Disruption of glucagon receptor signaling causes hyperaminoacidemia exposing a possible liver-alpha-cell axis

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INTRODUCTION

Glucagon is a peptide hormone of 29 amino acids processed from the prohormone, proglucagon, by prohormone convertase 2 (PC-2) in pancreatic alpha-cells (39). Activation of the hepatic glucagon receptor (GR) increases hepatic glycogenolysis and gluconeogenesis (10), and the physiological role of glucagon has been coupled to glucose metabolism with opposing actions to insulin (1, 28, 43). However, several studies have suggested that glucagon may play an equally important role in the regulation of hepatic acid metabolism (2, 5, 7, 13, 21, 41). Increased fasting and postprandial plasma concentrations of glucagon have been reported in clinical conditions including nonalcoholic fatty liver disease (NAFLD) and type 2 diabetes (33) raising the question whether this is related to amino acid metabolism.

Glucagon is secreted from the pancreatic alpha-cells is essential for regulation of blood glucose levels. However, glucagon may play an equally important role in the regulation of amino acid metabolism by promoting ureagenesis. We hypothesized that disruption of glucagon receptor signaling would lead to an increased plasma concentration of amino acids, which in a feedback manner stimulates the secretion of glucagon, eventually associated with compensatory proliferation of the pancreatic alpha-cells. To address this, we performed plasma profiling of glucagon receptor knockout (Gcgr−/−) mice and wild-type (WT) littermates using liquid chromatography-mass spectrometry (LC-MS)-based metabolomics, and tissue biopsies from the pancreas were analyzed for islet hormones and by histology. A principal component analysis of the plasma metabolome from Gcgr−/− and WT littermates indicated amino acids as the primary metabolic component distinguishing the two groups of mice. Apart from their hyperaminoacidemia, Gcgr−/− mice display hyperglucagonemia, increased pancreatic content of glucagon and somatostatin (but not insulin), and alpha-cell hyperplasia and hypertrophy compared with WT littermates. Incubating cultured α-TC1.9 cells with a mixture of amino acids (Vamin 1%) for 30 min and for up to 48 h led to increased glucagon concentrations (~6-fold) in the media and cell proliferation (~2-fold), respectively. In anesthetized mice, a glucagon receptor-specific antagonist (Novo Nordisk 25–2648, 100 mg/kg) reduced amino acid clearance. Our data support the notion that glucagon secretion and hepatic amino acid metabolism are linked in a close feedback loop, which operates independently of normal variations in glucose metabolism.

alpha-cell; amino acids; glucagon; glucagon receptor; hyperglucagonemia
of GR disruption. In addition, we investigated the hepatic clearance of amino acids in vivo after prior pharmacological or genetically induced GR blockage. Finally, amino acids were administered to the alpha-cell line aTC1.9 to monitor glucagon secretion and alpha-cell proliferation.

MATERIALS AND METHODS

Animal studies. Animal studies were conducted with permission from the Danish Animal Experiments Inspectorate, Ministry of Environment and Food of Denmark, permit 2013-15-2934-00833, and in accordance with the EU Directive 2010/63/EU and guidelines of Danish legislation governing animal experimentation (1987), and the National Institutes of Health (Publication No. 85-23). All studies were approved by the local ethical committee.

Female C57BL/6JRj mice (12 wk of age) were obtained from Janvier Laboratories, Saint-Berthevin Cedex, France. Mice were housed in groups of six to eight in individually ventilated cages and followed a light cycle of 12 h (lights on 6 AM to 6 PM) with ad libitum access to standard chow (catalog no. 1319, Altromin Spezialfutter, Lage, Germany) and water. Glucagon receptor knockout (Gcgr<sup>-/-</sup>) mice C57BL/6<sup>jm</sup>-Gcgr<sup>-/-</sup> were previously described (14). Male and female homozygotes and wild-type (WT) littermates, age 9–29 wk, were used. During all Gcgr<sup>-/-</sup> and littermate studies, the investigator was blinded to the genotype of the mice to avoid bias.

Biochemical and morphometric characterization of Gcgr<sup>-/-</sup> mice. Eleven Gcgr<sup>-/-</sup> mice (6 females 18–28 wk of age and 5 males 15–25 wk of age) and 15 WT littermates (8 females 18–25 wk of age and 7 males 15–22 wk of age) were fasted for 4 h with free access to water. A 50-μl blood sample was collected from the tail vein, and subsequently the mice were anesthetized with isoflurane (Baxter, Søborg, Denmark). When the mice were sufficiently sedated (absence of reflexes) the abdominal cavity was opened with a midline incision. Tissue samples from the pancreas and liver were harvested and either fixed in formaldehyde 4% and methanol 1–2% (Hounisen Laboratory Equipment, Skanderborg, Denmark) or snap-frozen on dry ice. Finally, the mice were subjected to a total blood collection from the inferior vena cava. The blood was immediately transferred to prechilled EDTA-coated Eppendorf tubes and stored on ice until centrifuged (1,650 × g, 15 min, 4°C). Plasma was collected and stored in prechilled EDTA-coated Eppendorf tubes and stored on ice until centrifuged (1,650 × g, 15 min, 4°C). Plasma samples from 11 Gcgr<sup>-/-</sup> littermates (6 females 18–28 wk of age and 5 males 15–25 wk of age) were fasted and anesthetized as above and the inferior caval vein exposed. At time 0 min the mice received an injection of 1 μmol/g body wt. A control group received 100 μl vehicle. At time 0 min, 1 μmol/g body wt Vamin diluted in sterilized PBS (100 μl) was injected into the caval vein. At time 12 min, the mice were subjected to total blood collection from the inferior vena cava. Blood samples were handled as described above and analyzed for total L-amino acid, insulin, and glucagon concentrations. Clearance was defined as the incremental area under the curve (AUC<sub>0–20min</sub>).

Amino acid clearance after ligation of the kidneys. Female C57BL/6JRj mice (12 wk of age) were anesthetized with isoflurane and subsequently subjected to kidney ligation (n = 4); four mice were sham operated and served as controls. At time 0 min, 1 μmol/g body wt Vamin diluted in sterilized PBS (100 μl) was injected into the caval vein. At time 12 min, the mice were subjected to total blood collection from the inferior vena cava. Blood samples were handled as described above and analyzed for total L-amino acid concentrations.

Biochemical analysis. Plasma concentrations of total L-amino acids were quantified using an enzyme-linked immunosorbent assay (ELISA) (catalog no. ab65347; Abcam, Cambridge, UK). This kit determines concentrations of free L-amino acids, but neither protein bound nor D-amino acids. The assay was evaluated by recovery experiments using pooled (n = 4) mouse plasma with added known amounts of amino acids (product no. A6282; Sigma-Aldrich, Copenhagen, Denmark). Recoveries of amino acids were on average 79 ± 9% in mouse plasma.

Plasma concentrations of glucagon were measured using a validated (52) two-site enzyme immunoassay (catalog no. 10-1281-01; Merckodia, Uppsala, Sweden) according to the manufacturer’s protocol.

Plasma concentrations of ammonia/ammonium and of bile acids were quantified using enzymatic assays (catalog no. ab93360; Abcam; and catalog no. STA-361; Cell Biosats, San Diego, CA, respectively).

Plasma concentrations of corticosterone and insulin were quantified using ELISAs (catalog no. ADI-900–097; Enzo, AH Diagnostics, Aarhus, Denmark; and catalog no. 10-1247-10, Merckodia, Uppsala, Sweden, respectively).

To avoid bias when performing biochemical analysis, all samples were assigned a number, so that genotype and treatment were unknown at the time of analysis.

Plasma metabolomics. Metabolomic analysis was performed on plasma samples from 11 Gcgr<sup>-/-</sup> (6 females 18–28 wk of age and 5 males 15–25 wk of age), and 11 WT littermates (6 females 18–25 wk of age and 5 males 18–22 wk of age), also used for biochemical and morphometric characterization.

Using liquid chromatographic-triple quadrupole mass spectrometric (LC-MS/MS) measurements and the AbsoluteIDQ p180 Kit (BIO-

Table 1. The components and concentrations of Vamin 14 electrolyte-free

<table>
<thead>
<tr>
<th>Amino Acid</th>
<th>Concentration, g/l</th>
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<tbody>
<tr>
<td>Alanine</td>
<td>12.0</td>
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<tr>
<td>Aspartic acid</td>
<td>2.5</td>
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<tr>
<td>Glutamic acid</td>
<td>4.2</td>
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<tr>
<td>Glycine</td>
<td>5.9</td>
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<tr>
<td>Proline</td>
<td>5.1</td>
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<tr>
<td>Serine</td>
<td>3.4</td>
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<tr>
<td>Tryptophan</td>
<td>1.4</td>
</tr>
<tr>
<td>Arginine</td>
<td>8.4</td>
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<tr>
<td>Cysteine</td>
<td>0.4</td>
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<tr>
<td>Histidine</td>
<td>5.1</td>
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<tr>
<td>Threonine</td>
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<tr>
<td>Methionine</td>
<td>4.9</td>
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<tr>
<td>Phenylalanine</td>
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<td>Valine</td>
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<td>Lysine</td>
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<td>Leucine</td>
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<td>Glutamic acid</td>
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<td>Glycine</td>
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CRATES Life Sciences, Innsbruck, Austria). 188 metabolites were quantified out of 10 μl plasma, including free carnitine, 39 acylcarnitines, 21 amino acids, 21 biogenic amines, hexoses, 90 glycerophospholipids (14 lysophosphatidylcholines and 76 phosphatidylcholines), and 15 sphingolipids. The assay procedures of the AbsoluteIDQ p180 Kit as well as the metabolite nomenclature have been described in detail previously (40, 54). Sample handling was performed by a Hamilton Microlab STAR robot (Hamilton Bonaduz, Bonaduz, Switzerland) and a Ultravap nitrogen evaporator (Povair Sciences, Leatherhead, UK), besides standard laboratory equipment. Mass spectrometric analyses were done on an API 4000 triple quadrupole system (Sciex Deutschland, Darmstadt, Germany) equipped with a 1200 Series HPLC (Agilent Technologies Deutschland, Böblingen, Germany) and a HYC PAL auto sampler (CTC Analytics, Swissen, Switzerland) controlled by the software Analyst 1.6.1. Data evaluation for quantification of metabolite concentrations and quality assessment were performed with the software MultiQuant (Sciex) and the MetaIDO software package, which is an integral part of the AbsoluteIDQ Kit. Metabolite concentrations were calculated using internal standards and reported in micromoles per liter.

All data have been deposited at Figshare at doi:10.6084/m9.figshare.5364082.v1

Histology and immunohistochemistry. Pancreas and liver tissue samples from the 10 Geck−/− mice (5 females 18–25 wk of age and 5 males 15–25 wk of age), and 11 WT littermates (6 females 18–25 wk of age and 5 males 18–22 wk of age), also used for plasma metabolomics analysis, were fixed for 24 h, and then transferred to 70% ethanol. The tissue samples were embedded in paraffin at the Finsen Laboratory (Rigshospitalet, Copenhagen Biocenter, Copenhagen, Denmark).

Tissue sections were stained for insulin (in-house-developed guinea pig anti-insulin antibody 2006, 1:10,000) and glucagon (in-house-developed rabbit anti-glucagon 4304, 1:2,000). For antigen retrieval, sections were pretreated by boiling in triethylene glycol (TEG) buffer, pH 9, for 15 min. The sections were then incubated overnight with either glucagon, insulin, or somatostatin antibody and after washing in PBS buffer, pH 7.4, subsequently incubated with a mixture of Alexa568-labeled donkey-anti-rabbit antibody (red, 1:500, Abcam) and Alexa488-labeled goat anti-guinea pig antibody (green, 1:500, Life Technologies, Carlsbad, CA). The slides were mounted with DAKO fluorescence mounting medium (Agilent, Santa Clara, CA) and examined using an Axioscope 2 plus microscope (Zeiss, Jena, Germany). Images were taken using a CoolSNAP camera (Photometrics, Tucson, AZ).

For composition and size, 30 islets from each mouse (2 to 3 sections separated by 200–300 μm evaluated per mouse) were photographed and the relevant areas (insulin, glucagon, and total islet area) were measured using Image-Pro 7 software (Media Cybernetics, Rockville MD) as previously described (31). To measure the mean size of alpha-cells and beta-cells in the two groups of mice, 20 glucagon positive and 20 insulin positive cells were measured for each mouse using Image-Pro 7 software.

The pancreas sections were also double-stained for the glucagon-like-peptide-1 receptor (GLP-1R) (mouse GLP-1 receptor antibody 7F38, 1:200, generous gift from Charles Pyke, Novo Nordisk, Målev, Denmark) and insulin, glucagon, and somatostatin (in-house-developed rabbit somatostatin antibody 1759, 1:4,000), respectively. The sections were pretreated with pronase for 10 min (0.1%, Roche, Switzerland) at 37°C and incubated overnight with the GLP-1R antibody, and after washing in PBS buffer, pH 7.4, subsequently incubated with biotin-anti-mouse antibody, 1:200, Vectastain-complex according to the manufacturer’s instruction (Vector, Burlingame, CA), stained with DAB and counterstained lightly with hematoxylin. After this staining was completed, the sections were incubated overnight with either glucagon, insulin, or somatostatin antibody and after a wash in PBS, incubated with either Alexa 568 (1:500), or Alexa 488. The slides were then mounted with DAKO fluorescence mounting medium, the sections were examined using Axioscope 2 microscope, and images were taken using the CoolSNAP camera.

Pancreatic protein extraction and measurements of pancreatic concentrations of glucagon, GLP-1, insulin, and somatostatin. Snap-frozen pancreatic tissues from six WT mice (2 females 22 and 25 wk of age and 4 males 15–20 wk of age), and five Geck−/− mice (3 females 22–25 wk of age and 2 males 22 and 25 wk of age) were subject to peptide extraction carried out as described previously (51). The dipeptidyl peptidase-4 (DPP-4) inhibitor, valine-pyrrolidide, was added to all samples and standards (final concentration 0.01 mmol/l) to prevent NH2-terminal degradation of GLP-1 during the assay incubation. Total amidated GLP-1 concentrations (the sum of 1–36 NH2, 7–36NH2, and 9–36NH2) were quantified using an RIA (code name 89390) specific for the amidated COOH-terminal of the GLP-1 molecule (35). Active GLP-1 concentrations (7–36 NH2) were measured using well-established immunoassay specific for the NH2-terminus (47). Glucagon concentrations were measured using an RIA (code name 4305) (18) validated by ELISA (48). Insulin concentrations were measured using an RIA (code name 2068–3 (34). Somatostatin concentrations were measured using an RIA (code name 1758–5) (4).

Stimulation of the alpha-cell line, alphaTC1.9, with amino acids. The alpha-cell line, alphaTC1 Clone 9 (ATCC CRL-2350), was used (16). The cells were seeded in 24-well plates (Nunc, ThermoScientific) at a cell density of 4 × 10^4 per well. Cells 80% confluent were incubated for 2 h with PBS (controls) or 1% Vamin added to the media. The cell media were subsequently harvested and centrifuged (1,500 g, 15 min, 4°C) to remove any cells or debris and kept at −80°C until analysis. For long-term incubation (48 h) cells were treated with Vamin or PBS as described above and also treated with 5-bromo-2-deoxyuridine (BrdU) to assess cell proliferation (catalog no. C10337; Invitrogen) according to the manufacturer’s protocol. Concentrations of glucagon were normalized to total protein content assessed with a BCA kit from Thermo Fisher Scientific (catalog no. 23225).

Statistics. All bioinformatics analyses were done with the Perseus software of the MaxQuant computational platform. For the principal component analysis we included the data set derived from mass spectrometry and, in addition, all available biochemical data obtained in the current study. A false discovery rate of < 0.05 after Benjamini-Hochberg correction was used to correct for multiple testing. When more than two groups were compared, a one-way ANOVA corrected for multiple testing using the Sidak-Holm algorithm was applied. To analyze data from two independent groups unpaired t-tests were used. Calculations were made using GraphPad Prism (version 7.02 for Windows; GraphPad Software, La Jolla, CA) and STAT14 (SE) (StataCorp, College Station, TX). All data are presented as means ± SE unless otherwise stated.

RESULTS

Biochemical and morphometric characterization of Geck−/− and WT mice. The body weight of Geck−/− mice did not differ from WT littermates (31 ± 6 vs. 30 ± 4 g, P = 0.6) (Fig. 1A).

Blood glucose concentrations were lower in Geck−/− mice compared with WT littermates (7 ± 1 vs. 9 ± 0.8 mmol/l, P < 0.0001) (Fig. 1B). Plasma glucagon concentrations were increased ~20-fold in Geck−/− mice compared with WT littermates (378 ± 158 vs. 21 ± 15 pmol/l, P < 0.0001) (Fig. 1C).

Plasma concentrations of total t-amino acids were ~3-fold increased in Geck−/− mice compared with WT littermates (7.4 ± 1.4 vs. 2.5 ± 0.6 mmol/l, P < 0.0001) (Fig. 1D). Ammonia/ammonium concentrations were increased ~1.5 fold in Geck−/− mice compared with WT littermates (27 ± 4 vs. 18 ± 4 μmol/l, P = 0.0002) (Fig. 1E). Plasma concentrations of corticosterone and bile acids did not differ between Geck−/− mice and WT mice.
Fig. 1. Biochemical characterization of glucagon receptor knockout mice and wild-type littermates. A: the body weight of glucagon receptor knockout (Gcgr\(^{-/-}\)) male (gray empty circles) or female mice (gray full circles) did not differ significantly, \(P = 0.4\), from wild-type (WT) male mice (black empty squares) or female littermates (black full squares). The body weight of female WT differed from WT males (25 ± 3 vs. 33 ± 5 g, \(P = 0.0002\)). B: blood glucose concentrations in Gcgr\(^{-/-}\) mice were significantly lower compared with WT, \(P < 0.0001\), and did not differ between males and females. C: plasma concentrations of glucagon were significantly increased in Gcgr\(^{-/-}\) mice compared with WT, \(P < 0.0001\), and did not differ between males and females. D: plasma amino acid concentrations were significantly increased in Gcgr\(^{-/-}\) mice compared with WT, \(P < 0.0001\), and did not differ between males and females. E: plasma ammonia/ammonium concentrations were significantly increased in Gcgr\(^{-/-}\) mice compared with WT, \(P = 0.0002\), and did not differ between males and females. F: plasma concentrations of corticosterone did not differ between Gcgr\(^{-/-}\) mice and WT, \(P = 0.8\). Plasma corticosterone concentrations of Gcgr\(^{-/-}\) females did not differ from Gcgr\(^{-/-}\) males, \(P = 0.2\). Plasma corticosterone concentrations of WT females differed from WT males (251 ± 72 vs. 85 ± 25 ng/ml, \(P < 0.0001\)). G: plasma concentrations of bile acids did not differ between Gcgr\(^{-/-}\) mice and WT, \(P = 0.3\), and did not differ between males and females. Gcgr\(^{-/-}\) mice, \(n = 11\) (15–28 wk of age); WT littermates, \(n = 15\) (15–25 wk of age). Data are presented as means ± SD. ****\(P < 0.0001\).

mice and WT littermates (corticosterone: 176 ± 64 vs. 173 ± 100 ng/ml, \(P = 0.9\); Fig. 1F) (bile acids: 6 ± 4 vs. 4 ± 0.6 µmol/l, \(P = 0.3\)), but the concentration of bile acids showed greater variation in Gcgr\(^{-/-}\) (Fig. 1G).

Plasma metabolomics analysis. A principal component analysis identified pooled concentrations of amino acids as the primary component separating Gcgr\(^{-/-}\) mice from WT littermates (Fig. 2A) and was, furthermore, in an additional set of analysis, the predictor with most effect (~8-fold) as well as the most significant predictor (\(P = 7.7 \times 10^{-9}\)) across the 188 identified metabolites (Fig. 2B). Alanine (Ala), glutamine (Gln), glycine (Gly), threonine (Thr), and serine (Ser) were, listed in decreasing order, the amino acids found in the highest concentration in Gcgr\(^{-/-}\) mice (Fig. 2C).

Gcgr\(^{-/-}\) mice show alpha-cell hyperplasia and hypertrophy. The mean islet size (area) was significantly larger in the Gcgr\(^{-/-}\) mice compared with WT littermates (43.206 ± 2.054 vs. 1.1939 ± 697 µm\(^2\), \(P < 0.0001\)). In the WT mice, the mean
beta-cell area was 5,555 ± 423 μm², amounting to a mean 46% of the mean total islet area, and the mean beta-cell area in the Gcgr⁻/⁻ mice was 7,047 ± 465 μm², amounting to a mean 16% of the total islet area. The percent beta-cell areas was thus smaller in the Gcgr⁻/⁻ mice (P < 0.05). In the WT mice, the mean alpha-cell area was 1,150 ± 70 μm², amounting to a mean 10% of the mean total islet area. The mean alpha-cell area in the Gcgr⁻/⁻ mice was 23,013 ± 1,207 μm², corresponding to a mean 53% of the total islet area. The alpha-cell area in the WT mice was significantly smaller than in the Gcgr⁻/⁻ mice (P < 0.0001). Representative double-stained (insulin and glucagon) pancreatic islets from Gcgr⁻/⁻ mice and WT littermates are shown in Fig. 3, A and D. The individual beta-cell size was slightly larger in the Gcgr⁻/⁻ mice than in the WT littermates (142 ± 0.2 vs. 102 ± 0.1 μm², P < 0.001) whereas the mean individual alpha-cell size was much larger in the Gcgr⁻/⁻ mice than in the WT littermates (240 ± 0.3 vs. 87 ± 0.1 μm², P < 0.0001).

Using a specific antibody against the murine GLP-1 receptor (37), the GLP-1 receptor was identified in pancreatic islets of both WT and Gcgr⁻/⁻ mice. Staining for the GLP-1 receptor, combined with insulin, glucagon, and somatostatin immunohistochemistry, revealed that the GLP-1 receptor was localized to the beta-cells in both groups of mice; however, GLP-1 receptor staining of the somatostatin- and glucagon-producing cells could not be excluded. The GLP-1 receptor was confined to the plasma membrane of insulin positive cells in WT mice whereas in Gcgr⁻/⁻ mice both plasma membrane and cytosolic staining was detected (Fig. 4).

Pancreatic concentrations of glucagon, GLP-1, insulin, and somatostatin in Gcgr⁻/⁻ mice and WT littermates. Pancreatic concentrations of extractable glucagon were higher in Gcgr⁻/⁻ mice compared with WT littermates (6,851 ± 2,361 vs. 366 ± 285 pmol/g, P = 0.0003). Similarly, pancreatic concentrations of extractable, amidated total GLP-1 were significantly higher in Gcgr⁻/⁻ mice compared with WT littermates.
Amino acids stimulate glucagon secretion and proliferation of alphaTC1.9 cells. Incubation with amino acids stimulated secretion of glucagon 6-fold from the alphaTC1.9 cells compared with PBS stimulated controls (59 ± 6 vs. 9 ± 8 pmol/g, P = 0.0006) (Fig. 5A). A prolonged amino acid stimulation of the alphaTC1.9 cells led to an increased proliferation by 2-fold compared with proliferation of cells not exposed to amino acids (31 ± 8% vs. 17 ± 4% BrdU positive cells, P = 0.01) (Fig. 5B).

Genetic disruption of glucagon receptor signaling influence on amino acid clearance. Gcgr<sup>−/−</sup> mice and WT littermates received identical amino acid loads (1 μmol/g) at time 0 min. At time 12 min, the plasma Δ amino acid concentration in WT littermates reached 0.8 ± 0.5 mmol/l, whereas the Gcgr<sup>−/−</sup> mice had a Δ concentration of 6 ± 1 mmol/l (data not shown).

Effects of pharmacological blockade of glucagon receptor signaling on amino acid clearance in mice. In mice treated with a glucagon receptor antagonist (GRA), fasting plasma concentrations of amino acids tended to be higher than in vehicle-treated mice (3.7 ± 0.3 vs. 3.0 ± 0.3 mmol/l, P = 0.1). In GRA + Vamin-treated mice, plasma concentrations of amino acids, at time 12 min, were significantly higher compared with vehicle + Vamin-treated mice (9 ± 1 vs. 5 ± 1 mmol/l, P = 0.006). In the latter group, concentrations returned to near baseline (4.8 ± 0.6 mmol/l) 20 min after the Vamin injection whereas concentrations in GRA + Vamin-treated mice remained elevated (7 ± 2 mmol/l) (Fig. 6A). The incremental area under the curve (iAUC<sub>0–20 min</sub>) was significantly higher in GRA + Vamin-treated mice compared with vehicle + Vamin-treated mice (P = 0.006) (Fig. 6B).

Effect of pharmacological blockade of glucagon receptor signaling on insulin secretion in mice. In GRA-treated mice, fasting plasma concentrations of glucagon were significantly higher than in vehicle-treated mice (11 ± 1 vs. 7 ± 2 pmol/l, P = 0.02). Glucagon concentrations increased in Vamin-treated groups (vehicle + Vamin and GRA + Vamin). The glucagon response at time 2 min tended to be larger (~2-fold) in GRA-treated mice compared with vehicle-treated mice (39 ± 11 vs. 17 ± 4 pmol/l, P = 0.1), (Fig. 6C) and the glucagon secretory response was prolonged in GRA-treated mice compared with vehicle-treated mice (Fig. 6D). The iAUC<sub>0–20 min</sub> of GRA + Vamin-treated mice was 4-fold larger when compared with vehicle + Vamin-treated mice (82 min × pmol/L vs. 19 min × pmol/L, P = 0.02) (Fig. 6D).

Effect of pharmacological blockade of glucagon receptor signaling on insulin secretion in mice. Fasting plasma concentrations of insulin did not differ from GRA- and vehicle-treated mice (0.9 ± 0.2 vs. 0.6 ± 0.5 ng/ml, P = 0.3). Plasma concentrations of insulin increased in response to the amino acid stimulation in both groups. GRA-treated mice had a smaller response in insulin secretion compared with vehicle-treated mice, although the difference was not significant (13 ± 3 vs. 19 ± 5 pmol/l, P = 0.3) at time 2 min (Fig. 7A). The iAUC<sub>0–20 min</sub> was smaller (~20 min × ng/ml, P = 0.1) in the GRA-treated group compared with the vehicle-treated group after the amino acid stimulation (Fig. 7B).
concentrations of amino acids 12 min after Vamin injection did not differ significantly between mice subjected to ligation of both kidneys and control mice (4.4 ± 0.6 vs. 3.7 ± 0.3 mmol/l, \( P = 0.4 \)) (data not shown).

**DISCUSSION**

Here we demonstrate that normal clearance of amino acids in mice requires glucagon receptor signaling, and that disruption of the latter, by a glucagon receptor antagonist (GRA, 25–2648) or genetic deletion of the glucagon receptor (Gcgr\(^{-/-}\)), results in significantly higher plasma concentrations of amino acids (hyperaminoacidemia) possibly due to decreased hepatic ureagenesis.

First, we recharacterized the Gcgr\(^{-/-}\) mouse with a focus on amino acid metabolism using plasma metabolomics, histology, and measurements of pancreatic peptide hormones. As previously reported (11, 14), Gcgr\(^{-/-}\) mice had lower blood glucose concentrations, elevated plasma concentrations of glucagon, increased alpha-cell mass, increased pancreatic content of glucagon, total and active GLP-1, somatostatin, and an equal content of insulin compared with WT littermates. Importantly, we found that Gcgr\(^{-/-}\) mice had pronounced increases in plasma amino acid and ammonia concentrations. In line with this, a metabolomics-driven principal component analysis showed that among 188 metabolites measured after glucagon receptor signaling disruption, the amino acid concentrations showed the most dramatic changes (~41%). In particular, the glucogenic amino acids showed the largest changes. We were, in contrast to what has been reported previously (53), unable to detect significantly higher plasma concentrations of bile acids in Gcgr\(^{-/-}\) mice compared with WT littermates, but we did observe a greater variability in the Gcgr\(^{-/-}\) mice and bile acids were indeed higher in some animals. In this report, corticosterone plasma concentrations were found to be similar in Gcgr\(^{-/-}\) mice and WT mice after a short-term fast (4 h), and others (11, 14) have reported similar corticosterone concentrations under similar conditions. However, the corticosterone levels of Gcgr\(^{-/-}\) mice were found to be increased twofold compared with WT mice upon a prolonged fast (>12 h), suggesting that increased levels of corticosterone may function to prevent hypoglycemia during prolonged fasting, but are less important in the fed state.

Remodeling of the pancreatic islets has been suggested to be of importance for maintaining adequate metabolism and in particular normal glucose concentrations (6).

We found the average area of the pancreatic islets to be significantly larger in the Gcgr\(^{-/-}\) mice than in the WT littermates, in accordance with previous studies (8, 14). The larger alpha-cell area was due to both hyperplasia and hypertrophy, as the individual alpha-cells in the Gcgr\(^{-/-}\) mice were significantly larger than in WT littermates, and there were also more alpha cells. In the pancreatic alpha-cells, proglucagon is primarily processed to glucagon by prohormone convertase (PC) 2 (39), and very little, if any, of proglucagon is processed to GLP-1 (20). Active GLP-1 (7–36NH\(_2\)) is therefore not a product of the alpha-cell proglucagon processing under normal conditions, according to the present studies. However, we detected significant amounts of fully processed active GLP-1 in pancreatic tissue from Gcgr\(^{-/-}\) mice. A plasticity in the proglucagon producing cells may therefore exist, as has previously been reported (19, 39), which can be activated upon...
metabolic or anatomical alterations. However, GLP-1 remained a minor product of proglucagon amounting to only ~1% of glucagon. If active GLP-1 was present in the WT pancreas in the same proportion, the concentration of active GLP-1 would be below detection limit (0.6 pmol/g). The larger amount of active GLP-1 in the pancreas may result in increased concentrations of circulating GLP-1 (14) in Gcgr−/− mice, and this has been suggested as an underlying reason for the improved glucose tolerance of these mice (11, 22). In connection with this finding we thought that it would be interesting to investigate the expression of the GLP-1 receptor in the markedly abnormal islets of the Gcgr−/− mice compared with the WT littermates. For this we used specific murine GLP-1 receptor antibody (37), but the receptor was found exclusively on the beta-cells. The sensitivity of the immunohistochemical approach does not allow us to exclude expression of a small (and therefore undetectable) number of GLP-1 receptors on the glucagon and somatostatin producing cells, but at least the hyperplastic and hypertrophic alpha-cells do not seem to express the receptor in large amounts, and expression in the beta-cells was apparently not downregulated by the larger than normal exposure to the ligand. Interestingly, the GLP-1 receptor staining in WT mice was confined to the cell membrane of the beta-cells, whereas both membrane and cytoplasmic staining was found in Gcgr−/− islets, due to internalization of the GLP-1 receptor. We speculate that this may be due to increased secretion of pancreatic GLP-1 in the Gcgr−/− mice thereby resulting in an increased internalization of the GLP-1 receptor. However, further studies are needed to clarify this.

We found an increased density of somatostatin cells in the Gcgr−/−islets compared with the WT islets, as also reported in Ref. 14 and in accordance with the higher somatostatin content measured in the pancreatic extracts from these mice. The delta-cells may therefore contribute to the increased islet area detected in Gcgr−/− mice. Finally, it has been suggested that some alpha-cells, contributing to the hyperplasia, may trans-
the glucagon receptor is not expressed in human or mouse adipose tissue or in muscles [there is a single aberrant report of receptor expression in muscles (17)], and we were also able to demonstrate that ligation of the kidneys did not acutely influence amino acid clearance in WT mice. We therefore suggest that the observed effects of glucagon receptor blockade on amino acid metabolism are a consequence of disrupted glucagon signaling in the liver.

Cultured alpha cells (alphaTC1.9) incubated with amino acids showed increased proliferation and glucagon secretion, supporting a role for amino acids as growth factors for the alpha-cells (32, 41) and as stimulators of glucagon secretion as previously suggested (3, 26, 36, 38, 41, 45).

The amino acid clearance experiments were performed using mice anesthetized with isoflurane. Isoflurane has, in contrast to other rodent anesthetics, previously been shown not to attenuate arginine-stimulated glucagon secretion in mice (52), supporting that the glucagon responses observed are physiologically relevant. The use of anesthetized mice allowed us to obtain sufficient plasma to perform accurate measurements of amino acids, glucagon, and insulin. In addition, by using anesthetized mice we were also able to exclude the effects of muscular contractions on plasma amino acids (46).

Some of the findings in Gcgr<sup>−/−</sup> mice are reflected in human studies; patients with inactivating glucagon receptor knockout mutations show pancreatic swelling, hyperaminoacidemia, and hyperglucagonemia (27). Subjects with nonalcoholic fatty liver disease (NAFLD) and type 2 diabetes have been shown to have elevated concentrations of plasma amino acids and hyperglucagonemia (49). These findings suggest that, also in humans, impaired liver function leads to hyperglucagonemia, perhaps as a consequence of impaired glucagon action on hepatic amino acid clearance (5–7).

![Figure 7. Plasma insulin and blood glucose concentrations following amino acid stimulation during pharmacological disruption of the glucagon receptor. A: both glucagon receptor antagonist (GRA, red line and circles) and vehicle (blue line and upward triangles)-treated mice responded with a fast increase in plasma insulin concentration following amino acid infusion (Vamin, 1 μmol/g body wt). Control groups receiving phosphate-buffered saline (PBS) instead of amino acids (GRA + PBS, pink line and squares; and vehicle + PBS, light blue line and downward triangles) showed no change in insulin concentrations. B: the incremental areas under the curve (iAUC<sub>0-20 min</sub>) are shown for the 4 groups. The amino acid stimulation resulted in an increase in insulin concentrations in both groups. C: blood glucose concentrations increased in response to the amino acid stimulation in both groups. Blood glucose concentrations also increased in the control groups receiving PBS. D: the total areas under the curve (iAUC<sub>0-20 min</sub>) are shown for blood glucose levels. Data are presented as means ± SE (n = 3–7).](image-url)
Disruption of glucagon signaling causes hyperaminoacidemia

acids, turnover, leading to elevated concentrations of circulating amino acids that stimulate alpha-cells. In line with this, patients with glucagon-producing tumors (glucagonomas) show hyperaminoacidemia (9, 50). This may be caused by the high rate of glucagon accelerated hepatic amino acid turnover and ureagenesis.

In conclusion, both pharmacological and genetic disruption of glucagon receptor signaling lead to severely impaired amino acid clearance in mice, supporting an essential role for glucagon receptor signaling in acute amino acid turnover in mice. Furthermore, mice lacking glucagon receptor signaling (Gcgr−/− mice) have hyperaminoacidemia, hyperglucagonemia, and alpha-cell hyperplasia. These findings support the existence of a liver-alpha-cell axis with amino acid and glucagon feedback loops in mice. This feedback circuitry may be particularly important during ingestion of protein-rich meals that raise the concentration of circulating amino acids. Disruption of the axis, be it by liver dysfunction as seen in NAFLD patients or in defects in glucagon receptor signaling as seen in patients with inactivating glucagon receptor mutations or treatment with a glucagon receptor antagonist, leads to hyperaminoacidemia and is, in this way, responsible for the apparent hypersecretion of glucagon (hyperglucagonemia) rather than disturbances in glucose metabolism.

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Disclosures

No conflicts of interest, financial or otherwise, are declared by the authors.

Author Contributions


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