Responsible NLP Checklist

Paper title: Aspect-Oriented Summarization for Psychiatric Short-Term Readmission Prediction Authors: WonJin Yoon, Boyu Ren, Spencer Thomas, Chanhwi Kim, Guergana K Savova, Mei-Hua Hall, Timothy A. Miller

How to read the checklist symbols:	
the authors responded 'yes'	
X the authors responded 'no'	
the authors indicated that the question does not apply to their work	
\square the authors did not respond to the checkbox question	
For background on the checklist and guidance provided to the authors, see the Responsible NLF page at ACL Rolling Review.	Checklist

✓ 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?

 Limitation section: We mentioned that this is not an application ready for direct clinical applications.
- **B.** Did you use or create scientific artifacts? (e.g. code, datasets, models)
 - B1. Did you cite the creators of artifacts you used?

 We cite the authors of language model papers, including BiomedBERT and Clinical LongFormer in Section 4.1, Llama in Section 5.2, and Qwen models in Appendix C..
 - B2. Did you discuss the license or terms for use and/or distribution of any artifacts? We are not releasing any assets.
 - ☑ 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)?

 We described about IRB approval and the secure environment in Section 2.1.
 - ☑ 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?
 - Section 2.1. All experiments, including data processing, were conducted in HIPAA-compliant environments. Appendix B.
 - ☑ B5. Did you provide documentation of the artifacts, e.g., coverage of domains, languages, and linguistic phenomena, demographic groups represented, etc.?

 Section 2.1
 - ☑ 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?

 Table 1, Section 4.2, Appendix F (Table 4)

☑ 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?

 Appendix D
- C2. Did you discuss the experimental setup, including hyperparameter search and best-found hyperparameter values?

 Appendix A, D
- 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?

Figure 4 (left), Appendix E, Appendix F, Figure 8

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?

Section 3.2.1

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

- 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*)
- 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*)
- 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)?

 This is a retrospective study using Electronic Health Records, and we comply with IRB requirements and applicable regulations regarding dataset collection.
- ✓ D4. Was the data collection protocol approved (or determined exempt) by an ethics review board? *IRB approval mentioned in Section 2.1.*
- 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?

☑ E1. If you used AI assistants, did you include information about their use?

Under the ACL Policy on Publication Ethics, our use case does not need to be disclosed. Our use case: a. Assistance purely with the language of the paper.