Influential Factors of Patients' Serious Adverse Effects

Key words: Patient Safety, Adverse Effects, Logistic Regression

Team: Yuhang Qiu, Zehao Xie, Bowen Zhang, Yifan Zhu



Outline



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Motivation

Patient safety is the fundamental component in healthcare services. According to World Health Organization, the occurrence of adverse events due to unsafe care is likely one of the 10 main causes of death and disability in the world. Hartwig's scale states that 46.03% of adverse reaction (ADRs) were classified as unpreventable, while 48.81% as probably preventable ADRs, and 5.16% as definitely preventable ADRs. The objective of this project is to analyze the association between factors, aiming to develop an effective model to decrease possible preventable ADRs rate in practice.

Research Questions

- Could we identify the correlation between different features?
- Could we find determinants for patient's advert effect's seriousness level?
- Could we build the effective model to examine these effects?

Data

- There are 9990 observations with 17 variables in raw data
- After data cleaning and processing, the dataframe involved in this project contains
 7425 observations with 8 created variables.

df.	df.head()											
	patientonsetage	patientonsetageunit	patientagegroup	patientsex	reaction	drug	summary	patientweight	reaction_count	drug_count	seriousness	seriousnesso
0	42.0	801.0	5.0	1.0	Headache	[{'drugcharacterization': '1', 'medicinalprodu	NaN	NaN	2	5	0	
1	53.0	801.0	NaN	1.0	Headache	[{'drugcharacterization': '1', 'medicinalprodu	NaN	NaN	8	3	1	
2	69.0	801.0	6.0	2.0	Headache	[{'drugcharacterization': '1', 'medicinalprodu	NaN	NaN	1	2	0	
3	53.0	801.0	NaN	2.0	Headache	[{'drugcharacterization': '1', 'medicinalprodu	NaN	NaN	6	2	1	
4	44.0	801.0	5.0	2.0	Headache	[{'drugcharacterization': '1', 'medicinalprodu	{'narrativeincludeclinical': 'CASE EVENT DATE:	NaN	7	1	0	

Methodology









Data Cleaning

- Cleaned the dataset to ensure data integrity
- Convert json to csv
- Explore nested data in the dataset
- Calculate the drug and reaction count for each patient
- Remove outliers & null values

Data Visualization

Built visualizations to explore the relationship between seriousness and other variables (age, gender, reaction count, etc.)

Model Building

Built a Logistic Regression model to predict patient seriousness

Assumption testing

Calculated p-value of each
variable to backtest the
assumptions based on
visualization results

 $logit(P_i) = ln(P_i/(1-P_i)) = \beta_0 + \beta_1 Patient Age_i + \beta_2 Pateint Male + \beta_3 Patient Female + \beta_4 Reaction Count + \beta_5 Drug Count$

Motivatio

Questions

Data

Findings

Conclusion

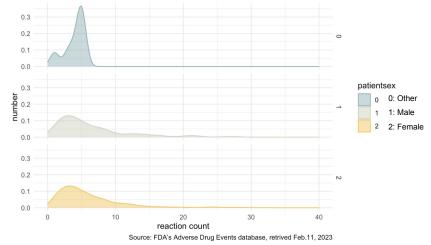
Discussion

Findings

Although the correlation between reaction count is weak related to seriousness level (left table), there is significant correlation with seriousness level = 1 when reaction count exceed

certain threshold (right image).

	Reaction Count	Drug Count	Seriousness
Reaction Count	1.0000	0.3509	-0.1113
Drug Count	0.3509	1.0000	-0.0631
Seriousness	-0.1113	-0.0631	1.0000



Motivatio

Ouestions

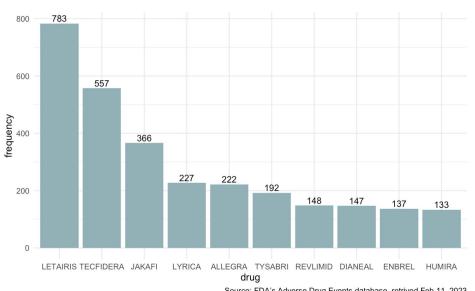
Data

Findings

Conclusion

Discussion

The Most Suspected Drugs that Caused Advert Effects by Frequence



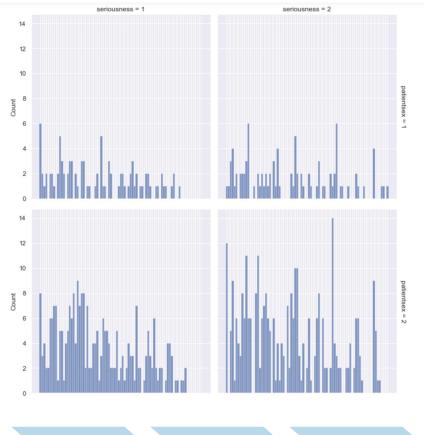
Source: FDA's Adverse Drug Events database, retrived Feb.11, 2023

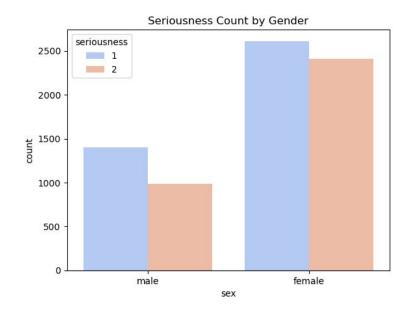
Some drugs are labeled as "Suspected" about causing the adverse effects in the dataset, we notice that some of them are significantly common than other drugs (For example, Letairis occurs ~8% in the dataset which covers more than 1000 different drugs).

Though we cannot explicitly deduce causal inference between certain drugs and adverse effects, these top drugs should be pay special attention [Might be included in future studies].

Findings

Seriousness Count by Age and Gender





Current function value: 0.677213

Iterations 5

Logit Regression Results

Dep. Variable:	seriou	isness I	No. Obse	rvation	s:	7401	
Model:	Logit		Df R	esidual	s:	7395	
Method:		MLE	D	f Mode	ıl:	5	
Date:	Sun, 12 Feb 2023		Pseud	o R-squ	0.0	0.01816	
Time:	05	:40:18	Log-Likelihood:		d: -50	-5012.1	
converged:		True		LL-Nul	l: -51	104.7	
Covariance Type:	nonr	obust	LLR p-value: 3.832e-38			e-38	
	coef	std err	z	P> z	[0.025	0.975]	
Intercep		std err 0.323	z -1.776	P> z 0.076	[0.025 -1.206	0.975] 0.059	
Intercep C(patientsex)[T.1.0	ot -0.5733				-	-	
	ot -0.5733 0] 0.2124	0.323	-1.776	0.076	-1.206	0.059	
C(patientsex)[T.1.0	0.2124 0.4715	0.323	-1.776 0.665	0.076 0.506	-1.206 -0.414	0.059	
C(patientsex)[T.1.0 C(patientsex)[T.2.0	0.2124 0.2715 0.0055	0.323 0.320 0.318	-1.776 0.665 1.482	0.076 0.506 0.138	-1.206 -0.414 -0.152	0.059 0.839 1.095	

Results of Logistic Regression

We mainly discuss effects of patient age, number of reactions and number of drugs on the seriousness level. All three of them have significant influences according to p-value, while an older patient would tend to have a higher seriousness level, the case is on the contrary for the other two factors.

Conclusion

- By investigating patient conditions and behaviors, we find some drugs appeared very frequently in patient's adverse reaction record, which should be take special attention in the future.
- Further, we successfully build a logistic regression model that indicates significant variables (total number of drug intake, total number of reaction); thus, could intentionally decrease the probability of preventable adverse reaction
- We recommend that patients pay extra attention to their health and other important features to reduce the preventable adverse reaction (ADRs) to the max extent, and we should have risk disclosure in advance

Discussion and Future Work

- We tried various approaches to tackle this challenging dataset and successfully defined our problem statements
- Though we draw some solid conclusions, there are lots of things we can do
- There are plenty of data in the original dataset we can further utilize
- The connection between various adverse effects and drugs can be explored if we have time to construct a proper mapping/connection relationship between these two factors
- We can try more powerful models in the future



Reference

[1] https://www.who.int/news-room/fact-sheets/detail/patient-safety

Data Source: https://open.fda.gov/apis/drug/event/how-to-use-the-endpoint/

Thanks for Listening

