

medIKAL: Integrating Knowledge Graphs as Assistants of LLMs for Enhanced Clinical Diagnosis on EMRs

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Abstract

Electronic Medical Records (EMRs), while integral to modern healthcare, present challenges for clinical reasoning and diagnosis due to their complexity and information redundancy. To address this, we proposed medIKAL (Integrating Knowledge Graphs as Assistants of LLMs), a framework that combines Large Language Models (LLMs) with knowledge graphs (KGs) to enhance diagnostic capabilities. medIKAL assigns weighted importance to entities in medical records based on their type, enabling precise localization of candidate diseases within KGs. It innovatively employs a residual network-like approach, allowing initial diagnosis by the LLM to be merged into KG search results. Through a path-based reranking algorithm and a fill-in-the-blank style prompt template, it further refined the diagnostic process. We validated medIKAL's effectiveness through extensive experiments on a newly introduced open-sourced Chinese EMR dataset, demonstrating its potential to improve clinical diagnosis in real-world settings.

1 Introduction

Electronic Medical Records (EMRs) are the digitized record of a patient's medical and health information and play an important role in the modern healthcare system. However, due to their complexity and information redundancy, clinical diagnosis based on EMRs extremely requires specialized medical knowledge and clinical experience. This demand has led to the development of automated methods to assist and support clinical diagnosis and decision-making.

Recently, large language models (LLMs) have demonstrated great potential in various medical domains (Lee et al., 2023; Lee, 2023; Ayers et al., 2023; Nayak et al., 2023). But directly applying LLMs to the medical field still has raised concerns about the generation of erroneous knowledge and hallucinations because of their lack of specific

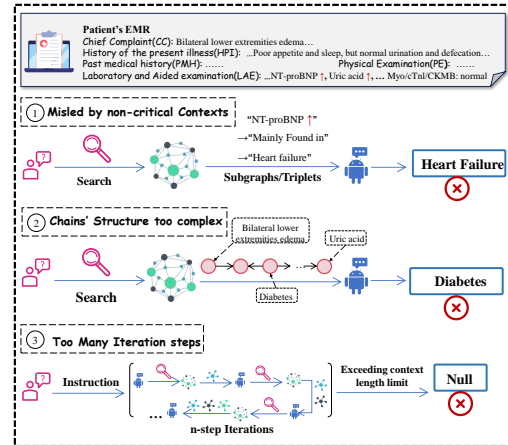


Figure 1: Limitations of existing methods using KG-augmented LLMs for application to EMR diagnostic tasks. ① use subgraphs/triplets to augment context. ② use reasoning chains to augment context. ③ use the iteration-based approach to involve LLMs in KG searching and reasoning.

medical knowledge (Bernstein et al., 2023). Training LLM in the medical domain requires a lot of high-quality data, and the best-performing LLMs available are often closed-source, making further training difficult ((Achiam et al., 2023)). Furthermore, considering that knowledge in the medical field is constantly being updated and iterated, for already trained LLMs, updating their parameters can only be done through retraining, which is extremely time-consuming and expensive (Baek et al., 2023b).

As a classic form of large-scale structured knowledge base, knowledge graphs (KGs) can provide explicit knowledge representation and interpretable reasoning paths and can be continually modified for correction or update. Therefore, KGs become an ideal complement to LLMs (Pan et al., 2024a). However, existing works on "LLM⊕KG" cannot be directly applied to EMR diagnosis tasks, mainly due to the following reasons: (1) Existing ap-

proaches rely on entity recognition in the input text to locate corresponding information in KGs, but they do not differentiate the contributions of different types of entities during searching on KGs. (2) They typically treat triplets or subgraphs obtained from KGs as direct context inputs or simply convert them into natural language, which can easily lead to the problem of exceeding the input length limit and hard to understand for LLMs when encountering complex structures and informative contexts. (3) It was found that when adopting a RAG paradigm, LLMs tend to overly rely on the knowledge in the provided context and fail to fully utilize their internal knowledge, making it easy to be misled by incorrect knowledge (Baek et al., 2023a).

In this paper, we propose a simple yet effective framework called medIKAL (Integrating Knowledge Graphs as Assistants of LLMs). Specifically, unlike other conventional approaches, we assign different weights to entities in the medical record based on their type, which enables us to more precisely localize possible candidate diseases in the KG. Meanwhile, in order to prevent the results from relying too much on the knowledge graph, we drew inspiration from the idea of "residual networks" to allow LLM to first diagnose without relying on external knowledge, and then merge the diagnosis results with the search results of the knowledge graph. Subsequently, we propose a path-based rerank algorithm to rank candidate diseases. Finally, we designed a special fill-in-the-blank style prompt template to help LLMs to better inference and error correction.

In summary, our contributions can be abbreviated as: (1) We raised the problem of a shortage of high-quality open-source Chinese electronic medical record data and we introduced an open-sourced Chinese EMR dataset. (2) We proposed an effective method that allows LLMs to handle information-dense and highly redundant electronic medical records to make effective diagnoses. (3) We conducted extensive experiments on our collected EMR dataset to demonstrate the effectiveness of medIKAL.

2 Related Work

2.1 Clinical Diagnosis and Prediction on EMRs

Electronic medical records (EMRs) provide detailed medical information about patients, including symptoms, medical history, test results, and

treatment records, and are widely used in patient care, clinical diagnosis, and treatment (Xu et al., 2024). Prior research has extensively focused on designing deep learning models for EMR data, addressing downstream tasks such as disease diagnosis and risk assessment (Gao et al., 2020; Xu et al., 2022; Wang et al., 2023b).

LLMs have demonstrated impressive performance in various medical tasks, including disease diagnosis and prediction in EMRs. Researchers have explored multiple approaches: Jiang et al. (2023a) used LLMs and biomedical knowledge graphs to construct patient-specific knowledge graphs, processed with a Bidirectional Attention-enhanced Graph Neural Network (BAT GNN); RAM-EHR (Xu et al., 2024) transformed multiple knowledge sources into text format, utilizing retrieval-enhanced and consistency-regularized co-training; DR.KNOWS (Gao et al., 2023) combined a knowledge graph built with the Unified Medical Language System (UMLS) and a clinical diagnostic reasoning-based graph model for improved diagnosis accuracy and interpretability; REALM (Zhu et al., 2024) integrated clinical notes and multivariate time-series data using LLMs and RAG technology, with an adaptive multimodal fusion network. Most studies focus on English EMR datasets like MIMIC-III (Johnson et al., 2016), which primarily contains ICU data and may not suffice for modeling mild cases, rehabilitation, or routine treatments. Research on Chinese EMR datasets remains limited.

2.2 Knowledge Graph Augmented LLM

Knowledge graphs have advantages in dynamic, explicit, structured knowledge representation and storage, and easy addition, deletion, modification, and querying (Pan et al., 2024b), which has led to increasing interest among researchers in exploring the integration of knowledge graphs with large language models. One typical paradigm is to incorporate knowledge graph triplets into the training data during the training phase and obtain their embedding representations through graph neural network modules (Zhang et al., 2019; Sun et al., 2021; Li et al., 2023; Huang et al., 2024). However, LLMs often have a large-scale requirement for pre-training corpora, making it difficult and costly to find or create knowledge graphs of a matching scale (Wen et al., 2023). More importantly, combining knowledge graphs with LLMs through embedding can result in the loss of their original ad-

vantages, such as interpretability of reasoning and efficiency of knowledge updates.

In recent studies, researchers have attempted to integrate KGs with LLMs through prompts (Wen et al., 2023; Wu et al., 2024; Yang et al., 2024; Wang et al., 2023a). They typically identify entities in the input text and locate the corresponding triplets or subgraphs in the KG, which are then transformed into natural language (Wen et al., 2023), entity sets (Wu et al., 2024), or reorganized triplets (Yang et al., 2024), etc., and concatenated with the input prompts to provide additional knowledge to LLMs. Another approach is to use an iterative strategy where the LLM acts as an agent to explore and reason step-by-step on the KG until it obtains sufficient knowledge or reaches the maximum number of iterations (Sun et al., 2023; Jin et al., 2024). However, this approach is more suitable for shorter questions. In scenarios with longer contexts, larger knowledge graph scales, and more complex structures, it can result in excessive interactions with the LLM and the inability to find the correct paths in the knowledge graph.

3 Method

3.1 EMR Summarisation and Direct Diagnosis via LLM

Considering that the EMRs contain a large amount of redundant information, direct use is easy to cause interference in the diagnostic process. So we first designed a series of questions to prompt LLM to summarize the key information in the EMR, such as patient symptoms, medical history, medication usage, medical visits, etc. Detailed prompt templates are shown in Table 10 and 11 in Appendix F. This process can be represented as:

$$\mathcal{M} = \text{LLM}([\text{Prompt}_{\text{sum}}, \mathcal{M}_{\text{orig}}]) \quad (1)$$

where $\mathcal{M}_{\text{orig}}$ represents the original input medical record, \mathcal{M} represents the medical record after decomposition and summarization, and $\text{Prompt}_{\text{sum}}$ is the textual prompt.

Based on the decomposed and summarized medical record, we allow the LLM to rely on its internal knowledge for preliminary diagnosis and obtain a set of potential diseases \mathcal{D}_{LLM} . This process can be represented as:

$$\mathcal{D}_{\text{LLM}} = \text{LLM}([\text{Prompt}_{\text{diag}}, \mathcal{M}]) \quad (2)$$

where $\text{Prompt}_{\text{diag}}$ denotes the textual instruction used to guide the LLM in performing preliminary

diagnosis and providing predicted diseases (see Table 12 in Appendix F).

3.2 Candidate Disease Localization and Reranking via KG

3.2.1 Entity Recognition and Matching

Before the knowledge graph search process, we perform entity recognition on the summarized EMR \mathcal{M} using a pre-trained NER model. This process can be represented as:

$$\mathcal{E}_{\mathcal{M}} = e_1, e_2, \dots, e_{|E|} = \text{NER}(\mathcal{M}) \quad (3)$$

Where the entity set extracted from the EMR is denoted as $\mathcal{E}_{\mathcal{M}}$, and NER denotes the pre-trained NER model.

Then for every $e_i \in \mathcal{E}_{\mathcal{M}}$, we link it to the corresponding node in the knowledge graph \mathcal{G} using dense retrieval methods. Specifically, given an entity $e_i \in \mathcal{E}_{\mathcal{M}}$, we use an encoding model to get the embedding of e_i , and calculate the similarity score between e_i and each entity node u_j in \mathcal{G} 's entity node set $\mathcal{E}_{\mathcal{G}}$, and the entity node with the highest similarity score is considered as a match. This process can be formulated as follows:

$$\hat{u}_i = \arg \max_{u_j \in \mathcal{E}_{\mathcal{G}}} \text{sim}(\text{enc}(e_i), \text{enc}(u_j)), \quad (4)$$

Where enc denotes the encoding model, and \hat{u}_i denotes the matched entity node. Finally, the set of matched entities is denoted as $\mathcal{E}_{\mathcal{Q}}$.

3.2.2 Candidate Disease Localization Based on Entity-Type Weights

Most of the previous work using KG to augment LLMs has not made a strict distinction between entity types when using entities for the knowledge graph search process. However, in the EMR, different types of entities are supposed to contribute differently to the diagnosis of a disease. For example, the association between a patient's current symptoms and the disease is more direct and closer.

So in this paper, we propose an entity type-driven method for candidate disease localization and filtering. For every entity $e_i \in \mathcal{E}_{\mathcal{Q}}$, we assign a contribution weight w_{t_i} according to its entity type t_i . Then we search for disease nodes in the 1-hop neighbors of e_i in \mathcal{G} and obtain the set of disease nodes \mathcal{D}_i , where the score of each disease in \mathcal{D}_i will be increased by w_{t_i} . The algorithm description of the above process can be found in Algorithm 1 in Appendix B. After getting the potential disease set $\mathcal{D}_{\mathcal{G}}$ generated by the KG search process, we

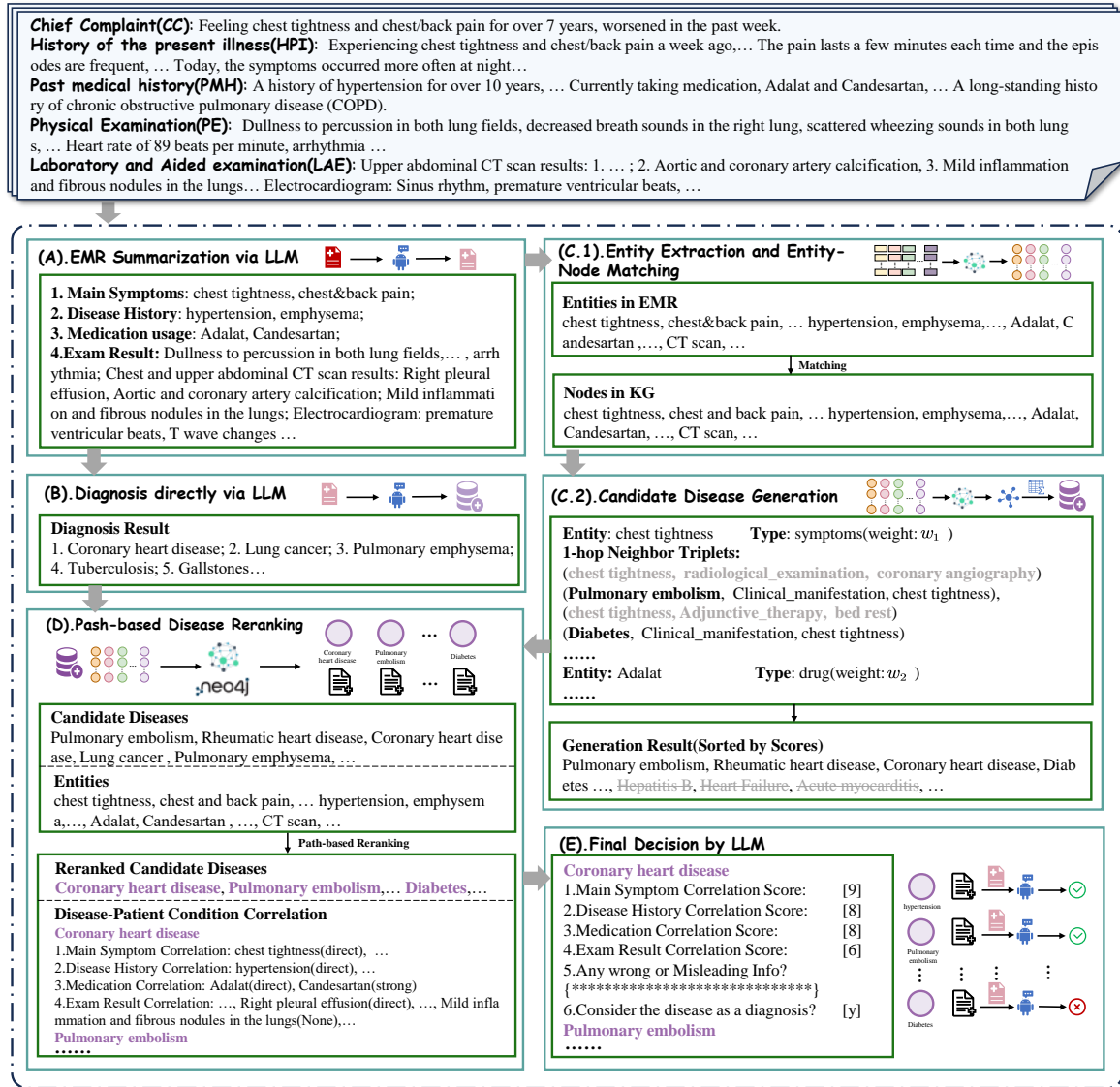


Figure 2: The overall workflow of medIKAL. It contains three main modules, namely: **Module 1.** preprocess before KG search (A, B, and C.1); **Module 2.** Candidate Disease Localization and Reranking via KG (C.2 and D); **Module 3.** Collaborative Reasoning for LLM and KG (E).

merge $\mathcal{D}_{\mathcal{G}}$ with the potential disease set \mathcal{D}_{LLM} obtained through LLM in Section 3.1, resulting in a candidate disease set $\mathcal{D}_{can} = \mathcal{D}_{LLM} \cup \mathcal{D}_{\mathcal{G}}$. Here we have drawn inspiration from the idea of residual networks (He et al., 2016). We hope to make more use of the LLM’s internal knowledge in this way, rather than relying solely on the knowledge graph for searching correct diagnosis.

3.2.3 Candidate Disease Reranking Based on Paths.

In actual clinical diagnosis, doctors usually make a diagnosis based on a series of information such as the patient’s symptoms, medical history, examination results, etc. Therefore, a correct diagnosis

should be correlated with most of the patient information. In order to model this correlation, we propose a path-based reranking algorithm. Specifically, we define $\text{dist}(\mathcal{D}_i, e_j)$ to denote the shortest path distance between disease \mathcal{D}_i and entity $e_j \in \mathcal{E}_{\mathcal{Q}}$ on \mathcal{G} . Diseases with closer total distances to the entity set $\mathcal{E}_{\mathcal{Q}}$ are considered to have a stronger association with the patient’s information, making them more likely to be the correct diagnostic results. The specific process of path-based reranking can be found in Algorithm 2.

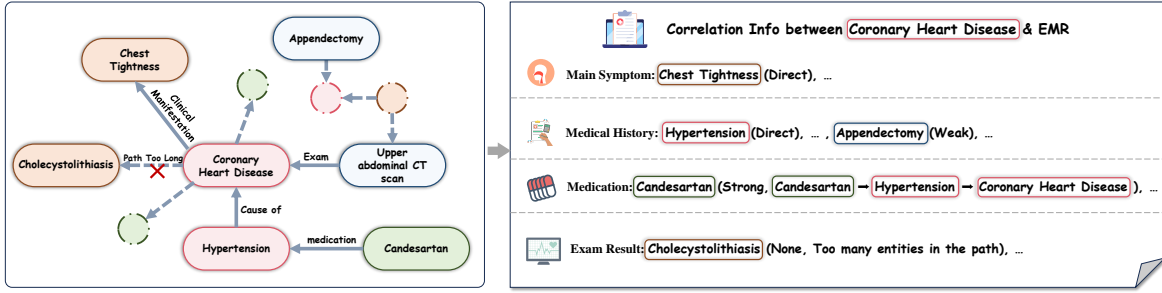


Figure 3: An illustration of how to combine rerank process with the knowledge construction process.

3.3 Collaborative Reasoning between LLM and KG Knowledge

After completing the search and reranking process based on the knowledge graph, we reconstructed the search results to provide additional contextual information for LLM for collaborative reasoning.

3.3.1 Reconstruction of KG Knowledge

EMRs are different from conventional medical QA tasks. Even though we have previously summarized them, they are still information-dense and complex-context structures, so the retrieved KG knowledge will also become extensive. If we still follow previous work and directly input triplets or knowledge chain paths as context knowledge, it would lead to overly chaotic structures that LLMs can hardly understand, which increases the possibilities of hallucination. Therefore, in this paper, we propose a way to reconstruct knowledge graph information. For each candidate disease $\mathcal{D}_i \in \mathcal{D}_{rerank}$, we classify and organize the information related to \mathcal{D}_i according to several aspects like the correlations between \mathcal{D}_i and the patient’s main symptoms, or between \mathcal{D}_i and the patient’s medical history, etc. An example illustration is shown in Figure 3.

In this way, we transform the information of paths and entities retrieved from the knowledge graph into a semi-structured representation of knowledge, which maximizes the manifestation of the association between each candidate disease and the content of the medical record, enabling the model to make more intuitive judgments and analyses. Moreover, since the association between the majority of entities and diseases has already been established during the processing of Section 3.2.2 and Section 3.2.3, the knowledge reconstruction process does not require re-searching \mathcal{G} , avoiding additional time consumption.

3.3.2 Clinical Reasoning and Diagnosis Based on Fill-in-the-Blank Prompt Templates

Based on the reconstructed knowledge described above, we designed a special prompt template in a fill-in-the-blank style to make the reasoning paths of LLM more rational. We guide LLM to quantitatively evaluate the degree of correlation between a specific disease \mathcal{D}_i and the aspects mentioned above, giving a score ranging from 0 to 10 (the higher the score, the higher the degree of correlation), and then calculate a total score. If the total score is higher than a pre-defined threshold θ , we consider the current candidate disease \mathcal{D}_i as one of the final diagnostic results. Additionally, to ensure the self-consistency of LLM, we also check the consistency between this total score and the prediction made by LLM. If they are inconsistent, we will check the original prediction \mathcal{D}_{LLM} to decide whether to drop \mathcal{D}_i . The specific prompt template can be found in Table 13 in Appendix F and relevant case studies can be found in Appendix 5.

4 Experiments

4.1 Experimental Setup

4.1.1 Datasets

CMEMR Dataset Construction: Considering the current lack of high-quality and widely covered EMR datasets in the Chinese community, we construct a dataset CMEMR (Chinese Multi-department Electronic Medical Records) collected from a Chinese medical website¹. We filtered the collected electronic medical records, excluding those with existing problems or missing key information. The details of the dataset can be seen in Table 5 in the Appendix. In order to ensure the correctness and usability of the collected medical records, we randomly sampled a batch of medical

¹ <https://bingli.iyyi.com/>

records in each department and consulted the corresponding department experts, mainly focusing on the correctness of the diagnosis results (i.e., the labels of our task).

In addition, to further validate our proposed method, we selected the following three datasets as supplements: (1) **CMB-Clin** (Wang et al., 2023c): The CMB-Clin dataset contains 74 high-quality, complex, real EMRs, each of which will contain several medical QA pairs. To be consistent with our approach, we simplify the task of this dataset to a pure disease diagnosis task. (2) **GMD** (Liu et al., 2022): The GMD dataset was constructed based on EMRs. Each sample in the dataset contains a target disease along with its explicit and implicit symptom information. (3) **CMD** (Yan et al., 2023): The CMD dataset is a follow-up to the GMD dataset. Its format is the same as the GMD dataset, and also sourced from EMRs. The only difference is that CMD contains a more variety of diseases and symptoms.

4.1.2 Baselines

We compared our proposed medIKAL with three series of baseline methods: LLM-only, LLM \oplus KG, and LLM \otimes KG (Sun et al., 2023):

LLM-only: They do not rely on external knowledge and only use the LLMs’ internal knowledge for reasoning, including CoT (Wei et al., 2022), ToT (Yao et al., 2024), and Sc-CoT (Wang et al., 2022)).

LLM \oplus KG: We selected four representative works, namely MindMap (Wen et al., 2023), ICP (Wu et al., 2024), HyKGE (Jiang et al., 2023b), and KG-rank (Yang et al., 2024), all of which are aimed at medical question-answering and reasoning tasks, so we believe they are highly relevant to our work in this paper.

LLM \otimes KG: This is the concept proposed by (Sun et al., 2023). It enables LLMs to participate in the search and reasoning process on KGs, check whether the current knowledge is sufficient to answer the question, and make decisions for the subsequent search process iteratively. We selected ToG (Sun et al., 2023) and Graph Chain-of-Thought (Jin et al., 2024) as baselines.

4.1.3 Evaluation metric

To enhance the scientific rigor and effectiveness of the evaluation, particularly in identifying disease diagnoses, following (Fan et al., 2024), we adopted the International Classification of Diseases (ICD-

10) (Percy et al., 1990) as the authoritative source and link standardized disease terminologies with natural language based diagnostic results. Initially, we extract disease entities from the diagnostic results and the label in the EMR. Then we implement a fuzzy matching process with a predefined threshold of 0.5 to link these disease entities with ICD-10 terminology, building two normalized disease sets $S_{\mathcal{D}}$ and $S_{\mathcal{R}}$. Finally we use these two sets to calculate the Precision, Recall and F1-score metrics. More details are shown in Appendix D.

4.1.4 Implementation Details

For the backbone model, we choose Qwen models with different parameter scales ([7B, 14B, 72B]). In all experiments, we set *do_sample* to false for consistent responses.

For the knowledge graph, we choose the CPubMed-KG. For the NER model mentioned in section 3.2.1, we utilize the RaNER (Wang et al., 2021) model released by Tongyi-Laboratory. For the Entity-node matching process in section 3.2.1, we choose CoROM (Long et al., 2022) model as our embedding model. The further implementation details are listed in Appendix C.

4.2 Experimental Results

4.2.1 Overall Performance

The main experimental results on CMEMR dataset are shown in Table 1. From the results, we can draw the following analysis:

(1) Our method significantly outperforms other baselines using LLM \oplus KG paradigm on CMEMR dataset, which demonstrated the effectiveness of our method on EMR-diagnosis task.

(2) The methods using LLM \otimes KG (i.e., ToG (Sun et al., 2023) and Graph-CoT (Jin et al., 2024)) perform poorly on EMR-diagnosis Tasks, since they are designed for short multi-hop QA task. The iteration steps and the complexity of beam search increase greatly as the amount of context and the size of KG increase, which makes it easily reach the upper limit of the number of iterative steps without collecting enough information, or exceed the input length limit of LLMs.

(3) As we expected, the performance of medIKAL improves with the scale of backbone models due to the increase of model’s reasoning and instruction-following ability. Considering the plug-and-play and train-free nature of our method, it can be flexibly deployed to backbone models of different sizes depending on the needs of different scenarios.

Methods		Qwen-7b-chat			Qwen-14b-chat			Qwen-72b-chat		
		R	P	F1	R	P	F1	R	P	F1
I	Direct	41.07	31.23	35.48	42.98	32.50	37.01	45.12	34.45	39.06
	CoT	41.24	31.06	35.43	42.58	31.67	36.32	46.01	33.19	38.56
II	ToT	39.25	31.77	35.11	43.19	32.56	37.12	45.45	34.87	39.46
	SC-CoT	41.99	31.69	36.12	42.34	32.90	37.40	45.49	34.59	39.29
III	MindMap	41.42	32.30	36.29	43.59	33.81	38.08	45.14	35.62	39.81
	KG-Rank	39.13	28.61	33.05	41.34	31.45	35.72	44.79	32.95	37.96
	ICP	40.13	30.67	34.76	41.58	30.23	35.00	44.00	32.38	37.30
	HyKGE	42.05	32.42	36.61	43.76	33.45	37.91	45.91	34.30	39.26
	ToG	38.78	26.94	31.79	39.09	27.31	32.15	40.39	27.81	32.93
	Graph-CoT	35.90	24.01	28.77	38.67	25.11	30.44	39.68	27.48	32.47
	Ours	42.16	32.86	36.93	43.96	33.65	38.12	46.43	35.72	40.37

Table 1: Experimental results on CMEMR dataset with different scale of backbone models. The best results are highlighted in bold.

Methods		CMB-Clin			GMD			CMD		
		R	P	F1	R	P	F1	R	P	F1
I	Direct	40.35	26.77	32.18	42.01	21.03	28.02	50.26	25.11	33.48
	CoT	40.66	27.23	32.62	42.44	21.30	28.36	51.02	25.49	33.99
II	ToT	39.94	25.90	31.42	41.68	20.80	27.75	49.39	24.48	32.73
	SC-CoT	41.10	26.31	32.08	42.73	21.37	28.49	51.14	25.57	34.09
III	MindMap	39.26	29.24	33.51	41.44	21.18	28.03	49.75	25.62	33.82
	KG-Rank	41.70	27.12	32.86	38.16	19.54	25.84	47.91	23.92	31.90
	ICP	40.27	25.54	31.25	39.38	19.63	26.20	46.26	23.15	30.85
	HyKGE	41.53	28.21	33.59	40.33	21.36	27.92	48.67	24.35	32.45
	ToG	35.41	19.18	24.88	41.76	20.85	27.81	50.73	25.24	33.70
	Graph-CoT	36.35	20.66	26.07	38.13	19.06	25.54	49.07	24.51	32.69
	Ours	41.89	27.68	33.33	42.37	21.43	28.46	51.26	25.74	34.27

Table 2: Experimental results on CMB-Clin, GMD, and CMD datasets using Qwen-7B-chat. The best results are highlighted in bold.

We also tested our method on three additional datasets and the experimental results are shown in Table 2. Our method performs stably on the CMB-Clin dataset, whose data format is also standard EMRs. On the GMD and CMD datasets, there is a slight degradation in the performance of our method. This is because although GMDs and CMDs are also constructed using EMRs, they contain too little patient information (only symptoms), which can easily localize to other related diseases on the knowledge graph leading to errors.

4.2.2 In-depth Analysis

How do different knowledge graph augmented prompts affect medIKAL’s performance? In order to verify our proposed special prompt template’s superiority, we compare it with sev-

eral knowledge graph-augmented prompt templates, including entities (Wu et al., 2024), relevant triplets (Yang et al., 2024), natural language, reasoning chains (Jiang et al., 2023b), and mindmap (Wen et al., 2023). The experimental results are shown in Table 3. According to the results, using relevant entities is very ineffective as it does not utilize the relational information contained in the knowledge graph at all. For the reasoning chains and mindmap, due to the information-intensive nature of EMR data, they can easily form overly large and complex-structure prompt contexts, making it difficult for LLMs (especially models with small parameters) to reason.

Does medIKAL integrate KG and LLM better compared with other baselines? The problem with most of the existing work based on knowledge

Methods	R	P	F1
Relevant Entities	39.22	28.74	33.17
Natural Language	39.88	28.92	33.52
Relevant Triples	40.26	29.61	34.12
Reasoning Chains	40.97	31.16	35.39
MindMap	41.10	31.41	35.60
FBP(ours)	42.16	32.86	36.93

Table 3: Performances of medIKAL using different knowledge graph-augmented prompt templates on CMEMR dataset. Note that we kept all the rest parts of the medIKAL and only replaced the final “fill-in-the-blanks” prompts with other methods to conduct this experiment.

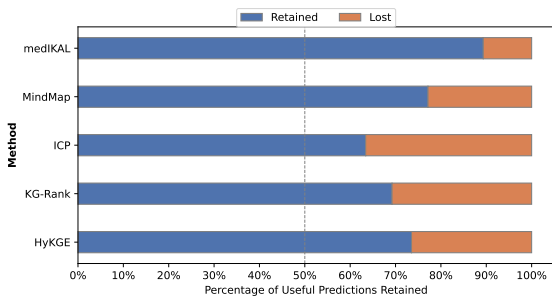


Figure 4: Evaluation results for medIKAL and other baseline methods' capabilities of utilizing LLM's internal knowledge. "Retained" denotes that the useful diagnosis from LLM's original predictions are kept as final results, and "Lost" denotes the opposite.

graphs is that the models can be overly dependent on the information obtained from KG and fail to use their own knowledge. Therefore, we counted the proportion of useful predictions in the original predictions of the model retained by medIKAL and other baseline methods. From the experimental results in Figure 4, medIKAL is able to minimize the model's over-reliance on knowledge graph knowledge and retains the majority of useful predictions compared to other baselines.

Moreover, from the case study in Figure 5, we can find that medIKAL can not only complement (Figure 5-(a)) and correct (Figure 5-(d)) the predictions of LLM using KG, but also effectively guide LLM to analyze and reason (Figure 5-(b)). Besides, the cross-validation approach through quantitative assessment and model judgment can also effectively improve the fault tolerance for LLMs' hallucination(Figure 5-(c)).

4.2.3 Ablation Study

We conduct the following ablation studies to demonstrate the importance of different modules

Method	R	P	F1
medIKAL	42.16	32.86	36.93
w/o SUM	41.56	32.37	36.39
w/o ETW	41.19	29.88	34.63
w/o PR	41.91	32.44	36.57
w/o RI	40.16	30.32	34.55

Table 4: Ablation study results on CMEMR dataset. *w/o* indicates removal of the corresponding module. "SUM" denotes "summarization". "ETW" denotes "Entity Type Weight". "PR" denotes "Path-based Reranking". "RI" denotes "Resnet-like Integration".

in medIKAL.

(a).*w/o* SUM (summarization): Remove the summarization step when pre-processing medical records and instead use the raw content directly.

(b).*w/o* ETW (Entity-Type Weight): Remove the entity-type weight when performing entity-based candidate disease searches, with all entities contributing equal weights.

(c).*w/o* PR (Path-based Reranking): Remove the reranking process for candidate diseases.

(d).*w/o* RM (Resnet-like Merging): Do not integrate the LLM's direct diagnosis result into the candidate disease.

The results in Table 4 show that both removing the "SUM" module and the "ETW" settings can seriously interfere with the performance, as the former leads to the introduction of a lot of redundant information in the original EMRs, while the latter leads to unimportant entities overly influencing the results. Removing the "RM" module would result in results that are entirely dependent on the KG search process, while the internal knowledge of the LLM is almost completely unused, thus causing a severe performance decrease.

5 Conclusion

In this paper, we proposed medIKAL, a framework that seamlessly integrates LLMs with knowledge graphs to enhance clinical diagnosis on EMRs, with its key innovation being the weighted importance assignment to medical entities and a resnet-like integration approach. Experimental results showed that medIKAL significantly outperforms baselines, demonstrating its potential to improve diagnostic accuracy and efficiency in real-world clinical settings. medIKAL offers a promising direction for AI-assisted clinical diagnosis, paving the way for more advanced healthcare applications.

549 Limitations

550 The limitations of collected CMEMR dataset.

551 Although we have meticulously examined, desensi-
552 tized, and verified the CMEMR dataset with medi-
553 cal experts, occasionally, the quality of the medical
554 records may still fall short in actual experiments.
555 Additionally, due to the limited sources of data, our
556 medical record dataset exhibits an uneven distribu-
557 tion across departments.

558 The limitations of proposed medIKAL frame- 559 work.

560 Although medIKAL has demonstrated its
561 effectiveness and great potential in the healthcare
562 field, it still has some limitations. Firstly, while it
563 is not strictly limited to EMR format inputs, it re-
564 quires a high amount of information from the input
565 data samples. When the input data information is
566 sparse, the improvement in model reasoning per-
567 formance by medIKAL decreases, and there is also
568 an increased risk of hallucinations. Furthermore,
569 medIKAL is unable to fully utilize numerical types
570 of medical test results through calculation. Ad-
571 dressing this issue is a key problem that needs to
be solved in our future work.

572 References

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A Detailed Information of the CMEMR dataset

Specific information on the CMEMR dataset is shown in Table 5.

B Algorithms for medIKAL

We summarize the comprehensive algorithmic procedure of ToG and ToG-R, as shown in Algorithm 1 and 2.

C Detailed Setting-ups for Different Modules in medIKAL Workflow

C.1 Details of the NER Model

The RaNER (Wang et al., 2021) model we use in this paper is released by Tongyi-Laboratory, which is trained on the CMeEE dataset (Zhang et al., 2022). RaNER adopts the Transformer-CRF model, using StructBERT as the pre-trained model base, integrating the relevant sentences recalled by external tools as additional context, and employing Multi-view Training for training. It can recognize a total of 9 types of entities, including body (bod), department (dep), disease (dis), drugs (dru), medical equipment (equ), medical examination items

Algorithm 1 Entity Type-driven Candidate Disease Localization and Filtering

Require: Entity Set \mathcal{E}_Q , Knowledge graph \mathcal{G} , Number of candidate diseases $topm$

Ensure: Candidate disease set \mathcal{D}_{can}

- 1: Initialize the set of diseases $\mathcal{D} \leftarrow \emptyset$
- 2: **for** each entity $e_i \in \mathcal{E}_Q$ **do**
- 3: Assign a contribution weight w_{t_i} according to its entity type t_i
- 4: Obtain 1-hop neighbor triplets in \mathcal{G} to locate relevant diseases $\mathcal{D}_i = \{d_{i1}, d_{i2}, \dots, d_{in}\}$
- 5: **for** each disease $d_{ij} \in \mathcal{D}_i$ **do**
- 6: **if** $d_{ij} \in \mathcal{D}$ **then**
- 7: Add w_{t_i} to the score of d_{ij}
- 8: **else**
- 9: Add d_{ij} to \mathcal{D} with an initial score w_{t_i}
- 10: **end if**
- 11: **end for**
- 12: **end for**
- 13: Sort the diseases in \mathcal{D} in descending order based on their scores
- 14: Select the $topm$ diseases to form \mathcal{D}_G
- 15: Merge \mathcal{D}_G with \mathcal{D}_{LLM} to form $\mathcal{D}_{can} \leftarrow \mathcal{D}_{LLM} \cup \mathcal{D}_G$
- 16: **return** \mathcal{D}_{can}

Algorithm 2 Candidate Disease Reranking Based on Paths

Require: Subgraph $\mathcal{G}_s = (V, E)$, Set of candidate diseases \mathcal{D}_{can} , Set of entities \mathcal{E}_Q , Number of reranked candidate diseases $topn$

Ensure: Reranked candidate diseases \mathcal{D}_{rerank}

- 1: Initialize an empty list scores
- 2: **for** each disease $\mathcal{D}_i \in \mathcal{D}_{can}$ **do**
- 3: Initialize score $\leftarrow 0$
- 4: **for** each entity $e_j \in \mathcal{E}_G$ **do**
- 5: Compute the shortest path $\text{dist}(\mathcal{D}_i, e_j)$
- 6: **if** $\text{dist}(\mathcal{D}_i, e_j) = \infty$ **then**
- 7: score \leftarrow score + 0
- 8: **else**
- 9: score \leftarrow score + $\frac{1}{\text{dist}(\mathcal{D}_i, e_j)}$
- 10: **end if**
- 11: **end for**
- 12: Append $(\mathcal{D}_i, \text{score})$ to scores
- 13: **end for**
- 14: Sort scores by the second element (score) in descending order
- 15: $\mathcal{D}_{rerank} \leftarrow$ Select the first $topn$ elements from scores
- 16: **return** \mathcal{D}_{rerank}

Department	Num	Avg Len
Gynaecology	411	627.46
Otolaryngology	212	967.99
Obstetrics&Gynecology	1316	489.15
Nursing	52	584.88
Emergency	87	552.96
Psychiatry	127	867.66
Rehabilitation	284	631.13
Dentistry	130	342.56
Anesthesiology	232	634.25
Internal Medicine	3590	528.72
Dermatology	286	518.08
Neurosurgery	3152	531.82
Ophthalmologic	100	453.24
Oncology	471	855.66
Total	10450	558.60

Table 5: Departments distribution of the collected EMRs. "Num" denotes the total number of EMRs of the department. "Avg Len" denotes the average number of words per record.

Retriever	R	P	F1
bm25	40.37	29.86	34.32
tf-idf	40.25	29.68	34.16
m3e	41.95	32.63	36.70
all-mpnet	42.01	32.75	36.80
bge	42.20	32.81	36.91
corom	42.16	32.86	36.93
bge + bm25	41.62	30.57	35.24
corom + bm25	41.75	30.46	35.22

Table 6: Performances of medIKAL using different retrieval methods during entity-node matching on CMEMR dataset.

(ite), microorganisms (mic), medical procedures (pro), and clinical symptoms (sym).

C.2 Retrieval Method

In entity-node matching process mentioned in section 3.2.1, we used a dense retrieval method to link EMR’s entities to KG’s nodes. In order to better explore the appropriate retrieval method, we implemented three types of retrieval methods based on the retriv library²: sparse retrieval, dense retrieval, and hybrid retrieval.

- Sparse Retrieval: We evaluated two representative methods, namely bm25 and tf-idf.

² <https://github.com/AmenRa/retriv>

- Dense Retrieval: We evaluated several representative embedding models, namely m3e-large (Wang Yuxin, 2023), all-mpnet-base-v2, bge-large-zh-v1.5, and CoROM.
- Hybrid Retrieval: We evaluated two combinations: "bge + bm25" and "corom + bm25".

The results are shown in Table 6. As we expected, the effect of dense retrieval is better than that of sparse retrieval and hybrid retrieval, because when the entity to be retrieved contains a large number of Chinese characters, sparse retrieval methods are very prone to mis-matching due to the lack of consideration of word order and semantics. According to the results, we choose the CoROM model as embedding model of the dense retrieval process.

The CoROM Chinese-medical text representation model we use in this paper is also released by Tongyi-Laboratory. It employs the classic dual-encoder text representation model and is trained on medical domain data with Multi-CPR (Long et al., 2022). The training process is divided into two stages – in the first stage, negative sample data is randomly sampled from the official document set, and in the second stage, difficult negative samples are mined via Dense Retrieval to augment the training data for retraining.

C.3 The Number of Candidate Diseases Set

To explore the influence of the number of candidate diseases $Top-k$ on medIKAL’s performance, we conduct experiments under settings with $Top-k$ ranging in [1, 2, 3, 5]. The results are shown in Table 7. According to the results, the Recall gradually decreases with the increase of $Top-k$, while the Precision increases. When the $Top-k$ is set very large or very small, although it can get a higher recall or precision rate accordingly, but from the practical clinical application scenario, too large or too small $Top-k$ is not conducive to assisting doctors in clinical diagnosis and decision-making. Therefore, in this paper we set $Top-k$ to 3 on CMEMR dataset, and 2 on CMB-Clin, GMD and CMD datasets.

C.4 Detailed Settings about Knowledge Graph

The knowledge graph we use in this paper is CPubMedKG-v1(Large-scale Chinese Open Medical Knowledge Graph)³ developed by Harbin Institute of Technology (Shenzhen). It is currently

³ <https://cpubmed.openi.org.cn/graph/wiki>

<i>Top-k</i>	R	P	F1
1	27.27	56.74	36.83
2	34.15	41.21	37.34
3	42.16	32.86	36.93
5	49.42	24.27	32.55
10	60.85	13.92	22.74

Table 7: Performances of medIKAL with different numbers of candidate diseases (denoted as *Top-k*) on CMEMR dataset.

the largest fully open Chinese medical knowledge graph in China. The knowledge is derived from over 2 million high-quality Chinese core medical journals under the umbrella of the Chinese Medical Association. It is regularly updated and conforms to mainstream Chinese medical standards in terms of entity and relationship specifications. The sources of entities and relationships are clearly defined, traceable, and easily distinguishable. The graph contains a total of 4,383,910 disease-centered triples. It includes 523,052 disease entities, 188,667 drug entities, 145,908 symptom entities, and a total of 1,728,670 entities. There are more than 40 types of relationships covering drug treatment, complications, laboratory tests, indications, risk factors, affected populations, mortality rates, and more. The total number of structured knowledge triples reaches 3.9 million.

For the entity type weights, we obtain the entity type weight allocation scores through the following two methods:

- We extract paragraphs related to diagnosis from the medical textbooks provided by (Jin et al., 2021). Specific example can be found in Table 8-(1).
- We selected 500 medical records with detailed diagnostic evidence from our collection and collected all diagnostic evidence. Specific example can be found in Table 8-(2).

We calculate the entity type proportions of all the segments above, obtaining initial entity type weights. We then fine-tune on randomly sampled medical record samples, the setting in our experiments can be found in Table 9. It is important to note that entity type weights are not fixed and can be adjusted according to different tasks, which is also the advantage of the method we propose.

For the shortest path algorithm in path-based

(1) Example:

[Diagnosis]: History of vitamin D overdose. Early elevation of blood calcium > 3 mmol/L (12 mg/dl), strong positive urinary calcium (Sulkowitch reaction), routine urinalysis shows positive urinary proteins, and in severe cases, red blood cells, leukocytes, and tubular patterns are seen.

(2) Example:

[Diagnostic Evidence]: 1.history of prior radiotherapy for esophageal cancer, long history of hypertension, history of smoking. 2.left limb weakness for 1 day. 3.Examination revealed hypertension, decreased muscle strength of the left limb, and decreased tenderness. 4.Ancillary tests showed immediate elevated blood glucose, ECG T-wave abnormality, cervical vascular ultrasound and cranial CT and MRI suggestive of cerebral infarction.

Table 8: (1).A specific example of paragraphs related to diagnosis from the medical textbooks provided by (Jin et al., 2021). (2).A specific example of diagnostic evidences in our collected EMRs.

reranking, we use GraphDataScience⁴ framework to implement it.

Type	Weight
dis	.1638
pro	.0043
sym	.6297
dru	.1391
bod	.0212
ite	.0372
equ	.0029
mic	.0009
dep	.0004

Table 9: Entity-type weight settings in our experiments.

D Evaluation Metrics Calculation

Firstly, for the disease entities in the diagnosis results \hat{D} and the reference diagnosis results \mathcal{R} in the medical records, we employed a fuzzy matching process (with a predefined threshold of 0.5) to associate these disease entities with ICD-10 terms, thus mapping \hat{D} and \mathcal{R} to two standardized disease

⁴ <https://neo4j.com/product/graph-data-science/>

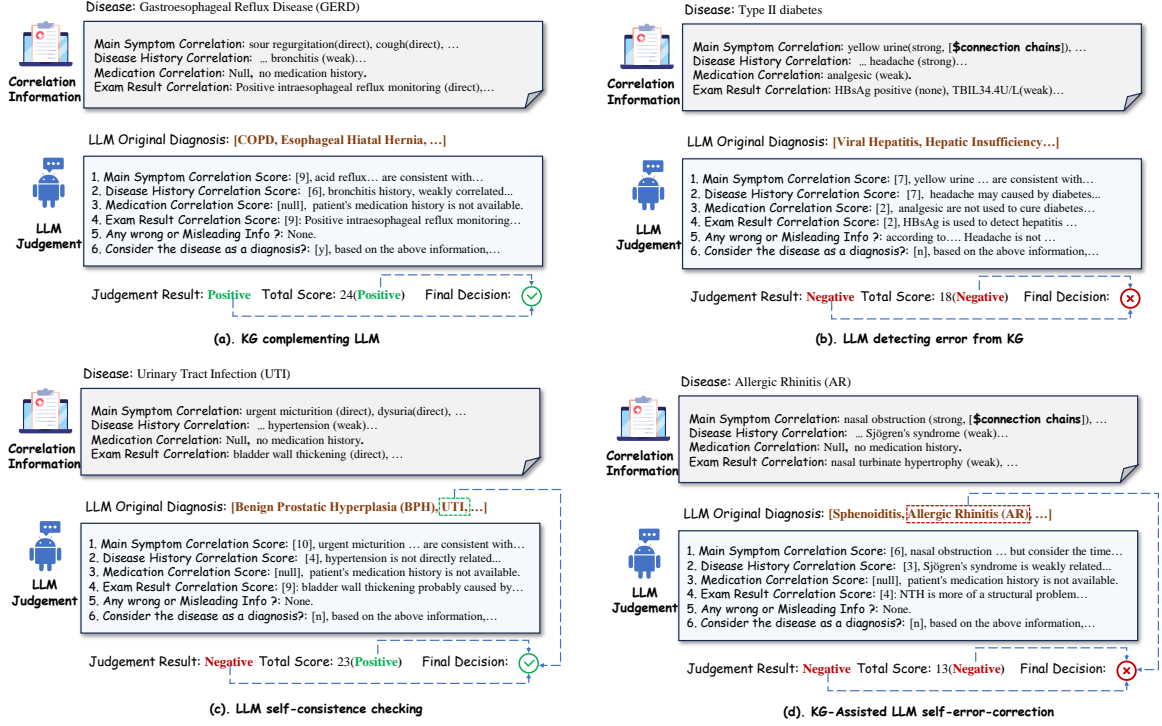


Figure 5: Case study.

sets $S_{\hat{D}}$ and $S_{\mathcal{R}}$ respectively. We then define: **True Positives (TP)**: The number of disease entities in the predicted result $S_{\hat{D}}$ that correctly match with the reference diagnosis $S_{\mathcal{R}}$.

False Positives (FP): The number of disease entities that appear in the predicted result $S_{\hat{D}}$ but do not match correctly with the reference diagnosis $S_{\mathcal{R}}$.

False Negatives (FN): The number of disease entities in the reference diagnosis $S_{\mathcal{R}}$ that do not appear in the predicted result $S_{\hat{D}}$. Based on the above statistical values, we calculate the following evaluation metrics:

$$\text{Recall (R)} : R = \frac{TP}{TP + FN} \quad (5)$$

$$\text{Precision (P)} : P = \frac{TP}{TP + FP} \quad (6)$$

$$\text{F1 Score (F1)} : F = \frac{2 \times P \times R}{P + R} \quad (7)$$

E Case Study

We show representative case studies in Figure 5 to demonstrate the effectiveness of our proposed medIKAL.

F The prompt templates used in this paper

[Role]<SYS>

You are an outstanding AI medical expert. You can summarize critical information for diagnosis based on the content of the patient's medical records.

[Role]<USR>

Below is a portion of the electronic medical record of a real patient. Please read the following content carefully to understand the patient's basic condition.

Patient Medical Record Content

""

"History of Present Illness": \${HPI}

"Past Medical History": \${PMH}

""

Task:

Based on the above content, please summarize the key information useful for diagnosis and treatment and generate a summary report.

Report Format Requirements:

Please fill in the "[]" sections according to the following format to complete the report. Use concise language whenever possible.

""

1. Main symptoms: []
2. Recent medical visits: [] (if none, write "none")
3. Past medical history: [] (if none, write "none")
4. Past surgical history: [] (if none, write "none")
5. Medication usage: [] (if none, write "none")

""

Output:

\${}

Table 10: The default prompt for the LLM Summarization module (for the patients' basic condition) .

[Role]<SYS>
You are an excellent AI medical expert. You can summarize key information useful for diagnosis based on the patient's examination results.

[Role]<USR>
Task:
Please summarize and generalize the key information useful for diagnosis based on the patient's examination results.
Example
""
[Patient's Examination Results]
"Physical Examination": Bilateral waistline symmetry, no tenderness in the bilateral ureteral regions, bladder area distended, no palpable mass, no redness or abnormal discharge at the urethral opening, no abnormalities in the scrotum, and no abnormalities in the bilateral testicles and epididymis. Digital rectal exam: Prostate approximately 4.0×5.0cm in size, soft, central area slightly shallow, small nodules palpable.
"Laboratory and Aided Examination": Ultrasound results show 1. Bilateral kidney cysts 2. Prostatic hyperplasia 3. No abnormalities in the ureters and bladder.
[Summary]
"Physical Examination": Digital rectal exam: Prostate approximately 4.0×5.0cm in size, central area slightly shallow, small nodules palpable. "Laboratory and Aided Examination": Ultrasound results show 1. Bilateral kidney cysts 2. Prostatic hyperplasia.
""
Please refer to the above example to summarize the patient's examination results.
[Patient's Examination Results]
"Physical Examination": \${PE}
"Laboratory and Aided Examination": \${LAE}
##Output:
\${}

Table 11: The default prompt for the LLM Summarization module (for the patients' exam results).

[Role]<SYS>
You are an outstanding AI medical expert. You can perform a preliminary disease diagnosis based on the patient's condition.

[Role]<USR>
##Patient Information
""
[General Condition]: \${summary_1}
[Examination Findings]: \${summary_2}
""
##Task
Based on the patient's symptoms, medical visit history, past medical history, and examination results, predict the possible diseases the patient may have (you can provide the top- $\{n\}$ possible predictions). Please only output the prediction results, do not output any other content.
##Prediction Results
Predicted Disease 1: \${} Predicted Disease 2: \${} Predicted Disease 3: \${} ...

Table 12: The default prompt for the LLM Direct Diagnose Module.

[Role]<SYS>

You are an experienced medical expert. You can evaluate the reasonableness of existing diagnostic results by considering the patient's symptoms, medical history, medication usage, and examination results.

[Role]<USR>

##Patient Information

""

[General Condition]: \${summary_1}

[Examination Findings]: \${summary_2}

""

A doctor has made a preliminary diagnosis based on the above information, with the diagnosis being: \${disease}

You need to consider whether this diagnosis is correct. To do this, you queried a medical knowledge graph and obtained the following information:

##Correlation Information

""

Correlation between diagnosis \${disease} and patient's main symptoms: \${correlation_1}

Correlation between diagnosis \${disease} and patient's medical history: \${correlation_2}

Correlation between diagnosis \${disease} and patient's medication usage: \${correlation_3}

Correlation between diagnosis \${disease} and patient's examination results: \${correlation_4}

""

##Task

Based on the patient's condition and the above information, and in combination with your own knowledge, please quantitatively evaluate the reasonableness of the diagnosis \${disease}.

##Requirements

""

1.Consistency with the patient's chief complaint score: [?] (out of 10)

2.Correlation with the patient's medical history score: [?] (out of 10)

3.Correlation with the patient's medication usage score: [?] (out of 10)

4.Correlation with the patient's examination results score: [?] (out of 10)

5.Are there any errors or misleading information in the "Correlation Information" section ?

6.Can this disease be used as a diagnostic result: [?] (y/n)

""

##Output:

\${}

Table 13: The default prompt for the LLM Diagnosis Evaluation Module.