Supplemental Online Content

Zupa MF, Vimalananda VG, Rothenberger SD, et al. Patterns of telemedicine use and glycemic outcomes of endocrinology care for type 2 diabetes. *JAMA Netw Open*. 2023;6(12):e2346305. doi:10.1001/jamanetworkopen.2023.46305

eTable 1. ICD-10 Codes Used to Identify Comorbid Conditions

eTable 2. Characteristics of Modeled vs Excluded Patients According to Presence of Follow-Up HbA_{1c}

eTable 3. Supplemental Analysis: Adjusted Difference HbA1c Change for Each Care Modality Cohort by Insulin Use

This supplemental material has been provided by the authors to give readers additional information about their work.

Cardiovascular	
Coronary artery disease	I20.X, I21.X, I22.X, I23.X, I24.X, I25.X
Congestive heart failure	I50.X, I42.X, I110.X, J19.X
Arrhythmia	I47.X, I48.X, I49.X
Cerebrovascular disease	I60.X, I61.X, I62.X, I63.X, I64.X, I65.X, I66.X, I67.X, I68.X, I69.X,
	DG45.X, DG46.X
Vascular disease	I70.X, I71.X, I72.X, I73.X, I74.X, I75.X, I76.X, I77.X, I78.X, I79.X
Gangrene	I96.X, E11.5X
Psychologic	
Psychotic disorders	F20.X, F21.X, F22.X, F23.X, F24.X, F25.X, F28.X, F29.X
Mood disorders	F30.X, F31.X, F32.X, F33.X, F34.X, F39.X
Post Traumatic Stress Disorder	F43.X
Anxiety disorders	F40.X, F41.X
Alcohol use disorder	F10.X
Substance use disorder	F11.X, F12.X, F13.X, F14.X, F15.X, F16.X, F18.X, F19.X

eTable 1. ICD-10 Codes Used to Identify Comorbid Conditions^a

a. Based on prior work by Pentakota et al²⁵

Characteristic	Total (N=4,275)	Included (N=3,778)	Excluded (N=497)	P- Value ^a	
Age (years); Mean (SD)	59.4 (13.3)	60.3 (12.7)	52.8 (15.8)	<0.001	
Gender Female N (%) Male N (%)	2,547 (60%) 1,728 (40%)	2,201 (58%) 1,577 (42%)	346 (70%) 151 (30%)	<0.001	
Race N (%) White Black Asian Other/Missing	3,759 (88%) 347 (8%) 87 (2%) 82 (2%)	3,332 (88%) 300 (8%) 81 (2%) 65 (1%)	427 (86%) 47 (9%) 6 (1%) 17 (3%)	0.02	
Hispanic or Latino N (%) Not specified/missing	78 (2%) 197 (5%)	66 (2%) 158 (4%)	12 (2%) 39 (8%)	<0.001	
SDI Score; Mean (SD)	40.7 (24.0)	40.5 (23.9)	41.5 (25.0)	0.40	
RUCA; N (%) Urban Suburban Rural	3,024 (71%) 921 (22%) 330 (8%)	2,645 (70%) 836 (22%) 297 (8%)	379 (76%) 85 (17%) 33 (7%)	0.02	
HbA1c (%); Mean (SD)	7.5 (1.8)	7.6 (1.7)	7.2 (2.1)	<0.001	
Insulin N (%) No insulin Basal only MDI	1,763 (41%) 703 (16%) 1,809 (42%)	1,476 (39%) 652 (17%) 1,650 (44%)	287 (58%) 51 (10%) 159 (32%)	<0.001	
Non-insulin med count; Mean (SD)	1.8 (1.1)	1.9 (1.1)	1.5 (1.0)	<0.001	
BMI categorized; N (%) 18.5-24.9 25-29.9 30-34.9 35-39.9 ≥40 Missing	217 (5%) 693 (16%) 965 (23%) 757 (18%) 867 (20%) 776 (18%)	196 (5%) 639 (17%) 880 (23%) 688 (18%) 770 (20%) 605 (16%)	21 (4%) 54 (11%) 85 (17%) 69 (14%) 97 (20%) 171 (34%)	<0.001	
Comorbid conditions N (%) Macrovascular Psychologic	1,526 (36%) 1,413 (33%)	1,393 (37%) 1,246 (33%)	133 (27%) 167 (34%)	<0.001 0.78	
Appointments/12 months; Mean (SD)	2.5 (1.0)	2.6 (1.0)	1.7 (0.7)	<0.001	
Number of follow-up HbA1c test results/12 months; Mean (SD)	1.5 (0.9)	1.7 (0.8)	0 (0.0)	<0.001	

eTable 2. Characteristics of Modeled vs Excluded Patients According to Presence of Follow-Up HbA_{1c}

a. For continuous variables, Kruskal-Wallis test was used; for categorical variables, Chi-square test was used

eTable 3. Supplemental Analysis: Adjusted Difference HbA1c Change for Each Care Modality Cohort by Insulin Use

Clinical Characteristic	Difference in Estimated HbA1c change at 12- months ^a (95% CI)	P value	Difference in Estimated HbA1c change at 24- months ^a (95% CI)	P- value			
No insulin (n=1,476)							
Mixed vs. telemedicine only Office vs telemedicine only	-0.09 (-0.36, 0.17) -0.22 (-0.53, 0.09)	0.990 0.324	-0.03 (-0.41, 0.34) -0.11 (-0.52, 0.31)	>0.999 >0.999			
Basal insulin only (n=652)							
Mixed vs. telemedicine only Office vs telemedicine only	-0.15 (-0.45, 0.15) -0.23 (-0.58, 0.12)	0.652 0.382	-0.34 (-0.78, 0.09) -0.36 (-0.85, 0.13)	0.250 0.297			
Multiple daily injections (n=1,650)							
Mixed vs. telemedicine only Office vs telemedicine only	-0.24 (-0.51, 0.02) -0.47 (-0.78, -0.15)	0.146 0.007	-0.03 (-0.39, 0.33) -0.19 (-0.60, 0.22)	>0.999 0.734			

a. Adjusted differences in HbA1c change are model-based estimates of the difference in HbA1c change from baseline between each category compared to the reference group, as indicated. These model-based estimates were obtained from linear mixed modeling of repeated measures of HbA1c adjusted for patient age, gender, race, ethnicity, social deprivation index, rurality, baseline HbA1c, and BMI; patients were nested within providers