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► **To cite this version:**

Nesrine Bannour, Christophe Servan, Aurélie Névéol, Xavier Tannier. A Benchmark Evaluation of Clinical Named Entity Recognition in French. The 2024 Joint International Conference on Computational Linguistics, Language Resources and Evaluation (LREC-COLING 2024), May 2024, Torino, Italy. hal-04523267

**HAL Id: hal-04523267**

**<https://hal.science/hal-04523267v1>**

Submitted on 27 Mar 2024

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# A Benchmark Evaluation of Clinical Named Entity Recognition in French

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## Abstract

**Background:** Transformer-based language models have shown strong performance on many Natural Language Processing (NLP) tasks. Masked Language Models (MLMs) attract sustained interest because they can be adapted to different languages and sub-domains through training or fine-tuning on specific corpora while remaining lighter than modern Large Language Models (LLMs). Recently, several MLMs have been released for the biomedical domain in French, and experiments suggest that they outperform standard French counterparts. However, no systematic evaluation comparing all models on the same corpora is available. **Objective:** This paper presents an evaluation of masked language models for biomedical French on the task of clinical named entity recognition. **Material and methods:** We evaluate biomedical models `CamemBERT-bio` and `DrBERT` and compare them to standard French models `CamemBERT`, `FlauBERT` and `FrALBERT` as well as multilingual `mBERT` using three publicly available corpora for clinical named entity recognition in French. The evaluation set-up relies on gold-standard corpora as released by the corpus developers. **Results:** Results suggest that `CamemBERT-bio` outperforms `DrBERT` consistently while `FlauBERT` offers competitive performance and `FrALBERT` achieves the lowest carbon footprint. **Conclusion:** This is the first benchmark evaluation of biomedical masked language models for French clinical entity recognition that compares model performance consistently on nested entity recognition using metrics covering performance and environmental impact.

**Keywords:** named entity recognition, domain adaptation, masked language models, clinical narratives

## 1. Introduction

The recent development of Clinical Data Warehouses in many French hospitals (Jannot et al., 2017; Madec et al., 2019; Pressat-Laffouilhère et al., 2022) is making unstructured clinical data, including narratives, available for secondary use. As a result, there is a growing need in the biomedical community for Natural Language Processing tools that facilitate the extraction of clinical information from text to support epidemiological studies. Many epidemiological indicators can be modeled as named entities to be extracted from the raw text of clinical reports.

For this reason, the task of Named Entity Recognition, or NER, has attracted a lot of attention in the past decades, in particular through shared tasks aiming at direct comparison of methods. Earlier challenges offered tasks for English (Uzuner et al., 2007, 2011) but more recently, other languages have also been addressed (Marimon et al., 2019; Intxaurreondo et al., 2018), including French (Névéal et al., 2015; Cardon et al., 2020).

The evaluation resources released in the shared tasks continue to be used for evaluating new methods and tools. However, individual efforts often come with adaptations of the data sets or metrics so that comparability is not possible across the board.

Herein, we address this issue by presenting a systematic evaluation that offers comparability across systems as well as with the literature introducing the reference corpora. The main contributions of this paper are:

- A benchmark evaluation of clinical named entity recognition in French based on original gold standard annotations, including nested entities
- A comparison of freely available masked language models for general and biomedical French on the NER task
- A comparison to strong symbolic baselines

## 2. Corpora

This section briefly presents the clinical French corpora used to train NER models and evaluate the systems considered in this benchmark.

- **DEFT** (Cardon et al., 2020) is a subset of 167 clinical cases from the CAS corpus (Grabar et al., 2018), introduced in the DEFT challenge in 2020<sup>1</sup>. This corpus is annotated with 13 types of clinical entities and five attributes. It

<sup>1</sup><https://deft.limsi.fr/2020/index-en.html>.

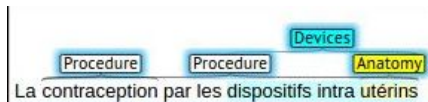


Figure 1: 3 layers of nested entities in an excerpt of the MEDLINE corpus

is divided into a training set of 85 documents, a validation set of 20 documents, and a test set of 62 documents.

- **E3C** (Magnini et al., 2021) is a European corpus of clinical cases. We use the French subcorpus, which comprises 1,615 clinical cases collected in the public domain. It is annotated with 6 types of named clinical entities, including `CLINENTITY`, which we disaggregate into subtypes in order to have an annotation scheme with a diversity approaching that of other corpora. Each entity of this type is associated with a Concept Unique Identifier from the UMLS (Unified Medical Language System) metathesaurus, which can be used to retrieve semantic groups (McCray et al., 2001). In our experiments, we use the gold-standard annotations in the first layer. We use 20% of the training set for validation in the NER models.
- **QUAERO French Med** (Névéol et al., 2014) comprises documents belonging to two text genres, which we treat separately. The EMEA subcorpus is a collection of 13 patient information leaflets supplied by the European Medicines Agency that describes drugs marketed in Europe. The MEDLINE subcorpus consists of 2,500 titles of scientific articles indexed in the MEDLINE database<sup>2</sup>. The entire corpus is annotated with 10 types of clinical entities derived from UMLS semantic groups.

Table 1 presents general descriptive statistics of the study corpora. Table 2 presents descriptive statistics of the distribution of entities in layers in the study corpora. Figure 1 present an excerpt of the MEDLINE corpus containing annotations over three layers: on Layer 1, "contraception par les dispositifs intra utérins" (*contraception with intra uterine devices*) is annotated with the entity type "PROCEDURE", on layer 2 "dispositifs intra utérins" (*intra uterine devices*) is annotated with the entity type "DEVICE" while "contraception" is annotated as a "PROCEDURE" and on layer 3, "utérins" (*uterine*) is annotated with the entity type "ANATOMY".

<sup>2</sup><http://pubmed.ncbi.nlm.nih.gov/>

	train	dev	test
	<i>DEFT</i>		
<b>Tokens</b>	31,752	5,076	20,360
<b>Entities (all)</b>	7,584	1,432	5,140
<b>Entities (Unique)</b>	5,037	1,230	3,809
	<i>E3C<sub>FR</sub></i>		
<b>Tokens</b>	19,808	-	4,671
<b>Entities (all)</b>	3,406	-	706
<b>Entities (Unique)</b>	2,197	-	566
	<i>EMEA</i>		
<b>Tokens</b>	14,944	13,271	12,042
<b>Entities (all)</b>	2,695	2,260	2,204
<b>Entities (Unique)</b>	923	756	658
	<i>MEDLINE</i>		
<b>Tokens</b>	10,552	10,503	10,871
<b>Entities (all)</b>	2,994	2,977	3,103
<b>Entities (Unique)</b>	2,296	2,288	2,390

Table 1: Number of tokens and entity annotations in each split of the study corpora.

Layer 1	Layer 2	Layer 3	Layer 4
<i>DEFT</i>			
65.13%	30.84%	4.02%	0%
<i>E3C<sub>FR</sub></i>			
94.47%	5.36%	0.12%	0.06%
<i>EMEA</i>			
85.73%	13.60%	0.64%	0.03%
<i>MEDLINE</i>			
74.69%	23.75%	1.50%	0.06%

Table 2: Distribution of entity annotations in each layer of the study corpora.

### 3. Named Entity Recognition models

We trained named entity recognition (NER) models using the python library `NLstruct` (Wajsbürt, 2021)<sup>3</sup>. `NLstruct` NER models comprise a text encoder, a word tagger, and a bounds matcher. In addition to state-of-the-art performance on NER tasks, `NLstruct` features the ability to address nested entities, which can be found in QUAERO French Med. `NLstruct` also supports annotations in the BRAT standoff format<sup>4</sup>, which is used by all three corpora in our study.

The text encoder component in `NLstruct` relies on embeddings produced by a BERT language model, a char-CNN encoder, and static French Fast-

<sup>3</sup><https://github.com/percevalw/nlstruct>

<sup>4</sup>The Brat Rapid Annotation Tool - BRAT (Stenertorp et al., 2012) produces annotations in the so-called standoff format described in the BRAT online manual <https://brat.nlplab.org/standoff.html>

Text embeddings. To compare French MLMs, we train one NER model using each of the language models in our benchmark. These MLMs are available in the HuggingFace transformers library (Wolf et al., 2020). However, it can be noted that some level of adaptation was needed at the tokenization step to use some of the models within `NLstruct`, especially for `frALBERT` and `FlauBERT` models<sup>5</sup>.

**General models.** We used the major French MLMs freely available for our experiments and a multilingual model.

- CamemBERT (Martin et al., 2020): a language model for French based on the RoBERTa (Liu et al., 2019) model that was pretrained on the OSCAR French corpus (Suárez et al., 2019). We use the `camembert-base` model.
- FlauBERT (Le et al., 2020): a language model for French based on the BERT (Devlin et al., 2019) model that was pretrained on a large multiple-source French corpus. We use the `flaubert-base-uncased` model.
- FrALBERT (Cattan et al., 2021): a compact model based on the ALBERT (Lan et al., 2020) model that was pretrained on 4GB of French Wikipedia part. We use the `fralbert-base` model.
- Multilingual BERT (Devlin et al., 2019): a language model that was pretrained on the 102 languages with the largest Wikipedia, including French. We use the `bert-base-multilingual-uncased` model. We denote Multilingual BERT as `mBERT`.

**Domain-specific models.** We used the French MLMs dedicated to the biomedical domain that were freely available for our experiments.

- CamemBERT-bio (Touchent et al., 2023): an adapted CamemBERT model for the biomedical domain that was built using continual-pretraining from `camembert-base` model and trained on a created French biomedical corpus from three sources: The ISTE database, the CLEAR corpus (Grabar and Cardon, 2018) and the third unannotated layer of the E3C corpus (Magnini et al., 2021). we use the `camembert-bio-base` model.
- DrBERT (Labrak et al., 2023): a RoBERTa-based French biomedical model that was trained from scratch on a web-based medical corpus. We use the `DrBERT-4GB` model.

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<sup>5</sup>The source code is available here: <https://gitlab.lisn.upsaclay.fr/nlp/deep-learning/nlstruct>

We also considered `ALiBERT` (Berhe et al., 2023) but did not find a publicly accessible version. Similarly, the model described by (Le Clercq de Lannoy et al., 2022) was not available to us.

**Baseline.** As a baseline, we use a symbolic method that builds a dictionary of entities found in the training and development splits of a corpus and simply matches these entities in the test split. In practice, we use the BRAT propagation tool introduced by (Grouin, 2016) to "propagate" BRAT annotations to the test sets<sup>6</sup>.

## 4. Evaluation metrics

We evaluate the performance of our models at the entity level by measuring the micro Precision, Recall, and F-measure. Confidence intervals at 95% confidence level were computed using the empirical bootstrap method (Dekking et al., 2007, p. 275). Each test corpus is sampled with replacement 1000 times, and evaluation metrics are calculated for each sample. Baseline scores were computed using `brateval` (Verspoor et al., 2013). To measure the carbon footprint of training and testing our models, we use the `Carbon tracker` tool (Anthony et al., 2020). These estimates are approximative and are computed by using an average carbon intensity of 58.48 gCO<sub>2</sub>/kWh corresponding to our location in France.

## 5. Results and discussion

Tables 3 to 6 present the results of our NER experiments on the study corpora.

### 5.1. NER performance

**Overall entity extraction performance.** The NER models trained using masked language models outperform the symbolic baseline by at least 10 points of F-measure (up to 40 points for DEFT), although the symbolic baseline can exhibit high precision (e.g., on EMEA, MEDLINE). Interestingly, the knowledge-based approach proposed by Van Muligen et al. (2016) continues to achieve the best results on the MEDLINE and EMEA corpora, with an F-measure of 0.7 and 0.75 respectively.

The performance of the general French and multilingual models is quite similar, including the multilingual model<sup>7</sup>, and, on average, lower compared to the performance of the biomedical models, except for the E3C corpus, where the `FlauBERT` model performs better. The `CamemBERT-bio`

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<sup>6</sup><https://github.com/grouin/propa>

<sup>7</sup>Copara et al. (2020) found `mBERT` to be outperformed by `CamemBERT-large`

Models	DEFT			
	Precision	Recall	F-measure	CO <sub>2</sub> eq (g.)
CamemBERT	0.73 [0.71-0.75]	0.75 [0.73-0.77]	0.74 [0.73-0.76]	7.7
FlauBERT	0.73 [0.72-0.76]	0.75 [0.73-0.77]	0.74 [0.73-0.76]	8
mBERT	0.72 [0.70-0.74]	0.74 [0.72-0.76]	0.73 [0.71-0.75]	8.3
frALBERT	0.68 [0.66-0.69]	0.67 [0.65-0.69]	0.67 [0.66-0.69]	4.5
CamemBERT-bio	<b>0.75</b> [0.73-0.77]	<b>0.77</b> [0.75-0.78]	<b>0.76</b> [0.74-0.78]	8.7
DrBERT	0.71 [0.68-0.73]	0.72 [0.70-0.74]	0.71 [0.69-0.73]	5.5
Baseline	0.38	0.32	0.35	-
Copara et al. (2020)	-	-	0.73	-

Table 3: Performance of nested entity extraction on the DEFT test set.

Models	E3C			
	Precision	Recall	F-measure	CO <sub>2</sub> eq (g.)
CamemBERT	0.52 [0.42-0.63]	0.50 [0.45-0.56]	0.51 [0.46-0.56]	3.6
FlauBERT	<b>0.54</b> [0.49-0.60]	<b>0.53</b> [0.46-0.60]	<b>0.54</b> [0.51-0.57]	4.1
mBERT	0.51 [0.45-0.58]	0.52 [0.47-0.57]	0.52 [0.48-0.54]	4.8
frALBERT	0.50 [0.41-0.59]	0.55 [0.51-0.59]	0.52 [0.47-0.58]	2.5
CamemBERT-bio	0.52 [0.44-0.61]	0.52 [0.45-0.59]	0.52 [0.48-0.55]	3.6
DrBERT	0.47 [0.40-0.57]	0.52 [0.46-0.60]	0.49 [0.46-0.53]	3.6
Baseline	0.24	0.37	0.29	-

Table 4: Performance of nested entity extraction on the E3C test set.

Models	MEDLINE			
	Precision	Recall	F-measure	CO <sub>2</sub> eq (g.)
CamemBERT	0.64 [0.62-0.66]	0.66 [0.64-0.67]	0.65 [0.63-0.66]	1.9
FlauBERT	0.67 [0.65-0.68]	0.69 [0.67-0.71]	0.68 [0.66-0.69]	2.2
mBERT	0.63 [0.61-0.65]	0.67 [0.65-0.69]	0.65 [0.63-0.67]	3
frALBERT	0.53 [0.51-0.54]	0.52 [0.49-0.54]	0.52 [0.50-0.54]	1.1
CamemBERT-bio	0.66 [0.65-0.68]	0.70 [0.68-0.72]	0.68 [0.66-0.70]	2.2
DrBERT	0.63 [0.61-0.65]	0.65 [0.63-0.67]	0.64 [0.62-0.66]	2
Baseline	<b>0.73</b>	0.30	0.42	-
Van Mulligen et al. (2016)	0.68	<b>0.72</b>	<b>0.70</b>	-

Table 5: Performance of nested entity extraction on the MEDLINE test set.

Models	EMEA			
	Precision	Recall	F-measure	CO <sub>2</sub> eq (g.)
CamemBERT	0.66 [0.62-0.70]	0.65 [0.56-0.73]	0.65 [0.59-0.72]	3.9
FlauBERT	0.69 [0.67-0.72]	0.66 [0.59-0.73]	0.68 [0.63-0.71]	4.4
mBERT	0.67 [0.64-0.72]	0.67 [0.61-0.73]	0.67 [0.63-0.72]	4
frALBERT	0.62 [0.57-0.67]	0.65 [0.61-0.70]	0.63 [0.59-0.68]	2.4
CamemBERT-bio	0.70 [0.66-0.74]	0.68 [0.61-0.75]	0.69 [0.63-0.74]	5.2
DrBERT	0.69 [0.66-0.72]	0.64 [0.58-0.71]	0.66 [0.62-0.71]	3
Baseline	<b>0.73</b>	0.43	0.55	-
Van Mulligen et al. (2016)	0.72	<b>0.79</b>	<b>0.75</b>	-

Table 6: Performance of nested entity extraction on the EMEA test set.

biomedical model seems to perform better than the DrBERT biomedical model, suggesting that continual-pretraining from an existing French model on biomedical data might be beneficial in achieving good outcomes.

**Nested entity extraction performance.** A layer-by-layer evaluation would be difficult to perform, due to the difficulty of aligning system outputs and reference annotations at the layer level. However, we can report that the system outputs contain annotations with depth 3 or 4 depending on the specific models and corpora and exhibit a distribution

of annotations across layers that is similar to that of reference annotations. This suggests that the nesting of annotations by NLStruct is successful. Moreover, F-measure for the DEFT corpus exceed .65, which would be the ceiling score for a system performing flat-entity extraction of layer 1 entities. This also suggests that the nested entity extraction is performed successfully.

### Entity extraction performance per entity type.

Due to space constraints, we are not providing detailed performance per entity type over the study corpora. Nonetheless, we can notice that the performance of the models tends to vary following similar trends, with highest performance reached for entity types with either high support in terms of training instances and/or high regularity in their occurrence patterns (e.g., temporal entities).

## 5.2. Comparability of models and experiments

To evaluate their biomedical CamemBERT-bio model, Touchent et al. (2023) fine-tuned a NER model on the semi-annotated layer 2 of the E3C corpus and evaluated it on the first layer of this corpus, yielding an F-measure of 69.85. A direct comparison with our results is not possible since we train and evaluate our model using only the first layer containing the gold standard annotations. However, it suggests that silver standard annotations can be useful for training an NER model.

Touchent et al. (2023) and Labrak et al. (2023) evaluated their biomedical MLMs on the two sub-corpora of QUAERO French Med and compared them to general French models. However, the task was cast as direct token classification and did not address the nested named entities. Indeed, Touchent et al. (2023) removed nested entities by keeping only the coarse entities, whereas Labrak et al. (2023) concatenated the names of the nested entities to produce new entities and evaluated their results at the token level. These experiments can be seen as intrinsic evaluations of the masked language models.

In contrast, our experiments aim to address the entity recognition task in an extrinsic setting. Our results suggest that the size training data available to train the NER models had more impact on NER performance than the language models used: performance of all approaches is generally lower for E3C vs. other corpora. Similarly, when looking at performance on individual entity types, we generally note that entities with the highest prevalence in the training sets yield higher performance.

## 5.3. Carbon footprint

Tables 3 to 6 show the carbon footprint of our NER experiments in terms of CO<sub>2</sub> equivalent measure in grams. The highest CO<sub>2</sub> carbon emissions are observed when training and testing the DEFT NER models. This is partly due to the fact that this corpus has more tokens than the other corpora, as illustrated in Table 1. Overall, the frALBERT-based models have the lowest carbon footprint. These models offer a decrease of carbon emission between 20% and 63% compared to other models, depending on models and corpora. Note that Carbon tracker does not consider the execution environment or energy production. As a result, the obtained measures in our experiments remain approximative. Touchent et al. (2023) reported that the carbon emissions for pre-training their CamemBERT-bio model is estimated to 0.84 kg CO<sub>2</sub> eq. Labrak et al. (2023) reported the overall carbon emissions of their 7 DrBERT-based models, which is 376.45 kg CO<sub>2</sub> eq.

## 6. Conclusion

This is the first benchmark evaluation of masked language models for the biomedical domain on the clinical French NER task, using three publicly available clinical French corpora. The evaluation is based on released gold standard annotations, including nested entities. CamemBERT-bio consistently outperforms DrBERT, while FlauBERT offers competitive results. Overall, frALBERT offers a fair compromise between *F-measure* and carbon impact, with performance that exceeds the baseline consistently by at least 10 points, and a carbon impact that consistently represents a fraction of the impact of other models. On the QUAERO French Med corpus, MLMs fail to outperform the knowledge-based approach proposed by Van Muligen et al. (2016). This systematic evaluation compares model performance using metrics covering both performance and environmental impact.

## 7. Ethical considerations and limitations

**Limitations.** We did not consider all versions of models: for example, camembert-large and flaubert-base-cased are not covered in our experiments. While a full evaluation could cover more models, it would incur a higher carbon footprint and we decided to select representative models of the categories that we aimed to cover: general and domain-specific models that had been recently evaluated on similar corpora without direct comparability.

## 8. Acknowledgements

We thank Dr. Bastien Rance for fruitful discussions on the content of this manuscript. This work was supported by ITMO-Cancer and ANR under grant CODEINE ANR-20-CE23-0026-01.

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