## SUPPLEMENTARY DATA

## Supplementary Table S1. Adverse Events by Treatment Arm within Age Cohort

Adverse Event	4 to <8 years old			8 to <12 years old			12 to <17 years	
Group	IM	2mg IN	3mg IN	IM	2mg IN	3mg IN	old IM	IN
	N=6	N=12	N=12	N=6	N=11	N=12	N=12	N=13*
One or More Events	5(83)	6(50)	5(42)	6(100)	5(46)	6(50)	7(58)	9(69)
Gastrointestinal	- ()	- ( /	- ( )		- ( - /	- ( )	- ()	- ( )
Abdominal pain	1(17)	1(8)	0	0	0	1(8)	0	0
(upper)		` /				` '		
Diarrhea	0	0	0	1(17)	0	0	0	0
Nausea	4(67)	4(33)	2(17)	3(50)	1(9)	1(8)	1(8)	3(23)
Vomiting	1(17)	1(8)	3(25)	3(50)	3(27)	4(33)	5(42)	4(31)
Total (≥1 GI	5(83)	5(42)	5(42)	5(83)	4(36)	6(50)	6(50)	6(46)
events) <sup>a</sup>								
Head discomfort								
Headache	0	2(17)	1(8)	2(33)	2(18)	4(33)	1(8)	4(31)
Nasal								
Nasal congestion	0	0	0	0	0	0	0	2(15)
Nasal discomfort	0	0	2(17)	0	0	0	0	1(8)
Sneezing	0	0	0	0	0	1(8)	0	0
Rhinalgia	0	1(8)	0	0	0	0	0	0
Total (≥1 nasal	0	1(8)	2(17)	0	0	1(8)	0	3(23)
events) <sup>a</sup>								
Ocular		0	0		0	0	0	1(0)
Eye irritation	0	0	0	0	0	0	0	1(8)
Lacrimation	0	0	0	0	1(9)	0	0	0
increased Ocular discomfort	0	0	0	0	0	0	0	1(8)
Total (≥1 ocular	0	0	0	0	1(9)	0	0	2(15)
events) <sup>a</sup>		O	O		1())	O		2(13)
Sensory/Pain								
Catheter site pain	0	1(8)	0	1(17)	0	0	0	0
Injection site	2(33)	0	0	3(50)	0	0	0	0
discomfort								
Total (≥1	2(33)	1(8)	0	3(50)	0	0	0	0
sensory/pain								
events) <sup>a</sup>								
Hypoglycemia	1/45	0	0	0	Ō	0	0	0
Hypoglycemia	1(17)	0	0	0	0	0	0	0
Cardiovascular		1(0)	0		0	0		0
Tachycardia	0	1(8)	0	0	0	0	0	0
Neurological		0	0	1/15>	0	0		0
Dizziness	0	0	0	1(17)	0	0	0	0

<sup>1</sup> serious adverse event was reported in which a 7 year old participant (intramuscular treatment) experienced a hypoglycemic event after receiving a bolus of insulin with lunch. The participant received 90 grams of oral carbohydrates and made a full recovery.

<sup>\*1</sup> participant in the 12-<17 years old had a repeat 3 mg intranasal glucagon dosing visit due to a device malfunction leading to insufficient receipt of glucagon during the initial visit; both dosing visits were included in the safety analysis

<sup>&</sup>lt;sup>a</sup> Number (%) of participants with at least 1 occurrence of the adverse event group

# SUPPLEMENTARY DATA

# Supplementary Figure S1. Intranasal Glucagon device



#### SUPPLEMENTARY DATA

## **List of Investigators**

A listing of the T1D Exchange Clinic Network sites with participating principal investigators (PI), coinvestigators (I), and coordinators (C) ordered by the number of participants recruited per site is included below:

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