Eureka – Finding a way to harvest the data in the medicinal product information

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1. Objective
Build and validate a natural language processing (NLP) tool that extracts adverse drug reactions (ADR) from the English Summary of Product Characteristics (SmPC) of Centrally Authorised Products (CAPs) in the European Union (EU).

2. Methods
A symbolic NLP application titled EurEKA - EU's Product Information Entity extraction and Knowledge acquisition – was developed using R and Python. It downloads and parses SmPCs in PDF format and uses a bespoke dictionary based on the Medical Dictionary for Regulatory Activities (MedDRA) to identify ADRs. Entities are mapped to MedDRA. Results were compared to a database of manually curated ADRs published by the Pharmacoepidemiologic Research on Outcomes of Therapeutics by a European Consortium (PROTECT).

3. Results
A total of 1136 SmPCs were processed and performance was assessed on 910 of these. The macro-averaged performance was F1 0.81, Precision 0.80, and Recall 0.84. Micro-averaged performance was F1 0.84, Precision 0.82, Recall 0.85.

4. Discussion
EurEKA achieved a performance equivalent to the best published performances. Analysis of a sample of SmPCs revealed missing terms in the PROTECT database suggesting performance is likely to be higher. The chosen method did not require annotating data and facilitates the implementation of multilingual methods by utilising available MedDRA translations. EurEKA was built with a human-in-the-loop module (EurEKA Extract) to correct extraction errors, and an analytical/query module (EurEKA Explore) (Fig. 1).

3. Conclusions
EurEKA's performance is at the level of the best performing tools to extract ADR data and can be used to facilitate the extraction of ADR data.