GABA induces a hormonal counter-regulatory response in subjects with long-standing type 1 diabetes

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ABSTRACT

Introduction Experimentally, gamma-aminobutyric acid (GABA) has been found to exert immune-modulatory effects and induce beta-cell regeneration, which make it a highly interesting substance candidate for the treatment of type 1 diabetes (T1D). In many countries, including those in the European Union, GABA is considered a pharmaceutical drug. We have therefore conducted a safety and dose escalation trial with the first controlled-release formulation of GABA, Remygen (Diamyd Medical).

Research design and methods Six adult male subjects with long-standing T1D (age 24.8±1.5 years, disease duration 14.7±2.2 years) were enrolled in a 11-day dose escalation trial with a controlled-release formulation of GABA, Remygen. Pharmacokinetics, glucose control and hormonal counter-regulatory response during hypoglycemic clamps were evaluated at every dose increase (200 mg, 600 mg and 1200 mg).

Results During the trial there were no serious and only a few, transient, adverse events reported. Without treatment, the counter-regulatory hormone response to hypoglycemia was severely blunted. Intake of 600 mg GABA more than doubled the glucagon, epinephrine, growth hormone and cortisol responses to hypoglycemia.

Conclusions We find that the GABA treatment was well tolerated and established a counter-regulatory response to hypoglycemia in long-standing T1D. Further studies regarding not only the clinical potential of Remygen for beta-cell regeneration but also its potential use as hypoglycemic prophylaxis are warranted.

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INTRODUCTION

Gamma-aminobutyric acid (GABA) is a mediator in the central nervous system (CNS) and in peripheral tissues with effects exerted through ionotropic GABA-A and GABA-C, as well as metabotropic GABA-B receptors.1 GABA is synthesized from glutamate by glutamic acid decarboxylase, a well-known autoantigen in type 1 diabetes (T1D).2 In the adult brain, GABA is the main inhibitory neurotransmitter, while its peripheral effects vary depending on the tissue. Outside the CNS, GABA is found at the highest concentrations in beta-cells and immune cells.3 4

Most immune cells express GABA receptor subunits and the metabolic machinery required for GABA metabolism. GABA exerts a predominantly inhibitory effect on immune cells by inhibiting the production of inflammatory cytokines.5 6 GABA receptor activation also enhances regulatory T cells, contributing to its immunosuppressive functions.7 In addition to beta-cells, GABA-A receptors are also expressed in a number of peripheral tissues, including the pituitary, chromaffin cells in the adrenal medulla and hepatocytes.8 9 10 In experimental studies with rodent and human beta-cells, GABA has been shown to reverse diabetes by stimulating beta-cell regeneration, both through beta-cell proliferation and, although more controversial, transdifferentiation.11 13 The combined immune-modulatory and beta-cell effects make GABA

Significance of this study

What is already known about this subject?
▶ Gamma-aminobutyric acid (GABA) has, in experimental studies, been found to exert immune-modulatory effects and induce beta-cell regeneration, which make it a highly interesting substance candidate for the treatment of type 1 diabetes (T1D).
▶ Activation of the GABA-A receptor by benzodiazepines has been found to blunt the counter-regulatory response to hypoglycemia.

What are the new findings?
▶ GABA treatment is well tolerated in subjects with T1D.
▶ GABA can paradoxically establish a counter-regulatory response to hypoglycemia in long-standing T1D.

How might these results change the focus of research or clinical practice?
▶ GABA has a potential use as hypoglycemic prophylaxis in clinical practice in addition to its potential beta-cell regenerative effects.
a highly interesting candidate for the treatment of T1D. However, in many countries, including those in the European Union, GABA is considered a pharmaceutical drug. In this initial safety and dose escalation study, we have evaluated short-term GABA treatment in subjects with T1D as a first step toward testing the potential of GABA as a beta-cell regenerative treatment.

METHODS
Ethics and study design
GABA is considered a pharmaceutical drug within the European Union. All participants were provided oral and written information and signed a written consent prior to inclusion in the study. The study was an open-label, investigator-driven phase I clinical trial, primarily assessing the safety of a controlled-release formulation of GABA, Remygen (Diamey Medical, Stockholm, Sweden), in adult male subjects with long-standing T1D. GABA was dosed in three steps: low (200 mg), medium (600 mg) and high (1200 mg) dose, with subsequent pharmacokinetic evaluation. Remygen was administered orally as a single daily dose to be taken under fasting conditions in the morning and each dose was administered for three consecutive days. The study period consisted of visits at the hospital for 11 consecutive days. Sample size was decided after dialogue with the Swedish Medical Products Agency. The study included two mixed meal tolerance tests (MMTT), four hypoglycemic clamps and three pharmacokinetic evaluations. All study visits were conducted at Uppsala University Hospital. A detailed study design and all inclusion/exclusion criteria can be found at ClinicalTrials.gov.

Safety monitoring
All subjects underwent general and neurological examinations, as well as blood sampling for laboratory testing, at multiple time points during the safety trial. The study was monitored by Uppsala Clinical Research Center. An external Data Safety Monitoring Board reviewed the data.

Laboratory testing
Laboratory tests included metabolic parameters (glucose, hemoglobin A1c (HbA1c) and C peptide), hematology status, renal function and markers of liver injury (aspartate transaminase (AST), alanine aminotransferase (ALT), alkaline phosphate, bilirubin). During the hypoglycemic clamp, the counter-regulatory hormones glucagon, growth hormone (GH), cortisol, epinephrine and norepinephrine were analyzed. Routine blood samples were analyzed at the central laboratory of Uppsala University Hospital. Epinephrine and norepinephrine concentrations were analyzed at Karolinska University Hospital laboratory. C peptide concentrations were also analyzed with an ultrasensitive C peptide ELISA (Mercodia, Uppsala, Sweden). Glycemic control was monitored by flash glucose monitoring (FGM) (FreeStyle Libre, Abbott Laboratories, Chicago, Illinois, USA).

Pharmacokinetics of GABA
GABA concentrations in plasma were determined the first day of every dose increase at 0, 30, 60, 120, 180, 300 min and 24 hours (nadir value) after GABA administration. GABA levels were analyzed by mass spectrometry at the Mass Spectrometry Based Metabolomics Facility at Uppsala University with a method approved by the US Food and Drug Administration.

Mixed meal tolerance test
An MMTT was performed after overnight fasting at baseline and at day 11, that is, last day of treatment with 1200 mg GABA. C peptide and glucose levels were measured at 0, 15, 30, 60, 90 and 120 min after ingestion of 6 mL/kg (maximum 360 mL) Resource Protein (Nestle Health Science, Switzerland).

Hypoglycemic clamp
A hypoglycemic clamp was performed after overnight fasting at baseline and at each GABA dosage step. An individualized insulin infusion dose was calculated based on body weight (2 mIE/kg/min) and a separate glucose infusion was titrated to achieve and maintain euglycemia (5.5 mmol/L) for 30 min, followed by hypoglycemia (2.5 mmol/L) for 30 min. Blood samples were collected at the end of each plateau. GABA was administrated 30 min before start.

Endpoints
The primary outcomes of the trial were (1) number of serious and adverse events (SAE/AE) and (2) changes in laboratory parameters, physical examinations and vital signs. The secondary outcomes were (1) C peptide response during MMTT determined as area under the curve (AUC), mean and peak values; and (2) hormonal counter-regulatory response. Additional variables such as fasting plasma C peptide, HbA1c, autoantibodies and number of self-reported hypoglycemic events were included in the general assessment.

Statistical analysis
Safety laboratory tests were compared with baseline with repeated measurements one-way analysis of variance (ANOVA) with Dunnett’s post-hoc test. For FGM data a non-parametric one-way ANOVA with Dunn’s multiple comparisons test was applied. Hormonal counter-regulatory response under normoglycemic and hypoglycemic conditions was computed by multiple t-tests and corrected for multiple testing with the Holm-Sidak method. Data are presented as mean±SEM. P values <0.05 were considered statistically significant.

RESULTS
Characteristics of the subjects
Ten male subjects with T1D were screened, of whom four were excluded from participation in the study (C peptide >0.12 nmol/L (n=2), elevated AST/ALT (n=1) and difficult venous access (n=1)). Six subjects were
The patient with remaining C peptide displayed a normal hormonal counter-regulatory response to hypo-glycemia at baseline, which was maintained under the GABA treatment (data not shown). In contrast, the five patients without detectable C peptide (<0.01 nmol/L) displayed absence of counter-regulatory response at baseline (figure 1B–F). During GABA treatment (600 mg) a counter-regulatory response of glucagon, epinephrine, GH, as well as cortisol occurred in response to hypoglycemia (figure 1B–F).

DISCUSSION

Short-term oral GABA treatment was not associated with any SAEs. AEs were registered in four subjects, but they were all mild and transient, mainly neurological sensations. In previous studies, fatigue and mild weakness have also been reported.14 In another study, 3 g of GABA caused mild transaminase increases in 2% of study subjects.15 Glucose control was maintained during the study period and the frequency of hypoglycemia did not increase.

In contrast to previous studies in which activation of the GABA-A receptor by benzodiazepines has been found to blunt the counter-regulatory response,16 we paradoxically found that the hormonal counter-regulatory response to hypoglycemia was enhanced by GABA treatment. A tentative explanation for this difference in results is that GABA per se has limited capacity to cross the blood–brain barrier, while benzodiazepines easily cross to exert their effects in the CNS. The mechanism for the counter-regulatory failure induced by benzodiazepines has previously been shown to be lactate release in the ventromedial hypothalamus.17 Also, the metabolism of GABA and benzodiazepines differs, which could in turn impact the counter-regulatory response. The present study shows that if the peripheral effects of GABA are isolated, the hormonal counter-regulatory response instead become enhanced by yet unknown mechanisms. A possible explanation could be the activation of GABA-A receptors in other peripheral organs/tissues of importance for the hormonal counter-regulatory response. For instance, GABA-A receptor activation in chromaffin cells within the adrenal glands induces the secretion of catecholamines.9 The study period and sample size (number of subjects included) were, however, too small to assess potential clinical benefits in hypoglycemia severity and frequency.
We conclude that the controlled-release formulation of GABA (Remygen) can be considered a safe treatment in T1D and that a larger, long-term clinical trial is warranted in order to further study the clinical potential of GABA as a drug for beta-cell regeneration and, in view of the findings in this study, hypoglycemia protection.

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Contributors P-OC and DE designed the study. HL, DE, HH, AE, JC-C (until February 28, 2019) and P-OC performed the clinical assessments in the study. DE, HH and P-OC researched the data, and DE, HL, and P-OC wrote the manuscript. All authors critically reviewed the manuscript. P-OC is the guarantor of this work.

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Competing interests DE and P-OC are listed as coinventors for a patent application submitted by Diamyd Medical, but do not have any financial interests in Diamyd Medical. None of the other authors have any conflicts of interest to declare.

Patient consent for publication Not required.

Ethics approval The study was approved by both the Uppsala County Ethics Board (Dnr 2018/200/1) and the Swedish Medical Products Agency. The reported investigations were carried out in accordance with the principles of the Declaration of Helsinki as revised in 2013.

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Data availability statement Data are available upon reasonable request. The data sets generated during the current study can be made available from the corresponding author upon reasonable request.

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Figure 1 GABA treatment improves the counter-regulatory hormone response to hypoglycemia in type 1 diabetes. (A) Flash glucose monitoring for all subjects (n=6) at baseline and during short-term oral treatment with GABA in increasing doses. (B–F) Hormonal counter-regulatory response during hyperinsulimemic hypoglycemic clamps. Glucose levels were regulated by an intravenous continuous rate of insulin and a variable rate of glucose in order to reach first a 30 min plateau with glucose levels of 5.5 mmol/L, followed by a 30 min plateau at 2.5 mmol/L. Blood samples were collected at the end of each plateau. The analysis contains data from the five subjects without detectable C peptide (<0.01 nmol/L) in which the counter-regulatory response was blunted at baseline. Black bars represent normoglycemia (5.5 mmol/L) and gray bars hypoglycemic conditions (2.5 mmol/L). Data are presented as means±SEM. *P<0.05 and **P<0.01 compared with the normoglycemic levels at the respective occasion. GABA, gamma-aminobutyric acid.
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REFERENCES

1. Watanabe M, Maemura K, Kanbara K, et al. Gaba and gaba receptors in the central nervous system and other organs. *Int Rev Cytol* 2002;213:1–47.
2. Baekkeskov S, Aanstoot HJ, Christgau S, et al. Identification of the 64K autoantigen in insulin-dependent diabetes as the gaba-synthesizing enzyme glutamic acid decarboxylase. *Nature* 1990;347:151–6.
3. Taniguchi H, Okada Y, Seguchi H, et al. High concentration of gamma-aminobutyric acid in pancreatic beta cells. *Diabetes* 1979;28:629–33.
4. Jin Z, Mendu SK, Bir nir B. Gaba is an effective immunomodulatory molecule. *Amino Acids* 2013;45:87–94.
5. Bergeret M, Khrestchatisky M, Tremblay E, et al. Gaba modulates cytotoxicity of immunocompetent cells expressing gabaA receptor subunits. *Biomed Pharmacother* 1998;52:214–9.
6. Tian J, Dang H, W allner M, et al. Homotaurine, a safe blood-brain barrier permeable GABA(A)-specific agonist, ameliorates disease in mouse models of multiple sclerosis. *Sci Rep* 2018;8:16555.
7. Hanssen SL, Fjalland B, Jackson MB. Modulation of gabaA receptors and neuropeptide secretion by the neurosteroid allopregnanolone in posterior and intermediate pituitary. *Pharmacol Toxicol* 2003;93:91–7.
8. Harada K, Matsuoka H, Fujihara H, et al. Gaba signaling and neuroactive steroids in adrenal medullary chromaffin cells. *Front Cell Neurosci* 2016;10:100.
9. Erlitzki R, Gong Y, Zhang M, et al. Identification of gamma-aminobutyric acid receptor subunit types in human and rat liver. *Am J Physiol Gastrointest Liver Physiol* 2000;279:G733–9.
10. Tian J, Dang H, Chen Z, et al. γ-Aminobutyric acid regulates both the survival and replication of human β-cells. *Diabetes* 2013;62:3760–6.
11. Untereiner A, Xu J, Bhattacharjee A, et al. γ-aminobutyric acid stimulates β-cell proliferation through the mTORC1/p70S6K pathway, an effect amplified by Ly49, a novel γ-aminobutyric acid type A receptor positive allosteric modulator. *Diabetes Obes Metab* 2020;22:2021–31.
12. Ben-Othman N, Vieira A, Courtney M, et al. Long-term gaba administration induces alpha cell-mediated beta-like cell neogenesis. *Cell* 2017;168:e11:73–85.
13. Cavagnini F, Invitti C, Pinto M, et al. Effect of acute and repeated administration of gamma aminobutyric acid (gaba) on growth hormone and prolactin secretion in man. *Acta Endocrinol* 1980;93:149–54.
14. Otomo E, Araki G, Mori A, et al. Clinical evaluation of gaba in the treatment of cerebrovascular disorders. multi-center double-blind study in comparison with pyrithioxine and placebo. *Arzneimittelforschung* 1981;31:1511–23.
15. Hedrington MS, Mikeladze M, Tate DB, et al. Effects of γ-aminobutyric acid a receptor activation on counterregulatory responses to subsequent exercise in individuals with type 1 diabetes. *Diabetes* 2016;65:2754–9.
16. Chan O, Paranjape SA, Horblitt A, et al. Lactate-induced release of gaba in the ventromedial hypothalamus contributes to counterregulatory failure in recurrent hypoglycemia and diabetes. *Diabetes* 2013;62:4239–46.