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## Background/Objective

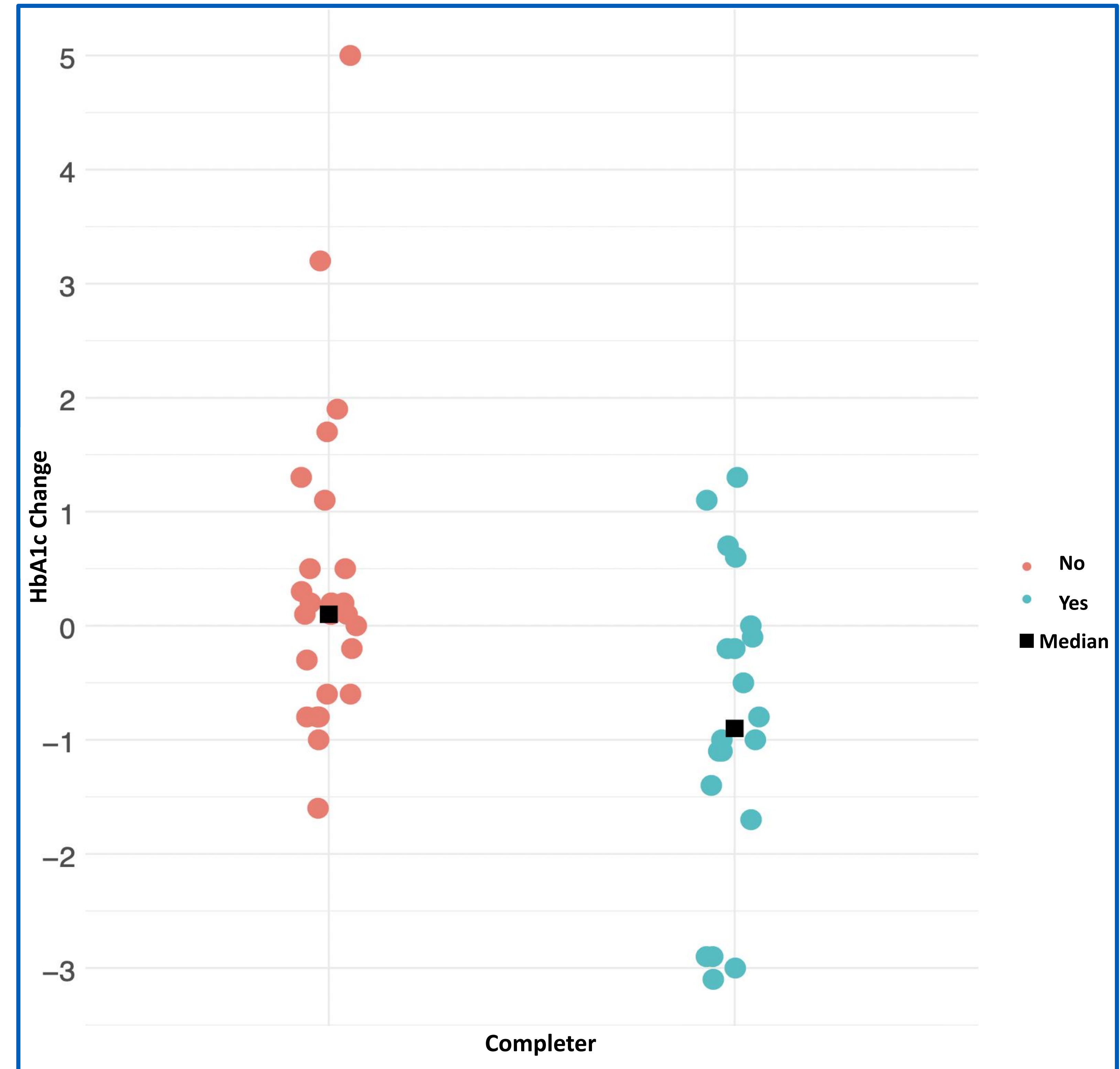
Deterioration of glycemic control between diabetes clinic visits is common in youth and young adults (YYA) with T1D. Remote Patient Monitoring (RPM) is a telehealth intervention that offers patients additional one-on-one visits with healthcare providers between standard clinic visits, focusing on optimizing diabetes self-management. We investigated whether YYA with elevated HbA1c who participate in RPM, versus those who do not, experience improved HbA1c.

## Methods

All patients had  $\geq 2$  elevated HbA1c results ( $\geq 9.0\%$ ) within 12 months and were offered RPM. Of these, 20 patients enrolled and completed at least one 15–20-minute RPM visit; 26 patients either declined, did not attend their appointment or were lost to follow-up. We evaluated change in HbA1c by comparing baseline HbA1c and HbA1c obtained 3 months after RPM was offered.

## Results

HbA1c improved in 75.0% (15/20) of completers and 34.6% (9/26) of non-completers. In completers, median change in HbA1c was  $-0.9\%$ ; in non-completers, median change was  $+0.1\%$ . In completers whose HbA1c improved, median change was  $-1.1\%$ .



## Data Management

- Patients are screened based off their most recent clinic visit; of those qualifying, approximately 5 are offered RPM each week.

### HbA1c results included in analysis

- If patients had a subsequent HbA1c result recorded prior to RPM outreach than their qualifying HbA1c, this was considered their baseline.
- If patients did not have a HbA1c result recorded within 3 months ( $\pm 2$  weeks), their baseline HbA1c and a subsequent HbA1c result were used to interpolate an approximate 90-day result.

## Conclusions and Future Work

Completers experienced a substantial decrease in median HbA1c. By facilitating proactive diabetes management, RPM for YYA with elevated HbA1c ( $\geq 9.0\%$ ) may lead to significant improvements in glycemic control. Future research with larger matched patient cohorts is warranted, to enable rigorous evaluation of intervention efficacy and ideal dose.

## Acknowledgements

Thank you to the Helmsley Charitable Trust for funding our work and to the Rising T1de Alliance Team for their hard work and dedication.

	Total	Completers	Non-Completers
<b>Total No</b>	46	20	26
<b>Sex</b>			
Female	17 (37%)	7 (35%)	10 (38.5%)
Male	29 (63%)	13 (65%)	16 (61.5%)
<b>Race</b>			
White	28 (60.9%)	12 (60%)	16 (61.5%)
Black	8 (17.4%)	4 (20%)	4 (15.4%)
American Indian or Alaska Native	1 (2.2%)	1 (5%)	0 (0%)
Multiracial	5 (10.9%)	2 (10%)	3 (11.5%)
Asian	1 (2.2%)	0 (0%)	1 (3.8%)
Hispanic	2 (4.3%)	0 (0%)	2 (7.7%)
Declined	1 (2.2%)	1(5%)	0 (0%)
<b>Age at T1D Onset</b>	8.3 $\pm$ 3.8	8.0 $\pm$ 4.4	8.5 $\pm$ 3.4
<b>Duration of T1D at baseline (years)</b>	3.8 (2.3-9.5)	3.4 (2.2-9.3)	4.2 (2.4-9.5)
<b>CGM Present at Baseline</b>	38 (82.6%)	17 (85%)	21 (80.1%)
<b>Pump Present at Baseline</b>	27 (58.7%)	13 (65%)	14 (53.8%)
<b>Baseline HbA1c</b>	10 (9.1-12.2)	10.2 (9.2 - 12.6)	9.8 (9.1-11.7)