

Abstract

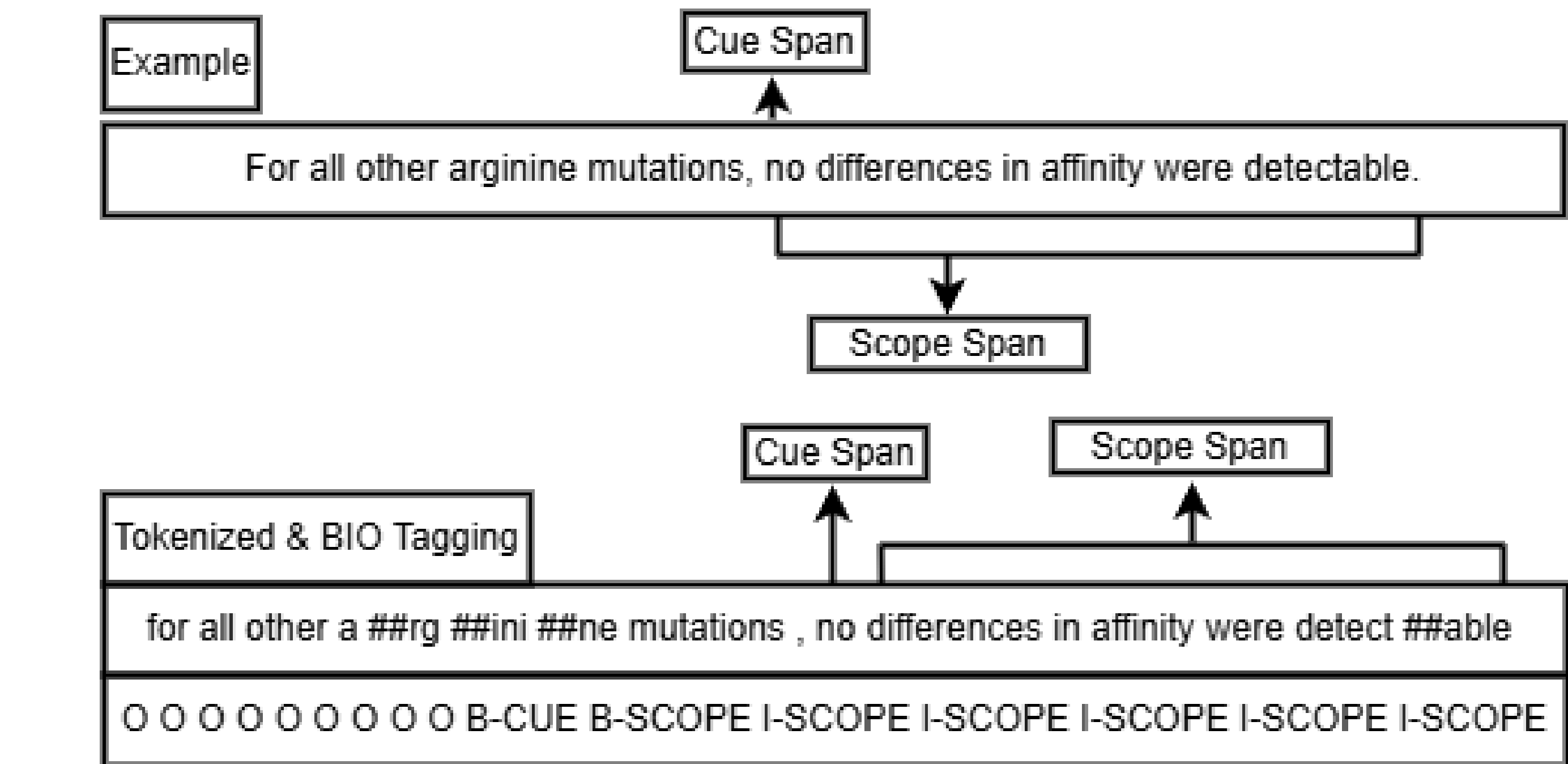
- Negation and speculation in biomedical texts affect the interpretation of clinical findings and and their downstream analysis.
- This project explores how contextual embeddings from BERT-based models can detect negation cues and scopes in sequence labeling tasks.

Project Objective

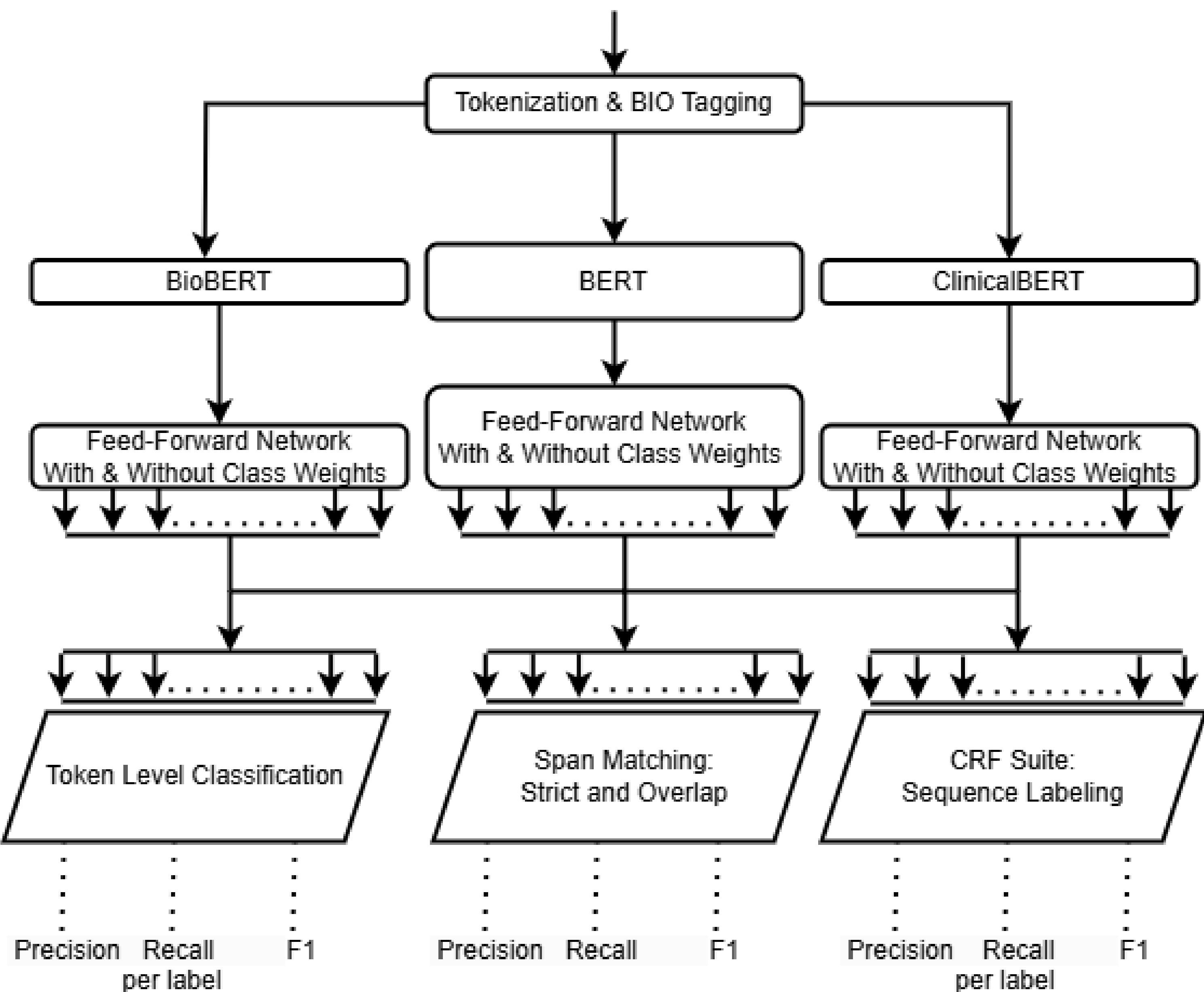
- Convert cue and scope spans in the BIOSCOPE dataset [1] into sequence-labeled BIO tags: (B-CUE, I-CUE, B-SCOPE, I-SCOPE, O).
- Fine-tune BERT-based models on BIO tagged data to evaluate their ability to detect negation cues and scopes.

Methodology

- Cue spans were removed from scope spans following [2].
- Fine-tuned BioBERT, BERT, and ClinicalBERT using BIO-tagged labels with a Feed Forward Network.
- Loss was calculated without and with class weights (Methods 1 and 2) to address label imbalances.
- Performance Evaluated on Token-Level Classification, Span Matching, and CRF Suite.



Example: Original sentence, tokenized text, and BIO-tagged labels.



Proposed architecture for fine-tuning and evaluation.

Results

| Model | Method | Token-Level | Span Matching | | CRF Suite |
|--------------|--------|-------------|---------------|--------------|------------|
| | | Micro F1 % | Strict F1 % | Overlap F1 % | Micro F1 % |
| BioBERT | 1 | 98.01 | 18.12 | 53.82 | 60.92 |
| BERT | 1 | 98.14 | 21.03 | 57.24 | 63.07 |
| ClinicalBERT | 1 | 97.85 | 16.77 | 52.10 | 57.52 |
| BioBERT | 2 | 96.34 | 12.65 | 41.59 | 50.86 |
| BERT | 2 | 96.59 | 15.42 | 44.93 | 53.81 |
| ClinicalBERT | 2 | 96.31 | 13.43 | 40.78 | 50.62 |

Performance of BERT-based models.

Sample Predictions: True vs. Predicted Labels

True Labels:
This, recognition, does, not (B-CUE), require (B-SCOPE), any (I-SCOPE), of (I-SCOPE), the (I-SCOPE), known (I-SCOPE), roX (I-SCOPE), RNAs (I-SCOPE).

Predicted Labels:
This, recognition, does, not (B-CUE), require (I-SCOPE), any (I-SCOPE), of (I-SCOPE), the (I-SCOPE), known (I-SCOPE), roX (I-SCOPE), RNAs (I-SCOPE).

Evaluation

- Token-Level:** Measures the F1-score for each token in the sequence.
- Span Matching:** Evaluates spans using Strict Match (exact) and Overlap Match (partial) criteria.
- CRF Suite:** Assesses sequence-level F1-scores by comparing predicted sequences to ground truth.

Analysis

- BERT** (Method 1) achieves the highest scores across tasks.
- Class weights improve performance on underrepresented labels but reduce overall scores.
- BioBERT** ranks second, while **ClinicalBERT** underperforms.
- BERT** embeddings are most effective for detecting negation cues and scopes.

Limitations and Future Work

- Limitations:**
- ClinicalBERT was used due to the unavailability of a pretrained NegBERT model.
 - Dataset imbalance, particularly for negation-related labels, impacted model performance.
- Future Work:**
- Introduce a CRF layer for enhanced sequence labeling..
 - Adopt [2] unified framework to integrate multiple negation datasets.

References

[1] V. Vincze, G. Szarvas, R. Farkas, G. Móra, and J. Csirik, "The bioscope corpus: Biomedical texts annotated for uncertainty, negation and their scopes," *BMC bioinformatics*, vol. 9, pp. 1–9, 2008. [Online]. Available: <https://doi.org/10.1186/1471-2105-9-S11-S9>.

[2] A. Yoshida, Y. Kato, and S. Matsubara, "Negation scope conversion: Towards a unified negation-annotated dataset," Torino, Italia: ELRA and ICCL, May 2024, pp. 12 093–12 099. [Online]. Available: <https://aclanthology.org/2024.lrec-main.1057>.

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