Improvement of Diabetes Mellitus Management in a Residentrun Clinic by using Continuous Glucose Monitoring (CGM)

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Introduction

- Diabetes Mellitus (DM) is a metabolic disorder that affects more than 34 million people of all ages in the USA, which is around 9-10% of the US population. The main subtypes are Type 1-DM and Type 2-DM [1].
- Self-monitoring of blood glucose does not provide an accurate picture of the glucose trends throughout a 24 hour day [2].
- Continuous glucose monitoring (CGM) is a more accurate and advanced method for monitoring 24-hour blood glucose.
- Randomized & observational studies of real-time CGM systems have demonstrated improved glucose control measured by time in range (TIR), decreased glucose variability measured by coefficient of variation (CV), decreased episodes of hyperglycemia and hypoglycemia compared to SMBG [3].
- CGM was started at our Internal Medicine residency primary care clinic for better glycemic control.

Objective

Would implementation of CGM improve Type 1 and 2 DM management in the MountainView Clinic employed by Internal Medicine and Transitional Year Residents?

	Methods				
Inclusion Criteria:	Exclusion Criteria:				
Age 18-80 years old	Patients wearing their CGM < 70% of the time	Re			
Diagnosis of Type 1 or Type 2 DM	Patients unresponsive to calls from the clinic				
Patients with uncontrolled blood glucose levels while using SMBG \geq 4 times daily	Patient non-compliant with dietary and exercise recommendations	Mc			
Patients on 3-4 Insulin injections medications	Patients unable to understand the	Nu			
Patients with HbA1c > 7% and who were only	instructions for titration of insulin based on CGM data	Pa In-o 3 n			
receiving their primary care in the Resident-run MountainView Clinic in Nevada	Patient with impaired decision-making capacity	up p wee die we			
Patient can use a CGM device	Patient missing > 2 scheduled clinic visits	a			
Patient can adjust their insulin based on CGM device					

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Type of Study: letrospective study

> Study Period: 2020 - 2023

Study Location:

Resident-run IountainView Clinic

umber of patients: 51

atient counseling:

office visits every 2months and followphone calls every 2 eks. Counseling on et and exercise, as ell as insulin-dose adjustments were performed.

Methods continued

- Data was extracted from the eClinicalWorks electronic medical record system and CGM devices (Freestyle Libre 3, DEXCOM G6/7). Paired t-test & Wilcoxon signed-rank test were used for the following data on HbA1c and BG pre- and post-CGM:
- - HbA1c
 - Time-in-Range (TIR) of Blood Glucose 70-180 mg/dL
 - Average blood glucose
 - Mild hypoglycemia 54-70 mg/dL
 - Pronounced hypoglycemia <54 mg/dL
- Initial measurements were taken and the final measurements were averages over time, calculated at end of study period.

Doculte										
Results										
Table 1										
Descriptive Statistics of Measured Variables										
Variable Me			Median	ledian Mean			SD			
PreA1c	9.2			9.914		2.217				
PostA1c		7.4			7.625		1.34			
PreTIR		31			0.336		0.207			
PostTIR		67			0.669		0.202			
PreAvgB	3G 230			242.255		65.485				
PostAvgE	vgBG 167			169.431		34.581				
PreMildH	eMildHypo 5			0.047		0.006				
PostMild	PostMildHypo 0			0.008			0.014			
PreProHy	/ро		3.1 0.031 0.013		013					
PostProH	PostProHypo 0 0.002 0		0.0	0.006						
Note: TIR = Time in Range (BG 70-180 mg/dL); AvgBG = Average Blood Glucose; MildHypo										
= Mild Hy	poglycemia	a (54-70 m	g/dL); Prol	Hypo = Pro	onounced H	lypoglyce	mia (< 54	mg/dL)		
Table 2										
Results o	f Paired T-	Test and W	licoxon Si	gned-Ranl	k Test					
Variable		Mean Diff	95% CI	of Mean				Cohen's		
Pair	Mean Diff	SD	D	iff	t	df	р	D		
PreA1c										
PostA1c	-2.288	-2.216	-1.665	-2.912	-7.373	50	0	-1.032		
PreTIR		~~ ~~~								
PostTIR	33.333	39.723	44.505	22.216	5.993	50	0	0.839		
PreAvgB										
G PostAvg										
BG	-72.824	-68.563	-53.540	-92.107	-7.585	50	0	-1.062		
Variable	12:021	Med Diff		02.101	1.000		0	11002		
Pair	Med Diff	SE			Z	df	р			
PreMildH										
уро										
PostMild										
Нуро	-3.1	-102.375	-	-	-6.193	50	0	-		
PreProH										
ypo De et Dree										
PostPro	ΓO	10/ 100			6 001	50	0			
Hypo Noto: Diff	-5.0	-104.199	- Andian: TI	- D - Tima :	-6.224	50 20 70 180	•			
		-	-		n Range (E emia (54-7)		• / ·	•		
Average Blood Glucose; MildHypo = Mild Hypoglycemia (54-70 mg/dL); ProHypo = Pronounced Hypoglycemia (< 54 mg/dL)										

Pronounced Hypoglycemia (< 54 mg/dL)

- 10% of participants discontinued their insulin due to better diabetic control and continued their oral diabetic medications.
- The quality of life improved in patients.
- Decreased frequency of unproductive office visits for DM follow-ups due to accessible and convenient data extraction from CGM devices.
- Successful implementation of CGM in a Resident-run clinic with proficiency of Residents to manage Type 1 and 2 DM patients on CGM.



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- events, and time in range (Table 1 and 2).
- responsible for.
- (i.e. anemia, CKD).
- type 1 and type 2 DM.
- care clinics.

- 2 2020,43(Suppl 6).
- 3
- Jan(1):91-97





Discussion

In our three-year study, we found switching from SMBG to CGM led to a significant improvement in the health outcomes of the 51 patients with type 1 or type 2 DM. We observed significant improvements in HbA1c levels, blood glucose levels, hypoglycemic

Our findings demonstrates that using CGM instead of SMBG methods can lead to better glycemic control, fewer side effects, and lifestyle improvements [4, 5]. To our knowledge, this is first time CGM is successfully implemented into a resident-run Internal Medicine primary care clinic (instead of an endocrine clinic), managed by Internal Medicine and Transitional Year residents. These results are likely due to the combination of medical resident vigilance and relatively smaller number of patients each resident is

CGM can also help with decreasing the frequency of unproductive office visits for DM management due to data accessibility and convenience. This leads to streamlined management, better provider experience, improved patient quality of care and satisfaction, and improved client health outcomes by maximizing health-related quality of life and reducing costs.

Some limitations of our project includes: small sample size of 51 patients, relative short project timeline of 3 years, differences of patient compliance with diet and lifestyle interventions, presence of other potential confounding diagnoses that may affect HbA1c levels

Conclusion

We have demonstrated a model that other Internal Medicine residency programs can implement to improve the quality of care for difficult-to-treat patients on multiple injections of Insulin per day with

Randomized prospective trials with larger sample sizes are needed to fully assess the potential of CGM implementation and guided treatment of diabetes mellitus in Internal Medicine residency primary

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