
Towards Human-AI Collaboration in Healthcare: Guided Deferral Systems with Large Language Models

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Abstract

Large language models (LLMs) present a valuable technology for various applications in healthcare, but their tendency to hallucinate introduces unacceptable uncertainty in critical decision-making situations. Human-AI collaboration (HAIC) can mitigate this uncertainty by combining human and AI strengths for better outcomes. This paper presents a novel *guided deferral* system that provides intelligent guidance when AI defers cases to human decision-makers. We leverage LLMs' verbalisation capabilities and internal states to create this system, demonstrating that fine-tuning small-scale LLMs with data from large-scale LLMs greatly enhances performance while maintaining computational efficiency and data privacy. A pilot study showcases the effectiveness of our proposed deferral system.

1. Introduction

Implementing artificial intelligence (AI) in decision-sensitive fields such as healthcare involves balancing the benefits of autonomy with the risks and costs of errors. Human-AI Collaboration (HAIC) aims to find this balance, by combining human and AI efforts. One approach to HAIC is using deferral systems which allow AI to handle straightforward cases while deferring complex ones to humans.

Clinicians often make decisions using their expertise in addition to the *intelligent guidance* of colleagues, which we define as task-based recommendations and informed reasoning rooted in logic. Current deferral systems lack this guidance, isolating human decision-makers. We propose that effective deferral systems should additionally simulate providing this intelligent guidance to decision-makers. This paper explores using LLMs to achieve this.

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The main challenge in building such systems is the computational expense of LLMs. Proprietary LLMs offer advantages such as state-of-the-art performance and easy implementation without needing high-performance hardware. However, they are impractical for data-sensitive applications due to lack of internal state access and privacy concerns. Open-source LLMs can perform well with relatively large amounts of parameters, but are slow and require high-performance hardware. Smaller LLMs are faster but less effective. These issues affect all real-world LLM applications, not just deferral systems. In this paper, we propose methodology for developing efficient and accurate LLMs of which are capable of guided deferral and suitable for healthcare applications.

In summary, the contributions of this paper are:

- We propose a novel deferral system, *guided deferral*, for large language models (LLMs) in computer-aided clinical diagnosis. This system not only defers cases to human decision-makers, but also provides *intelligent guidance*. We detail its practical application in healthcare and evaluate its efficacy through a pilot study.
- We evaluate the classification and deferral performance of two distinct sources of predictions; the *verbalised* and *hidden-state* predictions. Additionally, we demonstrate how a combination of these sources leads to a significant prediction in terms of classification and deferral performance.
- We demonstrate that instruction-tuning an open-source, efficient, and small-scale LLM on the *guard-railed* generation of a large-scale version of the same LLM leads to significantly improved classification and deferral performance in this task, surpassing even that of the larger model through the use of *guardrails*. This methodology ensures data privacy, which is critical for high-stakes environments such as healthcare.

2. Related Work

Human-AI Collaboration with Large Language Models. Few studies explore the use of LLMs in HAIC. [Wiegreffe et al. \(2022\)](#) first examined LLMs for explaining classification decisions using a human-in-the-loop approach to train a filter assessing explanation quality. [Rastogi et al. \(2023\)](#)

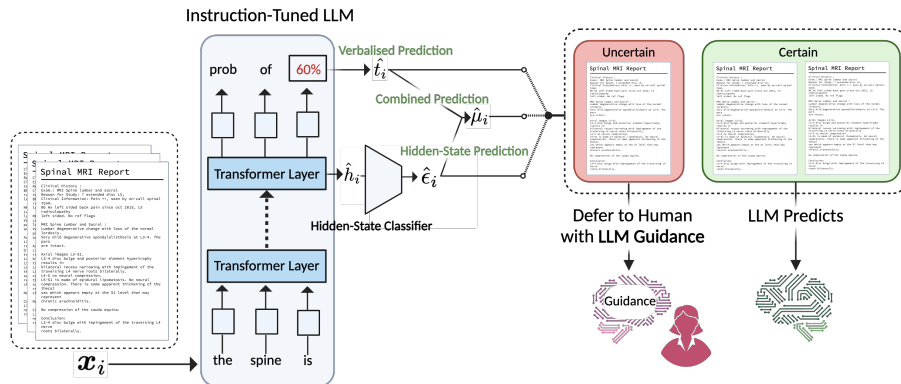


Figure 1. Our guided deferral system. Reports are parsed by an instruction-tuned LLM for clinical disorders. From the text output, we extract a verbalised prediction \hat{t} . We calculate a hidden-state $\hat{\epsilon}$ prediction from the final hidden-layer of the LLM, and its combination with \hat{t} through their mean $\hat{\mu}$. Uncertain predictions, determined by either \hat{t} , $\hat{\epsilon}$, or $\hat{\mu}$, are deferred to humans with guidance. Certain predictions are autonomously handled by the LLM.

used HAIC to audit error-prone LLMs with other LLMs. Dvijotham et al. (2023) proposed CoDoC, a deferral system with a black-box classifier and learned deferral AI for healthcare. Our system differs by using LLMs for guided deferrals. Banerjee et al. (2023) used LLMs to provide textual guidance for clinicians in decision-making on clinical imaging tasks and argued that deferral systems are sub-optimal due to anchoring bias and lack of decision-maker support. They proposed learning-to-guide, which trains LLMs to provide decision-making guidance without deferring cases. We contend that this approach still burdens decision-makers with the time and fatigue issues deferral systems address. Our work combines LLMs in deferral systems with valuable guidance for decision-makers on deferred cases, laying the foundation for advanced deferral algorithms like Learning-to-Defer (Madras et al., 2018) or Learning with Rejection (Cortes et al., 2016). Few studies (Mozannar & Sontag, 2021; Liu et al., 2021) have evaluated the efficacy of HAIC on text classification tasks but are not LLM focused.

Selective Prediction of Large Language Models. Existing research on LLMs in selective prediction includes methods to measure uncertainty in model responses after generation (Varshney & Baral, 2023). Chen et al. (2023) proposed improving selective prediction by incorporating self-assessment. Ren et al. (2023) explored detecting out-of-distribution instances in summarisation and translation tasks. Our paper applies selective prediction to deferral systems in clinical parsing. We show that combining the model’s internal state with its generated prediction enhances selective prediction without post-generation methods. We provide a comprehensive evaluation of selective prediction performance in clinical classification using real-world data, applied in deferral systems with in-distribution data.

Instruction-Tuning of Large Language Models. Finally, whilst there exists literature demonstrating improved zero-

shot performance of instruction-tuned (IT) LLMs on unseen tasks (Wei et al., 2022), our work studies this improvement specifically on an in-domain task through the use of *guardrails*. Additionally, there exists works on the use of IT in the medical domain (Liu et al., 2023), but these focus on report summarisation. No works have researched the applications of IT LLMs for the use of deferral systems. Zhang et al. (2023) provide a recent survey for additional insight in this area.

3. Methods

3.1. Sources of Predictions

Utilising LLMs in clinical parsing for disorder classification presents a unique challenge in determining the classification approach. We focus our study on the top-performing methods of two distinct sources of classifications; one from the internal-states of the LLM and another from the generated textual output. Additionally, we experiment with a third through combining these sources. Specifically, the *verbalised*, *hidden-state* and *combined* sources. Next, we formally define these sources.

Verbalised Prediction Source. The *verbalised probability* is the probability of the positive class extracted from the generated text of the LLM. We denote these probabilities for input x_i as \hat{t}_i .

Hidden-State Prediction Source. The second prediction probability is defined based on the hidden representations of LLM to implicitly detect classifications. This is inspired by Ren et al. (2023), which utilises the final-layer hidden embedding of LLMs for out-of-distribution detection. The output embedding $\hat{h}_i \in \mathcal{H}$ for input x_i is computed as the average of the decoder’s final-layer hidden-state vectors $g_{ik} \in \mathbb{R}^d$ over all K output tokens with a hidden dimension

of $d = 5120$ for the small-scale LLM:

$$\hat{h}_i := \frac{1}{K} \sum_{k=1}^K g_{ik}^n.$$

Then, an MLP $g : \mathcal{H} \mapsto \mathbb{R}$ is trained as a hidden-state classifier to learn the probability of the disorder from the LLM hidden representation, the *hidden-state prediction probabilities*, denoted $\hat{\epsilon}_i$, i.e. $\hat{\epsilon}_i = g(\hat{h}_i)$. We experimented with different models for the hidden-state classifier in ambitions of fully utilising the information embedded in \hat{h} (see Appendix D.1 for details).

Combined Prediction. Additionally, we combine the verbalised prediction and the hidden state prediction through their mean to form the *combined prediction*, of which we denote $\hat{\mu}_i$.

3.2. Instruction-Tuning Methodology

In generating well-calibrated verbalised probabilities, we use the “*verb. 1S top-k*” prompting strategy (Tian et al., 2023), prompting the LLM to provide the top k guesses and their probabilities in a single response. Adapting this to $k = 1$ for a binary setting, we prompt the LLM to return the probability of the positive class. Additionally, we prompt for the top reason the disorder *might* and *might not* be present, using *dialectic reasoning* (Hegel, 2018) to provide intelligent guidance for decision-makers. This technique has proven effective in decision support (Jarupathirun & Zahedi, 2007). An example output is in Figure 2.

Example Generated Guidance

TOP REASON FOR: The MRI report indicates that there is narrowing of the right exit foramen at the L5-S1 level, which is causing compression of the exiting right L5 nerve root. This suggests that there is foraminal stenosis (FS) at this level.

TOP REASON AGAINST: There is no mention of foraminal stenosis specifically at the L5-S1 level.

CONCLUSION: Based on the information provided, there is a possibility of foraminal stenosis at the L5-S1 level due to the narrowing of the right exit foramen and compression of the exiting right L5 nerve root. However, the report does not explicitly mention foraminal stenosis at this level.

PROBABILITY OF FS PRESENT AT L5-S1: 60%

Figure 2. Example guidance based on a spinal MRI radiological report. The instruction-tuned LLM is able to intelligently infer a diagnosis with sound logic without explicit textual diagnosis.

3.3. Deferral Mechanism

Our deferral strategy is based on the confidence of predictions that is determined by its distance to the chosen decision boundary of 0.5. Formally, predictions $\hat{p} \in \{\hat{t}, \hat{\epsilon}, \hat{\mu}\}$ are transformed to sorted relative confidence probabilities $\tilde{p} = \text{sorted}(2|\hat{p} - 0.5|)$ for equal comparison between the positive and negative classes. The resulting \tilde{p} determines the hierarchy of cases to be deferred.

The deferral performance is measured through recursively iterating through all elements \tilde{p} , deferring this prediction (without replacement) and measuring the classification performance of the LLM on the remaining cases. This procedure describes the AUARC metric (Nadeem et al., 2009) when measuring accuracy. Intuitively, good deferral behaviour should demonstrate a monotonically increasing accuracy with increasing deferral rate.

3.4. Pilot Study in Investigating the Effectiveness of Generated Guidance

We conduct a pilot study with 20 participants to evaluate the effectiveness of our guidance in the scenario of deferring 30 ($\approx 5\%$) of the most uncertain test predictions. Participants received background information, including clinical details, examples of prediction outcomes with associated MRI reports and the LLM’s performance on a validation set to help participants develop a sufficient mental model of the LLM, of which has been shown to be important in HAIC systems (Kulesza et al., 2012; Bansal et al., 2019). The ordering of the 30 questions were randomised for each participant to reduce order bias.

Participants moved to the next question if their prediction matched the LLM’s. If it differed, they received guidance and could either change their prediction or keep it based on the guidance and their understanding of the LLM. This process allowed us to assess human performance with and without guidance. Effective guidance should help participants recognise both the accuracy and inaccuracy of their judgements. Full details are in Appendix C.

4. Experiments

Data. We use the OSCLMRIC (Oxford Secondary Care Lumbar MRI Cohort) dataset, containing professionally annotated lumbar MRIs and radiological reports for various types of stenosis at different spinal levels, of which serve as our ground-truth labels. An example report is shown in Listing 2. The dataset is highly imbalanced, with $\approx 95\%$ of labels negative. Each report is parsed to detect the binary presence of three types of spinal stenosis (foraminal stenosis [FS], spinal canal stenosis [SCS], and lateral recess stenosis [LRS]) at six lumbar spine levels, resulting in 1,800 examples. The data is randomly split into 30% for generating an instruction-tuning dataset, 20% for training the hidden-state classifier, 20% for validation, and 30% for testing.

SOTA Baseline. We use Tulu V2 70B as the SOTA baseline model for our experiments, of which is the highest-performing open-source LLM against several benchmarks at the time these experiments were conducted (Beeching et al., 2023). Proprietary LLMs are omitted as baselines due to the aforementioned data privacy issues.

Table 1. Classification, calibration, deferral performance and LLM efficiency against test split (N=540) of 9 setups: 13B base and instruction-tuned models on their verbalised ($\hat{t}_{\text{BASE-13B}}$ and $\hat{t}_{\text{INSTRUCT-13B}}$ respectively), hidden-state ($\hat{e}_{\text{BASE-13B}}$ and $\hat{e}_{\text{INSTRUCT-13B}}$ respectively) and combined probability predictions ($\hat{\mu}_{\text{BASE-13B}}$ and $\hat{\mu}_{\text{INSTRUCT-13B}}$ respectively). We include Tulu-70B experiments as the SOTA baselines highlighted in grey. Statistically significant best results in bold.

SETUP	CLASSIFICATION PERF.			CALIBRATION PERF.		DEFERRAL PERF.	LLM EFFICIENCY		
	RECALL↑	PRECISION↑	F1-SCORE↑	ECE↓	ACE↓	AUARC↑	REL. S/GEN.↓	MEM.↓	E.R.↓
(1) $\hat{t}_{\text{BASE-13B}}$	0.85±0.02	0.61±0.01	0.61±0.01	0.26±0.01	0.32±0.01	0.897460±0.0063			
(2) $\hat{e}_{\text{BASE-13B}}$	0.73±0.02	0.94±0.03	0.80±0.02	0.03±0.01	0.04±0.01	0.994715±0.0006	0.08±0.02	6.02	0.29
(3) $\hat{\mu}_{\text{BASE-13B}}$	0.78±0.03	0.87±0.04	0.82±0.03	0.13±0.01	0.18±0.01	0.993318±0.0010			
(4) $\hat{t}_{\text{BASE-70B}}$	0.90±0.02	0.78±0.01	0.83±0.01	0.27±0.01	0.33±0.01	0.973529±0.0052			
(5) $\hat{e}_{\text{BASE-70B}}$	0.87±0.01	0.97±0.01	0.91±0.01	0.02±0.01	0.05±0.01	0.997191±0.0002	1.00	32.08	0.04
(6) $\hat{\mu}_{\text{BASE-70B}}$	0.90±0.01	0.95±0.02	0.92±0.01	0.14±0.01	0.19±0.01	0.997260±0.0003			
(7) $\hat{t}_{\text{INSTRUCT-13B}}$	0.85±0.02	0.92±0.02	0.88±0.02	0.27±0.01	0.32±0.01	0.987501±0.0024			
(8) $\hat{e}_{\text{INSTRUCT-13B}}$	0.89±0.01	0.98±0.01	0.93±0.01	0.01±0.01	0.05±0.01	0.997409±0.0002	0.08±0.02	6.02	0.0002
(9) $\hat{\mu}_{\text{INSTRUCT-13B}}$	0.91±0.01	0.98±0.01	0.94±0.01	0.14±0.01	0.19±0.01	0.996978±0.0013			

Abbreviations. We abbreviate IT and base (non-IT) models as “INSTRUCT” and “BASE”, respectively. Tests for statistical significance (SS) are conducted using a one-sided non-parametric Mann-Whitney U test (Mann & Whitney, 1947), unless stated otherwise.

Evaluation Metrics. The Area Under Accuracy-Rejection Curve (AUARC) and F1-score are used to measure the deferral and classification performance, respectively. We further evaluate the calibration of our setups with ECE (Guo et al., 2017) and ACE (Nixon et al., 2020). LLM efficiency is evaluated based on *Rel. s/Gen.* (seconds/generation relative to the baseline), *Mem.* (GPU VRAM required in GB), and *E.R.* the error rate (the number of unsuccessful generations for each successful generation).

4.1. Computational Results

Instruction-Tuned Models Are More Accurate, Equally Calibrated, Better Deferral Systems, and More Efficient. We find a SS improvement in F1-Score of verbalised predictions of $\hat{t}_{\text{INSTRUCT-13B}}$ in comparison to the baseline $\hat{t}_{\text{BASE-70B}}$, demonstrating instruction-tuning methodology to be successful. The greatest deferral performance results from the hidden-state predictions of the 13B setup, surpassing that of the 70B baseline. The IT model is seen to significantly improve in all LLM efficiency metrics.

Access to Internal States of LLMs Improves Deferral Performance. Utilising open-source LLMs allows for hidden-state predictions which are shown to be the best calibrated and are utilised in the greatest deferral performance (Table 1, Row 8). This finding suggests deferral systems built utilising proprietary LLMs without access to internal states and relying solely on verbalised predictions are inadequate.

The Combined Prediction Leads to Greatest Classification Performance. $\hat{\mu}_{\text{INSTRUCT-13B}}$ is the greatest classification source (Table 1, Row 9), surpassing that of

$\hat{e}_{\text{INSTRUCT-13B}}$ and $\hat{t}_{\text{INSTRUCT-13B}}$ ($p < 0.01$). The improvement in $\hat{\mu}$ over \hat{t} and \hat{e} is seen in all setups. This implies an interesting finding; in that the verbalised and hidden-state sources contain valuable and distinct pieces of information contributing to classification. There exists a positive correlation of $\rho = 0.53 \pm 0.07$ between these predictions. When focusing our analysis when $\hat{\mu}$ is correct and the predictions of \hat{t} and \hat{e} differ, a correct \hat{e} prediction is combining with an incorrect \hat{t} prediction in the majority of cases, at an average of $(75 \pm 0.14)\%$ of these cases. When $\hat{\mu}$ is incorrect, a correct \hat{t} prediction is combining with an incorrect \hat{e} prediction in the majority of cases, at an average of $(75 \pm 0.25)\%$ of these cases. See Figure 3 for confusion matrices.

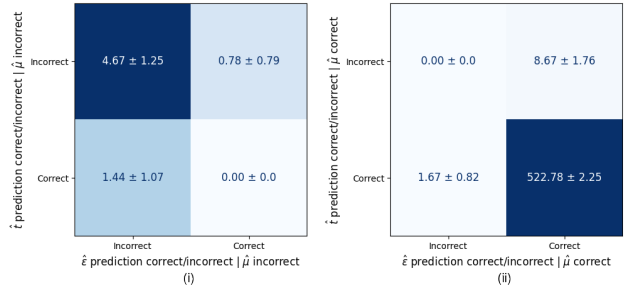


Figure 3. Confusion matrix illustrating the number of test cases for \hat{t} and \hat{e} precision, given (i) the incorrect prediction of their combination $\hat{\mu}$, and (ii) correct prediction.

The Combined Prediction Has The Greatest Deferral Performance in Mostly-Autonomous Systems. We find a SS ($p < 0.1$) improvement in the deferral strategy of deferring on $\hat{e}_{\text{INSTRUCT-13B}}$ and classifying on $\hat{\mu}_{\text{INSTRUCT-13B}}$, in comparison to the next-best strategy of deferring and classifying solely on $\hat{e}_{\text{INSTRUCT-13B}}$, in situations where $< 5\%$ of cases are deferred. In practice, this would be cases where users would require the system to be mostly autonomous. This implies that the combined prediction is a greater pre-

dictor of disease in high-uncertainty situations. In deferral rates $\geq 5\%$, there exists no SS difference between the two strategies, but a SS over all other setups.

4.2. Pilot Study Results

Following the computational deferral results in high uncertainty, we defer 5% of the test cases based on the hidden-state prediction and provide the classification prediction of the combined prediction.

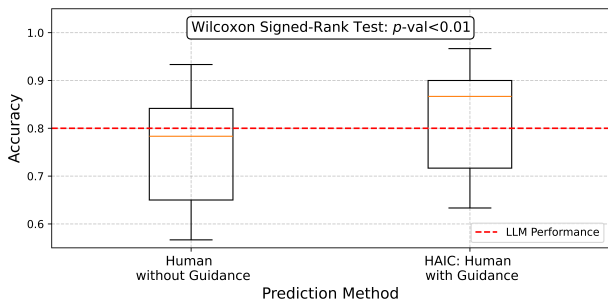


Figure 4. **Human accuracy without guidance is lower and more variable than with guidance (HAIC).** Accuracy of prediction methods from pilot study. On average, guided humans outperform both unguided humans and the LLM.

Guidance is Effective in Aiding Human Decision-Making. Figure 4 displays the accuracy of participant decision-making with and without guidance, compared to LLM performance. All participants’ accuracy improved with guidance. We rejected the null hypothesis that performance without guidance equals performance with guidance (paired one-sided Wilcoxon signed-rank test, $p < 0.01$). When confronted with AI disagreement, the provided guidance proved effective in assisting participants to arrive at the correct final decision. This was true not only when the participant was incorrect, but importantly also when the LLM was incorrect. We rejected the null hypothesis that the proportion of humans changing their prediction is independent of the AI’s prediction correctness (χ^2 -test, $p < 0.01$).

5. Conclusion

We develop a guided deferral framework using LLMs for clinical decisions, with HAIC as a key focus. Our study shows that IT small-scale LLMs can achieve significant deferral and classification performance while maintaining computational efficiency through instruction-tuning on supervised data from a large-scale LLM. We utilise two prediction sources—verbalised and hidden-state, yielding valuable insights for classification and deferral. The combination of these sources results in a significant classification and high-uncertainty deferral prediction. Finally, we validate the effi-

cacy of our proposed guided deferral system through a pilot study, with the results showing a significant improvement in human decision-making performance under the LLM guidance.

Limitations. In this paper, we prioritise healthcare in our analysis due to its greater ethical significance in research. However, there exists a severe lack of high-quality report data with labels in medical domain, making it difficult for LLM evaluation. When implementing our system, extra efforts are required to educate clinicians about its capabilities to prevent unexpected errors. This involves addressing human biases such as anchoring and confirmation bias, as well as briefing them on the system’s training data distribution, including concepts such as data drift and out-of-distribution instances. Such understanding is vital, particularly in scenarios where variations in report style or format might affect the system’s performance.

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A. Additional Listings

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### QUESTION: Is there <disorder> at the <ivd> level?

### INSTRUCTIONS: You are an expert spinal surgeon, giving guidance to novices in diagnosing <disorder> at the <ivd> level based on the
contents of a report. Firstly, start by giving the single top reason both for and against, considering information ONLY
regarding the <ivd> level. Finally, at the end of your response, give a final detailed conclusion including a definitive answer
to the question, based solely on your top reasons, and importantly including a probability that <disorder short> is present at
the <ivd> level in this format: 'CONCLUSION: <your conclusion>. PROBABILITY OF <disorder short> PRESENT AT <ivd>: <your
probability>% </s>'. You must end your response after this.

### RULES:
1. Your probability must not be 50%.
2. If you think it is present, your probability must be greater than 50%.
3. If you think it is not present, your probability must be less than 50%.
4. Even if the question is not answerable, you must still give a definitive answer and probability.
5. Give your answer only in English.

### ASSUMPTIONS:
1. Assume that any severity or of <disorder> indicates that it is present and your probability should be greater than 50%.
2. The absence of information about <disorder> at the <ivd> level indicates that it is not present and your probability should be less
than 50%.
3. Assume that you can only use information at the <ivd> level to diagnose <disorder>.
4. Foraminal Stenosis is the narrowing of the foramen, Lateral Recess Stenosis is the narrowing of the lateral recess, and Spinal Canal
Stenosis is the narrowing of the central spinal canal.
5. The presence of <disorder1> or <disorder2> does not imply the presence of <disorder>.

### REPORT: <report>
```

Listing 1. Base-prompt used in generating data to instruction-tune and in parsing of reports. The relevant level, disorder, question, and report were inserted to make the final prompt.

```
Clinical History :
Exam.: MRI Spine lumbar and sacral
Reason for Study: ? Right S1 Nerve Root Pain suitable for
injection/surgery
Clinical Information: 7/12 severe and worsening right leg
pain S1. Normal neurology. High disability, 2
crutches/wheelchir outdoors. IIEitis/crohns.

MRI Spine Lumbar and Sacral :
L5 vertebra plana in addition to diffuse infiltration of L4,
with minor collapse. There is also patchy involvement of the
S1 vertebral body, and of the right ilium and to a lesser
extent the left ilium. Small lesion in L1. Satisfactory
vertebral alignment.

Axial images L3-S1.
L3-4 no neural compression.
L4-5 narrowing of the right lateral recess due to tumour
with impingement of the traversing right L5 nerve root.
L5-S1 narrowing of the right exit foramen due to tumour with
compression of the exiting right L5 nerve root. Minor
displacement of the right S1 nerve root.

Conclusion:
Features of metastatic disease. Right-sided nerve root
compression as described.
```

Listing 2. An example spinal MRI radiological report.

B. Experimental Details

B.1. Inference

Inference hyperparameters are as follows:

- max_new_tokens: 1000
- temperature: 0.8
- top_k: 50
- top_p: 0.95

B.2. Training

We instruction-tune the 13 billion parameter LLM *Tulu V2 DPO* (Iverson et al., 2023) using QLoRA (Dettmers et al., 2023) on generations solely. The base model can be downloaded from <https://huggingface.co/allenai/>

tulu-2-dpo-13b. Training time on a single GeForce RTX 4090 varied by number of generated sets, but for no longer than 6 hours, with CUDA v12.4.

The hyperparameters used for training are listed below:

- **Model Preparation:**

- rank (r): 128
- target_modules: {q_proj, k_proj, v_proj, o_proj, gate_proj, up_proj, down_proj}
- lora_alpha: 16
- lora_dropout: 0
- bias: none
- use_gradient_checkpointing: True
- random_state: 3407
- max_seq_length: 4096

- **Training Configuration:**

- Training Batch Size: 10
- warmup_steps: 10
- logging_steps: 1
- learning_rate: 2e-4
- Optimiser: adamw_8bit
- bf16: True
- weight_decay: 0.1
- warmup_ratio: 0.01
- lr_scheduler_type: linear

B.3. Guard-rail Implementation

Algorithm 1 details the implementation of the guard-rail used in generating our instruction-tuning data.

Algorithm 1 Guard-rail Algorithm in Generating Instruction-Tuning Data

```
1: Set failure_count = 0
2: repeat
3:   Generate a response
4:   if response contains non-English text or does not end with an EOS token or contains multiple probability predictions
     or contains no firm answer then
5:     Retry generation
6:   else
7:     if prediction matches the label then
8:       Generation is successful
9:     End Algorithm
10:  else
11:    Increment failure_count by 1
12:  end if
13: end if
14: until failure_count = 10
15: Ignore label check and save the prediction.
```

C. Pilot Study

We detail the instructions given to pilot study participants in this section.

Background on the task: Spinal Stenosis

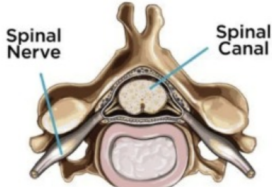
In this task, you will be asked to provide a **positive or negative diagnosis** for the presence of one of **three types of spinal stenosis** at one of **six specific intervertebral levels** based on the contents of an MRI report. Spinal stenosis is a medical condition characterised by the narrowing of the spaces within the spine, which can put pressure on the nerves that travel through the spine.

The three specific types of spinal stenosis you will be diagnosing are:

1. **Spinal Canal Stenosis (SCS)**: Also known as "central stenosis". This is the narrowing of the central spinal canal, the space in the vertebrae through which the spinal cord travels.
2. **Forminal Stenosis (FS)**: This is the narrowing of the passageways (foramina) in the spine through which nerves exit the spinal canal.
3. **Lateral Recess Stenosis (LRS)**: This is the narrowing of the openings (recesses) in the sides (lateral) of the spinal canal on the sides of the vertebrae through which the nerves travel. When there is lateral recess stenosis in both sides of the vertebrae, you will see this referred as "bilateral" recess stenosis.

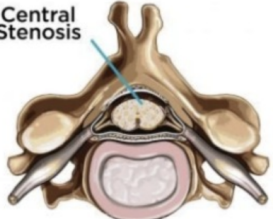
You'll only be diagnosing these disorders based on the contents of MRI textual reports, but below are some images of the 3 types of stenosis compared to a healthy vertebrae.

Healthy Vertebrae



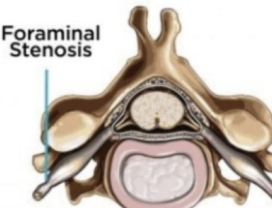
(a) Healthy vertebrae

Central Stenosis



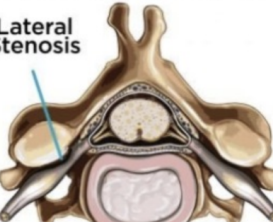
(b) Central Stenosis (Canal Stenosis)

Foraminal Stenosis



(c) Foraminal Stenosis

Lateral Stenosis



(d) Lateral Recess Stenosis

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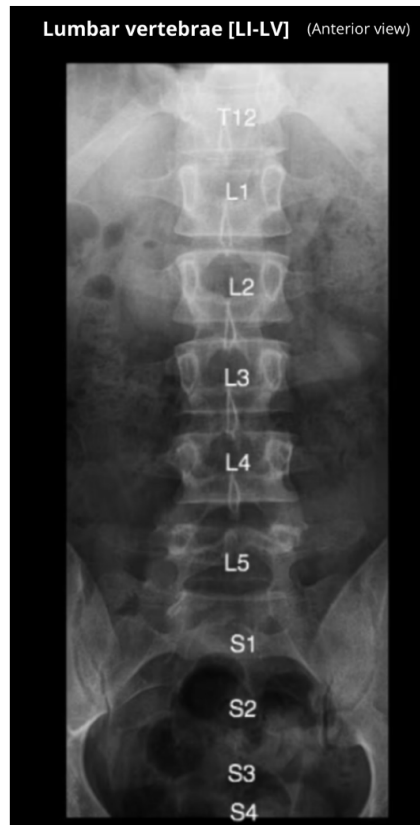
Figure 5. Page 1: Participants are first briefed on the clinical diagnosis task at hand, including the three types of spinal stenosis.

Background on the task: Intervertebral Levels

You will be specifically asked to diagnose these three types of spinal stenosis occurring at **one of six specific intervertebral levels** on the lumbar spine. Lumbar intervertebral levels refer to the specific segments of the lower back where the vertebrae are located. The lumbar spine consists of five vertebrae, numbered L1 to L5, which are stacked on top of each other. Each intervertebral level represents the space between two adjacent vertebrae.

In this task, you will be asked to diagnose at one of 6 intervertebral levels from the contents of the MRI reports. Inside the MRI reports, the intervertebral levels are (mainly) represented as follows:

1. T12-L1.
2. L1-L2.
3. L2-L3.
4. L3-L4.
5. L4-L5.
6. L5-S1.



Continue

Go Back

Figure 6. Page 2: Next, participants are first briefed on the notion of intervertebral levels.

Background on the task: The AI Model

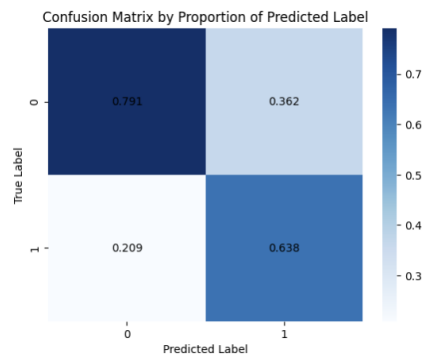
In this study, you will be classifying cases of which an AI model is uncertain about. Specifically, these cases are the 5% least certain cases of a test dataset.

The cases of which you will see were specifically chosen because an AI model deemed it was uncertain about classifying them itself. This does not mean that the cases are necessarily difficult for a human, but rather that the AI model was not confident in its diagnosis for its own reasons.

The confusion matrix below shows the performance of this AI model against a subset of a validation dataset. You can assume that this subset is representative of the test dataset that you will be diagnosing against.

The main points to take from the confusion matrix below are:

1. **When the model predicts a positive diagnosis, it is correct roughly 3 times out of 4.**
2. **When the model predicts a negative diagnosis, it is correct roughly 9 times out of 10.**
3. **The model is correct more often when making a negative diagnosis than a positive diagnosis.**



Continue

Go Back

Figure 7. Page 3: We give participants a mental model of the classification performance of the LLM of which they should expect.

Background on the task: AI Guidance

The AI not only provides a prediction, but also provides textual **guidance** on how to interpret its predictions. The guidance can be used to understand the model's decision-making process and to identify potential flaws in its prediction, and is designed to propose both **reasons for and against** making a positive diagnosis.

You will only be given this guidance if your prediction disagrees with the AI's prediction.

The ultimate prediction that the model makes is based off the guidance that it produces. The AI model has been fine-tuned to solely improve the classification performance of this guidance. During this process, the guidance will have been indirectly improved, **but not directly to a point where it is perfect** - the guidance can have poor reasoning.

It's up to you to decide whether the ultimate prediction is correct or not. You should incorporate its **confidence, guidance, the task assumptions, and your perception of how good the model is** to make a final decision.

Below is some example guidance of which has the same structure as the guidance you will see in the real study.

Example AI Guidance:

Reasoning For Foraminal Stenosis at L4-L5:

The report mentions "narrowing of the central canal and neural exit foramina with associated impingement on the exiting left L4 and both traversing L5" which indicates the presence of foraminal stenosis at the L4-L5 level.

Reasoning Against Foraminal Stenosis at L4-L5

The report does not explicitly mention foraminal stenosis at the L4-L5 level.

AI prediction:

Foraminal Stenosis is **present** at the L4-L5 level.

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Figure 8. Page 4: We brief participants on the type of guidance they should expect during the study.

Background on the task: True Positive Example

Here's an example. In the real test, you will only be given the AI-guidance if your prediction differs from the AI. You will then be asked if you want to change your decision.

Question: Is there Foraminal Stenosis at the L4-L5 intervertebral level?

Report:

AI Guidance:

TOP REASON FOR:

TOP REASON AGAINST:

AI Prediction: Positive Diagnosis

You'll be asked to record either a positive or negative prediction, and how confident you are on your decision.

Choose Prediction:

Negative Positive

Low Confidence Moderate Confidence High Confidence

Answer: Positive - Foraminal Stenosis is present at the L4-L5 intervertebral level.

Continue - True Negative Example Go Back

Figure 9. Pages 5-9: Participants are given 4 examples; a true a positive, a false positive, a true negative and a true positive. Sensitive data is censored behind grey boxes.

Are you ready?

Thanks again for participating in this pilot study!

Before beginning please keep in mind some assumptions that the expert annotator, the model and also you should make when diagnosing:

- **If there are no signs (either explicit or implicit) of a disorder, you should assume it is not present**
- **The presence of a disorder at one intervertebral level does not imply it is present at another**
- **Any severity of a disorder indicates that it is present, i.e. even mild symptoms warrants a positive diagnosis.**

After this page, the study will begin. The study will take approximately 15-20 minutes. Please make sure that you are able to concentrate uninterrupted for this time, as you cannot pause the study and your time to answer questions is recorded!

Any problems or questions, please message

If you are completely unsure about a prediction - have a go and reflect this in your recorded confidence!

Figure 10. Page 10: Before starting the study, users are briefed on the assumptions they should make when predicting.

Study: 1 of 30

Assumptions you should make:

- If there are no signs (either explicit or implicit) of a disorder, you should assume it is not present
- The presence of a disorder at one intervertebral level does not imply it is present at another
- Any severity of a disorder indicates that it is present, i.e. even mild symptoms warrants a positive diagnosis.

Question: Is there **Spinal Canal Stenosis** at the **L4-L5** level?

Report:

Choose Prediction:

Please press only one button per row!

Prediction Mismatch

Your prediction does not match the AI prediction. AI predicted Negative whilst you predicted Positive

Why it could be positive:

Why it could be negative:

Figure 11. Page 11-41: The study consists of 30 of the most uncertain predictions from the LLM. Participants are first asked to make their own prediction. If their prediction matches with the LLM, they continue to the next question. Otherwise, they are told they are in disagreement with the AI and are provided guidance. They are then given the opportunity to change their decision. Additionally, user's time to completion and self-recorded confidence are recorded during the study. Confidential data is censored behind grey boxes.

D. Additional Experiments

D.1. Hidden-State Classifier Experiments

In order to maximise the predictive capabilities of the final-hidden layer of the LLM, we experiment with different hidden-state MLP classifier architectures. We choose the architecture of which maximises the deferral performance (AURAC) on the validation split. Table 2 details the results of these experiments. We choose a three fully-connected layer as our hidden-state classifier due to joint statistically significant deferral performance and F1-Score.

Setup	F1-Score	AURAC
One fully-connected layer	0.88 \pm 0.03	0.995734 \pm 0.0004
Three fully-connected layers	0.91 \pm 0.01	0.995976 \pm 0.0003
Five fully-connected layers	0.90 \pm 0.01	0.996068 \pm 0.0003

Table 2. Comparison of classification performance (F1-Score) and deferral performance (AURAC) of two setups: one fully-connected layer and three fully-connected layers.