



STUDY OF A TELE-PHARMACY INTERVENTION FOR CHRONIC DISEASES TO IMPROVE TREATMENT ADHERENCE

THE STIC2IT RANDOMIZED CONTROLLED TRIAL

Niteesh K. Choudhry, MD, PhD

on behalf of:

Thomas Isaac, MD, MBA, MPH; Julie C. Lauffenburger, PharmD, PhD; Chandrasekar Gopalakrishnan, MD, MPH;
Nazleen F. Khan, BS; Marianne Lee, PharmD; Amy Vachon, PharmD; Tanya L. Iliadis, PharmD;
Whitney Hollands, PharmD; Scott Doheny, PharmD; Sandra Elman, PharmD; Jacqueline M. Kraft, PharmD;
Samrah Naseem, PharmD; Joshua J. Gagne, PharmD, ScD; Cynthia A. Jackevicius, PharmD, MSc;
Michael A. Fischer, MD, MS; Daniel H. Solomon, MD, MPH; Thomas D. Sequist, MD, MPH

Divisions of Pharmacoepidemiology and Pharmacoeconomics, Rheumatology, and General Internal Medicine, and Center for Healthcare Delivery Sciences, Department of Medicine, Brigham and Women's Hospital and Harvard Medical School; Atrius Health; Western University of Health Sciences; the Institute for Clinical Evaluative Sciences; Harvard Department of Health Care Policy

BACKGROUND

Medication non-adherence is extremely common

- **One-half of patients with cardiometabolic conditions do not adhere to their prescribed medications**
 - Leads to adverse clinical consequences and \$100-\$300 billion in preventable health spending each year in the U.S. alone
- **Interventions to improve adherence have been modestly effective**
 - Do not adequately address each individual's unique adherence barriers
 - Imprecisely targeted to patients who do not need adherence assistance
- **Even effective interventions are difficult to sustain**
 - Often Require new infrastructure and/or are expensive

OBJECTIVE

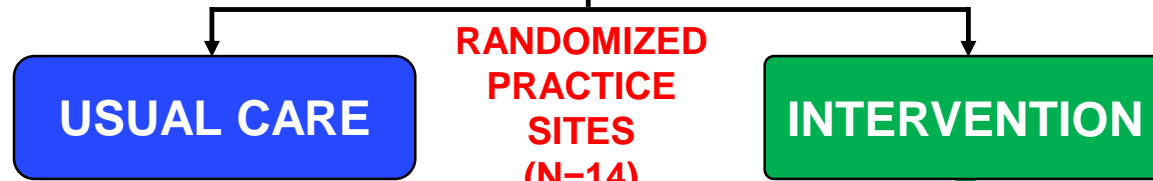
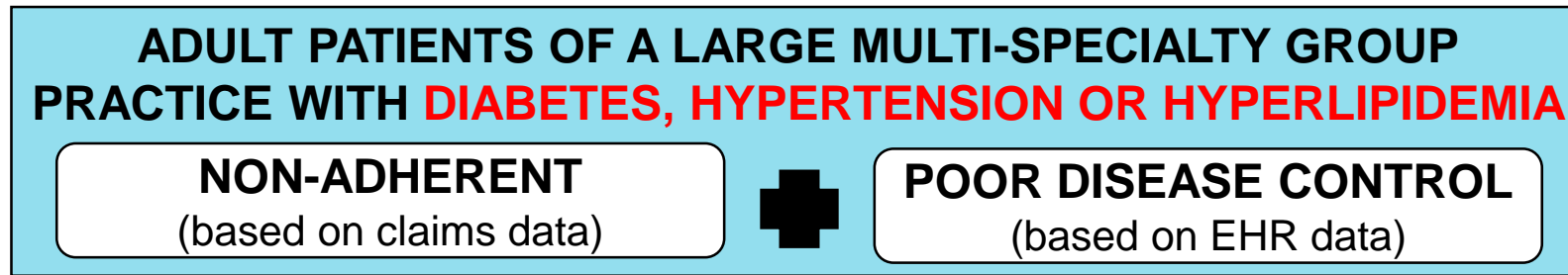
STIC2IT: Study of a Tele-pharmacy Intervention for Chronic diseases to(2) Improve Treatment adherence

- To evaluate the effect of a medication adherence intervention for diabetes, hypertension, and hyperlipidemia that was:



DESIGN

Open-label, pragmatic cluster-randomized trial



-
- CONTACTED AND OFFERED:**
- pharmacist telephone consultation (*using brief negotiated interviewing*)
 - text messages (*reminders or motivation*)
 - automated individual progress reports

Content tailored to “patient activation” + adherence barriers

- **ENROLLMENT:** Aug 2015-July 2016
- **END OF FOLLOW-UP:** July 2017

METHODS

Outcomes assessed using routinely-collected data

- Outcomes assessed during the 12 months after randomization

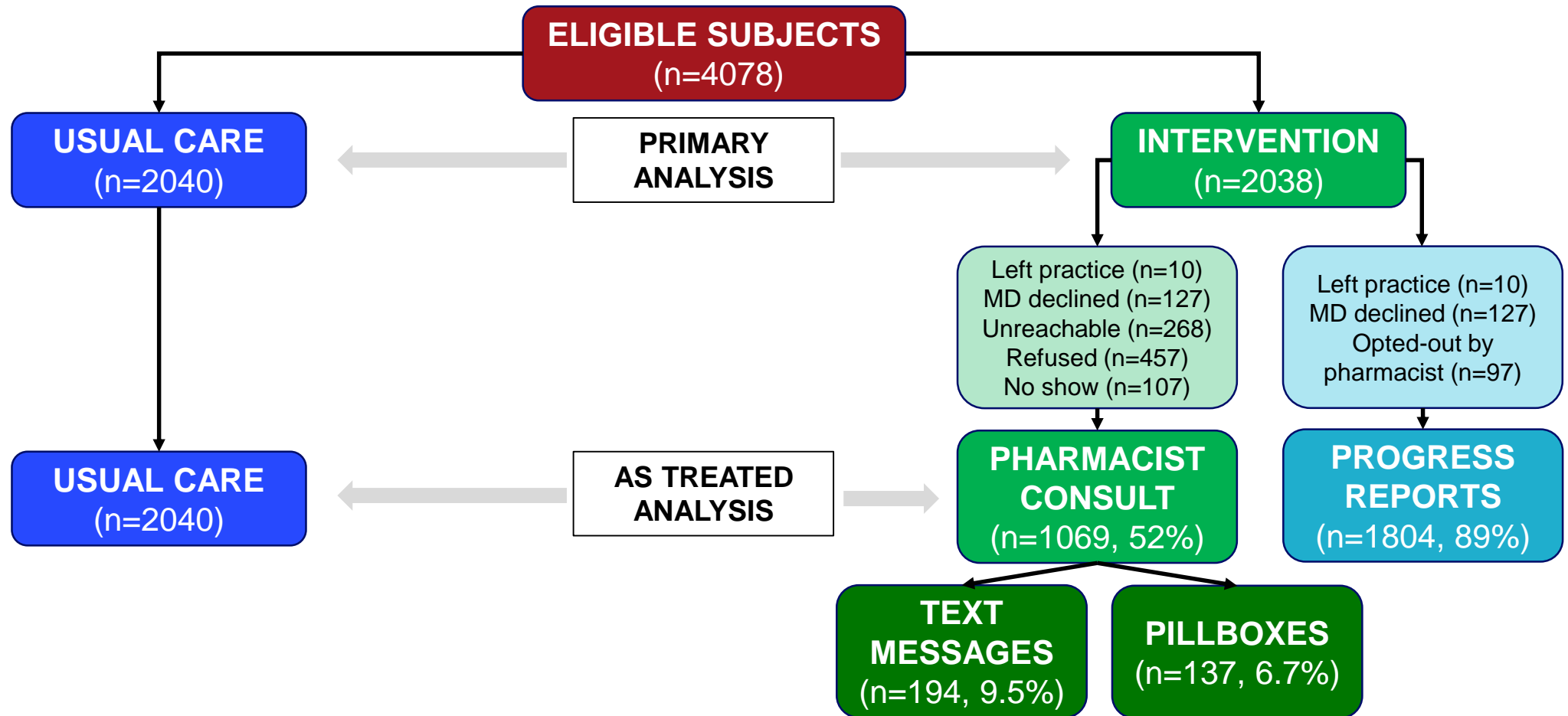
	Outcome	Data Source	Definition
1°	Medication adherence	Prescription health insurance data	Average adherence (“proportion of days covered”) for eligible medications at the time of randomization
2°	Disease control	Electronic health record data	Proportion of patients meeting guideline targets for: (a) all eligible conditions and (b) at least 1 eligible condition

- Primary analyses conducted on an intention-to-treat basis

- Powered for a 2.5% mean improvement in adherence assuming that <50% of patients would agree to a pharmacist consultation

RESULTS

Enrollment



Clinical pharmacist telephone consultations lasted a mean of 24.9 minutes; 1050 (98.2%) patients completed at least 2 calls and 175 (16.4%) patients received 3 or more calls



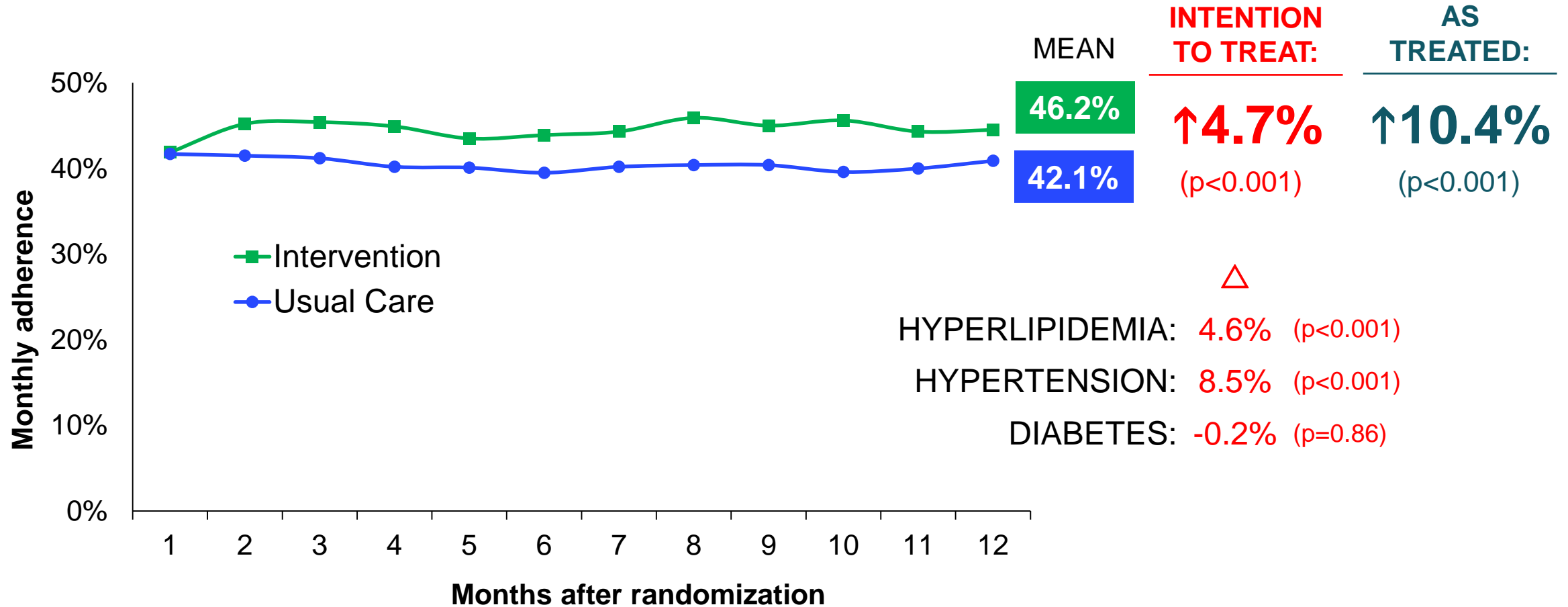
RESULTS

Baseline characteristics

CHARACTERISTIC	USUAL CARE (N=2040)	INTERVENTION (N=2038)
Age, mean years*	60.4	59.2
Male sex	54.7%	55.0%
White race*	53.6%	60.6%
Qualifying conditions		
Hyperlipidemia	72.0%	73.7%
Hypertension	25.9%	23.8%
Diabetes	12.1%	11.9%
Charlson comorbidity score, mean	0.90	0.74
Baseline disease control		
LDL cholesterol, mean mg/dL,	204.8	207.8
Systolic blood pressure, mean mmHg	149.9	149.2
Hemoglobin A _{1c} , mean	9.8	9.5
Baseline adherence, mean	57.0%	57.2%

* Standardized mean difference for age and race/ethnicity were >0.1; there were no other significant differences

PRIMARY OUTCOME
Adherence

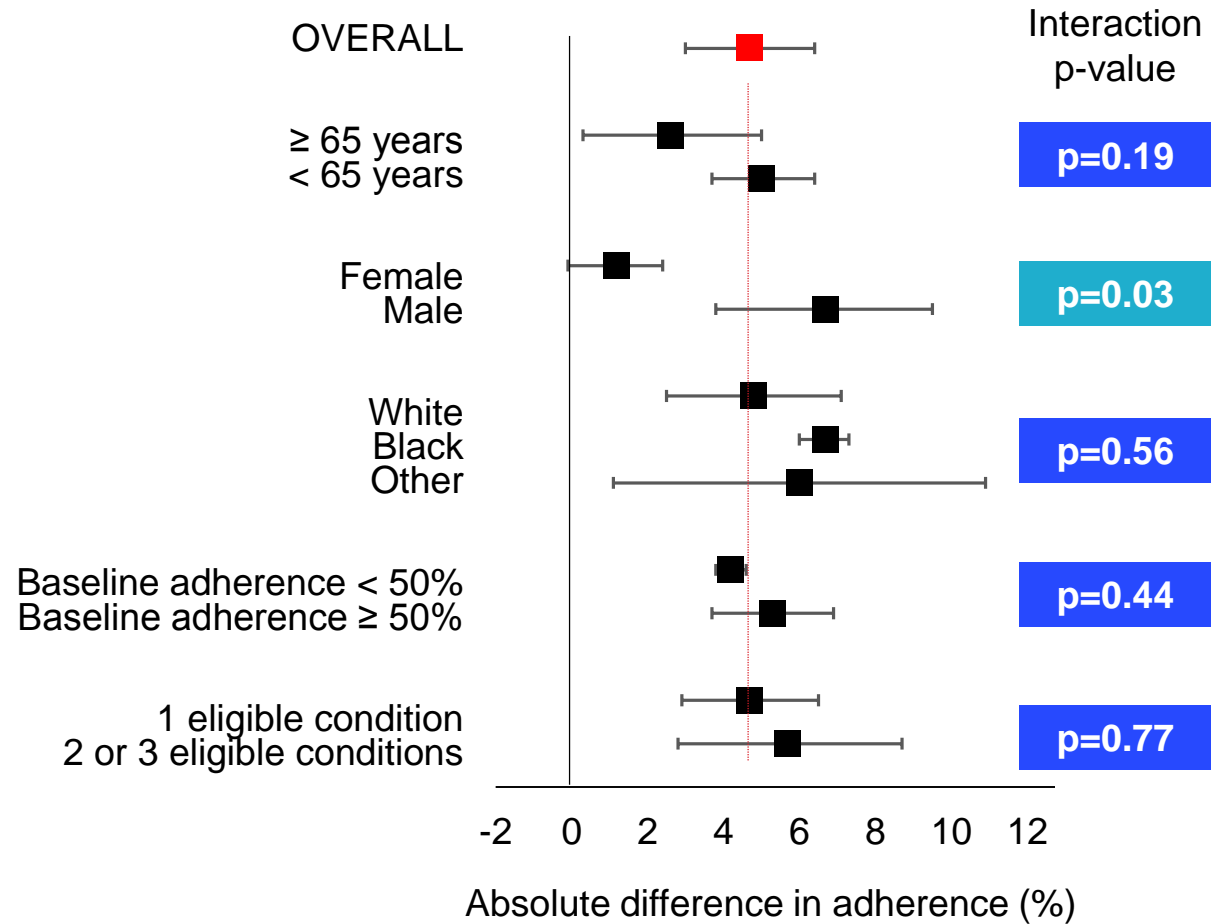


Median (IQR) time from randomization to pharmacist call (when it occurred): 22 (17 to 32) days



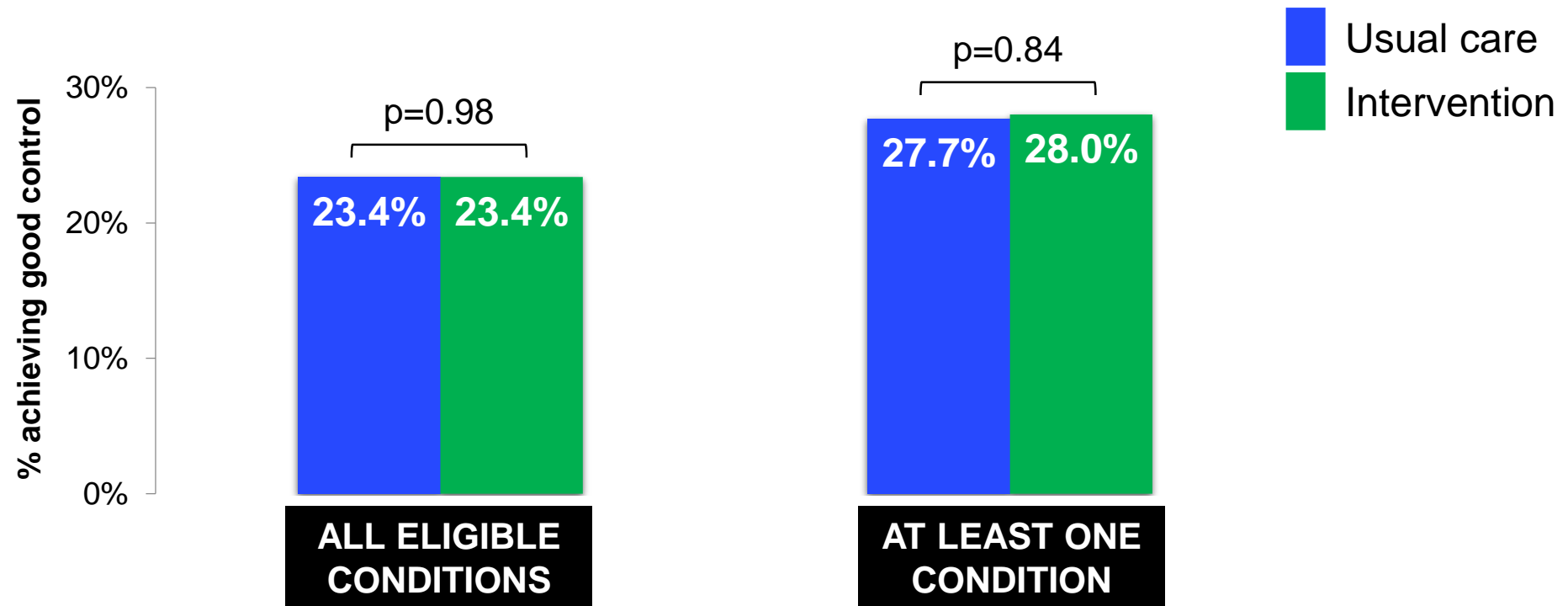
SUBGROUP ANALYSES

Adherence



SECONDARY OUTCOMES

Good disease control



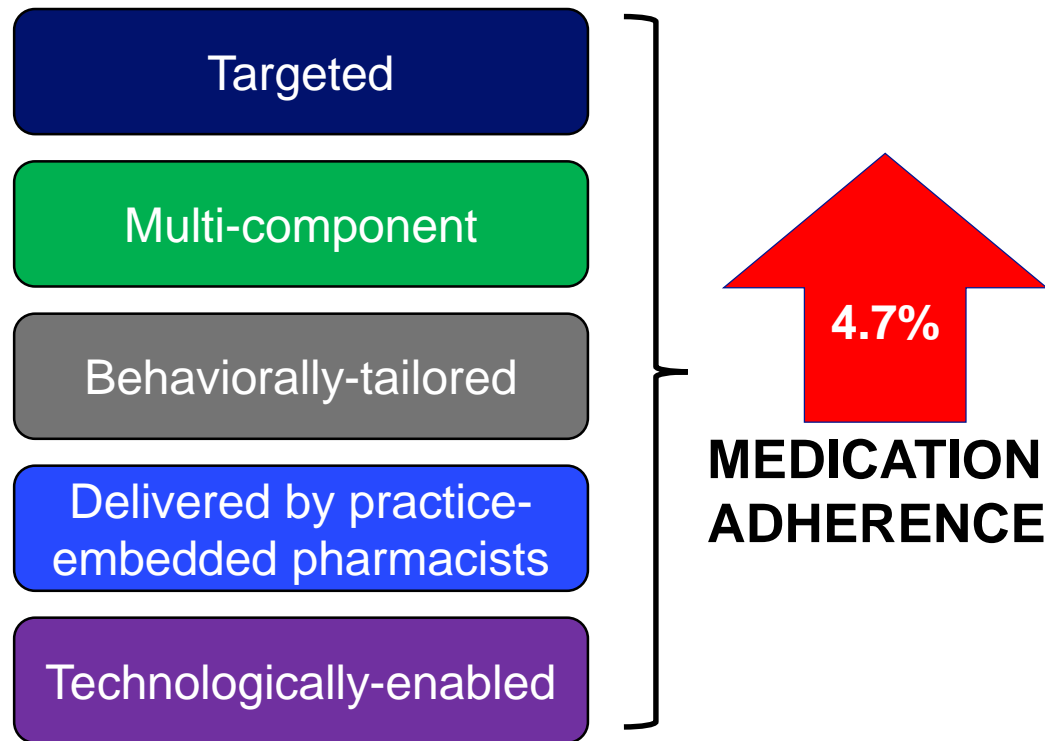
Mean duration between randomization and outcome assessment: 229.2 days



SUMMARY

The STIC2IT intervention improved adherence

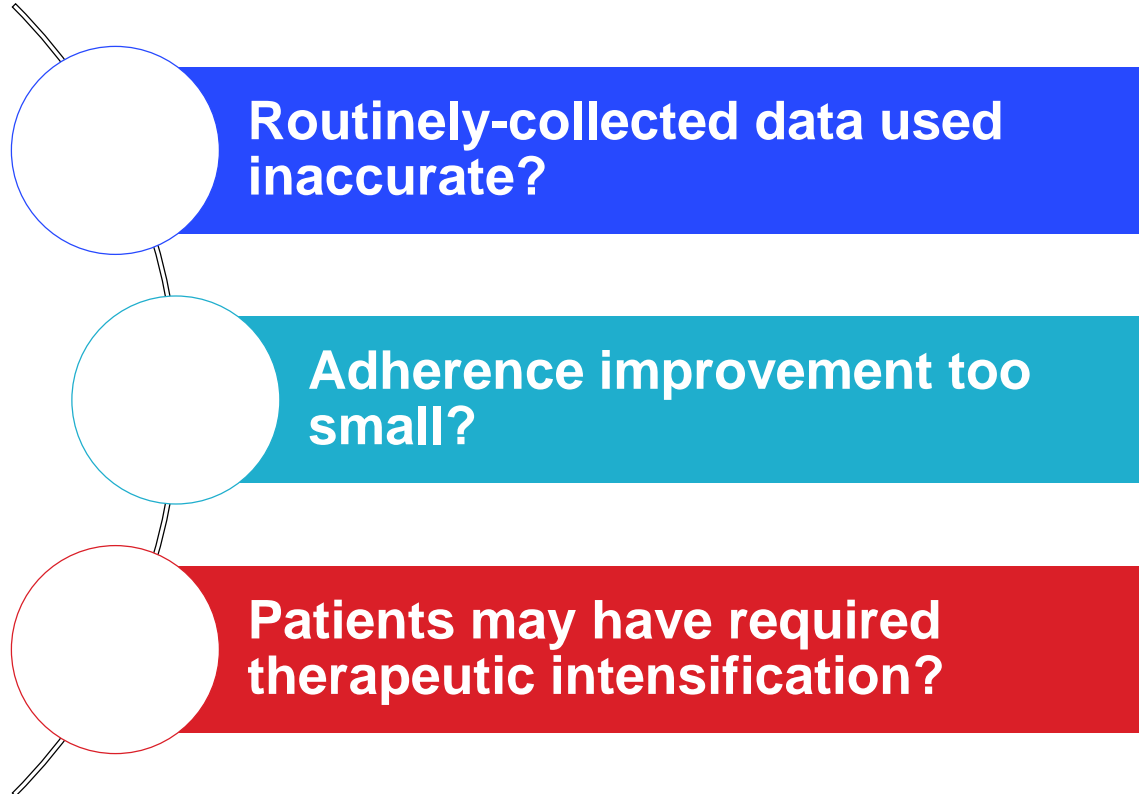
- An intervention for patients with diabetes, hypertension, and hyperlipidemia with poor medication adherence and suboptimal disease control:



- Effect size was similar to those achieved by more labor intensive interventions
- Used highly-pragmatic research methods to facilitate the generalizability of the results

SUMMARY AND IMPLICATIONS

Intervention did not improve secondary clinical outcomes



FUTURE INTERVENTIONS MAY NEED TO:

- Be more intensive while still pragmatic
- Focus on a more impactable patient population
- Simultaneously address adherence and other barriers to optimal disease control

Niteesh K. Choudhry, MD, PhD
Brigham and Women's Hospital/Harvard Medical School

E: nkchoudhry@bwh.harvard.edu