Responsible NLP Checklist

Paper title: *Med-VRAgent: A Framework for Medical Visual Reasoning-Enhanced Agents*
Authors: *guangfu guo, Xiaoqian Lu, Yue Feng*

How to read the checklist symbols:

☑ the authors responded 'yes'

☒ the authors responded 'no'

N/A the authors indicated that the question does not apply to their work

☐ the authors did not respond to the checkbox question

For background on the checklist and guidance provided to the authors, see the Responsible NLP Checklist page at ACL Rolling Review.

☑ **A. Questions mandatory for all submissions.**

☑ A1. Did you describe the limitations of your work?
This paper has a Limitations section.

☑ A2. Did you discuss any potential risks of your work?
See Section Ethical Considerations. We discuss that hallucinations and reasoning errors in medical tasks may lead to potential clinical risks. Our work is limited to research datasets and not directly deployed in clinical practice.

☑ **B. Did you use or create scientific artifacts? (e.g. code, datasets, models)**

☑ B1. Did you cite the creators of artifacts you used?
See Section 5.1 (Experimental Datasets). We cite IU-Xray, MIMIC-CXR, VQA-RAD, and GMAI-MMbench datasets, and all models used (e.g., LLaVA-Med, DeepSeek-VL, MiniCPM-V2).

☑ B2. Did you discuss the license or terms for use and/or distribution of any artifacts?
See Section 5.1 (Experimental Datasets). All datasets used are publicly available under research-only licenses (e.g., MIMIC-CXR requires credentialed access).

☑ B3. Did you discuss if your use of existing artifact(s) was consistent with their intended use, provided that it was specified? For the artifacts you create, do you specify intended use and whether that is compatible with the original access conditions (in particular, derivatives of data accessed for research purposes should not be used outside of research contexts)?
See Section 5.1. All datasets are used only for research purposes, consistent with their intended use and license conditions.

☑ B4. Did you discuss the steps taken to check whether the data that was collected/used contains any information that names or uniquely identifies individual people or offensive content, and the steps taken to protect/anonymize it?
See Section 5.1. All datasets (e.g., MIMIC-CXR) have been de-identified by their providers, ensuring no personally identifiable information remains.

N/A B5. Did you provide documentation of the artifacts, e.g., coverage of domains, languages, and linguistic phenomena, demographic groups represented, etc.?
(left blank)

The Responsible NLP Checklist used at ACL Rolling Review is adopted from NAACL 2022, with the addition of ACL 2023 question on AI writing assistance and further refinements based on ARR practice.

☑ B6. Did you report relevant statistics like the number of examples, details of train/test/dev splits, etc. for the data that you used/created?
See Section 5.1. We report dataset sizes, splits, and evaluation metrics (BLEU, ROUGE-L, METEOR, accuracy).

☑ **C. Did you run computational experiments?**

☑ C1. Did you report the number of parameters in the models used, the total computational budget (e.g., GPU hours), and computing infrastructure used?
See Section 5.3 (Model Implementation). We report model sizes, training setup, GPU types (Nvidia A6000), hours, and hyperparameters.

☑ C2. Did you discuss the experimental setup, including hyperparameter search and best-found hyperparameter values?
See Section 5.3. We specify batch size, learning rate, LoRA parameters, PPO hyperparameters, etc.

☑ C3. Did you report descriptive statistics about your results (e.g., error bars around results, summary statistics from sets of experiments), and is it transparent whether you are reporting the max, mean, etc. or just a single run?
See Section 6 (Experiments and Ablations).

☑ C4. If you used existing packages (e.g., for preprocessing, for normalization, or for evaluation, such as NLTK, SpaCy, ROUGE, etc.), did you report the implementation, model, and parameter settings used?
See Section 5.3. We list software frameworks used (peft, trl, FAISS, Grounding DINO, BioClinicalBERT, etc.), with parameter settings.

☒ **D. Did you use human annotators (e.g., crowdworkers) or research with human subjects?**

N/A D1. Did you report the full text of instructions given to participants, including e.g., screenshots, disclaimers of any risks to participants or annotators, etc.?
(left blank)

N/A D2. Did you report information about how you recruited (e.g., crowdsourcing platform, students) and paid participants, and discuss if such payment is adequate given the participants' demographic (e.g., country of residence)?
(left blank)

N/A D3. Did you discuss whether and how consent was obtained from people whose data you're using/curating (e.g., did your instructions explain how the data would be used)?
(left blank)

N/A D4. Was the data collection protocol approved (or determined exempt) by an ethics review board?
(left blank)

N/A D5. Did you report the basic demographic and geographic characteristics of the annotator population that is the source of the data?
(left blank)

☒ **E. Did you use AI assistants (e.g., ChatGPT, Copilot) in your research, coding, or writing?**

N/A E1. If you used AI assistants, did you include information about their use?
(left blank)