain knowledge assists in the entailment task. The results conclude two important claims: (1) the modularity of the knowledge infusion process that can be combined with any LMs is witnessed, and (2) Sem-KGN proves its effectiveness in having a local as well global understanding of the premise and hypothesis. We also compare the results against the best system (Zhu et al., 2019) at MEDIQA-RQE task that was based on the ensemble of LMs. However, to have a fair comparison and understand the role of our Knowledge-enriched graph encoder, we only utilize their model that have introduced "linear projection" over BERT. The proposed Sem-KGN has outperform the "linear projection" mechanism described in Zhu et al. (2019).

**Analysis:** Table-4 depicts the qualitative analysis of Sem-KGN, w.r.t baseline models on 50 randomly sampled DEV set. The first entry shows the effectiveness of Sem-KGN in being able to understand the focus of the premise and the hypothesis which is achieved by the dual encoding mechanism. The second entry in the table affirms the importance of domain-specific KGs in assimilating medical information.

**Error Analysis:** Table-4 entry 3 and 4 shows the leading cause of the error in the proposed model. We found that major misclassification occurred when the premise was complex and have a multiple questions. We also observed in some cases when there is high ambiguity between premise and hypothesis, model fail to recognize the correct label. For e.g., in the Table-4 entry 4, with the presence of term '*hyperthyroidism*' and '*diagnosed*' both in premise and hypothesis it is very difficult to recognize the true label: not-entailed.

### 4 Conclusion

In this paper, we proposed a framework Sem-KGN to recognize medical textual entailment. Our framework utilized the local context from BERT based document encoder and global context by expanding the vocabulary graph with the medical entities obtained from the medical knowledge-bases. We present an efficient aggregator scheme to fuse the multiple encoding. The proposed Sem-KGN framework outperformed the competent pre-trained language model and knowledge-graph enabled language model architectures with fair margin. In future, we plan to explore multi-domain knowledge-graph and efficient graph embedding based techniques for medical textual entailment.

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