Glucose-inducible hypertrophy and suppression of anion efflux in rat beta cells

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Abstract

Hypertrophy of beta cells from obese fa/fa rats is associated with increased sensitivity to basal glucose. Exposure to glucose in culture distorts insulin secretion more in beta cells from large than small islets from fa/fa rats. The aim of the present study is to investigate whether increased beta cell volume is associated with both glucose hypersensitivity and altered activity of the glucose-sensitive anion conductance. Beta cells from fa/fa rats had increased volume compared with those from lean rats after 24 h culture. Three-day exposure to 25 mM glucose in culture induced 10–15% hypertrophy in beta cells from lean rats and basal secretion from intact islets was increased tenfold. Estimates of ion channel activity were made from measurement of radiolabeled ion efflux. Taurine efflux, a marker of glucose-regulated anion channel activity, was reduced after high glucose exposure but no alterations in glucose-dependent K⁺ efflux were detected. The reverse hemolytic plaque assay was used to determine the contributions of the number of secreting cells (recruitment) versus secretion per cell in beta cells from enlarged (>250 µm diameter), intermediate (125–250 µm) and small (<125 µm) islets from lean and obese rats exposed to conditions mimicking hyperglycemia. After overnight culture, basal secretion was twofold greater from beta cells of large fa/fa islets compared with all other groups. Recruitment at low glucose was increased in all lean or fa/fa beta cells derived from >125 µm islets. When beta cells from small islets were exposed to supra-physiological glucose for 3 days, recruitment was increased at basal glucose and blunted at high glucose. Glucose exposure converts the recruitment profile of beta cells from small islets to resemble that of beta cells from large islets while inducing cellular hypertrophy and reduced anion conductance. However, hypertrophy alone did not predict functional characteristics of overnight-cultured beta cells from fa/fa rats.

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Introduction

Short-term changes in cell volume are postulated to modulate cellular function, for example by altering second messenger system activity (Haussinger & Lang 1991). Longer-term cell swelling might, however, be associated with pathological processes. In diabetes-related conditions, cell swelling caused by exposure to high glucose is implicated in retinopathy (Wakisaka et al. 1999). Volume regulation of pancreatic beta cells can also be induced by glucose (Best 1997) or osmotic changes (Britsch et al. 1994). Exposure to glucose but not non-metabolized analogues leads to increased cell volume coincident with increased insulin secretion (Miley et al. 1997). The mechanism by which changes in cell volume affect beta cell function is proposed to be through activation of a volume-sensitive anion (Cl⁻) conductance (Best et al. 1996a,b). Combined with inactivation of the K_ATP-dependent channel, outward movement of anions would contribute to beta cell depolarization (Best 1997).

In obesity, enlargement of pancreatic islets is a classic observation in both human subjects and rodent models. In the fa/fa Zucker rat, such enlargement has been attributed to both hyperplasia and hypertrophy of the beta cells (Hayek & Woodside 1979). If volume is inferred from published data on average beta cell area from islets isolated from lean and obese rats, an outcome of 30–60% increased volume per beta cell is calculated (Chan et al. 1999). Since beta cell mass of similar-aged fa/fa rats is reported to be approximately double that of controls (Pick et al. 1998), then as much as half of the increase might be attributable to an increase in cell volume. Previously, cellular hypertrophy in fa/fa rats was associated with basal glucose hypersensitivity and reduced expression of exocytotic proteins (Chan et al. 1999). This study was undertaken to determine if changes in the volume-sensitive anion conductance might...
contribute to beta cell hypertrophy associated with altered glucose responsiveness.

Materials and Methods

Animals

Lean and obese (fa/fa) 8-week-old Zucker rats of both sexes were obtained from Charles River Laboratories (St Constant, QC, Canada) and cared for in accordance with the principles of laboratory animal care and the regulations of the Canadian Council on Animal Care, vol. I and II. Standard chow (Purina 5001, Ralston Purina, St Louis, MO, USA) and water were provided and the animals allowed to feed ad libitum. The rats were fasted overnight prior to any experimental procedures, which were performed under sodium pentobarbital anesthesia (60 mg/kg i.p.). A fasted blood sample was obtained by cardiac puncture just prior to islet isolation. A fed blood sample was obtained from the tail vein of the rats 3–4 days prior to islet isolation.

Islet isolation and culture

Pancreatic islets were isolated as described (Kibenge & Chan 1995). Islets from fa/fa rats were divided into small (<125 μm, faSM), intermediate (<250 μm but >125 μm, faINT) or large (>250 μm, faLG) diameter groups, as in previous studies (Chan et al. 1999). Although 8-week-old lean rats have some islets >250 μm in size, these account for a very small proportion of the islets liberated by the isolation process. Thus, lean rat islets were sub-divided into <125 μm (LnSM) and >125 μm (LnINT) groups. The culture medium was Dulbecco’s modified Eagle’s medium (Gibco/BRL, Burlington, ON, Canada) containing 12.5 or 25.0 mM glucose, supplemented with 10 mM HEPES (Sigma), 1% antibiotic/antimycotic solution (Sigma), and 10% calf serum (Gibco/BRL) (final concentrations). These glucose concentrations were previously shown to induce more rapid alterations in secretory function in islets from fa/fa rats compared with lean controls (Chan et al. 1996). The islets were maintained in an environment of humidified 95% air and 5% CO₂ at 37 °C. Insulin secretion from intact islets was measured as described previously (Kibenge & Chan 1995). For studies of individual beta cells, islets were exposed to PBS (pH 7.4) containing trypsin (0.016%) and EDTA (0.2 mM) for 11 (small) to 15 min (large). The digestion was stopped by thoroughly washing the cells in fresh PBS.

Beta cell area measurement

Beta cells were mounted on glass slides as described below for the reverse hemolytic plaque assay. At least 50 cells from each islet size group per rat were quantified using computer-assisted morphometry of video images transmitted via a Zeiss microscope at a magnification of 1026 ×. The software was from Bioquant (R & M Biometrics, Nashville, TN, USA). Cell volumes were computed from the area measurements, assuming a spherical shape.

Estimation of ion channel activity

Efflux from pre-loaded islets of ⁸⁶Rb⁺ as a tracer for K⁺ was used to assess activity of ATP-dependent K⁺ (KATP) channels as described previously (Chan & MacPhail 1996). The activity of volume-sensitive anion channels was assessed by estimating efflux from pre-loaded cells of [³H]taurine, essentially as described by Best & Benington (1998). Briefly, batches of 100 islets incubated in buffer containing (mM) NaCl (135), KCl (5), MgSO₄ (1) NaH₂PO₄ (1), CaCl₂ (1.2), glucose (0–5) and HEPES (10, pH 7.4) and 4 µCi [³H]taurine for 2 h at 37 °C. After washing extensively to remove extracellular tracer, the islets were resuspended in fresh medium containing glucose and other reagents as indicated in the Results. Test glucose concentrations were the same as those used by others (Best & Benington 1998) in order to facilitate comparisons. Isotonicity was maintained by adding isosmotic glucose solution (308 mM) and reducing the volume of NaCl accordingly. At 5-min intervals, the supernatant was removed and fresh solutions added for a total of 20 min. At termination of the experiment, supernatant and islet pellets were resuspended in scintillation medium for liquid scintigraphy.

Reverse hemolytic plaque assay (RHPA) for beta cells

Insulin secretory activity of individual beta cells was assessed using a RHPA as described previously (Chan et al. 1998). This assay permits assessment of the activity of individual hormone-secreting cells (Smith & Neill 1987). Beta cells were exposed to glucose (1.4–16.5 mM) plus insulin antibody for 2 h, then guinea pig complement for 30 min. At the termination of the experiment, the cells were fixed with 1.0% glutaraldehyde in 0.04 M phosphate buffer (pH 7.5) for 15 min, then transferred to 0.01 M PBS (pH 7.4) and stored at 4 °C until immunostained for insulin. For each slide, recruitment, defined as the proportion of secreting beta cells, was assessed by light microscopy. Depending on cell density, 50–200 cells per slide were categorized. Insulin-positive cell doublets or clumps were not included in the totals. Computer-assisted measurement of plaque and cell areas was carried out using the Bioquant IV (R & M Biometrics) software package. Plaque areas were quantified only for 2.8, 8.3 and 16.5 mM glucose, representing baseline, half-maximal, and maximal glucose concentrations with respect to recruitment for the majority of the populations tested.
Images were transmitted via a Zeiss microscope and video camera for viewing at a final magnification of 1026 × on a computer screen.

Statistical analysis of data

Data in the Results section are reported as means ± S.E. unless otherwise stated. For beta cell studies, the number of individual islet donor rats and the total number of cells quantified are indicated as ‘n: x’. Significant differences in the recruitment were determined using χ² analysis with the Bonferroni adjustment for multiple comparisons using Minitab version 8.3 software (Minitab Inc., State College, PA, USA). The EC₅₀ values for concentration–response curves were calculated using GraphPad Prism (version 3.0 for Windows, GraphPad Software, San Diego, CA, USA, www.graphpad.com). All other comparisons were conducted using ANOVA (Minitab). Differences were considered significant if P<0.05 or better.

Results

Plasma glucose concentrations

There was no significant difference in plasma glucose concentrations of fasted lean vs fa/fa rats (6·4 ± 0·7 vs 7·0 ± 0·3 mM, P>0.05). However, in unstarved rats, plasma glucose concentration of fa/fa rats was 8·6 ± 0·4 mM, compared with 7·7 ± 0·4 mM in lean animals (P<0.05).

Beta cell size

Cell area in 24 and 72 h cultured beta cells was measured (Table 1). After 24 h exposure to 12·5 mM glucose, cell area was elevated by 15–30% in faINT and faLG compared with LnSM cells, consistent with previous findings (Chan et al. 1999). A similar trend to hypertrophy was noted in faSM beta cells, but this was not significant. Computed volume increases ranged from 25 to 40%. After high glucose exposure for 72 h, the areas of both faSM and LnSM beta cells became significantly enlarged. The volume increase was 15–25%. In contrast, there was no significant increase in beta cell area after 72 h culture in 12·5 mM glucose.

Ion fluxes

Metabolism of glucose activates a volume sensitive anion channel that transports mainly Cl⁻ and potentiates cellular depolarization (Miley et al. 1997). Induction of cellular hypertrophy might alter the activity of this channel and contribute to the adaptation to glucose observed in lean rats. After culture for 24 h in 12·5 mM glucose, ³H-taurine efflux was ~30% over 20 min in the presence of 0·5 mM glucose. Efflux was elevated by 20 mM glucose to 50% but in the presence of the anion channel blocker, 4,4′-dithiocyanotostilbene-2,2′-disulphonic acid (DIDS, 0·1 mM), was not different than at low glucose (Fig. 1A). Three-day culture in 12·5 mM glucose resulted in an overall suppression of taurine efflux, but the cells were still responsive to both glucose and DIDS. Exposure to chronically elevated glucose for 72 h markedly reduced channel activity in the presence of both 0·5 mM and 20 mM glucose (Fig. 1A) and DIDS did not effectively block taurine efflux.

Previously we showed that the altered glucose metabolism in islets of fa/fa rats is paralleled by changes in Kᵦᵦᵦ channel activity, such that by 5 mM glucose ⁸⁶Rb⁺ efflux was minimal and 15 mM glucose had no further effect. In

Table 1 Effect of culture conditions on individual beta cell area and calculated volume. 

<table>
<thead>
<tr>
<th>Islet donor</th>
<th>Cultured overnight – 12.5 mM glucose</th>
<th>Cultured 3 days – 12.5 mM glucose</th>
<th>Cultured 3 days – 25.0 mM glucose</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>A</td>
<td>V</td>
<td></td>
</tr>
<tr>
<td>LnSM</td>
<td>117·7 ± 1·6 (8:1130)</td>
<td>126·5 ± 1·5 (8:540)</td>
<td>136·5 ± 3·0 (7:425)*</td>
</tr>
<tr>
<td></td>
<td>961</td>
<td>1070</td>
<td>1200</td>
</tr>
<tr>
<td>LnNT</td>
<td>128·54 ± 2·1 (9:457)</td>
<td>ND</td>
<td>ND</td>
</tr>
<tr>
<td></td>
<td>1096</td>
<td></td>
<td></td>
</tr>
<tr>
<td>faSM</td>
<td>134·7 ± 3·0 (5:629)</td>
<td>ND</td>
<td>147·2 ± 4·7 (4:227)*</td>
</tr>
<tr>
<td></td>
<td>1176</td>
<td></td>
<td>1343</td>
</tr>
<tr>
<td>faNT</td>
<td>148·1 ± 2·8 (6:367)*</td>
<td>ND</td>
<td>ND</td>
</tr>
<tr>
<td></td>
<td>1356</td>
<td></td>
<td></td>
</tr>
<tr>
<td>faLG</td>
<td>137·4 ± 1·9 (7:931)*</td>
<td>ND</td>
<td>ND</td>
</tr>
<tr>
<td></td>
<td>1212</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

*P<0.05 compared with LnSM islet donors. ND, not done.
contrast, in lean rat islets a concentration-dependent closure of channels was evident from 0 to 15 mM glucose (Chan & MacPhail 1996). Therefore, we hypothesized that glucose exposure of the lean islets for 72 h might have either decreased $^{86}\text{Rb}^+$ efflux at basal glucose that could explain elevated insulin release or that higher glucose would fail to suppress $^{86}\text{Rb}^+$ efflux, consistent with the small degree of stimulation of insulin secretion observed. As shown in Fig. 1B, $^{86}\text{Rb}^+$ efflux was similar to control (overnight) values after 72 h of 12·5 or 25 mM glucose exposure. Similar sensitivity to the channel-opening agent diazoxide was also observed.

Insulin secretion from intact islets was also compared using glucose and the various channel modulators. As expected, basal insulin was elevated and glucose-stimulated insulin secretion was blunted (50% over basal) in islets cultured for 72 h in high glucose compared with 24 h cultured islets (85% over basal) (Fig. 1C). Three-day culture in 12·5 mM resulted in slightly elevated basal secretion, and a glucose-stimulated insulin response of $\sim$325%.

**Insulin secretory activity of beta cells**

Examination of insulin secretion by individual beta cells can reveal whether secretory changes are due to a change in the number of cells responsive to a particular glucose concentration (recruitment) or to altered secretion by each cell. The latter is assessed by measurement of plaque area in the RHPA. Secretory data can be correlated with individual beta cell size. The effects of culture glucose were compared using beta cells from small (SM) islets as for the ion flux experiments. These data were compared with those using beta cells from intermediate (INT) and large (LG) islets of both lean and $fa/\bar{fa}$ rats.

Recruitment of beta cells from small islets of either lean or $fa/\bar{fa}$ rats was $\sim$6% during exposure to basal glucose (1·4–4·2 mM) after 24 h culture in 12·5 mM glucose. In comparison, recruitment of beta cells in the faINT/LG and lnINT groups was significantly elevated (12–18% at 2·8 mM) (Fig. 2A and B). Increased sensitivity to low glucose was reflected by a left shift in the EC$_{50}$ to glucose in faINT/LG beta cells (Table 2). In contrast, the maximum recruitment was suppressed in all $fa$-derived populations compared with lean-derived cells (Fig. 2).
predicted that by increasing the duration of culture and the glucose concentration, beta cells from small islets might exhibit functional properties more like those of larger islets. After 72 h culture in 25.0 mM glucose, faSM and LnSM beta cells had a marked increase in recruitment to basal glucose (25–40% at 2.8 mM) (Fig. 2C) compared with 72 h culture in 12.5 mM glucose. Maximal recruitment reached about 45%, which was higher than in 12.5 mM cultured beta cells. Nonetheless, this represents a blunting of the glucose response, since the ratio of stimulated to basal secretion was approximately 1.5 for culture in 25 mM glucose versus 3.5 for 24 h culture in 12.5 mM glucose or 6.2 for 72 h culture in 12.5 mM glucose. There was no change in the EC50 for glucose after 72 h culture at either glucose concentration (Table 2).

The plaque sizes (Fig. 3) were assessed to provide indices of secretion per cell (Smith & Neill 1987). Mean plaque area was about 4000–5000 µm² for all groups except faLG beta cells, which averaged 7000–9000 µm². Except in LnSM beta cells there was no relationship between plaque area and glucose concentration. After 72 h culture, there was no discernible pattern in plaque areas (not shown).

Discussion

In obese animal models including the Zucker fa/fa rat, pancreatic islet hypertrophy and hyperplasia are a classical finding (Shino et al. 1973, Larsson et al. 1977). In young adult Zucker rat islets, about half of the increase in islet volume may be attributed to beta cell hypertrophy (Chan et al. 1999). Why beta cells in obese rodents become enlarged is not clear but ambient hyperglycemia, such as was observed in this study, may be a factor. Moreover, hypertrophied beta cells from freshly isolated islets of fasted rats exhibit functional distinctions from smaller cells that appear to contribute to inappropriate glucose-stimulated insulin secretion. These differences include increased secretion at basal glucose levels and a reduction in insulin...
content after exposure to high glucose (Chan et al. 1999). Interestingly, larger beta cells from normal rats are more likely to secrete at physiological glucose levels (Giordano et al. 1993, Chan et al. 1999) but there is no size difference in secretory versus non-secretory cells from the largest islets of fa/fa rats (Chan et al. 1999). In this study we examined the hypothesis that beta cell enlargement might be due to cellular swelling caused by environmental factors, such as glucose exposure. Our data show that extreme glucose exposure induces hypertrophy of normal beta cells that correlates with an increase in basal insulin secretion. The function of such beta cells was then compared with those of fa/fa rats, which were already hypertrophied.

Three-day exposure to supra-physiological glucose concentrations induced swelling in lean rat beta cells (25%) that was comparable to that measured in fa/fa rats (22–41%). Acute exposure of beta cells to 20 mM glucose invoked a 10–15% increase in cell volume that was maintained for the duration of the test (Miley et al. 1997).

### Table 2

Glucose sensitivity (EC50) of cultured beta cells. Units are mM ± SE for (n) islet donors

<table>
<thead>
<tr>
<th>Islet donor</th>
<th>Cultured overnight – 12.5 mM glucose</th>
<th>Cultured 3 days – 12.5 mM glucose</th>
<th>Cultured 3 days – 25.0 mM glucose</th>
</tr>
</thead>
<tbody>
<tr>
<td>LnSM</td>
<td>6.35 ± 0.97 (8)</td>
<td>8.34 ± 0.54 (8)</td>
<td>7.73 ± 0.94 (7)</td>
</tr>
<tr>
<td>LnINT</td>
<td>NAC (9)</td>
<td>ND</td>
<td>ND</td>
</tr>
<tr>
<td>faSM</td>
<td>8.09 ± 2.40 (5)</td>
<td>ND</td>
<td>7.21 ± 1.99 (4)</td>
</tr>
<tr>
<td>faINT</td>
<td>3.66 ± 0.92* (6)</td>
<td>ND</td>
<td>ND</td>
</tr>
<tr>
<td>faLG</td>
<td>5.00 ± 1.40 (7)</td>
<td>ND</td>
<td>ND</td>
</tr>
</tbody>
</table>

*P<0.05 compared with faSM beta cells by ANOVA. NAC, not able to calculate; ND, not done.

### Figure 3

Estimation of secretion per beta cell, by quantification of plaque area (µm², mean ± S.E.) generated by exposure of overnight-cultured beta cells to 2.8, 8.3 or 16.5 mM glucose for beta cells from LnSM (open bars), faSM (solid bars), LnINT (cross-hatched bars), faINT (dark stipple) and faLG (light stipple) islet donors after overnight culture in 12.5 mM glucose. Numbers of donor animals and cells are as for Table 2. *P<0.05 compared with 2.8 mM glucose, §P<0.05 compared with LnSM cells.
Basal insulin secretion from isolated islets was increased by tenfold. While the insulin response to higher glucose was not different from 24 h cultured islets, the fold increase over basal was blunted. These results are typical of sustained activation of beta cells (Ling & Pipeleers 1996).

Because cell swelling can be induced by glucose to activate an anion conductance (Best 1997, Miley et al. 1997) and is correlated with an increase in insulin secretion (Britsch et al. 1994, Miley et al. 1997), we considered that activity of the glucose-sensitive anion channel might be increased after chronic glucose exposure.

A role for a glucose-sensitive anion conductance in beta cell signaling has just started to become appreciated, although the notion that Cl⁻ fluxes can modulate insulin secretion was advanced more than 20 years ago (Sehlin 1978). This ATP-dependent channel was first described in 1978 by Kinard & Satin and is likely Cl⁻ conducting, based on pharmacological and electrical properties. The mechanism by which the anion conductance leads to beta cell depolarization is presumed to be a consequence of Cl⁻ efflux (Best et al. 1996a, b). By adding to the effects of K_{ATP} channel inactivation, the intensity of electrical activity could be regulated (Best 1997). Therefore, sustained cell swelling induced by chronic exposure to high glucose might increase the activity of the volume-sensitive anion channel and contribute, in particular, to enhanced basal insulin secretion. In this study, the opposite effect was observed in that [³H]taurine efflux was decreased by about 30% at basal glucose and was not increased by elevated glucose in the chronically cultured cells. However, the generalized suppression of activity of the volume-sensitive anion channel by glucose exposure might contribute to the reduction in glucose-stimulated insulin secretion observed, since its role is to potentiate the depolarization of the beta cell induced by K_{ATP} channel closure. Support for this conclusion is given by studies in which glucose-stimulated Cl⁻ efflux was blunted in islets from obese diabetic (db/db) mice (Berglund & Sehlin 1980) concomitant with impaired insulin secretion. Since db/db mice are markedly hyperglycemic, the notion that anion conductance is adversely affected by chronic glucose exposure is supported. However, K_{ATP} channel activity was also abnormal in those mice (Berglund et al. 1978) but was not altered in the current study. In other cell types such as hepatocytes, cell swelling induces changes in metabolic processing of glucose by inhibiting glycolysis and glycocalyx (Haussinger & Lang 1991). The effects of volume changes on such processes in beta cells has not been examined.

The fa/fa Zucker rat is a model of obesity that maintains relatively normal fasting glucose and glucose tolerance by compensatory increases in both basal and stimulated insulin secretion. Like db/db mice, a genetic mutation renders the leptin receptor non-functional in fa/fa rats (Chua et al. 1996) but the phenotype is not as severe. The obese rats we obtain from Charles River, Inc. consistently exhibit fasting glucose levels that are not significantly different from age-matched lean controls, yet average about 0.5 mM higher (Chan et al. 1999, Kibenge & Chan 2001). However, single fasting measurements fail to capture information about 24 h glucose exposure of the beta cells. An increase in ambient plasma glucose concentrations was also observed in this study. These measurements were obtained from plasma collected near the beginning of the light cycle. The fa/fa Zucker rat may have a disrupted diurnal rhythm such that serum glucose is more elevated during the dark hours (Martin et al. 1978). The possibility that glucose-induced swelling may contribute to beta cell hypertrophy in fa/fa rat islets cannot be ruled out. To compare beta cell size with secretion patterns of beta cells from various populations of islets from both lean and obese rats, the RHPA was used.

After short-term culture in moderate glucose, beta cells isolated from larger islets from both lean and fa/fa rats had elevated recruitment (number of actively secreting cells) at low glucose. There was no direct correlation with beta cell area, because faSM and faLG cells had a similar size distribution but faLG cells were two times more likely to secrete at 2.8 mM glucose. Moreover, secretion per cell (plaque area) was similar in all beta cell groups except the faLG cells, which secreted approximately twofold more insulin independent of glucose concentration. Although extreme glucose exposure can induce hypertrophy and a secretory phenotype similar to that observed in enlarged beta cells from obese rats, it appears unlikely that the two phenomena are closely linked. Indeed, associations between insulin secretory patterns and beta cell volume are more closely linked in freshly isolated islets from fasted rats where glucose was reduced for approximately 18 h, according to data from our previous study (Chan et al. 1999).

In conclusion, these studies have shown that chronic exposure to glucose does induce beta cell swelling. Unexpectedly, this was associated with a blunted response of the glucose-sensitive anion conductance. Therefore, chronic swelling is unlikely to account for enhanced basal secretion but may contribute to the failure of glucose to stimulate insulin release. Moreover, while exposure to glucose of normal beta cells from lean islets leads to a phenotype similar to that of beta cells from enlarged islets from obese rats, this congruence appears not to be related to hypertrophy-induced mechanisms per se. It is possible that beta cell enlargement is caused by glucose suppression of cell death but since beta cell turnover is normally very low, quantitative assessment of a further reduction is technically difficult.

Acknowledgement

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