When Raw Data Prevails: Are Large Language Model Embeddings Effective in Numerical Data Representation for Medical Machine Learning Applications?

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Abstract

 The introduction of Large Language Models (LLMs) has advanced data representation and analysis, bringing significant progress in their use for medical questions and answering. De- spite these advancements, integrating tabular data, especially numerical data pivotal in clini- cal contexts, into LLM paradigms has not been thoroughly explored. In this study, we examine the effectiveness of vector representations from 010 last hidden states of LLMs for medical diag- nostics and prognostics using electronic health record (EHR) data. We compare the perfor- mance of these embeddings with that of raw numerical EHR data when used as feature in-**puts to traditional machine learning (ML) algo-rithms that excel at tabular data learning, such** as eXtreme Gradient Boosting. We focus on instruction-tuned LLMs in a zero-shot setting to represent abnormal physiological data and evaluating their utilities as feature extractors to enhance ML classifiers for predicting diag- noses, length of stay, and mortality. Further- more, we examine prompt engineering tech- niques on zero-shot and few-shot LLM embed- dings to measure their impact comprehensively. Although findings suggest the raw data features still prevails in medical ML tasks, zero-shot LLM embeddings demonstrate competitive re- sults, suggesting a promising avenue for future research in medical applications.

031 1 Introduction

 Numerical data plays a pivotal role across var- ious domains. For instance, much of the data used for analytics from electronic health records (EHRs) are numerical values in tabular formats, documenting patient demographics (e.g., age), vi- tal signs, laboratory tests, and nurse assessments. Utilizing numerical data for predictive modeling has been instrumental in facilitating accurate diag- [n](#page-10-1)oses [\(Pang et al.,](#page-10-0) [2021\)](#page-10-0), risk stratifying [\(Zeiberg](#page-10-1) [et al.,](#page-10-1) [2019;](#page-10-1) [Green et al.,](#page-9-0) [2018\)](#page-9-0), and outcome pre-dictions [\(Akel et al.,](#page-8-0) [2021;](#page-8-0) [Chang et al.,](#page-8-1) [2019\)](#page-8-1) in

Example Question: What is the reference range and units of measurement for
Systolic Blood Pressure? (Answer: 90-120 mm Hg.) Likert Scale for Evaluation: (1) Strongly Disagree; (2) Disagree; (3) Neither Agree

Figure 1: Physician Evaluation of LLMs' Knowledge on Normal Vital Sign and Lab Test Values. This experiment probes Mistral-7B-Instruct and Llama2-13B-Chat on reference ranges for 24 vital signs and lab tests. Results show these models have a strong understanding of normal medical values, crucial for clinical applications. Table [1](#page-2-0) listed all 24 feature names, and more output examples are in Appendix [B.](#page-10-2)

healthcare. Machine learning (ML) classifiers like **043** gradient boosted [\(Chen and Guestrin,](#page-9-1) [2016\)](#page-9-1) have **044** excelled in these tasks for making accurate clini- **045** cal predictions [\(Churpek et al.,](#page-9-2) [2024;](#page-9-2) [Lolak et al.,](#page-9-3) **046** [2023;](#page-9-3) [Moore and Bell,](#page-10-3) [2022\)](#page-10-3). **047**

Recent work shows Large Language Models **048** (LLMs)' vast potential on text generation over **049** structured data input, including Chain-of-Thought **050** (CoT) reasoning over tabular data [\(Zheng et al.,](#page-10-4) **051** [2023\)](#page-10-4), classification on diseases [\(Hegselmann et al.,](#page-9-4) **052** [2023\)](#page-9-4). LLMs have also exhibited exceptional **053** promise in medical NLP tasks, evident in their stel- **054** lar performance in the United States Medical Li- **055** censing Examination (MedQA) [\(Nori et al.,](#page-10-5) [2023\)](#page-10-5). **056** However, the use of embedding representations, **057** particularly for medical diagnostics and outcome **058** predictions using standard EHR numerical data, **059** remains largely unexplored. In these areas, raw **060** data inputs have traditionally dominated feature **061** representation for ML algorithms before the era **062** of LLMs. This is exemplified by their use in **063** critical applications such as mortality prediction **064** [a](#page-9-6)nd early sepsis warnings [\(Deng et al.,](#page-9-5) [2022;](#page-9-5) [Hou](#page-9-6) **065** [et al.,](#page-9-6) [2020\)](#page-9-6), and patient infection [\(Bashiri et al.,](#page-8-2) **066**

Figure 2: This study investigates the feasibility of using LLM embeddings for numerical EHR data features representation in medical machine learning applications. To use LLMs, raw features are transformed into queries via templates. Under a zero-shot setting, these queries are encoded into embeddings for ML classification. We explore the effects of prompt engineering, few-shot learning using synthetic data generation, and parameter efficient tuning on LLM embeddings.

067 [2022;](#page-8-2) [Bhavani et al.,](#page-8-3) [2020\)](#page-8-3). The potential of LLM-**068** derived features as a viable alternative to raw data **069** features in ML applications is still unclear.

 This study aims to address this knowledge gap by examining the use of LLM embeddings for EHR numerical data representation in ML algorithms. Although LLMs are renowned for text generation, their embeddings may offer multiple advantages, such as leveraging LLMs' pre-trained knowledge and sophisticated text understanding to enhance domain-specific tasks. Moreover, using LLMs to represent tabular data allows for a unified model that encodes both structured and unstructured text in EHRs, seamlessly integrating and contextualiz- ing information across modalities, such as embed- ded tables in clinical notes [\(Soenksen et al.,](#page-10-6) [2022;](#page-10-6) [Kline et al.,](#page-9-7) [2022\)](#page-9-7).

 Our work presents novel examination of the im- pact of different formats and embedding methods on LLM last layers and ML classifiers. We focus on open-source, zero-shot LLMs suitable for single- GPU systems, considering the resource limitations prevalent in many hospitals and academic research settings. To establish a foundation for this work, we probed Mistral-7B-Instruct and Llama2-13B, two open-source, general-domain LLMs, for their knowledge of reference ranges for vital signs and lab test values. We directly asked about the stan- dard physiological values and units of measurement for 24 EHR features identified as critical predictor [v](#page-8-0)ariables for detecting clinical deterioration [\(Akel](#page-8-0) [et al.,](#page-8-0) [2021\)](#page-8-0). As in Figure [1,](#page-0-0) physician judgment indicates that LLMs possess this knowledge, pro-viding initial evidence for further investigation.

Our study utilizes three clinical prediction tasks **101** derived from two independent EHRs and four ML **102** classifier input settings. We investigate the im- **103** pact of table-to-text conversion formats, embed- **104** ding extraction methods, prompt engineering, and **105** few-shot techniques, along with early results from **106** parameter-efficient fine-tuning, on the quality of **107** LLM embeddings. Our main contributions are **108** threefold: **109**

- We present a comprehensive study exploring var- **110** ious factors that influence the performance of **111** numerical EHR feature embeddings generated by **112** LLMs for medical ML applications. **113**
- Our findings show that while LLM embeddings **114** paired with XGB classifiers can achieve perfor- **115** mance comparable to traditional raw data fea- **116** tures on some tasks, performance gaps persist, **117** necessitating further improvements to maximize **118** their effectiveness. **119**
- We discuss the efficiency and robustness of LLM **120** feature representation for numerical data versus **121** raw data in training ML classifiers. **122**

Results show that, despite external evidence indi- **123** cating that LLMs possess extensive knowledge of **124** medical facts, extracting usable representations of **125** this knowledge for downstream tasks will require **126** significant additional methodological progress. **127**

2 Related Work **¹²⁸**

Recent studies highlight LLMs in tabular data **129** analysis: [Hegselmann et al.](#page-9-4) [\(2023\)](#page-9-4) introduces **130** TableLLM, which converts tables to text using a **131** manual template. [Zheng et al.](#page-10-4) [\(2023\)](#page-10-4) studies CoT **132** reasoning over tables. [Akhtar et al.](#page-8-4) [\(2023\)](#page-8-4) ex- **133**

 amines the abilities of LLMs on numerical data understanding. [Zhu et al.](#page-10-7) [\(2024\)](#page-10-7), closest to our work, explores zero-shot LLM for structured longi- tudinal EHR data and finds that GPT-4 can outper- form XGB on clinical prediction tasks. Our study, however, uniquely focuses on open-box LLM em-beddings for enhancing ML algorithms.

 Raw EHR data are commonly used in medical ML applications, as found by a survey on medi- cal ML research [\(Si et al.,](#page-10-8) [2021\)](#page-10-8). They noted that labs and vital signs as frequent data types for pa- tient representation learning. [Churpek et al.](#page-9-2) [\(2024\)](#page-9-2) introduces an XGB algorithm predicting clinical de- teriorations using EHR features like demographics and lab values. [Wang et al.](#page-10-9) [\(2020\)](#page-10-9) used 104 clini- cal EHR features across various ML algorithms to establish baselines for clinical tasks such as mor- tality predictions. Our work uses the same dataset and tasks as [\(Wang et al.,](#page-10-9) [2020\)](#page-10-9) to compare LLM embeddings against traditional ML classifier out-comes on the same raw data feature baseline.

¹⁵⁵ 3 Datasets and Tasks

156 3.1 Diagnosis prediction for clinical **157** deterioration

 Early warning systems often use rule-based and ML algorithms to identify patients at risk of deteriora- [t](#page-9-8)ion or death without providing diagnoses [\(Churpek](#page-9-8) [et al.,](#page-9-8) [2014;](#page-9-8) [Kipnis et al.,](#page-9-9) [2016\)](#page-9-9). To address this, experts from multiple hospitals created a dataset that labels the diagnoses for patients who had a clin- ical deterioration event during their hospitalization. These expert-annotated diagnoses were performed with a full review of the EHR and served as the la- bels for our training data. Twenty-four tabular data features including demographics, vital signs, labs, interventions, and nursing assessments were ex- tracted from the structured EHR (eg. tabular data). They were previously identified as critical variables for clinical deterioration [\(Akel et al.,](#page-8-0) [2021\)](#page-8-0). The final datasets encompassed EHR data from 660 adult patients in medical-surgical ward within a U.S. health system. The primary diagnoses were Sepsis, Arrhythmia (Arrhy.), and Congestive Heart Failure (CHF) volume overload, with prevalence rates of 43.18% for Sepsis, 15.30% for Arrhyth- mia, and 11.82% for CHF, respectively. We used 5-fold validation on all 660 samples to generate five distinct test sets. $¹$ $¹$ $¹$ </sup>

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Table 1: Raw clinical data features from the EHR for diagnosis prediction task.

Table [1](#page-2-0) outlines the structured input features **182** from the cohort EHR dataset and target diagnoses **183** utilized in our analysis. The input features com- **184** prised a comprehensive set of clinical data points **185** including demographic information like age, vital **186** signs such as Systolic and Diastolic Blood Pres- **187** sure, and body Temperature, as well as a range of **188** serum laboratory tests including electrolytes, liver **189** function panel, renal function, red blood counts, **190** etc. These inputs served as predictors and are rele- **191** vant findings in making diagnoses like Sepsis, Ar- **192** rhythmia, and CHF. Despite its smaller sample size, **193** this EHR dataset includes physicians' manual chart **194** reviews and carefully curated data, providing accu- **195** rate annotations for patient diagnoses. **196**

3.2 Mortality and length-of-stay prediction **197**

The MIMIC-III dataset, derived from the EHR of **198** the Critical Care Units (ICU) at Beth Israel Dea- **199** coness Medical Center, has been utilized exten- **200** [s](#page-10-9)ively in research [\(Johnson et al.,](#page-9-10) [2016\)](#page-9-10). [Wang](#page-10-9) 201 [et al.](#page-10-9) [\(2020\)](#page-10-9) further developed an open-source **202** pipeline for extracting, preprocessing, and repre- **203** senting data from the MIMIC-III database, namely 204 MIMIC-Extract. This pipeline aggregates various **205** data types, such as tabular demographic data avail- **206** able at admission, vital signs with repeated mea- **207** sures, laboratory test results, time-varying inter- **208** vention signals, and prediction labels needed for **209** clinical tasks. MIMIC-Extract introduces two clini- **210** cal prediction tasks: mortality and length-of-stay **211** (LOS) predictions. The mortality prediction task **212** uses tabular data from the first 24-hour window of **213** a patient's ICU stay to predict mortality as a binary **214** classification task. The LOS prediction task, in **215** contrast, determines whether a patient's stay will **216** exceed three (LOS 3) or seven days (LOS 7) based **217** on the same 24-hour data period. Importantly, to **218** avoid competing risk outcomes between death and **219**

¹The dataset used in this study has been detailed in a clinical journal article currently under review, with a preprint also

available. To maintain the anonymity of this paper, references to the journal preprint were omitted. Details about the demographic characteristics of the patients, including gender, age, and race, are included in Appendix.

Diagnosis dataset Hospitalized patient of age *[value]* getting worse has labs and vitals values of systolic blood pressure *[value]* mmHg, diastolic blood pressure *[value]* mmHg, oxygen saturation*[value]* %, body temperature *[value]* celsius degree, ... total protein *[value]*, white blood cell *[value]*. What are the diagnoses for this patient?

MIMIC-Extract Hospitalized patient with lab and vital signs available: in the past 24 hours, the observed alanine aminotransferas values are [*list of unique values sorted by temporal order*], albumin values are [*list of unique values sorted by temporal order*], anion gap values are [*list of unique values sorted by temporal order*]...Predict if the patient mortality will occur in-hospital.

Table 2: The template for NARRATIVE serialization method for diagnosis prediction dataset (top) and MIMIC-Extract dataset (bottom).

220 LOS, patients who died within the 3- or 7-day LOS **221** window were excluded from the LOS prediction.

 We adopted the same data partitioning used in [\(Wang et al.,](#page-10-9) [2020\)](#page-10-9), comprising 16,700, 2,394, and 4,790 patient records for the training, develop- ment, and testing sets. Each patient record includes 104 time-varying tabular data features. More de- tailed demographic information can be found in the MIMIC-Extract study [\(Wang et al.,](#page-10-9) [2020\)](#page-10-9). The labels in the MIMIC-Extract dataset are highly skewed, with positive label distributions of 42.82% for LOS 3, 7.66% for LOS 7, 10.27% for Mort Hosp, and 7.10% for Mort ICU.

²³³ 4 Methods and Experiment Setup

 Figure [2](#page-1-0) illustrates the study overview and experi- ment setup. We began with a patient's tabular data input, represented using the Pandas DataFrame data structure (*raw data*). This raw data was converted to text using four distinct conversion methods, de- tailed in [§4.1,](#page-3-0) and LLM encoded the converted text, with the last hidden states extracted to generate em- bedding features ([§4.2\)](#page-3-1). These embeddings were subsequently used to train various ML classifiers on two datasets for binary prediction tasks.

 We started with zero-shot, off-the-shelf LLMs for experiments ([§4.3\)](#page-4-0). We then investigated the impact of prompt-engineering techniques and few- shot learning configurations on the embeddings and subsequent predictions ([§4.4\)](#page-4-1). An initial inves- tigation was also conducted to assess the effects of parameter-efficient fine-tuning on LLM embed- dings for ML tasks, focusing on two of the models **252** ([§4.5\)](#page-4-2).

253 As baselines, we included traditional ML classi-

fiers trained directly on raw tabular data inputs. To **254** benchmark the effectiveness of LLM embeddings, **255** we used randomly initialized embeddings of the **256** same size as the LLM-generated embeddings. **257**

4.1 Table-to-text conversion **258**

We employed four different methods to convert **259** EHR tables into input formats for LLMs: NAR- **260** RATIVES, JSON, HTML, and MARKDOWN. NAR- **261** RATIVES provide a continuous text description of **262** patient data, offering context and readability simi- **263** lar to clinical notes [\(Yu et al.,](#page-10-10) [2023\)](#page-10-10). JSON struc- **264** tures the data hierarchically, making it easy to parse **265** and interpret programmatically [\(Zhao et al.,](#page-10-11) [2023\)](#page-10-11). **266** HTML format leverages web-based structures to **267** present the data with tags [\(García-Ferrero et al.,](#page-9-11) **268** [2024\)](#page-9-11). MARKDOWN offers a lightweight markup **269** language that provides formatting while remaining **270** readable in plain text [\(Zhao et al.,](#page-10-11) [2023\)](#page-10-11). **271**

Table [2](#page-3-2) includes two NARRATIVES templates **272** used to format these varied clinical measurements **273** into a standardized query. These templates detail **274** the format in which data from the EHR dataset **275** are presented, integrating both laboratory results **276** and vital signs into a single descriptive snapshot of **277** a patient's current state. Each placeholder in the **278** template is populated with actual data points from **279** patient records, facilitating the transformation of **280** tabular EHR data into a format suitable for LLM **281** input, from which we then generate embeddings. **282**

The primary distinction between the templates **283** for the diagnosis prediction dataset and the MIMIC- **284** Extract dataset lies in the types of values incor- **285** porated. For diagnosis prediction, data are values **286** collected immediately before the early warning sys- **287** tem triggers for clinical deterioration. In contrast, **288** MIMIC-Extract tasks include laboratory and vital **289** signs data from the 24 hours prior to the event. We **290** extracted all unique values observed during the first **291** 24 hours of ICU admission in chronological order, **292** compiling these into a list format. If a feature has **293** no observations, it is omitted, resulting in variable **294** length sequences. **295**

4.2 Embedding extraction methods **296**

This section introduces the methods used to con- **297** vert input text to fixed-size vector for ML input. **298** We focused on the *last hidden states* of LLMs (as **299** in [\(Lu et al.,](#page-9-12) [2021\)](#page-9-12)), and employed three different **300** embedding extraction methods: Max Pooling cap- **301** tures the most salient features by taking the maxi- **302** mum value across all token embeddings for each **303**

 dimension [\(Bao et al.,](#page-8-5) [2023\)](#page-8-5); Mean Pooling com- putes the average value of the token embeddings, providing a balanced representation reflecting the overall content [\(Ram et al.,](#page-10-12) [2023\)](#page-10-12); Last Token uses the embedding of the last token as the repre- sentation, capturing the concluding context or final summary [\(Shani et al.,](#page-10-13) [2023;](#page-10-13) [Fu et al.,](#page-9-13) [2023\)](#page-9-13). We included embeddings extracted from first token as a reference point despite it is not ideal due to the nature of decoder-only models.

 Our choice of ML classifiers comprised two tree- based methods and a linear model to provide a comprehensive assessment of various predictive approaches. Specifically, we utilized eXtreme Gra- dient Boosting (XGB)[\(Chen and Guestrin,](#page-9-1) [2016\)](#page-9-1) and Random Forest (RF)[\(Breiman,](#page-8-6) [2001\)](#page-8-6) as our tree-based classifiers due to their robustness and ef- ficiency in handling diverse datasets with accuracy. Additionally, Logistic Regression with regulariza- tion (LR) as our linear model was chosen for its effectiveness in preventing overfitting via Ridge and Least Absolute Shrinkage and Selection Opera- tor regularization[\(Zou and Hastie,](#page-10-14) [2005\)](#page-10-14). Together, these classifiers form a balanced baseline setup that caters to both non-linear and linear decision boundaries in our data.

330 4.3 Selection of LLMs

 We assessed a mix of general-domain models and models trained on medical text. Three widely-used, general-domain LLMs that have been instruction-finetuned are Mistral-7B-Instruct- v0.1 [\(Jiang et al.,](#page-9-14) [2023\)](#page-9-14), Llama2-13B-chat-hf, Llama2-70B-chat-hf [\(Touvron et al.,](#page-10-15) [2023\)](#page-10-15), and Llama3-8B-instruct [\(LLaMa,](#page-9-15) [2024\)](#page-9-15). These mod- els are compatible with one Nvidia 80GB A100 GPU, making them popular choices among avail- able LLMs. For the domain specific LLM, We se- lected Meditron-7B [\(Chen et al.,](#page-9-16) [2023\)](#page-9-16), a Llama2- 7B based model continuously pretrained on medi- [c](#page-8-7)al text. We also included ClinicalBERT [\(Alsentzer](#page-8-7) [et al.,](#page-8-7) [2019\)](#page-8-7), pre-trained on MIMIC EHR text, rep- resenting encoders pre-trained on clinical text base-line compared to decoder-only LLMs.

347 4.4 Prompt design and few-shot learning

 Because the majority of LLMs we tested are instruction-tuned and require varying input formats, we utilized the chat templates to ensure proper integration of input data [\(Zheng,](#page-10-16) [2024\)](#page-10-16). In our study, the default setting involves including only the task-relevant question (shown on the right side of Fig [2\)](#page-1-0) in the system message and the converted **354** EHR data in the user input, without additional **355** system instructions, predefined personas, or other **356** context. Given that instruction-tuned LLMs are **357** known to be sensitive to system instructions, we **358** designed four system instructions that vary by per- **359** sona (medical professional, AI system), tasks (assess pa- **360** tients, generate embeddings for ML classifiers), thinking **361** style (chain-of-thoughts), and question type (general as- **362** sessment, binary question), enabling us to explore the **363** influence of prompt characteristics on the embed- **364** dings. All prompts were paraphrased for better **365** perplexity scores, following prompt optimization **366** strategies [\(Gonen et al.,](#page-9-17) [2023;](#page-9-17) [Lu et al.,](#page-9-18) [2023\)](#page-9-18). **367**

Two few-shot settings were explored besides **368** zero-shot prompt engineering. We generated syn- **369** thetic data for diagnosis prediction, by prompting **370** GPT-4 to generate values based on the attribute **371** names in Table [1.](#page-2-0) For each target diagnosis, GPT-4 **372** generated one example confirming the diagnosis **373** (positive) and one example negating it (negative). **374** Moreover, GPT-4 was asked to generate CoT ex- **375** planations identifying abnormal values and their **376** clinical significance. An expert physician and clini- **377** cal informaticist reviewed these synthetic data pairs **378** for quality assurance. The complete set of prompts **379** are presented in Table [8.](#page-12-0) **380**

4.5 Parameter efficient fine-tuning **381**

While our paper primarily focuses on evaluating **382** zero-shot LLMs for numerical feature representa- **383** tion, we included parameter-efficient fine-tuning **384** experiment to suggest future directions for improve- **385** ment. We employed QLoRA [\(Dettmers et al.,](#page-9-19) [2024\)](#page-9-19) **386** on Mistral-7B-Instruct and Llama3-8B-Instruct, us- **387** ing the MIMIC-Extract dataset due to its larger **388** training set compared to the diagnosis dataset. We **389** trained Mistral with a sequence classification head **390** on top, saving checkpoints with the lowest valida- **391** tion loss. Based on validation performance, we **392** optimized the (q, k, v, o) layers with $r = 16$, a 393 learning rate of 3e-5, and a LoRA dropout of 0.1. **394** Each model was trained for 3 epochs with early **395** stopping to prevent overfitting. **396**

4.6 Experiment setup 397

We used a 5-fold cross-validation on the diagnosis **398** dataset (660 patient records), resulting in 528 pa- **399** tients for training and 132 for testing per fold. For **400** mortality and LOS prediction tasks from MIMIC- **401** [E](#page-10-9)xtract data, we followed the data split from [\(Wang](#page-10-9) 402 [et al.,](#page-10-9) [2020\)](#page-10-9). We evaluated performance using Area **403**

Model Sepsis AUROC		Arrhythmia AUROC	CHF AUROC	Average	
	$(95\% \text{ CI})$	$(95\% \text{ CI})$	$(95\% \text{ CI})$	$(95\% \text{ C}I)$	
Raw Data Features Baseline					
LogisticRegression	74.40 (69.35, 79.56) 71.10 (67.01, 75.18)		54.79 (47.74, 61.79)	66.76 (61.37, 72.18)	
RandomForest	65.26 (61.79, 68.48)	53.07 (50.58, 55.80)	50.89 (49.01, 53.43)	56.41 (53.79, 59.24)	
XGB	71.17 (67.06, 75.11)	76.49 (71.32, 84.13)	58.47 (51.36, 65.15)	68.71 (63.25, 74.80)	
LLM embedding $+ XGB$ classifier					
Random	54.01 (49.89,58.44)	49.65(44.02,54.62)	50.02 (47.13, 52.29)	51.22 (47.01, 55.19)	
Mistral-7b-Instruct _{hest}	71.12 (67.54, 74.92)	68.00 (61.52, 73.93)	51.80 (44.48, 58.65)	63.40 (57.73, 68.77)	
Llama3-8b-Instruct $_{best}$	63.84 (57.31, 69.87)	71.08 (65.69, 75.87)	63.84 (56.77, 70.37)	66.25 (60.15,72.35)	
Llama2-13 b_{best}	66.02 (61.64, 70.32)	58.62 (52.62, 64.46)	49.69 (48.83, 62.58)	58.11 (54.36, 65.79)	
Llama2-70b-chat $_{best}$	68.57 (63.88, 71.53)	69.15 (67.08, 71.17)	53.87 (49.83, 58.52)	63.86 (60.93, 67.07)	
$Meditron_{best}$	66.74 (62.30, 66.15)	72.26 (65.28, 77.43)	58.11 (50.64, 64.48)	63.90 (58.28, 65.45)	
ClinicalBERT	58.80 (54.44, 63.04)	64.91 (61.84, 70.27)	49.67 (41.94, 57.51)	57.79 (52.74, 63.11)	
LLM embedding + Logistic Regression classifier					
Random	49.58 (47.68, 51.12)	49.22 (48.09, 50.43)	49.36 (47.12 51.06)	49.39 (47.63, 50.87)	
Mistral-7b-Instruct _{best}	62.61 (58.17, 66.95)	69.59 (64.67, 74.71)	48.98 (42.96,55.62)	60.39 (55.27, 65.76)	
$Llama3-8b-Instruct_{best}$	66.54 (62.32, 70.62)	70.22 (64.82, 74.11)	63.52 (55.91,69.20)	66.76 (61.50, 72.02)	
Llama2-13b-chat- hf_{best}	66.95 (62.82, 70.88) 66.04 (60.04, 71.22)		58.54 (52.09, 65.09)	63.84 (58.32, 69.06)	
Llama2-70b-chat- hf_{best}	69.50 (65.37, 73.43) 68.11 (61.75, 70.57)		62.72 (56.47, 68.39)	66.78 (61.20, 70.80)	
$Meditron_{best}$	66.91 (62.83, 71.09)	68.61 (63.49, 73.72)	57.60 (51.02, 63.89)	64.37 (59.11, 69.90)	
ClinicalBERT	47.28 (43.07, 51.63)	44.62 (38.79, 50.29)	46.98 (42.96, 55.62)	46.29 (41.61, 52.51)	

Table 4: Comparing raw data features and LLM embeddings features for ML classifiers on Diagnosis dataset. We report the best AUROC scores from LLM embedding across various embedding extraction and table-to-text conversion methods. The "Random" row indicates the randomly initialized embedding input. For ClinialBERT, we used [CLS] token embedding as the final representation. We use green color to highlight the LLM+ML results where it has CI overlapping with the best results (in bold fonted text).

404 Under the Receiver Operating Characteristic (AU-**405** ROC) with 95% confidence intervals (CI).

 For all ML classifiers, we determined the best pa- rameters through grid search on the validation set. Specifically, we tuned the number of estimators, maximum depth, learning rate, and minimum child weight for XGB classifiers, and alpha and L1 ra- tio for LR classifiers (see Appendix [E\)](#page-12-1). For LLMs under 13B, the maximum input length was 1042 for the diagnosis dataset and 3076 for the MIMIC dataset, resulting in a 4096-dimensional embed- ding. For 70B LLM, the max input length was 1500 and 4-bit quantization was set to avoid GPU memory errors, producing an 8192-dimensional embedding. All experiments ran on an Ubuntu server with an Nvidia 80GB A100 GPU.

⁴²⁰ 5 Results

421 5.1 Main results for diagnosis prediction

 Table [4](#page-5-0) presents AUROC scores for predicting Sep- sis, Arrhythmia, and CHF with different ML mod- els, demonstrating the effects of using LLM em- beddings compared to raw data features. XGB with raw data features stood out in the baseline, demon- strating the highest AUROC for Sepsis and Ar- rhythmia, and the highest average AUROC across all diagnoses. LR and RF, while using raw data input, showed moderately lower effectiveness.

431 For LLM embeddings with zero-shot setting, we **432** observed performance gain over a randomly initialized embedding approach into XGB with substan- **433** tial gains in all decoder LLMs. ClinicalBERT was **434** the only model, as an older pre-trained encoder, **435** that did not show notable performance gains over **436** the randomly initialized embedding model. Further, **437** Mistral embedding with XGB classifiers achieved **438** a competitive AUROC of 71.12 (vs. 71.16 of raw **439** data with XGB). Llama2-13B scored an AUROC of **440** 58.54 on CHF prediction, the best among all mod- **441** els. This demonstrates that LLM embeddings can **442** match or nearly match the performance of models **443** trained with raw data inputs. **444**

5.2 Main results for mortality prediction and **445** length-of-stay **446**

Figure 3: Accuracy (left) and AUROC (right) for in-ICU mortality (mort ICU), in-Hospital morality (mort Hosp), hospital LOS exceeding 3 days (LOS 3) and 7 days (LOS 7). The Logistic Regression (LR) and Random Forest (RF) baselines are reported from [\(Wang et al.,](#page-10-9) [2020\)](#page-10-9). The LLM results are from LLM embeddings + XGB settings. The CIs mostly overlap; for clarity in presentation, they were omitted from this figure.

Figure [3](#page-5-1) displays performance for various mod- **447** els on tasks of in-ICU mortality (mort ICU), in- **448**

 hospital mortality (mort Hosp), and hospital length- of-stay for more than 3 (LOS 3) and 7 days (LOS 7). The raw data features with XGB model con- sistently outperforms others with an AUROC of 92.02 in mort ICU and 89.83 in mort Hosp. LLM embeddings from Mistral, Llama3-8b, and Med- itron, while slightly lagging behind the raw data features with ML classifiers in the mortality tasks, performed comparably in the LOS 7 tasks. Mistral with XGB achieved accuracy of 92.34 and AU- ROC of 72.36 on LOS 7 task, showing competitive performance to XGB with raw data features, with accuracy of 92.32 and an AUROC of 76.93. The gap between LLM embeddings and raw data for mortality and LOS tasks suggests a need to improve time-varying feature representation.

465 5.3 Comparisons across different embedding **466** methods and data conversion methods

 Figure [4](#page-6-0) presents AUROC values for different embedding methods and data conversion formats across three models: Mistral, Meditron, and Llama3-8b. Max pooling achieves the highest per- formance for Mistral (64.62) and Meditron (62.54), while mean pooling is most effective for Llama3- 8b (64.69). The last token method yields moder- ate performance across all models, with AUROCs around 57, while first token embeddings result in the lowest AUROC values, indicating a less effec-tive representation for these models.

Figure 4: Comparison across different embedding methods and different format on the Diagnosis dataset. For simplicity, we used NARRATIVE and max pooling for the other analysis after this section.

 When encoding data with different formats, Mis- tral shows preference for NARRATIVES, JSON, and HTML. The MARKDOWN format generally yielded the lowest performance across the models, particu- larly for Mistral. JSON and HTML formats showed competitive performance, with JSON being slightly more effective for Meditron and Llama3-8b. No-tably, Llama3-8b exhibited the highest variability

across formats, with AUROCs ranging from 57.40 **486** (HTML) to 66.13 (NARRATIVES). **487**

5.4 Impact of prompt engineering and **488** few-shot learning **489**

We compared performance of Mistral and Llama3 **490** using different system instructions under zero-shot **491** and few-shot settings, as well as CoT examples. **492** Mistral, under 0-shot with a system instruction with **493** persona of medical professional and the task of as- **494** sessing patient condition (prompt 1 in Table [8\)](#page-12-0), 495 achieved an AUROC of 71.35 on Sepsis prediction, **496** the highest of all models. Llama3 with zero-shot **497** prompting using prompt 1 in Table [8](#page-12-0) showed re- **498** ported AUROC of 73.51 on Arrhythmia, surpass- **499** ing its counterpart at 71.08 but still below raw data **500** XGB baseline (76.49). CoT and few-shot exhibited **501** various performance and often resulted in lower **502** AUROC scores compared to Table [4.](#page-5-0) Full results **503** are provided in Appendix [C.](#page-11-0) **504**

5.5 Parameter efficient fine-tuning results **505**

Table 5: AUROC comparison before and after training LLM with QLoRA on MIMIC tasks.

Table [5](#page-6-1) presents results of Mistral and Llama3- **506** 8b under the QLoRA across all four tasks from **507** MIMIC-Extract. The performance drops are no- **508** ticeable, especially in the two mortality predictions. **509** To further understand the reason behind the perfor- **510** mance drops, we plotted the confusion matrices for 511 LOS 3 and Mort ICU, comparing Mistral's predic- **512** tions before and after QLoRA in Figure [5.](#page-7-0) For LOS **513** 3 prediction, the Mistral model with QLoRA shows **514** an increase in true negatives and a decrease in false **515** positives. However, the false negatives rises from **516** 1133 to 1473, and true positive drops from 918 to **517** 578. On the Mort ICU task, the Mistral model with **518** QLoRA correctly predicts no false positives, but **519** it fails to predict any positive cases (0 true posi- **520** tives). The performance drop can be attributed to **521** the imbalanced class distribution in the dataset, as **522** the models show a tendency to favor the majority **523** class (negative cases). During QLoRA, the LLM **524** might learn the class prevalence, biasing its rep- **525** resentation and making it challenging to correctly **526**

3 and Mort ICU tasks. Right: Mistral without QLoRA; left: Mistral after QLoRA.

527 identify the minority class.

528 5.6 LLM Embedding vs LLM Generation

 The final experiments compare the performance of LLM embeddings combined with ML classifiers against direct outputs from LLMs. This compar- ison shows that, although LLM embeddings gen- erally do not outperform raw data features, they offer a more robust and reliable solution than rely- ing on LLMs to directly answer Yes or No ques- tions. Our exploration revealed significant limita- tions in LLM generation for binary prediction tasks. For instance, Mistral frequently predicted 'Yes' for sepsis, arrhythmia, and CHF AORC, resulting in AUROC scores being 50, whereas LLM embed- dings achieved AUROCs of 71.12 for sepsis, 72.26 for arrhythmia, and 63.54 for CHF AUROC. Simi- lar patterns were observed from Llama3-8b results (Table [12\)](#page-13-0). On MIMIC-Extract tasks with highly skewed class distributions, Mistral and Llama3- 8b, when generating direct Yes/No answers, again showed reduced ability to discriminate between positive and negative cases (Table [13\)](#page-13-1). These find- ings underscore the need for embeddings, which provide a more nuanced and effective approach for clinical predictions. We refer readers to Ap-pendix [D](#page-11-1) for more details.

⁵⁵³ 6 Discussion

 To understand the discrepancy between the two data representations, we examined the training ef- fectiveness of raw data features and LLM embed- dings by controlling the training set size. Figure [6](#page-7-1) compares the performance of the raw data XGB baseline model with the Mistral and Meditron em-beddings across different training set sizes for two

Figure 6: AUROC comparison between Raw data, embeddings from Mistral and Meditron with XGB classifiers, by controlling the training set size on two MIMIC tasks.

tasks in the MIMIC dataset. The raw data XGB **561** baseline model shows a significant increase in AU- **562** ROC scores with larger training sets, achieving **563** high performance. In contrast, both the Mistral and **564** Meditron embeddings paired with XGB models **565** exhibit much smaller improvements, consistently **566** performing lower than the raw data XGB baseline. **567** This highlights the greater effectiveness of XGB **568** when learning from raw data features compared to **569** LLM embeddings for these prediction tasks. **570**

Our findings suggest that raw data features pro- **571** vide more informative input for ML models com- **572** pared to LLM-generated embeddings. While LLM **573** embeddings capture complex representations, they **574** may not be as tailored for binary medical predic- **575** tion tasks. Additionally, computing efficiency is an **576** important consideration, as LLMs require signifi- **577** cantly more GPU memory than raw data features. **578**

However, zero-shot LLM embeddings achieve **579** comparable performance in certain scenarios, high- **580** lighting their potential for rapid deployment with- **581** out extensive training. A promising direction is **582** distilling these embeddings into a smaller space **583** [w](#page-9-20)hile retaining their extensive knowledge [\(Lee](#page-9-20) 584 [et al.,](#page-9-20) [2024\)](#page-9-20). [BehnamGhader et al.](#page-8-8) [\(2024\)](#page-8-8) recently **585** proposes LLM2Vec, a method to train decoder-only **586** LLMs as text encoders with unsupervised training, **587** which merits further investigation. 588

7 Conclusion **⁵⁸⁹**

We present the first analysis of LLM embeddings **590** for numerical EHR data features in medical ML ap- **591** plications, showing the opportunity and challenges **592** of using LLM embeddings as a substitute of raw **593** data features. We hope to encourage future research **594** on improving LLM embeddings, particularly for **595** imbalanced label prediction, and advancing health **596** predictions with multi-modal data, while address- **597** ing interpretability and bias. **598**

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⁵⁹⁹ 8 Limitation

 In our study, we focused on investigating some of the most common LLMs, including Meditron, Mistral, Llama2, and Llama3. Due to GPU con- straints, some experiments, such as Qlora, were conducted on only one or two models, limiting the comprehensiveness of our analysis. We did not include black-box LLMs via API because, despite using fully de-identified data, both EHR datasets are protected under Data Use Agreement, restrict- ing us sharing with third parties. Additionally, we acknowledge that we did not explore all possible methods of prompting LLMs, which may have in- fluenced our results. Furthermore, our examination was restricted to the last layers of the LLMs, poten- tially overlooking valuable information encoded in other layers.

 Regardless of these limitations, our findings are consistent across models: zero-shot LLM embed- dings paired with machine learning classifiers gen- erally underperform compared to raw data features, though they sometimes achieve comparable perfor-**621** mance.

⁶²² 9 Ethical Statement

 Following the ACL's ethical review guidelines, our study on leveraging LLMs for medical diagnosis within EHR emphasizes ethical integrity by priori- tizing harm avoidance, privacy protection, fairness, transparency, and respect for intellectual property. While our research aims to advance medical di- agnostics through LLMs, there is a potential risk that misinterpretations of model predictions could inadvertently lead to diagnostic errors or bias in clinical decision-making. Therefore, rigorous vali- dation protocols, including expert medical review and bias detection mechanisms are needed to en- sure that model predictions are both accurate and equitable across diverse patient populations.

 We have rigorously ensured data de- identification, obtained ethical approvals, actively mitigated biases, and maintained openness in our methodologies and findings to uphold honesty and reproducibility. Our commitment extends to respecting intellectual property through proper attribution and license adherence, with the overarching goal of contributing positively to healthcare outcomes and societal well-being. This approach underscores the importance of robust, secure research practices in developing computational tools for healthcare, aligning with

our ethical responsibility to advance the field for **649** the public good. 650

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A Diagnosis Prediction Dataset Patient **⁹⁰⁵ Demographics** 906

Table 6: Patient demographic description of diagnosis prediction dataset. Note that in this work, we exclude the demographic information from ML input.

B Probing LLMs for Inherent Knowledge **⁹⁰⁷** of Normal Ranges **⁹⁰⁸**

A foundational question for using LLM embed- **909** dings for numerical data representation is whether **910** they possess inherent knowledge about the normal range of values for clinical data. To assess this, we first asked the LLMs about standard physiological ranges, measurement units, and reasoning from the feature set of tabular data outlined in Table [1.](#page-2-0) A board-certified physician assessed the LLM gen- erations using a 1 to 5 Likert scale across three dimensions: correctness of the range, accuracy of measurement units, and quality of explanations ("Reasoning"). Our probing experiments were con- ducted on Mistral and Llama2-13b, as these are general-domain LLMs that have been fine-tuned to follow instructions. This experiment was done prior to the release of Llama3 models, therefore we did not include them. We did not employ auto- mated metrics because there is no single set of ref- erence ranges for some features, as they are highly dependent on patients' ages, sex, and other demo-graphic factors.

 In our exploration of language model capabilities within the healthcare domain, we specifically probe the model's understanding of standard ranges and measurement units for EHR data features. The following table illustrates Mistral's performance on tasks designed to evaluate its ability to accurately identify and articulate the normal physiological ranges and corresponding units for 24 distinct EHR features. This assessment not only tests the model's grasp of medical terminology and concepts but also its potential application in clinical settings where precise knowledge of such parameters is crucial.

⁹⁴² C Prompt Design

 To test the impact of prompt engineering on LLM embeddings, we designed four distinct prompts, varying by persona, thinking style, and question types. Our prompt engineering strategy builds on prior work that utilizes perplexity to select opti- mal prompts [\(Gonen et al.,](#page-9-17) [2023\)](#page-9-17). Initially, we crafted meta-language prompts delineating clinical scenarios and tasks. We then employed GPT-4, a state-of-the-art LLM, to generate five paraphrases for each prompt to capture a range of expressions. We evaluated each paraphrase's naturalness and fluency by calculating its perplexity score, select- ing the most coherent paraphrase for each scenario for our experiments. The optimized prompts are detailed in Table [8.](#page-12-0)

958 We also used these prompts in few-shot experi-**959** ments. Given the complexity of the MIMIC-Extract **960** dataset, which includes 104 features with data colQuestion: What is the reference range for the Pulse Pressure Index?

Reference: 0.1, 1

LLM (Mistral): The reference range for Pulse Pressure Index (PPI) is not a commonly used vital sign or lab result label. It measures the ratio of the pulse pressure to the systolic blood pressure, used to assess cardiovascular risk. The PPI is typically reported as a percentage, with a normal range being less than 10%. However, the specific reference range may vary by population and clinical context. It is important to consult healthcare professionals for personalized PPI interpretations.

Physician Evaluation (Range, Unit, Reasoning): 3, 5, 5

Table 7: Mistral's probing output examples for Normal Ranges and Units of EHR data features

lected over 24 hours, creating high-quality syn- **961** thetic data is challenging. Thus, we focused our **962** efforts on the diagnosis prediction use case, which **963** involves simpler inputs of twenty-four clinical fea- **964** tures, facilitating quality control over the synthetic **965** data generation. **966**

As illustrated in Figure [2,](#page-1-0) we used GPT-4 to cre- **967** ate synthetic data depicting patient cases of clinical **968** deterioration with features in Table [1.](#page-2-0) This data set **969** includes both positive and negative diagnosis cases, **970** which were reviewed by an expert physician and **971** clinical informaticist for quality assurance. Our **972** few-shot experiments varied in complexity: the **973** first modified Prompt 1 to include example input- **974** output pairs ("Simple" few-shot setting), while the **975** second added a CoT explanation detailing the di- **976** agnostic reasoning into Prompt 3. The CoT was **977** structured to identify and reason over abnormal **978** values to conclude diagnoses, enhancing the data's **979** interpretability and educational value. **980**

D Results of LLM Direct Generation **⁹⁸¹**

We tested the ability of Mistral and Llama3 to di- **982** rectly predict Yes or No answers to questions from **983** the Diagnosis and MIMIC-Extract datasets. To **984** achieve this, we added specific instructions direct- **985** ing the LLMs to respond only with "Yes" or "No," **986** then parsed the outputs to 1 or 0 labels and com- **987** puted AUROC and Accuracy. For this experiment, **988** we set the maximum token limit to 25 and the top **989**

Prompt Description

Prompt 1 -Persona: Medical Professional As a healthcare provider, please assess the patient's condition provided below and outline the likely causes or diagnoses for their clinical worsening. List only the diagnoses and keep your response brief.

Prompt 2 -Persona: AI System You are an AI with medical expertise. Create an embedding for the probable problems or diagnoses that are causing clinical deterioration, based on the patient's condition detailed below, to aid in training a diagnostic prediction machine learning classifier. Be brief in your description.

Prompt 3 -Persona: Medical Professional (Chain-of-Thought) As a medical expert, please examine the patient's condition by first identifying any abnormal values. Next, critically analyze these values to assess their impact, and clearly state your final diagnosis regarding what might be causing the clinical deterioration. Keep your summary brief.

Prompt 4 -Persona: Medical Professional (Binary Question) You are a medical doctor. Based on the patient's condition, determine the likelihood that diagnosis X is causing their clinical deterioration. Be aware that diagnosis \bar{X} occurs in Y% of similar cases.

Model	Setting	AUROC CI $(\%)$
Mistral-7b-instruct	sys1	54.85 [48.18, 62.11]
	sys2	53.88 [47.37, 60.67]
	sys3	51.16 [44.34, 57.67]
	sys4	54.04 [46.84, 61.56]
	Fewshot	54.43 [46.62, 61.05]
	CoT	57.96 [60.72, 69.24]
Llama2-13b-chat	sys 1	56.49 [49.90, 63.09]
	sys2	55.61 [48.43, 62.31]
	sys3	50.41 [43.19, 57.33]
	sys4	60.24 [53.28, 67.09]
	Fewshot	53.12 [46.38, 59.84]
	CoT	54.10 [51.84, 60.59]
Llama3-8b-instruct	sys 1	52.81 [46.47, 59.21]
	sys2	51.11 [44.64, 57.27]
	sys3	49.03 [42.19, 55.74]
	sys4	55.23 [48.28, 61.79]
	Fewshot	53.24 [46.60, 59.99]
	CoT	51.44 [48.58, 53.77]

Table 8: System prompts for medical diagnosis assistance with different persona settings.

Table 9: AUROCs for various models and settings on CHF Volume Overload prediction.

990 k to 50.

 Table [12](#page-13-0) presents results of Mistral directly gen- erating "Yes/No" answers for the Diagnosis dataset. For all tasks (Sepsis, Arrhythmia, CHF), Mistral achieved an AUROC of 50.00, indicating no dis- criminatory ability. Accuracy varied across tasks, with Sepsis at 43.18%, Arrhythmia at 15.30%, and CHF at 11.82%, corresponding to the positive class distribution, demonstrating poor performance in direct prediction. Llama3 exihibited similar perfor- mance: it reported AUROC scores between 47.12 (Arrhythmia) to 50.28 (Sepsis), underperforming its embedding counterparts reported in Table [4.](#page-5-0) Ad- mittedly, extra effort in prompt engineering and pa- rameter searching could improve direct generation results. However, compared to their embedding + ML classifier counterparts in the same zero-shot setting and input format (NARRATIVES, without additional system instructions such as personas), their performance is significantly lower.

Table 10: One Time AUROC and Confidence Intervals for various models and settings on Arrhythmia prediction. Scores are multiplied by 100.

On the MIMIC-Extract tasks, table [13](#page-13-1) shows **1010** the results of Mistral-7b-Instruct and Llama3-8b- **1011** Instruct in directly generating "Yes/No" answers **1012** for various tasks in the MIMIC-Extract dataset. **1013** Both models demonstrated no discriminatory abil- **1014** ity, with AUROC scores close to 50 for all tasks. **1015** Accuracy varied, with notable high accuracy for 1016 MORT ICU and MORT HOSP tasks, particularly **1017** for Llama3-8b-Instruct (92.88% and 89.71%, re- **1018** spectively). However, these high accuracy scores **1019** likely reflect class imbalance rather than model 1020 performance. The contrast between the LLM di- **1021** rect prediction performance and LLM embedding **1022** + classifier performance further suggests that LLM **1023** embeddings provide a more robust method. **1024**

E Parameter Grids for ML Classifiers **¹⁰²⁵**

We conducted a comprehensive grid search **1026** for hyperparameter optimization on two classi- **1027** fiers: XGBoost (XGB) and Logistic Regression. **1028** For the XGB classifier, the parameter grid in-
1029

Model	Setting	AUROC CI $(\%)$
Mistral-7b-instruct	sys 1	71.35 [67.39, 75.73]
	sys2	67.63 [63.53, 71.80]
	sys3	65.67 [61.50, 69.78]
	sys4	67.87 [63.29, 71.88]
	Fewshot	67.32 [63.26, 71.20]
	CoT	64.29 [60.72, 69.24]
Llama2-13b-chat	sys1	68.79 [64.87, 72.59]
	sys2	69.82 [65.66, 73.92]
	sys3	68.92 [64.64, 73.09]
	sys4	64.62 [60.40, 68.73]
	Fewshot	66.49 [62.21, 70.59]
	CoT	65.13 [62.30, 69.25]
Llama3-8b-instruct	sys 1	67.05 [62.96, 71.11]
	sys2	66.07 [61.94, 70.28]
	sys3	64.80 [60.74, 69.11]
	sys4	66.81 [63.08, 70.90]
	Fewshot	66.87 [62.90, 70.84]
	CoT	62.12 [58.37, 66.96]

Table 11: AUROC Confidence Intervals for various models and settings on Sepsis prediction. Scores are multiplied by 100.

Model	Task	AUROC	Accuracy
Mistral-7b-instruct	Sepsis	50.00	43.18
	Arrythmia	50.00	15.30
	CHF	50.00	11.82
Llama3-8b-instruct	Sepsis	50.28	54.69
	Arrythmia	47.12	73.63
	CHF	47.61	77.12

Table 12: Results of Mistral and Llama3-8B directly generating "Yes/No" to the Diagnosis dataset. To align with the results reported for emebdding+ML classifiers settings, the LLMs are zero-shot, and no additional system instructions are included in the chat template.

Model	Task	AUROC	Accuracy
Mistral-7b-Inst	LOS 3	49.72	56.05
	LOS ₇	49.86	87.93
	MORT ICU	50.04	91.98
	MORT HOSP	49.79	86.93
Llama3-8b-Inst	LOS 3	50.73	57.24
	LOS ₇	50.10	92.29
	Mort ICU	49.99	92.88
	Mort Hosp	49.99	89.71

Table 13: Results of Mistral and Meditron direct generation of "Yes/No" to the MIMIC-Extract dataset

 cluded nestimators set to [50, 100, 250, 500], maxdepth ranging from [2, 5, 10, 15, 20], learningrate values of [0.005, 0.01, 0.05, 0.1], and minchildweight values of [1, 2, 3]. This ex- tensive search aimed to identify the best combina- tion of hyperparameters to enhance model perfor-**1036** mance.

1037 For the Logistic Regression classifier, we varied

Values
50, 100, 250, 500
2, 5, 10, 15, 20
0.005, 0.01, 0.05, 0.1
1, 2, 3

Table 14: Parameter grid for XGBoost (XGB) classifier.

Parameter	Values
alpha	0.1, 0.5, 1.0
ratio	0.1, 0.5, 0.9

Table 15: Parameter grid for Logistic Regression (LR).

alpha with values of $[0.1, 0.5, 1.0]$ and $11 ratio$ with 1038 values of [0.1, 0.5, 0.9]. This grid search was de- **1039** signed to fine-tune the regularization parameters to 1040 achieve optimal balance between model complexity **1041** and performance. **1042**

Grid-searching on XGB parameters took 25-40 1043 minutes on GPU. On LR, it took about 25 minutes **1044** to search for the best parameters. Training both **1045** classifiers took less than 5 minutes, even on the **1046** MIMIC-Extract dataset where there are more than **1047** 16000 samples. **1048**