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UCL

IQWiG Information Retrieval Meeting (IRM 2024): Software and data skills for information specialists

Introduction to OpenAlex tools for efficient automated updating of systematic reviews and maps

Ian Shemilt

EPPI Centre, University College London, United Kingdom

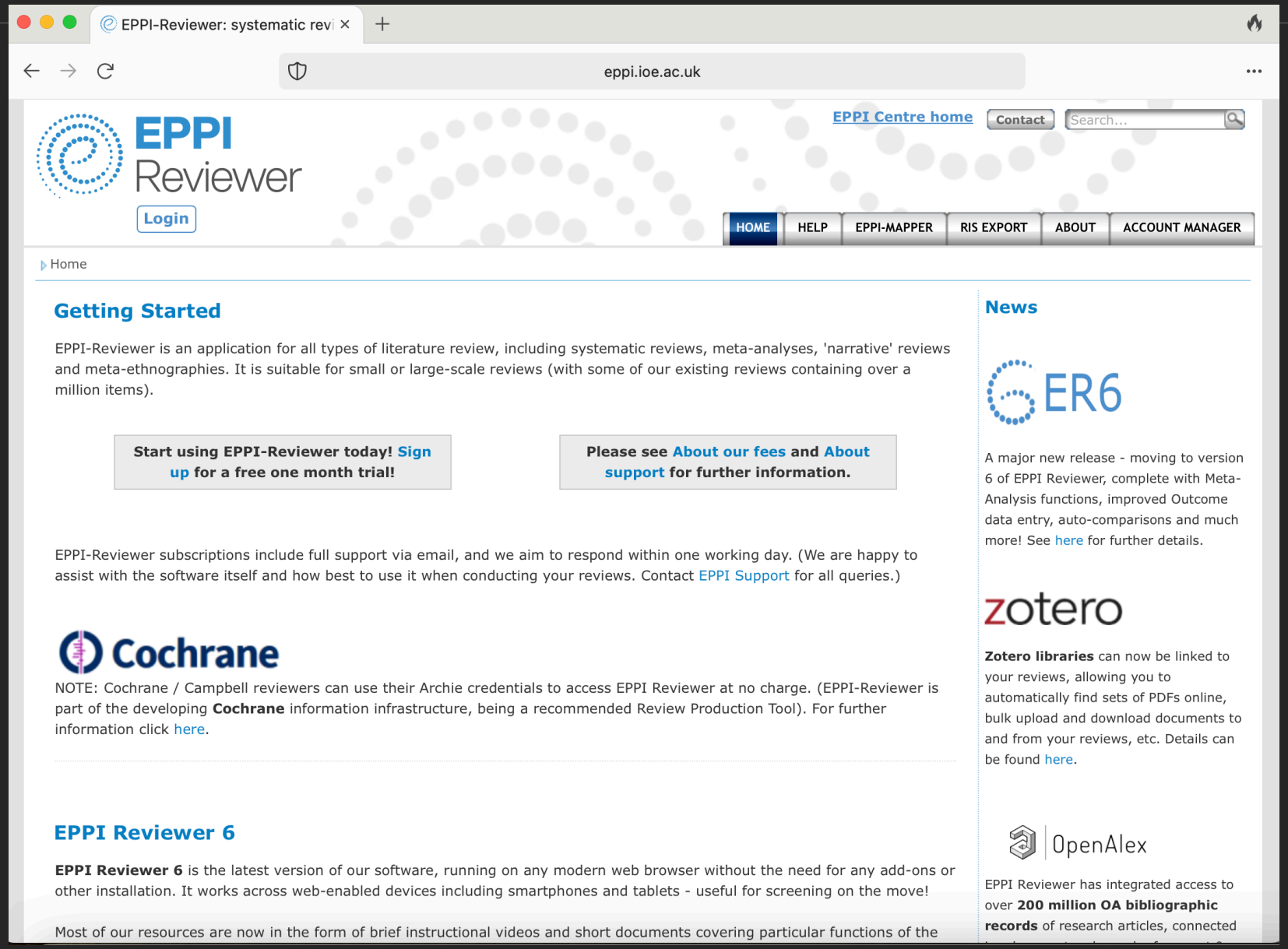
Conflict of interest disclosure

I have an interest that could be perceived as a direct conflict of interest in the context and content of this training workshop:

OpenAlex tools are hosted in EPPI Reviewer

I work at the EPPI Centre and help to develop and support this not-for-profit software

- Advanced web-based software for systematic reviews and other evidence synthesis
- Ordinarily licensed on a not-for-profit, software as service, subscription basis
- Some users charged fees for access, shareable reviews, direct support or training
- Free at the point of use to all Cochrane authors for use to produce Cochrane reviews
- Open source = “coming soon” (late Summer 2024🤞)



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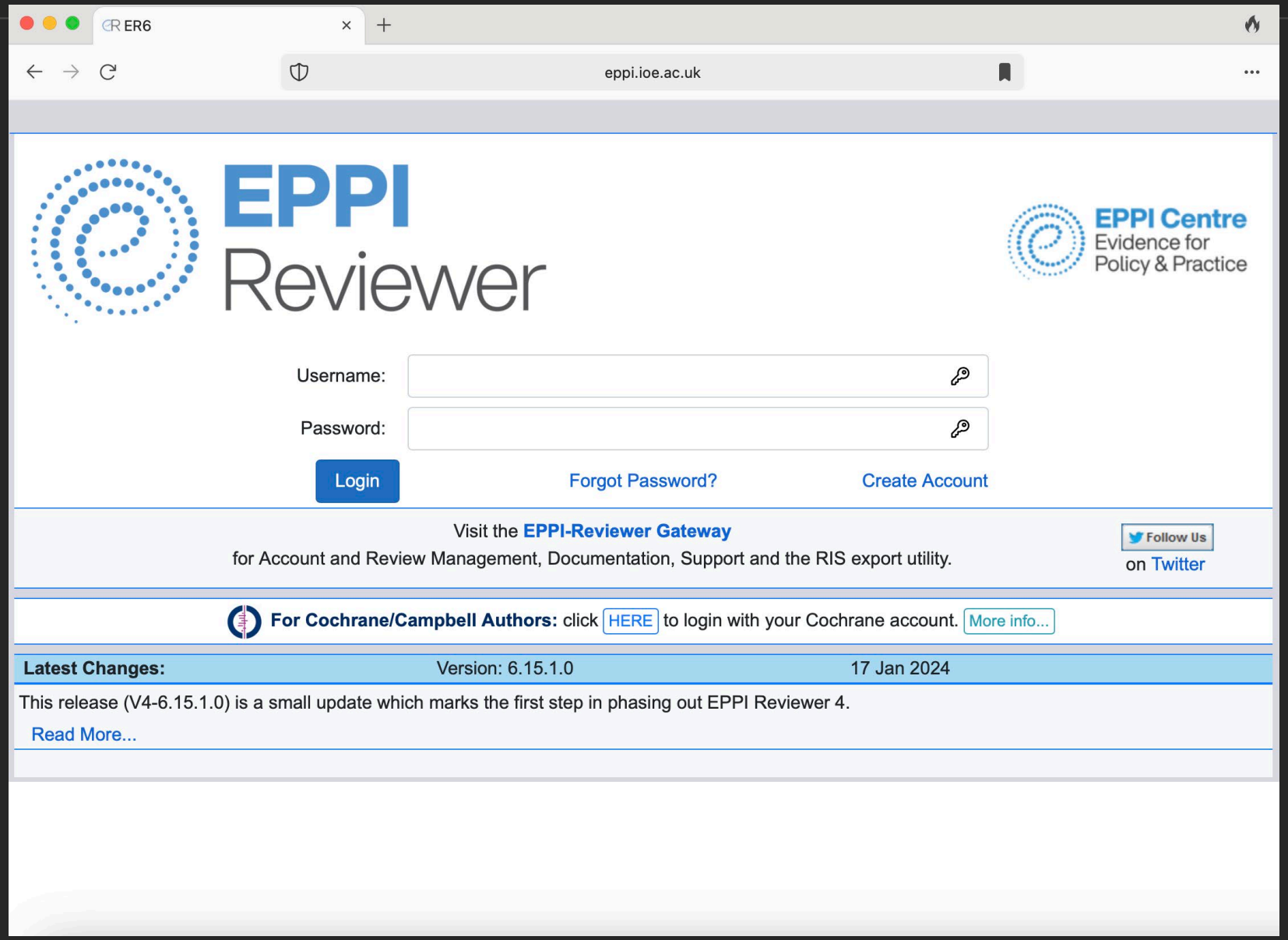
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Follow Us
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Latest Changes:
Version: 6.15.1.0
17 Jan 2024
This release (V4-6.15.1.0) is a small update which marks the first step in phasing out EPPI Reviewer 4.
Read More...

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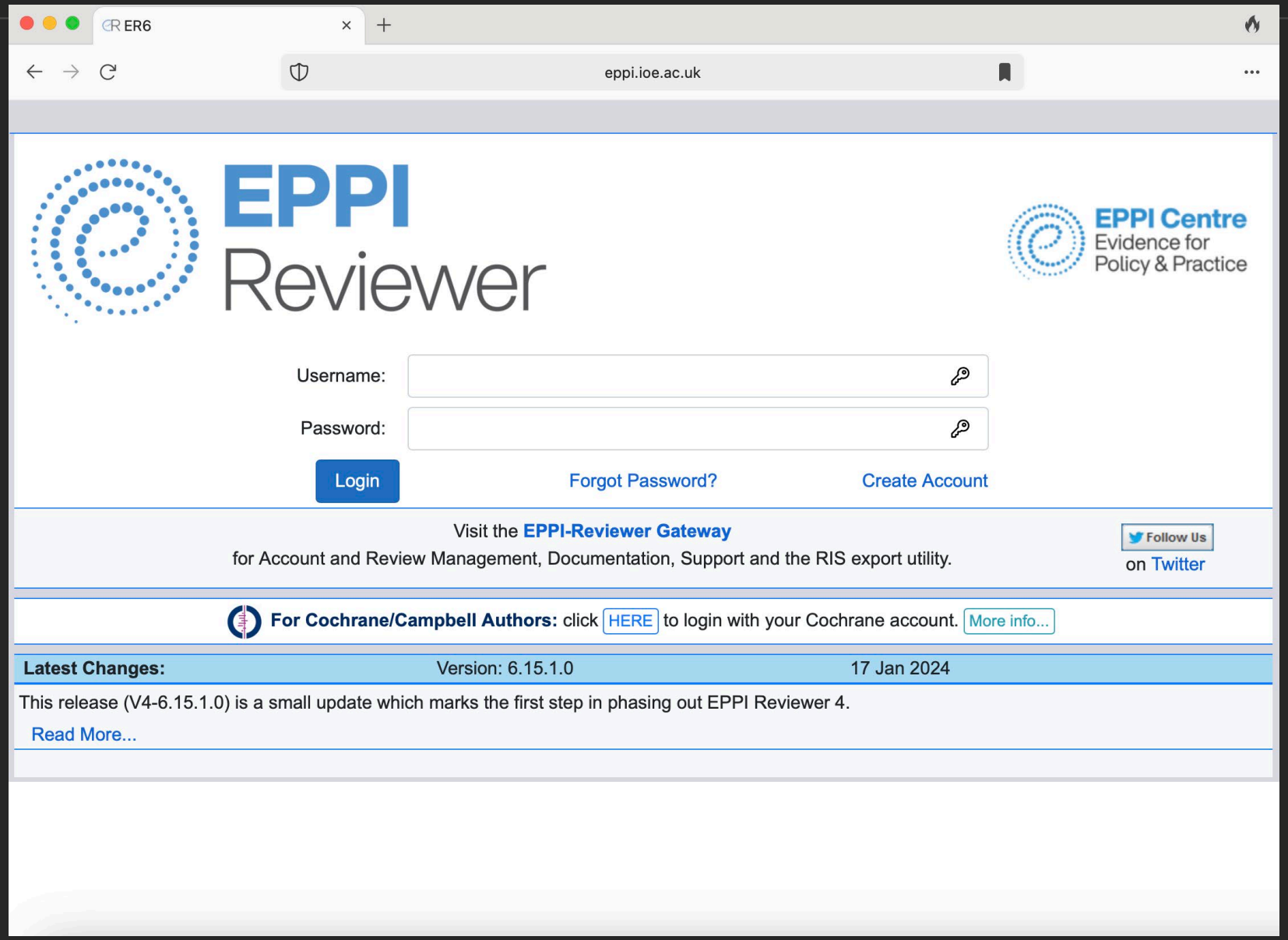
- Katya Mustafini
- Ehsan Zare Borzeshi

EPPI Centre

About me

- EPPI Centre at University College London (UCL)
- Methods for systematic reviews and other forms of evidence synthesis ➔ policy & practice decisions
- Automation tools and new technologies for increasing the efficiency of rigorous systematic review and evidence synthesis processes (≠ ‘rapid reviews’)



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About you

- Ever seen EPPI Reviewer?
- Ever used EPPI Reviewer?
- Regular user of EPPI Reviewer?
- Ever seen or used OpenAlex tools?

Outline

1. Introduce OpenAlex tools (~*15 mins*)
2. Demonstrate OpenAlex tools for updating an IQWiG evidence search (~*30 mins*)
3. Try to answer your questions! (~*15 mins*)

Systematic reviews

"The latest research shows that we really should do something with all this research."

- Identify, appraise and synthesize **all** the evidence *(research studies)* that meets pre-specified eligibility criteria to **answer a specific research question**
- Explicit, systematic methods aimed at minimising bias, to produce more reliable findings to inform decision-making
- Stages of the SR process:
  - Formulating questions and eligibility criteria
  - Searching for eligible studies
  - Selecting eligible studies
  - Coding included study characteristics & extracting data
  - Assessing risk of bias / quality of included studies
  - Synthesising data from included studies
  - Assessing certainty of evidence
  - Interpreting results and summarising findings

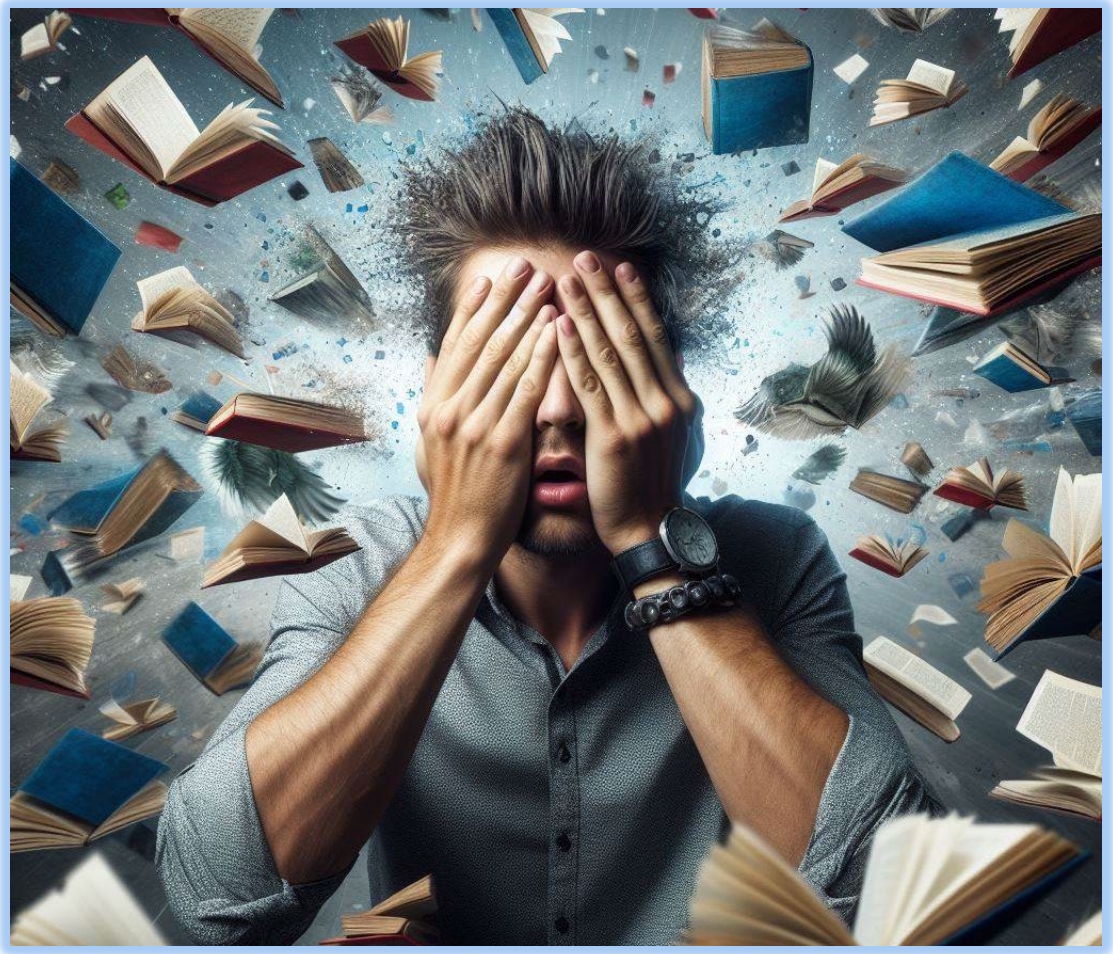
Evidence and gap maps

"The latest research shows that we really should do something with all this research."

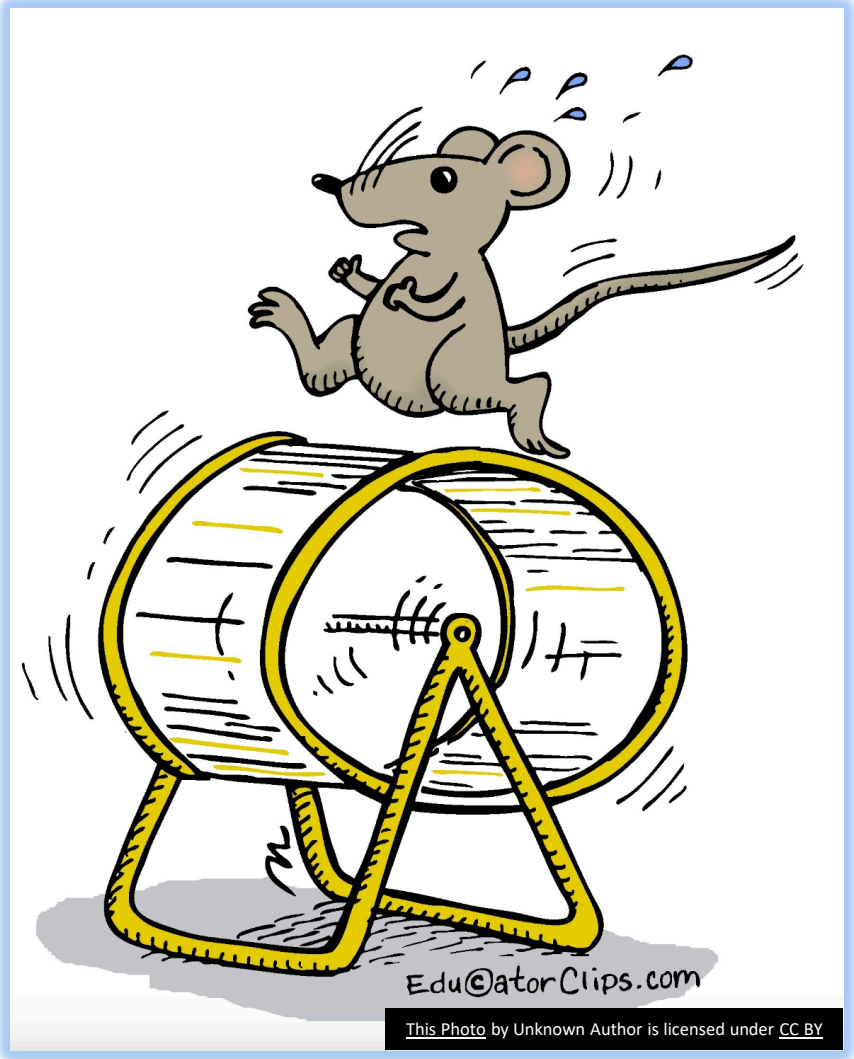
- Identify and describe a representative majority of (**almost all**) the evidence that meets pre-specified eligibility criteria to provide **an overview of a broad area of research**
- Use explicit, systematic methods aimed at minimising bias, to produce more reliable findings to inform decision-making
- Stages of the mapping (EGM) process:
  - Formulating questions and eligibility criteria
  - Searching for eligible studies
  - Selecting eligible studies
  - Coding included study characteristics & extracting data
  - Assessing risk of bias / quality of included studies **???**
  - ~~Synthesising data from included studies~~
  - ~~Assessing certainty of evidence~~
  - Visualising results and summarising findings

Automating evidence synthesis

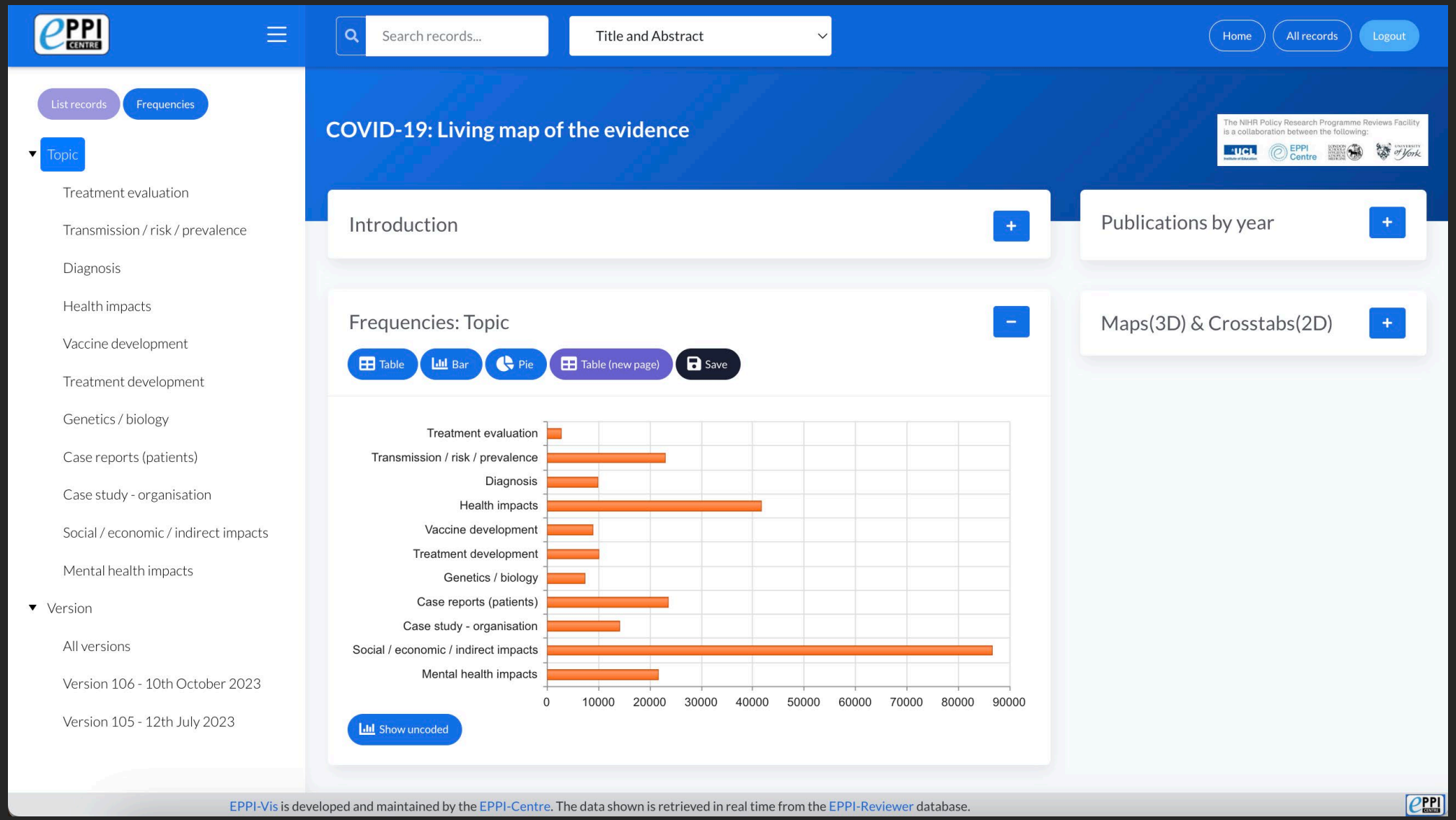
- Conventional evidence synthesis production processes, undertaken manually, are time and resource intensive
- They can therefore take a long time to produce, making it difficult to ‘land’ evidence at the ‘point’ of decisions
- Findings can also become quickly out of date - especially where bodies of evidence are rapidly emerging
- More efficient = reducing time and costs, while maintaining rigor and reliability (≠ ‘rapid reviews’) ‘earning the grounds for the evidence claims’



Keeping living reviews & maps up to date



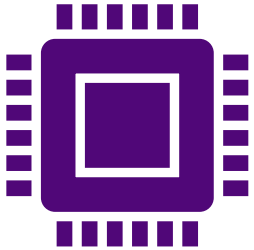
- Increasing awareness of the need to keep evidence synthesis up to date
- Evidence is rapidly emerging, current knowledge is uncertain, new research might change policy or practice
- Requires continual evidence surveillance
- Need for living evidence synthesis (capacity and tools!) was highlighted during the COVID-19 pandemic



https://eppi.ioe.ac.uk/cms/Projects/DepartmentofHealthandSocialCare/Publishedreviews/COVID-19Livingsystematicmapoftheevidence/tabid/3765/Default.aspx

Enablers of a new generation of digital (living) evidence synthesis tools

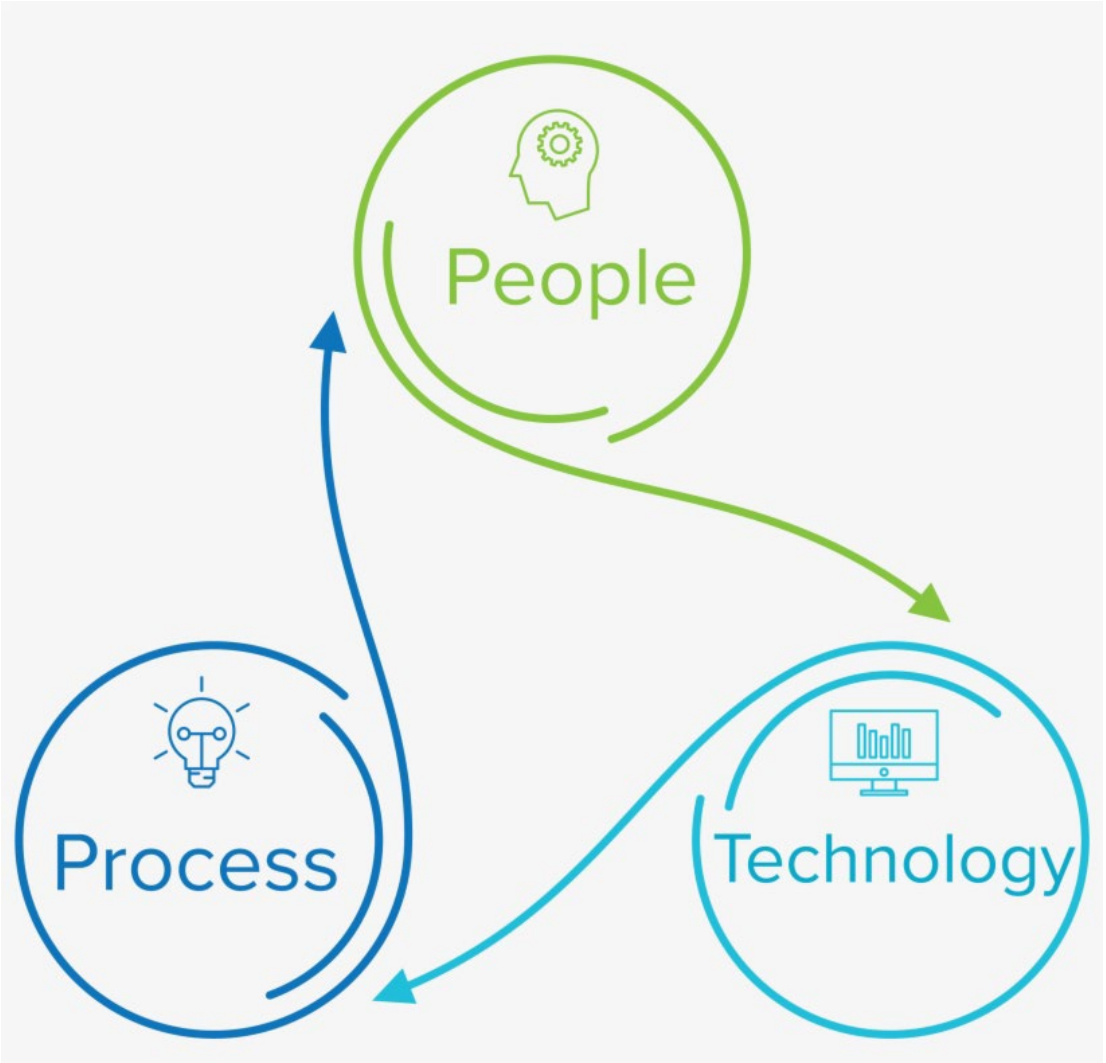
Increased availability of open access research



Increased computing power (both memory + compute)

Advances in machine learning technology

Automation ≃ semi-automation



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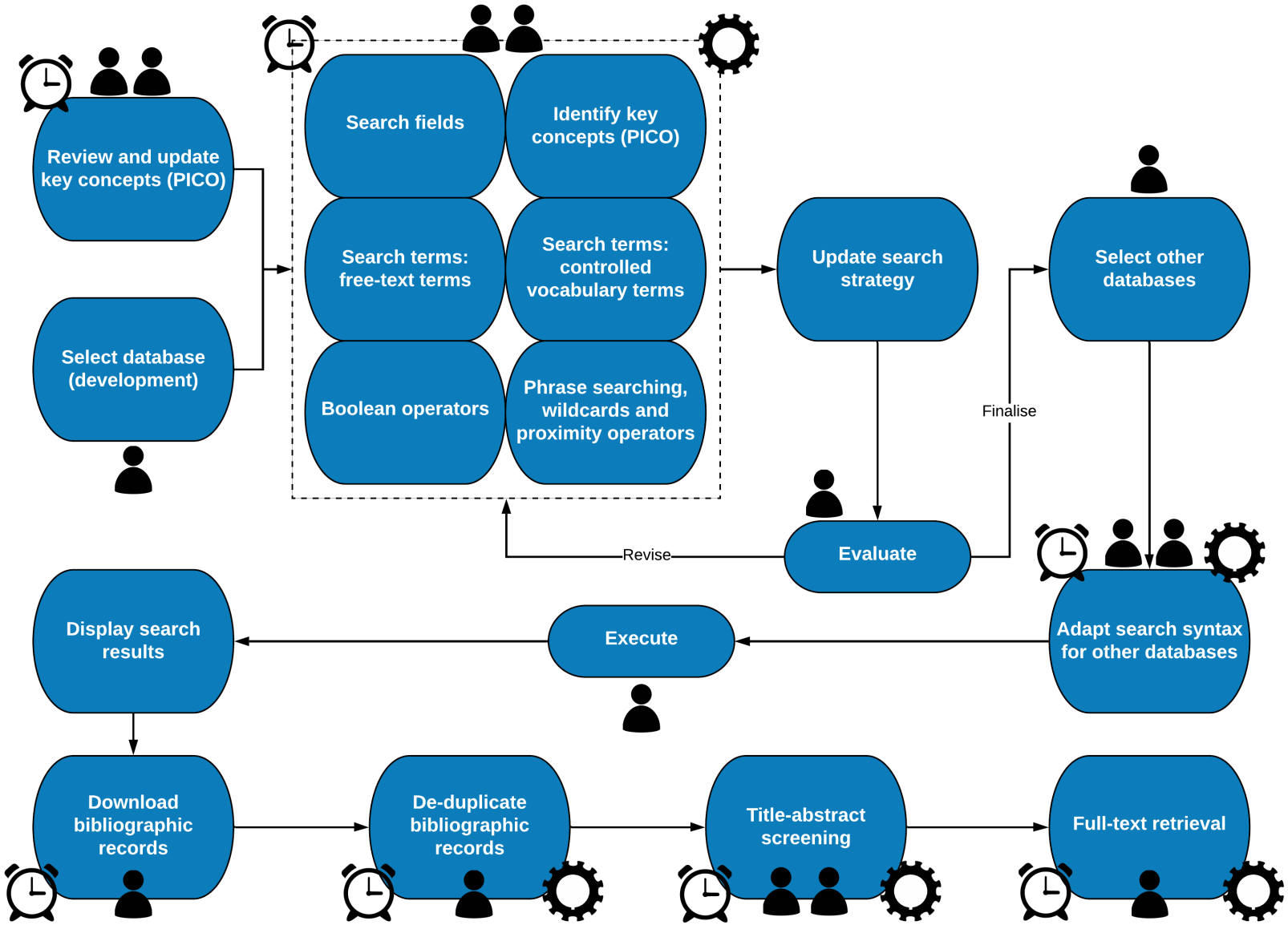
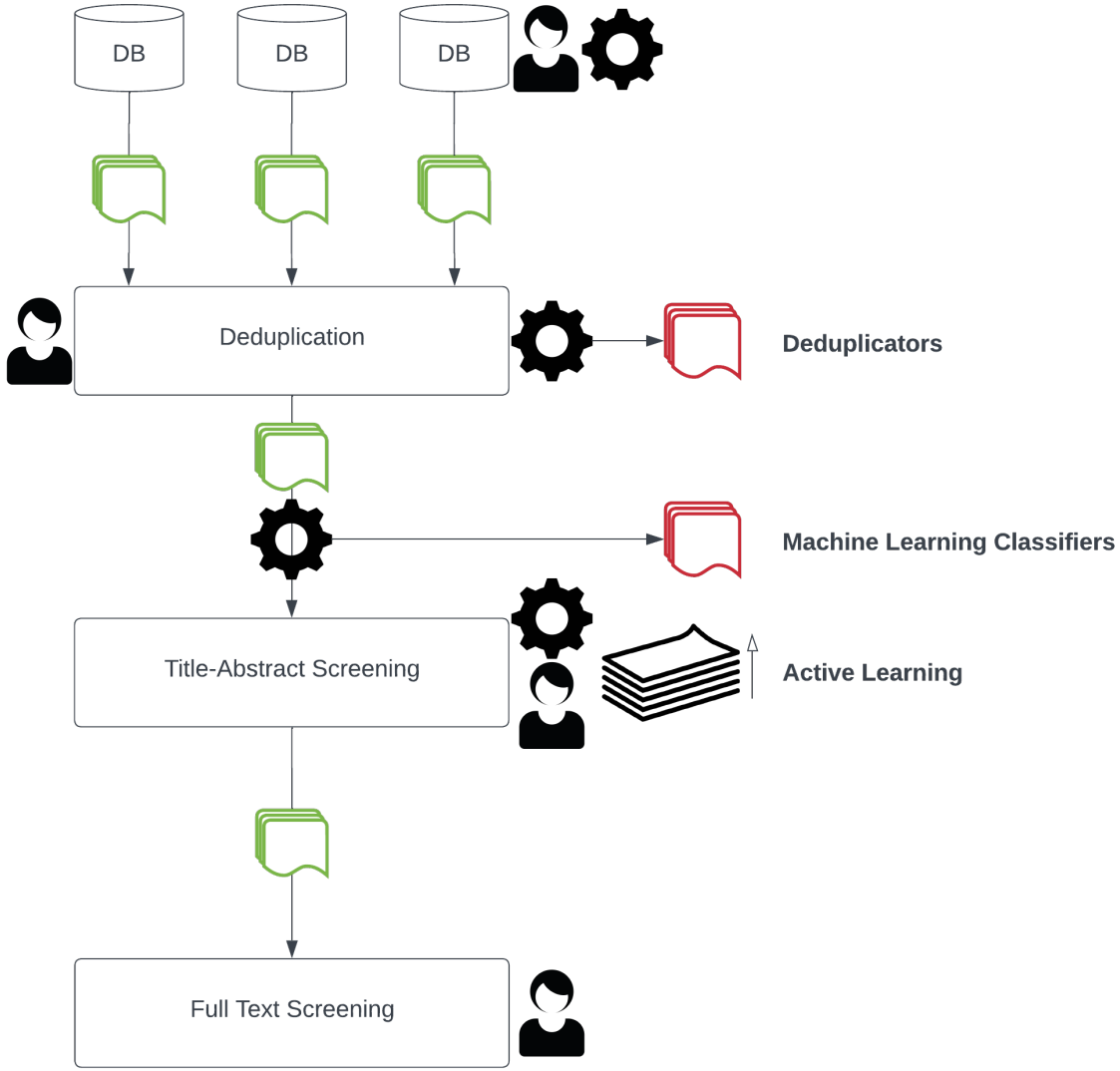
Standard information retrieval workflow for updating an (L)SR



Automation tools for study identification



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OpenAlex

Large ‘network graph’ dataset comprising >250M bibliographic records of research articles from across science

Open access – updated daily

Close-to-comprehensive coverage of journal articles

Key gaps for SRs and related use scenarios

- Trials registry records
- Conference abstracts
- Dissertations and Theses
- Grey literature

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Current stats

Last updated Tue Apr 23 2024

252M Works

56M Open Access works

31M from the Global South

4M datasets

92M Authors

5M with ORCIDs

12M from the Global South

255K Sources

45K that are Open Access

10K Publishers

32K Funders

108K Institutions

65K Concepts

5K Topics

252 Subfields

26 Fields

4 Domains

17 Sdgs

247 Countries

7 Continents

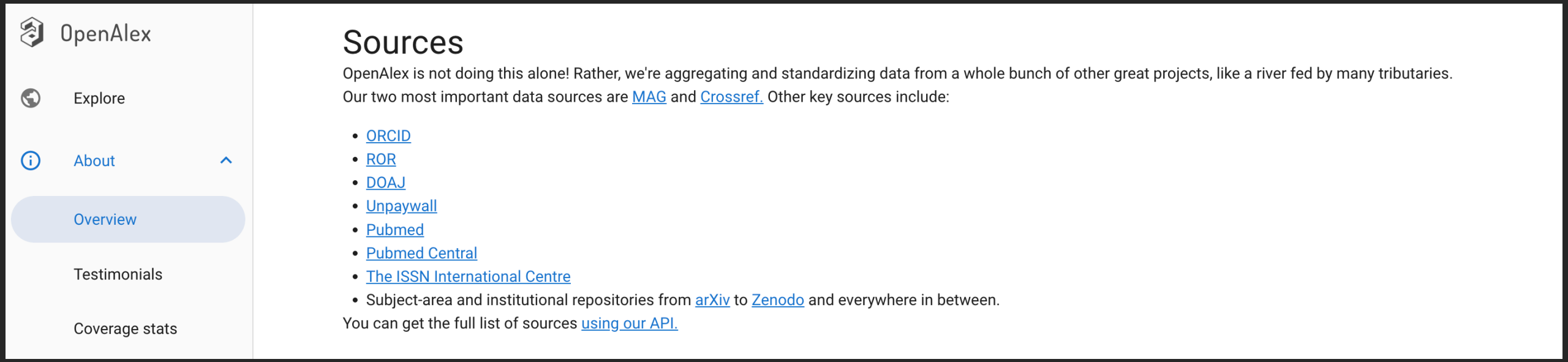
184 Languages

18 Types

6 Source-Types

8 Institution-Types

https://openalex.org/stats



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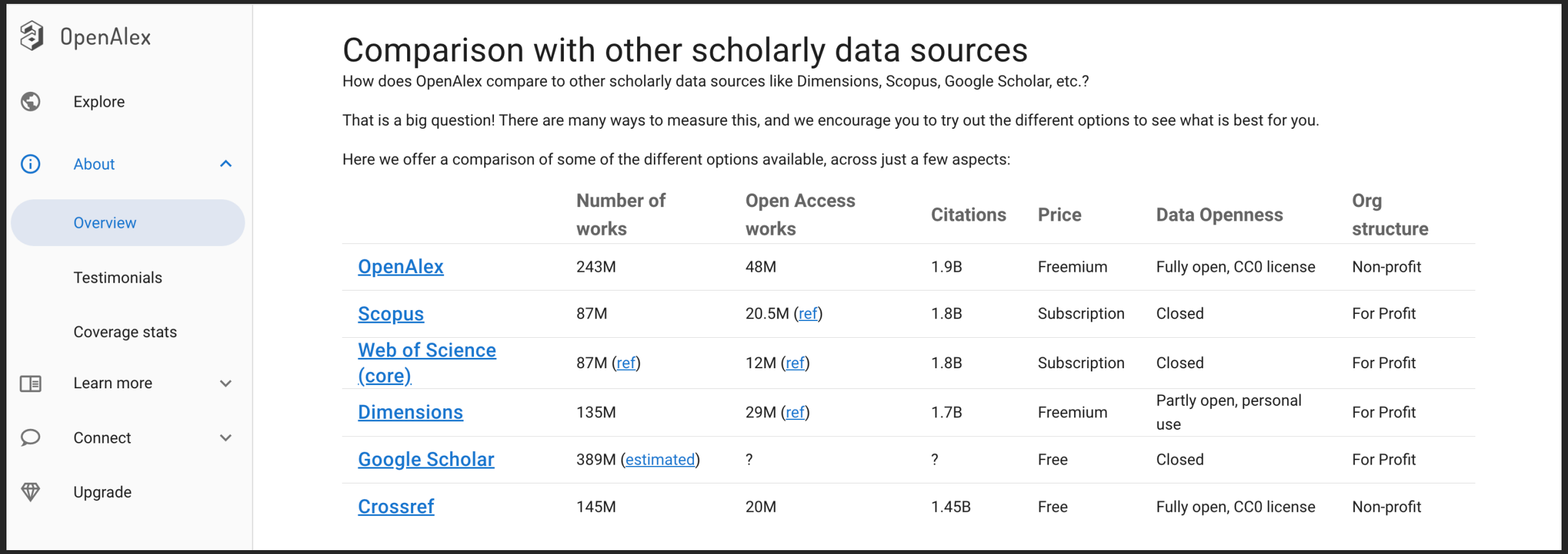


Comparison with other scholarly data sources

How does OpenAlex compare to other scholarly data sources like Dimensions, Scopus, Google Scholar, etc.?

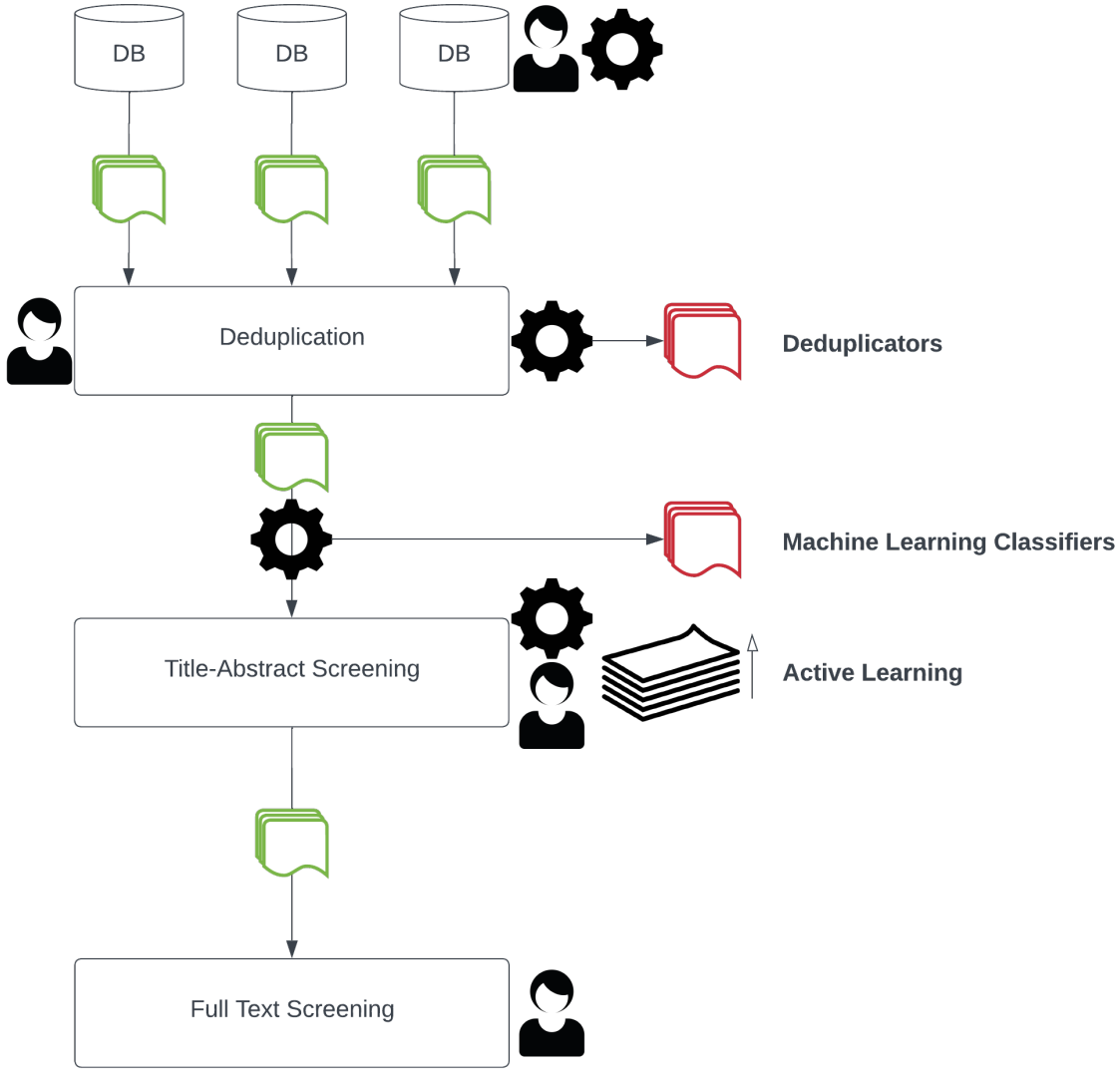
That is a big question! There are many ways to measure this, and we encourage you to try out the different options to see what is best for you.

Here we offer a comparison of some of the different options available, across just a few aspects:

	Number of works	Open Access works	Citations	Price	Data Openness	Org structure
OpenAlex	243M	48M	1.9B	Freemium	Fully open, CC0 license	Non-profit
Scopus	87M	20.5M (ref)	1.8B	Subscription	Closed	For Profit
Web of Science (core)	87M (ref)	12M (ref)	1.8B	Subscription	Closed	For Profit
Dimensions	135M	29M (ref)	1.7B	Freemium	Partly open, personal use	For Profit
Google Scholar	389M (estimated)	?	?	Free	Closed	For Profit
Crossref	145M	20M	1.45B	Free	Fully open, CC0 license	Non-profit

https://openalex.org/about

Automation tools for study identification



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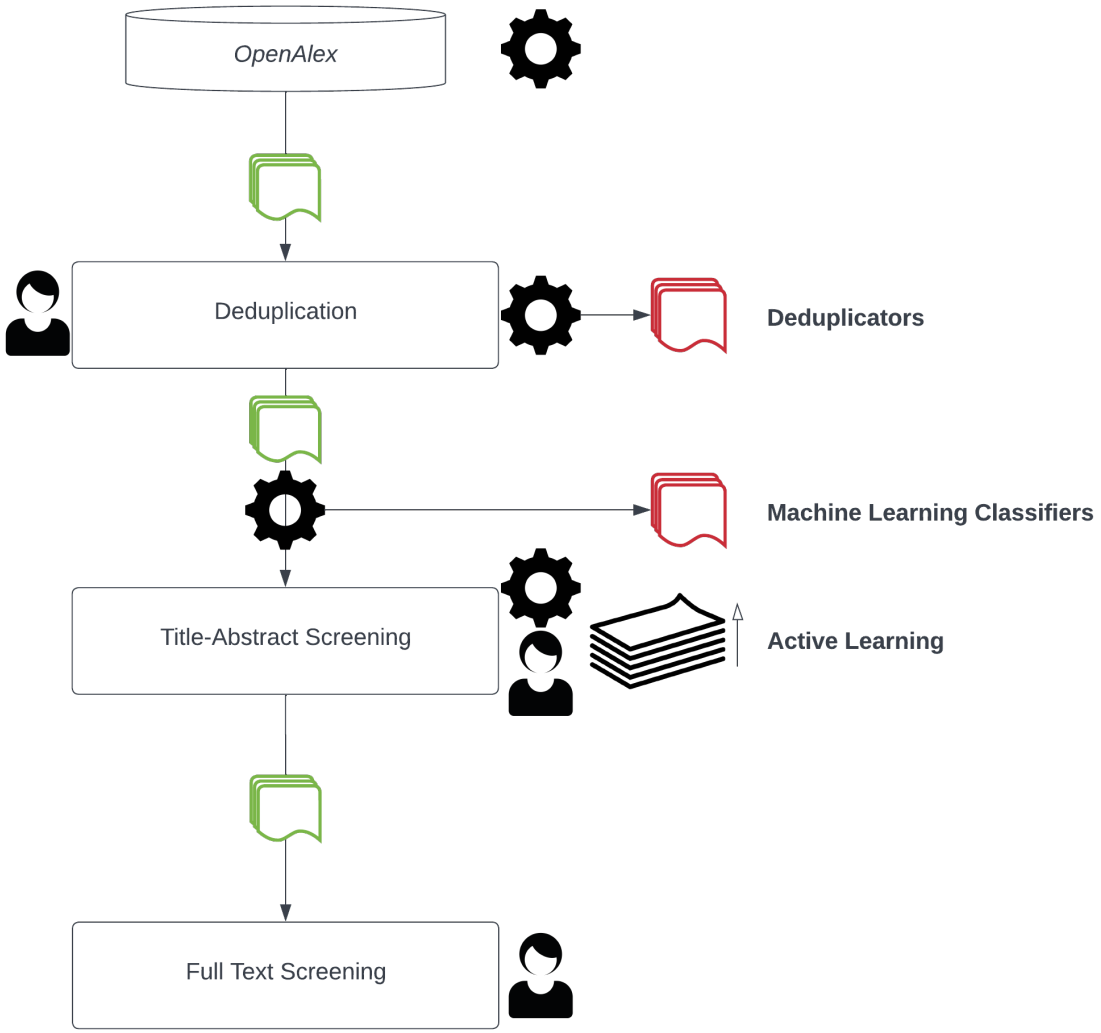

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OpenAlex methods research questions

1. Does OpenAlex contain the study reports we need to identify for systematic reviews, maps and other evidence syntheses? (*Recall*)

2. Can we use OpenAlex to efficiently identify the study reports we need for systematic reviews, maps and other evidence syntheses? (*Precision*)

OpenAlex = A large knowledge graph for open access research

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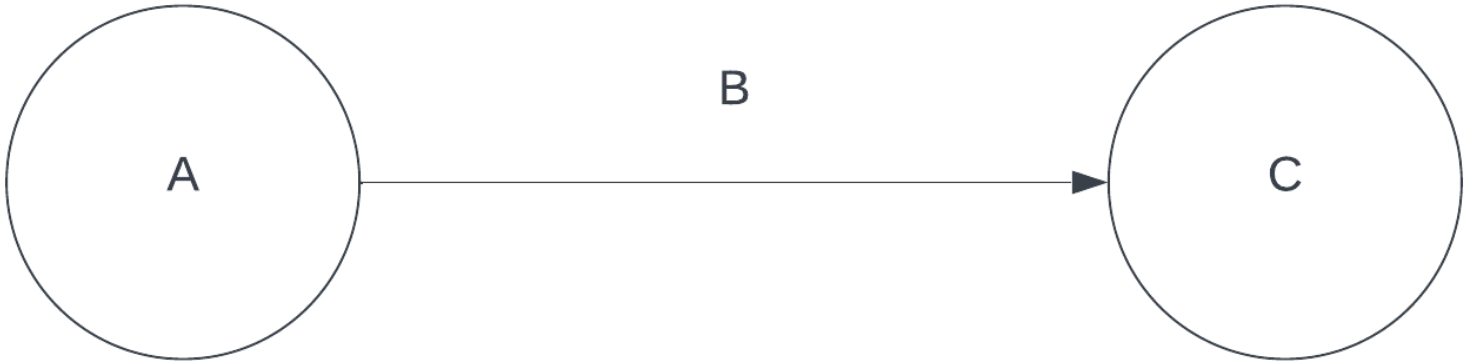
Knowledge graphs for open access research



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Knowledge graphs for open access research

RHEUMATOLOGY

Original article

The validation of a diagnostic rule for gout without joint fluid analysis: a prospective study

Laura B. E. Kienhorst[1], Hein J. E. M. Janssens[2,3], Jaap Fransen[4] and Matthijs Janssen[1]

Abstract

Objective. The gold standard for diagnosing gout is the identification of MSU crystals in joint fluid. In secondary care, the facilities or expertise to analyse joint fluid are not always available and gout is diagnosed clinically. To improve the predictive value of the clinical diagnosis of gout in secondary care, a diagnostic rule developed in primary care could be helpful. The aim of this study was to validate this diagnostic rule in a secondary care population with the gold standard as reference test.

Methods. Three hundred and ninety patients with monoarthritis were included. The variables of the diagnostic rule (male sex, previous arthritis attack, onset <1 day, joint redness, involvement of the first MTP joint, hypertension or one or more cardiovascular disease, and serum uric acid >5.88 mg/dl) were collected and scored. The affected joint was aspirated and joint fluid was analysed for the presence of MSU crystals.

Results. In 219 patients (56%) MSU crystals were found. The positive predictive value of a score of ⩾8 points was 0.87, the negative predictive value of a score of ⩽4 points was 0.95. The area under the receiver operating characteristic curve for the diagnostic rule was 0.86 (95% CI 0.82, 0.89). The Hosmer–Lemeshow goodness-of-fit test showed that the difference between the expected and the observed probability was non-significant ($P = 0.64$), indicating good agreement.

Conclusion. An easy-to-use diagnostic rule for gout developed in primary care shows good performance in secondary care and improves the predictive value of the clinical diagnosis of gout when joint fluid analysis is not available.

Key words: gout, diagnosis, diagnostic rule, joint fluid, synovial fluid.

Introduction

Gout is a common problem that affects 1.4% of the population [1]. The gold standard for diagnosing gout is the identification of MSU crystals in joint fluid by polarization microscopy [2, 3]. Although this is the gold standard, in some patients joint fluid analysis is not performed. Joint fluid analysis requires skills in arthrocentesis, the availability of polarization microscopy and familiarity with the use of this microscope. In primary care [4], but also in secondary care at emergency or other non-rheumatology departments, the facilities or expertise to analyse joint fluid are not always available. Even rheumatologists often lack these facilities or expertise [5, 6]. In that case gout is diagnosed based on clinical signs and symptoms only.

However, it has been shown that the validity of the clinical diagnosis of gout in primary care is limited [positive predictive value (PPV) 0.64 and negative predictive value (NPV) 0.87] [7]. To improve the predictive value of the clinical diagnosis of gout in primary care, a diagnostic rule (gout calculator) was developed [7]. This diagnostic rule includes scores for the following seven variables: male sex, previous patient-reported arthritis attack, onset within 1 day, joint redness, involvement of the first MTP joint (MTP-1), hypertension or one or more cardiovascular diseases and serum uric acid exceeding 5.88 mg/dl (equal to 0.35 mmol/l). The total score of the

Cited By

Clin Rheumatol (2014) 33:555–559
DOI 10.1007/s10067-014-2505-6

ORIGINAL ARTICLE

Additive value for ultrasonographic signal in a screening algorithm for patients presenting with acute mono-/oligoarthritis in whom gout is suspected

F. B. G. Lamers-Karnebeek · P. L. C. M. Van Riel · T. L. Jansen

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Abstract Crystal arthritides such as gout can be detected by ultrasonography (US). This study reveals the performance of joint US (double contour sign (DCS), tophus (T), hyperechoic spots eg, "snow storm" (SS)) for diagnosing gout and calcium pyrophosphate dihydrate crystal deposition disease (CPPD) in patients with acute mono- or oligoarthritis (MOA). The gold standard is the presence of monosodium urate (MSU)/CPPD crystals. Fifty-four Dutch patients had an acute MOA. US was performed on the following six joints maximum: the arthritic joint, the contra lateral side, metatarsophalangeal (MTP)-1, and knees bilaterally in case of arthritis in one of these joints. In case of wrist/PIP/MCP-arthritis, the knees and MTP-1 were scanned. These were examined for DCS, T, SS, and intercartilage rim (CPPD). Synovial fluid was aspirated from the affected joint for MSU proof. Twenty-six of the 54 (48 %) patients with MOA had MSU-proven gout. Sensitivity of DCS and any US abnormality (DCS, T, SS) was 77 and 96 %, respectively. The positive likelihood ratio (LR+) for DCS and any ultrasonographic abnormality (USabn) was 3.08 and 2.99, respectively, and the LR− was 0.31 and 0.06, respectively. In MSU-proven gout patients where the affected joint is not MTP-1, MTP-1 still showed USabn in 42 % of the patients. None of the CPPD patients had an intercartilage rim. In dedicated hands, ultrasonography deserves a place early in a screening algorithm of MOA patients, particularly if specificity is high enough to make punctures abundant or when microscopy is not available. In 86 % of the MSU-proven gout patients, the DCS is not present in another joint other than the affected or MTP-1 joint.

Keywords Double contour sign and diagnostic imaging · Gout · Mono- or oligoarthritis · Ultrasonography

Introduction

Acute arthritis deserves direct rheumatologic attention to establish the cause as soon as possible. Concerns have been raised for potential over diagnosis of rheumatoid arthritis (RA) with the use of newer diagnostic algorithms emphasizing sensitivity over specificity [1–4]. In populations with a high prevalence of crystal arthritis, this may be the case. With arthrocentesis and additional analysis of the synovial fluid/punctate, the crystal diagnosis and management can easily be obtained if microscopes are available. The identification of monosodium urate (MSU) crystals in the synovial fluid or tophaceous material is still the gold standard. This is ideally performed in the acute phase of arthritis or by puncturing a lesion suspected to be tophaceous [5]. Often, arthrocentesis and polarization microscopy is not easily organized, as the ubiquitous availability of polarized light microscopy is not optimal. Next to that, medicological complications may hamper performing intraarticular punctures in certain parts of the world. The advantages of an ultrasonography scan are no radiation for the patient, hardly time-consuming, and detecting early tophaceous accumulation subcutaneously and in deeper tissues. Conventional radiography shows only late effects of gout (established disease). So it will not help to diagnose gout in the early stage of the disease. During the last decade, several studies have suggested potentials for ultrasonography (US) in rheumatologic assessment of arthritis [6–11].

The following question is therefore important: Is ultrasonography a valuable option in a diagnostic strategy of arthritis? In gout and calcium pyrophosphate dihydrate crystal

Report ('Work') 1

Report ('Work') 2

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Concept (Topic)

‘Gout’

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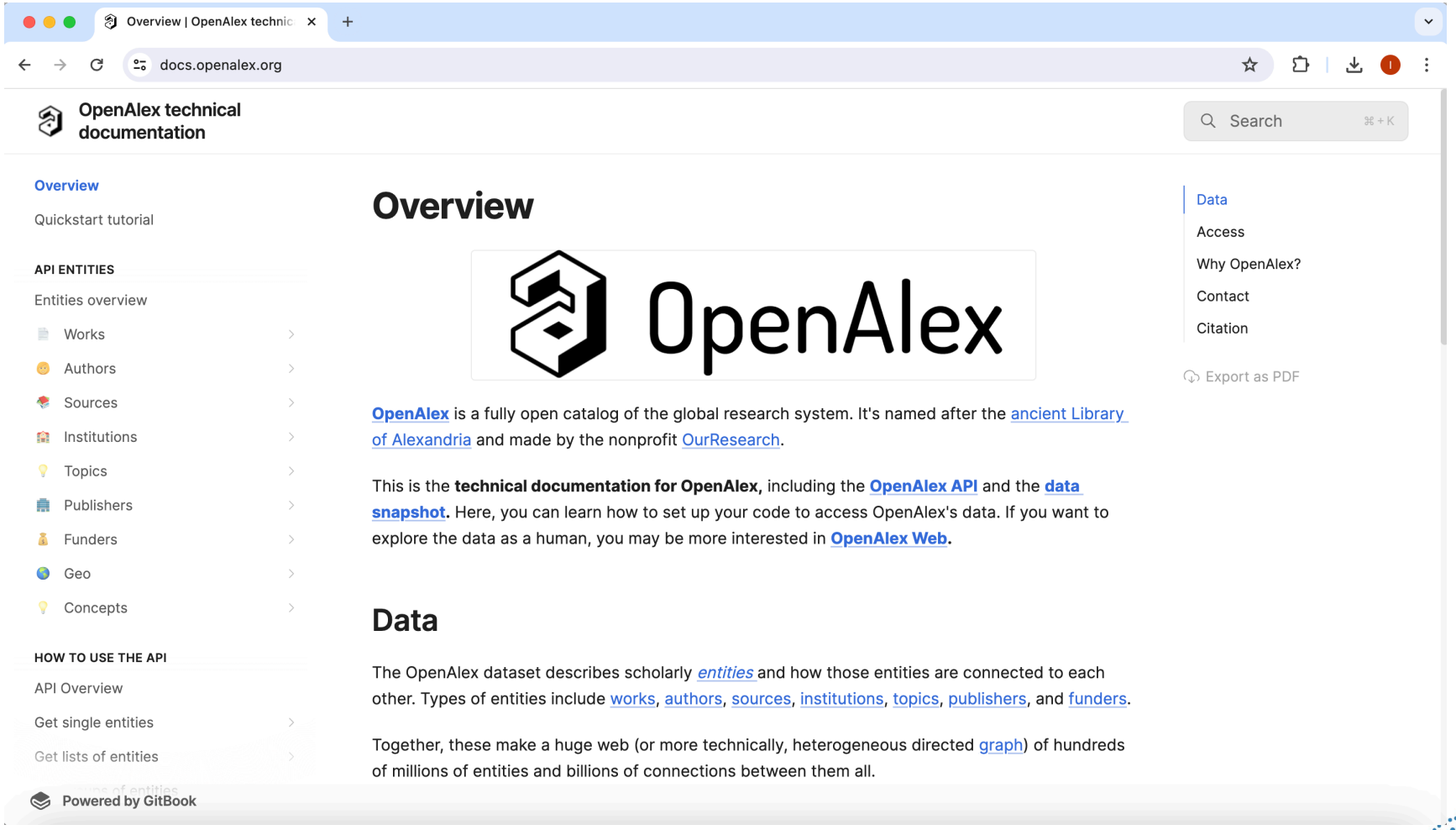

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Policy & Practice

OpenAlex methods research questions

1. Does OpenAlex contain the study reports we need to identify for systematic reviews, maps and other evidence syntheses? (*Recall*)

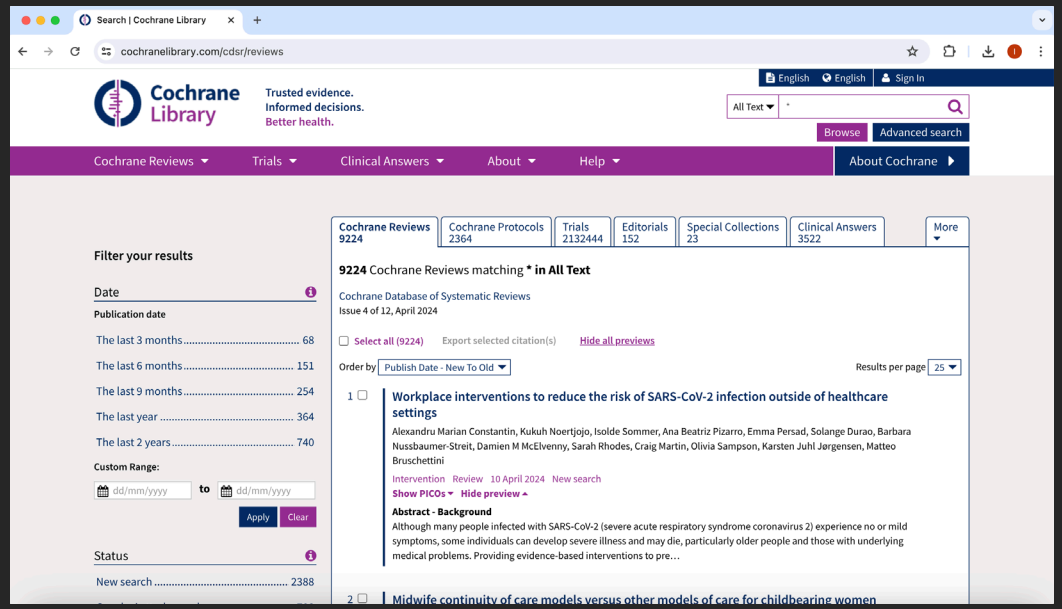
2. Can we use OpenAlex to efficiently identify the study reports we need for systematic reviews, maps and other evidence syntheses? (*Precision*)

OpenAlex methods research questions

1. Does OpenAlex contain the study reports we need to identify for systematic reviews, maps and other evidence syntheses? (*Recall*)

2. Can we use ***machine learning tools to exploit the network graph and text features of*** OpenAlex ***records*** to efficiently identify the study reports we need for systematic reviews, maps and other evidence syntheses? (*Precision*)

OpenAlex auto-update model for continual updating of larger Cochrane Reviews



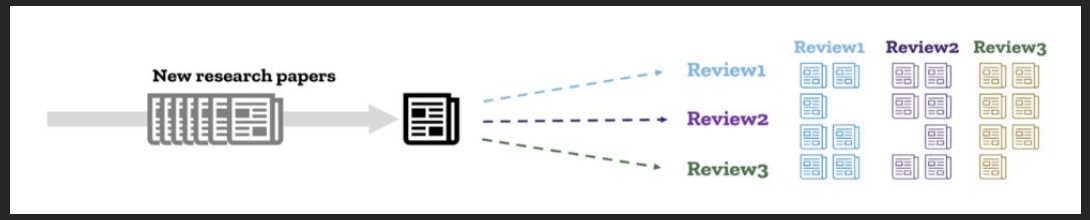


Table 1: Dataset statistics

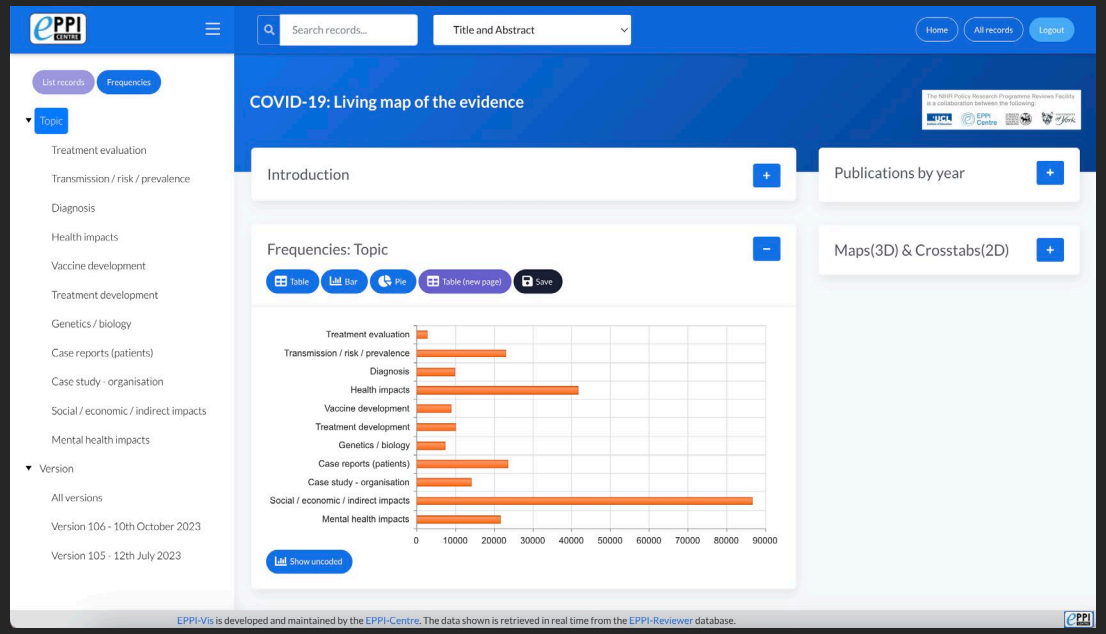
Num. unique reviews	644
Num. review domains	52
Num. unique publications	14574
Min publications per review	40

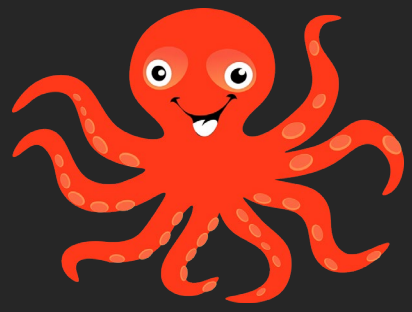
Table 2: **Experimental results.** Rows correspond to model runs with a certain set of features. Columns represent the best result for a certain threshold, in terms of the best precision with 0.97 recall. Training time is measured over a cloud virtual machine with a single GPU.

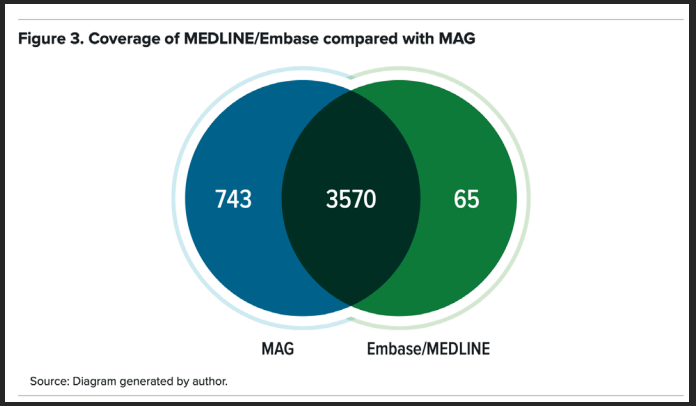
#	Features	Precision @97	Recall	Training time
	Baseline			
1	Auth-Cit-Tit	.593	.996	na
2	Auth-Cit-Tit-Abs	.352	.999	na
3	Auth-Cit-Tit-Abs-FoS-UMLS	.909	.988	2h29’
	With embeddings			
4	Auth-Cit-Tit-eTit	.947	.983	5h43’
5	Auth-Cit-Tit-eTit-Abs-eAbs	.96	.979	9h59’
6	Auth-Cit-Tit-eTit-Abs-eAbs-FoS-UMLS	.888	.989	na

doi.org/10.2139/ssrn.4406117

Automated OpenAlex searches with other ML tools for continual updating of a living map of COVID-19 research







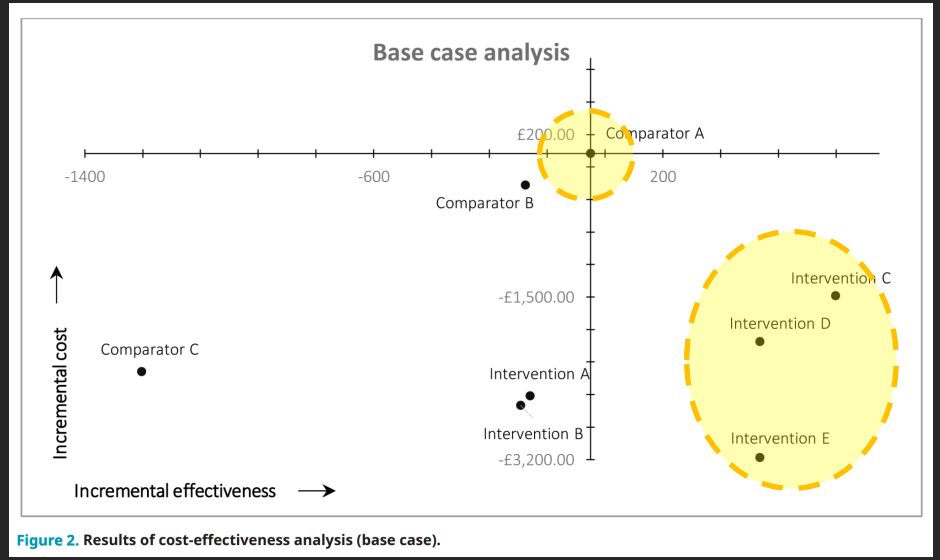
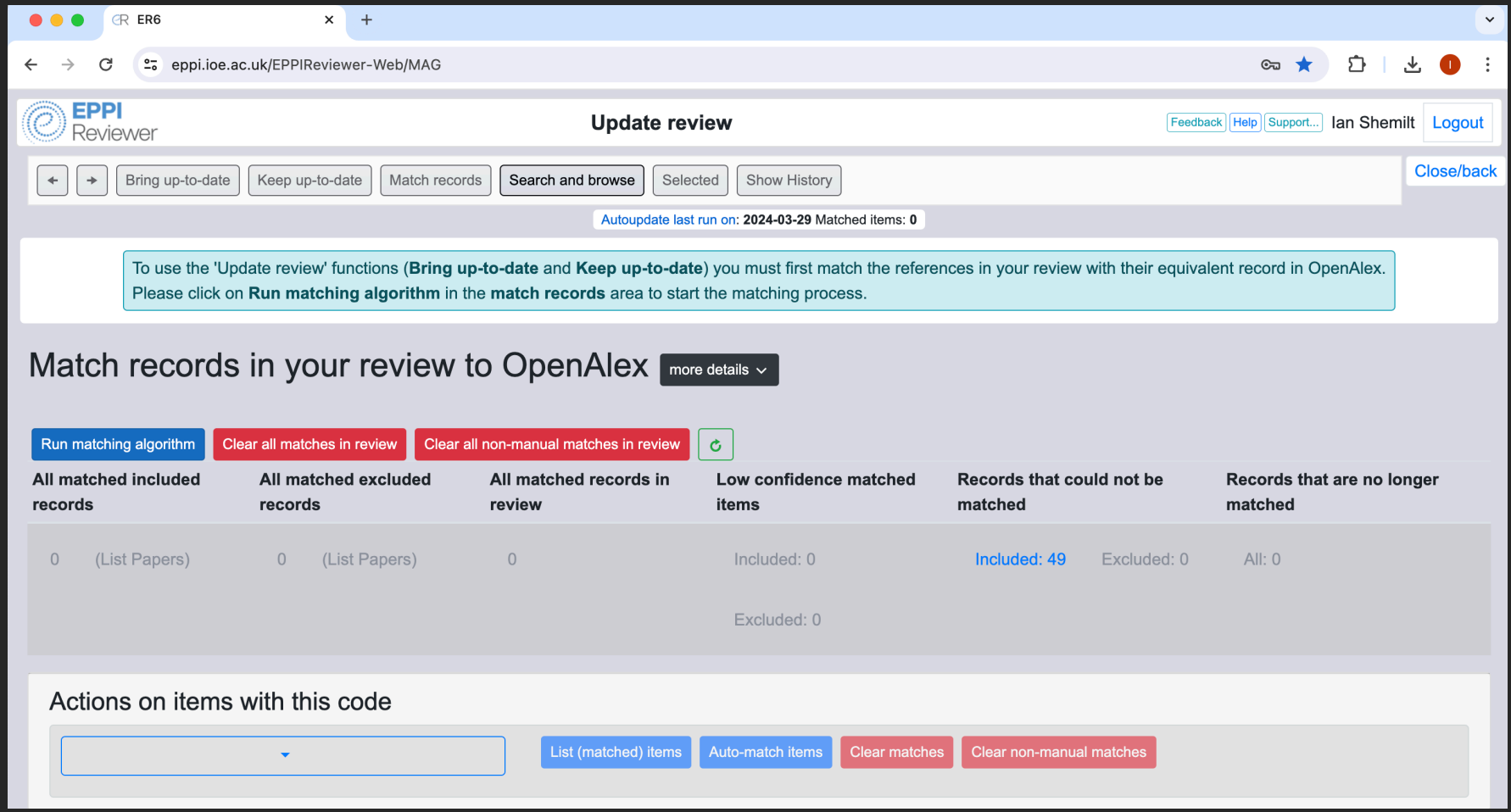


Figure 2. Results of cost-effectiveness analysis (base case).

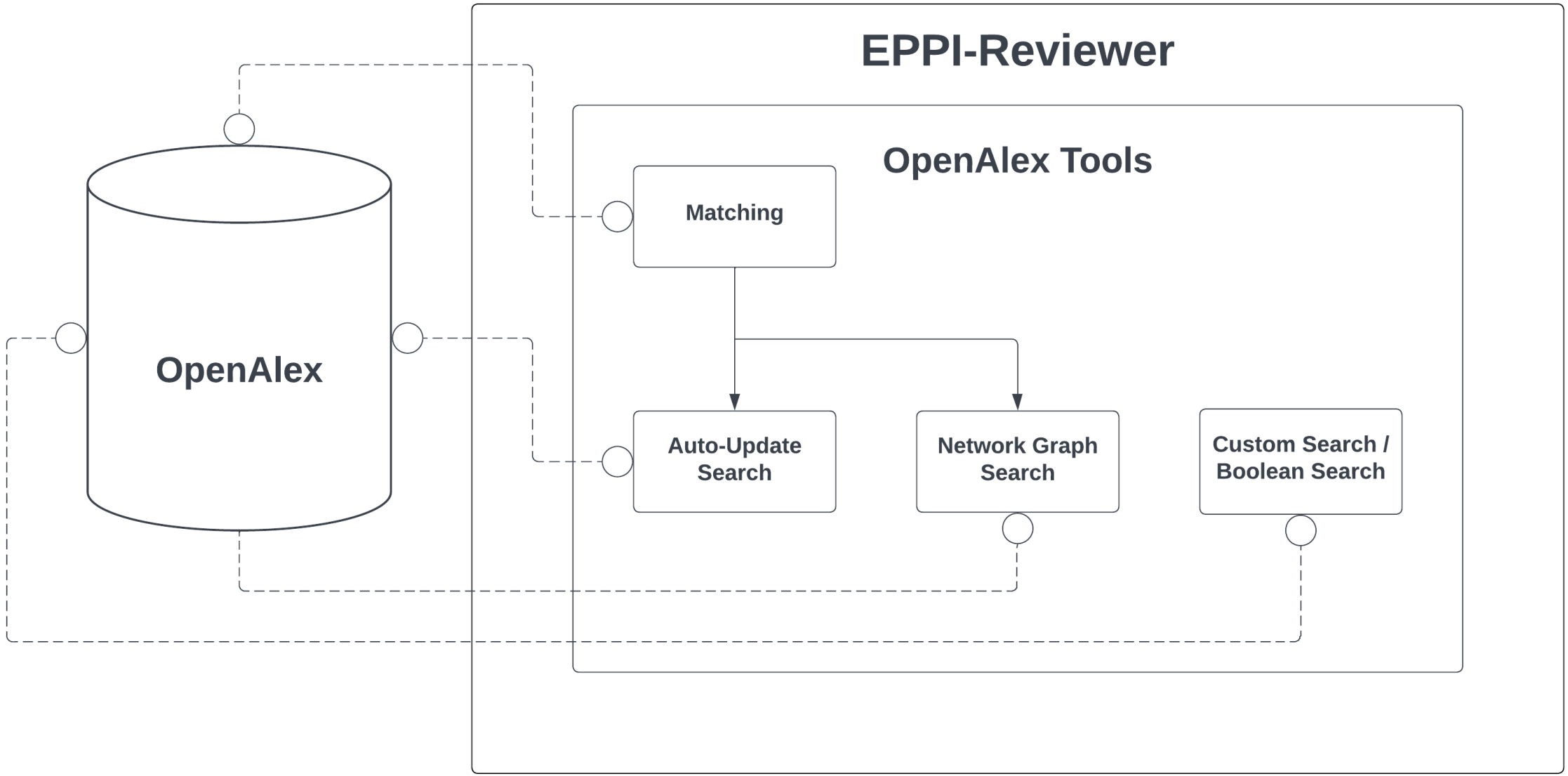
doi.org/10.12688/wellcomeopenres.17141.1

Further rigorous evaluations needed spanning a wide range of (L)SRs and (living) maps!

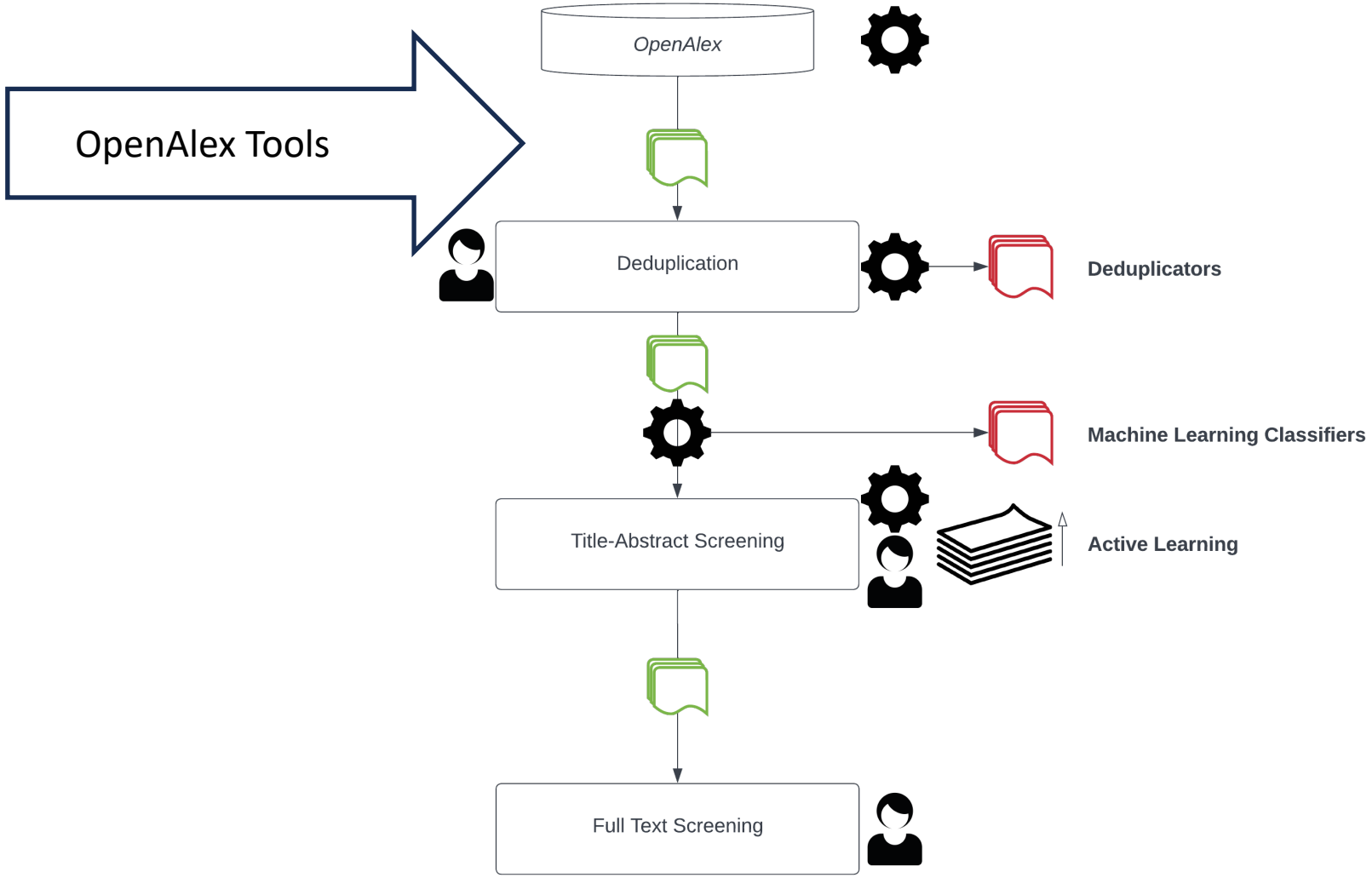
OpenAlex tools in EPPI Reviewer



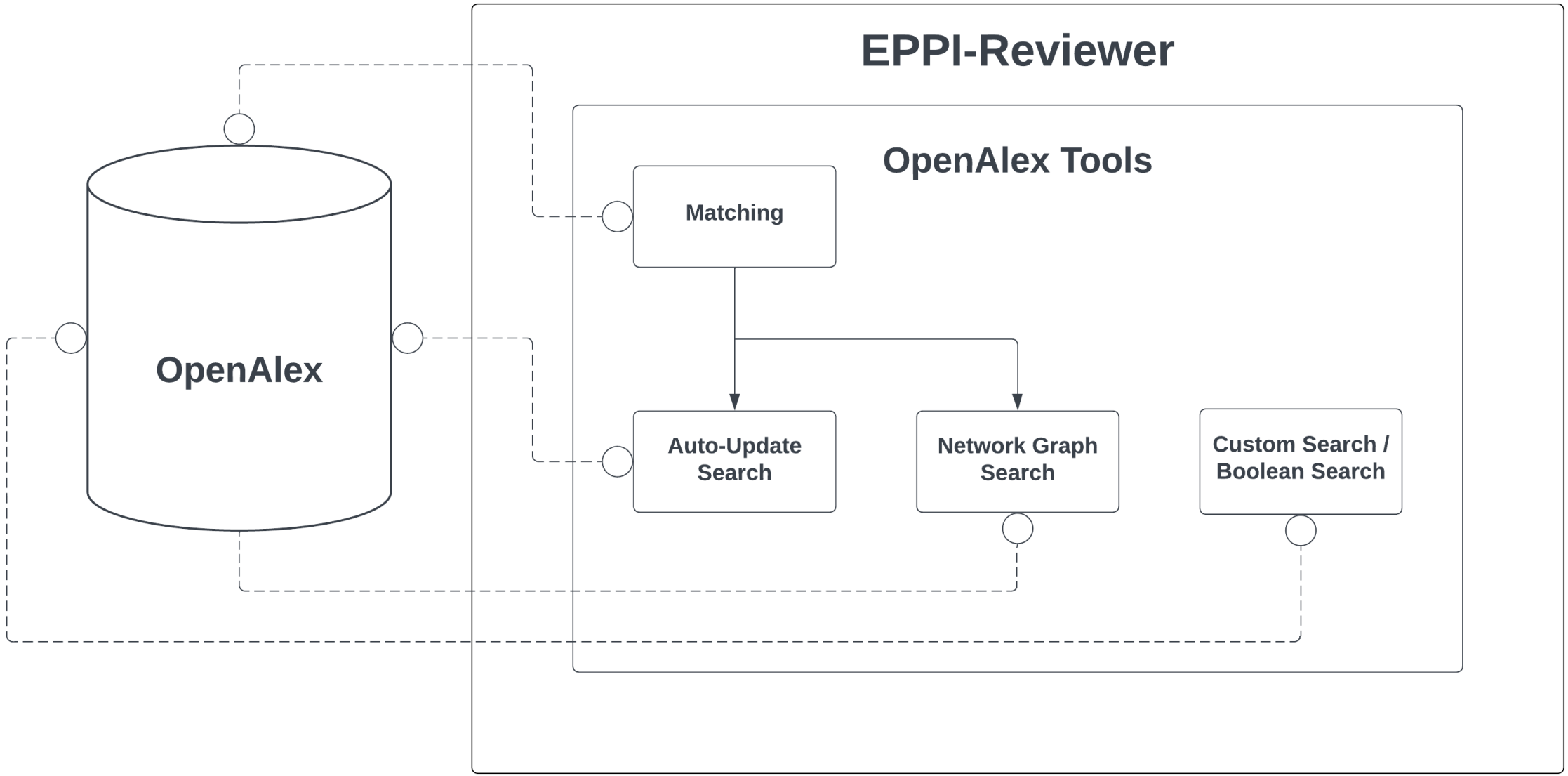
See eppi.ioe.ac.uk/cms/Default.aspx?tabid=3754 for further details


EPPI-Reviewer
OpenAlex Tools
Matching
OpenAlex
Auto-Update Search
Network Graph Search
Custom Search / Boolean Search

OpenAlex and ML tools for automated study identification



EPPI Centre
Evidence for
Policy & Practice

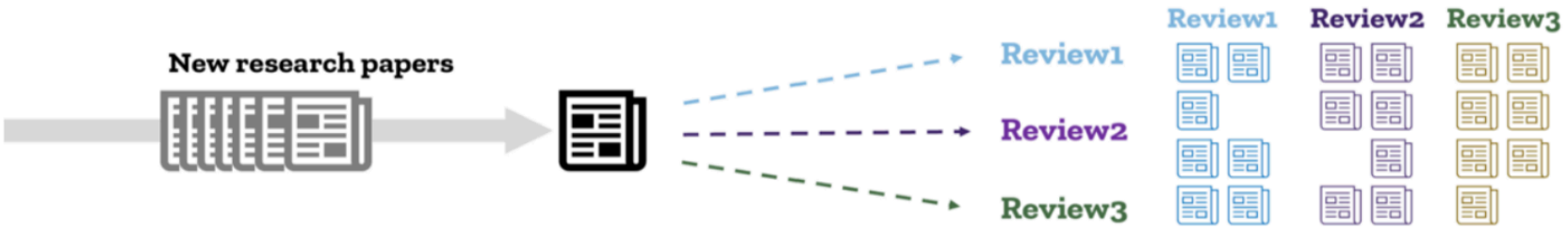

EPPI-Reviewer
OpenAlex Tools
Matching
OpenAlex
Auto-Update Search
Network Graph Search
Custom Search / Boolean Search

Auto-update

Problem Formulation

The domain entities are **papers** and **reviews**. Specifically, a review comprises a curated collection of papers that are relevant to a certain scientific question. The task is about recommending brand new papers to reviews owners based on their relevance to the scientific questions. More formally, the problem to address is the following:

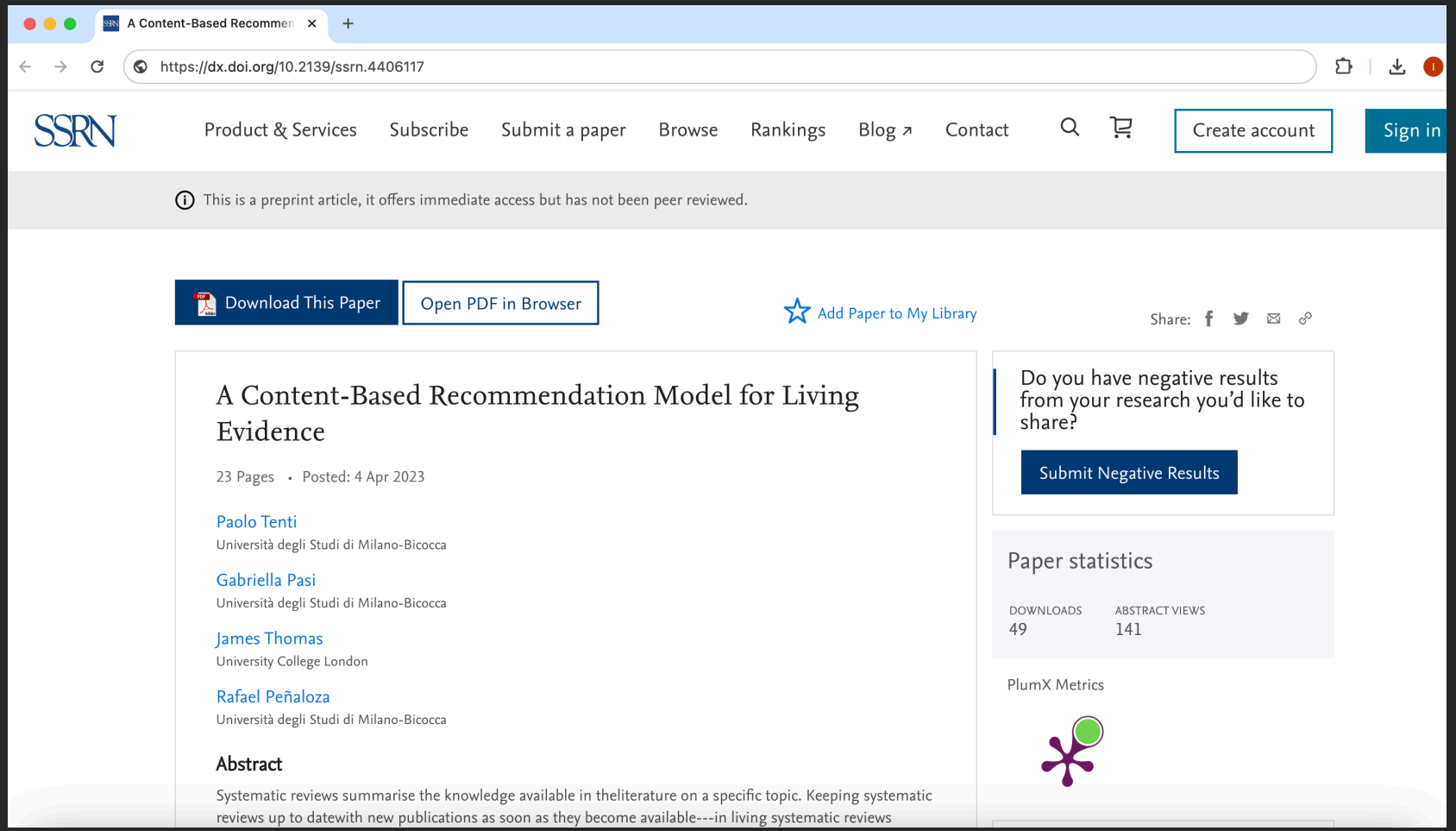
Given a set of reviews R, a set of papers P_r for each review $r \in R$ and a set of new papers P_{new}, the task is about finding a set of pairs (p, r) such that $p \in P_{new}$, $r \in R$ and p is relevant to r based on P_r.



Review1: "Empirical studies in Pair Programming for CS/SE Teaching in Higher Education"
Review2: "Interventions for recruiting smokers into cessation programmes"
Review3: "Immunotherapy (oral and sublingual) for food allergy to fruits"

devblogs.microsoft.com/ise/developing-and-deploying-a-recommender-model-for-continuous-systematic-literature-reviews

Auto-update – ML model technical details



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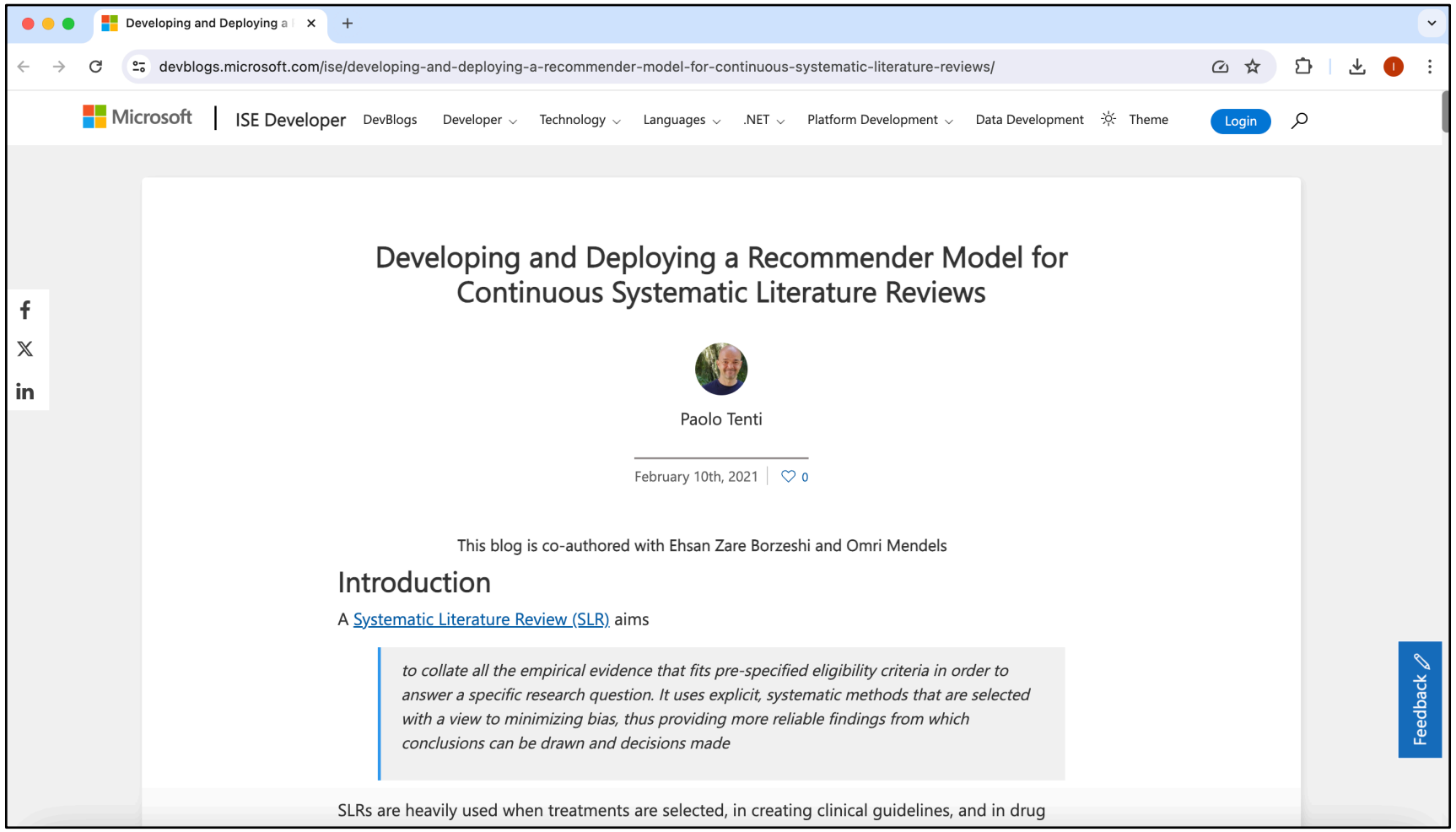
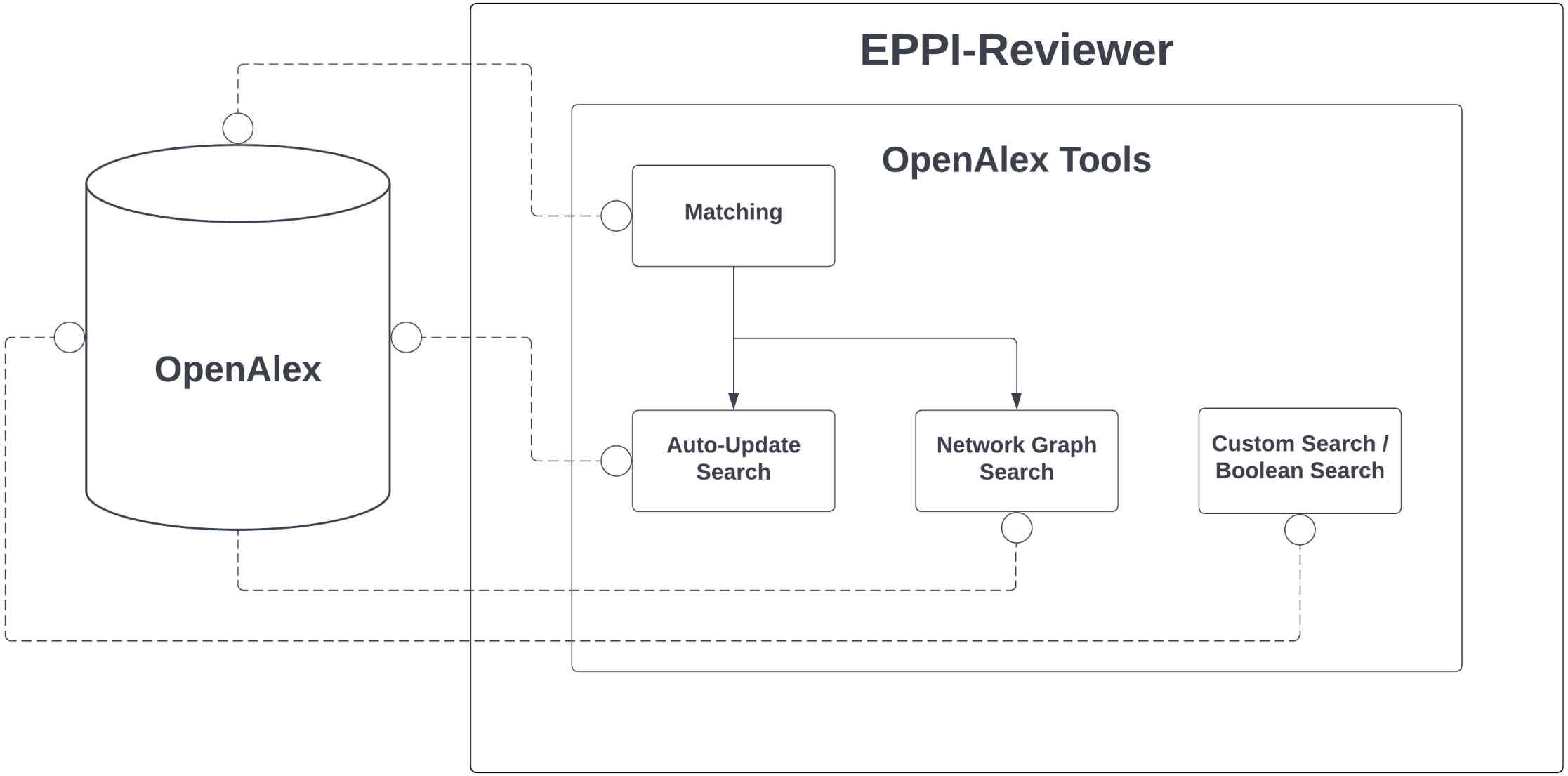
Auto-update – ML model technical details



devblogs.microsoft.com/ise/developing-and-deploying-a-recommender-model-for-continuous-systematic-literature-reviews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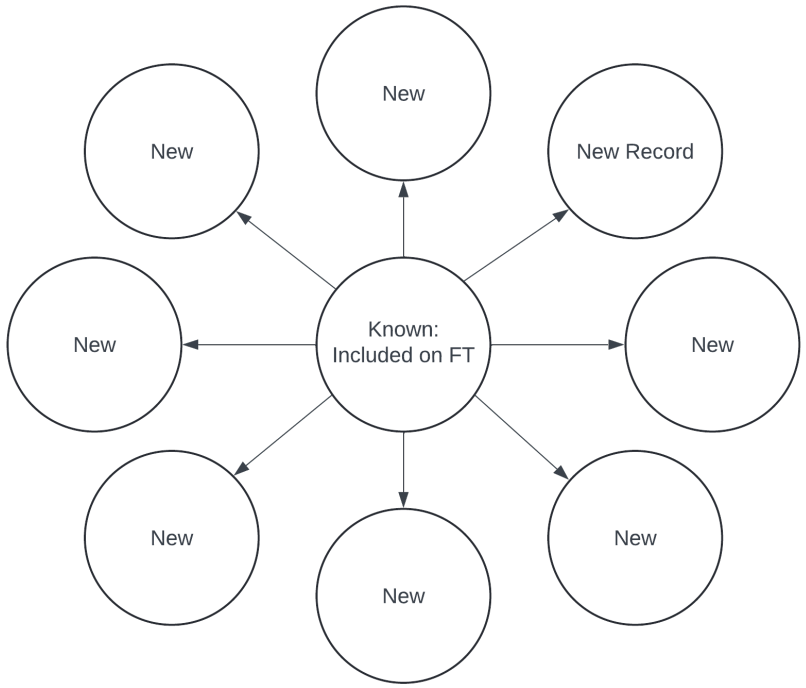
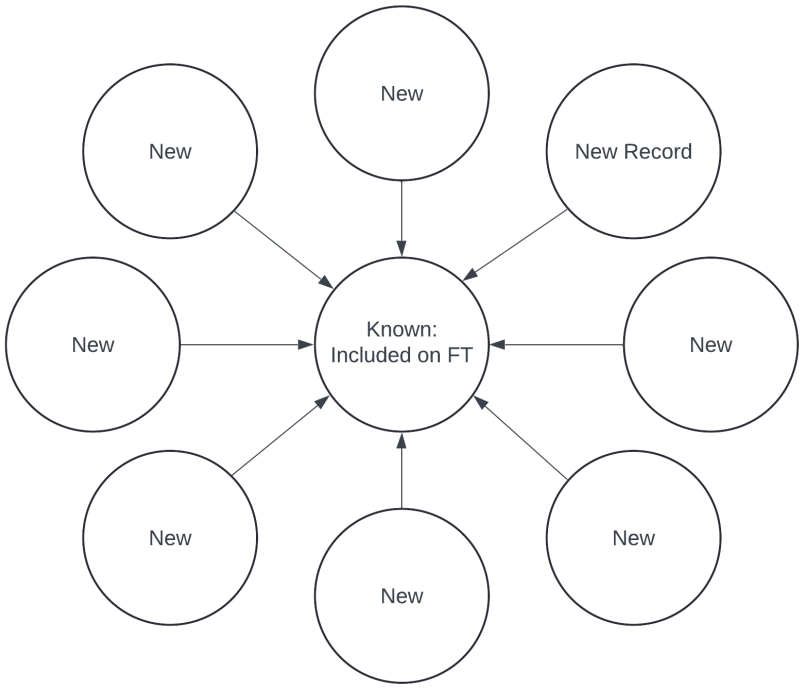
Network graph search – three modes



Bibliography
'New' in the bibliography of 'Known'



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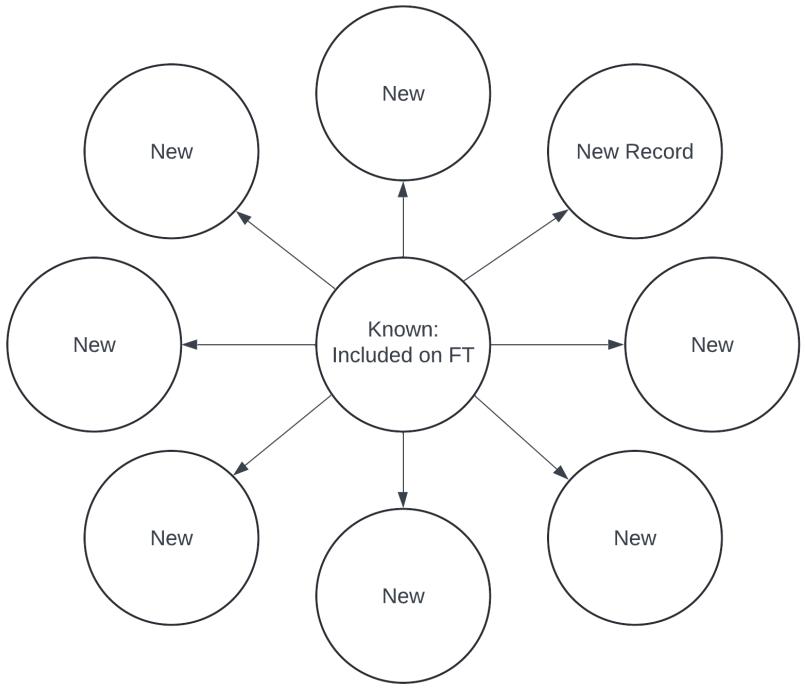

Recommended by
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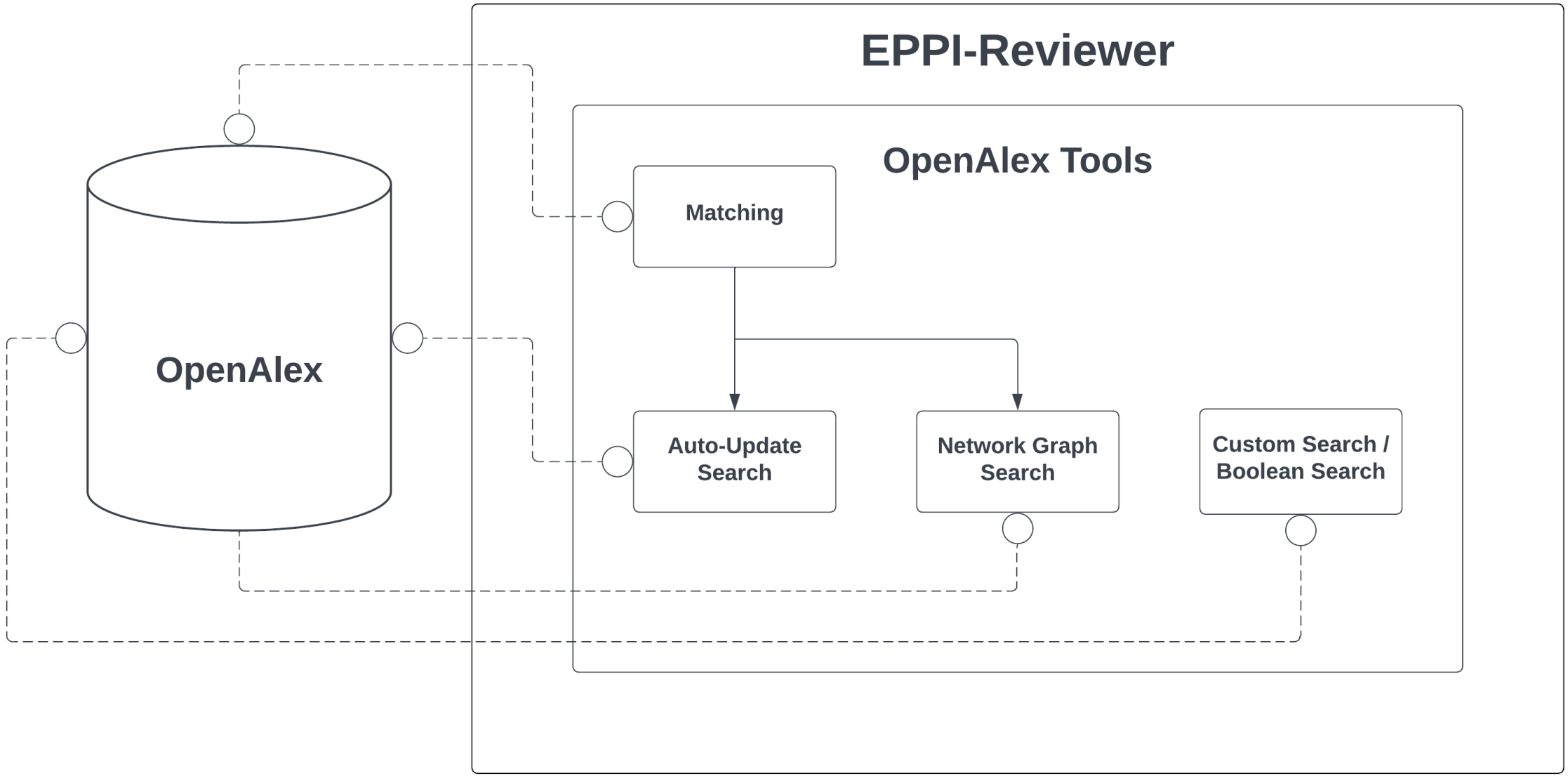
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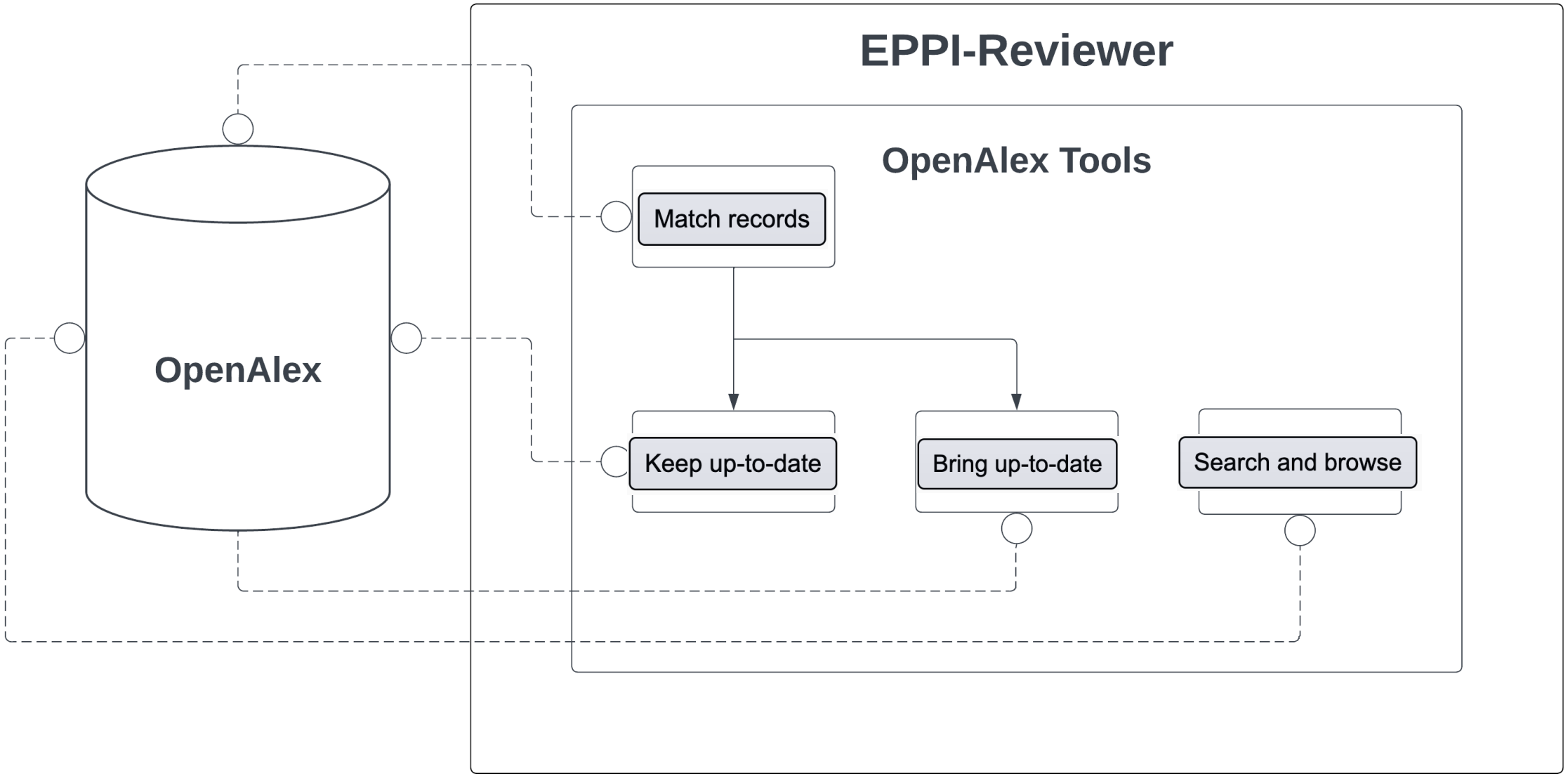

EPPI-Reviewer
OpenAlex Tools
Match records
OpenAlex
Keep up-to-date
Bring up-to-date
Search and browse

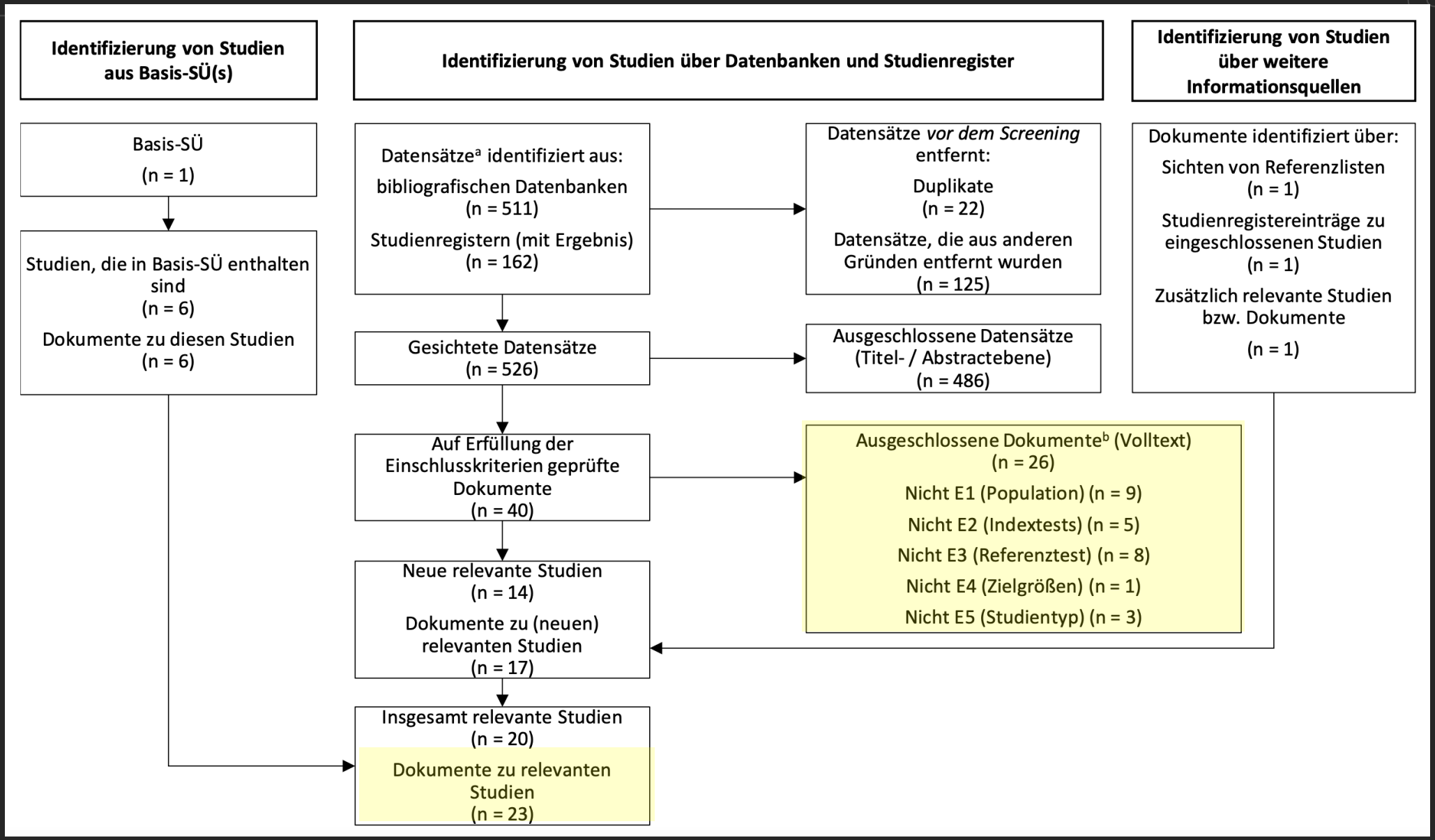
- Recent IQWiG evidence search for the S3 guideline on diagnosis and treatment of gout (v1.0)
- Search is for diagnosis (not treatment): reports of studies of accuracy of various procedures for diagnosing gout among adult patients with suspected gouty arthritis
- Conventional searches for primary studies:
  - Ovid MEDLINE(R) ALL 1946 to April 21, 2023
  - CENTRAL (The Cochrane Library): Issue 4, April 2023
  - 2 x CTRs
- Hypothetical use scenario = Update this evidence search using OpenAlex tools only: April 2023 to present day

https://www.iqwig.de/en/projects/v22-06.html

Tabelle 1: Übersicht über die Kriterien für den Studieneinschluss

Einschlusskriterien	
E1	Population: erwachsene Patientinnen und Patienten mit Verdacht auf eine Gichtarthritis
E2	Indextests: ▪ klinischer Diagnosescore (Gicht-Kalkulator), Cut-off: 8 [8] ▪ DECT ▪ Röntgen ▪ Ultraschall ▫ Doppelkontur-Zeichen und / oder Tophus[a] gemäß der OMERACT-Definition [9] – jegliche Auffälligkeit ▫ Doppelkontur-Zeichen und / oder Weichteil-Ansammlungen von Natriumuratkristallen und / oder Tophi[a] – jegliche Auffälligkeit ▫ Doppelkontur-Zeichen und / oder Schneesturm-Erscheinungsbild und / oder Tophus[a] – jegliche Auffälligkeit ▪ ACR- / EULAR-Klassifikationskriterien, Cut-off: 8 [10] Es werden ausschließlich Indextests herangezogen, die den Referenztest nicht beinhalten.
E3	Referenztest: mikroskopischer Nachweis von Natriumuratkristallen in der Gelenkflüssigkeit oder im periartikulären Gewebe
E4	Zielgrößen: personenbezogene Vierfeldertafel-Daten zur Berechnung der diagnostischen Güte (z. B. Sensitivität, Spezifität)
E5	Studientyp: prospektiv geplante Querschnitts- und Kohortenstudien[b]
E6	Publikationssprache: Deutsch oder Englisch
E7	Vollpublikation verfügbar[c]

https://www.iqwig.de/en/projects/v22-06.html



https://www.iqwig.de/en/projects/v22-06.html

Contact information

- Email: i.shemilt@ucl.ac.uk
- Profile: profiles.ucl.ac.uk/56390-ian-shemilt

- EPPI Centre: eppi.ioe.ac.uk/cms
- EPPI Reviewer: eppi.ioe.ac.uk/cms/er

EPPI Centre
Evidence for
Policy & Practice

UCL

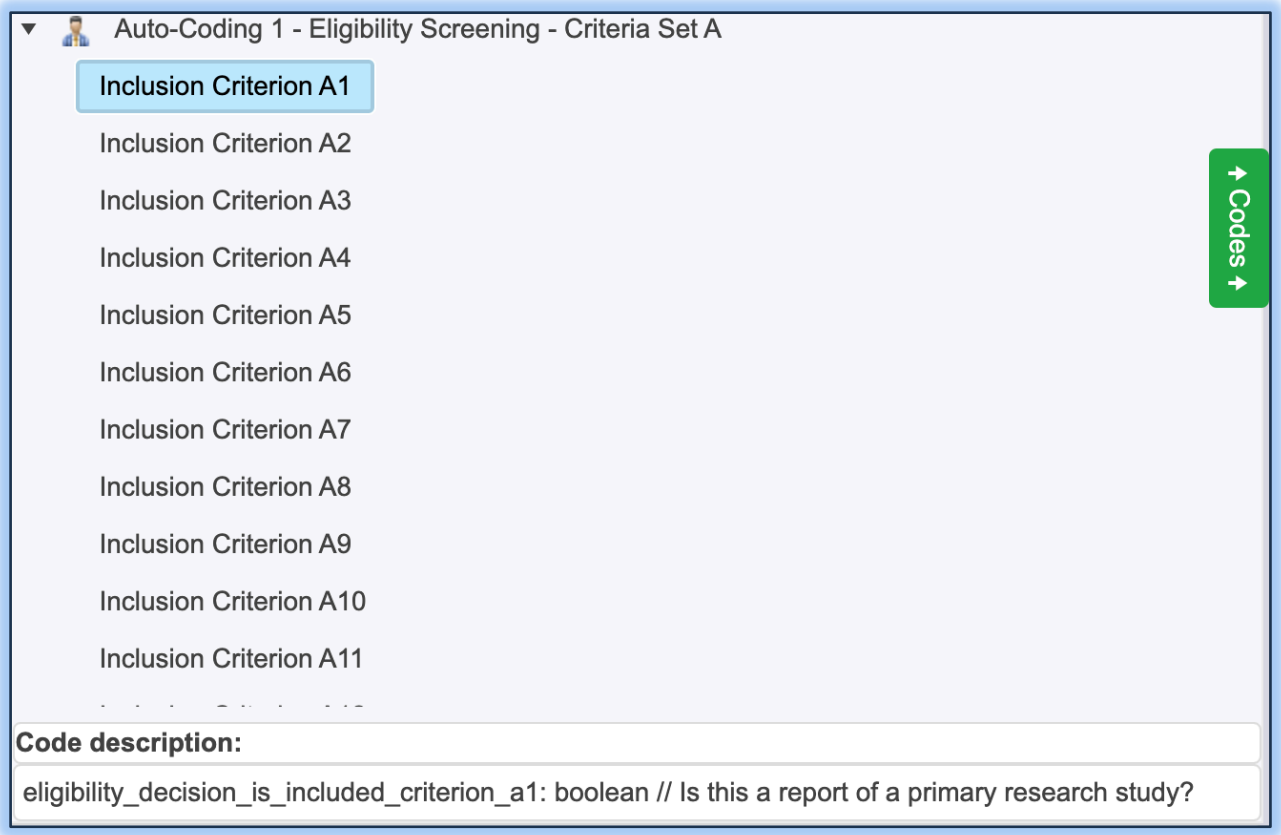
IQWiG Information Retrieval Meeting (IRM 2024): Software and data skills for information specialists

Introduction to OpenAlex tools for efficient automated updating of systematic reviews and maps

Ian Shemilt

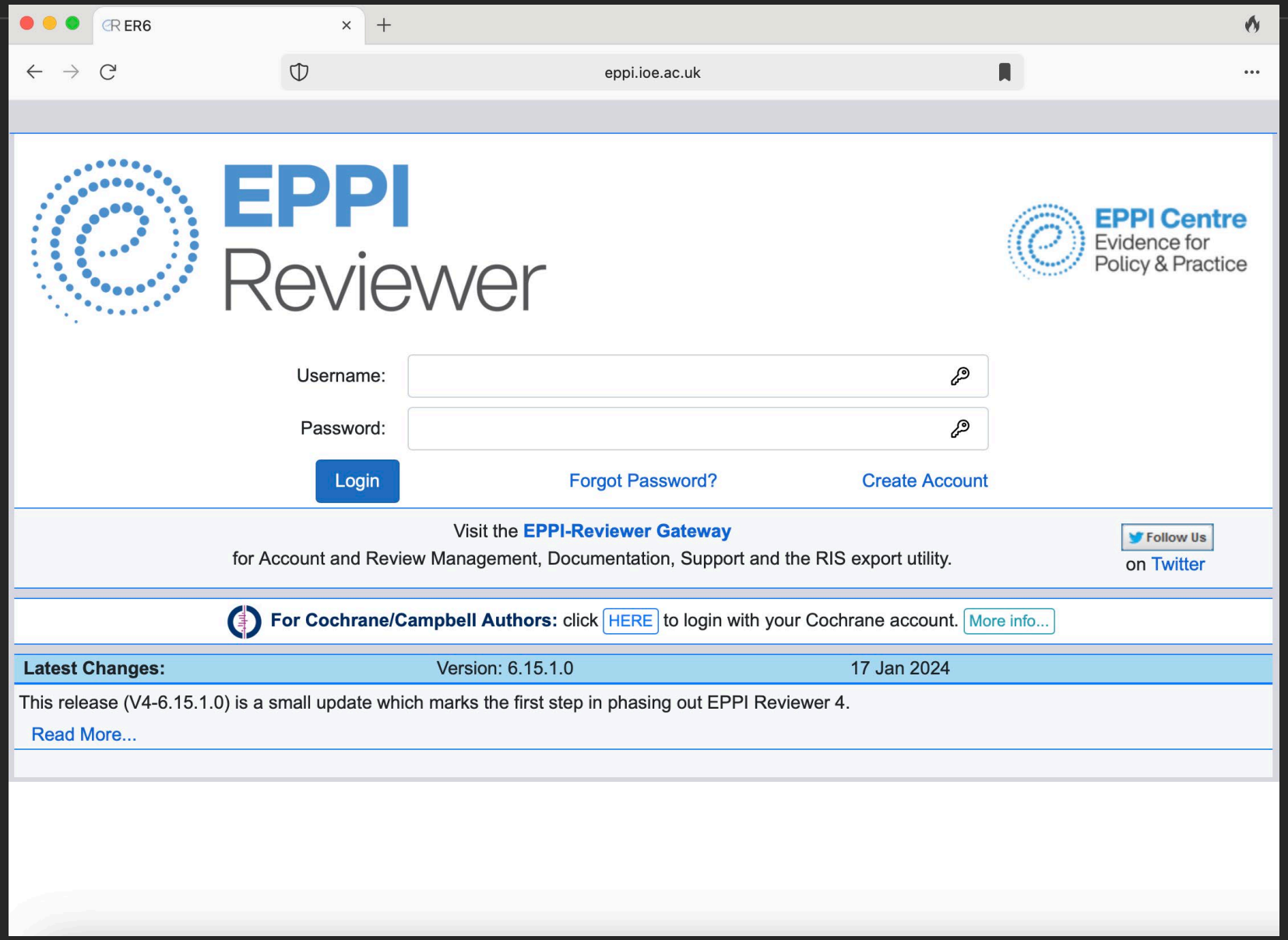
EPPI Centre, University College London, United Kingdom

GPT-4 auto-coding in EPPI Reviewer for eligibility screening



- ~82-85% agreement vs. human decisions (n=100)
- Recall ~0.95-0.98
- Precision ~0.86-0.87

- eligibility_decision_is_included_criterion_a1: boolean // Is this a report of a primary research study?
- eligibility_decision_is_included_criterion_a2: boolean // Is this a report of a systematic review?
- eligibility_decision_is_included_criterion_a3: boolean // Is this a report of a living systematic review?
- eligibility_decision_is_included_criterion_a4: boolean // Is this a report of a meta-analysis?



https://eppi.ioe.ac.uk/cms/er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REPOSITORIES

SWAT Store

SWAR Store

SWAR ID	Title	Link (Author(s) & Date)
1)	Citation screening in systematic reviews: Two approaches, two authors and time taken	SWAR01 Declan Devane (2022 Aug 25 1552)
2)	Effects of reading a written summary or listening to a summary podcast on knowledge and understanding of the findings of a systematic review.	SWAR02 Mike Clarke (2013 JUL 19 1210)
3)	To investigate whether supplying Cochrane Eyes and Vision review author teams	SWAR 03 Jennifer Evans and

https://www.qub.ac.uk/sites/TheNorthernIrelandNetworkforTrialsMethodologyResearch/SWATSWARInformation/Repositories/SWARStore/

New SWAR Registration – Forthcoming (Draft)

1. To assess the effectiveness and efficiency of conducting automated searches of the OpenAlex dataset – using OpenAlex tools in EPPI-Reviewer – compared with conventional electronic searches of multiple literature databases, for continual or regular updating of (living) systematic reviews.

2. To assess the effectiveness and efficiency of using bespoke, binary machine learning classifiers – trained, calibrated, evaluated and deployed using machine learning tools in EPPI-Reviewer – to automatically exclude some records prior to the citation (title-abstract) screening stage, compared with screening all citations, for continual or regular updating of (living) systematic reviews.