

Exploratory Data Analysis

Healthcare: Persistency of a Drug

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Submitted to: Data Glacier

Link to the Repo: https://github.com/shonjeeyeon/DG Week 11



Agenda

Executive Summary

Problem Description

Approach

Exploratory Data Analysis: Overview

Exploratory Data Analysis and

Hypothesis Testing

Conclusion

Recommendations

Executive Summary



- Using a dataset of 3,424 records and 69 features, Exploratory Data Analysis (EDA) was performed to analyze persistency of a certain medication.
- Visual comparisons as well hypothesis tests were used to determine differences in persistency levels among different sub-populations, grouped by features such as specializations of prescribers and comorbidities.
- Summary of the EDA and recommendations for further model development will be included in the presentation.



Problem Description

- Medication persistence refers to completing the medication treatment using the duration set by the prescriber. (Cramer JA et al., 2008)
- Therefore, persistence is important in patients' positive outcomes as well as in pharmaceutical industries' profits.
- Developing a model to automate prediction process will contribute to save time and cost spent by the company, and the prediction results can be used for marketing, patient education, or R&D purposes.



Approach

Dataset provided by the company (3,424 patient records)



EDA

Hypothesis Testing

Conclusion and Recommendations for Further Model Development

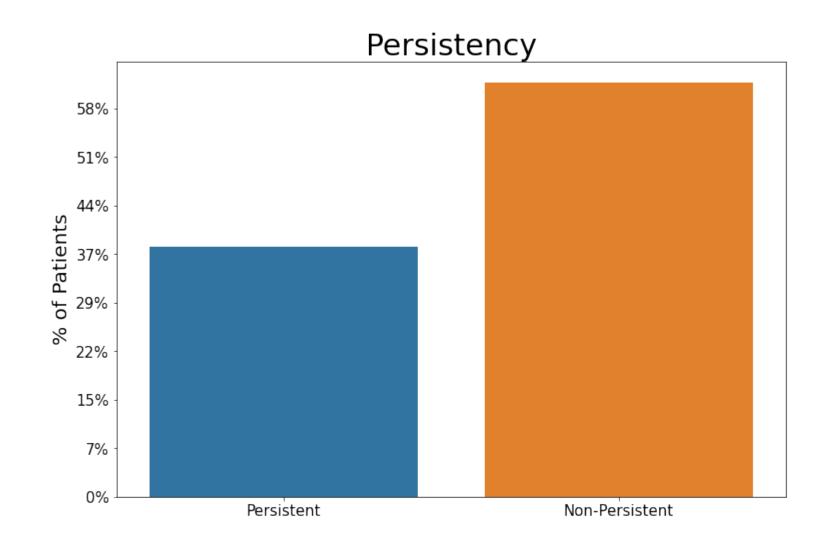


Exploratory Data Analysis:

Dataset Overview



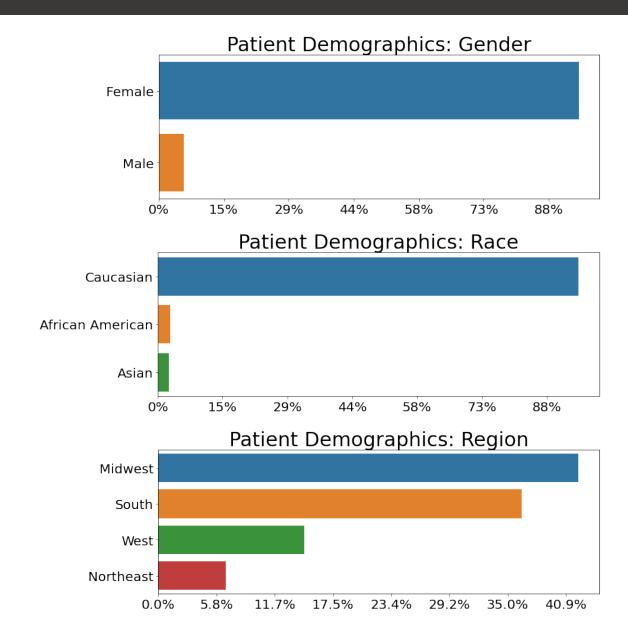
Dataset Overview: Persistency



 Among 3,424 patients in the dataset, approximately 40% of the population continued the medication persistently.



Dataset Overview: Patient Demographics

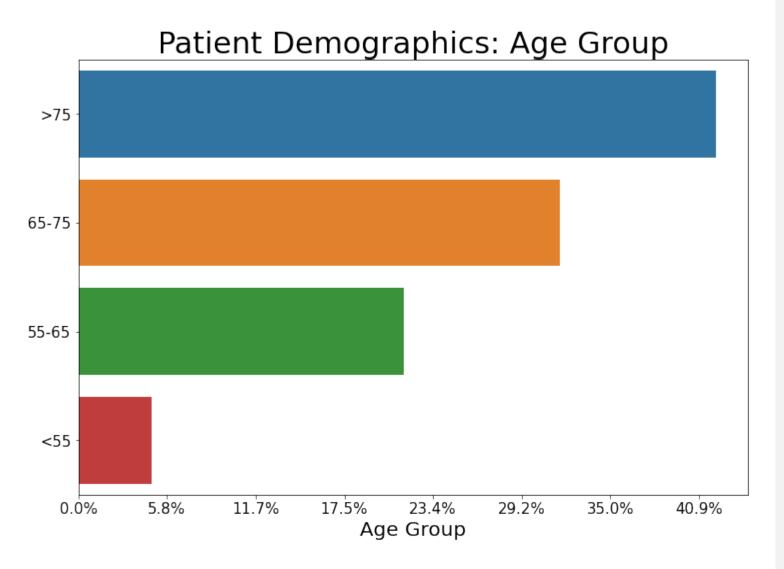


 Most of the patients were Caucasian or Female.

 Midwest was the most popular region, followed by South and West.



Dataset Overview: Patient Demographics

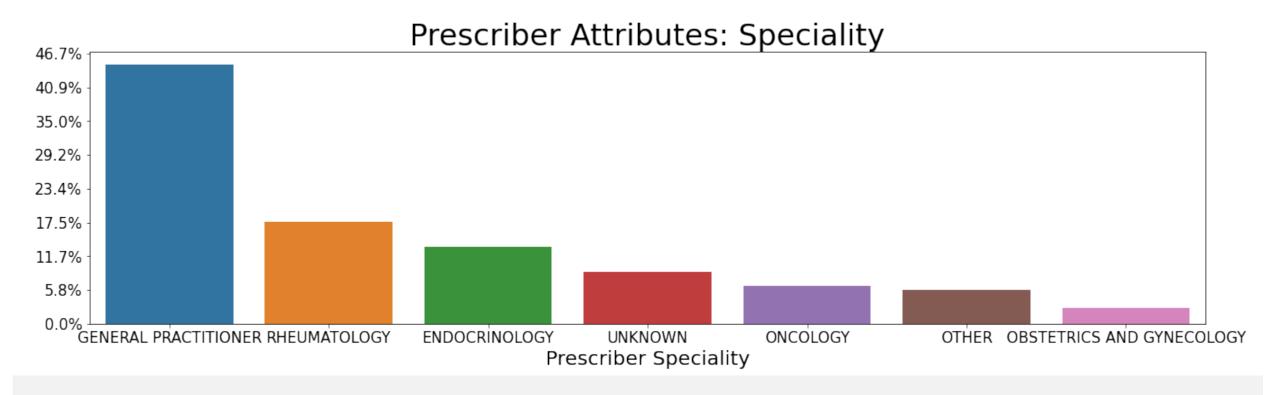


 Most of the patients in the dataset were over 75 years old, followed by patients aged 65 to 75 years old.

 Only about 5% of the population were under 55 years old.



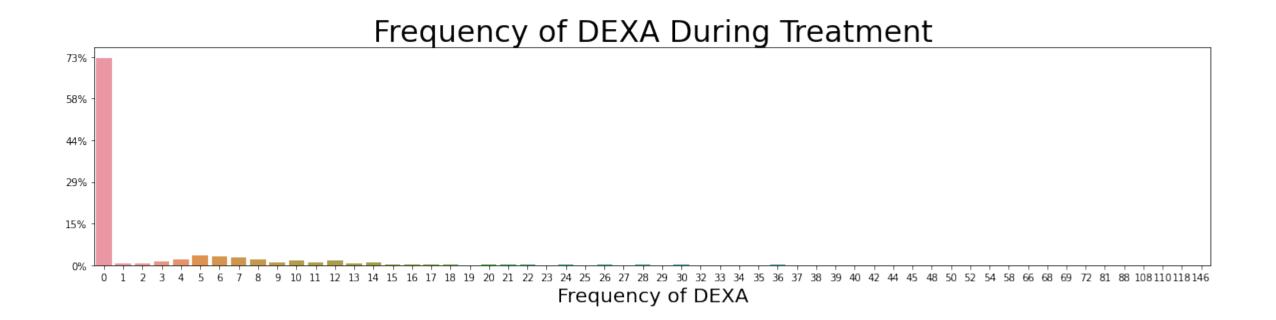
Dataset Overview: Prescriber Attributes



- The majority of the prescribers were General Practitioners.
- The second and third most common specialties were Rheumatology and Endocrinology.

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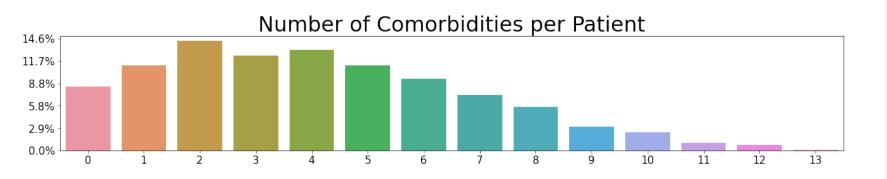
Dataset Overview: Frequency of DEXA

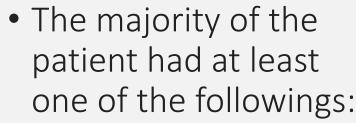


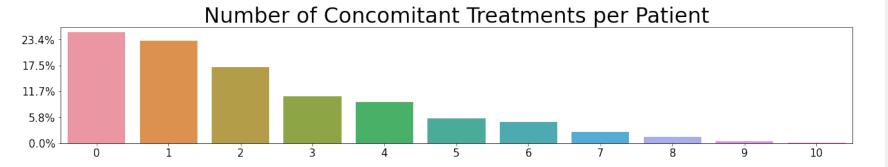
- About 75% of the patient did not take DEXA scan during treatment.
- The frequency DEXA scan during treatment varied substantially among patients, with patients taking the scan more than 140 times.



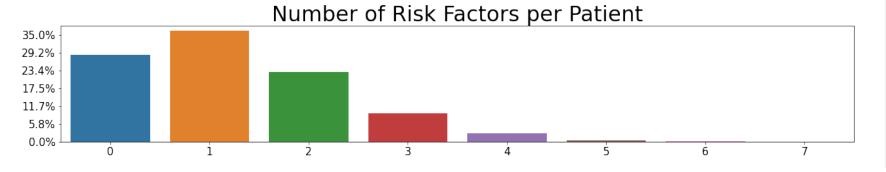
Dataset Overview: Comorbidities, Concomitant Treatments, and Risk Factors







- Comorbidities
- Concomitant treatments
- Risk factors



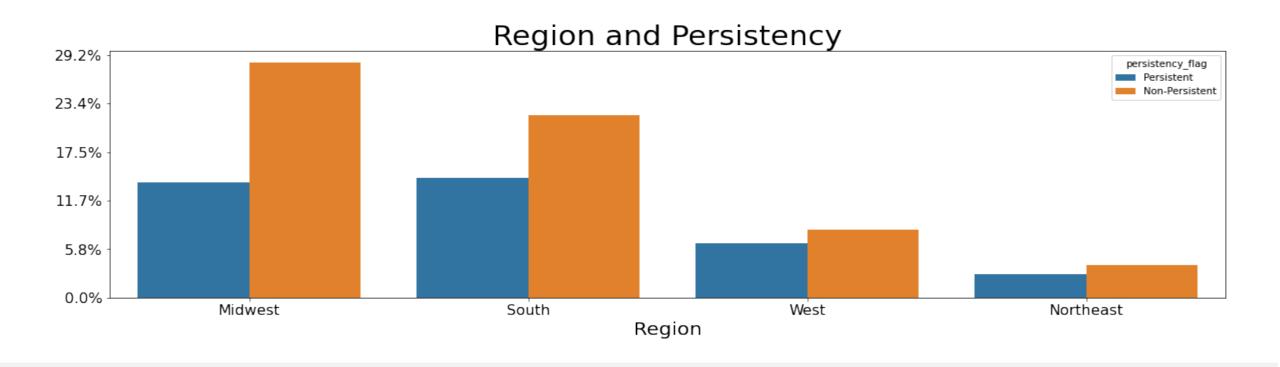


Exploratory Data Analysis:

Persistency Comparison and Hypothesis Testing



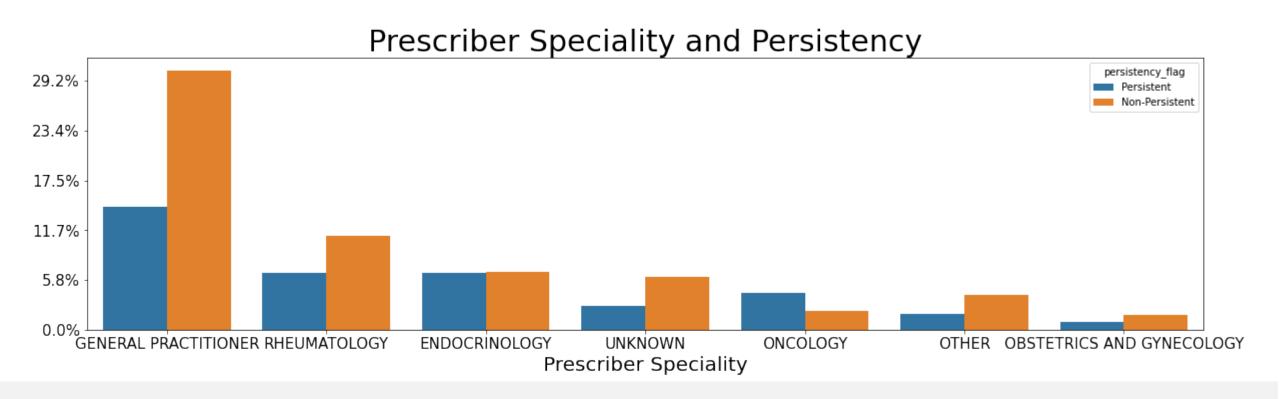
Region and Persistency



- Midwest has highest proportion of patients, but had the lowest persistency to nonpersistency ratio.
- There are significant differences in persistency level among regions (ANOVA, p< 0.05)



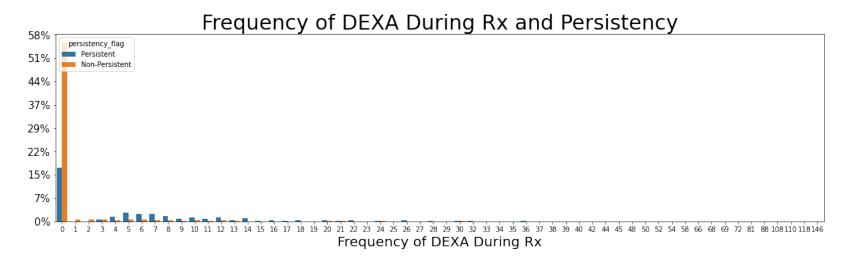
Prescriber Specialization and Persistency

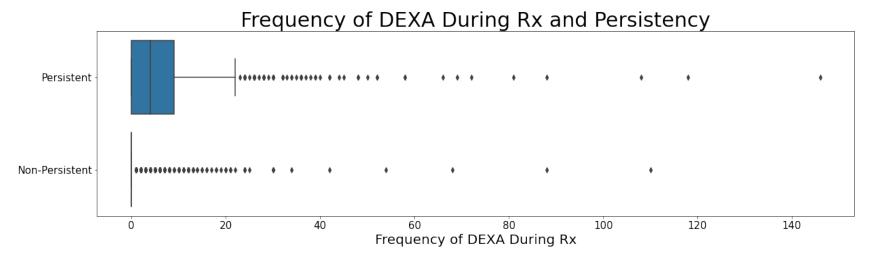


- Endocrinology and Oncology has higher persistent-to-non-persistent ratio
- There are significant differences in persistency level among specializations (ANOVA, p< 0.05)



DEXA Frequency and Persistency



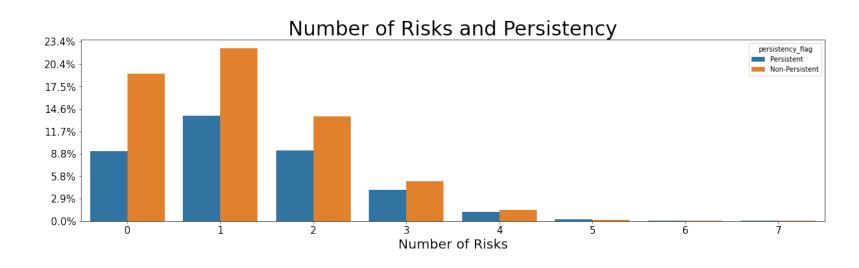


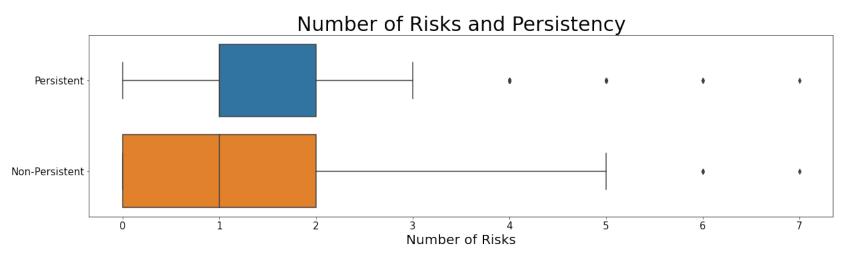
 Persistent group had higher mean number of DEXA scan during therapy

 There are significant differences in persistency level among different DEXA frequency groups
 (ANOVA, p< 0.05)



Number of Risks and Persistency



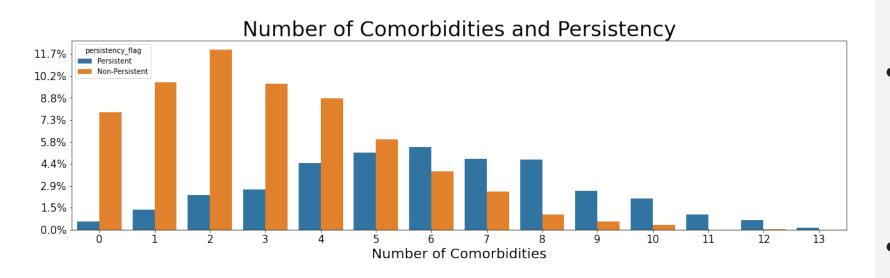


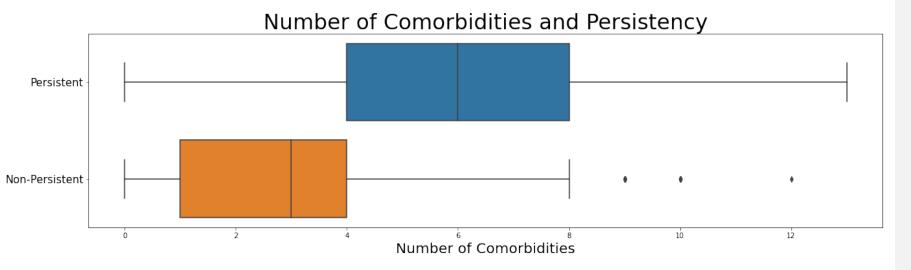
- In general, populations with more risk factors has higher persistent-to-nonpersistent ratio
- There are significant differences in persistency level across different risk counts

(ANOVA, p < 0.05)



Number of Comorbidities and Persistency



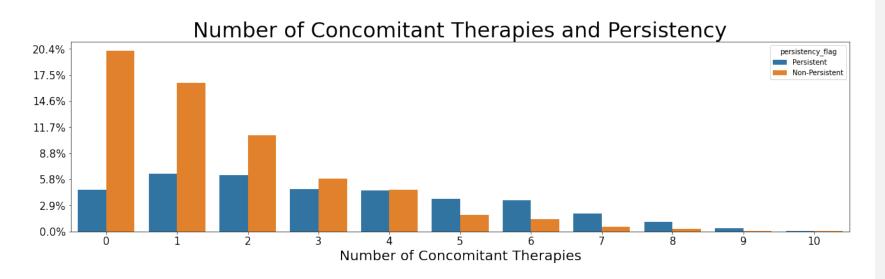


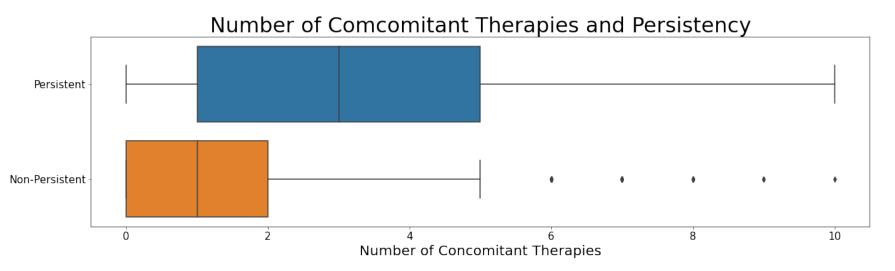
- Persistent group has higher mean in number of comorbidities.
- There are significant differences in persistency level across different numbers of comorbidities

(ANOVA, p < 0.05)



Number of Concomitant Therapies and Persistency





- Persistent group has higher mean in number of concomitant therapies.
- There are significant differences in persistency level across different numbers of concomitant therapies

(ANOVA, p < 0.05)



Other Features and Hypothesis Testing

- There were significant differences in persistency between:
 - Patients who had used glucocorticoid during therapy and who had not
 - Patients who had experienced fractures during therapy and who had not
 - Patients who had used injectables during therapy and who had not (Chi-square tests, p values <0.05)
- Every comorbidity and concomitant therapy was associated with significant difference in persistency (Chi-square tests, p values <0.05)
- Among risk factors, rheumatoid arthritis, chronic hypogonadism, tobacco use, chronic malnutrition, vitamin D insufficiency, and immobilization were associated with significant difference in persistency (Chi-square tests, p values <0.05)



Summary of the EDA

- Region played a role in differences in persistency.
- Prescribers with certain specializations were associated with higher persistency.
- Taking DEXA scan during therapy were associated with higher persistency.
- Having at least one comorbidities, concomitant therapies, and/or risk factors were associated with higher persistency.
- All of the comorbidities and concomitant therapies, as well as select risk factors, were associated with difference in persistency.



Recommendations

- Classification should be used for model and application development.
- Simpler models such as decision tree or linear regression should be tried first, before moving on to more complex and time-consuming models.
- The dataset has a substantial number of features. Using PCA or RFE to select important features is highly recommended.



Reference

• Cramer JA, Roy A, Burrell A, Fairchild CJ, Fuldeore MJ, Ollendorf DA, Wong PK. Medication compliance and persistence: terminology and definitions. Value Health. 2008 Jan-Feb;11(1):44-7. doi: 10.1111/j.1524-4733.2007.00213.x. PMID: 18237359.

Thank You

