Can Zero-shot LLMs save additional work in machine learning prioritised screening for systematic reviews?

Max Callaghan



October 30, 2024

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 - Wang et al. (2024) showing impressive results for the zero-shot setting

	Model	Setting	P	R	B-AC	F3	Suc	WSS
	Bi ^o RT BERT	$\begin{array}{c} { m Unc} \\ { m Cal}(0.95) \\ { m Cal}(1) \end{array}$	0.06 0.06 0.06	$0.95 \\ 0.92 \\ 0.97$	$0.61 \\ 0.64 \\ 0.60$	$0.30 \\ 0.31 \\ 0.29$	$0.74 \\ 0.50* \\ 0.82$	$0.26 \\ 0.34 \\ 0.23$
2017	Thritis	Unc Cal(0.95) Cal(1)	0.08 0.06* 0.05*	0.87 0.92* 0.99 *	0.72 0.69* 0.60*	0.35 0.32 0.28	0.26 0.52 0.96	$0.56 \\ 0.44 \\ 0.20$
CLEF-2017	13b-in	Unc Cal(0.95) Cal(1)	0.19 0.06* 0.05*	0.41 0.93 0.98	0.66 0.59* 0.53*	0.31 0.28 0.26	0.04 0.50* 0.88*	0.91 0.25 0.08
	Engenib	Unc Cal(0.95) Cal(1)	0.31 0.08 0.06	0.13 0.93* 0.97*	0.56 0.72 0.63	0.13 0.35* 0.30	0.00 0.52* 0.90 *	0.98 0.50 0.29
	Bio RE	Unc Cal(0.95) Cal(1)	0.06 0.07 0.06	0.97 0.91* 0.97	0.59 0.63 0.59	0.29 0.30 0.29	0.87 0.57* 0.87	0.19 0.33 0.21
2018	Thring	Unc Cal(0.95) Cal(1)	0.09 0.08* 0.06*	0.88 0.94* 0.99 *	0.75 0.71* 0.62*	0.37 0.35* 0.30	0.27 0.50 1.00	0.59 0.46 0.24
CLEF-2018	13b-in	Unc Cal(0.95) Cal(1)	0.26 0.06 0.05	0.36 0.94* 0.97	0.66 0.59* 0.53*	0.30 0.29 0.27	0.00 0.47* 0.80*	0.94 0.22 0.08
	Engenib	Unc Cal(0.95) Cal(1)	0.35 0.09* 0.06	0.12 0.94* 0.99*	0.54 0.75 0.64	0.12 0.38* 0.32*	0.00 0.50* 0.93*	0.95 0.54 0.28
	BERT	Unc Cal(0.95) Cal(1)	0.07 0.08 0.08	0.99 0.89 0.91	0.58 0.59 0.59	0.30 0.26 0.27	0.88 0.50 0.62	0.18 0.27 0.25
119-dta	Thring	Unc Cal(0.95) Cal(1)	0.09 0.10* 0.08*	0.92 0.91* 0.97*	0.71 0.71* 0.66	0.35 0.34 0.32	0.62 0.50 0.75	0.49 0.50 0.34
CLEF-2019-dta	13brins	Unc Cal(0.95) Cal(1)	0.19 0.08 0.07	0.49 0.95 0.99	0.69 0.56 0.51	0.32 0.29 0.28	0.00 0.50* 0.88*	0.87 0.16 0.03
· .	agemio	Unc Cal(0.95)	0.31	0.21 0.91	0.59 0.73	0.19 0.34*	0.00 0.50*	0.96



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Table 2
Performance of GPT 4o mini in screening titles and abstracts against a human reviewer's

Data set	k-value	Precision	F1-score	Kappa	PABAK	AUC-ROC	Recall
Hall	0.0	0.575	0.672	0.667	0.662	0.900	0.808
Hall	0.5	0.649	0.670	0.666	0.663	0.844	0.692
Hall	1.0	0.653	0.644	0.640	0.636	0.815	0.635
PTSD	0.0	0.129	0.226	0.216	0.210	0.924	0.895
PTSD	0.5	0.161	0.270	0.260	0.255	0.892	0.816
PTSD	1.0	0.176	0.286	0.277	0.271	0.868	0.763
Virus	0.0	0.339	0.485	0.446	0.417	0.882	0.851
Virus	0.5	0.375	0.508	0.473	0.446	0.861	0.789
Virus	1.0	0.436	0.557	0.527	0.502	0.860	0.772

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- There are a lot of bad papers, which will continue to proliferate, each tweaking some parameters to achieve a high performance that cannot be generalised

Table 2

Performance of GPT 40 mini in screening titles and abstracts against a human reviewer's good truth using different k-value

Data set	k-value	Precision	F1-score	Kappa	PABAK	AUC-ROC	Recall
Hall	0.0	0.575	0.672	0.667	0.662	0.900	0.808
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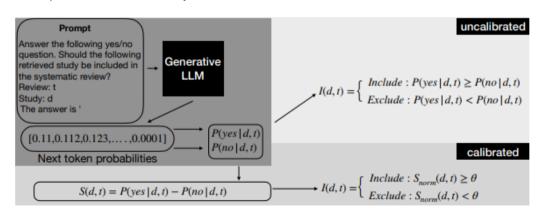
Realistic and useful evaluations of LLMs for screening

Despite the hype, LLMs might still be helpful, but we need to know

- How do LLMs compare against commonly used (and much cheaper!) baselines?
- How could they safely be used in the real world (Hint: with stopping criteria and prioritised screening)

Zero-shot Generative Large Language Models for Systematic Review Screening Automation

Wang et al. (2024) propose a method to extract probability-like scores from LLMs for inclusion/exclusion decisions in a systematic review



```
1 def binary_probs(tokenizer, model, prompt, no_words=['no'], yes_words=['yes'], return_all
       =False):
      device = 'cuda' if torch.cuda.is available() else 'cpu'
2
      encoded text = tokenizer(prompt, return tensors="pt").to(device)
3
      #1. step to get the logits of the next token
      with torch inference mode():
5
6
          outputs = model(**encoded text)
7
      next token logits = outputs.logits[0, -1, :]
8
Q
      # 2. step to convert the logits to probabilities
10
      next_token_probs = torch.softmax(next_token_logits, -1)
11
13
      topk next tokens = torch.topk(next token probs. 50)
      tokens = [tokenizer.decode(x).strip().lower() for x in topk next tokens.indices]
14
      p = topk next tokens.values
15
16
      df = pd.DataFrame.from_dict({'t': tokens,'p': p.cpu()})
      y = df[df['t'].isin(yes_words)]['p'].sum()
18
      n = df[df['t'].isin(no words)]['p'].sum()
19
20
21
      if return all:
          return df.groupby('t').sum().reset_index().sort_values('p', ascending=False).
22
       reset_index(drop=True)
      return v-n. v+n
23
```

Result: <s>[INST] «SYS» You are a systematic review helper tasked with finding out whether a study is relevant to the review Drug Class Review: Pharmacologic Treatments for Attention Deficit Hyperactivity Disorder Answer 'yes' if the study is relevant, or 'no' if not «/SYS» Study: Diuretics and beta-blockers do not have adverse effects at 1 year on plasma lipid and lipoprotein profiles in men with

htudy: Diuretics and beta-blockers do not have adverse effect at 1 year on plasma lipid and lipoprotein profiles in men with hypertension. Department of Veterans Affairs Cooperative Study Group on Antihypertensive Agents.

Concern based on the reported short-term adverse effects of antihypertensive agents on plasma lipid and lipoprotein profiles (PLPPs) has complicated the therapy for hypertension.

Should the study be included? Answer yes or no. [/INST] No, this study should not be included in the drug class review for Pharmacologic Treatments for Attention Deficit Hyperactivity Disorder. The study is focused on the effects of diuretics and beta-blockers on plasma lipid and lipoprotein profiles in men with hypertension, and does not directly relate to the treatment

of Attention Deficit Hyperactivity Disorder (ADHD). The review is focused on pharmacologic treatments for ADHD, and the study does not address any of the included drug classes or outcomes of interest for the review.



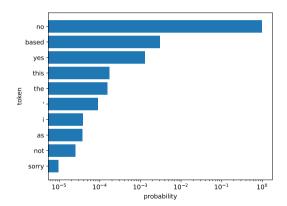
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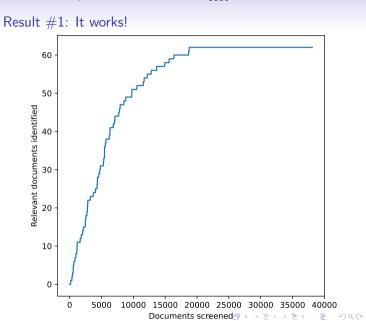
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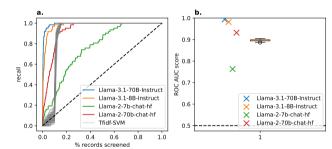
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- We turn these predictions into a prioritised list of documents to be screened

 The method produces a ranking that identifies all relevant documents before all documents have been screened



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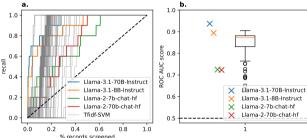
Brouwer_2019 (N=38114, p=0.2%)
Psychological theories of depressive relapse and recurrence: A systematic review and meta-analysis of prospective studies



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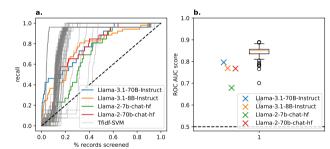
Bos 2018 (N=4878, p=0.2%)
Cerebral small vessel disease and the risk of dementia: A systematic review and meta-analysis of population-based evidence

a. b.



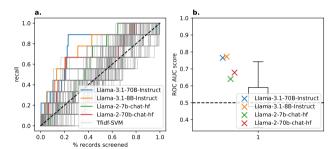
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- ROC AUC scores offer one way to compare rankings

Appenzeller-Herzog_2019 (N=2873, p=0.9%)
Comparative effectiveness of common therapies for Wilson disease: A systematic review and meta-analysis of controlled studies



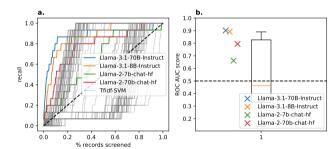
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Chou_2004 (N=1630, p=0.6%)
Comparative efficacy and safety of skeletal muscle relaxants for spasticity and
musculoskeletal conditions: a systematic review



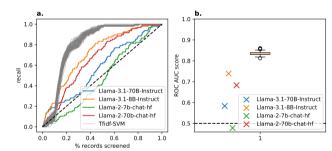
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Donners_2021 (N=258, p=5.8%)
Pharmacokinetics and Associated Efficacy of Emicizumab in Humans: A Systematic
Review

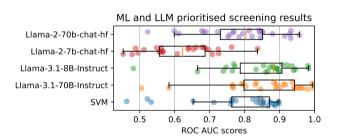


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Jeyaraman_2020 (N=1175, p=8.2%)
Does the Source of Mesenchymal Stem Cell Have an Effect in the Management of
Osteoarthritis of the Knee? Meta-Analysis of Randomized Controlled Trials



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Stopping criteria

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- Stopping when $p<\alpha$ gives a recall > target in the α^{th} percentile, in 97/100 settings

r target	model confidence	SVM	Llama-2-7b	Llama-2-70b	Llama-3.1-8B	Llama-3.1-70B
0.80	0.50	0.938	0.758	0.893	0.911	0.911
	0.80	0.924	0.797	0.866	0.845	0.889
	0.90	0.900	0.818	0.872	0.836	0.876
	0.95	0.867	0.806	0.888	0.878	0.878
	0.99	0.811	0.838	0.865	0.876	0.886
0.90	0.50	0.963	0.905	0.936	0.957	0.954
	0.80	0.957	0.900	0.958	0.961	0.933
	0.90	0.947	0.937	0.959	0.953	0.939
	0.95	0.933	0.943	0.944	0.947	0.939
	0.99	0.928	0.941	0.947	0.926	0.958
0.95	0.50	0.979	0.947	1.000	0.981	0.987
	0.80	0.974	0.973	0.976	0.975	0.979
	0.90	0.973	0.968	0.981	0.978	0.980
	0.95	0.964	0.973	0.982	0.976	0.968
	0.99	0.973	0.975	0.986	0.983	0.975
0.99	0.50	1.000	1.000	1.000	1.000	1.000
	0.80	0.994	1.000	1.000	1.000	0.991
	0.90	0.993	0.996	1.000	0.999	0.996
	0.95	0.993	0.997	1.000	0.999	0.996
	0.99	0.996	0.992	0.999	0.997	0.996

Stopping criteria

- Next, we calculate scores for our stopping criteria with a range of different recall targets
- Stopping when $p<\alpha$ gives a recall > target in the α^{th} percentile, in 97/100 settings
- In most settings, total work savings were greater for SVMs

r target	model confidence	SVM	Llama-2-7b	Llama-2-70b	Llama-3.1-8B	Llama-3.1-70B
0.80	0.50	126,898	100,248	121,777	127,401	132,808
	0.80	113,708	64,568	97,147	111,411	114,748
	0.90	105,053	57,678	90,327	101,871	106,668
	0.95	98,503	48,608	83,338	94,921	101,288
	0.99	87,803	37,838	71,397	79,471	85,058
0.90	0.50	114,988	65,778	99,217	112,011	115,498
	0.80	93,923	42,768	79,807	83,661	97,438
	0.90	84,658	34,158	65,007	76,701	82,268
	0.95	76,483	28,748	59,447	69,251	73,108
	0.99	62,350	23,118	49,790	55,091	59,088
0.95	0.50	98,258	44,408	82,907	94,731	101,318
	0.80	74,098	27,238	58,587	64,281	68,678
	0.90	63,245	21,738	44,470	53,091	57,388
	0.95	54,860	18,781	37,370	45,954	48,671
	0.99	41,676	13,545	27,171	34,855	35,592
0.99	0.50	63,573	22,638	42,897	54,811	55,378
	0.80	31,323	11,158	20,597	29,191	30,398
	0.90	21,160	5,825	13,480	13,641	18,478
	0.95	16,229	4,065	8,837	9,435	11,938
	0.99	7,980	1,985	4,355	4,934	6,625

Table: absolute work savings across datasets (N records=169,288)

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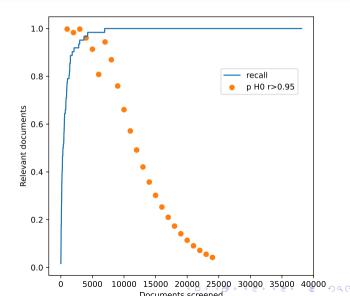
Note that although the LLM implementation is simplistic, so is the comparator

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- They do not have magic properties that fix problems of traditional ML, they are simply likely to make them worse
- We need to be on our guard!
- Do not forget stopping criteria!
- But they have some advantages, such as the fact that we only need to calculate scores once!



- Callaghan, M. and Müller-Hansen, F. (2020). Statistical Stopping Criteria for Automated Screening in Systematic Reviews. Systematic Reviews.
- De Bruin, J., Ma. Y., Ferdinands, G., Teijema, J., and Van de Schoot, R. (2023). SYNERGY Open machine learning dataset on study selection in systematic reviews.
- Wang, S., Scells, H., Zhuang, S., Potthast, M., Koopman, B., and Zuccon, G. (2024). Zero-shot Generative Large Language Models for Systematic Review Screening Automation.
- Xia, Z., Ye, J., Hu, B., Qiang, Q., and Debnath, R. (2024). LLMscreen: A Python Package for Systematic Review Screening of Scientific Texts Using Prompt Engineering.