A Simple Machine Learning Framework to Aid Citation Screening in Systematic Reviews and Meta-Analyses of Aging and Longevity Research Studies

Marko Lalović March 10, 2021

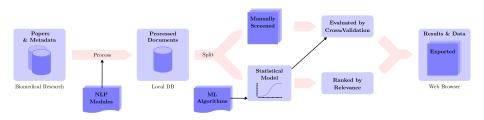


Figure 1: General overview of proposed framework.

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Background

- A systematic review typically addresses a specific clinical question by collecting and analyzing data from all the relevant and unbiased set of studies.
- Citation screening is the first yet tedious task of narrowing down the large set of citations retrieved via a broad database query to those relevant for the review.

Idea

Develop a machine learning framework to semi-automate citation screening in systematic reviews and meta-analyses, thereby reducing reviewers workload (and screening errors).

Related Work

- Wallace et al. developed a semi-automated citation screening algorithm for systematic reviews of biomedical literature.
- Bannach-Brown et al. described their approaches to aid citation screening for a systematic review of pre-clinical animal studies.
- Howard et al. deployed a general software system that automate the required methodologies called "SWIFT-Review".
- Przybyła et.al introduced a web-based software system called "RobotAnalyst".
- O'Mara-Eves et al. performed a systematic review of current approaches.
- To date, no use of any tools related to automating (or semi-automating) the screening process of systematic reviews or meta-analyses of aging and longevity research was reported.

Our Contribution

- Present a simple machine learning framework that can be used in the screening stage of systematic reviews or meta-analyses of aging and longevity research studies.
- Present our experimental results on dataset related to Dasatinib and Quercetin Senolytic Therapy Risk-Benefit Analysis¹(D&Q Analysis).

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¹The analysis is part of "Rejuvenation Now" non-profit initiative that: "seeks to continuously identify potential rejuvenation therapies and systematically evaluate their risks, benefits, and associated therapeutic protocols to create transparency" published by Forever Healthy Foundation

Outline

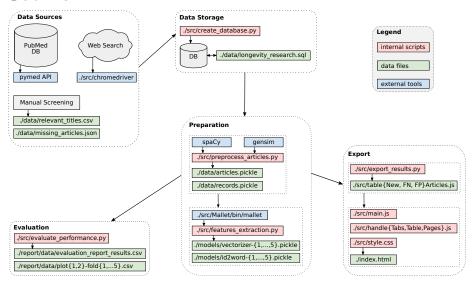


Figure 2: Technical Overview.

Features

Extracted features are based on:

- Provided list of search terms: dasatinib, senolytic, senescent, ...
- Possible publication types: case report, clinical trial, review, ...
- TF-IDF scores of terms: chronic myeloid, adverse event, tyrosine kinase, ...
- LDA model probabilities of belonging to topics:

Topic Terms

- trial, clinic, efficaci, report, safeti, assess, evalu, show, advers, ...
- 2. week, placebo, treatment, group, extract, symptom, score, patient, hypericum, ...
- 3. dasatinib, patient, case, treatment, report, chronic, leukemia, therapi, myeloid, ...
- 4. cvd, risk, cardiovascular, prevent, factor, profil, diseas, lipid, import, ...
- 5. flavonoid, anthocyanin, individu, adult, genistein, dietari, isoflavon, flavanon, intak, ...
- 6. sunitinib, sorafenib, target, imatinib, includ, cancer, erlotinib, anticanc, malign, ...
- 7. bone, marrow, chromosom, abnorm, deriv, prognost, signific, aberr, delet, ...
- 8. lifespan, elegan, longev, stress, effect, life, span, extend, increas, ...
- 9. muscl, smooth, vsmc, skelet, aortic, havsmc, resist, accordingli, folfiri, ...
- 10. quercetin, effect, cell, concentr, human, dose, depend, studi, increas, ...

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Model

- Used L1 regularized logistic regression model from scikit-learn.
- Fitted using Liblinear solver with balanced class weights.
- Placed more emphasis on recall by using recall scorer.

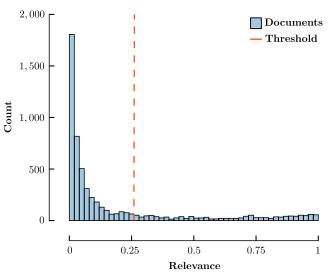
Definition (Relevance)

The model estimates the conditional probability, called *relevance score*, that a given document d is relevant given feature vector X^d :

 $Pr(d \text{ is relevant}|X^d)$

Threshold Selection

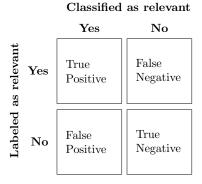
The selected cut-off threshold for D&Q Analysis was 0.26 where the binary classifier achieved 95% recall:



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Errors

There are four possible outcomes of comparing the classification result with human assigned label:



Performance Evaluation

Performance was assessed by:

Definition (Statistical measures)

Precision P is the fraction of documents labeled as relevant among documents classified as relevant:

$$P = \frac{TP}{TP + FP}$$

Recall R (also known as *sensitivity*) is the fraction of documents labeled as relevant that were also classified as relevant:

$$R = \frac{TP}{TP + FN}$$

Work Saved over Sampling WSS@R is the reduction of documents that need to be screened compared to a random ordering of the documents to achieve a level of recall R:

$$WSS@R = \frac{TN + FN}{N} - (1 - R)$$



Evaluation Results

Fold	Recall	Precision	PR-AUC	WSS@R
1	0.94	0.13	0.54	0.53
2	0.90	0.16	0.33	0.61
3	1.00	0.14	0.48	0.63
4	0.94	0.19	0.54	0.67
5	0.97	0.20	0.43	0.71
Mean	0.95 (0.03)	0.17(0.03)	0.46 (0.08)	0.63 (0.06)

Table 1: Summarized results of 5-fold cross-validation for D&Q Analysis

Evaluation of Results

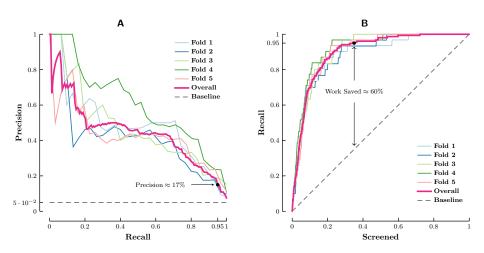


Figure 3: Visualized results of 5-fold cross-validation for D&Q Analysis

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Export

Results for Dasatinib and Quercetin Senolytic Therapy Risk-Benefit Analysis

		New Articles	False Negatives	False Positives				
	Te	able 2: Estimated Relevance for False Negatives un	il 17. April, 2020 included	in the Risk-Benefit Analysis but no	ot classified as relevant.			
Relevance	Date	Title		Abstract		Expand	URL	
0.3	22. März, 2013	Bioavailability of quercetin: problems and pr	omises Que	ercetin (QC) is a typical plant fla	vonoid, possesses d	+	pubmed	
0.29	1. Januar, 2005	Cytotoxicity of flavonoids toward cultured n	ormal hum The	cytotoxicity of flavonoids, inclu	iding apigenin, erio	+	pubmed	
0.25	6. Februar, 2016	Association Between BCR-ABL Tyrosine Ki	nase Inhib Imp	ortance: A phase 3 trial with por	natinib in patients	+	pubmed	
0.24	22. August, 2016	Targeting Pro-Inflammatory Cells in	Idiopathic Pulmonary F	ibrosis: a Human Trial (IPF)		Zhu) -	clinicaltrials	
The study team hypothesizes that intermittent (3 doses administered over 3 consecutive days in 3 consecutive weeks) and administration of combination Dasatinib (100 mg/d) + Quercetin (1250 mg/d) will be safe and well tolerated in patients with PF. Treatment with D+Q will result in reduced abundance of pro-inflammatory cells within subjects over baseline. Finally, the reduction in biomarkers of cellular pro-inflammatory state will be related to no change in functional and patient reported outcomes.								
0.22	19. Januar, 2017	Identification of cellular targets involved in o	ardiac fail Air	ns: The aims of the present study	were to evaluate t	+	pubmed	
0.22	3. Oktober, 2017	Short-term High Dose of Quercetin and Resv	eratrol Alt Bao	kground: Hyperglycemia-media	ted oxidative stress	+	pubmed	
0.22	14. Februar, 2018	BCR-ABL Tyrosine Kinase Inhibitors: Which	h Mechani Ima	tinib, the first-in-class BCR-AB	L tyrosine kinase in	+	pubmed	
0.19	1. Januar, 2008	Quercetin pharmacokinetics in humans	The	purpose of this study was to exa	amine the pharmac	+	pubmed	
0.18	6. Mai, 2009	Tyrosine kinase inhibitor-induced platelet dy	sfunction i Das	atinib is associated with increase	ed risk of bleeding	+	pubmed	
0.18	17. März, 2004	Quercetin, an over-the-counter supplement, or	auses neur A 2	2-month-old boy, who regularly	consumed the oral	+	pubmed	
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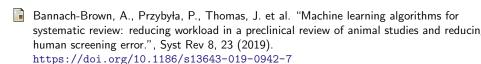
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Figure 4: Interactive tables of exported documents for D&Q Analysis.

Links

- Interactive tables of exported documents for D&Q Analysis: https://markolalovic.com/longevity-research-screening/
- Technical report: https://zenodo.org/record/4593957/files/zenodo.4593957.pdf
- Source code: https://github.com/markolalovic/longevity-research-screening/

References



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