

# Adherence to Disease-Modifying Anti-Rheumatic Drugs In Patients in an Integrated Clinic and Specialty Pharmacy

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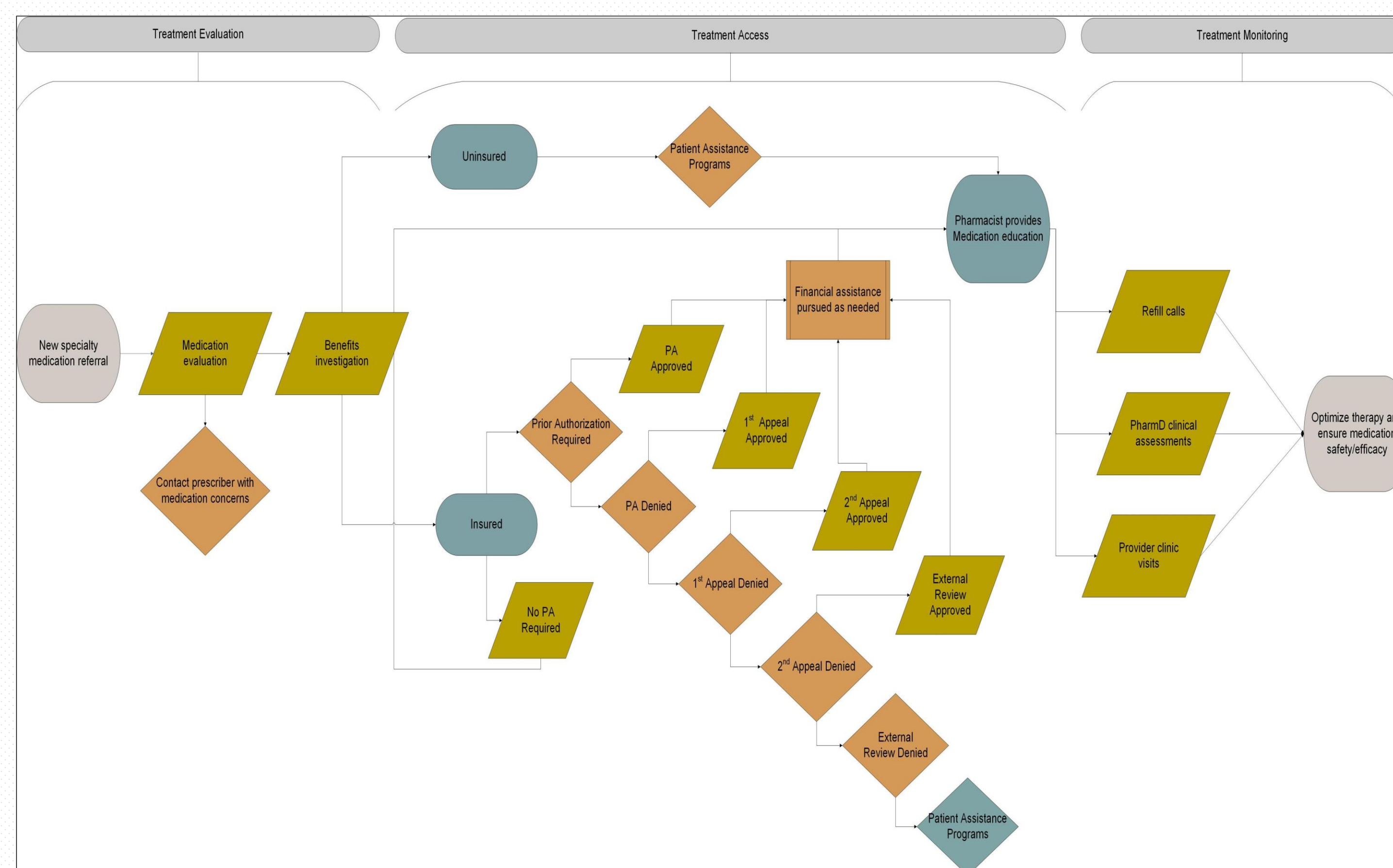
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## BACKGROUND

- Adherence to disease-modifying anti-rheumatic drugs (DMARDs) is associated with reduced rheumatoid arthritis (RA) disease activity and better radiological outcomes<sup>1</sup>
- Adherence to DMARDs is low, ranging from 30 to 85%<sup>2</sup>
- Older age, male gender, treatment non-naivete, and lower out-of-pocket costs have been associated with higher adherence<sup>2</sup>
- Health-systems integrated specialty pharmacy models have been shown to improve medication adherence in other specialty diseases<sup>3</sup>

**Objective: Evaluate rates and predictors of adherence to DMARDs within a health-system integrated specialty pharmacy**

Figure 1: Vanderbilt Integrated Specialty Pharmacy Model



## METHODS

- DESIGN:** Single center, retrospective cohort
- SAMPLE:** Adults with RA treated at Vanderbilt Rheumatoid Arthritis Clinic who filled 3+ DMARD prescriptions from Vanderbilt Specialty Pharmacy between July 2016 and June 2017
- OUTCOME:** Medication adherence, measured as proportion of days covered (PDC)
- ANALYSIS:** Descriptive statistics to summarize data and proportional odds logistic regression to assess associations between patient characteristics and PDC

## RESULTS

- Median PDC was 0.94 (IQR=0.82-0.99, mean  $\pm$  SD = 0.87  $\pm$  0.17)
- In multiple regression analyses, PDC was significantly higher in male and treatment naïve patients (table 2)

## STUDY SAMPLE

Table 1: Sample Characteristics (n=231)

Characteristic	% (n)
Gender (% female)	78% (179)
Race (% White)	90% (207)
Treatment naïvete* (% naïve)	24% (55)
Financial assistance use	71% (163)
Insurance category	
Commercial	64% (148)
Government	36% (83)
Prescribed DMARD**	
Etanercept	41% (100)
Adalimumab	38% (93)
Abatacept	11% (26)
Tofacitinib	6% (15)
Certolizumab	4% (9)
Tocilizumab	<1% (1)

\*Naivety: no pharmacy claim for a DMARD within the previous 6 months  
\*\*DMARD=disease modifying anti-rheumatic drug

## RESULTS

### ADHERENCE RATES AND PREDICTORS

Figure 2: Adherence Distribution

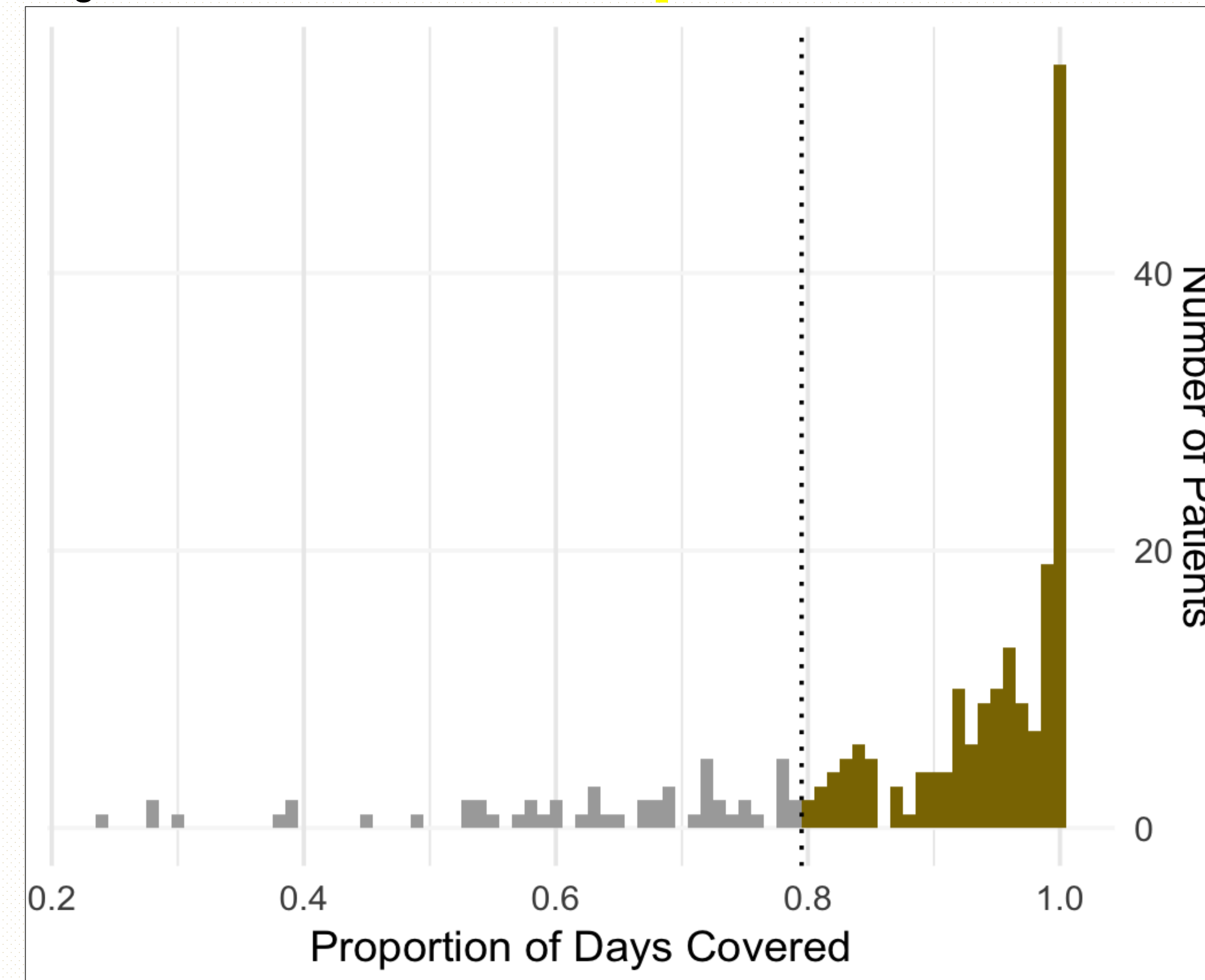


Figure 3: Median and Interquartile Range of PDC by Patient Characteristics

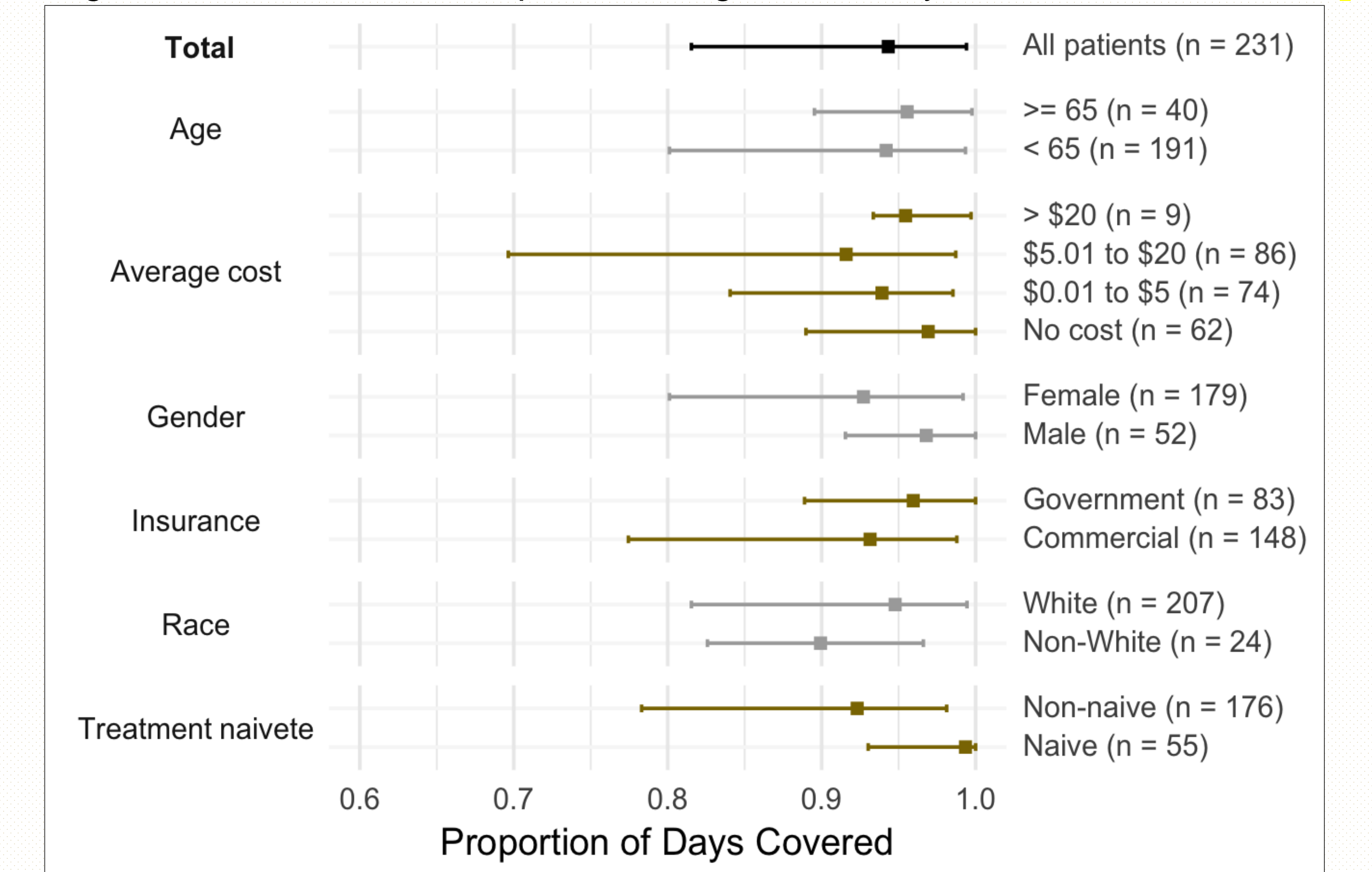
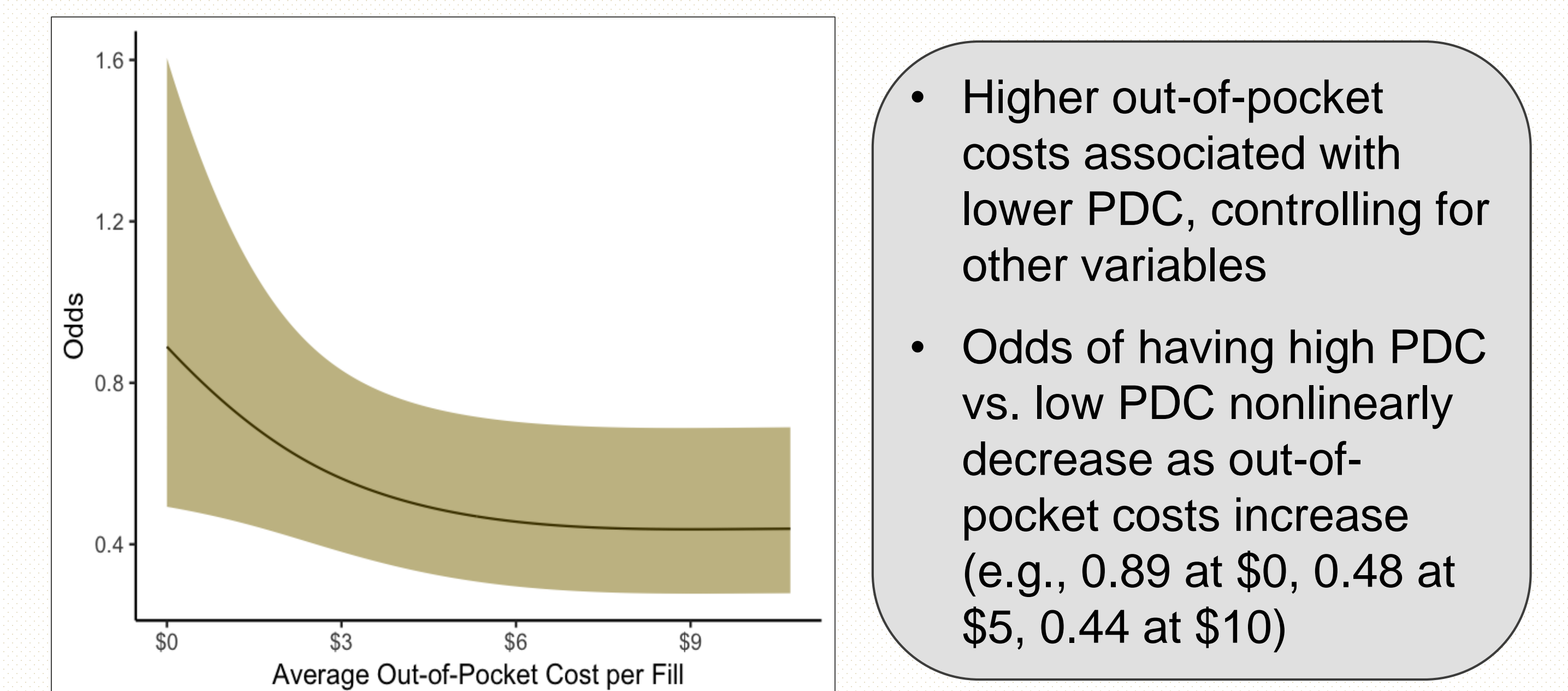


Table 2: Predictors of Higher Adherence (as Measured by PDC)

Variable	Odds Ratio	95% CI	p-value
Age (per 1 year)	1.01	0.98, 1.03	0.672
Female Gender (ref=Male)	0.47	0.27, 0.83	<b>0.009</b>
White Race (ref=non-White)	1.48	0.70, 3.15	0.312
Government Insurance (ref=Commercial)	1.20	0.66, 2.18	0.558
Non-naïve to Treatment (ref=naïve)	0.28	0.16, 0.50	<b>&lt;0.001</b>
Average Costs	See Figure 4		

- Males were 2.13 times more likely than females to achieve higher PDC
- Treatment naïve patients were 3.56 times more likely than non-naïve patients to achieve higher PDC

Figure 4: Association Between Adherence and Cost



- Higher out-of-pocket costs associated with lower PDC, controlling for other variables
- Odds of having high PDC vs. low PDC nonlinearly decrease as out-of-pocket costs increase (e.g., 0.89 at \$0, 0.48 at \$5, 0.44 at \$10)

## FINANCIAL OUTCOMES

Figure 5: Frequencies of Financial Assistance Use and Average Out-of-Pocket Costs per Fill

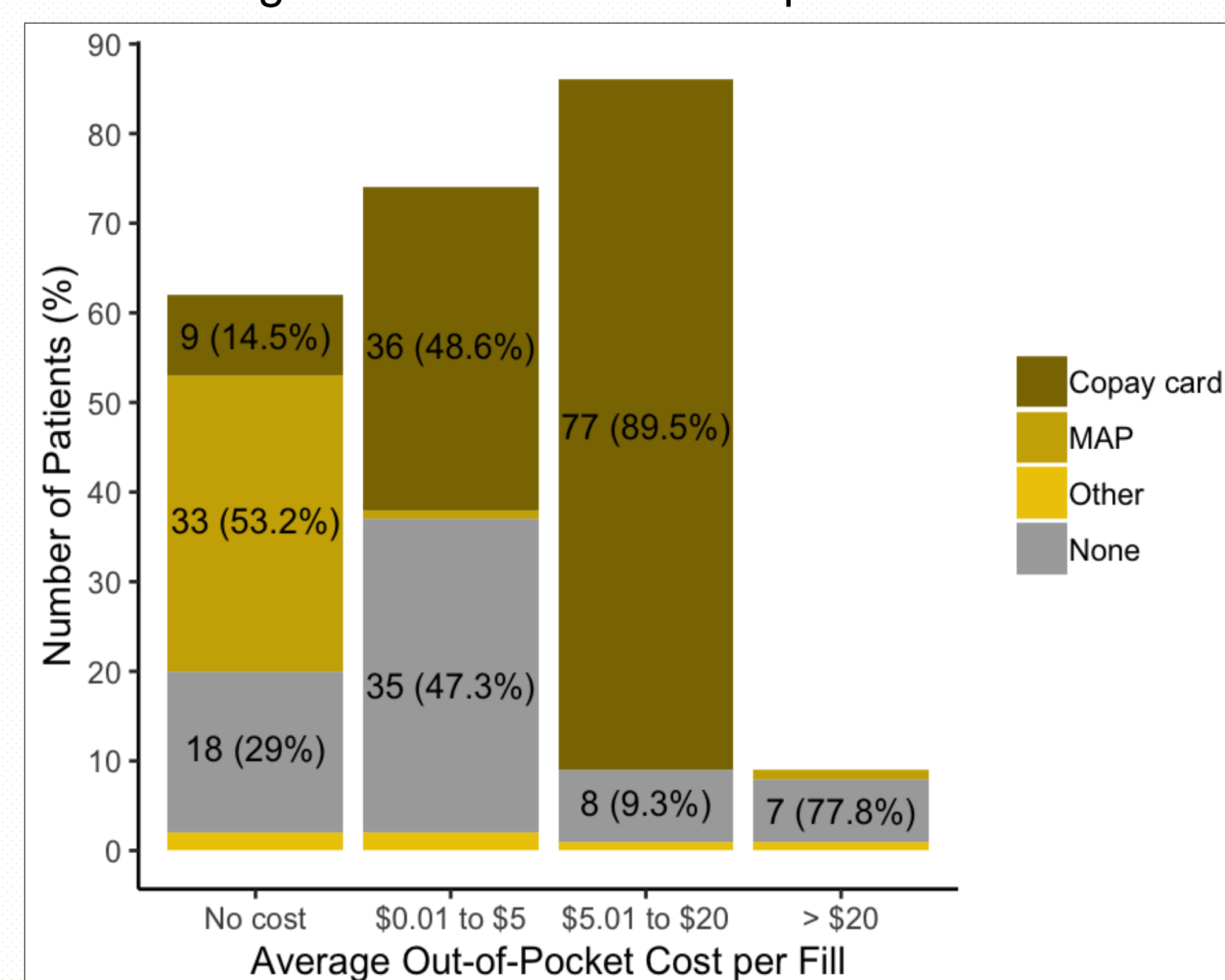
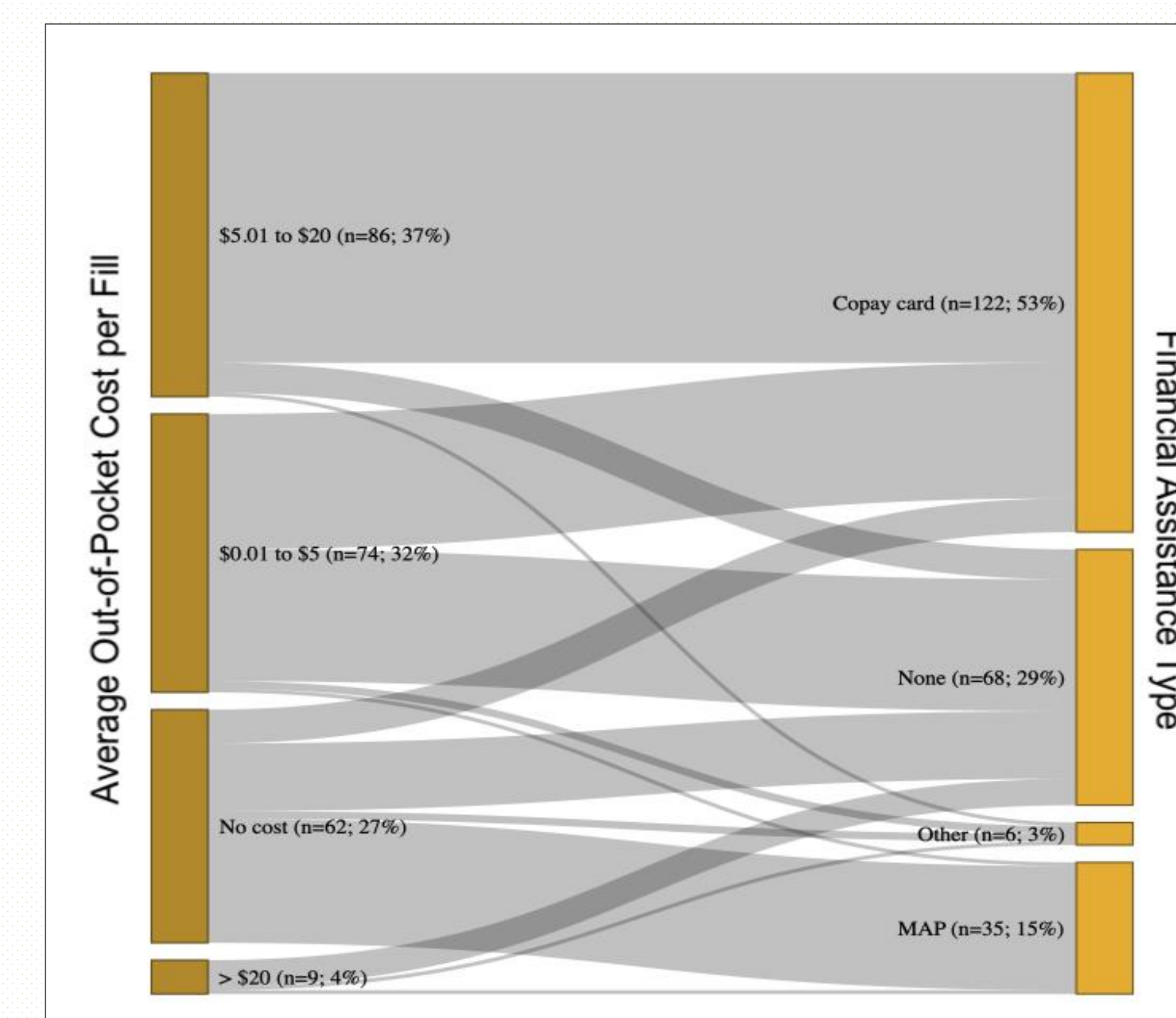


Figure 6: Average Out-of-Pocket Cost per Fill and Financial Assistance Distribution



- Average patient out-of-pocket cost was \$19.50 (SD=\$96.00) and 27% paid \$0.00, while only 4% paid more than \$20
- 71% of the sample received financial assistance (54% received copay card, 16% received medication assistance program, 1% received other type of assistance)

## CONCLUSIONS

- High rates of adherence to DMARDs were seen in a health-system integrated specialty pharmacy model
- Most patients received financial assistance to reduce out-of-pocket costs, which might contribute to high adherence
- Female patients and treatment non-naïve patients may be at higher risk for treatment non-adherence
- Further research is needed to quantify associations between specialty pharmacy interventions, DMARD adherence, and favorable clinical outcomes in patients with RA

## REFERENCES

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## DISCLOSURES

Authors of this presentation have the following to disclose concerning possible financial or personal relationships with commercial entities that may have a direct or indirect interest in the subject matter of this presentation. Autumn Zuckerman - Receives research support from Gilead Sciences, Inc. and Sanofi Inc. Nate Berger, Megan Peter, Joshua DeClercq, and Leena Choi - Nothing to disclose