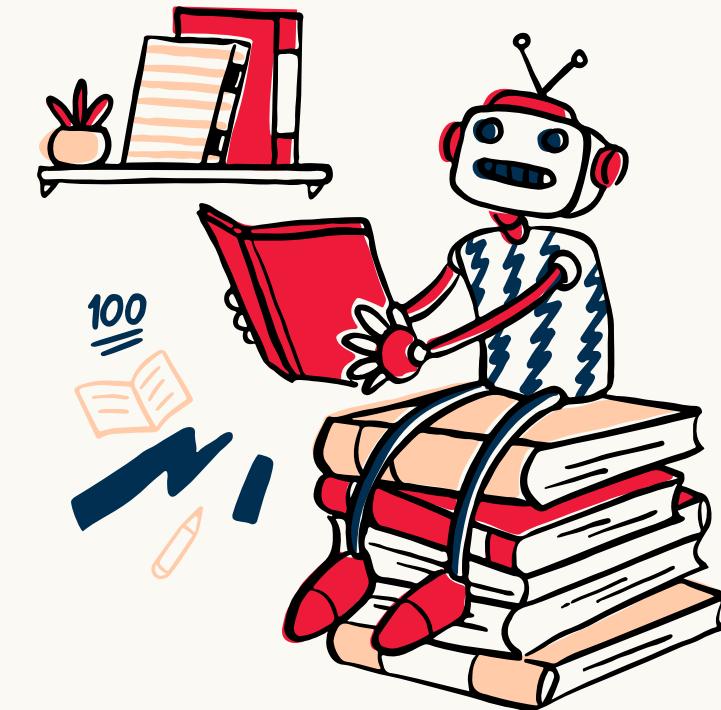


APPLIED DATA SCIENCE PROJECT

Patient Preference Studies Classification Project



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Cesar Augusto Seminario Yrigoyen
Francesco Giuseppe Gillio



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The Picture



"The urgent demand for tools that support efficient access, integration, and analysis of health data to derive actionable insights from **patient-reported outcome** and real-world evidence"

- European Commission



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The Project Value Proposition



For medical researchers who struggle to access true Patient Preference Studies (PPS) in PubMed's vast, noisy search results—where rule-based systems fall short—our AI-driven software harnesses LLM's semantic analysis to detect true PPS and uncover relevant clinical sub-areas



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The Project Value Proposition

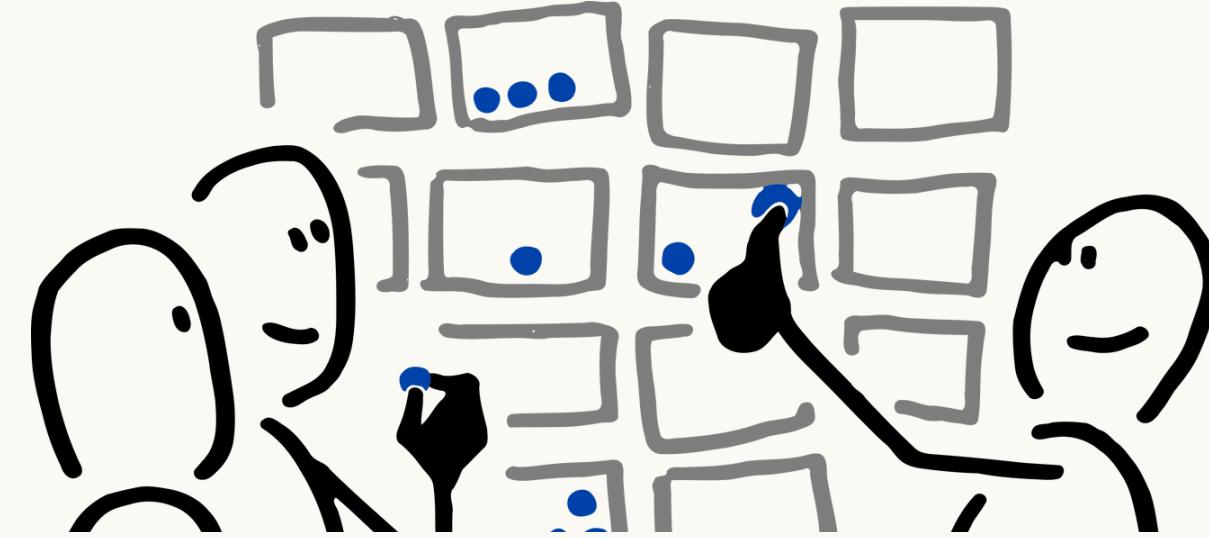
Our project **supports** evidence-based healthcare
and **enhances** research access and knowledge discovery



United Nations
Sustainable Development Goals
2030



The Objective



Support efficient access, collection,
and validation of Patient
PreferenceStudies

Develop an automated system for
effective classification of PPS
into relevant categories

An **easy-to-use AI software** to detect true Patient Preference Studies
and uncover relevant clinical sub-areas
from PubMed search results



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Data Collectors

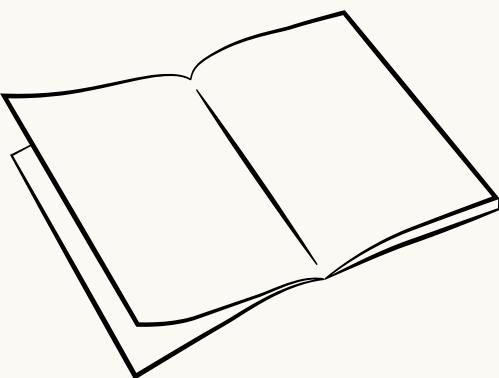


Medical Researchers



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Data Source



PubMed
links queries to records via
rule-based systems (MeSH)
and lightweight machine learning
algorithms (Best Match)

Textual Data

The head-to-head comparison of diabetic **patient preferences** for glucose-monitoring devices.

Abstract

A comparison of discrete choice experiment (DCE) and swing-weighting (SW) methods assesses diabetic **patient preferences** for glucose-monitoring devices. The analysis highlights critical attributes such as ease of use, accuracy, and cost, revealing differences in attribute prioritization and trade-offs. Insights from this evaluation inform the selection of preference-elicitation techniques in patient-centered healthcare research.



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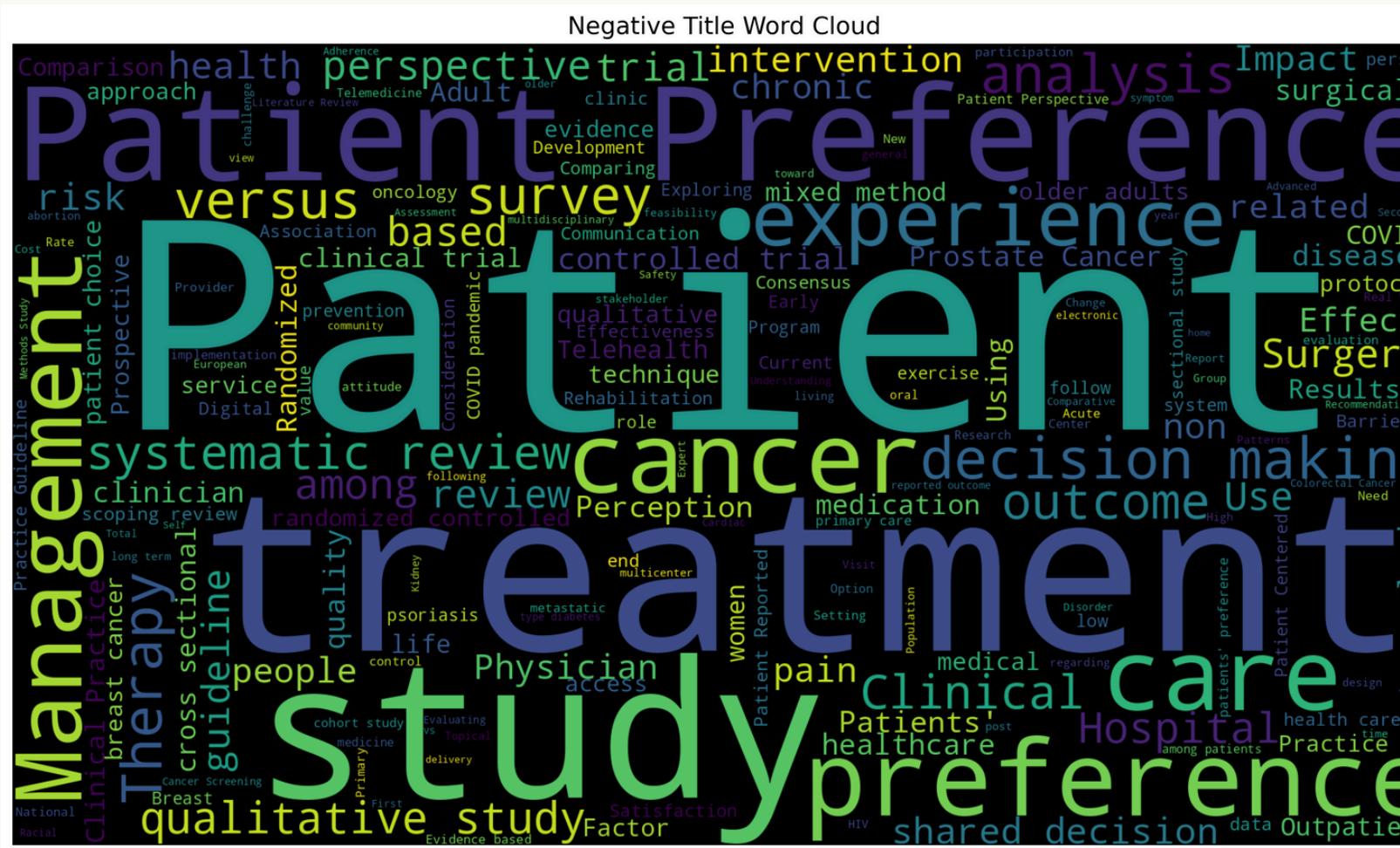
The Data

The head-to-head comparison of diabetic patient preferences for glucose-monitoring devices.

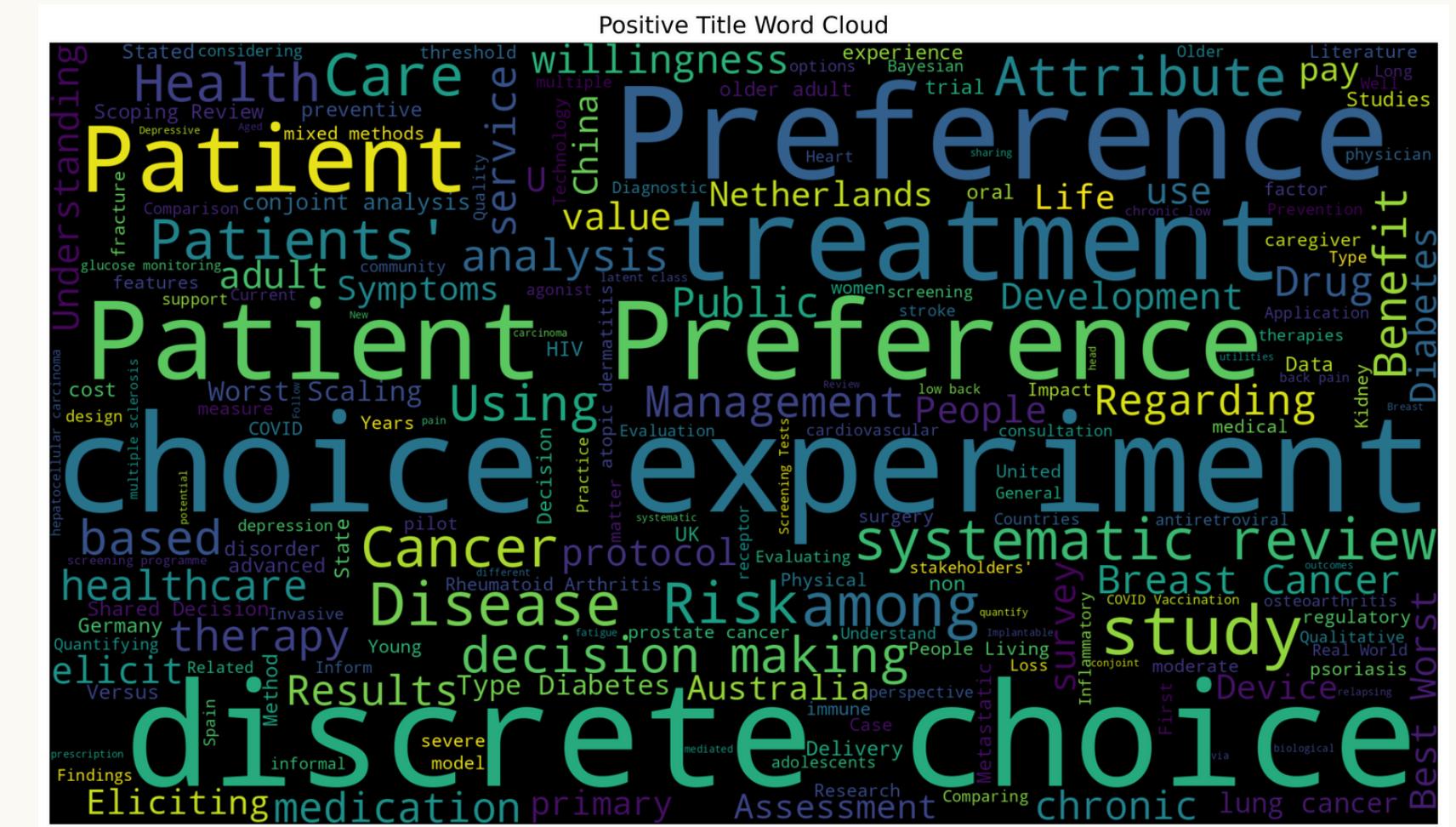
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Non Patient Preference Study



Patient Preference Study



The Binary Text Classifier

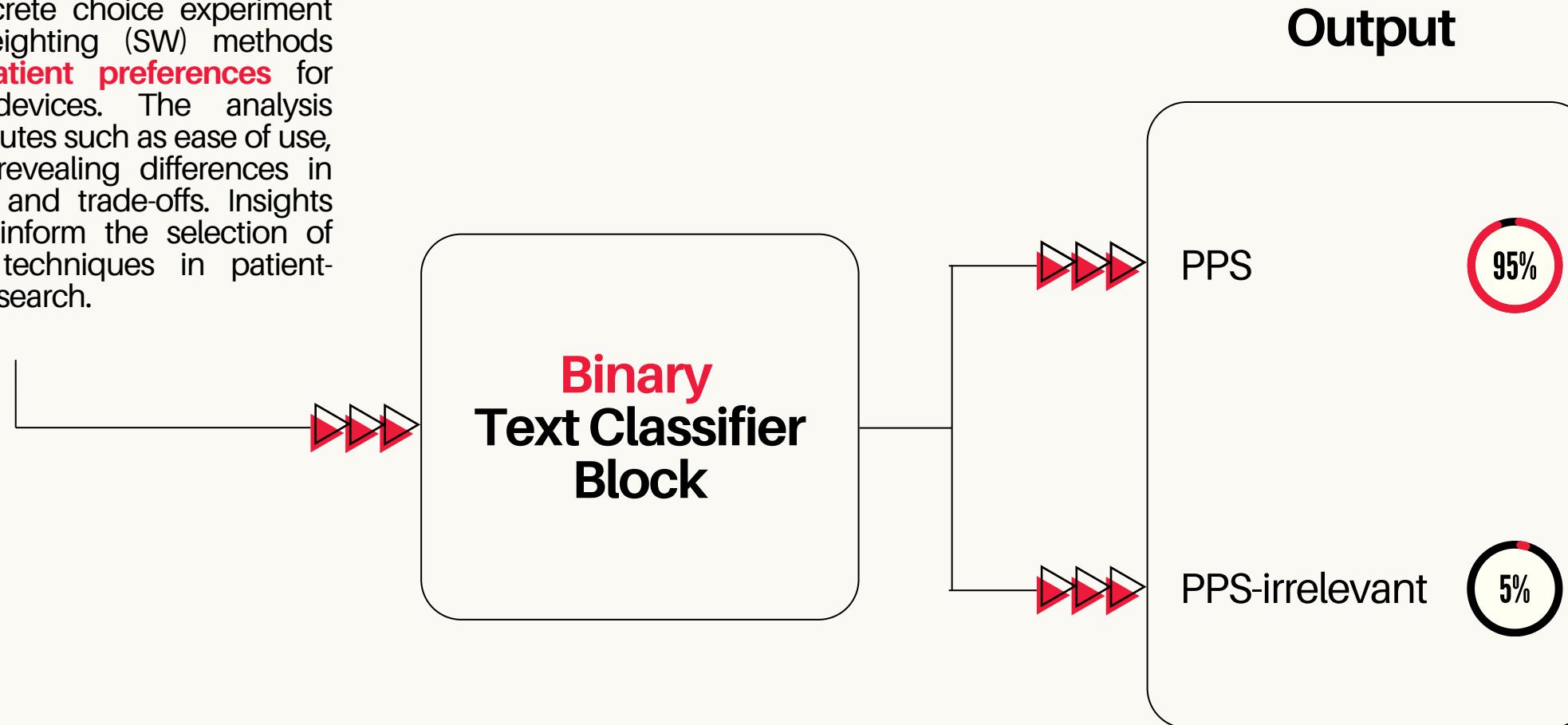


Binary Text Classification

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The Hypotheses and Research Questions

Large Language Models (LLMs) understand contextual relationships in text, enough to fool a human



How to adapt a Large Language Model for text classification by relevance to **Patient Preference Studies?**

Traditional ML Models (MLMs) continue to reign supreme in clinical prediction systems

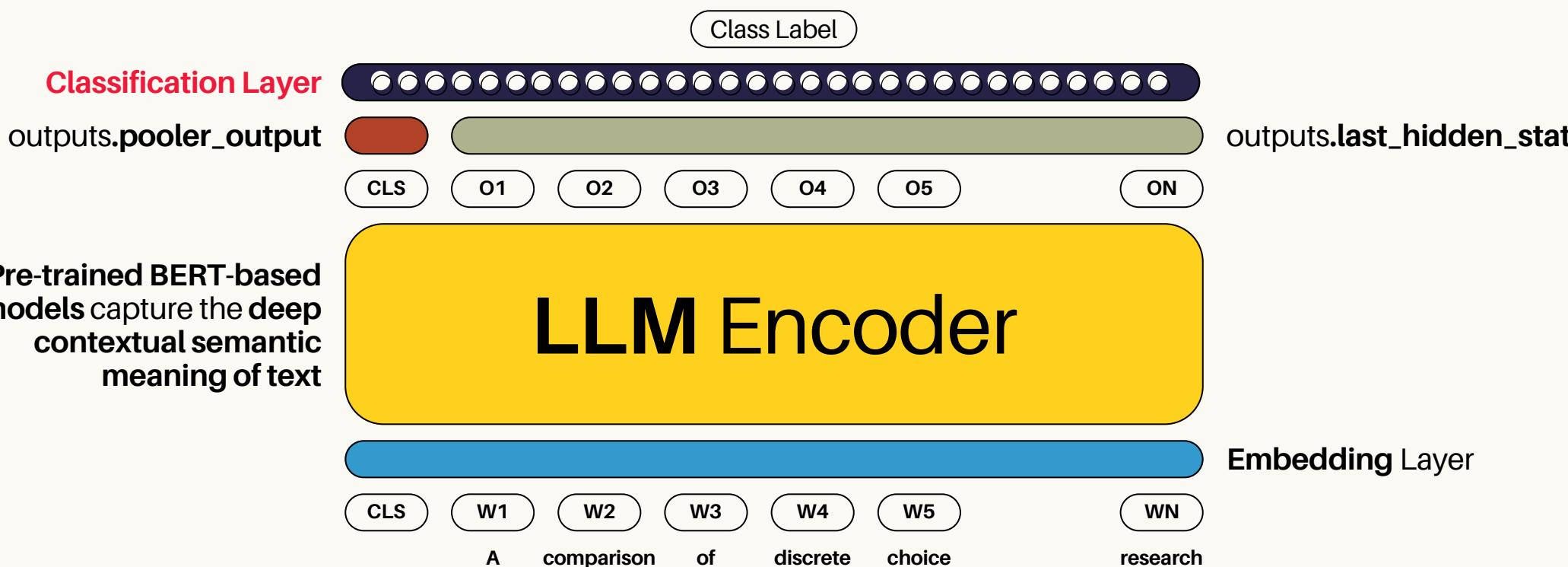


How to bring these two different worlds together into a single and innovative system?

Classification Layer on top

and hyper-parameters tuning to recognize the Patient Preference Studies pattern.

So train and test,
why not?

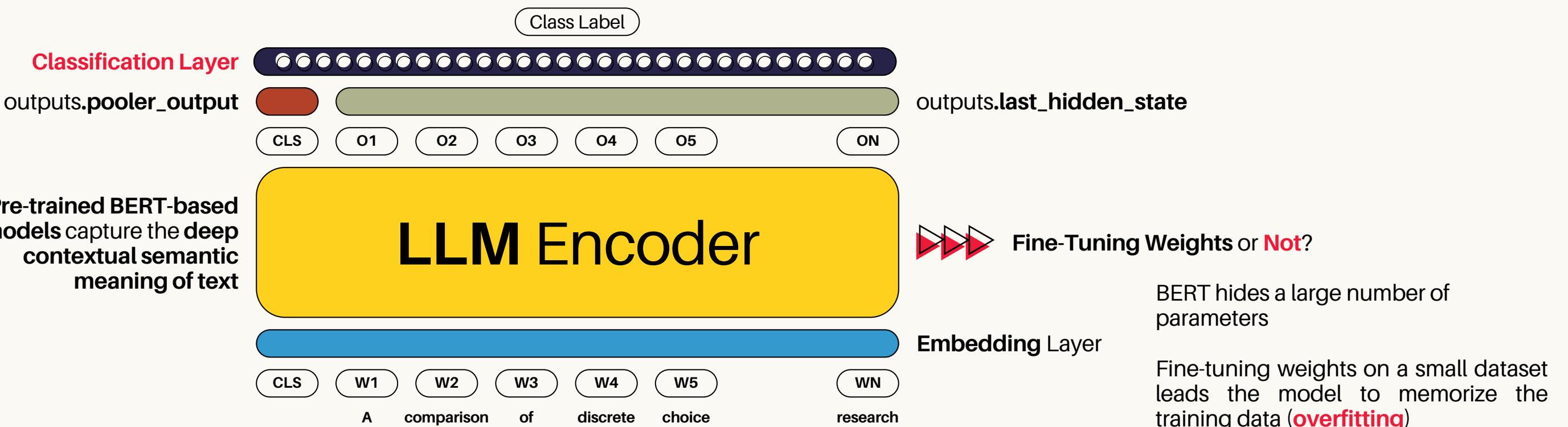


A comparison of discrete choice experiment (DCE) and swing-weighting (SW) methods assesses diabetic **patient preferences** for glucose-monitoring devices. The analysis highlights critical attributes such as ease of use, accuracy, and cost, revealing differences in attribute prioritization and trade-offs. Insights from this evaluation inform the selection of preference-elicitation techniques in patient-centered healthcare research.

LLM for Text Classification

Dataset Size: 1.200 Papers

with fewer than 200 Patient Preference Studies,
not enough data to train a LLM from scratch,
not enough to fine-tune a pre-trained LLM
without **overfitting**



A comparison of discrete choice experiment (DCE) and swing-weighting (SW) methods assesses diabetic **patient preferences** for glucose-monitoring devices. The analysis highlights critical attributes such as ease of use, accuracy, and cost, revealing differences in attribute prioritization and trade-offs. Insights from this evaluation inform the selection of preference-elicitation techniques in patient-centered healthcare research.

The Hybrid Solution

BERT-Based Model with frozen weights to encode

medical texts into machine-readable format

Train a Machine Learning Classifier

on top to draw and maximize the

margin between classes

Normalize and Train

```
pipeline = Pipeline([
    ('scaler', StandardScaler()),
    ('classifier', classifier)
])
pipeline.fit(x_train, y_train)
```

Freeze Weights

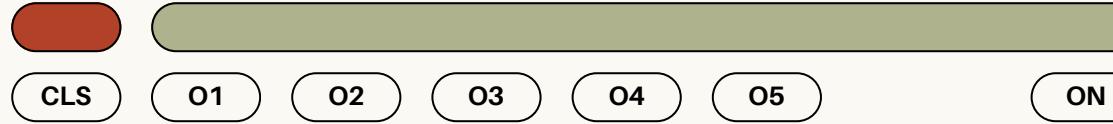
```
with torch.no_grad():
    output = model(**inputs)
```

Class Label

Machine Learning Classifier



outputs.pooler_output



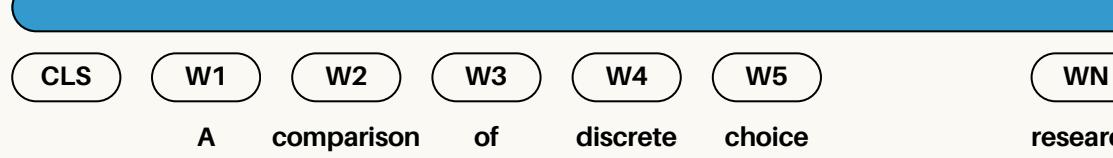
outputs.last_hidden_state



LLM Encoder



Embedding Layer



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What to evaluate?

The worst outcome:
researchers lose papers on PPS

The target outcome:
researchers retrieve papers on PPS alone



remove noise

True Positive Rate (TPR)

$$\frac{\text{TP}}{\text{TP} + \text{FN}}$$

for Class 1 (PPS)

The probability that an actual positive tests positive

Positive Predictive Value (PPV)

$$\frac{\text{TP}}{\text{TP} + \text{FP}}$$

for Class 1 (PPS)

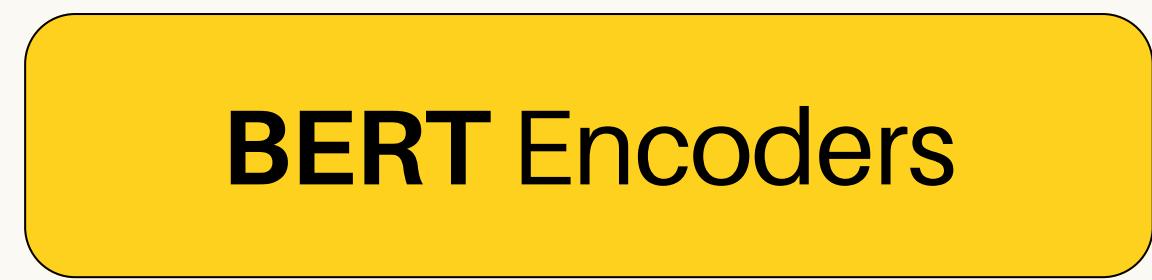
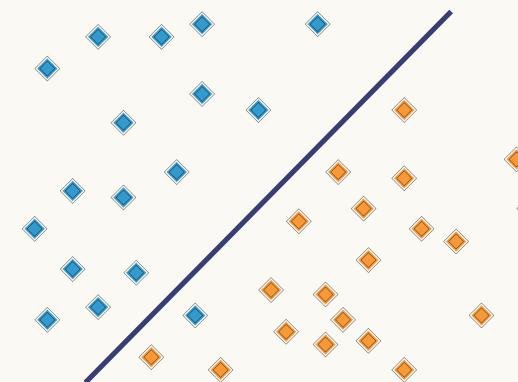
The probability that a positive test matches an actual positive

The Experimental Setup

The head-to-head comparison of diabetic patient preferences for glucose-monitoring devices.

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The head-to-head comparison of diabetic patient preferences for glucose-monitoring devices.

A comparison of discrete choice experiment (DCE) and swing-weighting...



Hugging Face

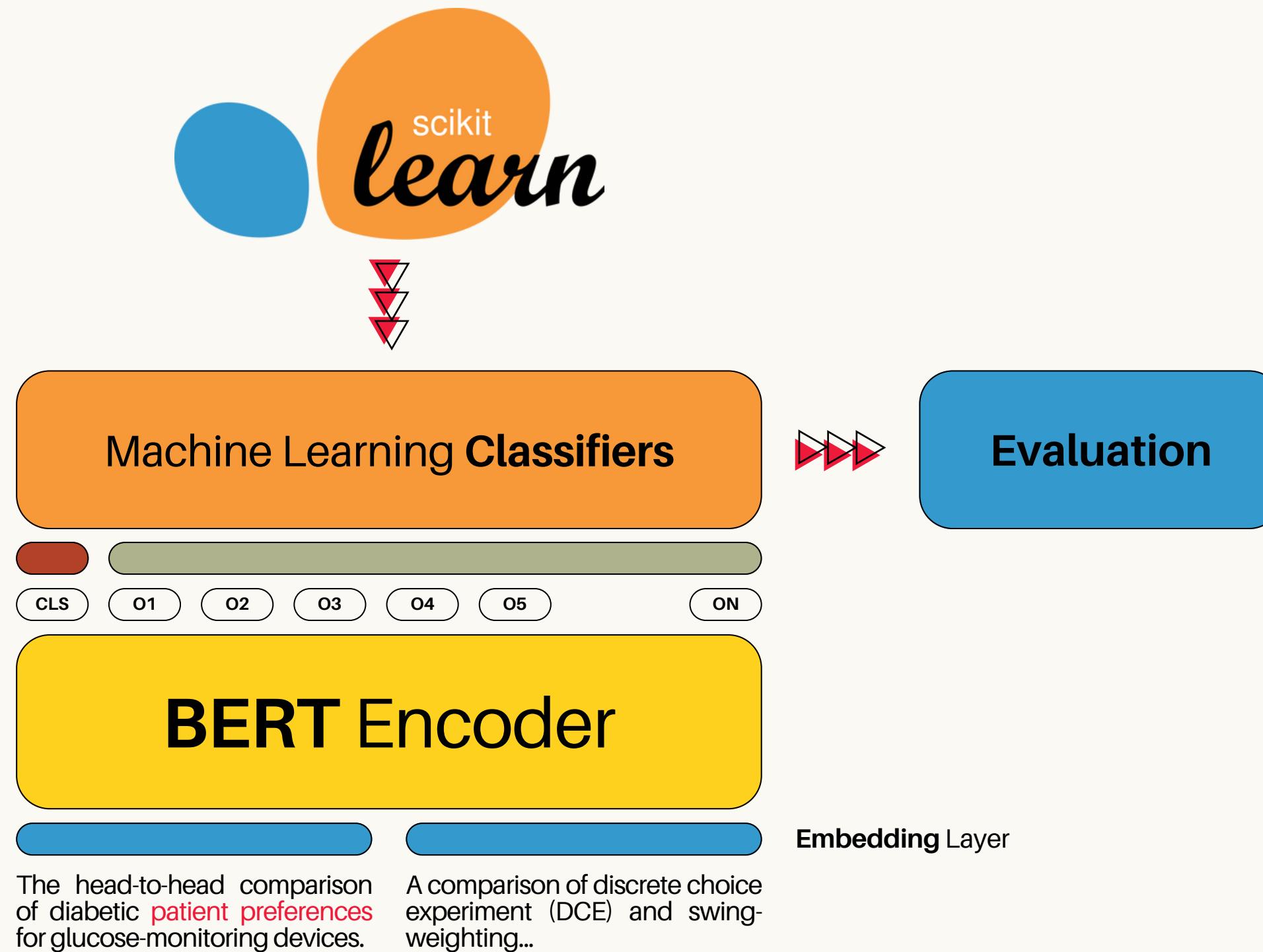
Embedding Layer

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The head-to-head comparison of diabetic **patient preferences** for glucose-monitoring devices.

A comparison of discrete choice experiment (DCE) and swing-weighting...

Embedding Layer

The Top Three



Hugging Face

BERT Models

BERT-Base Model	F2-PPS	F1-PPS	TPR	PPV
pubmedbert-base-embeddings	0.833	0.836	0.831	0.844
BiomedNLP-BiomedBERT-base-uncased-abstract	0.821	0.821	0.822	0.825
S-PubMedBert-MS-MARCO	0.816	0.81	0.819	0.803



pubmedbert-base-embeddings

Classifier Model	F2-PPS	F1-PPS	TPR	PPV
k-Nearest Neighbors	0.895	0.82	0.953	0.719
Deep Neural Network	0.893	0.892	0.894	0.882
Logistic Regression	0.891	0.891	0.889	0.884

BiomedNLP-BiomedBERT-base-uncased-abstract

Classifier Model	F2-PPS	F1-PPS	TPR	PPV
SVM (RBF Kernel)	0.906	0.91	0.903	0.918
Neural Network	0.9	0.906	0.896	0.914
Logistic Regression	0.899	0.9	0.898	0.899

The Top Parameters

k-Nearest Neighbours

Parameters	Values
K (neighbours)	3, 5, 8, 13, 21
Metric	euclidean, manhattan, minkowski



K-Nearest Neighbors Parameters	F2-PPS	F1-PPS	TPR	PPV
K = 5, Metric = euclidean	0.897	0.823	0.953	0.724

for `pubmedbert-base-embeddings`

High Recall Model

Support Vector Machine

Parameters	Values
C (penalty parameter)	1e0, 1e1, 1e2
Gamma (kernel coefficient)	1e-5, 1e-4, 1e-3, auto, scale

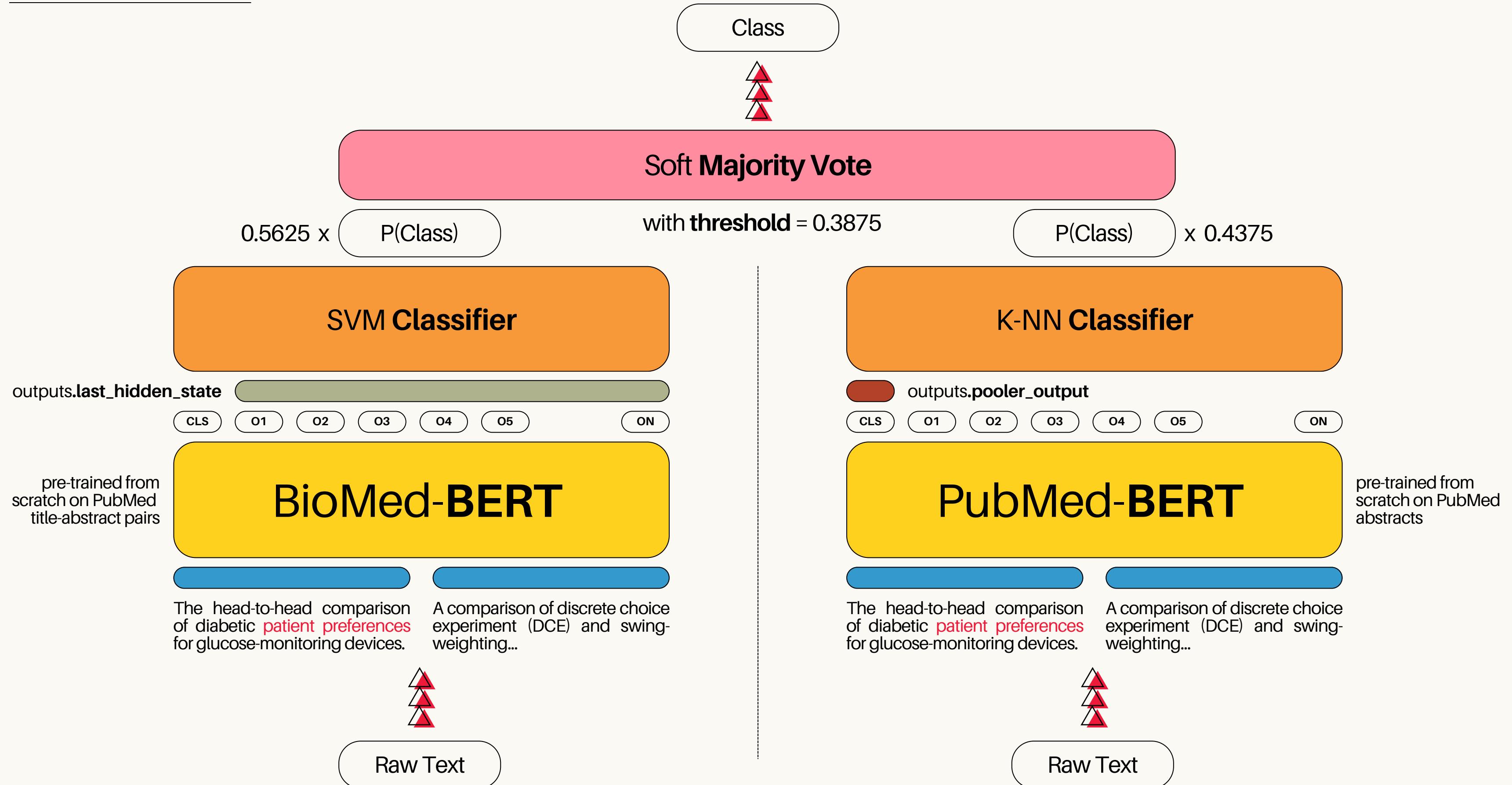


Support Vector Parameters	F2-PPS	F1-PPS	TPR	PPV
C = 1e1, Gamma = auto	0.901	0.921	0.888	0.957

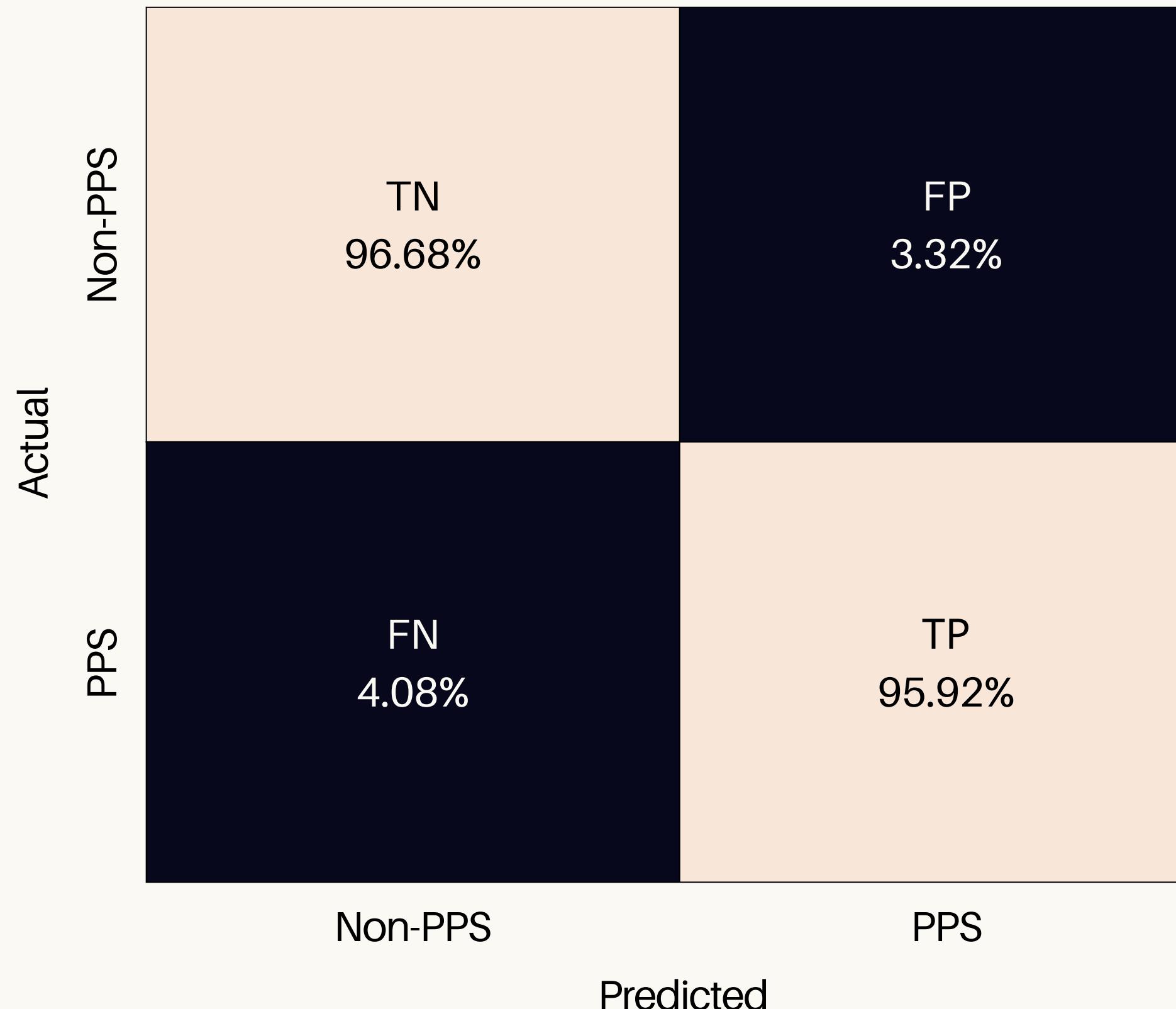
for `BiomedNLP-BiomedBERT-base-uncased-abstract`

High Precision Model

The Deliverable Model



The **Result**



The
model
**loses 5 PPS
out of 100 papers,
but removes more
than 95% of the noise**

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What's Next?



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The Multi-Label Text Classifier

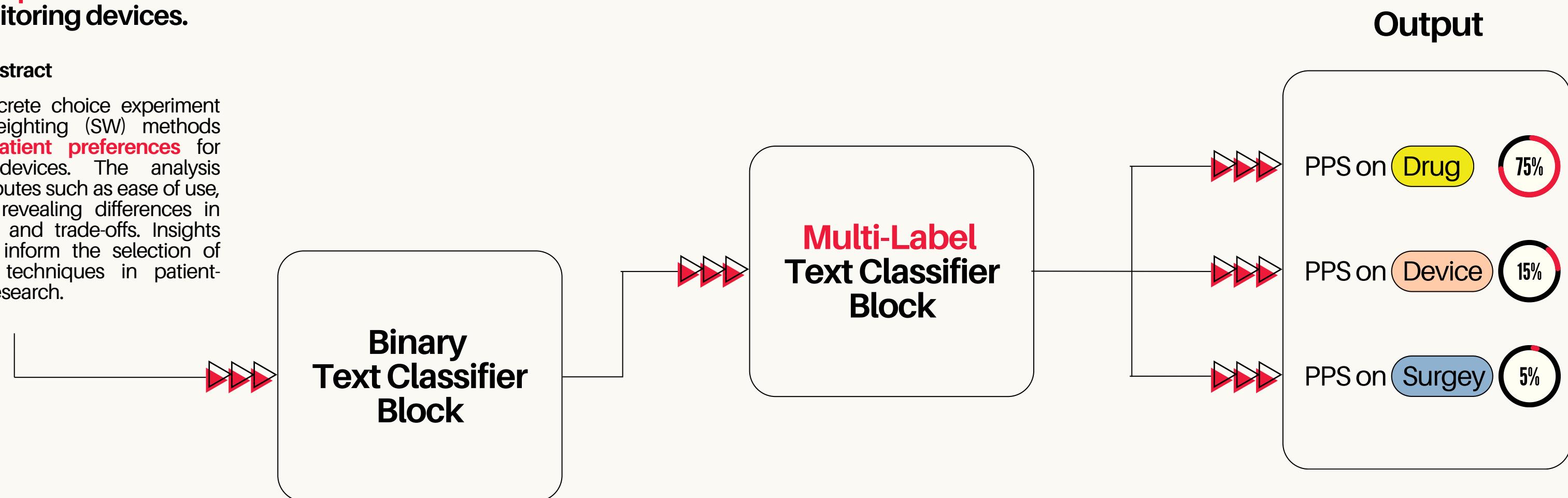


Multi-Label Text Classification

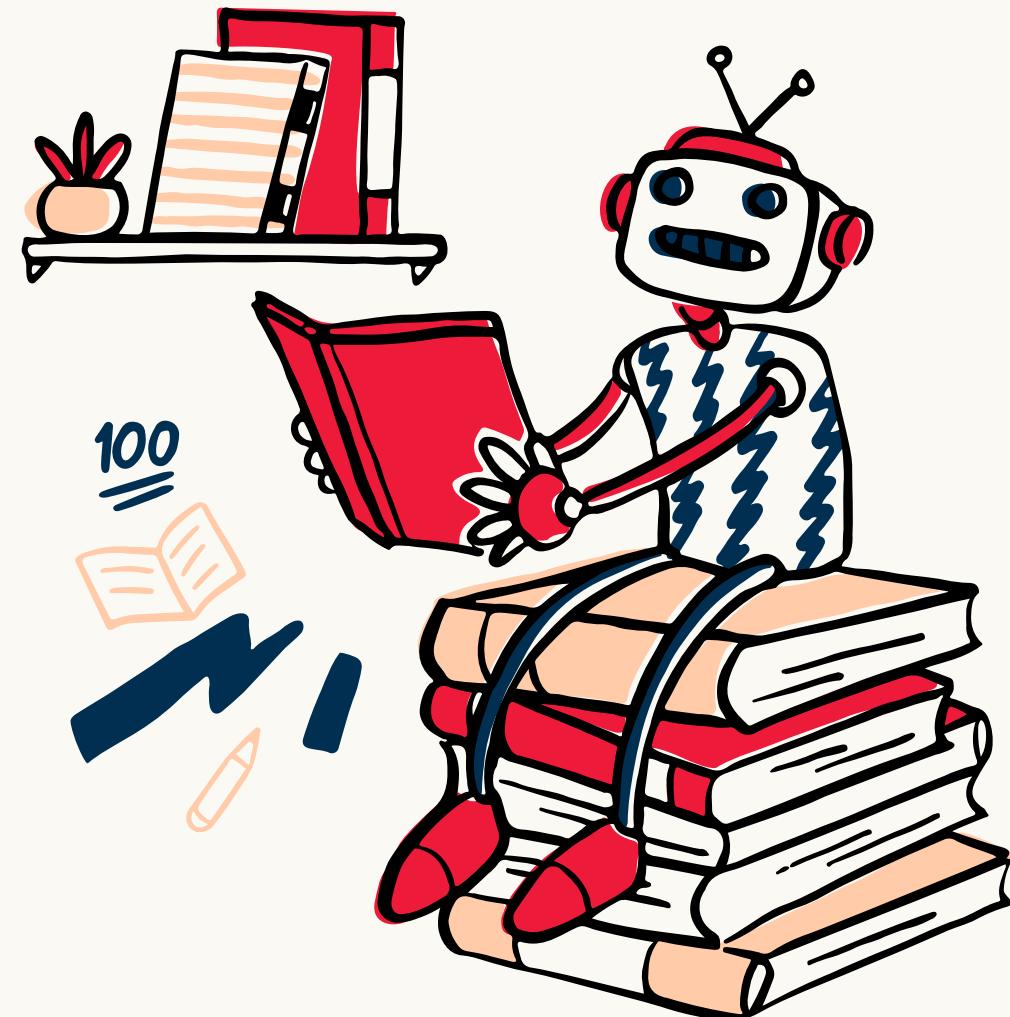
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The Multi-Label Classifier Methods



1. ML Dataset and Imbalance Analysis

- Mean imbalance ratio
- Coefficient of variation of IR
- Scumble index

2. Data-driven approach for model selection: comparison of most used MLC

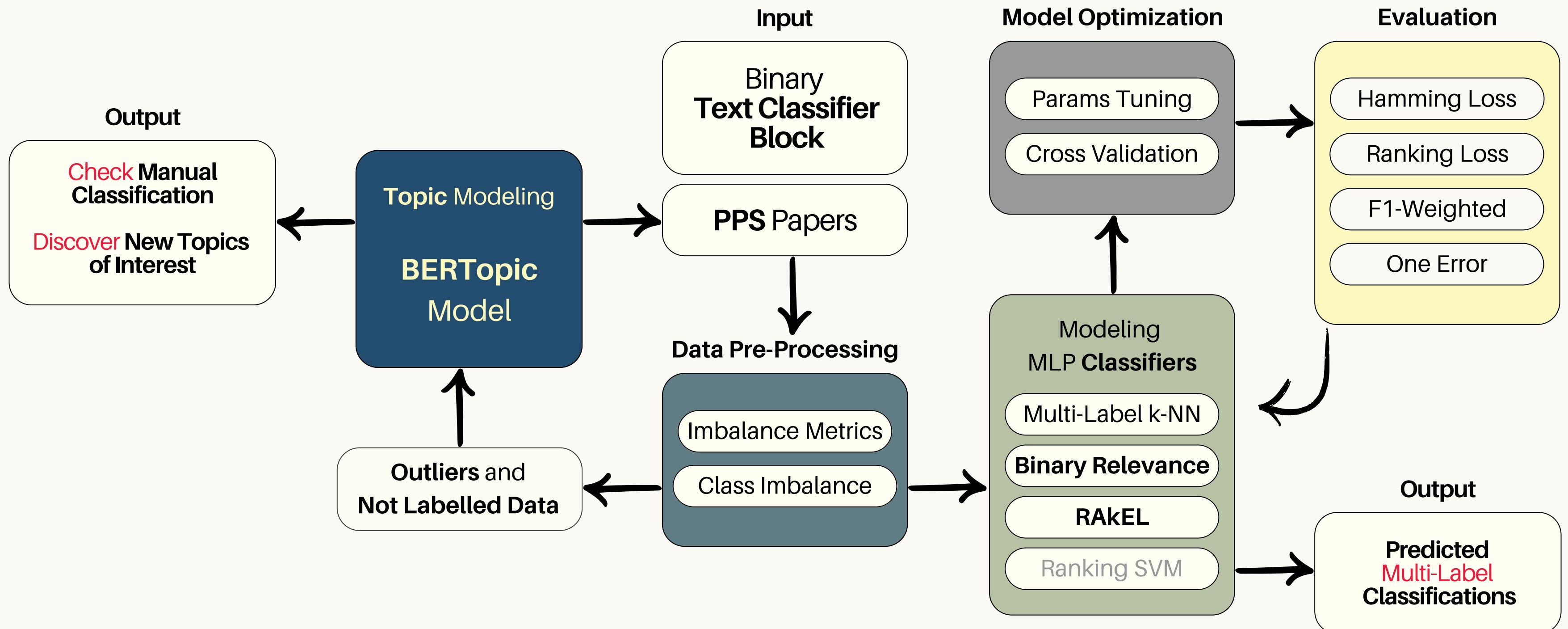
- Problem transformation (Binary relevance, RAkEL)
- Algorithm adaptation (multilabel kNN, VW ML kNN, ranking SVM)
- Ensemble models
- Cost sensitive (RAkEL)

3. Metrics for model comparison

- Hamming loss
- Ranking loss
- F1-score (micro, weighted)
- Coverage error
- One error

4. Best model selection

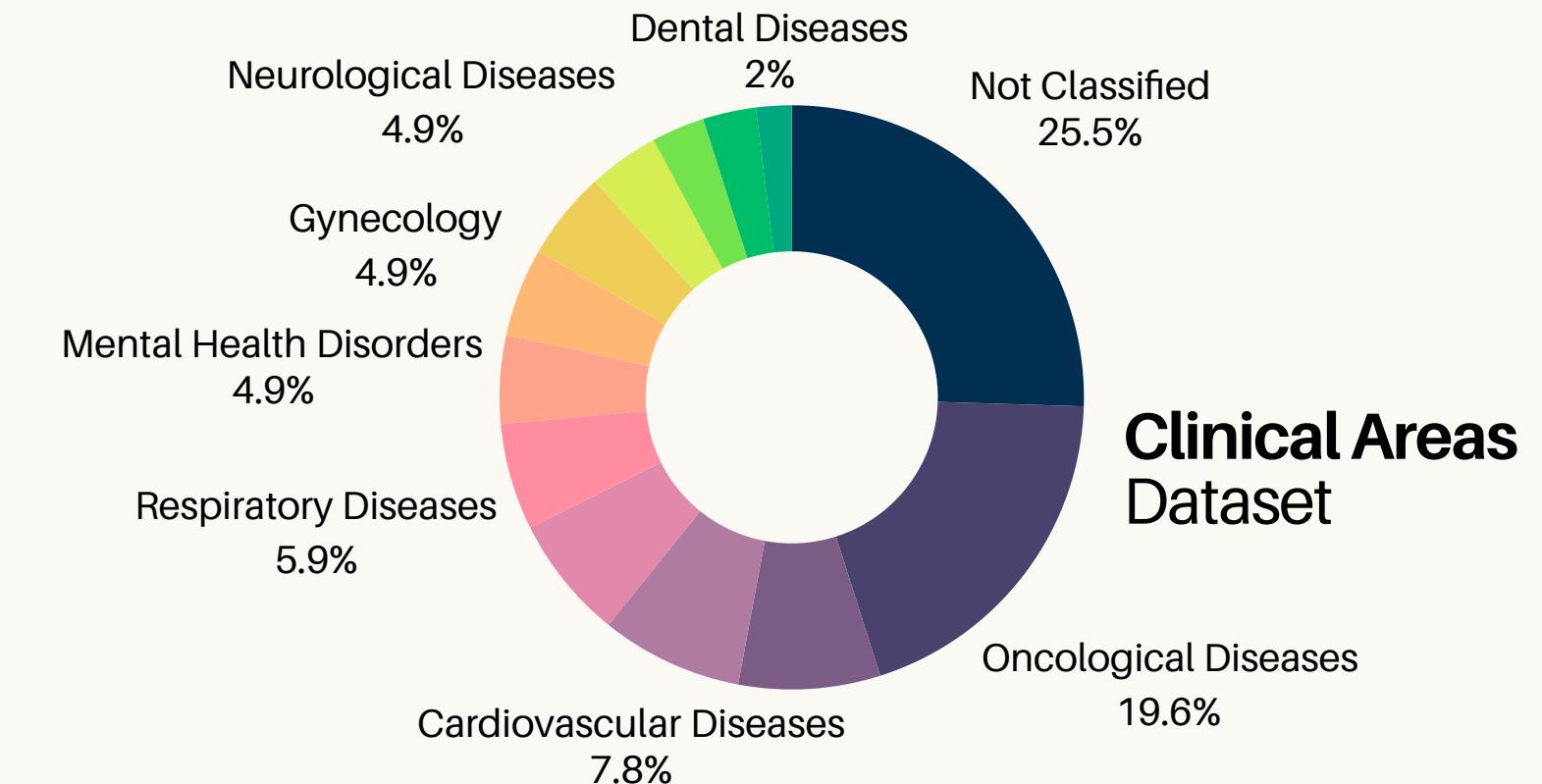
The Experimental Setup



Dataset Imbalance Metrics

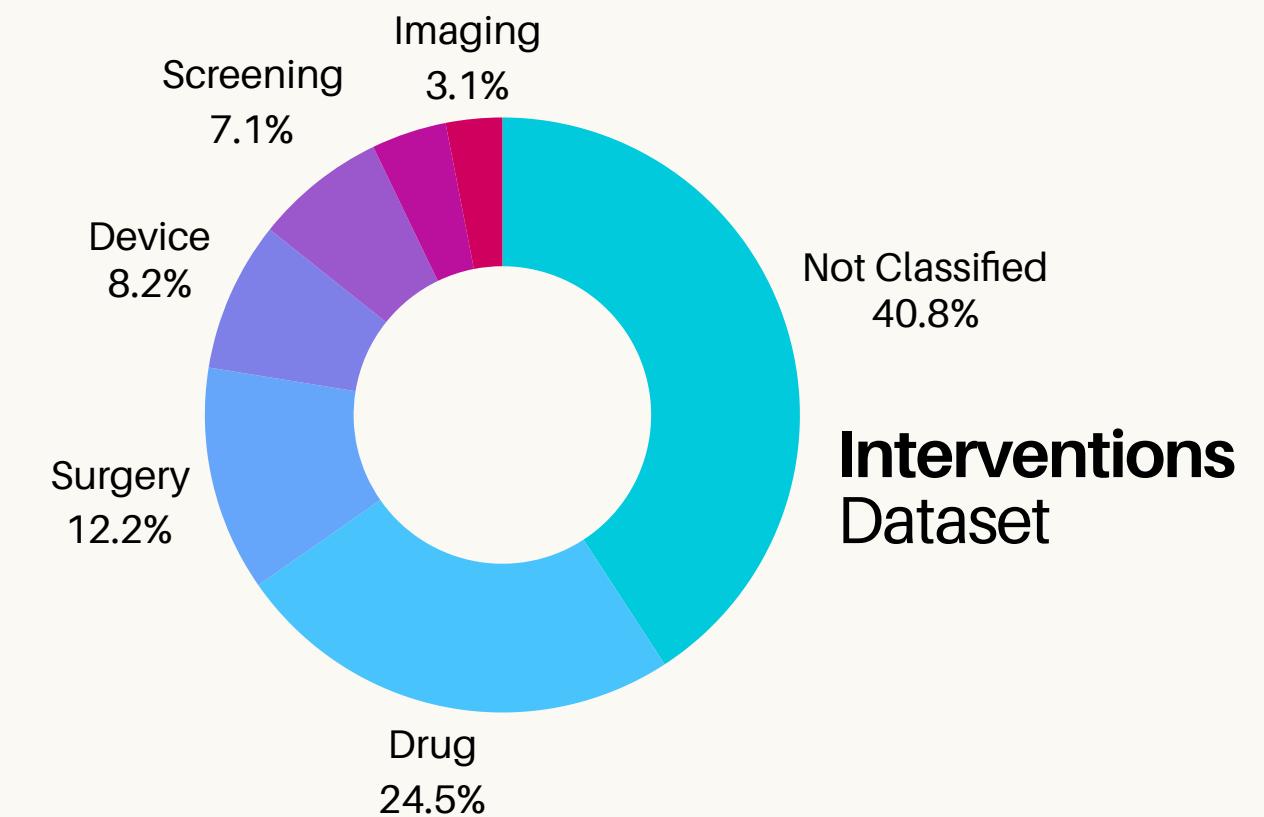
Dataset Imbalance

ML Dataset	Mean IR	Max IR	CVIR	Scumble
Clinical Areas	4,53	12,20	1,91	0,63
Interventions	4,63	8,82	2,37	0,44



Dataset Description

ML Dataset	# samples	Labels	Class Sets	Card	Dens	TCS
Clinical Areas	2192	12	88	0,87	0,07	13,19
Interventions	2192	6	23	0,63	0,11	11,12



The Experiments



Multi-Label k-Nearest Neighbours

Parameter	Values
K (neighbours)	3, 5, 7, 11
S (smooth)	10, 50, 100, 200



Dataset	K	s
Clinical Areas	11	1
Interventions	11	1

Value-Weighted Multi-Label k-Nearest Neighbours

Parameters	Values
K (neighbours)	1, 3, 5, 7, 11
a	0.3, 0.5, 0.7
b	0.3, 0.5, 0.7



Dataset	K	a	b
Clinical Areas	1	0.5	0.5
Interventions	1	0.3	0.3

The Experiments



RAkEL - Labelset: [2, 3, 4, 5, 6, 7]

Model	Parameter	Values
Gaussian NB	Smoothing	1e-9, 1e-8, 1e-7, 1e-6
Random Forest	N (estimators)	10, 50, 100, 200



Dataset	Labelset	s
Clinical Areas	6	1e-9
Interventions	3	1e-7

Dataset	Labelset	N
Clinical Areas	6	50
Interventions	7	200

Binary Relevance

Model	Parameter	Values
kNN	K (neighbours)	1, 3, 5, 7, 11
Multinomial NB	Alpha	0.3, 0.5, 0.7
SVC	Kernel	linear, rbf, sigmoid



Dataset	K
Clinical Areas	5
Interventions	11

Dataset	Alpha
Clinical Areas	0.3
Interventions	0.3

Dataset	Kernel	C
Clinical Areas	Linear	1e-2
Interventions	Linear	1e-2

The Multi-Label Classifier Results

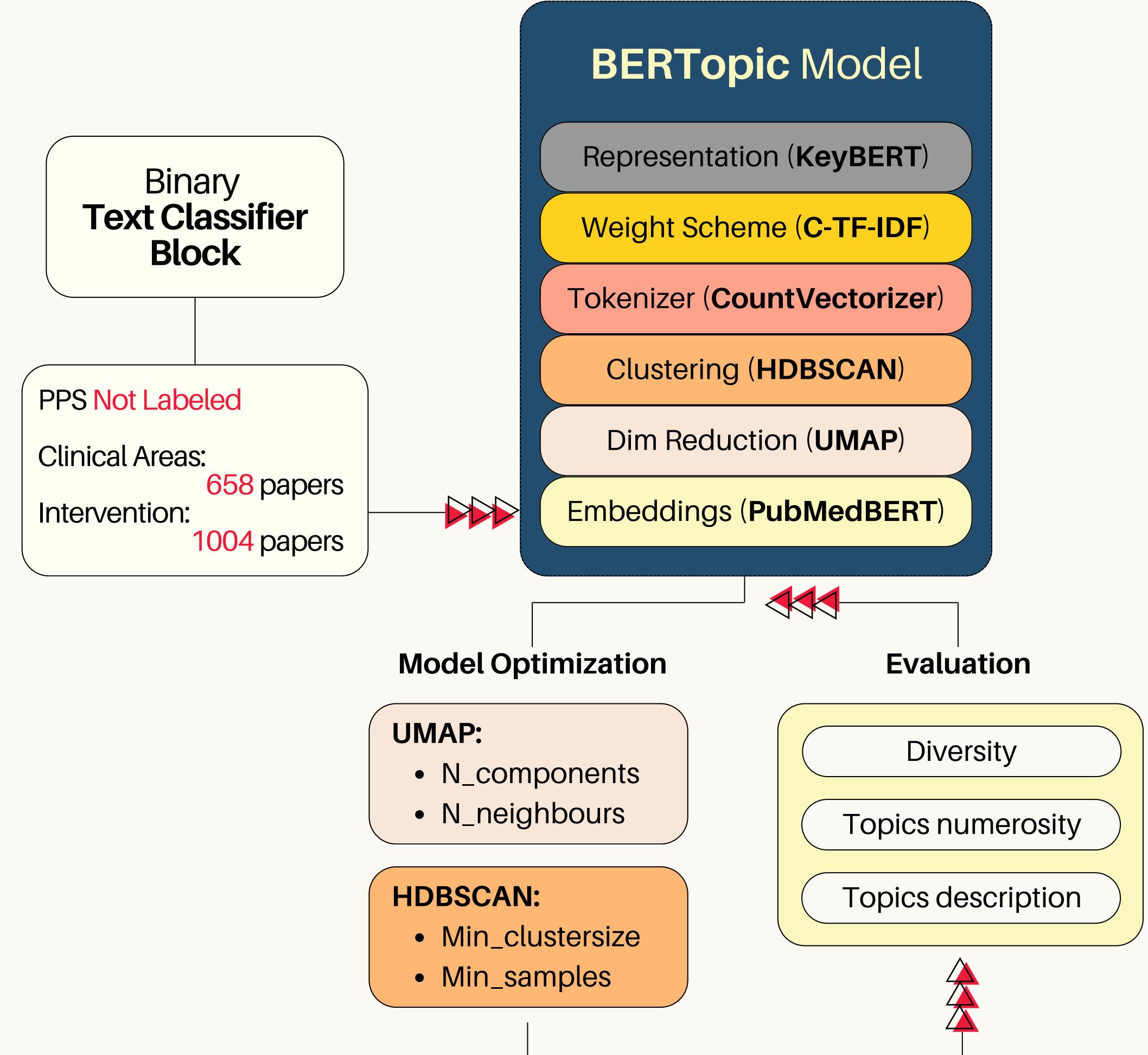
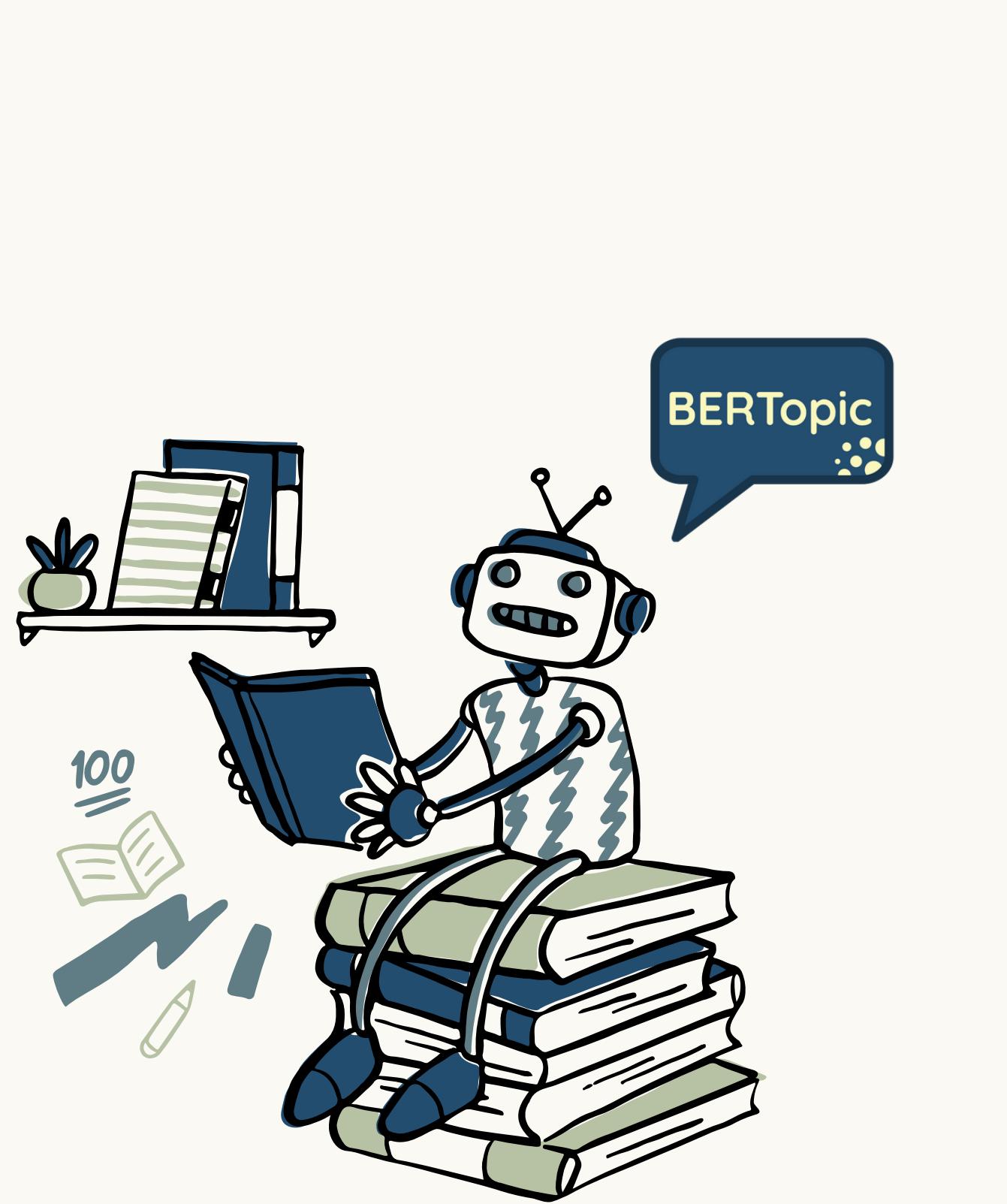
Clinical Areas

Model	F1-Micro	F1-Weighted	Hamming Loss	Ranking Loss	Coverage Error	One Error
ML kNN	0,757	0,748	0,030	0,076	1,714	0,614
VW MLkNN	0,750	0,747	0,034	0,171	2,877	0,470
BR (kNN)	0,778	0,771	0,027	0,200	3,216	0,479
BR (MultinomialNB)	0,704	0,720	0,047	0,032	1,242	0,409
BR (SVC)	0,823	0,815	0,022	0,158	2,735	0,437
RAkEL (Gaussian NB)	0,757	0,759	0,034	0,147	2,698	0,458
RAkEL (Random Forest)	0,426	0,361	0,050	0,510	6,660	0,742

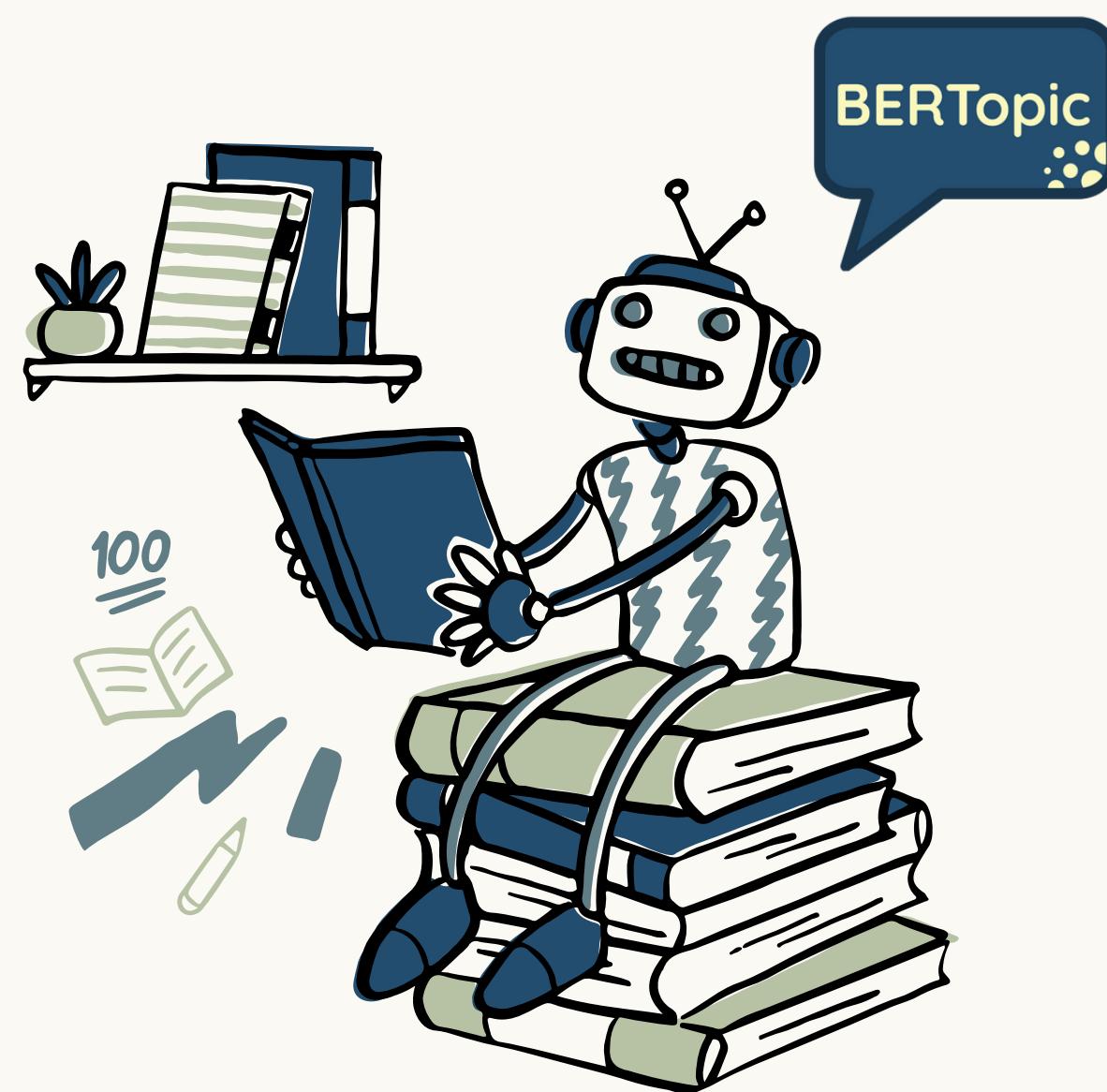
Interventions

Model	F1-Micro	F1-Weighted	Hamming Loss	Ranking Loss	Coverage Error	One Error
ML kNN	0,644	0,633	0,065	0,071	0,984	0,625
VW MLkNN	0,602	0,599	0,083	0,211	1,709	0,643
BR (kNN)	0,648	0,633	0,062	0,228	1,787	0,652
BR (MultinomialNB)	0,630	0,655	0,095	0,049	0,874	0,572
BR (SVC)	0,710	0,694	0,053	0,194	1,648	0,622
RAkEL (Gaussian NB)	0,661	0,675	0,080	0,145	1,396	0,606
RAkEL (Random Forest)	0,571	0,519	0,066	0,298	2,121	0,707

The BERTopic Experimental Setup



The BERTopic Results



Data	Topic	Count Papers	Suggested Topic	Label
CLINICAL AREAS	2	36	Psoriasis	Skin Diseases (NEW)
	9	21	Eye_cataract_macular	Visual System Diseases (NEW)
	15	16	Bladder_oab_urinary	Urinary Diseases (NEW)
INTERVENTIONS	13	23	Contraceptive_counseling	Drugs
	24	14	Research_testing_diagnonstic	Screening

The Conclusions



The **Binary Classifier** performs the first task (PPS identification): built on two ML models (SVC and KNN) on top of two BERT embeddings; a majority voting system drives the class prediction. The model runs with a TPR of **96%** and a PPV of **97%**.



The **Multi-Label Classifier** classifies papers of relevance (from the binary classifier) into different categories through a multi-label classification process.



The investigation explores various multilabel classifiers, identifying Binary Relevance with SVC as a classifier, ML kNN, and RAkEL with GaussianNB classifier as effective approaches. Data augmentation serves as the next step to enhance results and address low-frequency label-sets.



The **BERTopic** study identifies new labels for the clinical areas categorization as well as further items for specific labels in the intervention one.

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Share

High-Precision Binary Classifier: Patient Preference Study or Not?



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Input Options

Select Input Method

TXT File (PubMed)

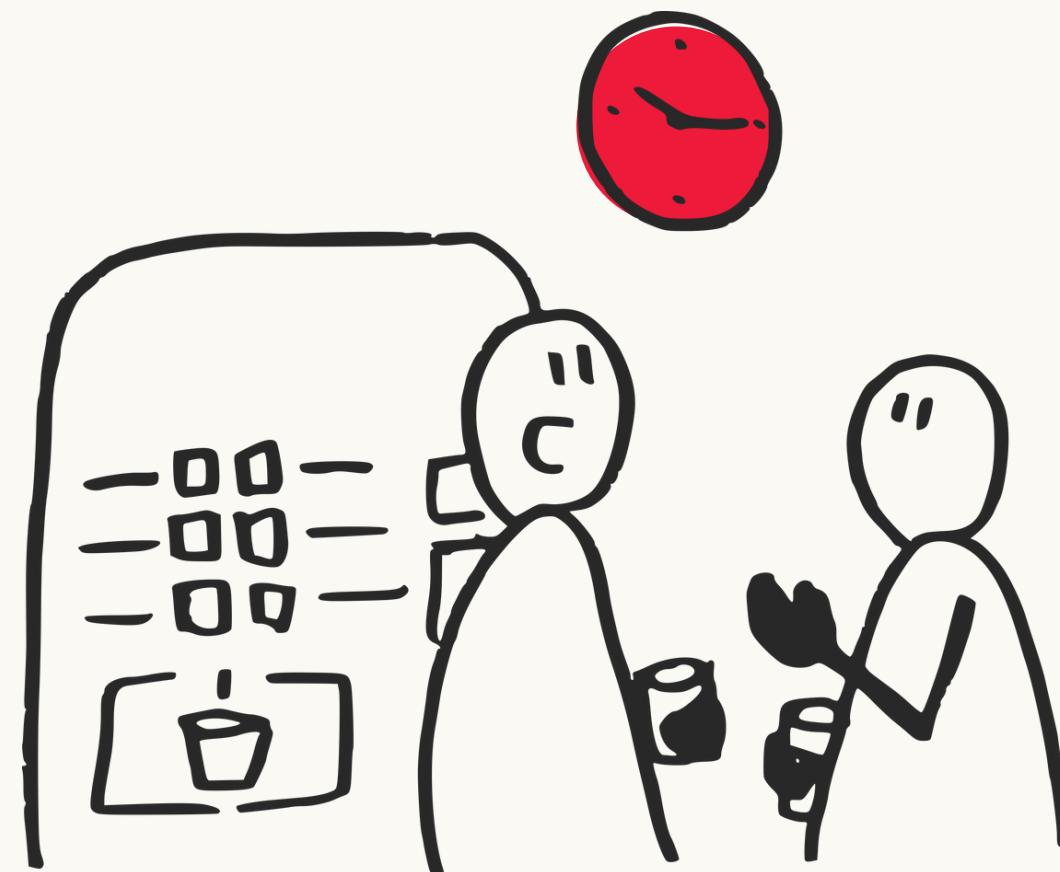
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3. Click the "Save" button at the top right of the results page.
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5. Click "Create File" to download the results in a `.txt` file.
6. Upload the `.txt` file here.

<https://pps-binary-classifier.streamlit.app>

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Thank You



Cesar Augusto Seminario Yrigoyen
Francesco Giuseppe Gillio



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