

University of Colorado

# Bromocriptine Improves Central Aortic Health in Adolescents with Type 1 Diabetes Mellitus

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Table 1. Baseline characteristics of T1DM

**Participants** 

Age (years)

Diabetes duration (years)

Lean body mass (kg)

Fat free mass (kg)

**Body fat percentage** 

Lean fat percentage

reported as mean ± SD or median with

Sex (male)

HbA1c (%)

BMI (kg/m²)

Placebo N = 34

**BMI Percentile** 

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# **Background**

- The incidence rate of youth-onset type 1 diabetes mellitus (T1D) is rising annually worldwide and is projected to increase over the next 40 years.
- The presence of peripheral vascular dysfunction accelerated by large arterial stiffness, and impaired cardiac function has been clearly demonstrated in youth with T1D, predisposing young people to increase risk of cardiovascular events.
- This predisposes young patients to the life-long exposure to elevated ventricular afterload and increased incidence of cardiovascular events.
- Therapeutic strategies to mitigate vascular dysfunction are urgently needed.

# **Hypothesis**

We hypothesized that bromocriptine quick release (BCQR) therapy would improve vascular health in youth with T1D.

## Methods

- This was a placebo-controlled, random-order, double-blinded, cross-over study investigating BCQR as adjunct therapy on central aortic stiffness as measured by phase-contrast MRI.
- Participants also underwent flow mediated dilation test and brachial distensibility evaluation using tonometry.
- Adolescents with T1D were randomized 1:1 to phase-1 of 4-week BCQR (minimum dose 1.6 mg daily) or placebo therapy after which all vascular measurements were performed.
- Following a 4-week washout period, phase 2 was performed in identical fashion with the alternate treatment.

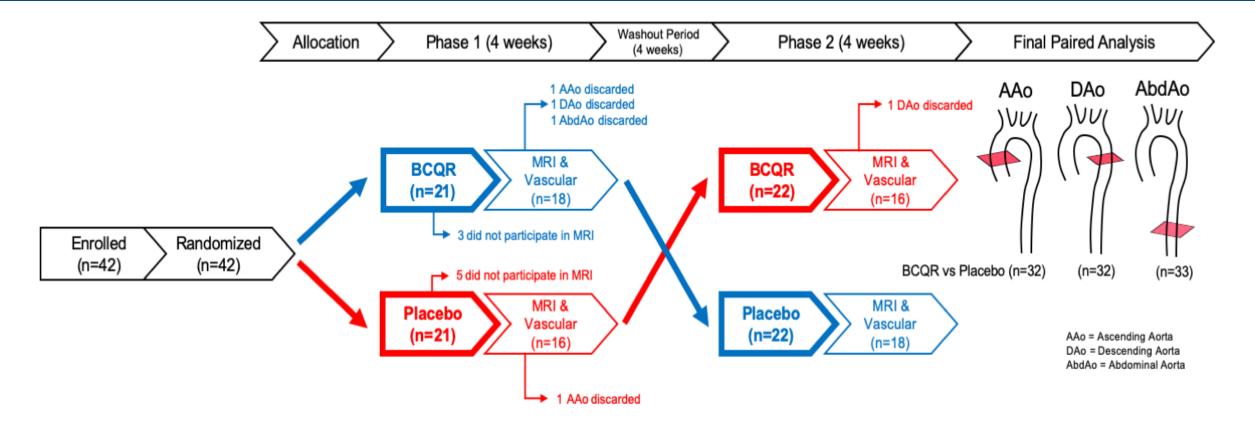


Figure 1. CONSORT (Consolidated Standards of Reporting Trials) diagram of cross-over clinical trial in adolescents with type 1 diabetes mellitus.

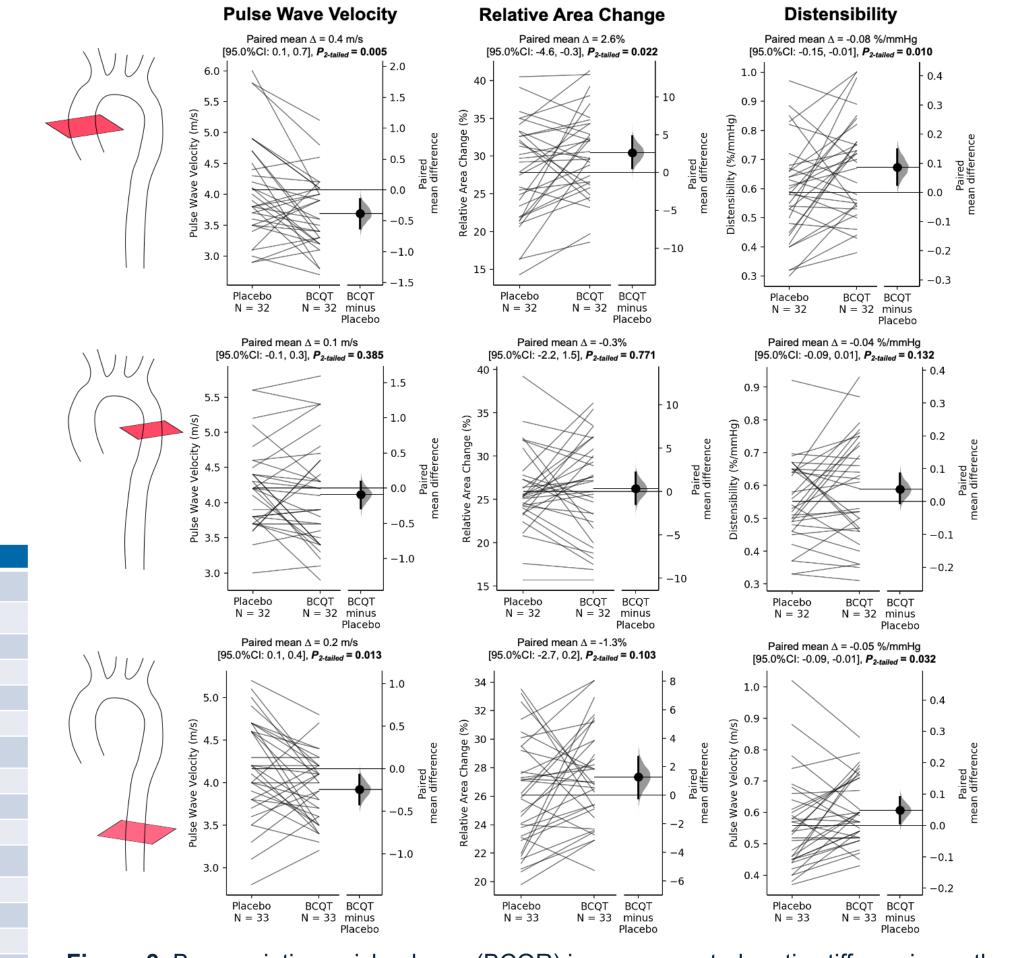
## Results

		Placebo	Bromocriptine	P-value
T1DM $(n = 34)$	Pressure Hemodynamics			
15.9 ± 2.6	Systolic blood pressure (mmHg)	118 ± 9	113 ± 11	< 0.001
13 (38.2%)	Diastolic blood pressure (mmHg)	$69 \pm 6$	66 ± 7	0.039
5.8 (3.5 - 9.5)	Pulse pressure (mmHg)	49 ± 7	46 ± 8	0.015
8.6 ± 1.1	Mean arterial pressure (mmHg)	88 ± 7	85 ± 8	0.002
24.6 ± 5.8	Ascending Aorta			
71.4 ± 26.1	Pulse Wave Velocity (m/s)	$4.0 \pm 0.8$	$3.7 \pm 0.6$	0.005
$43.8 \pm 9.4$	Relative Area Change (%)	28.1 ± 6.8	$30.7 \pm 5.7$	0.022
45.8 ± 9.8	Distensibility (%/mmHg)	$0.59 \pm 0.17$	$0.68 \pm 0.17$	0.010
32.7 ± 8.1	Decending Aorta			
	Pulse Wave Velocity (m/s)	4.2 ± 0.6	4.1 ± 0.8	0.385
$0.64 \pm 0.08$	Relative Area Change (%)	25.9 ± 4.5	26.2 ± 5.7	0.771
corresponding IQR.	Distensibility (%/mmHg)	0.55 ± 0.12	$0.59 \pm 0.16$	0.132
aired mean $\Delta$ = 2 mmHg	Abdominal Aorta			
%Cl: -4.0, 0], <b>P</b> <sub>2-tailed</sub> = <b>0.039</b>	Pulse Wave Velocity (m/s)	$4.2 \pm 0.6$	$3.9 \pm 0.4$	0.013
- 15	Relative Area Change (%)	26.1 ± 3.8	$27.3 \pm 3.4$	0.103
	Distensibility (%/mmHg)	0.56 ± 0.15	0.61 ± 0.11	0.032
10	<b>Endothelial Function</b>			
5 5	Reactive Hyperemia Index	$2.30 \pm 0.56$	$1.99 \pm 0.57$	0.006
alired differ	LN reactive hyperemia index	0.81 ± 0.23	$0.65 \pm 0.28$	0.003
Paired mean difference	Framingham RHI Risk Score	0.87 (0.55 - 1.04)	0.75 (0.25 - 0.99)	0.010
1	Peripheral Vascular Stiffness			
-10	Brachial distensibility (%/mmHg)	$0.059 \pm 0.012$	$0.061 \pm 0.013$	0.156

Data reported as mean ± SD or as median with corresponding IQR

Table 2. Arterial Hemodynamics of Participants with T1DI

Figure 2. Bromocriptine quick release (BCQR) improves overall pressure hemodynamics. BCQR reduced systolic A) and diastolic B) blood pressure as well as pulse pressure C) and mean arterial pressure D).



**Figure 3.** Bromocriptine quick release (BCQR) improves central aortic stiffness in youth with T1DM. Ascending aortic stiffness (top row) was improved by decreased pulse wave velocity (PWV), increased relative area change (RAC), and increased distensibility. There were no differences in the descending aorta (middle row). Thoraco-abdominal stiffness was improved as evidenced by decreased PWV and increased distensibility.

#### Conclusions

0.303

 $6.25 \pm 1.15$ 

- Arterial stiffness is a predictor of all cause cardiovascular events, as such, BCQR therapy might serve as a clinical intervention to reduce cardiovascular risk in youth onset-T1DM.
- Additionally, BCQR further improves standard pressure hemodynamics, also likely to improve long-term cardiovascular outcomes.
- Further study is needed to determine the significance of BCQR's ultimate impact on cardiovascular morbidity and mortality in T1DM