#### **REVIEW ARTICLE**



## Artificial Intelligence Based on Machine Learning in Pharmacovigilance: A Scoping Review

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#### Abstract

**Introduction** Artificial intelligence based on machine learning has made large advancements in many fields of science and medicine but its impact on pharmacovigilance is yet unclear.

**Objective** The present study conducted a scoping review of the use of artificial intelligence based on machine learning to understand how it is used for pharmacovigilance tasks, characterize differences with other fields, and identify opportunities to improve pharmacovigilance through the use of machine learning.

**Design** The PubMed, Embase, Web of Science, and IEEE Xplore databases were searched to identify articles pertaining to the use of machine learning in pharmacovigilance published from the year 2000 to September 2021. After manual screening of 7744 abstracts, a total of 393 papers met the inclusion criteria for further analysis. Extraction of key data on study design, data sources, sample size, and machine learning methodology was performed. Studies with the characteristics of good machine learning practice were defined and manual review focused on identifying studies that fulfilled these criteria and results that showed promise. **Results** The majority of studies (53%) were focused on detecting safety signals using traditional statistical methods. Of the studies that used more recent machine learning methods, 61% used off-the-shelf techniques with minor modifications. Temporal analysis revealed that newer methods such as deep learning have shown increased use in recent years. We found only 42 studies (10%) that reflect current best practices and trends in machine learning. In the subset of 154 papers that focused on data intake and ingestion, 30 (19%) were found to incorporate the same best practices.

**Conclusion** Advances from artificial intelligence have yet to fully penetrate pharmacovigilance, although recent studies show signs that this may be changing.

#### **Key Points**

We reviewed 393 papers in the intersection of pharmacovigilance (PV) and machine learning, and most involved signal detection as opposed to data intake or data analysis.

There has been a rapid rise in the use of deep learning in the PV literature, but corresponding dramatic success has not been seen in other fields such as computer vision, natural language processing, and healthcare.

There are opportunities to implement machine learning approaches throughout the PV pipeline.

Link to PRISMA-ScR Checklist: http://prisma-statement.org/Exten sions/ScopingReviews.

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## **1** Introduction

Pharmacovigilance (PV) is fundamentally a data-driven field as it requires the collection, management, and analysis of a large amount of data gathered from a wide range of disparate sources [1]. The primary type of data used in PV are individual case safety reports (ICSRs), which are records of suspected adverse events collected via multiple channels, aggregated and organized into large databases, and constantly monitored to detect safety signals [2]. ICSRs come from a multitude of sources, including chatbot interactions, electronic health records (EHRs), published literature, patient registries, patient support programs, or even directly from patients via social media [3]. Reports are collected worldwide and characterized by heterogeneity in format, language, and unique characteristics of the underlying healthcare systems. Adverse events must be identified and analyzed in order to find potential emerging safety issues in medicines and vaccines.

The central challenge of PV is how to make sense of these large and heterogeneous data to quickly and reliably find the 'needles in the haystack,' which are safety signals that require escalation and triage [4]. Given the rise of artificial intelligence (AI) powered by new advancements in machine learning (ML) across many fields of science [5-7] and medicine [8-11] over the last decade [12, 13], many have speculated [14, 15] that these same technologies could be brought to bear on the core problems of PV. The use of these methods for human safety data first appeared in the early 1990s [16] and has steadily increased since the 2000s. The goal of this review is to systematically identify works that use ML, broadly defined, for safety data to characterize the current state of ML in PV, and to provide clarity on ways that recent advances in AI and ML can be translated to improve various components of PV.

Care must be taken when attempting to extrapolate the success of ML in other areas compared with PV since there are specific factors that may account for the recent success of ML that may or may not be present for PV applications [15]. More so than any other ML technique, it has been the rise of 'deep learning' methods that have catalyzed the current AI revolution [13]. These methods are scalable and can train on petabytes of data through the use of graphical processing units (GPUs) [17] and continue to improve even when the performance of non-deep learning methods has saturated [18, 19]. In addition to scalability, deep learning's modular nature brings the added benefit of easily incorporating domain-specific knowledge (often called an *inductive bias*) to point the model in the direction of good or parsimonious solutions [20]. Although image recognition is not commonly a task in the current PV pipeline, deep learning models known as convolutional neural networks (CNNs) offer a particularly salient example of how potent the combination of large data and domain knowledge can be.

CNNs were introduced in 1988 [21] but it was not until 2010 when datasets [22] with millions of images became available that they began to transform the field of computer vision [23, 24]. Moreover, the convolution operator and the network structure (modeled loosely on the visual cortex [25]) in CNNs are powerful image-specific inductive biases that give the deep learning model a head start when learning a new image recognition task. Without either of these components (large data and the inductive bias of convolution), it is unlikely that deep learning would have caused the computer vision revolution of the 2010s. Indeed, numerous studies have found that without large data and inductive biases, deep learning is often no better than traditional statistical models [26–28]. These lessons have been born out repeatedly in subsequent applications of game playing [29, 30], biology [5, 6], natural language processing (NLP) [31, 32], and image generation [33–35].

Taken as a whole, it is thus reasonable to expect that a field is unlikely to experience a true paradigm shift from the current crop of deep learning-powered AI techniques without having at least some of these prerequisites in place. Despite the widespread interest in AI and its application to safety data [36, 37], including several review articles [14, 15], there are no scoping reviews that critically assess the extent to which PV is poised to be improved by AI under this framework. Previous reviews have focused on specific elements such as NLP techniques for clinical narrative mining in EHRs [38] or in reducing the frequency or impact of adverse events to patients [39]. Our review is unique in that it seeks to fill this gap to provide a clearer understanding of how current AI/ML practices and standards in PV align with the critical factors for success identified in adjacent areas such as biology and medicine.

To be as comprehensive as possible, we take a broad definition of ML for safety data and include traditional signal detection methods such as Bayesian Confidence Propagation Neural Networks (BCPNN) and related techniques [40, 41] given their roots in ML (see the Methods section for the full search details). We surveyed a 21-year period from the year 2000 to September 2021, reflective of the time before and after significant AI breakthroughs in 2012–2015 [23, 42-46], to see what effect, if any, these results had on PV. The scope of this review is limited only to (1) ingestion of safety data from all sources, including the safety data pipeline, social media, EHRs, and scientific literature, followed by (2) the processing and structuring of data, as well as (3) the processes of analyzing, understanding, linking, and disseminating or sharing, the safety data. While ML has made advances in healthcare more broadly [47–51] and ML research in these areas does have the potential to impact PV, we sought to characterize the use of ML that directly analyzed safety data (e.g. social media, forums, or ICSRs such as those in the FDA Adverse Event Reporting System [FAERS]) and excluded studies that performed adjacent kinds of tasks (e.g. ML research on biochemical pathways or meta research on drug safety). Thus, for the purposes of this review, we have chosen to retain our focus by limiting the review to those topics related specifically to the application of ML and human safety data (i.e. work that explicitly analyzes data on suspected adverse events of drugs and vaccines) for data ingestion or analysis.

#### 2 Methods

#### 2.1 Study Design

We queried four databases (PubMed, Embase, Web of Science Core Collection, and IEEE Xplore) for abstracts of fulltext research papers containing terms related to ML and PV. The searches were carried out on 9 September 2021 and were limited to articles published in the year 2000 or later. Furthermore, our review was limited to full-text English articles and conference abstracts (non-English papers were excluded). The full list of search terms and the search query used to identify the articles in this review are available as an electronic supplementary file.

We focused our search criteria on ML terms related to disproportionality analysis, common to PV research, as well as modern ML techniques (e.g. deep learning). This allowed us to compare traditional methods of PV alongside cuttingedge ML research. Articles solely focused on rules-based methods or knowledge graph- or ontology-based methods (e.g. Merrill et al. [52, 53]) were excluded from this review since, on their own, these are not direct ML methods per se.

Two independent reviewers determined if an abstract was in the scope of this review. A third reviewer adjudicated conflicts between reviewers or indecision by a reviewer (indicated by a 'maybe' vote). For studies that met the inclusion criteria, one reviewer conducted a full-text review to extract data. Analysis of extracted data was performed using the R statistical programming language [54]. Statistical significance for testing for a difference in proportions Chi-square test enabled subgroup analysis of studies that proposed methods for the intake and processing of safety data. This subgroup was enriched for ML models of interest. Topic modeling, described below, enabled an analysis of temporal trends in methods development.

#### 2.2 Evaluation Criteria

To understand the extent to which PV studies are amenable to current trends in the broader ML literature, we assessed each paper using the following criteria:

- *Task type:* We categorized each study into one of three categories reflective of the primary approach: signal detection, data intake, or data analysis.
  - Signal detection: Papers that are 'traditional' PV analyses that seek to estimate a statistical quantity (e.g. information component, odds ratio, etc.) for signal detection. This category could also include alternative ML methods for signal detection.
  - Data intake: Papers that use ML models to process safety data of various kinds for storage in databases or for downstream activities such as signal detection. Examples include adverse event detection, named entity recognition, and other preprocessing activities.
  - Data analysis: Papers that leverage safety data but do not fall into either of the previous categories. Examples include clustering of adverse events and topic modeling.

- Dataset and dataset size: We collected the name of the dataset and number of data points each study used to train and/or assess its methodology. In the case of multiple reported dataset sizes, we reported the 'most specific' number. For example, if a study reported using millions of safety reports from FAERS, but trained and assessed models on a subset of thousands of reports related to acute kidney injury, we went with the smaller number.
- Modeling approach: We identified the primary algorithm or model used in each study in addition to any secondary algorithms or techniques.
  - *Examples*: BCPNNs, reporting odds ratios (ROR), random forests, transformers, etc.
- Method novelty: Given the importance of domain adaptation seen in other fields such as computer vision and NLP, we subjectively assessed whether researchers in each study used an 'off the shelf' ML algorithm (e.g. random forests, support vector machines [SVMs]) or used a model tailored to the task, or otherwise made non-trivial modifications to an existing algorithm to improve performance (e.g. beyond hyperparameter tuning).
- Use of external information or pretrained models: One of the great benefits of current deep learning models is the ability to leverage external data and pretrained models when labeled data are scarce. We checked for additional information as inputs to the model (e.g. incorporation of known adverse effects or molecular structure). We also looked for the use of models that had been trained on other datasets then transported to the PV task at hand.
- *Reproducibility:* We searched for dataset and code availability to indicate whether or not a study was reproducible. We did consider social media data as publicly available, but acknowledge that it may be difficult to exactly reproduce a dataset based on a social media crawl. For code availability, we identified all papers that provided a link to Github or other web-hosted code, or provided supplementary materials with code. We manually assessed this subset of papers for currently available code (e.g. no dead links).
- Overall evaluation: We recorded a binary subjective evaluation indicating whether each study was reflective of the best practices in the broader ML literature (e.g. appropriate inductive biases, no obvious test-train leakage, tuning hyperparameters, cross validation). This determination was based on how well each study reflected the other criteria on this list.

#### 2.3 Topic Modeling

We trained a structural topic model (Latent Dirichlet Allocation [LDA]) using the 'stm' R package [55]. In order to process the documents included in our final screen, we preprocessed the text to remove all non-alpha-numeric text and removed references (e.g. '[1]'). We then removed punctuation and limited analysis to words between 2 and 20 characters. For each document, we only considered words that appeared at least 10 times in that respective document. After preprocessing by removing stop words and stemming words, we needed to select the number of topics, *K*, to instantiate the topic model with. We considered a range of 5–45 topics and chose K = 25 based on log-likelihood on held-out data, exclusivity, and semantic coherence. Finally, we fit the topic model using a semi-collapsed variational expectation-maximization algorithm regressed on the year of publication.

## **3 Results**

#### 3.1 Main Results

We manually reviewed 7744 unique abstracts that were identified by searching the PubMed, Embase, Web of Science, and IEEE Xplore databases. After manual screening by at least two reviewers, 672 (8.7%) abstracts passed inspection and had their full-text retrieved (Fig. 1). Of these, 279 (41.5%) did not meet the inclusion criteria due to lack of relevance after reviewing the full text, or did not have the full text available, resulting in 393 articles for analysis. Figure 2 displays summary information for the primary datasets and models in addition to the task classification for each study.

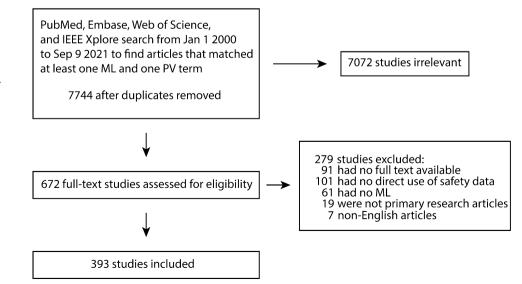
Overall, FAERS was the most popular single database (Fig. 2a) and was used by 24% of the studies. Social media data were used by 12% of studies, while EHR data were used by 11% of studies. Traditional statistical PV methods such as disproportionality scores remain very popular, with 144 (37%) of the included studies using one of them as the primary analysis model. Sample sizes varied greatly across studies and across datasets (Table 1 and Fig. 3), with FAERS being notable for having a mean sample size of 4.3 million and a median sample size of 243,510. Note that the notion of sample size is difficult to compare across data sources since the unit of analysis can be quite different. For example, studies using social media data often reported the number of posts (e.g. Tweets), while EHR studies often reported the number of patients. Ten percent of studies reported no explicit sample size at all.

With respect to method novelty, the vast majority (73%) used 'off-the-shelf' methods with little to no problem-specific adaptation or domain knowledge. Method novelty varied with respect to method type; 61% of papers using deep learning or other ML methods had novel adaptations, while only 10% of disproportionality papers did. Similarly, 92% trained a model 'from scratch', leaving only 8% of studies that leverage a pretrained model in some capacity, and only 18% explicitly used some kind of external information or data. Six-three percent of the studies used data that were publicly available, while 7% had code that was publicly accessible at some point in time. Our reviewers' subjective evaluation found that 42 (10%) studies were reflective of modern best practices in ML.

#### 3.2 Subgroup Analysis of Data Intake and Pipeline Studies

We performed a subgroup analysis of studies that proposed methods for the intake and processing of safety data. The use of transfer learning, methodological novelty, and the types of models used are shown in Fig. 4 and the sample size by dataset is shown in Table 2. Compared with all included studies of the previous section, this category had significantly higher levels of methodological innovation and novelty (40% vs.

Fig. 1 Summary of inclusion and exclusion process. Articles identified in one of the four databases using keyword and MeSH term searches were manually screened for inclusion. *MeSH* Medical Subject Heading, *ML* machine learning, *PV* pharmacovigilance



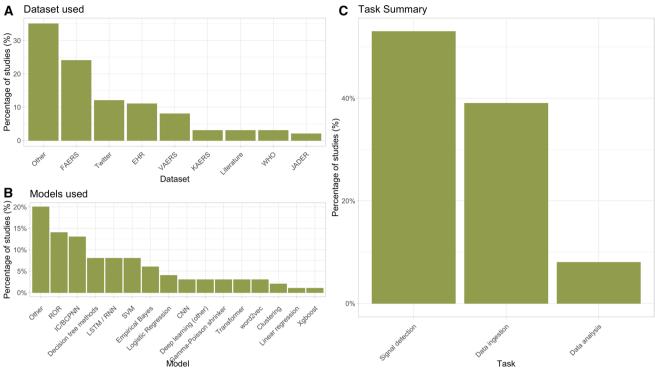


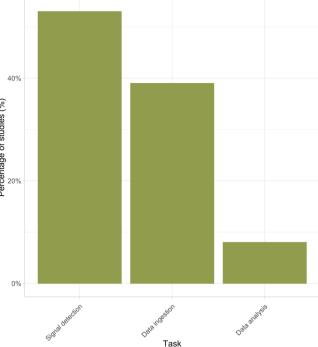
Fig. 2 Summary of datasets, primary algorithms, and task type of the included studies. a Primary dataset used in each study. b Primary analysis method or model used by each study. c Study task classification. EHR electronic health record, FAERS FDA Adverse Event Reporting System, JADER Japanese Adverse Drug Event Report, KAERS Korea Adverse Event Reporting System, VAERS Vaccine

Table 1 Summary statistics for types of data utilized and sample sizes used

Dataset	n	Mean	Median	SD	IQR
EHR	43	628,859	2633	2,004,196	63,516
FAERS	93	4,306,251	243,510	14,602,542	3,134,268
JADER	9	136,331	33,852	184,969	282,323
KAERS	12	254,293	4982	789,068	6993
Literature	11	17,200	3000	34,797	14,964
Other	138	3,067,200	26,508	14,025,930	259,820
Social media	47	46,685,103	14,570	304,970,307	100,183
VAERS	30	64,363	2619	234,839	13,714
Vigibase	10	365,710	11,144	941,166	14,944

SD standard deviation, IQR interquartile range, EHR electronic health record, FAERS FDA Adverse Event Reporting System, JADER Japanese Adverse Drug Event Report, KAERS Korea Adverse Event Reporting System, VAERS Vaccine Adverse Event Reporting System

27%; p = 0.03) and had higher uses of pretrained models (19% vs. 8%; p = 0.03). We found that 30 (20%) of these studies reflect current best practices, higher than the 10% estimate when considering all included papers.



Adverse Event Reporting System, WHO World Health Organization, ROR reporting odds ratio, IC information component, BCPNN Bayesian Confidence Propagation Neural Network, LSTM long shortterm memory, RNN recurrent neural network, SVM support vector machine, CNN convolutional neural network

#### 3.3 Temporal Trends

Next, we assessed how some of the patterns from the previous sections varied during the study period. Figure 5 shows trends for the number of publications, task type, and model use for each year in the study period. By and large, the volume of ML-related PV publications went up year-over-year (Fig. 5a). Starting in 2015, the number of studies leveraging ML for data intake (Fig. 5b) markedly increased, which coincided with a rapid increase in the number of studies using deep learning (Fig. 5c), and, by 2020, deep learning was the most popular technique used in the included PV studies. These trends suggest that, especially for data intake studies, ML may be starting to gain traction.

#### 3.4 Topic Model Analysis

We then performed a topic model analysis of the full text using Latent Dirichlet Allocation (LDA) [56] to see what high-level trends were present. In Table 3, we show four of the most prevalent topics discovered by LDA along with

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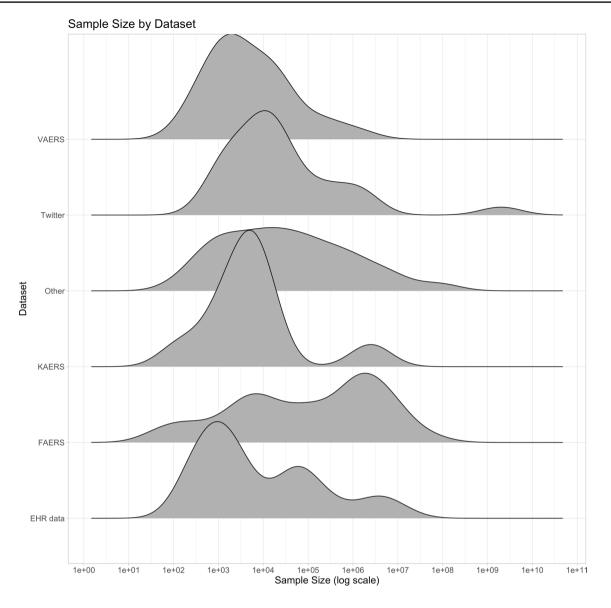
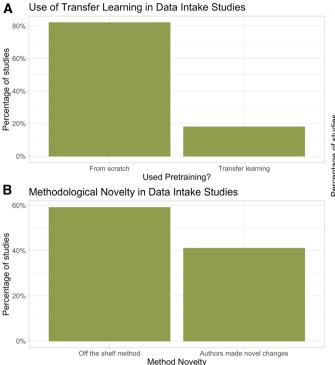


Fig. 3 Distribution of sample size for the most popular datasets. *EHR* electronic health record, *FAERS* FDA Adverse Event Reporting System, *KAERS* Korea Adverse Event Reporting System, *VAERS* Vaccine Adverse Event Reporting System

keywords that identify the topic and our subjective assessment of topic focus. We analyzed topic model results using the 'stm' and 'LDAvis' packages [55, 57]. Figures 6 and 7 show topics whose expected relative proportion increased and decreased, respectively, during the study period. These results align with those from the manual annotation presented in the previous section. Deep learning has seen relative gains in use recently and is likely to see further increases in coming years.

## 4 Discussion

Our scoping review revealed several interesting trends. First and most obvious, traditional signal detection methods in PV (e.g. BCPNN) and data sources remain quite popular and, until very recently, comprised the bulk of signal detection research. That is not to say that the use of these approaches has slowed, but that research development has shifted to other areas of method development. Interest at



Most popular models for data intake and processing С 20% 15% Percentage of studies 0% 5% 0% LSTMIRMM OR

Fig. 4 Breakdown of the use of a transfer learning, b methodological novelty, and c popular algorithms for data intake and pre-processing studies. LSTM long short-term memory, RNN recurrent neural net-

Table 2 Summary of sample sizes used in intake and processing pipeline studies

Dataset	п	Mean	Median	SD	IQR
EHR data	22	327,731	1237	1,082,972	6184
FAERS	4	1,158,504	70,357	2,223,398	1,146,282
Other	80	2,249,619	10,296	11,094,273	267,589
Social media	26	77,012,220	13,450	392,214,186	29,502
VAERS	3	5939	6034	3380	3379

SD standard deviation, IQR interquartile range, EHR electronic health record, FAERS FDA Adverse Event Reporting System, VAERS Vaccine Adverse Event Reporting System

the intersection of ML and PV is growing; Fig. 5a shows that the number of publications has increased approximately sixfold in the past 10 years alone. Figure 5c indicates that deep learning-based methods have recently eclipsed statistical methods in terms of publication numbers. This may have been enabled by new developments in deep learning for text analysis (e.g. transformers [32]), as initial deep learning progress focused on image recognition and was thus less relevant to PV tasks. Moreover, easy to use frameworks such as Tensorflow and PyTorch have enabled rapid development of ML models. In particular, there has been a rise in the amount of papers focusing on more sophisticated ML techniques. This indicates that the field is shifting towards classification

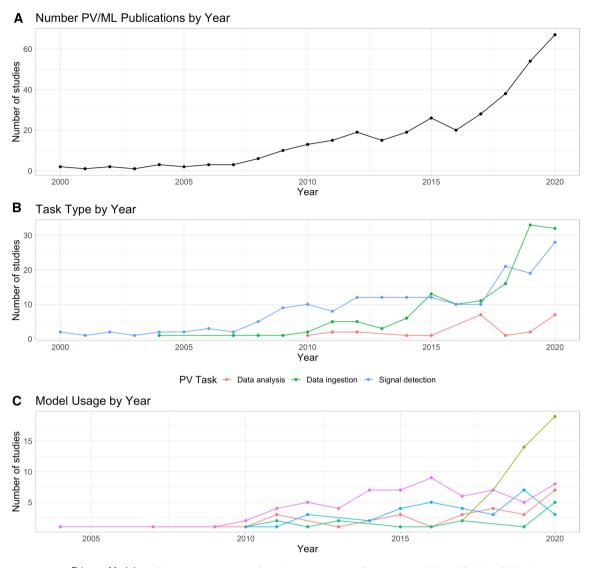
Mode work, SVM support vector machine, CNN convolutional neural net-

work, ROR reporting odds ratio

or regression tasks in addition to the more traditional safety signal statistical analyses. Figure 5b shows this fourfold increase in supervised tasks over the last 5 years alone.

In our subgroup analysis of studies that proposed methods for the intake and processing of safety data, we found articles within this category demonstrated higher levels of methodological innovation and novelty (40% vs. 27%; p = 0.03) and made more use of pretrained models (19% vs. 8%; p = 0.03). This could be the result of more freedom to define the task and model when compared with signal detection tasks and the ability to leverage existing pretrained models trained on other types of non-PV text data. Although these crucial ML ingredients were more frequently present in this task, this is lower than what would be expected for other areas where transfer learning is ubiquitous [11, 58].

One limitation of our review is a property of scientific publishing: only novel results are typically published in peerreviewed journals. For signal detection papers, that means our scoping review has covered both novel methods of signal detection and new drug/adverse event relationships identified by standard methods. In contrast, innovative data intake and analysis methods have been included in our review, but routine use of ML for these parts of PV are missing from the published record. This is partially reflected in our method novelty results; 61% of papers with deep learning or ML made novel changes, while only 10% of disproportionality



Primary Model 🔶 Decision tree methods 🔶 Deep Learning 🛥 Logistic Regression 🛥 SVM 🔶 Traditional PV Methods

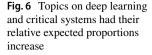
**Fig. 5** Temporal trends in the pharmacovigilance literature. **a** Total number of publications by year shows an increasing volume of articles that use ML for PV. **b** The type of task performed by each study.

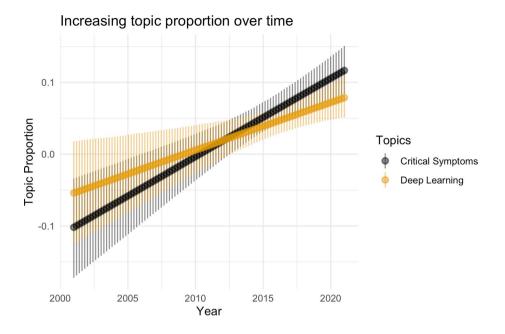
**c** Trends in usage for several classes of models. *ML* machine learning, *PV* pharmacovigilance, *SVM* support vector machine

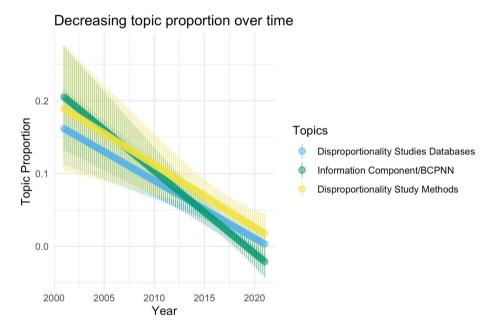
Table 3Top four topicsdiscovered by LDA byprevalence

Topic focus	Keywords	Prevalence (%)
Deep learning	Embed, LSTM, layer, model	7.01
Adverse events post-vaccination	Vaccine, VAERS, Guillain–Barré syn- drome, report	6.17
Signal detection	BCPNN, PRR, Gamma Poisson Shrinker, shrinkage, method	5.87
Information extraction/NLP	Annotate, entity, ADE, sentence	5.70

Topic focus is a subjective assessment of the content based on the most relevant keywords for each topic, as determined by setting the relevancy metric  $\lambda$  to 0.3 in the 'LDAvis' package (see electronic supplementary Table S1 for a full list of topics). *NLP* natural language processing, *VAERS* Vaccine Adverse Event Reporting System, *BCPNN* Bayesian Confidence Propagation Neural Network







**Fig. 7** Topics on disproportionality analysis, BCPNN had their relative expected proportions decrease. *BCPNN* Bayesian Confidence Propagation Neural Network

analyses made novel changes in our reviewers' estimations. While this bias is unavoidable, we believe that this scoping review likely captures most uses of ML for data ingestion and analysis due to the fact that the rapid rise of ML has been so recent. This bias will affect the ratio between signal detection and other papers, but the conclusions within subgroups should be unaffected.

However, returning to our original framing, it is not yet clear whether PV has assembled the critical mass of ingredients needed to benefit from the recent AI revolution powered primarily by large-scale deep learning methods, although it may be trending in that direction. Figure 3 shows that there is a large amount of human safety data that can serve as fodder for model training. However, most studies are bespoke, oneoff exercises that do not introduce methodological novelty beyond what has been published in other areas, and do not use pretrained models or external data despite an increasing use of deep learning methods. In contrast to other areas that have rapid growth and transformation, PV studies still mostly focus on a single task and use a narrow subset of available data, neglecting to use pretrained models or external data. Models that leverage multimodal data (e.g. text and structured data) have been particularly useful in other areas [34, 59, 60]. Another contrast with other areas is the availability of code. Only 7% of PV studies in our review had publicly available code at one point in time (approximately 1% of papers have dead links to code). This is in contrast to 2019 estimates of code availability, where 21% of studies in ML for healthcare more broadly, 39% of computer vision papers, and 48% of NLP papers provided code [61]. Indeed, sharing of code, data, and rapid dissemination of results through preprint servers have accelerated progress in other areas of ML [62–64]. Code-sharing enables others to build on previous work rapidly, which is extremely important when model complexity is high, as can be the case with many complicated deep learning models.

We wish to emphasize that we are not suggesting that PV researchers must follow the deep learning template, nor do we believe that deep learning is the only viable method for PV tasks. However, if PV tasks are to be improved by *current* approaches based on deep learning, then the criteria of large datasets, the use of pretrained models when appropriate, method novelty, and reproducibility are a reasonable set of requirements. By these criteria, we found that 10% of all studies and 20% of pipeline/intake studies were reflective of current trends based on deep learning. With the increasing use of large datasets and the rise of more modern ML techniques observed in Fig. 5, it is likely safe to project that this percentage will increase over the next several years.

#### 4.1 Recommendations

We provide some concrete recommendations that we believe could enhance AI and deep learning applications in PV.

- Incorporating domain knowledge: Incorporating domainspecific knowledge biases in ML PV models (e.g. onedimensional CNNs to detect symptoms next to medications in Tweets or graph neural networks to leverage molecular structure). Known relationships or ontologies that relate symptoms, diseases, and drugs could also be directly incorporated into the model to improve performance [65].
- *External information and pretrained models:* Incorporating external information about mechanisms of action, common adverse effects, or geographic location of reports. This could be used to help triage reports; if multiple reports with the same constellation of symptoms appear for a particular medication that one would not expect (e.g. as encoded by a prior distribution), this would be a clear sign for further investigation. There are numerous pretrained models available for text [66] and molecular data [67] that serve as good foundations for extracting information for PV text and could provide strong prior information when detecting adverse events in case reports.
- *Methodological innovation available in other areas of ML*: Incorporating new advances in ML literature, such as uncertainty quantification [68, 69], federated learning,

and fairness ideas. Causal inference [70], an emerging field of epidemiology and computer science, is another promising avenue improving ML PV by incorporating known information about causal relationships directly into the model.

- Data sharing and reproducibility: Common data formats, benchmarks, and code sharing to foster reproducibility. There are several established benchmark datasets that have been used in the literature (n2c2 2018, MADE 1.0, etc.), but there is no equivalent of MNIST or CIFAR-10 for PV that can objectively measure progress on a difficult but standardized task.
  - Many efforts (e.g. Lindquist et al. [71], Hochberg et al. [72], Harpaz et al. [73]) towards benchmark tasks have resulted in rich labels for true positive and true negative drug/ADR relationships. However, these datasets do not come with accompanying preprocessed data (e.g. safety reports or social media posts) that would provide an easy-to-use benchmark.
  - Few investigations release their data after they scrape a public social media platform or forum. Although in theory one could recreate the authors' work, it is nearly impossible to capture the exact same posts and process them in the same way.
  - Studies have, in general, not published their code or calculations. We appreciate this is more understandable and less problematic for disproportionality analysis with reporting odds ratio, proportional reporting ratio (PRR), or BCPNN, but even for such studies, there is value in code provision to enable reproducibility. For projects that include more complex/modern ML models, public code repositories are lacking. In contrast to PV, this has been an influential factor that has spurred rapid development in the ML community.

#### 4.2 Promising Near-Term Applications of Machine Learning in Pharmacovigilance

While our review found that there is still much room for improvement, we wish to offer some near-term tasks that could benefit from the well-executed use of ML today. We offer suggestions for areas across the PV pipeline [74], where ML may have impact in the near- to medium-term, but note that some of these tasks will likely still require substantial effort to achieve.

• *Translation and multi-language models:* Case reports and other safety data can be submitted from anywhere in the world and written in hundreds of different languages. Processing these often necessitates translation into a common language of record before further evalu-

ations can take place. In recent years, ML has become exceedingly good at translation [75, 76], even for lowresource languages that do not have large amounts of training text available [77, 78]. There are even individual models that have been trained on vast amounts of language data and are capable of processing hundreds of languages [79]. Moreover, many of these models are publicly available and could be easily repurposed for the PV intake and processing pipeline. Integrating the translation model directly into the PV pipeline in a trainable way will allow it to adapt the capabilities to a variety of tasks when compared with treating translation as an auxiliary and separate preprocessing task.

- Named Entity Recognition (NER): Automatic extraction
  of key phrases and nouns is a common task in PV data
  intake and is known in the NLP literature as named entity
  recognition. There has been rapid progress on this task
  in other areas of ML [31, 80–82], including scenarios
  with multiple languages [83, 84], biomedical applications [85–87], and when labeled data are scarce (see the
  example in Yao et al. [88] detailed in the next section).
- *Text summarization and generation:* Case reports can often contain large volumes of unstructured text that individual case examiners must sift through and synthesize. Abstractive summarization by deep learning has also experienced an impressive leap in capabilities in recent years [89, 90] and thus could easily be applied to the analogous task in PV. Likewise, reports must be generated using codified and structured data, and the generative capabilities of deep learning models could be used for this task.
- *Causal inference:* The critical question of PV is whether a drug is actually causing the adverse events that have been reported in safety reports. Causal inference [91] is a statistical field that provides estimates of treatment effects in real-world data. There has recently been heavy interest in the intersection of causal inference and ML [92]. There is a nearly one-to-one translation of the ideas of causal inference to PV and this could serve as another tool for signal detection and data analysis.

#### 4.3 Exemplar Studies

In this section, we wish to highlight several studies that were determined to reflect current ideas and trends in ML to provide good exemplars for how future studies might be conducted. Du et al. [88] provide an example of how an accurate adverse event annotation pipeline can be built, even when there are not large amounts of annotated data, using transfer and self-supervised learning. In the investigation, the authors only had a small set of labeled data in the form of 91 annotated VAERS reports, and the goal was to construct an ML system to automatically extract mentions of *named entities* (e.g. adverse events, procedures, social circumstances, etc.) from the reports. They accomplished this goal by leveraging a pretrained transformer model known as BioBERT [66] that was fine-tuned on an unannotated set of 43,240 VAERS reports. They show that this approach leads to significantly better performance for this task when compared with traditional NLP methods and when compared against deep learning methods that did not employ transfer learning. Additionally, the annotated dataset they created is publicly available so that others may build on their work [93].

Zhang et al. [94] showed how ML can be useful in complex adverse drug reaction recognition tasks. Adverse drug reactions can be found in all types of media, including scientific literature, EHR data, and Tweets. In these settings, the drug and reaction are not necessarily in the same sentence or near each other in text. Typical ML methods rely on local semantic information (e.g. words in a single sentence) and can struggle in identifying these adverse drug reactions. Zhang et al. leverage a novel mechanism known as multi-hop attention to endow models with the ability to focus across multiple words in a single sentence and between sentences. They used the publicly available benchmarks TwiMed and ADE to assess model performance and to compare with baselines. They demonstrate that their method outperforms well-established ML models such as SVM, CNNs, and LSTMs. Additionally, they show multi-hop attention is superior at identifying adverse drug reactions compared with self-attention and multi-head self-attention, two recent mechanisms found in transformer models. They also performed a comprehensive ablation study to isolate which of their innovations resulted in improved performance.

Finally, Wang et al. [95] demonstrate how ML can assist in determining causation from case reports. The authors utilize causal inference, which is a conceptual framework that, under certain assumptions, allows for estimation of causal effects. This means that one can answer counterfactual or 'what if' questions such as 'what if a patient took medication A rather than medication B?'. They combine causal inference with transformer models that are trained on FAERS safety reports. Wang et al. assess their proposed transformercausal inference model on two tasks: identifying causes of analgesic-induced acute liver failure and identifying causes of tramadol-related mortalities. Their model is able to recapitulate known risk factors for these adverse events (e.g. acetaminophen consumption for liver failure, and suicidality for tramadol mortalities). Moreover, the model was able to identify potential secondary risk factors that predispose individuals to liver failure. Importantly, Wang et al. published their code and preprocessed data (i.e. FAERS reports). This will enable future researchers to reproduce and extend their work.

## 5 Conclusions

We have conducted a scoping review of the use of ML for PV applications. Our aim was to assess the extent to which PV has been or is ready to be improved by current deep learning-based AI techniques. We found that while certain modern practices have begun to appear, many of the primary reasons for the recent success of AI have yet to be translated. We conclude that without certain structural changes, PV is unlikely to experience similar kinds of advancements from current approaches to AI.

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#### Declarations

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**Conflicts of Interest** Jeffery L. Painter and Andrew Bate were current full-time employees and stockholders of GSK at the time this study was completed. Benjamin Kompa, Joe B. Hakim, Anil Palepu, Kathryn Grace Kompa, Michael Smith, Paul A. Bain, Stephen Woloszynek, and Andrew L. Beam have no conflicts of interest to declare.

Ethics Approval Not applicable.

Consent to Participate Not applicable.

Consent for Publication Not applicable.

Availability of Data The datasets generated and/or analyzed during this scoping review are available on Github (https://github.com/beamlabhsph/drug\_safety\_scopingreview). All references included in the final analysis are available in the 'references' folder in the repo. Additionally, a companion website to our review is available at https://beamlab. shinyapps.io/MLforPVReview.

**Code Availability** The code to reproduce our figures and LDA analysis is available at https://github.com/beamlab-hsph/drug\_safety\_scopingreview

Authors' Contributions Conceptualization: AB, ALB. Methodology: BK, JBH, PAB, SW, JLP, AB, ALB. Query formation: BK, JBH, PAB, JLP, AB, ALB. Abstract review: BK, JBH, AP, KGK, MS, SW, ALB. Abstract review conflict adjudication: BK, ALB. Full-text review: BK, JBH, AP, KGK, MS, SW, ALB. Full-text conflict adjudication: BK, ALB. Data extraction: BK, JBH, AP, KGK, MS, ALB. Formal analysis and investigation: BK, SW, ALB. Writing – original draft preparation: BK, ALB. Writing – review and editing: BK, JLP, AB, ALB. Funding acquisition: AB, ALB. Resources: AB, ALB. Supervision: AB, ALB. All authors read and approved the final version.

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