



Applying and Improving a Publicly Available Medication NER Pipeline in a Clinical Cancer EMR

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Background and Objectives



Larger Project (CaVa)

Oral systemic therapy usage not captured in current database

Aim to reliably extract oral systemic therapy usage from clinical notes using NLP

Avoids the need for linkage to administrative dispensing data to capture this information



Current Study

Identify all medication information targets in clinical notes using NLP

Not specific to cancer therapies or oral agents

Foundation towards developing the NLP infrastructure for the larger project



Clinical NLP Advancements

Pre-trained language models on biomedical and/or clinical text increasingly available

- less so pre-trained complete pipelines for specific clinical NLP tasks

Improved success of transfer learning with Transformer architecture

Downstream tasks (e.g. NER, classification) still require annotated text

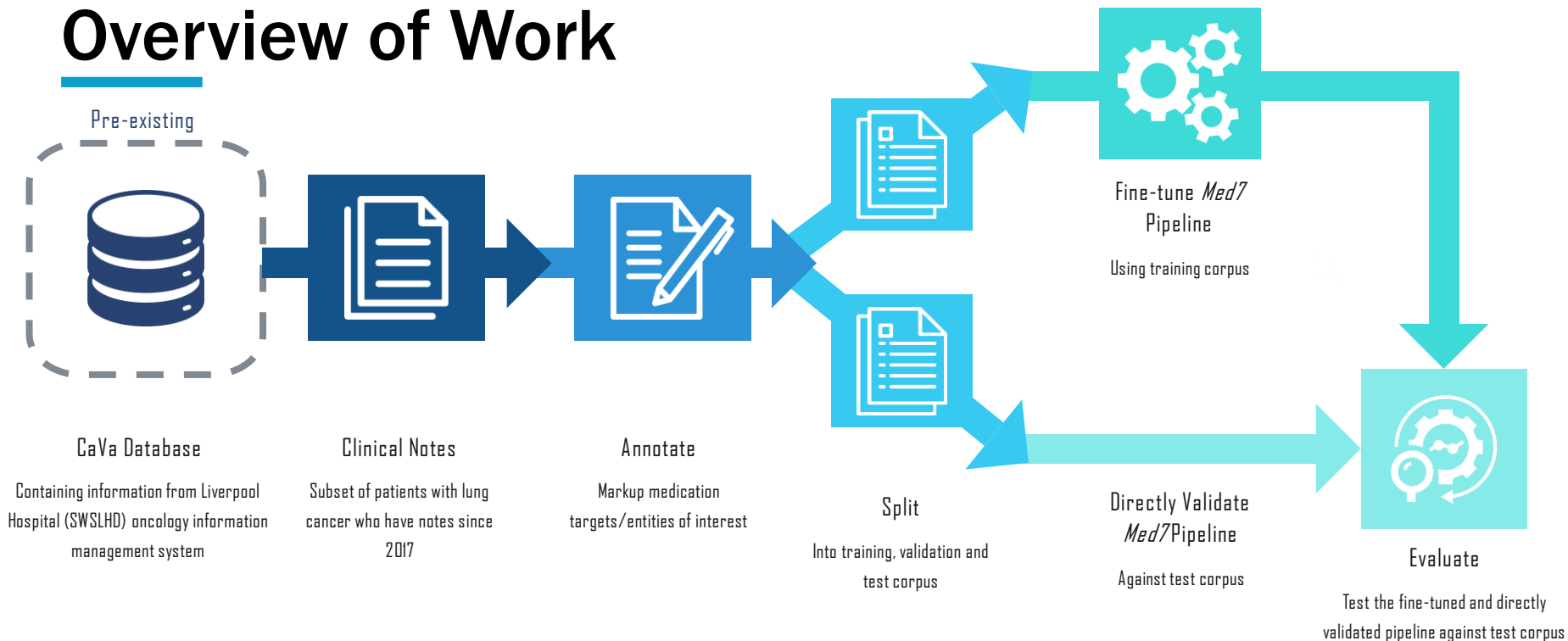
We examined *Med7* - one of the few complete publicly available, pre-trained pipelines (with RoBERTa and NER weights) for medication targets in clinical text, with and without further fine-tuning on a small annotated corpus.

Pre-training and fine-tuning performed with the open-source Python library `med7`.

spaCy

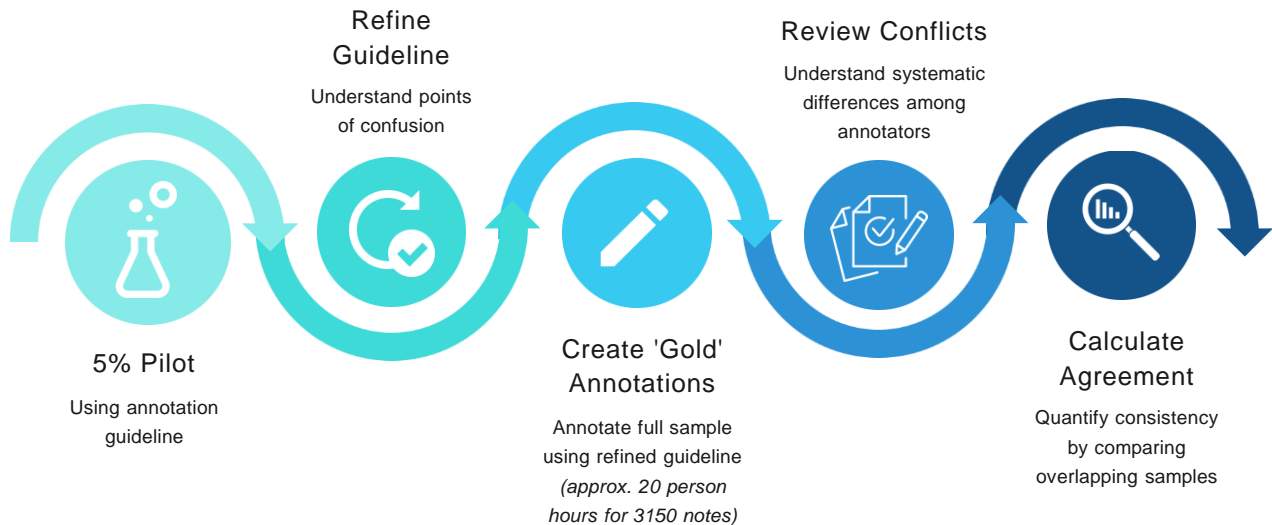


Overview of Work





Annotation



Entities

Drug

Drug Information

Drug Class
Strength
Dosage

DRUG | DRUGCLASS | DOSAGE | STRENGTH | FORM |
FREQUENCY | ROUTE | DURATION |

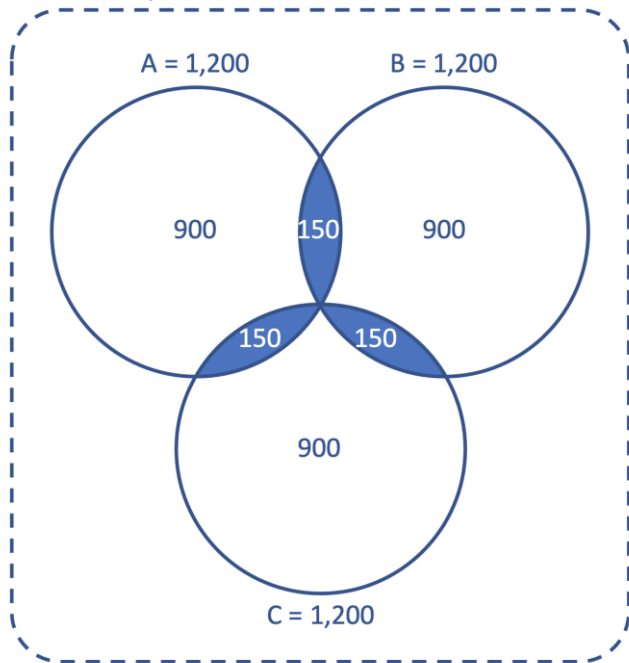
OncHx: NSCLC T3N2M0. PHx: MSSA bacteraemia 2018. SHx: Ex-smoker 20 pk yrs, EtOH 10-12 std/wk premorbidly, denies IVDU, HPC : 48M presented today for day 1 cycle 2 of vinorelbine DRUG PO ROUTE 60mg/m2 STRENGTH + carboplatin DRUG 5 AUC STRENGTH IV ROUTE - Vinorel DRUG switched from IV ROUTE in cycle 1 due to parasthesias. Plan: (1) PO ROUTE dex DRUG 4mg STRENGTH one DOSAGE tab FORM BD FREQUENCY for next 48 hrs DURATION (2) Return in 1/52 for 2nd dose Navelbine DRUG - consider switch back to IV ROUTE if well-tolerated today (3) Present to ED if unwell/febrile.

✓ ✗ ⌕ ↶

*Fabricated text

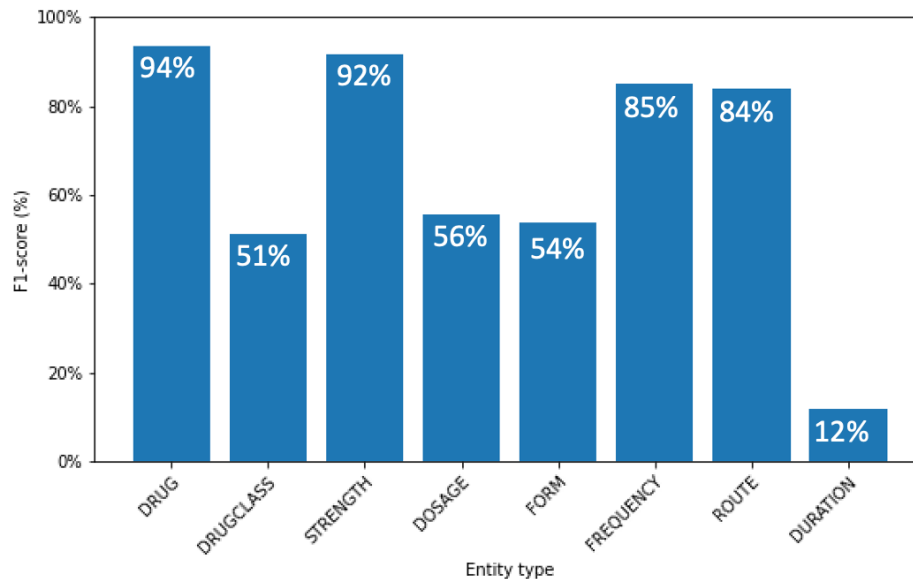


Total corpus = 3,150



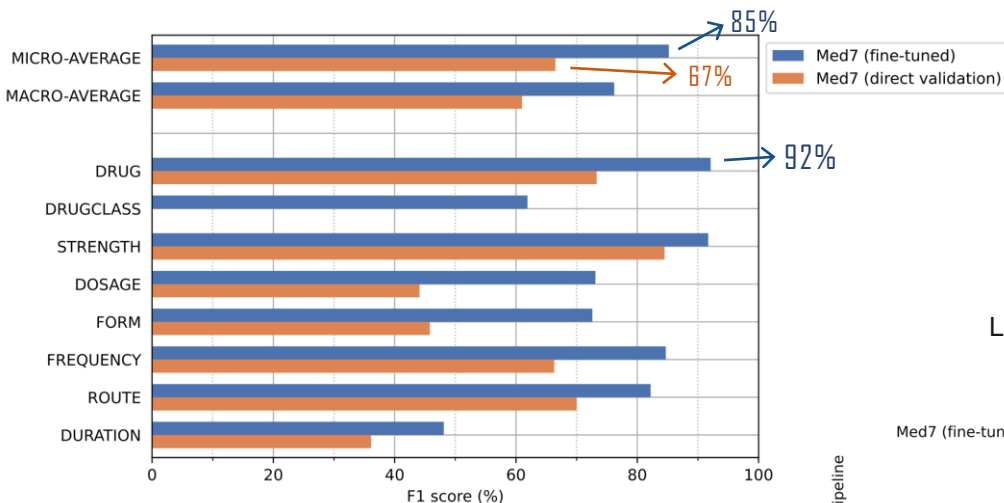
Mean F1-score for all pairs of overlapping samples

- Overall: 82% (strict word and entity matches)
- By entity:





Results



Effects of fine tuning the downstream NER component

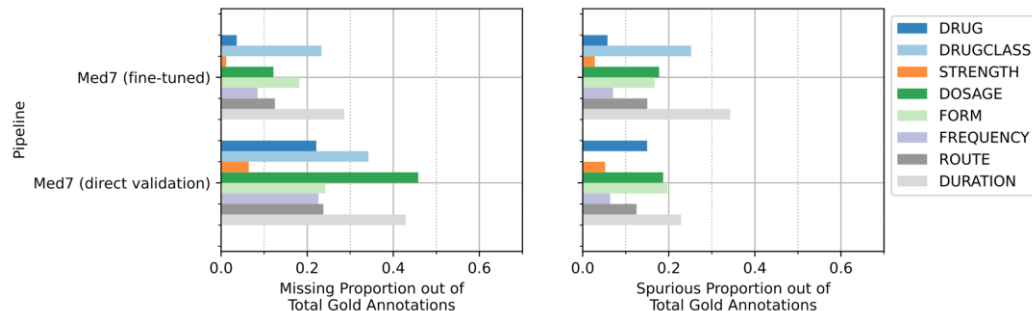
Superior performance per entity and overall

Smallest impact on performance for *strength*

Largest impact on performance for *dosage*

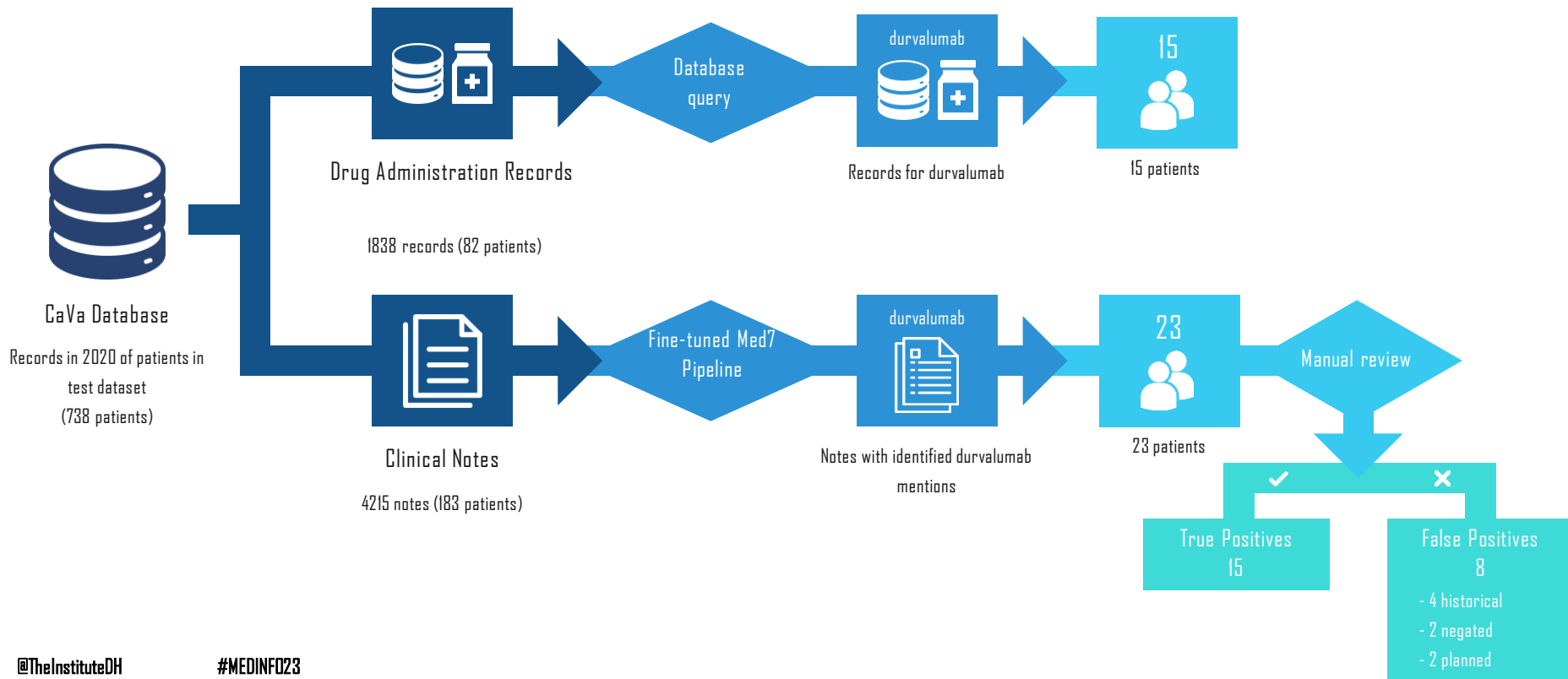
Large improvement on picking up false negative (missing) entities

Less improvement in reducing false positive (spurious) entities



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Key Takeaways

Similar training and target data are important for developing high quality clinical NLP solutions

High performance is achievable with low annotation budgets, especially if utilising pre-trained language models

Pre-trained pipelines for clinical targets are not ubiquitous. Pipeline components can be sourced from previous work in the same institution, particularly language models

NER performance was proportional to the quality and quantity of annotated data for that entity

Further dependency/context resolution is vital for clinical utility



Next Steps



Optimisation of Annotation

Learn from error analysis and optimise iteratively

Leverage other strategies e.g. weak supervision with labelling functions, active learning with 'human-in-the-loop' annotation



Dependency Resolution

Context of mention (planned/negated/considered/historical)

Linkage of annotated drug information to drug names



Patient-Level Validation

Against high-quality data e.g. linkage to administrative dispensing data