Impact of AllazoEngine-Targeted Interventions on Medication Adherence: Repeated Measures Difference-in-Differences Analysis

Brittanie Gracey¹, BS; Patricia Prince¹, MPH; Clifford Jones¹, BSE; Suzy Connor², BCACP, PharmD; Estay Greene², PharmD

¹AllazoHealth, New York, NY, United States ²Blue Cross and Blue Shield of North Carolina, Durham, NC, United States

Corresponding Author: Clifford Jones, BSE AllazoHealth New York, NY, United States Email: cliff@allazohealth.com

Abstract

Background: AllazoHealth utilizes predictive analytics to improve medication adherence by targeting patients whose behavior can be changed with intervention programs. The AllazoEngine utilizes Rx claims, previous intervention data, and demographics to predict future adherence, to prioritize the patients whose behaviors can be changed, and to select the intervention channel and messaging most effective for each individual patient. Blue Cross and Blue Shield of North Carolina commissioned AllazoHealth's predictive analytics and separately commissioned medication intervention delivery services for this adherence program.

Objective: This study aimed to evaluate the effectiveness of the AllazoEngine and targeted interventions to improve medication adherence.

Methods: This was a double-blind, randomized controlled trial (RCT) focused on RAS antagonists, oral anti-diabetics, and Statins. Patients were randomized to receive no intervention, traditional non-Allazo-targeted interventions, or interventions targeted by the AllazoEngine. All interventions consisted of live calls, direct mail to patients, and faxes to prescribers. Patients were defined as adherent in accordance with Medicare Star ratings methodology if their proportion of days covered (PDC) was greater than 80%. Patients' adherence status in 2015 was compared to their adherence status in 2016 after the intervention period. Difference-in-Differences (DiD) analysis was used to compare the effect of each intervention method. Statistical significance was set to 10%.

Results: The primary study population consisted of 14,377 controls, 5,423 traditional non-Allazo targeted-intervention patients and 24,527 Allazo targeted intervention patients. Patients had comparable characteristics at baseline and comparable decrease in medication adherence in the pre-intervention observation period across the intervention groups. Non-Allazo-targeted interventions did not statistically improve the likelihood of adherence. Patients who received Allazo-targeted interventions performed statistically better than both the non-Allazo targeted group and control group (P=.06 and .03). Assuming net positive uplift from non-Allazo interventions. Allazo interventions accounted for 7.7 times the per-patient uplift in adherence compared to non-Allazo interventions.

Conclusions: Due to the specific study design of not including new patients, adherence decreased in each intervention group over the intervention period. However, the decrease was significantly less for those in the Allazo group compared to both the non-Allazo and control groups.

(*iproc 2017;3(1):e13*) doi: <u>10.2196/iproc.8427</u>

KEYWORDS

diabetes; medication adherence; star-ratings; hypertension; predictive analytics



IPROCEEDINGS

Edited by T Hale; this is a non-peer-reviewed article. Submitted 12.07.17; accepted 25.08.17; published 22.09.17. <u>Please cite as:</u> Gracey B, Prince P, Jones C, Connor S, Greene E Impact of AllazoEngine-Targeted Interventions on Medication Adherence: Repeated Measures Difference-in-Differences Analysis iproc 2017;3(1):e13 URL: http://www.iproc.org/2017/1/e13/ doi: 10.2196/iproc.8427 PMID:

©Brittanie Gracey, Patricia Prince, Clifford Jones, Suzy Connor, Estay Greene. Originally published in Iproceedings (http://www.iproc.org), 22.09.2017. This is an open-access article distributed under the terms of the Creative Commons Attribution License (https://creativecommons.org/licenses/by/4.0/), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work, first published in Iproceedings, is properly cited. The complete bibliographic information, a link to the original publication on http://www.iproc.org/, as well as this copyright and license information must be included.

