



HHS Public Access

Author manuscript

IEEE J Biomed Health Inform. Author manuscript; available in PMC 2024 April 12.

Published in final edited form as:

IEEE J Biomed Health Inform. 2023 July ; 27(7): 3645–3656. doi:10.1109/JBHI.2023.3271580.

Forecasting User Interests Through Topic Tag Predictions in Online Health Communities

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Abstract

The increasing reliance on online communities for healthcare information by patients and caregivers has led to the increase in the spread of misinformation, or subjective, anecdotal and inaccurate or non-specific recommendations, which, if acted on, could cause serious harm to the patients. Hence, there is an urgent need to connect users with accurate and tailored health

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information in a timely manner to prevent such harm. This article proposes an innovative approach to suggesting reliable information to participants in online communities as they move through different stages in their disease or treatment. We hypothesize that patients with similar histories of disease progression or course of treatment would have similar information needs at comparable stages. Specifically, we pose the problem of predicting topic tags or keywords that describe the future information needs of users based on their profiles, traces of their online interactions within the community (past posts, replies) and the profiles and traces of online interactions of other users with similar profiles and similar traces of past interaction with the target users. The result is a variant of the collaborative information filtering or recommendation system tailored to the needs of users of online health communities. We report results of our experiments on two unique datasets from two different social media platforms which demonstrates the superiority of the proposed approach over the state of the art baselines with respect to accurate and timely prediction of topic tags (and hence information sources of interest).

Index Terms—

Topic tag prediction; healthcare information; time sensitive data; interest Forecasting; e-healthcare; health informatics

I. Introduction

PERSONAL healthcare management increasingly requires accommodating a higher degree of involvement of patients and informal caregivers. Consequently, patients and their caregivers expect healthcare providers to provide more detailed information about the disease, prognosis, and treatment. However, there is often a disconnect between healthcare providers and patients (and their caregivers) in terms of language used, and the information that is shared with the patients. The situation is not much better with information available through electronic health records (EHR) [6], [14], [19].

This situation is exacerbated by the fact that only 12% of U.S. adults have proficient health literacy [25], [55]. Health literacy is especially low among older adults, minorities, groups with low income and socio-economic status, leading to increased healthcare usage and cost and worse health outcomes [38]. Given the difficulty of comprehending health information available from healthcare providers or personal EHR, patients and caregivers increasingly turn to the internet and online communities. In such communities, patients and their caregivers interact with and seek information, e.g., regarding treatment options, side-effects of medications, etc., from their peers, in simple, lay terms.

While online communities can be useful sources of information and emotional support for patients, they also increase the risk of misinformation [35], or subjective [56], anecdotal and inaccurate or non-specific recommendations, which, if acted on by patients, could cause them serious harm. Hence, there is an urgent need to connect users with accurate and tailored health information in a timely manner to prevent such harm.

Typically, in an online community, a user seeking information posts a natural language query; Peers, or in some cases, subject matter experts (e.g., clinicians or nurses) respond

to the query or point to useful sources of information [50]. Somewhat more savvy users may be able to hone in on appropriate keywords or search tokens (tags) to use in order to seek relevant information on the web (Google, Bing, etc.), medical information boards (WebMD, Mayo, etc.) or specialized platforms (HealthUnlocked, PatientsLikeMe, etc.). The keywords used in such queries are suggestive of the respective user's information needs. However, due to low health literacy, users may not necessarily be able to identify the right keywords to use; and it can be tedious and often frustrating for patients to sift through the results retrieved in response to their queries. Against this background, we explore a solution that aims to lower the barriers for patients to obtain accurate and understandable sources, e.g., articles written for laypersons that meet their health information needs. In this study, we use data from an online community HealthUnlocked,¹ i.e., a social network designed to offer health-related information and emotional support for patients and their caregivers. A longitudinal study of HealthUnlocked has shown that it has positively impacted many patients [2] in terms of health outcomes and engagement in care. An example of the data extracted from HealthUnlocked is shown in Fig. 1. Though Fig. 1 shows only one post, a typical user interacts a lot more with multiple posts and provides their corresponding keywords (tags). We observe that in such a setting, information needs of a user may involve medications, side effects, treatment options and other time critical concerns. These needs depend on not only the patient demographics and health condition, but also, where the patient is with regard to progression of the disease (e.g., stage of cancer in the case of cancer patients), or treatment being given (e.g. chemotherapy), or response to treatment (e.g., side effects, cancer remission).

A. Research Question

This study is designed to answer the following research question: Can the user profiles together with the traces of interactions of the user with the community (e.g., keywords used in past queries) and information from similar users be used to effectively predict contextually relevant topic tags associated with the *future* information needs of users?

We hypothesize that patients with similar histories of disease progression or with similar courses of treatment would have similar information needs at comparable stages. In treating many diseases, e.g., cancer physicians follow a particular course of treatment that is customized based on the patient's profile and how they are responding to treatment. For example, the National Comprehensive Cancer Network (NCCN) has laid out guidelines for the treatment of non-small cell lung cancer [10]. If two users, say *A* and *B*, have similar histories of disease progression and treatment, we can expect that their information needs at comparable stages are likely to be similar. In Fig. 2, we illustrate an example with two users who have similar histories of disease progression and treatment. In this case, the topic recommendations for user *B* can be improved by utilizing user *A*'s past keywords. In this case, user *B* can be recommended articles related to "*concurrent chemotherapy and radiation*".

¹[Online]. Available: <https://healthunlocked.com/>

Based on the preceding discussion, we arrive at the following problem specification. For any given user U in the community, with a user profile B^U , temporally ordered sequence of n prior posts $\{P_{t-n}^U, \dots, P_t^U\}$, encoded by the corresponding keyword sets $\{G_{t-n}^U, \dots, G_t^U\}$, and indexed by their position in the sequence as well as the past information traces of all the similar users in the network, the task is to predict the topic tags G_{t+1}^U associated with a hypothetical future post P_{t+1}^U that describes the future information needs of the user U .

Disambiguation: “Tags” and “Keywords” can be used interchangeably in our work as they both describe the abstract token used to represent a larger body of text. However, we try to differentiate by using “Keywords” for tokens that have occurred prior to prediction and “Tags” as tokens that are yet to be predicted (future).

B. Contributions

We propose an innovative approach for predicting future information interests of participants in online communities as they move through different stages in their disease or treatment. We report results of our experiments on an expert curated data set from which we demonstrate the superiority of the proposed approach over the state of the art baselines with respect to accurate and timely prediction of topic tags (and hence information sources of interest).

The key contributions of this work include:

- A novel approach to predicting topic keywords associated with the future information needs of users (patients, caregivers) in an online health community. We step away from traditional bag-of-words classification and towards a new sentence generation based topic prediction.
- A novel methodology for collaborative-hybrid filtering based topic recommendation which leverages similarity of user disease progression timelines.
- A semantic evaluation of predicted tags along with the traditional metrics such as precision and recall.

C. Related Work

The vast amounts of healthcare information available on the internet has led to the requirement of recommendation systems that help users navigate their care by enabling them with accurate information [1], [24]. Additionally, Deng and Liu [8] discuss the role of such systems in reducing risks like misinformation and time delays in access to accurate medical information. Yu et al. [53] review the recommendation models including the usage of collaborative filtering like [23], [28] and [41] and finds its relevance in healthcare topic recommendation. Unlike other user-centric recommendation systems, healthcare based models are a lot harder to be subjected to collaborative filtering as there is high degree of individuality in the information sought by the users [7]. Sahoo et al. [39] however, propose a CNN based feature aggregation approach to perform user-content based recommendation and presents improvements over personalization metrics. Similarly, Jiang et al. [18] create

a similarity matching model based on heterogeneous networks to perform collaborative filtering for healthcare topic recommendation.

Extracting features from text documents is at the centre of our proposed topic recommendation system. Text documents are in essence a sequence of vectors placed together with context and grammar. Early approaches to model text included the usage of recurrent neural networks (RNNs) like in [15] and [36]. It became evident that the order (positions) of these vectors in relation to their value was vital in text based tasks. Bidirectional long-short term memory models (LSTMs) including [20], [33] and specifically healthcare applications like [17] emphasized the importance of positional embeddings of the word vectors and improved performance on tasks like translation and sequence-to-sequence text generation. Transformers, a recent advancement by [46] along with their attention modules have revolutionised feature extraction like in [49]. Bidirectional Encoder Representations from Transformers (BERTs) an advancement of transformers have additionally aided in text based tasks like classification [13], [30], summarization [29], [34] and text generation [5], [27]. They are ideal for sentence/token generation conditioned upon prior context [12].

Table I compares our work in view of current literature. The position of current healthcare recommendation literature along with advances in natural language processing technology pose the ideal setting for a novel medical information recommendation system that predicts the future interests of users through topic tags.

II. Data

We obtain our primary training data from [HealthUnlocked.com](https://www.healthunlocked.com) (HU), which is one of the largest online health communities. The website has several sub-communities organized around specific health conditions or diseases. In our work, we focus on three such communities designated for lung cancer patients and their caregivers: *Lung Cancer Support*, *The Roy Castle Lung Cancer Foundation* and *British Lung Foundation*. Our choice of the communities was motivated by three considerations. First, managing chronic conditions and treatment regimes associated with lung cancer presents information needs that can be met through sustained interactions with the community. Second, chronic conditions impact the information needs of users over time, which are not adequately addressed by existing recommendation systems. Lastly, these communities represent the largest fraction of the users of the online community.

Each user has an opportunity to create a profile which includes demographic data, e.g., age, gender, etc. as well as the health condition. Once registered, a user then may pick one or more communities to join. Once in a community, users can post a “Main-Post” or post a “Comment” on an existing “Main-Post”. Additionally, each “Main-Post” can have associated “keywords” at the time of posting. These user supplied keywords can be considered to be indicative of the information needs of the user at the time and act as ground truth for training and evaluating the proposed tag prediction system.

We acquired de-identified data from the lung cancer communities for the period between 2011 and 2020. From the three lung cancer communities, we have a total of 1711 unique users. Note that ethical (or IRB) approval was not needed for our study because the data were de-identified at source (at HealthUnlocked), there is neither direct interaction nor intervention with human subjects, and there is no means to link the de-identified data to specific subjects. For this study, we focus on the subset of anonymized users with at least one post and only valid posts with a minimum of three words or more are considered. The resulting data now filters down to 1032 unique users. Each user has an anonymized hashed User-ID that links their profile-bio, posts, and keywords to the user. Among the 1032 unique users, the average posts per user is about 2.4 (minimum is 1 and maximum is 68). Each post has on average 3.2 keywords (minimum is 2 and maximum is 10). About 55% (574) of the users have a profile-bio. An example of a user's information is provided in Fig. 1.

Due to the unstructured nature of the text in our data set, there is a need to create temporal traces for training which can be used by our model. For this purpose, we pose the user data as described below.

A. Structuring the Information Traces of Users

We organize the information associated with each user, including the profile bio and the traces of the user's interaction with the community (information seeking posts or comments) as shown in Fig. 3. Here we assume that a user U creates a profile-bio B before they post their first post P_0 . At time T_{-1} , we use the bio to predict tags \hat{G}_0 where $\hat{\wedge}$ indicates the prediction. This prediction is made before the first post has been created. As time progresses, the user posts P_0 at time T_0 with keywords G_0 . We then accumulate the profile bio B , the post P_0 and keywords G_0 to predict the tags \hat{G}_1 of a future unseen post P_1 . We continue to do this with a temporal window of Pn posts in the past along with the current post to predict tags of a future post. At time T_n , though we have n posts from the past, using all the posts and their keywords could add too much noise into the topic tag prediction system. We empirically question the influence of history (see Section IV-A2) and determine the ideal Pn . We then use only the last Pn posts, their associated keywords, and the profile bio which remains unchanged throughout time T_0 to T_n , as well as the current post and corresponding keywords to predict a future topic tag \hat{G}_{n+1} .

B. Example Generation

To create a training example, we need an input sequence (input text document) containing profile-bio, posts and keywords and a target sequence (Tags of future post). This is commonly known as sequence-to-sequence training. In our setting, we increase our training example pairs by extracting all possible pairs of training data as shown in Fig. 4. In order to maintain chronological patterns in the data, we pick pairs that have information from consecutive time steps. For example, if *Post-1* and *Keywords-1* are from the latest time step, the target can only be *Keywords-2* and cannot be used to predict *Keywords-3*. Using such a rule, a single user's timeline is converted into several chronological training pairs. Following this, we obtain a total of 10,791 training pairs from the 1032 unique users. From this dataset, we use 8415 random pairs (78%) for training and the remaining 2376 (22%) pairs for testing.

C. Tag Statistics

Our training set has 472 unique tags while the test set has 375 tags. Among the 375 tags, 18 of them are not in the training set and are completely new words. However, since these words occur in the posts of the users, they are a part of the vocabulary that the models can observe and thus are not deleted from the test set. Fig. 5 shows the distribution of the top 20 tags in the test set. It is evident that ‘*Chemotherapy*’, ‘*Cancer and Tumors*’ occur several times more frequently than other tags. Tag prediction models designed around classification could incorrectly pick frequently occurring tags leading to poorer performance. Additionally, their metrics tend to be sensitive to the tag frequency in the training data. We address this issue of skewness and the resulting bias in Section IV-C.

D. Validation on Facebook Data

In order to validate generalizability of our approach to data from other social media platforms, upon training our proposed model, we ran validation on data extracted from a Facebook group named *Chronic Pain Support Group*.² The data was extracted through web-crawling of posts between 2015 to 2021. All identifiable information like user-names, mentions and locations were removed. From the complete list of 11219 users who had posts, we selected those who have at least 4 consecutive posts. This resulted in a total of 1253 users. It should be noted that the training is only performed on the HU data and the Facebook data is reserved for testing. Unlike the HU dataset, the Facebook data does not contain ground truth tags provided by users. Here, we instead use a term frequency-inverse document frequency (tf-idf) method to extract 5 tags from each post and use them as the ground truth reference. Similar to the HU dataset, we first structure the information traces and generate examples for the purpose of validation.

III. Methodology

Recent advancements in the field of natural language processing (NLP) have opened a realm of possibilities due to a better understanding of context from text data. Bidirectional Encoder Representations from Transformers (BERT) [9] have further allowed text representations to be learnt from vast sources like the Book Corpus and the English Wikipedia with more than 3300 million words. Their utility has been recognized in the bio-informatics community through the proposal of Bio-BERT [26], an extension of the BERT, trained over the PubMed and PMC medical word corpora. The extension of the pre-trained models helps in inferring context, more effectively than the traditional frequency based models, from text containing medically relevant words including those from online health communities. We leverage the Bio-BERT to extract illness-relevant information and predict future topics of interest based upon the users’ needs.

Our pipeline is intended to extract context from the unstructured data and recommend accurate and time-relevant tags for each user in the online health community database. We make several modifications from works in the literature to accommodate our data and increase the overall accuracy of tag prediction. Among these, is our usage of Bio-BERT

² <https://www.facebook.com/groups/11864244228>

model as a sentence generator instead of a simple bag-of-words classifier. Additionally, we show the need for collaborative filtering but move away from traditional K-means method to a disease timeline based clustering method. In this section, we will describe our proposed tag prediction procedure, the model architecture, the objective function for performing the training and the metrics used to compare the models in our experiments.

A. Tag Prediction

Our primary objective in this work is to predict future topic tags for users in order to forecast their medical interests. Unlike existing tag prediction algorithms, we do not conform to a classification approach where a model picks the most probable tag from a set of tags. Instead, we perform an auxiliary sentence generation (ASG) task. The ASG task is similar to sentence completion where the last word is masked out using the *[MASK]* token. In our set-up, the sentence generated is a set of comma separated future tags with no grammar or punctuation. We fine-tune a pretrained Bio-BERT model to read a given input document and pick a word from its entire vocabulary that fits the context and position of the mask. A similar training approach has been employed by [5] for generating questions based on a paragraph of context and an answer sentence. Since we are using a Bio-BERT model Tag generation, we refer to our model as **BBERT**_{T_g}. An example of the training process has been shown in Fig. 6. First, we have a *[CLS]* token to indicate the start of the input document. The text in Orange is the profile-bio followed by a *[SEP]* separation token. Similarly, in Blue, we have the past and current posts, the Green text corresponds to tags from previous and current posts, the tags from neighboring users or shared tags (see Section III-D) are in Red and the predicted future tags are in Purple. We experimented with the positioning of these individual elements in the input document. However, they did not lead to a significant difference in the results. Having accumulated all the parts of the input document, we first clean the text through removal of stop words and lemmatization. Following this, the words are converted to vectors using a BERT word-embedding and tokenization module which converts the sentences into tokens based on the position of the words in the sentence and the position of the sentence in the input document. This encoded data is then fed to the Bio-BERT model along with the BERT-tokens like *[CLS]*, *[SEP]*, *[MASK]* etc.

The pre-trained Bio-BERT model uses the originally prescribed BERT_{base} model with 12 layers, 768 hidden dimensions and 12 attention heads. We followed the training procedure set by [5] with the usage of an Adamax optimizer set with an initial learning rate of 5e-5. The training is performed over 15 epochs with 8415 examples while 2376 examples are used for testing. We use softmax as the final layer of the model where the Bio-BERT computes the probabilities of 200K+ words. Following literature, we adapt the Beam search [11] method to pick the top words and decode the predicted Bio-BERT output into regular words (tags).

Modification for Bio-BERT with long text Documents—The traditional transformer model is meant for short sentences as computing attention over large input documents is computationally expensive and requires lengthy training. Our input documents range from about 14 words to 550+ words when multiple data sources like profile-bio, posts, keywords etc. are combined. We modify our fine-tuned model based on the recommendations provided

by [3] where a “longformer” is proposed. This reduces the attention computation from quadratic to linear space. Gated Recurrent Units (GRUs) and Long-Short Term Memory (LSTM) models, though not entirely bidirectional in attention, are however capable of handling such large documents without any modifications.

B. Objective Function

In our sequence-to-sequence based learning approach the input sequence is the input document containing profile bio and posts and the output sequence is the target tags set. Both the sequences are assumed to come from a single long sentence where the target sequence is masked. We first tokenize the input sequence x into n tokens, $\{x_i\}$, $i = 1, 2, \dots, n$, and predict the sentence fragment $x^{p:q}$ where p and q are positions such that $p < q < n$ using the specially masked input which can be denoted as $x^{/p:q}$. Unlike traditional tasks like translation, x does not have a paired y but we instead have to rely on the conditional probability of a token appearing next in the sequence given the past tokens (until the r -th position, $p \leq r < q$) of the sequence. Let us denote the domain from which x is obtained as χ , the log likelihood objective to learn the model parameters θ then becomes:

$$L(\theta; \chi) = \frac{1}{|\chi|} \sum_{x \in \chi} \log P(x^{p:q} | x^{/p:q}; \theta). \quad (1)$$

Expanding the conditional probability as a product of probabilities at each position between p and q , we obtain (2). Here, r is the position at which the [MASK] token is placed.

$$L(\theta; \chi) = \frac{1}{|\chi|} \sum_{x \in \chi} \log \prod_{r=p}^q P(x_r^{p:q} | x_{<r}^{p:q}, x^{/p:q}; \theta). \quad (2)$$

The position, or r , can be controlled to dictate the amount of prior information in the input document. In our experiments, we always have a non-empty set of tokens for prior data coming from either Profile Bios or Posts. However, (2) can be used with null prior data where it will generalize into the OpenAI GPT model for generating text [42].

C. Metrics

To evaluate the tag generation model, we can view it as a recommendation system which recommends topic tags. This is similar to medication recommendation evaluation by [31]. We can generate any number of tags once the model is trained as this would only involve generating longer sentences. Traditionally, if K is the number of tags recommended, Recall@ K and Precision@ K are used to evaluate the success of the recommendation. In order to enable fair comparison using both Recall and Precision, we use the popular F1@ K metric where K is the number of predictions made and F1 is calculated with equal weights to both recall and precision.

We go beyond the traditional evaluation and calculate cosine similarity in the feature space between the recommended tags and the ground truth because the tag generation network is not picking words from a list but instead generating new words with a certain degree of variety. To calculate the cosine similarity, we first extract the word embeddings (features) of the K predicted tags and individually check their similarity to the word embeddings of the ground truth tags. If the ground truth tag is ‘*radiotherapy*’ and the predicted tag is ‘*radiation therapy*’ the penalty will not be too large.

D. Model Improvement Through Collaborative Filtering

Our approach for model improvement comes from the intuition that “*similar users seek similar information*”. Collaborative filtering is an approach to view a user as a parametric function of interests defined by other similar users in the community. Multiple recommendation systems including [21] and [44] utilize collaborative filtering to extract a deeper understanding of user needs and provide recommendations based on other users in the group.

Identifying similar users is a crucial step in performing collaborative filtering. In our setting, with healthcare data, we identify similar users by matching relative disease timelines. Users with similar illness progression tend to search for similar information. Information search history of users who have already traversed the timeline can be utilized to recommend topics to users trailing along similar paths. We hypothesize that the similarity in treatment profiles for chronic conditions like non-small cell lung cancer (NSCLC), can be leveraged to recommend information at different time intervals based on the disease progression.

Neighbor Tags: For a query user U with a post P^U and keywords G^U posted on date T , we identify nearest neighbors h_1, h_2, \dots, h_n using the novel disease timeline matching method described below. For each neighbor h_j , we check for posts that have occurred prior to T . The keywords associated with such a post will be then referred to as *Neighbor Tags* or S_T and are appended to the input document after the posts and their corresponding keywords (see Fig. 6). The entire input document is then used for future topic tag prediction.

Disease Timeline Matching for Identifying Neighbors: We describe the procedure for our novel approach using the following steps:

- The unstructured profile-bios are converted into a structured 13-column disease timeline using a Named Entity Recognition (NER) Bio-BERT model.
- A Bio-BERT model was also used to extract features from the disease timelines and were clustered in the feature space using t-Stochastic Neighborhood Embedding (t-SNE).
- The h nearest neighbors are identified using Euclidean distance in the two dimensional t-SNE feature space.
- Keywords of neighbors which have occurred prior to query post are used as neighbor tags (S_T).

- Neighbor tags are then appended to the input document before being provided to the BERT_{T_g} model.

Conversion of Profile-Bios: In order to provide structure to the profile bios, we contextually sorted the text into various classes. The classes created are shown in Fig. 7. Where “Other” class corresponds to words that do not belong to any of the other 12 classes. “Current” refers to the time when the bio was written, “Past” and “Future” are with respect to the “current” time. As it can be observed, the data is forced to take a temporal shape in order to extract user-disease timeline. This annotation was done by two subject experts who manually read each profile bio and filled in the corresponding columns. We compared the similarity of the annotations and obtained a cosine similarity of 0.811 and a Cohen’s kappa of 0.784, which is only 0.16 short of “perfect agreement” [32]. An intersection of the labels between the annotators is used as ground truth for the NER text classification task.

In order to automate this conversion between unstructured profile bios to structured timeline data, we trained a NER model to perform a 13-class classification of profile-bio text. We perform classification only to obtain the temporal disease timeline and not the topic tags. We trained two models namely (i) Bio-BERT [26] and (ii) LUKE-Deep Contextualized Entity Representations with Entity-aware Self-attention [51]. The Bio-BERT is only fine-tuned on our dataset while LUKE was trained entirely from scratch on our dataset. This was done to evaluate the value of information provided by the Bio-BERT pre-training. For both models, we used the B,I,O labelling procedure (Beginning of a label, Inside a label and Outside a label) to maintain standardized practices followed in the NLP community and to increase reproducibility. In the training set, we used 414 profile bios with 44,320 total words. In the non overlapping test set, there were 19,014 words from 160 bios. The models were trained for 25 epochs on a Nvidia DGX Tesla V-100 GPU for approximately 3.5 hours for fine-tuning Bio-BERT and 18 hours for training the LUKE. The Bio-BERT model outperformed the LUKE classifier in the following metrics- (a) The overall accuracy of the Bio-BERT model was 80.59% while LUKE achieved 73.63%. (b) The average recall percentage for Bio-BERT was 74.98% while only 66.29% for LUKE. The imbalance in the dataset skews the predictions which lead to a high false positive rate (see Fig. 8). The precision for each of the classes is low due to the large number of words in the “Other (OUT)” category. The confusion matrix for Bio-BERT along with its class-wise precision and recall is shown as well. For all the downstream tasks that follow, we use the Bio-BERT model fine tuned on our dataset as the primary feature extractor. In cases when the user has not provided an input for any of the above mentioned classes, or if the model fails to pick the provided input, the model predicts an empty string for that particular class and we use an ‘NA’ token instead and proceed with feature extraction.

Feature Extraction and t-SNE: The next step is a clustering process to perform unsupervised clustering based on similarities in the column values. The 13 columns can be treated as a sentence vector for feature extraction. We employed the previously used Bio-BERT fine tuned on profile-bios to extract the features of the structured text data and map it to a two dimensional t-distributed stochastic neighbor embedding (t-SNE) space. Unlike K-means [54], with t-SNE, we do not have fixed clusters. Areas in the 2D plot (see

Fig. 9) can be loosely defined as topics and users can belong to more than one particular topic similar to soft-clustering [22]. Features extraction can be performed from our filtered temporal disease timelines as well as directly from the profile bios. The comparison of the two feature extractions is visualized as shown in the t-SNE plots acquired after 750 iterations in Fig. 9. As it can be seen, features extracted from temporal timelines following NER form natural clusters compared to features directly extracted from profile bios. This is primarily due to the noise removal inherently performed by the NER task. For illustration, we pick a random user (#248) and check the nearest 3 users' profile-bios. Inferred from Fig. 9(b), the clustering is around *COPD* and *Emphysema*. For the purpose of clustering, we find the nearest h users to the primary user using Euclidean distances in the t-SNE feature space. We accumulate all keywords of the neighbourhood users (S_T), occurring prior to time T of current post and append it to the input document of the primary user. At the end of clustering, a particular user has the following items in their input document: (a) The profile-bio B^U , (b) The previous posts and their corresponding keywords (c) The current post and its keywords and (d) Neighbor tags from similar users (S_T). This input document is used to predict future tags G_{T+1}^U .

IV. Experiments

A. Model Ablation

We first determine the best settings for our proposed $BBERT_{Tg}$ for the obtaining the highest tag prediction F1 score.

We experiment with 4 settings namely

- **Input Document Accumulation:** Accumulation of different elements of the input document- (1) Profile Bios only (B^U), (2) Profile Bios & Posts (B^U+P^U) (3) Profile Bios, Posts & Corresponding keywords ($B^U+P^U+G^U$).
- **Influence of History:** The number of past posts (P_n^U) to use for predicting future tags.
- Different types of user clustering.
- Ideal number of neighbors for collaborative-filtering.

1) Input Document Accumulation: Significance: To understand the importance of different components of the input document.

We conducted three experiments with increasing amount of information presented in the input document. We start with the profile bio if available and then add posts and their corresponding keywords in the subsequent trials. For this experiment, we use 2 past posts along with the current post ($P_n = 3$). The average results with 4 runs each are shown in Table II.

2) Influence of History: Significance: To understand the translation of user interests over time through posts.

Our dataset is primarily based on chronic health conditions, therefore, utilizing the past information through past posts is vital for predicting future topics of interest. In this experiment, we determine the ideal number of previous posts (P_n^U) required. $P_n^U = 1$ corresponds to using only the current post while $P_n^U = 4$ accumulates the last 3 posts along with the current post. For this experiment, we use the $B^U + P^U + G^U$, referred to as full mode of data accumulation. Table III details the average results for 4 runs of this experiment.

3) Type of Clustering: Significance: To identify the best performing user clustering method. We experiment with three forms of clustering namely:

- Grouping of user profile-bios using K-means clustering.
- Using profile-bio text features extracted from Bio-BERT model for feature space clustering (t-SNE).
- Novel disease timeline based similarity matching of users, which is our proposed and adopted approach.

For K-means clustering, we vectorized the words using scikit-learn [37] **TfidfVectorizer** package and used a gap statistic based elbow approach [43] to identify the ideal number of clusters. We experimented with 2 to 80 clusters and found the characteristic elbow at 34 clusters. Following this, we fit the profile bios into 34 multi-user clusters. The average number of users in each of these clusters were approximately 11. We chose clusters with at least 4 users (1 query user and 3 neighbors).

In the case of profile-bio based feature extraction, we replaced the TfidfVectorizer with a Bio-BERT model to improve the semantic representation of profile-bios in the feature space. We reduced the Bio-BERT feature vectors to a two dimensional t-SNE space and found the 3 ($h = 3$) nearest neighbors using simple euclidean distances.

The final method compared is our novel temporal disease timeline matching approach described in Section III-D. The comparison is performed through improvement in topic tag prediction F1 score as detailed in Table IV. Please note the improvement in F1 score at K=5 level with 5 tag predictions.

4) Ideal Number of Neighbors: Significance: To identify the ideal number of neighbors to be used for tag sharing.

Having established the ideal clustering model, we experiment with number of neighbors to determine influence of similar users in the community. Table V presents the results with 5 levels of neighbors.

B. Model Comparison

1) Comparison With HealthUnlocked Dataset: In the second set of experiments, we compare our best model (Full-Mode with shared tags, $P_n = 3$, $h = 3$) with the current state-of-the-art (SOTA) topic tag recommendation techniques using the same metrics. We also establish a baseline using Latent Dirichlet allocation (LDA) method of tag prediction

with an implementation similar to [4], [16]. We identified two SOTA models based on architectural and feature extraction differences. The first is HashRec [48] and the second is Attention-based Multimodal Neural Network Model for Hashtag Recommendation (AMNN) [52]. HashRec is a hashtag recommendation model for social media text, specifically, conversational text on Twitter and Weibo. The model uses two encoders to encode the user posts and the conversation individually, followed by a bi-attention module to capture their interactions. The extracted features are further merged and fed into the hashtag decoder consisting of sequential Gated Recurrent Units (GRUs). We provide the first encoder with Profile Bios and the second encoder with a combination of prior posts, their keywords and shared tags from the neighborhood. We train the model to produce future tags sequentially. The default (Twitter) hyper-parameters and settings were used. AMNN is an attention based model for predicting tags from multi-modal data. The model uses two individual encoders, one for text and one for images consisting of bi-directional Long- Short Term Memory (LSTM) modules followed by an attention unit. The features are concatenated and fed to a sequential GRU model to predict tags. We modified the network to have two text encoders by removing the image encoder and the remainder of the network is unchanged. Similar to the previous model, the profile bios are provided to one of the encoders and the remaining input document is given to the next text encoder. We used the default hyper-parameters and settings to obtain results. The SOTA models are trained and tested on the same training set and test set used for our Bio-BERT (BBERT_{Tg}) model. Table VI reports the results of this experiment. The best performance is in **bold** while the second best is underlined.

2) Validation on Facebook Data: In this experiment, we use the previously trained models to predict topics of interest for users of the Facebook group *Chronic Pain Support Group* (data described in Section II-D). Each model is provided with 2 previous posts and 1 current post (3 posts in total) and asked to predict the topics of the unseen 4th post. A total of 1310 examples from 1253 users were used for the validation. This out of distribution prediction is a challenging task as the models are only trained on lung cancer data while the validation is entirely on the chronic pain dataset. We compare our top model (BBERT_{Tg}) against other models with 5 ($K = 5$) tag predictions. The average results from 4 runs of this experiment are presented in Table VII.

C. Data Bias Experiments—Classification based models even with Bio-BERT embeddings do not perform as accurately as text generation models due to the imbalance in the prediction classes (tags). We show this by comparing a Bio-BERT classification (BBERT_C) model against our proposed tag generation model (BBERT_{Tg}). This is included in Table VI. Very frequently occurring tags can skew results as they make it easier for models to guess the most frequent tag multiple times. We checked for the robustness of LDA and the BBERT_{Tg} models by deleting the top two tags (*Chemotherapy* and *Cancer and Tumors*) from the train and test sets. This experiment is to measure the impact of tag frequency on the models. As each example in the test set had 5 or more tag targets, dropping two tags did not require omission of test examples. Table VIII shows the F1 scores when ($K = 3$) tags are predicted with the two modified models.

V. Interpretation of Results

A:

Modeling information seeking behavior in a healthcare setting requires identifying key variables like current stage of disease, disease similarity and history of interests. We designed our ablation studies to identify these factors and leverage the information to develop an accurate temporal tag prediction model.

A1: From Table II we observe that the amount of information presented to the $BBERT_{Tg}$ model largely affects the model's ability to understand topics of interest. While the profile bios are instrumental in capturing overarching context and history, they poorly correlate with the users' dynamic interests when considered individually, yielding the lowest F1 score. Thus utilizing multiple data sources like the posts and their corresponding keywords help in improving predictions.

A2: As anticipated, the number of previous posts in the input document causes variations in the tag prediction metrics (see Table III). The posts in our dataset are asynchronous which means the time gap between them can be as small as 5 minutes or as large as multiple years. As the disease progresses, users have varying interests and if the time gap is larger, two or more consecutive posts will have completely different topics discussed. This confuses the model to pay attention to incorrect topic-keywords which are no longer of interest. We notice that the F1 score is the highest when 2 previous posts along with the current post which is the ideal trade-off point.

A3-A4: Improvements to the model come through collaborative filtering, an additional source of information regarding similar users in the database. In Table IV, experimenting with random neighbors yielded lower F1 scores at all levels indicating that including search tags from random users only caused the model to pay attention to noisy inputs. While on the other hand, Bio-BERT features extracted from cleaned NER disease timelines yielded the cleanest set of neighbors who positively contributed to the tag prediction metrics. The shared tags (S) from neighbouring users helped the Bio-BERT model predict better as it is likely that they have queried about this topic in the past. An interesting observation from Table V is that the number of nearest users did not affect the cosine similarity significantly. We believe the reason for this is that since cosine similarity is calculated in the feature space, the user clusters are also defined in a similar feature space. Due to this, adding more users does not negatively affect cosine similarity. It did however point out that using more than 3 ($h > 3$) neighbors could induce noise in the model due to uniqueness in the timelines. Hence we use $h = 3$ or three neighbors for our final model.

B1: In Table VI, we illustrate comparison of our proposed $BBERT_{Tg}$ model against the state of the art networks. Tag prediction models tend to perform well when the tags to be predicted depend only on the input document provided. However, understanding context and predicting the a future tag is a non-trivial task that leads to low F1 score with the HashRec and AMNN models. To overcome this, we pose the problem as a sentence generation task instead of the traditional classification task and obtain benchmark performance. We even

use the Bio-BERT embeddings in a classification task ($BBERT_C$) to show that sentence generation performs better due to robustness to skewed tag distribution which is usually the case with social media based text data. To show that our model is truly understanding the context, we show an example of generated tags for a future post by providing the previous posts, tags, profile-bio etc to all the models in comparison. The *future* post **not** provided to the model was: *“My partner was diagnosed with stage 4 lung cancer (nslc) in June 2009. He has had Chemotherapy, Radiotherapy and Tarceva. He started taking the new unlicensed drug Afatinib yesterday. So far no side effects! He has been told that the side effects can be more severe than Tarceva. I really hope this drug works for him as we are fast running out of options.”* Our $BBERT_{T_g}$ model was the only one to predict the tag *Afatinib* and also understand that the patient is undergoing a drug trial.

B2: In this validation experiment using Facebook data, Table VII demonstrates the superiority of $BBERT_{T_g}$ across all metrics even without collaborative filtering as it extracts and understands context and relies on the vocabulary to generate tags unlike other models which are restricted to their trained keywords. Please note that due to the unavailability of Profile-Bios in the Facebook dataset, we do not perform collaborative filtering and thus we only use the $BBERT_{T_g}$ (full-mode) for experiments. The model’s ability is exemplified in the qualitative comparison presented in the second part of Table IX. The Facebook post from the Chronic Pain dataset that was **not** provided to the model was: *“Amitriptyline was most effective but once the disc tore it stopped working, and the dosage was raised but only caused fatigue and weight gain, and withdrawal caused the itching to start that has not gone away completely.”* In the validation trial, $BBERT_{T_g}$ was able to predict tags unrelated to lung cancer and closer to chronic pain while all other models had some cancer related tags.

C: Finally, we emphasize that our $BBERT_{T_g}$ model is robust to tag occurrence frequency and show that even when the 2 most frequent tags are removed, the performance does not change by much when compared to a traditional LDA model as shown in Table VIII.

VI. Summary AND Discussion

In summary, our topic recommendation system, a hybrid system combining collaborative-filtering and content based ideas is designed for online health community users and utilises patient timeline similarity to group similar users. We use a pretrained Bio-BERT to perform the tasks of classification and sequential sentence generation. Our predictions of future topics yield accurate and contextual tags. We compare our model against models proposed for similar tasks like HashRec and AMNN and observe a superior performance.

We have empirically presented the quantitative advantage of our methodology over existing models. This improvement in precision and recall metrics across two unique datasets is attributed to two specific contributions- (1) An auxiliary sentence generation task for tag generation instead of tag classification and (2) Usage of “disease timelines” to match similar users and perform collaborative filtering. Additionally, we also performed ablation studies to understand the importance of several factors like history and influence of similar users in the network. As a part of future work, we will focus on two specific aspects. First, we intend to improve the matching of similar users by leveraging interactions and graphs. Second, we

plan to utilize the predicted tags to retrieve tailored medical articles from trusted sources like WebMD and Mayo clinic.

Acknowledgment

We also would like to thank [HealthUnlocked.com](https://www.healthunlocked.com) for providing the data used in this work.

This work was supported in part by the NSF Center for Health Organization and Transformation (CHOT), in part by the Penn State Institute of Computational and Data Sciences (ICDS), and in part by the Federal Ministry of Education and Research (BMBF), Germany under the project LeibnizKILabor under Grant 01DD20003.

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De-Identified, Random User-ID	User Profile-Bio
c285f04c817cbc11e	"Husband has Stage IV lung cancer, currently on Zometa. Advice please"
User Post (Post #1 of 9)	User Provided Keywords for Post #1
"My husband has recently been diagnosed with Stage 4 lung cancer, with mets in nodes, liver, adrenal. Non-smoker. Tested negative for targeted chemo. Starting his chemo on X"	Lung Cancer Zometa Chemotherapy

Fig. 1.

Example input snippet from HealthUnlocked; The data is de-identified by HealthUnlocked where dates and names within a post are removed. The date of the post, however, is made available.

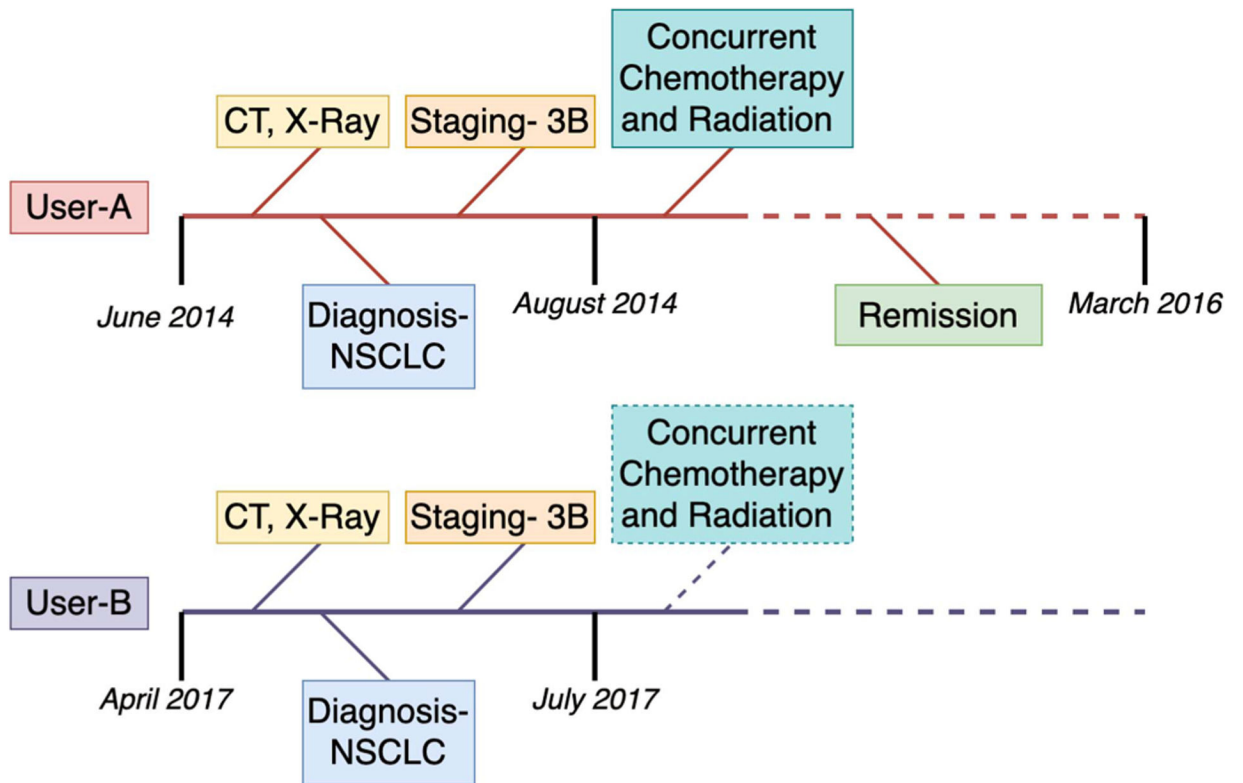


Fig. 2.
Similarity in relative disease timelines can help cluster similar users.

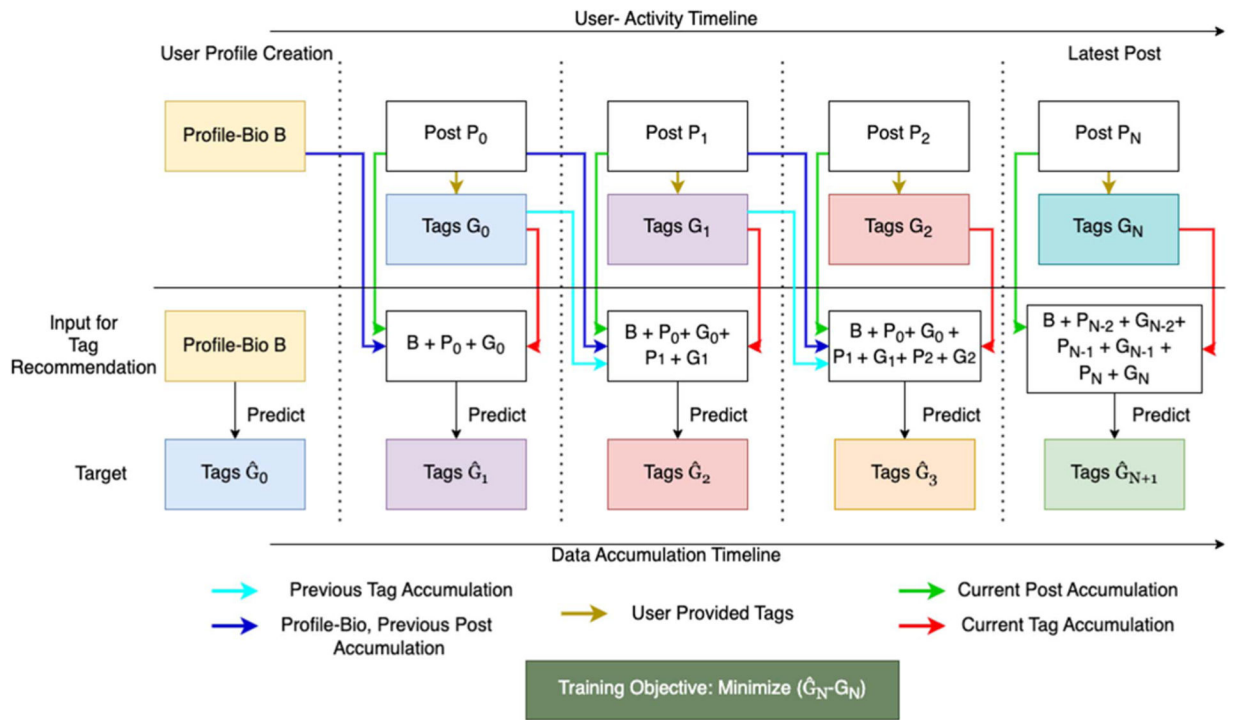
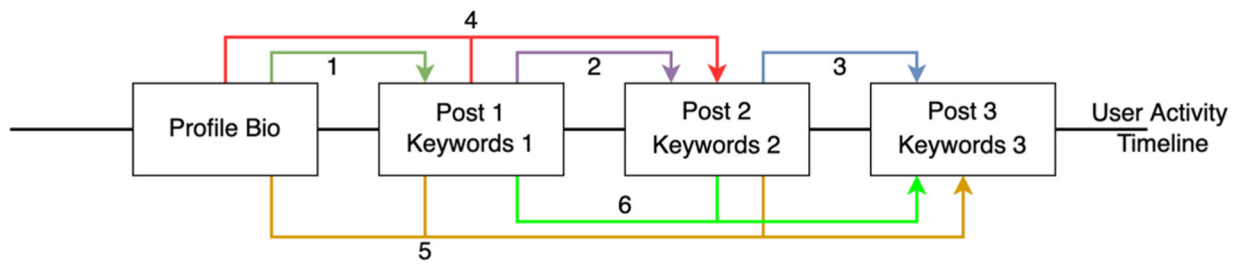


Fig. 3. User Timeline and Data accumulation for temporal Tag Prediction.









Training Example	Input Document	Training Target
1 	Profile-Bio	Keywords 1
2 	Post 1, Keywords 1	Keywords 2
3 	Post 2, Keywords 2	Keywords 3
4 	Profile-Bio + Post 1, Keywords 1	Keywords 2
5 	Profile-Bio + Post 1, Keywords 1 + Post 2, Keywords 2	Keywords 3
6 	Post 1, Keywords 1 + Post 2, Keywords 2	Keywords 3

Fig. 4. Timeline from a single user with a profile-bio and 3 posts is converted into 6 training examples.

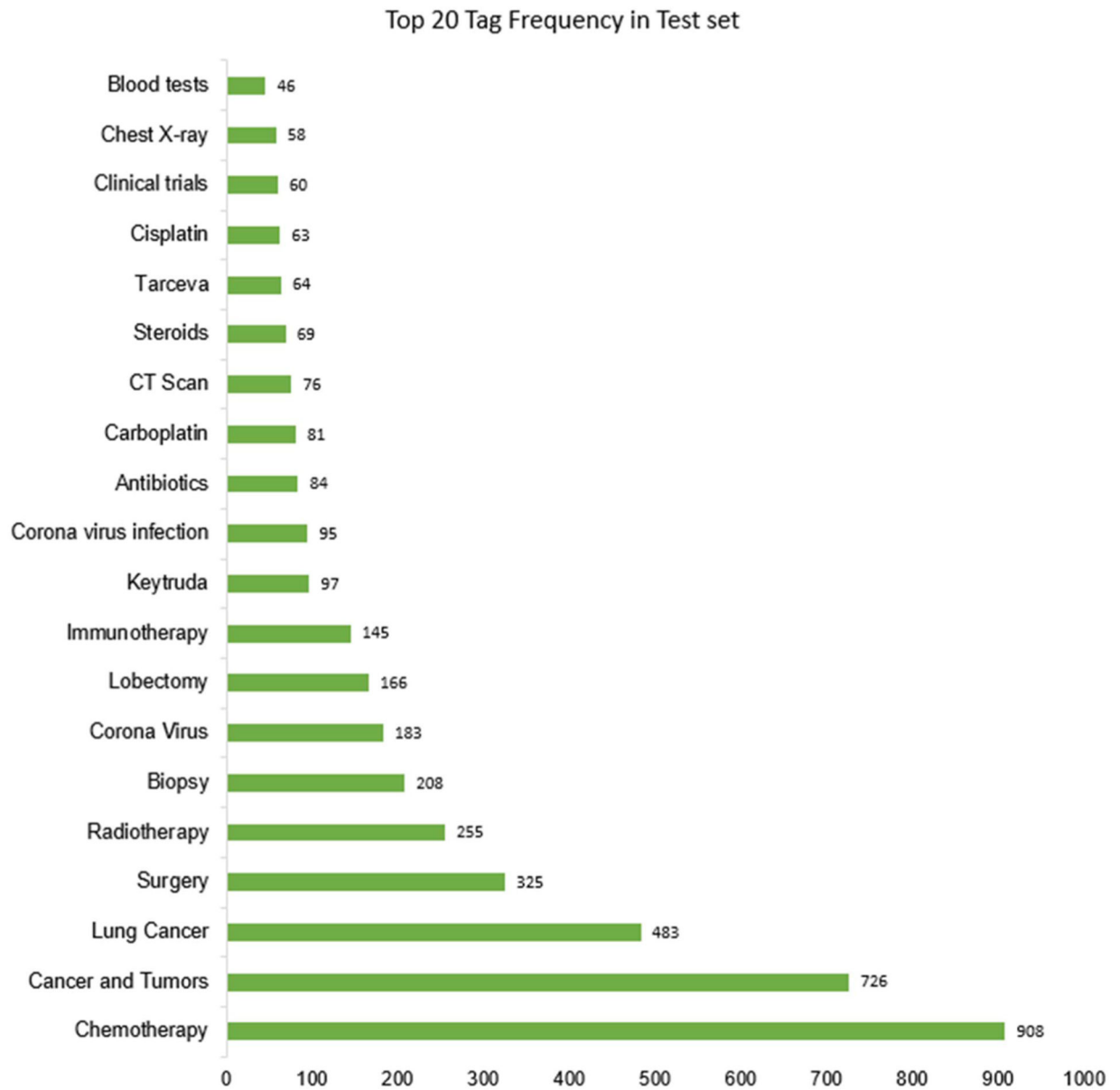


Fig. 5.
Plot of 20 most frequently occurring tags in test set.

Input Document D_i

Masked Target Token

Iteration 0	[CLS] Hi , X 39 diagnosed non small cell lung cancer stage 3b oct 2007 concurrent chemotherapy radiotherapy trial used hospitals following completion trial . working ward sister specializing pancreatic cancer unfortunately unable carry working although n't give easily [SEP] finished radiotherapy left bad cough nasty taste mouth suggestions to make cough easier part cancer tracher makes breathing quite hard [SEP] lung cancer, radiotherapy, drug trial [SEP] chemotherapy, non small cell lung cancer, radiotherapy, cancer [SEP] the future tags are: [MASK]	respiratory				
Iteration 1	[CLS] Hi , X 39 diagnosed non small cell lung cancer stage 3b oct 2007 concurrent chemotherapy radiotherapy trial used hospitals following completion trial . working ward sister specializing pancreatic cancer unfortunately unable carry working although n't give easily [SEP] finished radiotherapy left bad cough nasty taste mouth suggestions to make cough easier part cancer tracher makes breathing quite hard [SEP] lung cancer, radiotherapy, drug trial [SEP] chemotherapy, non small cell lung cancer, radiotherapy, cancer [SEP] the future tags are: respiratory [MASK]	tract				
Iteration 2	[CLS] Hi , X 39 diagnosed non small cell lung cancer stage 3b oct 2007 concurrent chemotherapy radiotherapy trial used hospitals following completion trial . working ward sister specializing pancreatic cancer unfortunately unable carry working although n't give easily [SEP] finished radiotherapy left bad cough nasty taste mouth suggestions to make cough easier part cancer tracher makes breathing quite hard [SEP] lung cancer, radiotherapy, drug trial [SEP] chemotherapy, non small cell lung cancer, radiotherapy, cancer [SEP] the future tags are: respiratory tract [MASK]	infection				
Iteration 3	[CLS] Hi , X 39 diagnosed non small cell lung cancer stage 3b oct 2007 concurrent chemotherapy radiotherapy trial used hospitals following completion trial . working ward sister specializing pancreatic cancer unfortunately unable carry working although n't give easily [SEP] finished radiotherapy left bad cough nasty taste mouth suggestions to make cough easier part cancer tracher makes breathing quite hard [SEP] lung cancer, radiotherapy, drug trial [SEP] chemotherapy, non small cell lung cancer, radiotherapy, cancer [SEP] the future tags are: respiratory tract infection [MASK]	[SEP]				
Iteration 4	[CLS] Hi , X 39 diagnosed non small cell lung cancer stage 3b oct 2007 concurrent chemotherapy radiotherapy trial used hospitals following completion trial . working ward sister specializing pancreatic cancer unfortunately unable carry working although n't give easily [SEP] finished radiotherapy left bad cough nasty taste mouth suggestions to make cough easier part cancer tracher makes breathing quite hard [SEP] lung cancer, radiotherapy, drug trial [SEP] chemotherapy, non small cell lung cancer, radiotherapy, cancer [SEP] the future tags are: respiratory tract infection [SEP][MASK]	radiotherapy				
Iteration 5	[CLS] Hi , X 39 diagnosed non small cell lung cancer stage 3b oct 2007 concurrent chemotherapy radiotherapy trial used hospitals following completion trial . working ward sister specializing pancreatic cancer unfortunately unable carry working although n't give easily [SEP] finished radiotherapy left bad cough nasty taste mouth suggestions to make cough easier part cancer tracher makes breathing quite hard [SEP] lung cancer, radiotherapy, drug trial [SEP] chemotherapy, non small cell lung cancer, radiotherapy, cancer [SEP] the future tags are: respiratory tract infection [SEP] radiotherapy [MASK]	[SEP]				
	Profile Bio	Post(s)	Corresponding Keywords	Neighbor Tags	Predicted Tags	BERT Tokens

Fig. 6.

Example of a sequential training routine of $BBERT_{T_g}$ with 2 tag generations. The input document has multiple sentences and each sentence is separated by $[SEP]$ token. Each iteration has a specific target. The target token at each iteration is appended at the position of the $[MASK]$ token in the next iteration. A $[SEP]$ token is added after each tag is predicted.

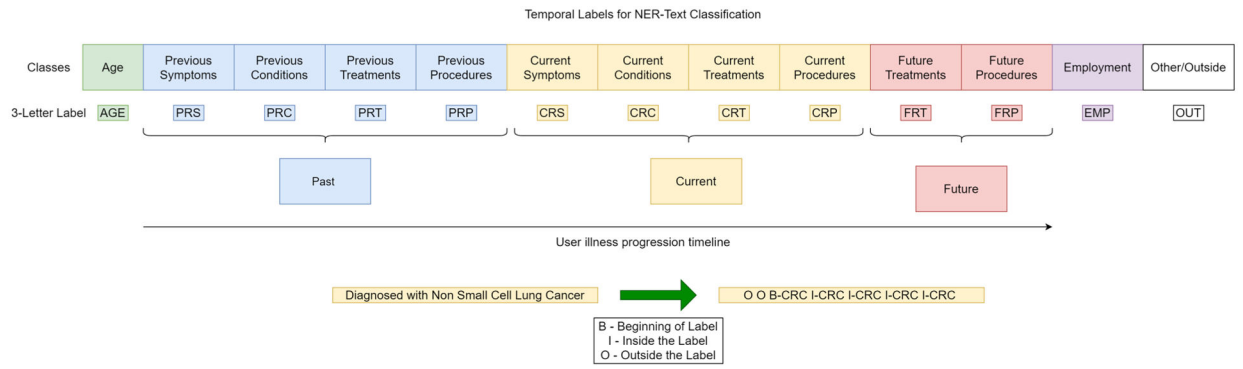
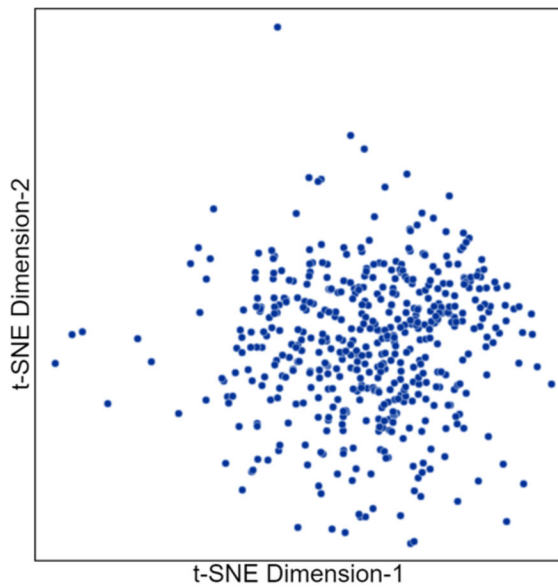


Fig. 7. Temporal Named Entity Recognition task Labels based on illness progression.

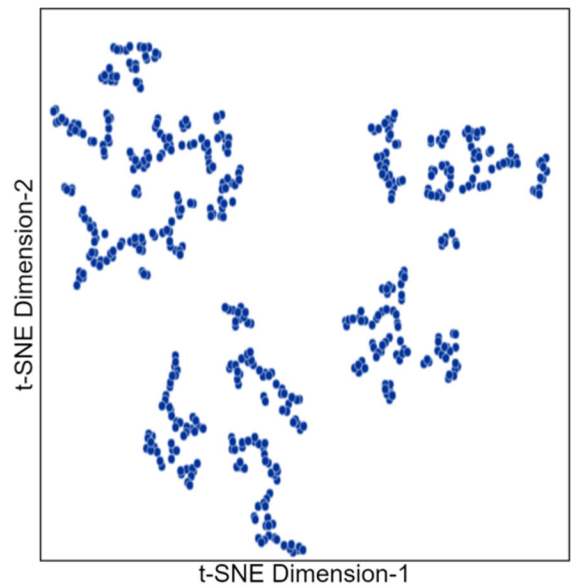
		True Labels													
		AGE	PRS	PRC	PRT	PRP	CRS	CRC	CRT	CRP	FRT	FRP	EMP	OUT	Precision %
Predicted Labels	AGE	100	1	0	0	0	2	1	1	0	0	0	0	331	22.93
	PRS	0	340	9	5	1	10	11	6	2	2	1	2	209	56.85
	PRC	0	16	1694	12	10	20	44	10	2	0	1	7	278	80.89
	PRT	2	5	24	417	6	11	17	20	2	3	0	0	319	50.48
	PRP	0	3	20	9	127	10	4	6	7	0	1	0	131	39.93
	CRS	0	43	18	17	2	301	7	3	2	5	0	1	310	42.45
	CRC	0	27	77	22	6	16	1487	8	3	2	1	7	296	76.17
	CRT	0	8	20	31	5	8	17	200	12	2	0	3	208	38.91
	CRP	0	3	12	9	11	5	13	1	57	7	1	3	207	17.32
	FRT	0	2	5	29	4	7	11	7	1	62	4	1	62	31.79
	FRP	0	2	18	9	1	5	1	1	2	10	22	2	35	20.37
	EMP	0	2	3	1	2	6	3	4	1	0	0	285	221	53.97
	OUT	28	7	41	8	2	24	33	11	4	1	3	12	10233	98.32
	Recall %		76.92	74.07	87.27	73.28	71.75	70.82	90.17	71.94	60	65.96	64.7	88.23	79.69

Fig. 8. Confusion Matrix from Bio-BERT NER classification of profilebio text in test set.



Closest 3 Bios to User #248
Married with three sons .Age 80 .Have Bronchiectasis and arthritis .
Lung disease due to building trade asbestos but still smiling living and enjoying
Had condition called oral lichen planus inside both cheeks

(a)



Closest 3 Bios to User #248
COPD sarcoidosis emphysema pulmonary hypertension
Hello , I came here to find out about COPD, emphysema in particular .This forum has helped me so much .
I have COPD (emphysema) , Asthma and allergic rhinitis among other health condition

(b)

Fig. 9. (a) t-SNE plot of features extracted directly from profile-bios of 414 users, Closest 3 bios (shortened for image) to random user #248; (b) t-SNE plot of features extracted from NER temporal-classified user timeline of 414 users, Closest 3 bios (shortened for image) to random user #248.

Differentiating Existing Models and Our Approach

TABLE I

#	Traditional Methods	Our Method
1	Tag Prediction	Generating new words from vocabulary
2	Tag Prediction Training	Synthesize auxiliary sentences
3	Objective	Prediction of tags for unseen future document
4	User Clustering	Based on disease progression similarity
5	Vulnerability	Robust to tag distribution
6	Evaluation Metrics	Cosine similarity in feature space along with P@K, R@K

Influence of Data Accumulation in Input Document, $P_n = 3$, Averaged Over 4 Individual Runs

TABLE II

Input Document	$K=1$				$K=3$				$K=5$			
	Recall	Precision	Cosine Similarity	F1	Recall	Precision	Cosine Similarity	F1	Recall	Precision	Cosine Similarity	F1
B^U (Only profile bio)	0.205	0.487	0.491	0.288	0.430	0.296	0.476	0.350	0.483	0.187	0.428	0.269
$B^U + P^U$	0.274	0.791	0.866	0.407	0.517	0.442	0.830	0.476	0.594	0.363	0.826	0.450
$B^U + P^U + G^U$ (Full mode)	0.296	0.874	0.909	0.442	0.590	0.584	0.891	0.586	0.681	0.402	0.887	0.505

Influence of Past Posts in Input Document With Full Mode Data Accumulation, Averaged Over 4 Individual Runs

TABLE III

Number of past posts in Input Document	K= 1				K=3				K= 5			
	Recall	Precision	Cosine Similarity	F1	Recall	Precision	Cosine Similarity	F1	Recall	Precision	Cosine Similarity	F1
$P_{p=1}$ (Only current post)	0.271	0.791	0.866	0.403	0.536	0.453	0.856	0.491	0.597	0.381	0.832	0.465
$P_{p=2}$ (One previous post)	0.277	0.820	0.909	0.414	0.545	0.567	0.879	0.555	0.657	0.395	0.876	0.493
$P_{p=3}$ (Two previous posts)	0.296	0.874	0.909	0.442	0.590	0.584	0.891	0.586	0.681	0.402	0.887	0.505
$P_{p=4}$ (Three previous posts)	0.286	0.856	0.887	0.428	0.593	0.577	0.887	0.584	0.671	0.381	0.852	0.486
$P_{p=5}$ (Four previous posts)	0.284	0.823	0.871	0.422	0.555	0.502	0.874	0.527	0.649	0.377	0.834	0.476

TABLE IV

Comparison of User Grouping Approaches With $h = 3$ Neighbors, $Pn = 3$ and Full Mode Data Accumulation, Averaged Over 4 Individual Runs; No Neighbors and Random Neighbors Are Included as Baselines

Clustering Method	$K=1$				$K=3$				$K=5$			
	Recall	Precision	Cosine Similarity	F1	Recall	Precision	Cosine Similarity	F1	Recall	Precision	Cosine Similarity	F1
No Clustering, No Neighbors	0.296	0.874	0.909	0.442	0.590	0.584	0.891	0.586	0.681	0.402	0.887	0.505
No Clustering, 3 Random Neighbors	0.277	0.813	0.763	0.413	0.510	0.526	0.730	0.517	0.597	0.383	0.700	0.466
K-Means of Profile-Bios [54]	0.290	0.865	0.812	0.434	0.572	0.545	0.742	0.558	0.610	0.384	0.765	0.471
BBERT Features of Profile-Bios	0.301	0.880	0.911	0.448	0.629	0.612	0.892	0.620	0.708	0.491	0.887	0.579
BBERT Features of NER Disease Timelines	0.317	0.952	0.916	0.475	0.661	0.649	0.892	0.654	0.739	0.524	0.890	0.613

TABLE V

Influence of Neighbourhood Users in Input Document With $P_n = 3$ and Full Mode Data Accumulation, Averaged Over 4 Individual Runs

Neighbourhood Users	K=1				K=3				K=5			
	Recall	Precision	Cosine Similarity	F1	Recall	Precision	Cosine Similarity	F1	Recall	Precision	Cosine Similarity	F1
$h=0$ (Full Mode)	0.296	0.874	0.909	0.442	0.590	0.584	0.891	0.586	0.681	0.402	0.887	0.505
$h=1$	0.302	0.898	0.911	0.451	0.601	0.597	0.891	0.598	0.712	0.437	0.889	0.541
$h=2$	0.319	0.957	0.916	0.478	0.648	0.636	0.891	0.641	0.730	0.495	0.889	0.589
$h=3$	0.317	0.952	0.916	0.475	0.661	0.649	0.892	0.654	0.739	0.524	0.890	0.613
$h=A$	0.316	0.948	0.916	0.474	0.655	0.640	0.890	0.647	0.733	0.501	0.877	0.595
$h=5$	0.309	0.912	0.908	0.461	0.652	0.638	0.888	0.644	0.725	0.460	0.871	0.562

TABLE VI

Model Comparison With State of the Art Works and Bbert C_I

Model	K= 1				K=3				K=5			
	Recall	Precision	Cosine Similarity	F1	Recall	Precision	Cosine Similarity	F1	Recall	Precision	Cosine Similarity	F1
LDA	0.137	0.419	0.642	0.206	0.320	0.445	0.619	0.372	0.480	0.203	0.606	0.285
AMNN	0.264	0.794	<u>0.847</u>	0.396	0.585	0.529	<u>0.795</u>	0.555	0.627	0.497	0.715	0.554
HashRec	<u>0.278</u>	<u>0.836</u>	0.771	<u>0.417</u>	<u>0.602</u>	<u>0.563</u>	0.736	<u>0.581</u>	<u>0.681</u>	<u>0.503</u>	<u>0.773</u>	<u>0.578</u>
BBERT C_I (Ours)	0.266	0.797	0.834	0.398	0.536	0.499	0.718	0.516	0.593	0.438	0.695	0.503
BBERT T_F (Ours)	0.317	0.952	0.916	0.475	0.661	0.649	0.892	0.654	0.739	0.524	0.890	0.613

TABLE VIIValidation of Models With Facebook “Chronic Pain” Dataset, $K = 5$

Model	$K=5$			
	Recall	Precision	Cosine Similarity	F1
LDA	0.197	0.184	0.208	0.190
AMNN	0.350	0.223	0.346	0.272
HashRec	0.398	0.275	0.341	0.325
BBERT _{T_g}	0.644	0.519	0.726	0.574

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TABLE VIII

Impact of Frequent Tags in Tag Prediction Model With $K = 3$ (Brackets Indicate the Change in Absolute Value)

Model	$K=3$			
	Recall	Precision	Cosine Similarity	F1
LDA	0.320	0.445	0.619	0.372
BBERT _{T_g}	0.661	0.649	0.892	0.654
LDA _{$-2T_g$}	0.257 (-0.063)	0.372 (-0.073)	0.429 (-0.190)	0.303 (-0.069)
BBERT _{$-2T_g$}	0.658 (-0.003)	0.643 (-0.006)	0.878 (-0.014)	0.650 (-0.004)

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TABLE IX

Qualitative Comparison of Tag Prediction

Ground Truth Tbgs for HealthUnlocked Dataset (Future Post): Chemotherapy / Radiotherapy / Tarceva / Afatinib / Cancer and Tumors	
Model	Predicted Tags
LDA	Chemotherapy / Cancer / Lung / Treatment / Medication
HashRec	Chemotherapy / Cough / Cancer / Infection / Medication
AMNN	Chemotherapy / Tumor / Medication / Respiratory / Cough
BBERT _{CI}	Chemotherapy / Infection / Cancer / Tumors / Cough
BBERT _{T_g} (Full Mode) (No-Neighbors)	Chemotherapy / Infection / Tumors / Radiotherapy / Cancer
BBERT _{T_g} (Ours) (3 Neighbors)	Chemotherapy / Radiotherapy / Afatinib / Cancer and Tumors / Drug Trial
Ground Truth Tbgs for Face book Chronic Pain Dataset (Future Post): Fatigue / Lumbar / Disc Tear / Weight Gain / Withdrawal	
Model	Predicted Tags
LDA	Back Pain / Cancer / Side Effects / Headache / Medication
HashRec	Back Pain / Cancer / Side Effects / Headache / Medication
AMNN	Sitting / Tumor / Back Pain / Metastasis / Surgery
BBERT _{T_g} (Full Mode) (No-Neighbors)	Tiredness / Side Effects / Weight Gain / Lower Back / Tylenol

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