

AI-Powered Clinical Decision Support Systems Using Retrieval-Augmented Generation: A Framework for Evidence-Based Medical Recommendations

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Abstract—Clinical Decision Support Systems (CDSS) promise to improve healthcare quality by facilitating evidence-based medicine; however, standalone Large Language Models (LLMs) suffer from hallucinations, outdated knowledge, and non-transparent reasoning. This paper presents a comprehensive framework for AI-powered CDSS that integrates Retrieval-Augmented Generation (RAG) to ground clinical recommendations in verified medical evidence. The proposed system architecture combines dense retrieval pipelines with transformer-based embeddings, curated clinical knowledge bases, and strategic hallucination reduction mechanisms. Our evaluation framework encompasses retrieval metrics (precision, recall, MRR), generation quality metrics (faithfulness, attribution, completeness), and clinical relevance measures aligned with healthcare standards. The system addresses the critical gap between generic LLM capabilities and domain-specific clinical safety requirements. Expected outcomes include 1.35x improvement in diagnostic accuracy compared to baseline LLMs, reduced hallucinations through context grounding, and transparent attribution of clinical recommendations to authoritative sources. This work contributes to bridging the gap between cutting-edge AI and trustworthy clinical deployment, with implications for guideline interpretation, diagnostic assistance, and clinical decision-making workflows.

Index Terms—Clinical Decision Support Systems, Retrieval-Augmented Generation, Large Language Models, Healthcare AI, Medical Knowledge Retrieval, Evidence-Based Medicine, Hallucination Reduction, Vector Embeddings

I. INTRODUCTION

The integration of artificial intelligence into clinical practice presents unprecedented opportunities to enhance diagnostic accuracy, reduce medical errors, and improve patient outcomes. Clinical Decision Support Systems (CDSS) have demonstrated significant potential in facilitating evidence-based medicine by providing timely, clinically relevant recommendations grounded in current medical literature and clinical guidelines [1]. However, conventional CDSS rely on manually curated rule-based systems or machine learning models trained on fixed datasets, limiting their adaptability to evolving medical knowledge and novel clinical scenarios [2].

The advent of Large Language Models (LLMs) such as GPT-4 and LLaMA has introduced powerful natural language understanding capabilities that could revolutionize clinical decision-making. Yet these models exhibit critical limitations in medical domains: their training data becomes progressively outdated, they generate plausible-sounding but factually incorrect responses (hallucinations), and they provide limited transparency regarding the sources of their recommendations [3].

Retrieval-Augmented Generation (RAG) has emerged as a promising paradigm to address these limitations by augmenting LLM capabilities with retrieval mechanisms that access external knowledge sources [4]. Rather than relying solely on parametric knowledge encoded in model weights, RAG systems retrieve relevant information from curated knowledge bases before generating responses, substantially reducing hallucinations and enabling continuous knowledge updates. Recent meta-analysis shows that RAG implementation improves clinical task performance by an odds ratio of 1.35 compared to baseline LLMs, with particular benefits in diagnostic accuracy and guideline interpretation [5].

A. Research Contributions

This paper presents a complete framework for RAG-enhanced CDSS that addresses the critical gap between cutting-edge AI and clinical deployment:

- Comprehensive system architecture integrating knowledge base development, retrieval pipelines, generation mechanisms, and hallucination reduction strategies
- Clinical-specific evaluation framework combining retrieval metrics, generation quality measures, and clinical relevance assessments
- Evidence-based methodology for clinical knowledge base construction from guidelines and literature
- Concrete hallucination mitigation strategies tailored to medical safety requirements
- Implementation roadmap with technology recommendations and scalability considerations

II. RELATED WORK AND LITERATURE REVIEW

A. Clinical Decision Support Systems

CDSS have evolved significantly since the 1970s, progressing from simple rule-based knowledge systems to sophisticated machine learning models [6]. A 2020 meta-analysis of 151 studies found that computerized CDSS improve clinical outcomes in approximately 60% of studies, with particular benefits in medication dosing, preventive care, and guideline adherence [1]. However, success depends critically on workflow integration, clinician acceptance, and knowledge base accuracy. Conventional CDSS exhibit several limitations: outdated knowledge bases, labor-intensive rule creation, challenges in capturing knowledge from unstructured literature, and poor handling of novel clinical presentations.

B. LLMs and Hallucination in Healthcare

Large Language Models demonstrate impressive performance across healthcare tasks: diagnostic reasoning, clinical note summarization, and patient education [7]. However, hallucinations—generation of plausible but unsupported information—are particularly acute in medical contexts. A systematic review identified that over 30% of LLM-generated clinical guidance lacked proper evidentiary grounding, raising serious patient safety concerns [3]. Hallucinations manifest as fabricated citations, invented drug interactions, and confident assertions about rare conditions unsupported by medical literature.

C. Retrieval-Augmented Generation

RAG augments LLMs with retrieval mechanisms accessing external knowledge sources [4]. The basic pipeline comprises three stages: indexing (documents to vector embeddings), retrieval (semantic similarity search), and generation (grounded response creation). Recent applications in medical domains include GPT-4 enhanced with medical literature for hepatology guideline interpretation, achieving 88% accuracy on challenging cases, and clinical trial patient screening reducing manual review time by 40% [5].

III. PROPOSED METHODOLOGY

A. System Architecture Overview

The RAG-CDSS system comprises five integrated modules operating within a comprehensive pipeline:

- 1) **Query Processing:** Clinical queries undergo medical entity recognition, normalization, and synonym expansion
- 2) **Retrieval Pipeline:** Hybrid dense and sparse retrieval with confidence-based filtering
- 3) **Context Management:** Reranking and evidence aggregation for LLM input
- 4) **Generation Module:** LLM-based response generation with prompt engineering and constraints
- 5) **Verification:** Faithfulness checking, hallucination detection, and confidence scoring

B. Clinical Knowledge Base Development

The knowledge base construction involves three coordinated stages:

Stage 1: Source Collection - Systematic collection of clinical practice guidelines from authoritative sources, peer-reviewed clinical literature from PubMed Central, and validated clinical references. Domain focus on selected medical specialty (cardiology, oncology, or internal medicine).

Stage 2: Document Preprocessing - Clinical text chunking respects semantic boundaries (clinical recommendations, evidence presentations, diagnostic criteria) rather than arbitrary token counts. Chunks are annotated with source type, medical domain, evidence level (Level I RCTs through Level V expert opinion), and clinical context.

Stage 3: Entity Extraction - Deployment of biomedical NLP models (BioBERT, PubMedBERT) for medical entity recognition, relationship extraction, and knowledge graph construction validated against medical ontologies (SNOMED CT, MeSH).

C. Retrieval Pipeline

The retrieval pipeline employs hybrid approach combining dense and sparse retrieval:

Dense Retrieval: Transformer-based embedding models (PubMedBERT, BioBERT) with vector database (FAISS or Pinecone). Query embedding retrieves top-k chunks based on cosine similarity.

Sparse Retrieval: BM25-based keyword retrieval complements dense retrieval for exact medical terminology matching. Hybrid fusion combines dense and sparse results using rank-based fusion strategies.

Reranking: Cross-encoder models improve relevance ordering. Confidence scoring incorporates cosine similarity, cross-encoder scores, metadata-based relevance, and clinical context matching.

D. Generation and Hallucination Reduction

The generation module includes six complementary hallucination reduction mechanisms:

- 1) **Retrieval Optimization:** Improved chunking and embedding strategies provide stronger evidence grounding
- 2) **Constrained Generation:** Structured output formats and keyword constraints reduce fabrications
- 3) **Citation Requirements:** Explicit source citation with automated verification
- 4) **Post-Generation Verification:** Faithfulness checking using entailment analysis
- 5) **Confidence Scoring:** Multi-component confidence incorporating retrieval quality and cross-reference alignment
- 6) **Human-in-the-Loop:** Clinician review before implementation with feedback loops

E. Prompt Engineering

Strategic prompt structure provides critical guidance to the LLM:

System Role: Define AI as clinical assistant constrained to evidence-based recommendations.

Context Inclusion: Patient demographics, relevant history, current presentation.

Evidence Specification: Retrieved clinical guidelines and literature chunks with source attribution.

Output Format: Structured format requiring recommended approach, evidence quality level, supporting citations, and confidence assessment.

Parameter Tuning: Temperature 0.3-0.5 reduces hallucination risk; top-p 0.9 maintains diversity while reducing unlikely outputs.

IV. EVALUATION FRAMEWORK

A. Retrieval Quality Metrics

- **Precision and Recall:** Target > 85% precision and > 80% recall
- **Mean Reciprocal Rank (MRR):** Measures ranking quality; Target MRR > 0.85
- **Normalized Discounted Cumulative Gain (NDCG):** Accounts for relevance gradations

B. Generation Quality Metrics

- **Faithfulness:** Percentage of statements supported by retrieved chunks; Clinical requirement: $\geq 90\%$
- **Attribution Accuracy:** Verification that cited sources support stated claims
- **Completeness:** Whether recommendations address all relevant aspects
- **Hallucination Detection:** False or unsupported claims; Target: < 5%

C. Clinical Relevance Metrics

- **Diagnostic Accuracy:** Concordance with gold-standard diagnoses (top-1, top-2, top-3)
- **Sensitivity and Specificity:** True positive and true negative rates
- **Guideline Concordance:** Conformance to evidence-based guidelines
- **Clinician Trust:** User perception of transparency and reliability

D. Safety and Performance Metrics

- **Adverse Event Risk:** Potential for harmful recommendations
- **Latency:** Response time acceptable for clinical workflow (< 10 seconds)
- **Cost-Efficiency:** API costs and deployment expenses

V. EXPECTED RESULTS AND DISCUSSION

A. Anticipated Performance

Based on literature precedent, we expect:

TABLE I
EXPECTED SYSTEM PERFORMANCE METRICS

Metric	Target	Baseline LLM
Dense Retrieval Precision	88-92%	N/A
Dense Retrieval Recall	82-88%	N/A
MRR	0.87-0.92	N/A
Faithfulness Score	91-96%	45-55%
Hallucination Rate	< 5%	20-30%
Attribution Accuracy	89-95%	30-40%
Diagnostic Accuracy (Top-1)	+1.35x	Baseline
Guideline Concordance	90-95%	60-70%
Sensitivity	87-93%	72-78%
Specificity	82-89%	68-75%
Response Latency	< 10 sec	2-5 sec

B. Clinical Impact

Benefits: Reduced diagnosis time in complex cases, improved guideline adherence, enhanced clinician confidence through transparent reasoning, continuous knowledge updates reflecting latest evidence.

Implementation Challenges: EHR system integration, clinician workflow disruption, privacy requirements, cross-setting validation.

C. Limitations

- Initial focus on single medical domain; generalization requires further work
- Knowledge base currency depends on systematic updates
- Performance contingent on retrieval quality
- Limited to English-language medical literature

D. Future Work

- 1) Integration with AI agents for multi-step clinical reasoning
- 2) Deep integration of knowledge graphs and medical ontologies
- 3) Seamless EHR system integration with real-time patient data
- 4) Multimodal extension incorporating medical imaging
- 5) Randomized controlled trials demonstrating patient outcome improvements

VI. IMPLEMENTATION ROADMAP

A. Technical Stack

- **Framework:** LangChain or LlamaIndex for RAG orchestration
- **Vector Database:** FAISS (on-premises) or Pinecone (cloud)
- **Embedding Model:** PubMedBERT or domain-adapted BERT variants
- **LLM Integration:** OpenAI API, Anthropic Claude, or LLaMA models
- **Evaluation:** DeepEval or Ragas frameworks

B. Development Phases

Phase 1 (Weeks 1-2): Literature review and requirements analysis.

Phase 2 (Weeks 3-4): Knowledge base development and quality assurance.

Phase 3 (Weeks 5-6): Retrieval pipeline implementation and optimization.

Phase 4 (Weeks 7-8): LLM generation module and hallucination reduction mechanisms.

Phase 5 (Weeks 9-10): Comprehensive evaluation and optimization.

Phase 6 (Week 11): Documentation, testing, and final refinement.

VII. CONCLUSION

This paper presents a comprehensive framework for AI-powered Clinical Decision Support Systems enhanced by Retrieval-Augmented Generation. By combining transformer-based embeddings, curated clinical knowledge bases, and multi-layered hallucination reduction mechanisms, the system bridges the gap between remarkable LLM capabilities and stringent clinical safety requirements. The proposed methodology addresses critical limitations of both traditional CDSS and unaugmented LLMs through continuous knowledge updates, explicit evidence grounding, and transparent source attribution. The comprehensive evaluation framework enables rigorous performance assessment before clinical deployment. Expected improvements of 1.35x in diagnostic accuracy combined with 60-80% hallucination reduction position RAG-enhanced CDSS as valuable tools for evidence-based clinical practice. Future work should focus on clinical validation studies, integration with existing workflows, and demonstration of patient outcome improvements.

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