



Improving ACEI/ARB Adherence Among Nonadherent Patients Using Motivational Interviewing Intervention

Mohan A¹, Majd Z¹, Johnson ML¹, Essien EJ¹, Barner JC², Serna O³, Gallardo E³, Fleming ML⁴, Ordonez N¹, Holstad MM⁵, Abughosh SM¹

¹University of Houston College of Pharmacy; ²The University of Texas at Austin; ³CareAllies; ⁴Department of Pharmacotherapy, Chapman University School of Pharmacy; ⁵Nell Hodgson Woodruff School of Nursing, Emory University

Contact Information:
Zahra Majd
University of Houston
Email:
zmajd@central.uh.edu

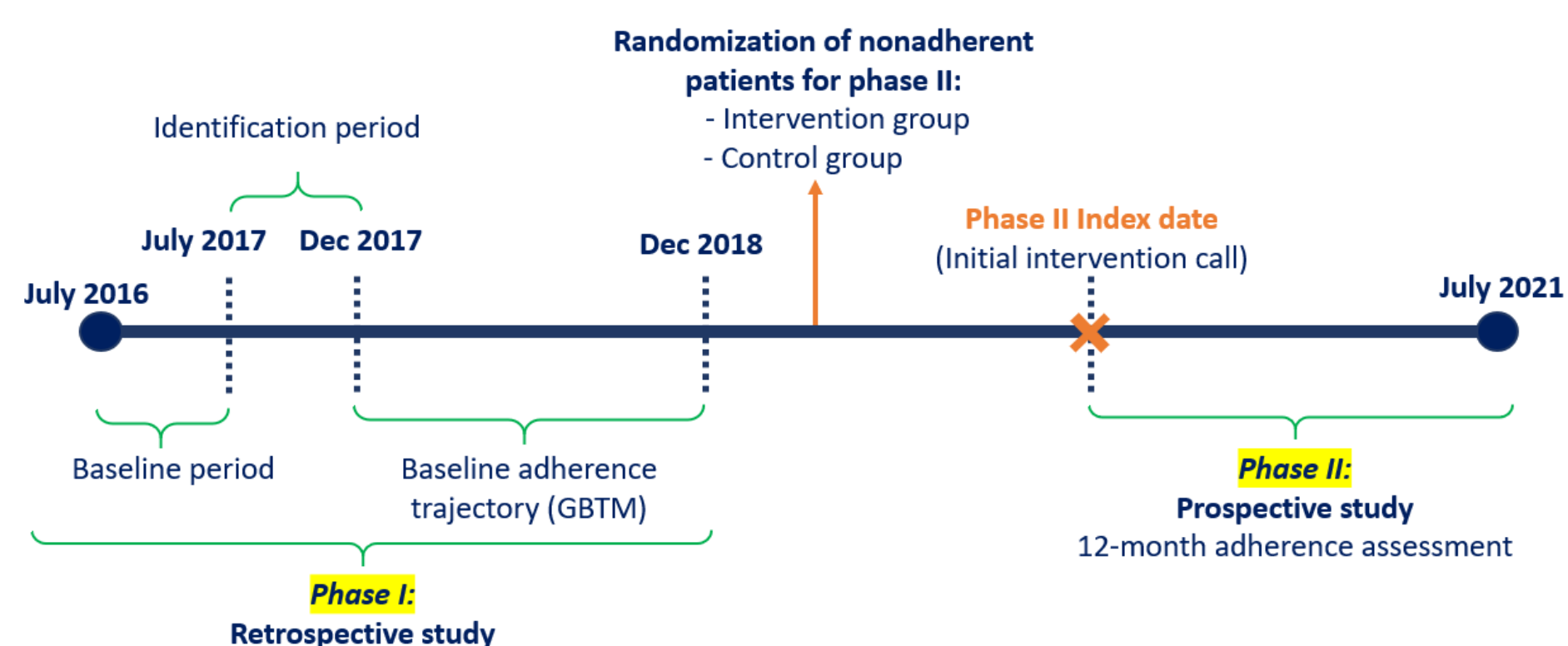
BACKGROUND

- Hypertension (HTN) is an important risk factor for cardiovascular (CV) diseases among patients with diabetes mellitus (DM).
- Despite demonstrated benefits of angiotensin-converting enzyme inhibitors (ACEI) and angiotensin receptor blockers (ARB), poor medication adherence is prevalent among DM patients.
- Poor adherence to ACEI/ARBs could lead to uncontrolled blood pressure, higher CV complications and mortality rate.
- Group-based trajectory modeling (GBTM) overcomes limitations of traditional adherence methods by describing longitudinal patterns of adherence.
- GBTM identifies clusters of patients with similar medication taking behavior over time which could be used in designing tailored adherence promotion interventions.
- Motivational interviewing (MI) is an effective patient-centered approach that improves medication adherence by identifying patient-specific barriers and exploring possible solutions.

OBJECTIVE

- To evaluate the effectiveness of a phone-based MI intervention conducted by pharmacy students and tailored by the past adherence trajectories on adherence to ACEI/ARBs among nonadherent patients with comorbid DM and HTN

Figure 1. Study design



METHODS

Data source:

- Administrative claims data from a Texas Medicare Advantage Plan

Inclusion criteria:

- ☑ Comorbid HTN and DM
- ☑ A refill for ACEI/ARB (July-Dec 2017)
- ☑ Continuous enrollment

Exclusion criteria:

- ☒ Diagnosis of dementia
- ☒ ACEI/ARB contraindications

MI Intervention:

- Initial call and 5 follow-up calls by MI-trained student pharmacists
- Customized by past adherence trajectories

Control group:

- Usual care

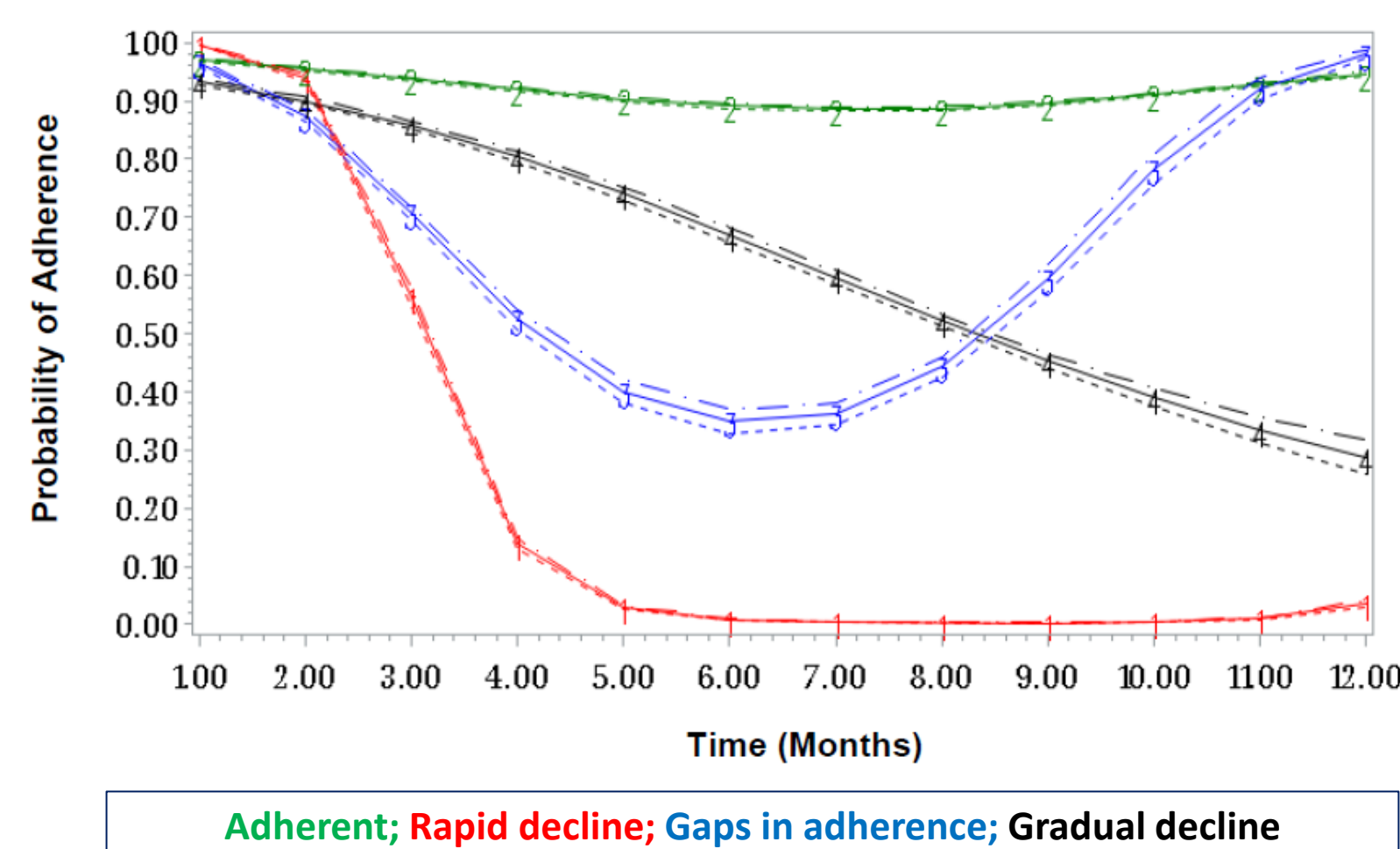
Adherence measurement:

- Proportion of Days Covered (PDC) measured following the initial call for the intervention and matched date for the control group
- PDC ≥ 0.80 considered as adherent

Statistical analysis:

- Descriptive statistics
- Multivariable logistic regression model:
 - Outcome: adherence during 12-month post-MI implementation
 - Primary predictor: intervention (≥ 4 calls, < 4 calls) vs control
- SAS version 9.4 (SAS Institute, Cary, NC)

Figure 2. Baseline adherence trajectories



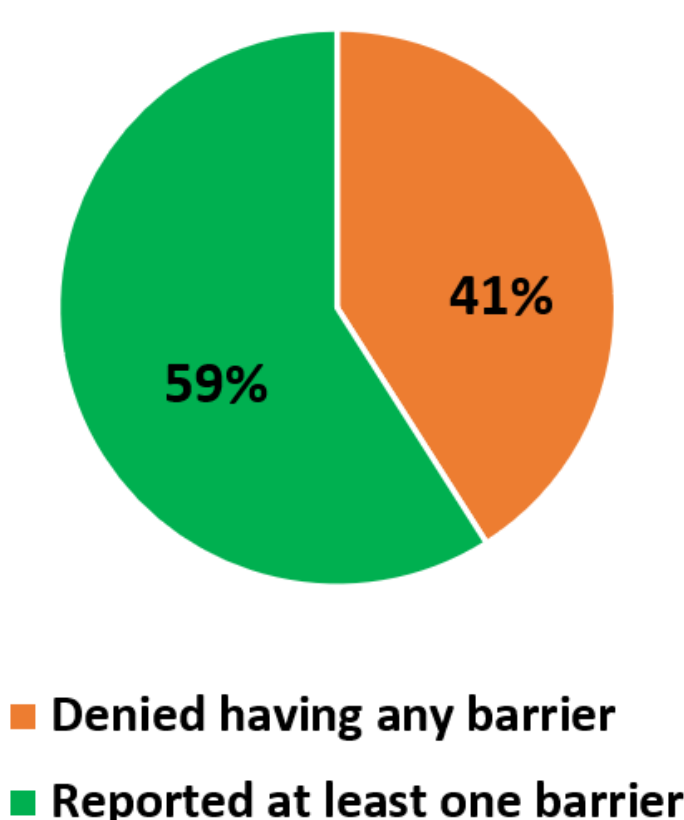
RESULTS

Table 1. Baseline characteristics of the study groups

Variable	Intervention (N=240)	Control (N=480)
Age, mean (SD)	68 (9.29)	69 (9.26)
Male, n (%)	145 (60.42)	270 (56.25)
Low-income subsidy beneficiaries, n (%)	118 (49.17)	247 (51.46)
General prescriber, n (%)	207 (86.25)	410 (85.41)
Myocardial infarction, n (%)	4 (1.67)	9 (1.88)
Depression, n (%)	31 (12.92)	43 (8.96)
Congestive heart failure, n (%)	26 (89.17)	441 (91.88)
Stroke, n (%)	10 (4.17)	13 (2.71)
Coronary artery disease, n (%) *	177 (73.75)	389 (81.04)
≥ 90 -day refill type, n (%)	214 (89.17)	438 (91.25)
≤ 2 co-medications, n (%)	169 (70.42)	361 (75.21)
No previous Hospitalizations	214 (89.17)	436 (90.83)
Prevalent use of ACEI/ARB, n (%) *	215 (89.58)	395 (82.29)
Regimen complexity, mean (SD) *	4.27 (4.32)	3.59 (3.76)
CMS risk score, mean (SD) *	1.50 (0.99)	1.31 (0.86)
Baseline adherence trajectories		
Gaps in adherence, n (%)	102 (42.50)	188 (39.17)
Gradual decline, n (%)	94 (39.17)	179 (37.29)
Rapid decline, n (%)	44 (18.33)	113 (23.54)

* Statistically significant difference

Figure 3. Adherence barriers identified from the initial calls



Most frequent barriers:

- Forgetfulness
- Side effects
- Comorbidities/polypharmacy
- Lack of knowledge about disease/medication
- Cost issues
- Family issues

Table 2. Logistic regression model (N=720)

Variables	Adherence vs nonadherence	
	OR (95% CI)	P value
No of calls		
<4 calls vs control	1.13 (0.77-1.65)	0.55
≥ 4 calls vs control	2.01 (1.17-3.44)	0.01
Previous trajectories		
Gaps in adherence vs rapid decline	3.19 (1.99-5.10)	<0.0001
Gradual decline vs rapid decline	2.63 (1.63-4.25)	<0.001
Gender (Male vs female)	1.03 (0.74-1.45)	0.82
Age	0.99 (0.97-1.01)	0.54
Health plan (No subsidy vs low-income subsidy)	0.88 (0.63-1.24)	0.49
Myocardial infarction (Yes vs no)	0.89 (0.22-3.49)	0.87
Depression (Yes vs no)	1.15 (0.67-1.96)	0.59
Congestive heart failure (Yes vs no)	1.22 (0.64-2.34)	0.54
Stroke (Yes vs no)	0.93 (0.34-2.46)	0.88
Coronary artery disease (Yes vs no)	0.92 (0.59-1.42)	0.71
Refill type (≥ 90 days vs. < 90 days)	1.26 (0.70-2.27)	0.42
Number of co-medications (> 2 vs ≤ 2)	1.94 (1.11-3.38)	0.01
Previous hospitalizations (≥ 1 vs none)	0.80 (0.43-1.51)	0.50
Prevalent users (Yes vs no)	1.68 (1.03-2.75)	0.03
Prescriber specialty (General vs Specialty)	0.69 (0.41-1.17)	0.17
Regimen complexity	0.93 (0.87-0.99)	0.03
CMS risk score	0.81 (0.66-0.99)	0.04

* 46.25% of the intervention and 42.09% of the control group were adherent during the follow-up period.

* Mean PDC (SD) for the intervention and control group were 0.63 (0.34) and 0.60 (0.33), respectively.

CONCLUSION

- Motivational interviewing with multiple patient encounters is a promising form of intervention that can improve adherence to ACEI/ARBs among nonadherent patients.
- Tailoring interventions based on past adherence trajectories may enhance intervention effectiveness as it provides more insight into the patient behavior.
- Future research should assess benefits of MI-based interventions on medication adherence over a longer follow-up period.

SPONSORSHIP

This study was funded by the National Heart, Lung, and Blood Institute (NHBLI), 1R15HL135700-01A1.