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Explainable Automated Coding of Clinical Notes using Hierarchical Label-wise Attention Networks and Label Embedding Initialisation

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Abstract

Background: Diagnostic or procedural coding of clinical notes aims to derive a coded summary of disease-related information about patients. Such coding is usually done manually in hospitals but could potentially be automated to improve the efficiency and accuracy of medical coding. Recent studies on deep learning for automated medical coding achieved promising performances. However, the explainability of these models is usually poor, preventing them to be used confidently in supporting clinical practice. Another limitation is that these models mostly assume independence among labels, ignoring the complex correlations among medical codes which can potentially be exploited to improve the performance.

Methods: To address the issues of model explainability and label correlations, we propose a Hierarchical Label-wise Attention Network (HLAN), which aimed to interpret the model by quantifying importance (as attention weights) of words and sentences related to each of the labels. Secondly, we propose to enhance the major deep learning models with a label embedding (LE) initialisation approach, which learns a dense, continuous vector representation and then injects the representation into the final layers and the label-wise attention layers in the models. We evaluated the methods using three settings on the MIMIC-III discharge summaries: full codes, top-50 codes, and the UK NHS (National Health Service) COVID-19 (Coronavirus disease 2019) shielding codes. Experiments were conducted to compare the HLAN model and label embedding initialisation to the state-of-the-art neural network based methods, including variants of Convolutional Neural Networks (CNNs) and Recurrent Neural Networks (RNNs).

Results: HLAN achieved the best Micro-level AUC and F_1 on the top-50 code prediction, 91.9% and 64.1%, respectively; and comparable results on the NHS COVID-19 shielding code prediction to other models: around 97% Micro-level AUC. More importantly, in the analysis of model explanations, by highlighting the most salient words and sentences for each label, HLAN showed more meaningful and comprehensive model interpretation compared to the CNN-based models and its downgraded baselines, HAN and HA-GRU. Label embedding (LE) initialisation significantly boosted the previous state-of-the-art model, CNN with attention mechanisms, on the full code prediction to 52.5% Micro-level F_1 . The analysis of the layers initialised with label embeddings further explains the effect of this initialisation approach. The source code of the implementation and the results are openly available at https://github.com/acadTags/Explainable-Automated-Medical-Coding.

Conclusion: We draw the conclusion from the evaluation results and analyses. First, with hierarchical label-wise attention mechanisms, HLAN can provide better or comparable results for automated coding to the state-of-the-art, CNN-based models. Second, HLAN can provide more comprehensive explanations for each label by highlighting key words and sentences in the discharge summaries, compared to the *n*-grams in the CNN-based models and the downgraded baselines, HAN and HA-GRU. Third, the performance of deep learning based multi-label classification for automated coding can be consistently boosted by initialising label embeddings that captures the correlations among labels. We further discuss the advantages and drawbacks of the overall method regarding its potential to be deployed to a hospital and suggest areas for future studies.

Keywords: Automated medical coding, Deep learning, Attention Mechanisms, Explainability, Natural Language Processing, Multi-label classification, Label correlation

1. Introduction

Diagnostic or procedural coding of medical free-text documents (e.g. discharge summaries) aims to derive a coded summary of disease-related information about patients, for clinical care, audit, and research. In hospitals, such coding is usually done manually, requiring much cognitive human effort, but could potentially be automated. An automated program could efficiently take a clinical note as input and then output medical codes from existing classification systems, e.g. ICD (International Classification of Diseases). This could facilitate coding professionals provide more accurate results.

This clinical task is technically challenging, due to (i) the explainability required to process long documents, in average about 2000 tokens in a discharge summary in MIMIC-III [1],

- and thus pose a "needle-in-a-haystack" issue to locate the key words and sentences relevant to each code; (ii) the complex label correlations in the multi-label setting, in average about 16 different ICD-9 (the Ninth Revision) codes per discharge summary in the MIMIC-III dataset [2], which inherently ex-
- hibit the complex relations among codes; and (iii) a large set ⁶⁵ of codes when using all the codes as candidates for prediction, e.g. around 13k unique codes in ICD-9 and many times further in ICD-10 [3] and ICD-11 [4].

Automated medical coding has been studied for more than

- ²⁵ a decade. Early studies mostly use systems based on rules, ⁷⁰ grammar, and string matching, as reviewed in [5]. Recent studies adapt deep learning based document classification methods, which commonly formalise the task as a *multi-label classifica-tion* problem [6, 7, 1]. Typically, they use variations of Recur-
- ³⁰ rent Neural Networks (RNNs) and Convolutional Neural Net-⁷⁵ works (CNNs) to derive a continuous representation of clinical notes matched to the high-dimensional coding space. However, few studies have tackled the above challenges above regarding explainability and label correlations.
- Explainability (or interpretability, used interchangeably in ⁸⁰ this paper) is a key requirement for models applied to the clinical domain, particularly regarding the ethical aspect and to build medical professionals' trust in machine learning models [8, 9]. Also, to facilitate the work of coding professionals, a
- ⁴⁰ desired automated coding system should be able to highlight 85 the most essential part of a long clinical note to support the assignment of medical codes. To address this, models based on CNNs can be adapted to highlight *n*-gram information to support the explanation, as in [7]. Solely the *n*-grams, however,
- ⁴⁵ may not be enough to provide accurate interpretation reflecting ⁹⁰

the document structure.

In this work, we propose to highlight the most essential words and sentences in a document for automated medical coding, inspired and adapted from Hierarchical Attention Networks (HAN) [10] and the recent model, Hierarchical Attention bidirectional Gated Re-current Units (HA-GRU) [1]. With attention mechanisms, HAN can highlight the salient words and sentences related to the overall prediction. However, HAN could not generate a specific interpretation for each label. HA-GRU [1] can provide a *sentence*-level explanation for each label, but still could not specify the most essential *words* leading to the decision of each code. We present a novel model, Hierarchical Label-wise Attention Network (HLAN), which has label-wise word-level and sentence-level attention mechanisms, so as to provide a richer explainability of the model.

We formally evaluated HLAN along with HAN, HA-GRU, and CNN-based neural network approaches for automated medical coding. With better or comparative coding performance in various data settings, HLAN can further generate more comprehensive explanations through key sentences and words for each label, as indicated from the analysis on model explainability. The analysis of the false positive predictions also shows that the explanation based on the hierarchical label-wise attention mechanisms in HLAN can serve as a reference for medical professionals and engineers to make reasonable coding decisions and system iterations even when the model seems to predict erroneously.

Apart from model interpretability, another issue not thoroughly studied in deep learning based multi-label classification is label correlation. Medical codes are related and can be predicted together, for example, the code 486 (ICD 9 for Pneumonia) commonly appeared for over 1.5k times (out of about 53k documents) with the code 518.81 (Acute respiratory failure) in the MIMIC-III dataset. Such co-occurrences are under-lied by the clinical, biomedical, and biological associations among different diseases. Deep learning for multi-label classification represents the label space with orthogonal vectors: each label as a one-hot vector and each label set as a multi-hot representation [11, 7]. This, however, assumes independence among labels.

We propose an effective label embedding initialisation approach to tackle the label correlation problem. We encode the label correlation using pre-trained label embeddings from the label sets in the training data, derived from the coding practice. Then the label embeddings are used to initialise the weights in the final hidden layer and label-wise attention layers. The idea is that the linear projection can automatically leverage the label similarity encoded in the continuous label embedding space. This approach shows consistent and significant improvement, while not requiring hyper-parameter tuning or further computa-135 tional complexity.

We evaluate our approach with three specific datasets based on the openly available, MIMIC-III database [2], containing clinical notes in the critical care sector in the US. The first two datasets, full code and top-50 code predictions, are the same as

in the work [7] for comparison. The third dataset was created to₁₄₀ simulate the task of identifying high-risk patients for shielding during the COVID-19 (Coronavirus disease 2019) pandemic by predicting the ICD-9 codes matched to the codes used in the UK NHS (National Health Service) patient shielding identification
 method¹.

Thus, the contribution of the paper includes:

• A novel, Hierarchical Label-wise Attention Network (HLAN) for automated medical coding. The proposed HLAN model provides an explanation in the form of attention weights on both the word level and the sentence level for the prediction of each medical code.

 An effective label embedding (LE) initialisation approach to enhance the performance of various deep learning models for multi-label classification. Analysis of the LE initialised layers shows the efficacy to leverage label correlations for medical coding.

A formal comparison of the main deep learning based methods for automated coding. Experiments on three datasets based on the MIMIC-III discharge summaries, i.e. full code prediction, top-50 code prediction, and the 160 NHS COVID-19 shielding-related code prediction, show the advantage of the proposed method over the state-of-the-art methods (CNNs, Bi-GRU) and downgraded baselines (HA-GRU, HAN). Label embedding initialisation significantly improved the performance of neural network 165 models in most evaluation settings. An analysis and comparison of the model interpretability demonstrate the most comprehensive explanations from the HLAN model.

The rest of the paper is organised as follows. First, we review
 the related work on automated medical coding with explainabil-¹⁷⁰
 ity, deep learning methods for multi-label classification, and

label correlation in Section 2. Then, we present the problem formulation, followed by the proposed model, HLAN, and the idea of LE initialisation in Section 3. The experiments, including datasets, experimental settings, main and per-label results, analysis and comparison of model explainability, and analysis on the layers initialised with LE, are in Section 4. We finally discuss the advantages and drawbacks of the overall methods in Section 5 and summarise the work in Section 6.

2. Related Work

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We will first present the task of automated medical coding with the methods used especially in most recent studies, then introduce in detail the mainstream breakthrough on deep learning-based multi-label classification for the task, and finally review the label correlation issue, particularly relevant to the medical and clinical domain.

2.1. Automated Medical Coding with Explainability

Automated medical coding is the task of transforming medical records, especially the natural language in the clinical notes, into a set of structured, medical codes to facilitate clinical care, audit, and research [5]. The applied alphanumerical codes in the clinical domain, such as ICD and SNOMED-CT, represent patients' diagnosis, procedures and other information with controlled clinical terminology.

One of the earliest reviews back in 2010 [5] surveyed 113 studies on coding or classification of clinical notes. Most of the studies applied tools with rule-based, grammar-based, and string matching methods, and they in overall suffered the challenges of reasoning and the lack of method generalisability. The field of automated medical coding has in more recent years been advanced with the open, benchmarking datasets like radiology reports in [12] and MIMIC-III [2] discharge summaries. With the datasets, deep learning based approaches have been proposed and tested, which have generally demonstrated better performance than traditional machine learning methods. The work in [6] compared the deep learning based method, CNN, with several traditional machine learning methods, support vector machine, random forests, and logistic regression, for ICD-9 code prediction (number of ICD-9 codes |Y|=38) from 978 radiology reports in [12]. The result showed comparable or improved results of the deep learning approach to the traditional methods, even without parameter tuning in the CNN model. The work in [7] adapted CNN with attention mechanisms and established a state-of-the-art performance in predict-

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¹https://digital.nhs.uk/coronavirus/ shielded-patient-list/methodology

ing the full set (|Y|=8,921) and the top-50 most frequent ICD-9₂₂₀ codes (|Y|=50) from MIMIC-III discharge summaries.

A key aspect of clinical applications is their requirement of the *explainability* of models. Users are entitled to a "right of explanation" when their data being used for AI algorithms, as potentially regulated by the General Data Protection Regulation²²⁵

(GDPR) [9]. For clinical applications, e.g. radiology, the Joint European and North American Multisociety Statement raises great ethical concern on AI algorithms regarding explainability, i.e. "the ability to explain what happened when the model made a decision, in terms that a person understands" [8, p. 438].290

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- While deep learning achieves better results in general, the approach is inherently less transparent than traditional methods due to its extremely complex networks of non-linear activation.
- Few studies explored the explainability of deep learning
 models for automated medical coding. A representative work²³⁵ is the study [7], which compared the ability of different models to highlight *n*-grams along with the models' ICD-9 code prediction. A manual evaluation showed that the CNN model with attention mechanisms can generate more meaningful *n*-grams
 relevant to the labels [7]. The study [1] proposed a Hierarchi-
- relevant to the labels [7]. The study [1] proposed a Hierarchical Attention bi-directional Gated Recurrent Unit (HA-GRU)²⁴⁰ to produce a sentence-level explanation for each code, instead of *n*-gram-level explanation. In this work, we propose an approach with enhanced interpretability, from both the label-wise
 word-level and the sentence-level attention weights, to support
- automated coding.

2.2. Deep Learning-based Multi-label Classification with Attention Mechanisms

Automated medical coding is mainly formulated as a multilabel classification problem [13, 14, 7, 1], where each object²⁵⁰ (e.g. clinical note) is associated with a set of labels (e.g. diagnosis or procedure ICD codes) instead of a single label in binary or multi-class classification.

Deep learning has become the main approach for multi-label document classification [11, 15] in recent years. The advantage²⁵⁵ of multi-label deep learning models lies in their straightforward problem formulation and strong approximation power on large datasets, resulting in better performance over traditional machine learning approaches, as compared in [16, 15, 6]. For

automated coding, some of the notable neural network mod-260
els adapted for multi-label classification are variations of CNNs
[6, 7] and RNNs [1] with attention mechanisms. Pre-trained models with multi-head self-attention blocks (e.g. BERT, Bi-directional Encoder Representations from Transformers) [17],

while substantially improved many NLP tasks, so far still are under-performing for automated coding with the MIMIC-III discharge summaries [18, 19].

The idea of the above mentioned *attention mechanism* is a key, recent advancement in deep learning for NLP, originated from machine translation to align (or attend to) words in the source sentence in one language to predict each of the target words in another language [20]. This inspires to jointly learn to represent the important words and sentences while classifying a document in HAN [10], thus also enables to *explain* the inner working of deep learning models. HAN was adapted to a multi-label classification setting to classify socially shared texts in [15] and for automated medical coding [1]. Founded on the studies above, our approach provides a richer label-wise attention mechanism at both the word and the sentence level for automated medical coding.

2.3. Label Correlation

In multi-label classification, labels are potentially correlated to each other. As the example in Section 1, the medical codes of "Pneumonia" and "Acute Respiratory Failure" tend to appear together in the MIMIC-III discharge summaries. In automated medical coding, the number of unique code |Y| is large (|Y| = 8921 in the MIMIC-III dataset) and further the possible label relations (e.g. the number of pairwise combinations is near to $|Y|^2$). Such correlations among the labels represent additional knowledge that could be exploited to improve performance [21].

This issue of *label correlation* (or "label dependence") remains an ongoing challenge [21] in multi-label classification, especially with deep learning models. Deep learning for multilabel classification mostly represents the label space with orthogonal vectors: each label as a one-hot vector and each label set as a multi-hot representation, in general domains [11] and clinical domains [7, 1]. Combined with the sigmoid activation and binary cross-entropy loss, this overall approach, effectively, assumes independence among labels.

One recent approach to address the problem is through weight initialisation [22, 23]: initialising higher weights for dedicated neurons (each encoding a co-occurrence relation among labels) in the final hidden layer. The approach showed performance improvement, however, it is not computationally efficient to assign each neuron in the final hidden layer to represent one of the massive (even the pairwise) patterns of label relations. An alternative method is through regularisation in [15] to enforce the output layer of the neural network to ²⁸⁵ satisfy constraints on label relations. This requires to further tune the hyper-parameters of the regularisers so that a relatively marginal improvement (0.5-1.5% example-based F_1 on scientific paper abstracts and questions in social Q&A platforms) could be achieved. In this study, we further propose a novel effective weight initialisation approach to tackle the label correlation problem, by initialising pre-trained dense label embeddings instead of the sparse co-occurrence representations.

3. Proposed Method

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We formalise automated medical coding from clinical notes
as a multi-label text classification problem [11]. With deep learning, multi-label classification mainly contains two, inte-300 grated parts, (i) a neural document encoder, representing documents into a continuous representation, and (ii) a prediction layer, matching the document space to the label space. We
present the problem formalisation and the deep learning based multi-label classification in Section 3.1. Then, regarding the neural document encoder, we propose the hierarchical label-305 wise attention network in Section 3.2, followed by the idea of label embedding initialisation in the prediction layer in Section 3.3.

3.1. Problem Formulation with Deep Learning Models

Formally multi-label classification can be defined as follows. Suppose X denoting the collection of textual sequences (e.g. clinical notes), and $Y = \{y_1, y_2, ..., y_{|Y|}\}$ denotes the full set of labels (i.e. ICD codes) of size |Y|. Each instance $x_d \in X$ is a word sequence of a document, where *d* is the document index.³¹⁵ Each $x_d \in X$ is associated with a label set $Y_d \subseteq Y$. Each label set Y_d can be represented as a |Y|-dimensional *multi-hot* vector, $\overrightarrow{Y_d} = [y_{d1}, y_{d2}, ..., y_{d|Y|}]$ and $y_{dl} \in \{0, 1\}$, where a value of 1 indicates that the *l*th label y_l has been used to annotate (is relevant to) the *d*th instance, and 0 indicates irrelevance. The task is to³²⁰ learn a complex function $f : X \to Y$ based on a training set $D = \{x_d, \overrightarrow{Y_d} | d \in [1, m]\}$, where *m* is the number of instances in the training set.

Neural document encoders in deep learning models (e.g. CNN, RNN, and BERT, as review in Section 2.2) represent each word sequence *x* as a continuous vector *h*, with matrix projection and non-linear activation. The representation *h* is projected to the label space and turned into $p_{dl} \in (0, 1)$ with the sigmoid function ($\sigma(x) = \frac{1}{1+e^x}$), as defined in Equation 1 below, where the weight w_l (a row vector in *W*) and the bias *b* are parameters to be learned during the training process. The obtained p_{dl} is

the probability of the label (e.g. ICD code) y_l being related to the document (e.g. discharge summary) d.

$$p_{dl} = \sigma(w_l h + b)$$
, or collectively as $p_d = \sigma(W h + b)$ (1)

The loss function is commonly the binary cross-entropy loss [11] as defined in Equation 2, which measures the sum of negative log-likelihood of the predictions p_{dl} of the actual labels. A large deviation between $\overrightarrow{Y_{dl}}$ and p_{dl} will cause a greater value in the L_{CE} and thus will be penalised during training.

$$L_{CE} = -\sum_{d} \sum_{l} (\overrightarrow{Y_{dl}} \log(p_{dl}) + (1 - \overrightarrow{Y_{dl}}) \log(1 - p_{dl}))$$
(2)

For inference, a calibration threshold *Th* (default as 0.5) is set to assign the label to the document when $p_{dl} > Th$.

3.2. Hierarchical Label-wise Attention Network

Following the framework above, the neural document encoder in HLAN (as illustrated in Figure 1) takes into input the word sequence $x_d = \{x_{d1}, x_{d2}, ..., x_{dn}\}$, where x_{di} denotes the sequence of tokens in the *i*th of all *n* sentences, and output the document representation. The distinction to HAN [10] is that HLAN represents the same document differently at both the word-level and the sentence-level regarding different labels. HLAN extends the contextual vectors in HAN to the label-wise contextual matrices, V_w and V_s . The document representation also becomes a matrix, C_d , where each row (corresponding to each label) has the same dimensionality as *h*.

As shown in Figure 1, the model consists of an embedding layer, hidden layers (hierarchical label-wise attention layers), and a prediction (or projection) layer. First, the embedding layer transforms the one-hot input representation u_{di} of each token in the sequence of the *i*th sentence x_{di} into a lowdimensional continuous vector, $e_{di} = W_e u_{di}$, where we used the neural word embedding algorithm, Word2vec [24], to pre-train W_e for its efficiency.

Second, we applied the Gated Recurrent Unit (GRU) [25], a type of RNN unit, to capture long-term dependencies in the clinical narrative. An RNN unit "reads" each token in the sequence one by one, every time producing a new hidden state $h^{(t)}$, corresponding to the token at time *t*. Different from the vanilla RNN unit, GRU additionally considers the previous tokens by using a reset gate $r^{(t)}$ and an update gate $z^{(t)}$. This allows to model the dependencies among tokens in long sequences. A GRU can be formally defined as in the Equations 3 below, where \vec{h} denotes the hidden states through forward processing,

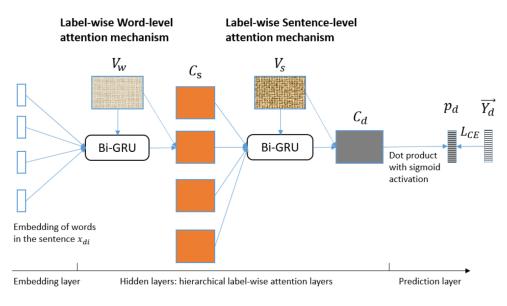


Figure 1: Hierarchical Label-wise Attention Network (HLAN)

 σ is a non-linear activation function (e.g. sigmoid function), $W_{hr}, W_{hz}, W_{h\tilde{h}} \in \mathbb{R}^{d_h \times d_h}$ are weights, and $b_r, b_z \in \mathbb{R}^{d_h}$ represent bias terms. A bi-directional adaptation was applied by concatenating the hidden states at each time after read the sequence both forwardly (\rightarrow) and backwardly (\leftarrow) to form a more comprehensive representation, $h^{(t)} = [\vec{h}^{(t)}; \vec{h}^{(t)}] \in \mathbb{R}^{2d_h}$. This subarchitecture is generally known as Bi-GRU [25].

$$r^{(t)} = \sigma(W_{er}e^{(t)} + W_{hr}\vec{h}^{(t-1)} + b_r)$$

$$z^{(t)} = \sigma(W_{ez}e^{(t)} + W_{hz}\vec{h}^{(t-1)} + b_z)$$

$$\tilde{h}^{(t)} = \tanh(W_{e\tilde{h}}e^{(t)} + W_{h\tilde{h}}(r^{(t)} \circ \vec{h}^{(t-1)}))$$

$$\vec{h}^{(t)} = (1 - z^{(t)}) \circ \vec{h}^{(t-1)} + z^{(t)} \circ \tilde{h}^{(t)}$$
(3)

For simplicity, we use the function $h = \text{Bi-GRU}(e, \Theta)$ to denote the whole process (with bi-directional concatenation of hidden states) above. Instead of applying one single Bi-GRU layer to represent the whole document, we applied a word-level Bi-GRU to represent each sentence and then a sentence-level one to represent the whole document, as illustrated in Figure 1. This captures the hierarchical structure of the document and relieves the burden of having a too lengthy sequence for each GRU [1] (e.g. from the original sequence length 2500 in the MIMIC-III discharge summaries to only 100 on the word level and 25 on the sentence level).

A common way is to represent the whole sequence as the concatenated hidden state $h^{(t)}$ of the last time *t*. This representation tends to emphasise the ending elements (i.e. words or sentences) and does not discriminate between the elements in a sequence. In fact, the key information for medical cod-345

ing is contained in a well selected part of the lengthy discharge summary. We therefore use an attention mechanism to learn a weighted average of the hidden states to form a final representation as in [10, 20]. The attention scores are based on an alignment (or a similarity computation) of each hidden representation in a sequence to a context vector. The context vector is usually *shared* for all labels as in [10, 15], whereas in medical coding, it is essential to interpret the amount of attention paid regarding a *specific* medical code to the clinical note.

$$h^{(i)} = \text{Bi-GRU}(e, \Theta_w)$$

$$v^{(i)} = \tanh(W_w h^{(i)} + b_w)$$

$$\alpha^{(i)}_{wl} = \frac{\exp(V_{wl} \bullet v^{(i)})}{\sum_{o \in [1, n_t]} \exp(V_{wl} \bullet v^{(o)})}$$

$$C_{sl} = \sum_{i \in [1, n_t]} \alpha^{(i)}_{wl} h^{(i)}$$
(4)

Thus, the adapted, *label-wise word-level attention mechanism* is defined in Equations 4 above. The context matrix for the word-level attention mechanism is denoted as $V_w \in \mathbb{R}^{|Y| \times d_w}$, where each row V_{wl} (of attention layer size d_w) is the context vector corresponding to the label y_l . The attention score $\alpha_{wl}^{(i)}$ for the label y_l is calculated as a softmax function of the dot product similarity between the vector representation $v^{(i)}$ (transformed from the *i*th hidden state $h^{(i)}$ with a feed-forward layer) and the context vector V_{wl} for the same label. n_t denotes the number of tokens in a sentence. The sentence representation C_{sl} , as a row vector in $C_s \in \mathbb{R}^{|Y| \times 2d_h}$, for the label y_l , is computed as the weighted average of all the hidden state vectors $h^{(i)}$.

In a similar way, we can compute the *label-wise sentencelevel attention mechanism* as defined in Equations 5, which encodes each row C_{sl} in the sentence representations C_s to a label-wise sentence representation $S_l^{(r)}$, to be non-linearly transformed to $U_l^{(r)}$ and aligned to the corresponding row V_{sl} in₃₇₀ sentence-level contextual matrix $V_s \in \mathbb{R}^{|Y| \times d_s}$, and outputs the sentence-level attention scores α_{sl} (for a label y_l) and the document representation matrix $C_d \in \mathbb{R}^{|Y| \times 4d_h}$. To note that the dimensionality of S_l^r and thus C_{dl} are further doubled to $4d_h$ through the Bi-GRU process.

$$S_{l}^{(r)} = \text{Bi-GRU}(C_{sl}, \Theta_{S})$$

$$U_{l}^{(r)} = \tanh(W_{S}S_{l}^{(r)} + b_{S})$$

$$\alpha_{sl}^{(r)} = \frac{\exp(V_{sl} \bullet U_{l}^{(r)})}{\sum_{q \in [1,n]} \exp(V_{sl} \bullet U_{l}^{(q)})}$$

$$C_{dl} = \sum_{r \in [1,n]} \alpha_{sl}^{(r)}S_{l}^{(r)}$$
(5)

Then, we use a label-wise, dot product projection with logistic sigmoid activation to model the probability of each label to each document, as defined in Equation 6, adapted from Equa-³⁸ tion 1. The parameters in w_l are row vectors in the projection matrix W.

$$p_{dl} = \sigma(w_l C_{dl} + b_l) \tag{6}$$

We finally optimise the binary cross-entropy loss function₃₉₀ in Equation 2 with L_2 regularisation using the Adam optimiser [26].

3.3. Label Embedding Initialisation

For automated medical coding, the diagnostic and proce-³⁹⁵ dural codes (or labels) have complex semantic relations, and can potentially be leveraged to improve prediction. Clinically, these code relations represent the correlation among diseases and medical procedures from the medical coding practice.

As we reviewed in Section 2.3, previous studies on weight⁴⁰⁰ initialisation to address the label correlation issue mostly focus on a co-occurrence based representation of labels [22, 23]. Both studies dedicate a neuron in the final hidden layer to initialise one single co-occurrence pattern. There are, however, very lim-

ited neurons to be assigned to initialise the massive number of₄₀₅ label relations, especially for the large label size in automated coding.

Instead of encoding the sparse co-occurrence patterns of labels, we learn low-dimensional, dense, label embeddings. For two correlated labels y_i and y_k , e.g. 486 (Pneumonia) and₄₁₀ 518.81 (Acute respiratory failure), one would expect that the prediction of one label has an impact on the other label for some clinical notes, i.e. p_{dj} is correlated or has a similar value to p_{dk} . To achieve this, according to Equations 1 or 6, we propose to initialise their corresponding weights w_j and w_k (corresponding to the labels y_j and y_k) in W with a label representation E which reflects the actual label correlation (e.g. similarity between y_j and y_k) in a continuous space.

A straightforward idea is thus to initialise the projection matrix W using E as pre-trained label embeddings, e.g. with a neural word embedding algorithm, learned from the label sets in the training data, $\{\overrightarrow{Y_d}|d \in [1,m]\}$. For initialisation, we pre-train the label embeddings E with dimensionality the same as W. We used the Continuous Bag of Words algorithm in word2vec [24] for its efficiency and its power to represent the correlations of the labels. Figure 2 shows an intuitive visualisation, for which we used an unsupervised technique, T-SNE (t-distributed Stochastic Neighbor Embedding), to reduce the dimensionality of the learned label embeddings while preserving the local similarity and structure of the labels [27]. It can be observed that the ICD-9 code learned from the MIMIC-III training label sets can capture the semantic relations that are distinct from the ICD-9 hierarchy. For example, 486 (Pneumonia) and 518.81 (Acute respiratory failure) appear closely on the bottom while they are not under the same parent in the ICD-9 hierarchy.

Besides, the label embedding initialisation can also be applied to the context matrices V_w and V_s (see Figure 1) in the label-wise attention mechanisms. Taking the word-level attention mechanisms in Equation 4 as an example, we can initialise V_{wl} with the pre-trained label embedding E_l for the label y_l . This imposes a tendency for context vectors of the correlated labels to align the Bi-GRU encoded token representation v_i in a geometrically similar way. Similarly, we can also initialise the label-wise attention layer in CNN+att [7] and in HA-GRU [1]. While the initialised layers are dynamically updated during the training, the tendency that imposed by label embeddings remains for most neural networks; we will empirically demonstrate this in the analysis of initialised layers in Section 4.8.

For automated coding, the approach can be extended by initialising label embeddings with the clinical ontologies and description texts of ICD codes. However, due to the different nature of the knowledge (i.e. embedded label relations), the external sources may bring contradictory label correlations to the ones in the dataset, as also discussed in [28]. In this research, we focus on leveraging label relations from the label sets alone

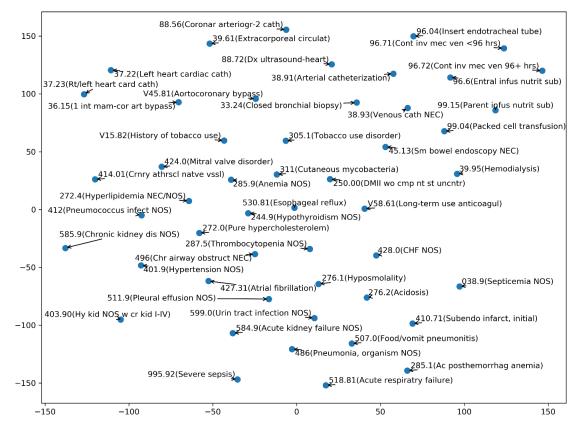


Figure 2: The 2-dimensional T-SNE plot of word2vec Continuous Bag of Words label embeddings of the 50 ICD-9 codes in MIMIC-III-50, trained on the whole training label sets, $|\vec{Y_d}| d \in [1, m]$, in MIMIC-III.

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as in [22, 23], as it directly reflects the label correlation of the coding practice that generated the dataset, and leave the integration of external knowledge for a future study.

4. Experiments

⁴¹⁵ We tested HLAN and several strong baseline models, with label embedding initialisation, on three data sets based on the MIMIC-III database. The main results show the comparative results of HLAN to other state-of-the-art models on the datasets and the consistent improvement with label embedding initiali-⁴⁴⁰

- sation. More importantly, through an analysis on model interpretability, we also show that HLAN can provide a more comprehensive explanation using the label-wise word and sentencelevel attention mechanisms. Analysis of the layers initialised with label embeddings further reveals the effect of the ini-445
- tialisation approach. The source code of our implementation and the results are openly available at https://github.com/ acadTags/Explainable-Automated-Medical-Coding.

4.1. Datasets

We used the benchmark dataset, MIMIC-III ("Medical Information Mart for Intensive Care") [2], which contains clinical data from adult patients admitted to the critical care unit in the Beth Israel Deaconess Medical Center in Boston, Massachusetts between 2001 and 2012, to validate our approach. The ICD-9 codes annotated by professionals in the dataset were used as labels. We focused on discharge summaries and followed the preprocessing and data split from [7]. The preprocessed full MIMIC-III dataset has 8,922 unique codes as labels assigned to 52,724 discharge summaries, where 47,724 of them (from 36,998 patients) were used for training, 1,632 for validation, and 3,372 for testing. We also used the same top-50 setting (termed as "MIMIC-III-50") from [7], which narrows down the labels to the top 50 by their frequencies (codes and their frequencies are available in Table S1 in the supplementary material). This has 8,067 discharge summaries for training, 1,574 for validation, and 1,730 for testing.

We further created a subset of discharge summaries annotated using the COVID-19 shielding related ICD codes. This simulates the application of identifying key patients for shielding during the pandemic. We used the ICD-9 codes matched

- to the ICD-10 codes selected by the NHS to identify patients485 with medium or high risks during COVID-19. The considered patients were related to solid organ transplant recipients, people with specific cancers, with severe respiratory conditions, with rare diseases and inborn errors of metabolism, on immunosup-
- ⁴⁵⁵ pression therapies, or who were pregnant with significant congenital heart disease², which is still in active use and under maintenance at the time of writing this paper. While the actual⁴⁹⁰ EHR data and the shielded patient list from the NHS are not easy to obtain, the ICD-10 codes are openly available for re-
- ⁴⁶⁰ use³. We thus used MIMIC-III to simulate the task of identifying patients for shielding during COVID-19. We selected those appeared at least 50 times in the MIMIC-III dataset, resulting in 20 ICD-9 codes (out of 79 matched codes), available in Table⁴⁹⁵ S2 in the supplementary material. After filtering the MIMIC-III
- dataset with the selected ICD-9 codes, there are 4,574 discharge summaries for training, 153 for validation, and 322 for testing.
 We name this dataset as "MIMIC-III-shielding".

Statistics of the three datasets are in Table 1. Denoted by Ave,⁵⁰⁰ the average number of labels per document (or label cardinality)

- ⁴⁷⁰ in the training set of MIMIC-III, MIMIC-III-50, and MIMIC-III-shielding are 15.88, 5.69, and 1.08, respectively. While all originated from MIMIC-III database, the three datasets represent different case scenarios in automated medical coding with₅₀₅ various scales of data and vocabulary size ("vocab"), number
- ⁴⁷⁵ of labels to predict, and the average number of labels per document. While the full MIMIC-III dataset has much more training instances, it is more complex as its number of labels |*Y*| and vocabularies are significantly greater than MIMIC-III-50 and₅₁₀ MIMIC-III-shielding.

Table 1: Statistics of the datasets

Dataset	Vocab	Train	Valid	Test	Y	Ave	
MIMIC-III-50	59,168	8,067	1,574	1,730	50	5.69	-
MIMIC-III-shielding	47,979	4,574	153	322	20	1.08	
MIMIC-III	140,795	47,724	1,632	3,372	8,921	15.88	515

Figures of ICD-9 code distributions by frequency in the three datasets are available in Figure S1 in the supplementary material, along with the list of the selected codes (and their frequencies) in the MIMIC-III and MIMIC-III-shielding datasets. The⁵²⁰ statistics show a high imbalanced characteristics of the labels in all three data settings. Most label occurrences are from a few labels and there is a long-tail of labels having very low frequencies. This is most pronounced in the full label setting ("MIMIC-III") and also presented in the other two datasets.

4.2. Experiment Settings

We implemented the proposed Hierarchical Label-wise Attention Network (HLAN) model and the other baselines for comparison:

- 1. CNN, Convolutional Neural Network, which is essentially based on [29] for text classification, and applied in [6, 30] for automated medical coding.
- CNN+att (or CAML), CNN with a label-wise attention mechanism, proposed in [7].
- 3. Bi-GRU, Bi-directional Gated Recurrent Unit [25] for multi-label classification. The document representation is set as the last concatenated hidden state $h^{(t)}$.
- HAN, Hierarchical Attention Network [10], which can be considered as a downgraded model of HLAN when the attention mechanisms are shared for all labels (see Figure 1, when V_w, V_s, and C_s, C_d become vectors, same for all labels).
- 5. HA-GRU, Hierarchical Attention bi-directional Gated Recurrent Unit, proposed in [1], which can be considered as a downgraded model of HLAN when the word-level attention mechanism is shared for all labels (see Figure 1, when V_w and C_s become vectors, same for all labels, while V_s and C_d are the same as in HLAN).

We applied the label embedding initialisation approach (denoted as "+LE") to all the models above. We pre-trained the label embeddings E from the label sets in the training data with the word2vec (Continuous Bag of Words with negative sampling) algorithm [24]. The label embeddings have the dimension same as the final hidden layer or the label-wise attention layer(s) in each neural network model. We applied the Python Gensim package [31] to train embeddings, by setting the window size as 5 and minimum frequency threshold ("min_count") as 0. Label embeddings were normalised to unit length for initialisation. Xavier initialisation [32] was used for labels not existing in the training data for faster model convergence. We used the same setting to train and initialise the 100-dimension word embeddings W_e from the documents.

²A clearer description of the "high risk" category is in https: //digital.nhs.uk/coronavirus/shielded-patient-list/ methodology/background

³To see the annexe B in https://digital.nhs.uk/coronavirus/ shielded-patient-list/methodology/annexes 525

The implementations of HLAN and HA-GRU were adapted from our previous implementation⁴ of HAN in [15] using the565 Python Tensorflow [33] framework, originated from brightmart's implementation⁵, all under the MIT license. We adapted

- HA-GRU with the sigmoid activation and binary cross-entropy as described in Section 3.1, instead of the softmax activation used in the original paper [1], for a controlled comparison with 570 other models. For CNN, CNN+att, and Bi-GRU, we adapted the implementation⁶ from [7] using the PyTorch framework
- [34] with the same parameters for MIMIC-III and MIMIC-50 535 from [7]. For MIMIC-III-shielding, we used the same hyperparameters as in MIMIC-50. We did not get the results with HA-GRU and HLAN for the MIMIC-III dataset, due to the memory limit caused by the large label size (|Y| = 8,921), while for MIMIC-III-50 and MIMIC-III-shielding, we obtained the 540 results of all models.

The input token length for the models was padded to 2,500 as in [7]. We optimised the precision@k or micro- F_1 metrics (defined in Section 4.3) during the training⁷, according to the implementation in [7]. The batch size for CNN, CNN+att, Bi-GRU were set as 16 as in [7], for HLAN and HA-GRU as 32, and HAN as 128. For HLAN, HAN, and HA-GRU, we tried both a customised rule-based parsing of real sentences with Spacy⁸ and using text chunks of fix length as "sentences"; for

- both ways, we set the sentence length as 25 and padded the 550 number of sentences to 100. The dimensions of the final document representation were 512, 500, 50, 400 for Bi-GRU, CNN, CNN+att, and HLAN (also HAN and HA-GRU), respectively. All models were trained using a single GeForce GTX TITAN
- X server, and the trained HLAN, HA-GRU, and HAN mod-555 els were further tested using a CPU server (4-core, Intel(R) Xeon(R) Platinum 8259CL CPU @ 2.50GHz). The detailed hyper-parameter settings, containing learning rate, dropout rate, and CNN specific parameters (kernel size and filter size), with the estimated training and testing times, are in Table S3 in the supplementary material. 580

We also experimented with BERT as the neural document encoder. Due to the GPU memory limit, we tested the nor-

mal size of a BERT model, i.e. BioBERT-base [35], which had been further pre-trained with PubMed paper abstracts⁹ and full texts¹⁰; we used a sliding window approach to address the token limit issue (512 tokens) in BERT. Our results from the BioBERT-base model were similar to the results in [19], significantly worse than HLAN and CNN¹¹. We believe further adaptations are necessary for BERT models on automated medical coding and leave the direction for a future study.

4.3. Evaluation Metrics

For comparison, we applied the same set of label-based metrics as in [7] and according to the evaluation of multi-label classification algorithms [14, 13]. The chosen metrics include micro- and macro-averaging precision (P), recall (R), F_1 score (F_1) , area under the receiver operating characteristic curve (AUC), an the precision @k.

The micro-averaging metrics treat each document-label as a separate prediction, whereas the macro-averaging metrics are an average of the per-label results. Micro- and macro-averaging applies to all the binary evaluation metrics including precision, recall, AUC. For example, the micro- and macro-averaged precision is defined in Equation 7 below. Recall is calculated in a similar way, but divided by all the true cases $(TP_1 + FN_1)$, and F_1 is then the harmonic mean of the calculated precision and recall, i.e. $F_1 = \frac{2 \times P \times R}{P + R}$. The AUC is defined by two metrics, the true positive rate (or recall) on the Y axis and false positive rate on the X axis, depicting the tradeoff between the two metrics when varying the calibration threshold Th [36]. The overall performance of a classifier (with a set of varied Th) can thus be reflected by AUC.

$$\text{Micro-P} = \frac{\sum_{l=1}^{L} TP_l}{\sum_{l=1}^{L} TP_l + FP_l} \qquad \text{Macro-P} = \frac{1}{|L|} \sum_{l=1}^{L} \frac{TP_l}{TP_l + FP_l}$$
(7)

In some clinical application or epidemiological studies, only one type of code (either the diagnosis or the procedure code) is favoured. Thus, for the MIMIC-III full label setting, we also report Micro- F_1 results on the diagnosis codes (F_1 -diag) and procedure codes (F_1 -proc) separately as in [7].

Furthermore, we report the example-based metric, precision@k as in [7], averaged over all the documents,

⁴https://github.com/acadTags/Automated-Social-Annotation/ tree/master/2%20HAN 585

⁵https://github.com/brightmart/text_classification ⁶https://github.com/jamesmullenbach/caml-mimic

⁷We optimised precision@k for CNN, CNN+att, and Bi-GRU for MIMIC-III and MIMIC-III-50, and micro- F_1 for all other models and for the MIMIC-III-shielding dataset.

⁸We parsed sentences using the rule-based pipeline component in Spacy with adding double newlines as another rule to segment sentences, see https: //spacy.io/usage/linguistic-features#sbd.

⁹https://pubmed.ncbi.nlm.nih.gov/

¹⁰https://www.ncbi.nlm.nih.gov/pmc/

¹¹We thus do not report the BERT results here but make the implementation details and results available on https://github.com/acadTags/ Explainable-Automated-Medical-Coding.

where each precision score is the fraction of the true positive in the top-k labels, having highest score p_{dl} , for the document d. The idea is to simulate the real-world scenario that the system recommending k predicted medical codes and to evaluate the percentage of them being correct. The number of top-ranked635 labels k were set as 8 for MIMIC-III, 5 for MIMIC-III-50 to be consistent to the study [7], and 1 for MIMIC-III-shielding, near to the average number of labels per document (see Table 1).

4.4. Main Results

- We report the mean and the standard deviation (i.e. the 595 square root of variance) of the testing results of 10 runs with randomly initialised parameters for each model. The results of the MIMIC-III-50, MIMIC-III-shielding, and MIMIC-III datasets are shown in Table 2, 3, and 4, respectively. 645
- For the top 50 label dataset (MIMIC-III-50, see Table 2), 600 HLAN performed the best among all experimental settings, achieved significantly better Micro-AUC (91.9%), Micro- F_1 (64.1%), and Precision@5 (62.5%) than the second best model, CNN. This shows the advantage of the hierarchical label-wise
- attention mechanisms for top-50 code prediction. With the 605 same calibration threshold, the precision of HLAN is better than CNN absolutely by 15% (73.2% vs. 57.7%), while recall is lower with a similar absolute value, indicating that tuning the threshold to balance precision and recall could further improve the F_1 scores. 610

For code related to high-risk patients for shielding during the COVID-19 pandemic (MIMIC-III-shielding, see Table 3), results (of Micro-AUC) show that HLAN (96.9%) and HAN (97.6%) performed comparably to the best performed model,

- CNN (97.9%). HLAN obtained a high value of precision@1,660 615 slightly below CNN by 1% (81.2% vs 82.2%), while the difference was not significant (p > 0.05). The better performance of CNN (or HAN) may be because that smaller datasets like MIMIC-III-shielding, with much fewer documents and labels
- (see Table 1), tends to favour models with simpler architectures.665 620 In both MIMIC-III-50 and MIMIC-III-shielding, HA-GRU did not perform better than HLAN, this shows that the labelwise word-level attention mechanisms in HLAN further improved the performance. Also, surprisingly, the HLAN or HAN
- models with the real sentence split did not perform better (up670 625 to 2.8% less Micro- F_1) than using text chunk "sentences" (of 25 continuous tokens) in all three datasets. This is probably because, with the sentence split setting, some tokens and sentences were lost during the padding procedure, which could sig-675

For the full label setting ("MIMIC-III"), HAN has better results of Micro-AUC and precision@8 than the vanilla CNN and Bi-GRU, but worse than the CNN+att approach specifically tuned for this dataset. With label embedding initialisation, CNN+att+LE achieved significant best results on MIMIC-III (an Micro-AUC of 98.6%). It is worth to further explore to enhance the scalability of HLAN so that it can process datasets with large label sizes. Also to note that results of the Macro-level metrics (averaging over labels) were dramatically lower than the Micro-level ones (calculated from documentlabel pairs), showing the strong imbalance of labels in MIMIC-III (see Section 4.1).

Injecting the code relations through label embedding consistently boosted the performance of automated medical coding. It is clear that most models were improved with label embedding initialisation ("+LE"). Models were affected to different extend by label embedding: CNN+att model was mostly improved with "+LE" (an increase of 6.6% Macro-AUC on MIMIC-IIIshielding), the rest models (CNN, Bi-GRU, HA-GRU) being relatively less affected, while there was no significant improvement for HLAN or HAN on the datasets. This may due to the fact that the prior layers, e.g. hierarchical layers and the label-wise attention layers, could already learn some of the label relations. We thus further analyse the LE-initialised layers in Section 4.8 to understand the effect of label embedding initialisation. Besides, most metrics with the "+LE" models also have higher stability (i.e. reduced variance); and low variance is an essential characteristic to deploy a model in the clinical setting.

4.5. Result for each label

Apart from the overall performance of the models, it is also essential to see how the models perform regarding each medical code. Figures 3 show the precision and recall of the five diagnosis codes having the highest and the lowest frequencies in MIMIC-III-50. For this analysis, we selected the three best performing models, CNN, CNN+att, and HLAN, all with label embedding initialisation ("+LE"), in terms of AUC metrics for MIMIC-III-50 (see Table 2). We provide the full per-label results of HLAN+LE with the MIMIC-III-50 and MIMIC-IIIshielding datasets in Table S1-S2 in the supplementary material.

In Figures 3, we can observe that the overall trend of performance is generally consistent to, while not solely dependent on, the label frequency in the training data. For the five most frequent labels, the models achieved around 70%-90% precision and recall. For example, in terms of precision, HLAN obtained

Table 2: Results on MIMIC-III-50 dataset (50 labels)

		Ma	icro			Mi	cro		Top-k
Model	AUC	Р	R	F_1	AUC	Р	R	F_1	P@5
CNN	88.1±0.3	51.5±0.9	67.4±1.0	58.4±0.5	90.9±0.2	55.6±1.1	71.2±0.9	62.4±0.6	61.8±0.3
+LE	88.3±0.3	$53.0 \pm 1.0^{*}$	66.7±1.5	59.1±0.5*	<u>91.3±0.1</u> *	$57.7 \pm 1.4^{*}$	70.4±1.4	$63.4 \pm 0.5^{*}$	<u>62.1±0.3</u>
Bi-GRU	80.6±1.1	47.2±3.2	36.7±2.6	41.2±2.3	85.5±1.0	58.1±3.2	45.8±2.2	51.2±1.9	51.3±1.7
+LE	80.9 ± 0.8	47.3 ± 2.0	$39.2 \pm 2.0^{*}$	42.8 ± 1.5	<u>85.8±0.7</u>	57.5 ± 2.2	$48.4 \pm 2.1^{*}$	$52.5 \pm 1.3^{*}$	<u>52.1±1.2</u>
CNN+att	88.1±0.0	63.1±0.1	$48.4 \pm 0.2^{*}$	$54.8 \pm 0.2^{*}$	91.1±0.0	70.9±0.2	53.1±0.2*	60.7±0.1	60.8±0.1
+LE	88.3±0.0*	$64.3 \pm 0.3^{*}$	46.0 ± 0.1	53.6±0.1	$91.3\pm0.0^{*}$	71.6±0.1*	52.5 ± 0.1	60.6±0.1	$61.6 \pm 0.1^*$
HAN	87.0±0.4	61.7±2.7	46.3±2.3	52.8±1.1	90.1±0.3	68.2 ± 3.1	52.9 ± 2.4	59.4±0.7	59.5±0.7
+LE	87.3±0.4	61.3±3.2	46.9 ± 2.9	53.0 ± 1.1	90.3±0.4	67.9 ± 4.1	54.2 ± 2.8	60.1±0.7	<u>59.9±0.8</u>
HA-GRU	85.3±1.3	59.3±2.5	43.1±4.0	49.9±3.4	89.2±0.9	69.5±0.6	48.7±4.1	57.2±2.8	57.9±1.7
+LE	$86.4 \pm 0.7^{*}$	$62.1 \pm 1.9^{*}$	44.3±2.3	51.7 ± 1.9	90.1±0.5*	$71.1 \pm 1.2^{*}$	50.7±2.3	$59.1 \pm 1.4^{*}$	$59.5 \pm 1.0^{*}$
HLAN	88.4±0.7	65.0±1.2	51.0 ± 2.6	57.1±1.6	91.9±0.4	72.9±0.8	57.3±2.5	64.1±1.4	62.5±0.7
+LE	88.4±0.5	65.5±1.5	50.2 ± 1.1	56.8±0.8	<u>91.9±0.3</u>	73.2±0.6	56.9±1.0	64.0 ± 0.7	62.4±0.6
+sent split	86.9±0.5	63.6±1.3	47.8 ± 2.4	54.5 ± 1.7	90.4±0.3	71.5±1.2	53.8 ± 2.1	61.4±1.2	60.2±0.7

The results of better metric score between the model with label embedding initialisation ("+LE") and the model *not* using LE initialisation are <u>underlined</u>, and the asterisk (*) further marks the paired two-tailed t-tests with .95 significant level (p < 0.05) between them. The best result for each metric (column) is in **bold**. The AUC, F_1 , and P@5 scores in HLAN models with *italics* indicates their significantly improved results (p < 0.05) over the second best model category (i.e. HLAN vs. CNN). The model with lower variance is preferred if the average scores are the same.

Table 3: Results on MIMIC-III-shielding dataset (20 labels)

		Ma	acro			Mi	cro		Top-k
Model	AUC	Р	R	F_1	AUC	Р	R	F_1	P@1
CNN	96.9±0.2*	59.8±1.2	59.6±1.3	59.7±0.9	97.9±0.4*	80.5 ± 1.3	76.2±0.9	78.3±1.0*	82.2±0.8
+LE	96.7±0.2	60.4 ± 2.8	60.4±2.3	60.4±2.3	97.6±0.3	78.8±2.5	76.4±1.8	77.5±0.7	81.6±0.9
Bi-GRU	91.9±1.4	57.4±3.1	43.4±2.2	49.4±2.2	93.6±0.8	77.9±2.8	58.5±1.9	66.8±1.2	72.2±1.6
+LE	92.0 ± 1.6	58.6 ± 1.5	$46.8 \pm 2.3^{*}$	$52.0\pm1.4^{*}$	95.1±0.7*	78.1±2.1	$61.8 \pm 2.7^{*}$	$68.9 \pm 1.6^{*}$	<u>75.1±2.0</u> *
CNN+att	88.9±1.3	46.7±4.6	37.6±2.4	41.7±3.3	93.5±0.2	86.9±1.2*	52.9 ± 2.8	65.7±2.0	70.0±2.6
+LE	$95.5 \pm 0.0^{*}$	62.1±2.2*	$48.4 \pm 1.9^{*}$	$54.4 \pm 2.0^{*}$	<u>96.1±0.0</u> *	83.3±0.5	$61.4 \pm 0.6^{*}$	70.7±0.3*	<u>77.7±0.3</u> *
HAN	96.0±1.4	66.4±2.7	58.2 ± 2.0	62.0 ± 2.0	97.4±0.3	82.9±1.8	68.7±2.4	75.1±1.5	78.1±1.7
+LE	<u>96.4±1.3</u>	65.2 ± 2.1	56.5 ± 2.9	60.5 ± 2.3	97.6±0.3	83.4 ± 1.2	68.2 ± 2.0	75.0±1.2	79.2 ± 1.7
HA-GRU	93.4±2.0	60.9 ± 3.9	51.6±2.8	55.8 ± 3.1	96.7±0.4	83.0±2.1	65.8±2.2	73.4±1.6	80.3±1.5
+LE	93.9 ± 2.0	59.2 ± 4.3	49.7±4.2	54.0 ± 4.1	96.8±0.9	81.3±4.0	66.3 ± 4.1	73.0±3.6	79.1±4.3
HLAN	93.5±2.5	59.8±2.9	53.2 ± 2.6	56.3 ± 2.4	96.9±0.7	81.4±1.8	69.0±2.9*	74.6±1.6	81.2±1.2
+LE	93.5±1.9	60.5 ± 4.2	52.7 ± 5.0	56.3 ± 4.6	96.5±0.4	81.8 ± 2.8	65.6 ± 4.0	72.7±3.1	79.8±3.0
+sent split	94.5 ± 1.2	60.9 ± 2.1	51.7±3.1	55.8 ± 2.3	96.3±0.2	81.4±2.2	64.4 ± 2.5	71.9±1.7	77.8±2.6

The results of better metric score between the model with label embedding initialisation ("+LE") and the model *not* using LE initialisation are <u>underlined</u>, and the asterisk (*) further marks the paired two-tailed t-tests with .95 significant level between them. The best result for each metric (column) is in **bold**. The AUC, F_1 , and P@1 scores in CNN models with *italics* indicates their significantly improved results (p < 0.05) over the second best model category (i.e. CNN vs. HAN or HLAN). The model with lower variance is preferred if the average scores are the same.

highest to 91.7% for 427.31 (Atrial fibrillation) and lowest to 71.4% for 584.9 (Acute kidney failure). For the five least frequent labels, the results were much worse due to the fewer training data for the labels and the imbalance issue. For precision,

- HLAN generally performs better than CNN and CNN+att, especially there is a significant gap for low frequent labels; while for recall, CNN outperforms the other two models. We also note that the precision and recall could be tuned in favour of only one of them through changing the calibration threshold Th
- (now set as the default value, 0.5), considering the need and the preference of the coding work when deploying the model to support coding professionals.

4.6. Model Explanation with Hierarchical Label-wise Attention Visualisation

A critical requirement of the clinical use of automated medical coding systems is their explainability or interpretability. We propose to use label-wise word-level and sentence-level attention mechanisms in HLAN to enhance the explainability of the model. The learned word-level and sentence-level attention scores for the label y_l are $\alpha_{wl} \in (0, 1)$ and $\alpha_{sl} \in (0, 1)$ (see Equations 4 and 5, respectively). For a more concise visualisation, we propose a *sentence-weighted* word-level attention score $\tilde{\alpha}_{wl}$ to only highlight the words from the salient sentences. This adapted word-level attention score is calculated as $\tilde{\alpha}_{wl} = \mu \alpha_{\bar{s}l} \alpha_{wl}$, where $\alpha_{\bar{s}l}$ is the attention score of the sentence where the word is belong to and μ is a hyperparameter to control the

Table 4: Results on MIMIC-III dataset (8,922 labels)

	Macro					Micro						
Model	AUC	Р	R	F_1	AUC	Р	R	F_1	F_1 -diag	F_1 -proc	P@8	
CNN	81.8±0.7	4.5±0.4	3.7±0.5	4.1±0.4	97.0±0.1	51.0±2.8	36.9±1.7	42.8±0.9	41.1±1.0	50.7±0.9	59.6±0.5	
+LE	$82.4 \pm 0.4^{*}$	4.7 ± 0.4	3.6±0.2	4.1 ± 0.2	97.1±0.1	53.0 ± 2.6	36.9±1.2	43.4 ± 0.6	41.7±0.6	51.3±0.9	$60.3 \pm 0.4^{*}$	
Bi-GRU	83.5±1.6	4.9±0.2	3.6±0.5	4.1±0.4	97.3±0.3	52.8±4.6	34.8±2.2	41.8±1.5	39.3±1.6	51.7±1.3	58.9 ± 2.2	
+LE	$84.9 \pm 0.7^{*}$	5.0 ± 0.4	3.6±0.5	4.2 ± 0.5	<u>97.6±0.1</u> *	<u>55.4±4.1</u>	34.8 ± 2.4	42.6±1.5	40 ± 1.6	52.7±1.1	60.3 ± 1.8	
CNN+att	88.6±0.2	7.7±0.2	6.4±0.3	7.0±0.2	98.4±0.0	62.8±0.3*	43.9±0.4	51.7±0.1	50.1±0.2	59.8±0.1	69.4±0.2	
+LE	90.2±0.0*	9.3±0.1*	8.0±0.1*	8.6±0.1*	<u>98.6±0.0</u> *	61.8 ± 0.4	45.6±0.1*	$52.5 \pm 0.1^{*}$	50.7±0.1*	$60.7 \pm 0.1^{*}$	69.7±0.1*	
HAN	88.5±0.1*	$5.4 \pm 0.2^{*}$	$2.7 \pm 0.2^{*}$	$3.6 \pm 0.2^{*}$	98.1±0.1	63.2±3.3	$30.0\pm1.1^{*}$	$40.7 \pm 0.7^{*}$	37.0±0.7*	52.6±0.9*	$61.4 \pm 1.3^{*}$	
+LE	88.2±0.2	5.1±0.2	2.4 ± 0.1	3.3 ± 0.2	<u>98.1±0.0</u>	<u>63.2±1.0</u>	27.6±1.1	38.4±1.0	34.8±1.2	50.6±1.0	59.6±0.6	
+sent split	87.4±0.7	4.7 ± 0.6	2.3±0.4	3.1±0.5	97.9±0.1	60.4 ± 2.2	25.3±2.5	35.6 ± 2.6	31.4 ± 2.6	49.1±2.3	56.3 ± 2.0	

The results of better metric score between the model with label embedding initialisation ("+LE") and the model *not* using LE initialisation are <u>underlined</u>, and the asterisk (*) further marks the paired two-tailed t-tests with .95 significant level between them. The best result for each metric (column) is in **bold**. The model with lower variance is preferred if the average scores are the same.

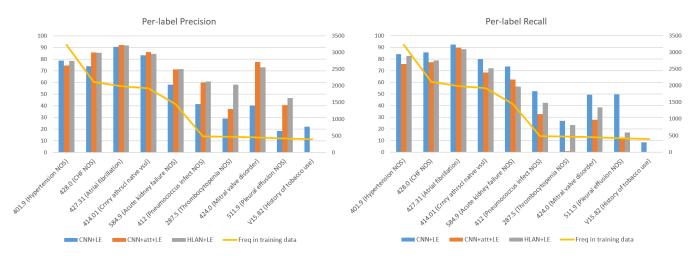


Figure 3: Precision and recall of the five most and the five least frequent ICD-9 diagnosis codes in the MIMIC-III-50 dataset. The bar chart (with the left y-axis) shows the metric score, while the line chart (with the right y-axis) shows the number of occurrences or the frequency ("Freq") of the label in the training data.

magnitude of the final weighted attention score. A greater μ will result in highlighting more words in the clinical note and we empirically set μ as 5. We clip the value of $\tilde{\alpha}_{wl}$ to 1 if it is above 1.

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An example attention visualisation for a random document⁷²⁵ (number 24) in MIMIC-III-50 using the model HLAN+LE with the parsed sentences ("+sent split") is shown in Figure 4. The two columns on the left visualise the sentence-level attention scores α_{sl} for the two codes, 427.31 (Atrial fibrillation) and 428.0 (Congestive heart failure, unspecified), respectively. The⁷³⁰ highlighted sentences are corresponding to the sections "past medical history" and "discharge diagnosis" in the discharge summary. This is in line with our intuition that the key diag-

⁷¹⁵ nosis information is likely to be contained in the two sections. The words are highlighted according to the adapted word-level⁷³⁵ attention score $\tilde{\alpha}_{wl}$. Words related to the code 427.31 is highlighted in yellow and for 428.0 in blue. It is clear that the most salient words are highlighted, and the model successfully ⁷²⁰ recognised the abbreviations and alternative short forms commonly used by clinicians in the clinical note, for example "a fib" as a short form of atrial fibrillation and "chf" as the abbreviation of Congestive Heart Failure. This shows that the proposed HLAN model can learn to recognise the strongly correlated words (e.g. "a fib") related to the label (e.g. the code 427.31) with label-wise attention mechanisms, even given the fact that the label description (i.e. the knowledge that 427.31 is "Atrial Fibrillation") were not fed into the model during training. Other relevant words are highlighted, e.g., "ef" (short for Ejective Fraction), "pressor", and "extremities", which show a correlation to the code 428.0 while not indicating a causal relation to the diagnosis. We also note that the highlighted words like "age" and "drugs" were too general, which could not be directly related to the diagnosis from a clinician's point of view. This may be related to the peaky distribution of the softmax (normalised exponential) function to form the attention scores (see Equation 4), paying the most of the attention to only a few (one or two) words in a long sentence.

For 427.3	1 For 428.0	Document #2	24 in MIMIC	-111-50								
0.02	0.01	admission	date	discharge	date	service	surgery					
0.01	0.01	allergies	patient	recorded	as	having	no	known	allergies	to	drugs	attending
0.01	0.02	major	surgical	or	invasive	procedure	ex	lap	r	hemicolectomy	mucous	fistula
0	0	history	of	present	illness	age	over	f	presented	to	location	un
0	0.01	admitted	to	hospital1	and	taken	directly	to	or	upon	arrival	
0.41	0.8	past	medical	history	pmhx	а	fib	aortic	stenosis	chf	last	ef
0	0	pertinent	results	00pm	blood	wbc	rbc	hgb	hct	mcv	mch	mchc
0	0.01	brief	hospital	course	age	over	f	transferred	from	location	un	and
0	0.02	patient	was	taken	directly	to	the	operating	room	for	an	exploratory
0	0	she	underwent	а	right	hemicolectomy	mucous	fistula	ileostomy	gj	tube	placement
0.01	0.01	fluid	balance	intraoperatively	included	units	ffp	units	plt	u	nits	prbc
0	0	patient	was	kept	intubated	and	taken	directly	to	the	surgical	intensive
0	0.02	she	required	maximum	pressor	support	to	maintain	sufficient	cardiac	index	
0	0.03	patient	did	show	signs	of	distal	ischemia	to	extremities	by	the
0.01	0	family	meeting	at	latter	evening	decided	to	make	patient	cmo	patient
0.01	0	medications	on	admission	last	name	un	amlodipine	mg	qd	benarepril	mg
0.49	0	discharge	diagnosis	cardiopulmonary	arrest	perforated	colon	atrial	fibrillation	ventilatory	support	discharge

Figure 4: An example of interpretation using attention visualisation from the Hierarchical Label-wise Attention Network (HLAN), the chosen example is a random document (index 24) in the MIMIC-III-50 dataset with two true positive labels, ICD-9 code 427.31 (Atrial fibrillation) and 428.0 (Congestive heart failure, unspecified). The two red columns show the sentence-level attention scores for the two codes respectively. The tokens highlighted by yellow (for code 427.31)

or blue (for code 428.0) show the importance of them based on the value of sentence-weighted word-level attention scores. The deeper the colour, the higher the (sentence-weighted) attention scores, and thus the more important the highlight words or sentences contributes to the model prediction. Only the first part (11 tokens) of each sentence was shown for a clearer display.

4.7. Comparison of Model Explanations

- Following the previous section, we further qualitatively anal-⁷⁶⁵ yse and compare the interpretability of the HLAN model and other baseline models. Table 5 shows how CNN, CNN+att, HAN, HA-GRU, and HLAN, all with label embedding initial-isation, highlight the "important" part of a random document
 ⁷⁴⁵ (number 24) to predict two different labels (427.31 and 428.0).⁷⁷⁰ The CNN¹² and CNN+att chose the most salient *n*-grams based
- on the max-pooling and the attention mechanism, respectively
 [7]. HLAN and its downgraded models, HA-GRU and HAN, alternatively, highlighted the important sentences and words.
 The distinction is that HAN has the same highlights of the same⁷⁷⁵
- document for different labels (columns in Table 5), and HA-GRU has the same word-level but different sentence-level highlights across labels, while HLAN can highlight the most salient words and sentences for different labels. This gives HLAN the most comprehensive interpretability among the models. 780

Compared to CNN, we observe that CNN+att generated a more relevant set of *n*-grams. This is in accordance with the conclusion in [7]. We also found that the attention weights from CNN are unstable, i.e. the suggested *n*-grams from CNN were not the same among different runs. Compared to the interpre-⁷⁸⁰ tation with *n*-grams, highlighting the key sentences and words can produce a more comprehensive interpretation, as the latter is based on the whole hierarchical structure of a document.

Especially with the sentence parsing ("HLAN+LE+sent split", see the last row in Figure 5, corresponding to the visualisation in Figure 4), we can clearly see which sections of the discharge summary, along with words, contribute more to predict the label.

It is also interesting to see how the proposed model interpret when it predicted a medical code not previously assigned by the coding professionals. We selected some representative "false positive" results from the HLAN+LE model with sentence splitting in Table 6. We presented the prediction results and the highlighted explanations to an experienced clinician to validate and deduce the potential reason for the error. In Table 6, we observe that the model can explain the predictions with key sentences and words, therefore it is easier for us to know where there may have been a problem. For example, for the first two rows, "doc-68" and "doc-19" in MIMIC-III-50, the highlighted words and sentences are quite relevant to the noncoded, "false positive" ICD-9 code, indicating that there might have been missed coding or the disease was a past disease of the patient.

The false positives in "doc-1" and "doc-65" in MIMIC-IIIshielding are errors related to the wrong correlations learned from the data, particularly regarding the high granularity and subtle difference among sub-type diseases. In "doc-1", the highlighted words "htn elev lipids" show that the patient has a certain type of hypertension, but does not necessarily mean the predicted code 416.0 for "Primary pulmonary hypertension". In "doc-65", the strongly highlighted word "metastasis" actu-

 $^{^{12}}$ We further normalised the scores of *n*-grams in CNN based on max-pooling₇₉₀ from [7] to probabilities, to be comparable to the attention scores in other models.

Table 5: Comparison of model interpretability across deep learning models of true positive predictions on a random document (index 24) in the MIMIC-III-50 dataset

Model	doc-24 to predict 427.31 (Atrial fibrillation)	doc-24 to predict 428.0 (Congestive heart failure, unspecified)						
	<i>n</i> -gram-1 (0.105): admission date discharge date service surgery	<i>n</i> -gram-1 (0.096):surgical intensive care unit she required max-						
CNN+LE	allergies patient recorded as having no known	imum pressor support to maintain sufficient cardiac index						
	<i>n</i> -gram-2 (0.083):surgical or invasive procedure ex lap r hemi-	<i>n</i> -gram-2 (0.075):past medical history pmhx a fib aortic stenosis						
	colectomy mucous fistula ileostomy gj tube placement history of	chf last ef in osteoporosis reflux						
	present illness							
	<i>n</i> -gram-3 (0.075):presented to location un with perforated vis-	<i>n</i> -gram-3 (0.071):on ventilation support family meeting at latter						
	cous hd stable upon transfer to location un	evening decided to make patient cmo patient						
CNN+att+LE	<i>n</i> -gram-1 (0.026):upon arrival past medical history pmhx a fib	n-gram-1 (0.017): pressor support to maintain sufficient cardiac						
CININ+all+LE	aortic stenosis chf last ef in osteoporosis reflux doctor first name	index patient did show signs of distal ischemia to extremities by						
	hx appendectomy many years ago social history non	the afternoon urine output post						
	n-gram-2 (0.023):diagnosis cardiopulmonary arrest perforated	n-gram-2 (0.011):past medical history pmhx a fib aortic stenosis						
	colon atrial fibrillation ventilatory support discharge condition	chf last ef in osteoporosis reflux doctor first name hx appendec-						
	death discharge instructions none followup instructions none	tomy many years ago social history non contributory						
	sent-1 (0.34): medical history pmhx a fib(0.374) aortic stenosis chf	(0.206) last ef in osteoporosis reflux(0.097) doctor first name hx						
HAN+LE	appendectomy many(0.28) years ago social history non contributory							
	sent-2 (0.18): arthritis fosamax q week(0.039) coumadin(0.282) qd discharge medications none discharge disposition expired dis-							
	charg diagnosis cardiopulmonary(0.019) arrest(0.109) perforated(0.134) colon(0.119) atrial(0.173) fibrillation(0.023) ventila-							
	tory(0.047) support discharge condition death							
	sent-3 (0.11): admission(0.201) date(0.263) discharge(0.05) date(0.055) service(0.075) surgery(0.118) allergies(0.062) pa-							
	tient(0.013) recorded(0.054) as having no known allergies to drugs attending first name3(0.021) lf(0.02) chief complaint perforated							
		attending first names(0.021) n(0.02) effet complaint perforated						
	bowel(0.046) major							
HA-GRU+LE	bowel(0.046) major sent-1 (0.62): arthritis fosamax q week coumadin(0.06) qd dis-	Did not predict 428.0 (i.e. false negative)						
HA-GRU+LE	bowel(0.046) major sent-1 (0.62): arthritis fosamax q week coumadin(0.06) qd dis- charge medications none discharge disposition expired discharge							
HA-GRU+LE	bowel(0.046) major sent-1 (0.62): arthritis fosamax q week coumadin(0.06) qd dis- charge medications none discharge disposition expired discharge diagnosis cardiopulmonary arrest perforated colon atrial fibrilla -							
HA-GRU+LE	bowel(0.046) major sent-1 (0.62): arthritis fosamax q week coumadin(0.06) qd dis- charge medications none discharge disposition expired discharge diagnosis cardiopulmonary arrest perforated colon atrial fibrilla- tion(0.94) ventilatory support discharge condition death	Did not predict 428.0 (i.e. false negative)						
	bowel(0.046) major sent-1 (0.62): arthritis fosamax q week coumadin(0.06) qd dis- charge medications none discharge disposition expired discharge diagnosis cardiopulmonary arrest perforated colon atrial fibrilla- tion(0.94) ventilatory support discharge condition death sent-1 (0.54): arthritis fosamax q week coumadin qd discharge	Did not predict 428.0 (i.e. false negative) sent-1 (0.71): medical history pmhx a fib aortic stenosis chf(1.0)						
	bowel(0.046) major sent-1 (0.62): arthritis fosamax q week coumadin(0.06) qd dis- charge medications none discharge disposition expired discharge diagnosis cardiopulmonary arrest perforated colon atrial fibrilla- tion(0.94) ventilatory support discharge condition death sent-1 (0.54): arthritis fosamax q week coumadin qd discharge medications none discharge disposition expired discharge diagno-	Did not predict 428.0 (i.e. false negative) sent-1 (0.71): medical history pmhx a fib aortic stenosis chf(1.0) last ef in osteoporosis reflux doctor first name hx appendectomy						
	bowel(0.046) major sent-1 (0.62): arthritis fosamax q week coumadin(0.06) qd dis- charge medications none discharge disposition expired discharge diagnosis cardiopulmonary arrest perforated colon atrial fibrilla- tion(0.94) ventilatory support discharge condition death sent-1 (0.54): arthritis fosamax q week coumadin qd discharge medications none discharge disposition expired discharge diagno- sis cardiopulmonary arrest perforated colon atrial(1.0) fibrillation	Did not predict 428.0 (i.e. false negative) sent-1 (0.71): medical history pmhx a fib aortic stenosis chf(1.0)						
	bowel(0.046) major sent-1 (0.62): arthritis fosamax q week coumadin(0.06) qd dis- charge medications none discharge disposition expired discharge diagnosis cardiopulmonary arrest perforated colon atrial fibrilla- tion(0.94) ventilatory support discharge condition death sent-1 (0.54): arthritis fosamax q week coumadin qd discharge medications none discharge disposition expired discharge diagno- sis cardiopulmonary arrest perforated colon atrial(1.0) fibrillation ventilatory support discharge condition death	Did not predict 428.0 (i.e. false negative) sent-1 (0.71): medical history pmhx a fib aortic stenosis chf(1.0) last ef in osteoporosis reflux doctor first name hx appendectomy						
	bowel(0.046) major sent-1 (0.62): arthritis fosamax q week coumadin(0.06) qd dis- charge medications none discharge disposition expired discharge diagnosis cardiopulmonary arrest perforated colon atrial fibrilla- tion(0.94) ventilatory support discharge condition death sent-1 (0.54): arthritis fosamax q week coumadin qd discharge medications none discharge disposition expired discharge diagno- sis cardiopulmonary arrest perforated colon atrial(1.0) fibrillation ventilatory support discharge condition death sent-2 (0.18): medical history pmhx a fib(1.0) aortic stenosis chf	Did not predict 428.0 (i.e. false negative) sent-1 (0.71): medical history pmhx a fib aortic stenosis chf(1.0) last ef in osteoporosis reflux doctor first name hx appendectomy						
	bowel(0.046) major sent-1 (0.62): arthritis fosamax q week coumadin(0.06) qd dis- charge medications none discharge disposition expired discharge diagnosis cardiopulmonary arrest perforated colon atrial fibrilla- tion(0.94) ventilatory support discharge condition death sent-1 (0.54): arthritis fosamax q week coumadin qd discharge medications none discharge disposition expired discharge diagno- sis cardiopulmonary arrest perforated colon atrial(1.0) fibrillation ventilatory support discharge condition death sent-2 (0.18): medical history pmhx a fib(1.0) aortic stenosis chf last ef in osteoporosis reflux doctor first name hx appendectomy	Did not predict 428.0 (i.e. false negative) sent-1 (0.71): medical history pmhx a fib aortic stenosis chf(1.0) last ef in osteoporosis reflux doctor first name hx appendectomy						
	bowel(0.046) major sent-1 (0.62): arthritis fosamax q week coumadin(0.06) qd dis- charge medications none discharge disposition expired discharge diagnosis cardiopulmonary arrest perforated colon atrial fibrilla- tion(0.94) ventilatory support discharge condition death sent-1 (0.54): arthritis fosamax q week coumadin qd discharge medications none discharge disposition expired discharge diagno- sis cardiopulmonary arrest perforated colon atrial(1.0) fibrillation ventilatory support discharge condition death sent-2 (0.18): medical history pmhx a fib(1.0) aortic stenosis chf last ef in osteoporosis reflux doctor first name hx appendectomy many years ago social history non contributory	Did not predict 428.0 (i.e. false negative) sent-1 (0.71): medical history pmhx a fib aortic stenosis chf(1.0) last ef in osteoporosis reflux doctor first name hx appendectomy many years ago social history non contributory						
HLAN+LE	bowel(0.046) major sent-1 (0.62): arthritis fosamax q week coumadin(0.06) qd dis- charge medications none discharge disposition expired discharge diagnosis cardiopulmonary arrest perforated colon atrial fibrilla- tion(0.94) ventilatory support discharge condition death sent-1 (0.54): arthritis fosamax q week coumadin qd discharge medications none discharge disposition expired discharge diagno- sis cardiopulmonary arrest perforated colon atrial(1.0) fibrillation ventilatory support discharge condition death sent-2 (0.18): medical history pmhx a fib(1.0) aortic stenosis chf last ef in osteoporosis reflux doctor first name hx appendectomy many years ago social history non contributory sent-1 (0.49): discharge diagnosis cardiopulmonary arrest perfo-	Did not predict 428.0 (i.e. false negative) sent-1 (0.71): medical history pmhx a fib aortic stenosis chf(1.0) last ef in osteoporosis reflux doctor first name hx appendectomy many years ago social history non contributory sent-1 (0.8): past medical history pmhx a fib aortic stenosis						
HLAN+LE	bowel(0.046) major sent-1 (0.62): arthritis fosamax q week coumadin(0.06) qd dis- charge medications none discharge disposition expired discharge diagnosis cardiopulmonary arrest perforated colon atrial fibrilla- tion(0.94) ventilatory support discharge condition death sent-1 (0.54): arthritis fosamax q week coumadin qd discharge medications none discharge disposition expired discharge diagno- sis cardiopulmonary arrest perforated colon atrial(1.0) fibrillation ventilatory support discharge condition death sent-2 (0.18): medical history pmhx a fib(1.0) aortic stenosis chf last ef in osteoporosis reflux doctor first name hx appendectomy many years ago social history non contributory sent-1 (0.49): discharge diagnosis cardiopulmonary arrest perfo- rated colon atrial fibrillation(1.0) ventilatory support discharge	Did not predict 428.0 (i.e. false negative) sent-1 (0.71): medical history pmhx a fib aortic stenosis chf(1.0) last ef in osteoporosis reflux doctor first name hx appendectomy many years ago social history non contributory sent-1 (0.8): past medical history pmhx a fib aortic stenosis chf(0.888) last ef(0.112) in osteoporosis reflux doctor first name						
	bowel(0.046) major sent-1 (0.62): arthritis fosamax q week coumadin(0.06) qd dis- charge medications none discharge disposition expired discharge diagnosis cardiopulmonary arrest perforated colon atrial fibrilla- tion(0.94) ventilatory support discharge condition death sent-1 (0.54): arthritis fosamax q week coumadin qd discharge medications none discharge disposition expired discharge diagno- sis cardiopulmonary arrest perforated colon atrial(1.0) fibrillation ventilatory support discharge condition death sent-2 (0.18): medical history pmhx a fib(1.0) aortic stenosis chf last ef in osteoporosis reflux doctor first name hx appendectomy many years ago social history non contributory sent-1 (0.49): discharge diagnosis cardiopulmonary arrest perfo-	Did not predict 428.0 (i.e. false negative) sent-1 (0.71): medical history pmhx a fib aortic stenosis chf(1.0) last ef in osteoporosis reflux doctor first name hx appendectomy many years ago social history non contributory sent-1 (0.8): past medical history pmhx a fib aortic stenosis						
HA-GRU+LE HLAN+LE +sent split	bowel(0.046) major sent-1 (0.62): arthritis fosamax q week coumadin(0.06) qd dis- charge medications none discharge disposition expired discharge diagnosis cardiopulmonary arrest perforated colon atrial fibrilla- tion(0.94) ventilatory support discharge condition death sent-1 (0.54): arthritis fosamax q week coumadin qd discharge medications none discharge disposition expired discharge diagno- sis cardiopulmonary arrest perforated colon atrial(1.0) fibrillation ventilatory support discharge condition death sent-2 (0.18): medical history pmhx a fib(1.0) aortic stenosis chf last ef in osteoporosis reflux doctor first name hx appendectomy many years ago social history non contributory sent-1 (0.49): discharge diagnosis cardiopulmonary arrest perfo- rated colon atrial fibrillation(1.0) ventilatory support discharge condition death discharge instructions none followup instructions none	Did not predict 428.0 (i.e. false negative) sent-1 (0.71): medical history pmhx a fib aortic stenosis chf(1.0) last ef in osteoporosis reflux doctor first name hx appendectomy many years ago social history non contributory sent-1 (0.8): past medical history pmhx a fib aortic stenosis chf(0.888) last ef(0.112) in osteoporosis reflux doctor first name						
HLAN+LE	bowel(0.046) major sent-1 (0.62): arthritis fosamax q week coumadin(0.06) qd dis- charge medications none discharge disposition expired discharge diagnosis cardiopulmonary arrest perforated colon atrial fibrilla- tion(0.94) ventilatory support discharge condition death sent-1 (0.54): arthritis fosamax q week coumadin qd discharge medications none discharge disposition expired discharge diagno- sis cardiopulmonary arrest perforated colon atrial(1.0) fibrillation ventilatory support discharge condition death sent-2 (0.18): medical history pmhx a fib(1.0) aortic stenosis chf last ef in osteoporosis reflux doctor first name hx appendectomy many years ago social history non contributory sent-1 (0.49): discharge diagnosis cardiopulmonary arrest perfo- rated colon atrial fibrillation(1.0) ventilatory support discharge condition death discharge instructions none followup instructions none sent-2 (0.41): past medical history pmhx a fib(1.0) aortic stenosis	Did not predict 428.0 (i.e. false negative) sent-1 (0.71): medical history pmhx a fib aortic stenosis chf(1.0) last ef in osteoporosis reflux doctor first name hx appendectomy many years ago social history non contributory sent-1 (0.8): past medical history pmhx a fib aortic stenosis chf(0.888) last ef(0.112) in osteoporosis reflux doctor first name						
HLAN+LE	bowel(0.046) major sent-1 (0.62): arthritis fosamax q week coumadin(0.06) qd dis- charge medications none discharge disposition expired discharge diagnosis cardiopulmonary arrest perforated colon atrial fibrilla- tion(0.94) ventilatory support discharge condition death sent-1 (0.54): arthritis fosamax q week coumadin qd discharge medications none discharge disposition expired discharge diagno- sis cardiopulmonary arrest perforated colon atrial(1.0) fibrillation ventilatory support discharge condition death sent-2 (0.18): medical history pmhx a fib(1.0) aortic stenosis chf last ef in osteoporosis reflux doctor first name hx appendectomy many years ago social history non contributory sent-1 (0.49): discharge diagnosis cardiopulmonary arrest perfo- rated colon atrial fibrillation(1.0) ventilatory support discharge condition death discharge instructions none followup instructions none	Did not predict 428.0 (i.e. false negative) sent-1 (0.71): medical history pmhx a fib aortic stenosis chf(1.0) last ef in osteoporosis reflux doctor first name hx appendectomy many years ago social history non contributory sent-1 (0.8): past medical history pmhx a fib aortic stenosis chf(0.888) last ef(0.112) in osteoporosis reflux doctor first name						

* CNN and CNN+att suggested top *n*-grams, while HAN, HA-GRU, and HLAN suggested key sentences ("sent-") and words in the sentences. "+sent split" denotes the HLAN model using real sentence splits. The numbers in the parentheses are the attention scores (e.g. for HLAN, α_w and α_s) in the models. ** For CNN and CNN+att, some of the suggested top-3 *n*-grams were combined together if any of them overlapped; up to five tokens before and after the top *n*-grams were also displayed.

*** For HAN, HA-GRU, and HLAN, the sentences were selected by those with sentence-level attention scores above 0.1 and the words were selected by those with the word-level attention scores above 0.01. Both HAN and HA-GRU predicted 427.31 but not 428.0. The word- and sentence-level attention weights of HAN are shared for all labels, therefore the interpretation is the same for both columns.

ally is related to pancreatic cancer, rather than the more common lung cancer as predicted. This wrong correlation may₈₀₅ be due to the imbalance of vocabularies in the training data: there are 94 (about 2% out of 4,593) discharge summaries in the training data where "pancreatic" and "metasta" appeared together in MIMIC-III-shielding, while there are significant

more discharge summaries (661, about 14%) where "lung" and 810

795

The error in the last example was due to the subtle difference between two codes (280.00 vs 280.03). We noticed some unexpected highlights (e.g. the local oncologist's name code) in the last example ("doc-95" in MIMIC-III-shielding). This₈₁₅

"metasta" appeared together.

may be related to that "hypercalcemia" (appeared in both sent-1 and sent-2) can be caused by cancer, while neutropenia can be caused by treatments like cancer chemotherapy. The word "chemotherapy" was highlighted in another sentence with an attention weight 0.09 (not presented as below the 0.1 threshold) and the word "neutropenia" in the document was not included during the padding process. While it is very likely that the neutropenia was induced by the drug for chemotherapy (that is, 280.03, Drug induced neutropenia), we did not find direct words in the report to point the cause of the disease (thus 280.00, Neutropenia, unspecified, is also appropriate).

In general, we observe that the label-wise attention mecha-

Document index (dataset)	False positive ICD-9	Explanation with the most relevant sentences and words by at-	potential reason
	code	tention scores	
doc-68 (MIMIC-III-50)	427.31 (Atrial fibrillation)	sent-1 (0.32): discharge diagnosis septic shock due to as- cending cholangitis choledocholithiasis atrial fibrillation(1.0) with rapid ventricular response pulmonary emboli deep venous	missed coding
		thrombosis upper gi bleed peptic ulcer	
	-	sent-2 (0.31): past medical history recent pe dvt afib(1.0) htn hypotension hypothyroidism cad mild chf	
	-	sent-3 (0.25): she was found to have bilateral pe s and new afib(1.0) and started on couradin	
doc-19 (MIMIC-III-50)	401.9 (Hypertension NOS, or Unspecified essential hypertension)	sent-1 (0.84): decision made to proceed with primary right to- tal knee arthroplasty past medical history htn(1.0) asthma al- lergies diabetes social history nc family history nc	past disease or missed coding
doc-1 (MIMIC-III-shielding)	416.0 (Primary pulmonary hypertension)	sent-1 (0.45): brief hospital course year old female with h o mild alzheimer s disease cea in htn(0.177) elev(0.145) lipids(0.659) bladder ca who presents as a transfer sent-2 (0.36): past medical history mild alzheimer s disease	subtle difference in language (regarding the type of hypertension)
		l cea in htn(0.284) elev(0.167) lipids(0.518) bladder ca no known metastasis	
doc-65 (MIMIC-III-shielding)	197.0 (Secondary malignant neoplasm of lung)	sent-1 (0.31): brief hospital course yo man with history of metastatic(1.0) pancreatic cancer was admitted with dyspnea new ascites and profound hyponatremia sent-2 (0.3): history of present illness yo cantonese and span-	subtle difference in language (regarding the type of secondary cancer) imbalance of vocabularies
	-	ish speaking male with metastatic(1.0) pancreatic cancer was admitted from the ed with dyspnea altered mental status and sent-3 (0.1): metastatic(1.0) pancreatic cancer evidence of progression of ct abdomen pelvis	or diseases in the training data
doc-95 (MIMIC-III-shielding)	280.00 (Neutropenia, unspecified)	sent-1 (0.24): she has since been found to have a rising ldh and hypercalcemia and decided with her local(0.538) oncologist dr first name8(0.448) namepattern2 name stitle to sent-2 (0.17): at presentation on she developed hypercalcemic with a calcium(1.0) of an elevated ldh	subtle difference between the predicted label 280.00 and the ground truth label 280.03 (Drug induced neutropenia)

Table 6: Examples of false positives of HLAN (with label embedding "+LE" and sentence splitting "+sent split") on MIMIC-III-50 and MIMIC-III-shielding

nisms in the HLAN model can provide a more comprehensive explanation to support the predictions. For wrong or non-coded predictions, the explanations through highlighted sentences and⁸⁴⁰ words can help us better understand the problem. This provides

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an essential reference to help coding professionals use the system and help engineers fix the problems for the next system iteration.

4.8. Analysis of Label Embedding Initialisation

We previously visualised the label embedding from the MIMIC-III dataset reduced to two dimensions using T-SNE, in Figure 2. The visualisation intuitively shows how the label embedding can capture the correlations among ICD-9 codes derived from the coding practice in the clinical setting.

It is also interesting to know, after the dynamic update during training, how the weights in the initialised layers (the final projection layer and the attention layer) preserve the semantics of the label embedding, and why, in a few cases, LE did not result in a significant improvement. We thus extracted the weights in⁸⁵⁵ the learned layers and measured their similarity to the original

label embedding. Based on the idea of label similarity, we calculated the top-10 similar labels for every label based on the pairwise cosine similarity of the rows in the initialised layer weights (e.g. rows such as w_j , w_k in W in Equation 1 or rows V_{wl} in V_w in Equations 4), and also the top-10 similar labels from original label embedding E, and then to see to what extent the two sets of "top-10 similar labels" overlap. We used the Jaccard Index to measure the degree of overlap between the two sets for each label, which is the size of the intersection divided by the size of the union of the two sets. We averaged the Jaccard Index over the labels. Thus the final metric reflects how the layers can retain the semantics, i.e. label similarities, of the label embedding E. We also used the models without the LE initialisation as the control group and calculated this averaged Jaccard Index from their layers for comparison.

The results are displayed in Figure 5. We selected several representative models that either were significantly improved with LE initialisation approach or did not improve with LE according to the results in Tables 2-4. The experiment shows that weights in the final projection layer (and the label-wise attention layer, if applied) with LE initialisation ("+LE") can capture further label similarities from the label embedding. We also observed a strong correlation between the performance improvement with LE (see Tables 2-4) and the increase of averaged Jaccard Index with LE initialisation (i.e. the extent that the ini-

- tialised layers captures the semantics of LE after training, as reflected in Figure 5). The models which are more enhanced by LE (for example, CNN+att, with 6.6% improvement of Macro-905 AUC with "+LE" in Table 3) have a greater averaged Jaccard Index compared to the models without LE (0.76 vs. 0.42-0.43)
- in Figure 5. On the contrary, the models which were *not* improved with LE, e.g. HLAN and HAN, for automated coding, also, was also *not* affected by LE in terms of the averaged Jac-910 card Index. The less effect of LE on HLAN and HAN may be because the hierarchical attention layers (especially with the
- B70 label-wise attention mechanisms) could already model certain label correlations through the document-level matching process. In overall, the analysis supports the idea that LE initial-915 isation, capturing the label correlations, is a key factor to enhance automated coding with deep learning based multi-label
- er5 classification. Since LE can be visualised after dimensionality reduction (see Figure 2), this further serves as a mean to help explain the overall model.

5. Discussion

We have presented the results on the three datasets and analysed the interpretability of models and the layers initialised with label embeddings.

The main advantage of HLAN lies in its model explainability, based on the label-wise word-level and sentence-level attention mechanisms. The qualitative comparison of model explanation suggests that the highlighted key words and sentences 885 from HLAN tend to be more comprehensive and more accurate than, those from HAN or HA-GRU and the n-gram explanation from the CNN related models. Such explainable highlights can be particularly helpful when medical coding professionals need to locate the essential part of a long clinical note. When the model suggests a code, its accompanying explanation could be served as a reference for professionals to validate whether the code should be included. This has the potential to build the users' trust in the deep learning model and help identify missed and erroneous coding. 895

The label embedding initialisation approach boosted the performance and reduced the variance of most models. The method is efficient, not requiring further model parameters. It is independent of the neural encoders, and can thus be applied to₉₄₅ various deep learning models for multi-label classification. Our

various deep learning models for multi-label classification. Our analysis on the LE initialised layers show that they can preserve the semantics in the pre-trained label embeddings and therefore can better capture the label similarity from the data. This further contributes to the explainability of the overall approach. There are a few exceptions that LE did not improve the performance, this may be due to the fact that the hierarchical layers can already model certain label correlations when optimising the document-label matching.

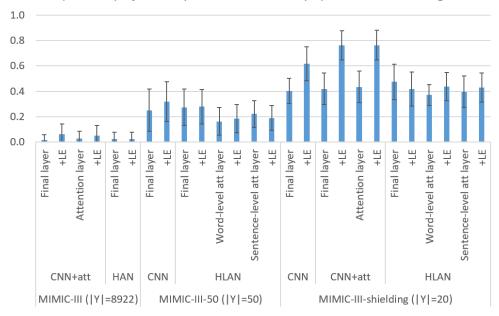
In terms of the performance, for MIMIC-III-50, the HLAN model with LE achieved significant better micro-level AUC (91.9%) and F_1 score (64.1%) than the previous state-of-the-art models; for MIMIC-III-shielding, HLAN and HAN performed comparably to CNN (all around 97-98% micro-level AUC); for MIMIC-III, the previous state-of-the-art model CNN+att was significantly boosted by LE initialisation, achieving best AUC and F_1 scores (Micro-level AUC of 98.6% and Micro-level F_1 of 52.5%).

It is worth nothing that the higher comprehensiveness in explanation from HLAN is at the cost of further memory requirements and the training time¹³. Thus, in practice, if there are only limited computational resources (e.g. a single GPU with 12GB memory), we suggest training HLAN with a fewer number of codes, e.g. equal or less than 50, in a sub-disease domain or for specific tasks (i.e. shielding-related diseases during COVID-19) that require higher model explainability for decision making. We also notice that the vanilla CNN can be trained relatively faster with significantly less memory requirement; HAN and HA-GRU can also be applied as "downgraded" alternatives of HLAN for tasks with larger label sizes. It is also worth to explore to optimise the implementation and to distil the model of HLAN to enable its application to large label sizes.

While training deep learning models can be slow, during the testing phase, the trained models perform reasonably efficient for real-time inference. On average, it requires less than 1/3 second (330 milliseconds) to assign ICD codes with explainable highlights for a discharge summary with a CPU server using HLAN trained from MIMIC-III-50; and the CNN related models can process even faster (see Table S3 in the Supplementary Material). This allows the efficient use of the models in real-time for automated coding.

Also, the calibration threshold (default as 0.5) could be tuned to adjust the precision and recall of the system when deploying it to a coding department. While high precision is obtained when suggesting a few top-ranked predictions, a system with a higher recall can help the coding professionals to prevent

¹³The estimated training and testing time of the models are in Table S3 in the Supplementary Material.



Averaged Jaccard Index of top-10 similar labels between model layers (the final projection layer or the attention layer) and label embedding

Figure 5: Averaged Jaccard Index between the sets of top-10 similar labels derived from the layers (final projection layer and label-wise attention layers with or without label embedding initialisation) and from the label embedding (LE). The higher the averaged Jaccard Index, the more similar the overall semantics between the layer weights and the pre-trained label embedding. Error bars show the standard deviation over the labels. Representative models are selected for all the three datasets. "+LE" means label embedding initialised for the layer indicated in the closest left bar.

missed coding. A higher recall can be achieved by using a lower970 calibration threshold, e.g. 0.3-0.4. The results are also highly varied across labels, as seen in the per-label results in Figures 3. A domain for future studies is therefore to investigate few-shot or zero-shot learning for the rare labels and we noticed one recent related work in [37], which is based on ICD-9 hierarchies975 and descriptions to better predict rare labels.

The results demonstrate the usefulness of label embedding to boost coding performance for most models. In this work, the label embedding was trained with the label sets in the 955 training data using the Continuous Bag of Words algorithm in980 word2vec. Thus, it encodes the similarity of medical codes derived from the real-world coding practice in the critical care unit of the US hospital. This knowledge is distinct from the ICD-9

hierarchy, as visualised in Figure 2 and there may be contradictions between them. The advantage of the former is that it directly learned the label correlation from the existing hospital and does not require external knowledge. There are recently₉₈₅ more studies on leveraging the hierarchy for medical coding as in [38, 39]. One future direction is thus to combine the local

knowledge with external knowledge for the task.

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The analysis of false positives in the model (see Section 4.6 and Table 6) suggests further research in the area of automated990 medical coding. The errors are likely due to missed coding,

past medical history rather than present diseases, nuances of language variations, imbalanced vocabularies, and high label granularity. The highlighted sentences and words helped us better determine the cause of the problems. Since missed coding is very common in real-world practice, as also pointed out recently in [40], it is worth to adapt the current algorithms to capture missing labels and emerging new labels. Information on the report template may further help the model select the relevant part of a discharge summary and differentiate a present disease from a past disease. Unable to capture the subtle variations or labels is potentially related to wrong correlations learned from the imbalance of vocabularies and labels in the dataset. This may be addressed by incorporating various external knowledge.

6. Conclusion

In this paper, we examined the existing deep learning based automated coding algorithms and introduced a new architecture, Hierarchical Label-wise Attention Network (HLAN) and a label embedding initialisation approach for automated medical coding. We tested the approaches on the benchmark datasets extracted from the MIMIC-III database, with the simulated task to predict ICD-9 codes related to the high-risk diseases selected by the NHS for shielding during the COVID-19 pandemic. The experiment results showed that HLAN has a more comprehensive explainability and better or comparative results to the pre_{1035}

- vious state-of-the-art, CNN-based approaches and the downgraded models, HAN and HA-GRU. The proposed label embedding initialisation effectively boosted the performance of the state-of-the-art deep learning models, capturing label correla¹⁰⁴⁰ tions from the dataset, which reflects the coding practice.
- Analyses on the experiment results of this work suggest that future studies are required in several areas: incorporating external knowledge, learning to capture missed coding, rare la¹⁰⁴⁵ bels, and emerging new labels. In particular, automated medical coding work requires to be tested in real-world clinical settings
- and iteratively improved with inputs from relevant professionals such as coders, nurses and clinicians. Thus an open ared⁰⁵⁰ to work on in the future is to adapt automated coding models with human corrections in real-time, which is mostly related to human-in-the-loop machine learning and active learning [41].
- ¹⁰¹⁰ Inspire by these, we plan to further test and develop the ap¹⁰⁵⁵ proach to support the coding department in the NHS. We will consult processionals to identify and address the issues involved in deploying the system to facilitate coding staff and improve efficiency, accuracy, and overall satisfaction.

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