Domain-Specific NLP

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Domain-Specific Models

Domain-Specific Models

Pretrained (Large) Language Models are trained on content crawled over the internet, books, reports and news papers and are, hence **are open-domain**.

A textual domain is the distribution over language characterizing a given topic or genre [1].

- You are more likely to see the word "integer" in computer science than in news papers.
- An (L)LM will be more perplex to the word "integer" even though the input comes from a StackOverflow post.

Don't Stop Pretraining: Adapt Language Models to Domains and Tasks. [1]

Table 1: List of the domain-specific unlabeled datasets. In columns 5 and 6, we report ROBERTA's masked LM loss on 50K randomly sampled held-out documents from each domain before $(\mathcal{L}_{ROB.})$ and after (\mathcal{L}_{DAPT}) DAPT (lower implies a better fit on the sample). ‡ indicates that the masked LM loss is estimated on data sampled from sources similar to ROBERTA's pretraining corpus.

Figure 2: Vocabulary overlap (%) between domains. PT denotes a sample from sources similar to ROBERTA's pretraining corpus. Vocabularies for each domain are created by considering the top 10K most frequent words (excluding stopwords) in documents sampled from each domain.

Table 5: Results on different phases of adaptive pretraining compared to the baseline RoBERTa (col. 1). Our approaches are DAPT (col. 2, §3), TAPT (col. 3, §4), and a combination of both (col. 4).

"We show that pretraining the model towards a specific task or small corpus can provide significant benefits. Our findings suggest it may be valuable to complement work on ever-larger LMs with parallel efforts to identify and use domain and task relevant corpora to specialize models."

BioBERT

"[..] the word distributions of general and biomedical corpora are quite different, which can often be a problem for biomedical text mining models." [2]

BioBERT

 Table 1. List of text corpora used for BioBERT

BioBERT

"We showed that pre-training BERT on biomedical corpora is crucial in applying it to the biomedical domain. Requiring minimal task-specific architectural modification, BioBERT outperforms previous models on biomedical text mining tasks such as NER, RE and QA."

"[...] while both BERT and ELMo have released pretrained models, they are still trained on general domain corpora such as news articles and Wikipedia." [3]

 Table 1: Test performances of all BERT variants on all tasks and datasets. [...]

Table 2: Comparing SciBERT with the reported BioBERT results on biomedical datasets.

NB: SciBERT was trained on curated textual data; not trained on code or script for example --at leat not trained directly and purposefully on this kind of data

"Computing has indeed revolutionized how research is conducted, but information overload remains an overwhelming problem [...]. In this paper, we argue for a better way through large language models. Unlike search engines, language models can potentially store, combine and reason about scientific knowledge." [4]

- Galactica was trained on a rather small highly curated dataset.
- All the data was standardized as markdown text.

Table 1: Tokenizing Nature. Galactica trains on text sequences that represent scientific phenomena.

- 1. **Citations:** we wrap citations with special reference tokens [START_REF] and [END_REF].
- 2. **Step-by-Step Reasoning:** we wrap step-by-step reasoning with a working memory token, mimicking an internal working memory context.
- 3. **Mathematics:** for mathematical content, with or without LaTeX, we split ASCII operations into individual characters. Parentheses are treated like digits. The rest of the operations allow for unsplit repetitions. Operation characters are !"#\$%&'*+,-./:;<=>?^_'| and parentheses are ()[]{}.

- 4. **Numbers:** we split digits into individual tokens. For example $737612.62 \rightarrow 7,3,7,6,1,2,..,6,2$.
- 5. **SMILES formula:** we wrap sequences with [START_SMILES] and [END_SMILES] and apply characterbased tokenization. Similarly we use [START_I_SMILES] and [END_I_SMILES] where isomeric SMILES is denoted. For example, C(C(=O)O)N → C,(,C,(,=,O,),O,),N.
- 6. **Amino acid sequences:** we wrap sequences with [START_AMINO] and [END_AMINO] and apply character-based tokenization, treating each amino acid character as a single token. For example, MIRLGAPQTL -> M,I,R,L,G,A,P,Q,T,L.

7. **DNA sequences:** we also apply a character-based tokenization, treating each nucleotide base as a token, where the start tokens are [START_DNA] and [END_DNA]. For example, CGGTACCCTC -> C, G, G, T, A, C, C, C, T, C.

Figure 5: Prompt Pre-training. Pre-training weighs all tokens equally as part of the self-supervised loss. This leads to a weak relative signal for tasks of interest, meaning model scale has to be large to work. Instruction tuning boosts performance post hoc, and can generalize to unseen tasks of interest, but it risks performance in tasks that are distant from instruction set tasks. Prompt pre-training has a weaker task of interest bias than instruction tuning but less risk of degrading overall task generality.

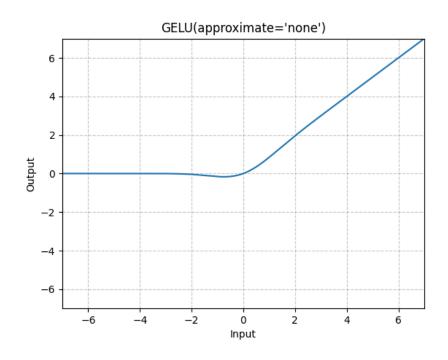
- GeLU Activation GeLU activations for all model sizes.
- Context Window a 2048 length context window.
- **No Biases** following PaLM, we do not use biases in any of the dense kernels or layer norms.
- Learned Positional Embeddings learned positional embeddings for the model.
- **Vocabulary** vocabulary of 50k tokens using BPE. The vocabulary was generated from a randomly selected 2% subset of the training data.

Gaussian Error Linear Units function (GeLu)

$$GELU(x) = x * \Phi(x)$$

Where $\Phi(x)$ is the Gaussian function.

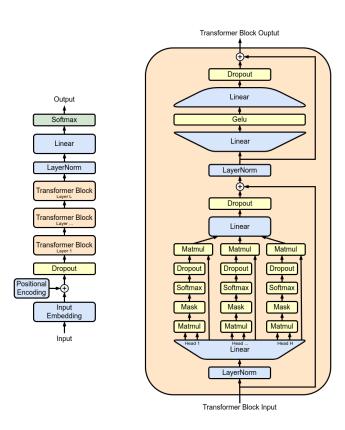
$$GELU(x) pprox x*rac{1}{2}(1+Tanh(rac{2}{\pi}*(x+0.044715*x^3)))$$



- Allows small negative values when x < 0.
- Avoids the dying ReLU problem.

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Why no biases?



Unsupervised Classification Models

[6]

[5]

[6]

- (1) The data is being compressed mutliple time -> challeging document can be hard to embed.
- (2) Slow to process, as we need to chunk the inputs to make multiple inferences.

Can we do better?

Contrastive learning uses similar data point and opposite ones in order for the model build close representations for the first ones and and more separated ones for the latter. [7]

- Unsupervised SimCSE: standard dropout as data augmentation
- Supervised SimCSE: use pairs in NLI datasets

$$\mathcal{L}_{uns} = -lograc{exp(rac{sim(ext{ h}_{i}, ext{ h}_{i}^{+})}{ au})}{\sum_{j=1}^{N} exp(rac{sim(ext{ h}_{i}, ext{ h}_{i}^{+})}{ au})}$$

$$\mathcal{L}_{sup} = -lograc{exp(rac{sim(old h_i,old h_i^+)}{ au})}{\sum_{j=1}^N exp(rac{sim(old h_i,old h_j^+)}{ au}) + exp(rac{sim(old h_i,old h_j^-)}{ au})}$$

[8]

- The pretrained embeddings are being regularized to be more uniform.
- Semantically close pairs are better aligned.

Better performances, hence solving (1).

See also [9] [10]

Learning Long-Range Dependencies

Long-range attention models

Sliding window attention: Longformer [11]

Long-range attention models

Sliding window attention: Mistral 7B [12]

State-space models: Mamba

Gu, Albert, and Tri Dao. "Mamba: Linear-Time Sequence Modeling with Selective State Spaces." arXiv, December 1, 2023. https://doi.org/10.48550/arXiv.2312.00752.

Questions?

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