Critical Role of Gap Junction Coupled K_{ATP} Channel Activity for Regulated Insulin Secretion

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Pancreatic β-cells secrete insulin in response to closure of ATP-sensitive K⁺ (K_{ATP}) channels, which causes membrane depolarization and a concomitant rise in intracellular Ca^{2+} (Cai). In intact islets, β-cells are coupled by gap junctions, which are proposed to synchronize electrical activity and Cai oscillations after exposure to stimulatory glucose (>7 mM). To determine the significance of this coupling in regulating insulin secretion, we examined islets and β-cells from transgenic mice that express zero functional K_{ATP} channels in approximately 70% of their β-cells, but normal K_{ATP} channel density in the remainder. We found that K_{ATP} channel activity from approximately 30% of the β-cells is sufficient to maintain strong glucose dependence of metabolism, Cai, membrane potential, and insulin secretion from intact islets, but that glucose dependence is lost in isolated transgenic cells. Further, inhibition of gap junctions caused loss of glucose sensitivity specifically in transgenic islets. These data demonstrate a critical role of gap junctional coupling of K_{ATP} channel activity in control of membrane potential across the islet. Control via coupling lessens the effects of cell–cell variation and provides resistance to defects in excitability that would otherwise lead to a profound diabetic state, such as occurs in persistent neonatal diabetes mellitus.

Introduction

β-cells within the intact islet of Langerhans exhibit synchronous glucose-dependent bursts of electrical activity. Dissociated β-cells also show increased electrical activity at elevated glucose concentrations, but this activity is variable from cell to cell, with some cells being electrically “silent” and others continuously “bursting” at any given glucose concentration [1–4]. Synchronization of electrical activity via gap junctions has long been argued as essential for normal glucose-dependent insulin secretion in the intact islet [5–8]. In particular, islets devoid of gap junction–forming Connexin 36 (Cx36) exhibit limited synchronization of glucose-stimulated intracellular Ca^{2+} (Cai) [8]. To test the extent and role of this coupling in the islet electrical response, it would be desirable to examine within the intact islet the behavior of cells that would be silent or bursting as isolated cells. To date, it has not been possible to know which cell is which in the setting of the intact islet. However, we have generated transgenic mice containing two electrically distinct types of β-cells that uniquely permit us to track the behavior of specific individual cells within the intact islet [9]. As previously shown, these mice express a β-cell–specific, dominant-negative Kir6.2[AAA]-GFP transgene, in which the pore-forming subunit of the ATP-sensitive K⁺ (K_{ATP}) channel is rendered nonfunctional. Mosaic expression of the Kir6.2[AAA]-GFP transgene results in a high-level expression in approximately 70% of the β-cells, and these expressing cells are distributed randomly throughout the islet [9,10]. Thus, K_{ATP} channels are functionally “knocked out” of 70% of cells, yet the intact islets still show glucose-dependent insulin secretion, with an even steeper concentration dependence of insulin secretion than wild-type (WT) islets [9].

Results/Discussion

To examine the behavior of individual cells within the intact islet, we developed a microfluidic device to hold pancreatic islets stationary while under continuous fluid flow (see Figure S1) [11]. We compared the glucose-dependent Cai responses of WT and Kir6.2[AAA] transgenic islets (Figure 1). Near the periphery of the islet, fluorescence from a Ca^{2+} sensor (Fura Red) was relatively uniform in all cells (Figure 1A), and spectrally separable from green fluorescent protein (GFP) fluorescence (Figure 1B). Both the GFP-positive and negative populations of β-cells (K_{ATP}-absent and K_{ATP}-present cells, respectively) within a Kir6.2[AAA] transgenic islet showed identical Cai-responses (Figure 1C). WT islets were “Cai inactive” across the tissue below 8 mM glucose (i.e.,...
Ca_i remained low and no oscillations were observed; Figure 1D), consistent with previous studies [9,11]. Kir6.2[AAA] islets were also completely Ca_i inactive across the tissue at low glucose concentrations (2 and 4 mM), but they exhibited Ca_i activity at slightly lower glucose levels than WT (6 mM). This leftward shift in glucose dependence of Ca_i in Kir6.2[AAA] islets is consistent with the leftward shift in whole-islet glucose-dependent insulin secretion (Figure 1E) [9]. Kir6.2[−/−] and SUR1[−/−] mice, which completely lack K_ATP channels, show essentially no glucose dependence of electrical activity, Ca_i, or insulin secretion [12–14]. Thus, the near-normal glucose response of Ca_i (Figure 1D) and insulin secretion (Figure 1E) by Kir6.2[AAA] islets are initially somewhat surprising, since K_ATP is functionally knocked out in approximately 70% of the β-cells. Paracrine signaling could influence the electrical synchronization between islet cells, for instance by δ-cell secretion of somatostatin [15]. However, these data suggest that K_ATP channel activity in the residual 30% of cells is coupled through gap junction conductance that is strong enough to hyperpolarize the knockout cells. This suggestion, in turn, predicts that glucose dependence will be lost in dispersed Kir6.2[AAA]-GFP cells. Accordingly, we measured the glucose-stimulated Ca_i response and insulin secretion from dispersed cell preparations (Figure 2). Image fields of isolated Kir6.2[AAA] β-cells show uniform loading of the Ca_2þ indicator dye (Fura-2; Figure 2A), and no overlap
with GFP fluorescence (Figure 2B). Both the Ca\textsubscript{i} response and insulin secretion were essentially glucose independent in dispersed Kir6.2[AAA] cells (Figure 2C and 2D). This is in striking contrast to the Cai response and insulin secretion from dispersed WT cells which are strongly glucose dependent, but is consistent with previous results from SUR knockout mice [16].

It was previously shown that dispersed GFP-positive Kir6.2[AAA] \(\beta\)-cells are continuously depolarized in non-stimulatory glucose concentrations [9]. To examine whether cell–cell coupling reconstitutes glucose-dependent membrane potentials in intact islets, we imaged intact islets using a membrane potential–sensitive dye (Figure 3A). At 2 mM glucose, a few cells on the islet periphery were noticeably more depolarized than others. These bright cells were never

![Figure 2](image1)

**Figure 2.** Dispersed Cell Ca\textsubscript{i} and Insulin Responses

(A) Dispersed Kir6.2[AAA] cells labeled with Fura-2 (340-nm image). These are mainly \(\beta\)-cells, with the expected minority of non-\(\beta\)-cells.

(B) GFP image of the same field of view of (E). A fraction of cells (four of eight) are GFP positive.

(C) The glucose-stimulated Ca\textsubscript{i} response of dispersed cells from WT (black square) and Kir6.2[AAA] (red circle) islets measured using Fura-2 fluorescence microscopy. Kir6.2[AAA] cell preparations were a combination of GFP-labeled and non-labeled cells. The percentage of Ca\textsubscript{i}-active cells was measured in 2, 6, and 10 mM glucose on three separate days in WT and Kir6.2[AAA] dispersed cells, respectively (mean ± standard error of the mean [SEM], \(n = 3\)). The WT cells show a strong glucose dependence, whereas the Kir6.2[AAA] cells show similar numbers of Ca\textsubscript{i} active cells at all glucose concentrations.

(D) Insulin secretion from dispersed WT and Kir6.2[AAA] cells (mean ± SEM, \(n = 4\)) from static 1-h incubations. This incubation time shows K\textsubscript{ATP} channel–dependent insulin secretion, but does not encompass K\textsubscript{ATP} channel–independent insulin secretion shown to occur in knockout islets beyond 3 h of incubation [13]. The difference in means ± 95\% confidence interval for (C) and (D) are shown in Figure S2B and S2C, respectively.

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![Figure 3](image2)

**Figure 3.** Membrane Potential and Metabolic Response of Kir6.2[AAA] Islets and Cells

(A) The membrane potential of GFP-positive and -negative cells within a Kir6.2[AAA] islet. Islets were bathed in 500 nM of membrane potential dye (DiSBAC\textsubscript{2}(3)) (\(n = 4\)). This oxanol dye enters cells in a membrane potential–dependent manner with increasing intensity as a cell becomes depolarized, and is spectrally distinct from GFP. The GFP-positive and -negative cells showed similar intensity (membrane potential) at 2 and 15 mM glucose stimulation, and had responses consistent with the responses observed in WT islets (data not shown).

(B) The NAD(P)H glucose-dose response of WT (\(n = 9\)) and Kir6.2[AAA]-GFP (\(n = 10\)) islets. Device-trapped islets were treated to the indicated glucose concentrations for more than 5 min prior to image collection. The normalized change in NAD(P)H fluorescence intensity (\(\Delta F/\Delta F_0\)) is plotted versus glucose stimulation. The black and red dotted lines are the data fitted to a Hill equation with Km values of 9.0 ± 0.7 and 7.6 ± 0.7 mM for the WT and Kir6.2[AAA] islets, respectively.

(C) The NAD(P)H responses of dispersed GFP-negative and -positive Kir6.2[AAA] cells. The cells were incubated in 2 mM glucose (open bars) prior to incubation for 5–10 min with 15 mM glucose (closed bars). GFP fluorescence was used for the distinction as GFP negative or positive. NAD(P)H-intensity measures were taken from cells in four fields of view from nine separate dishes of Kir6.2[AAA] dispersed cells (\(n = 9\)).

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The oscillating active area and total islet area were measured in mM glucose (WT: 8 mM glc). The black bar in each panel indicates 30 s. Relative Ca²⁺ traces from the Fluo-4 response, and each panel shows the islets exposed to 0, 10, or 50 μM (n = 4 and 5, respectively) or 4 mM (n = 5 and 6, respectively) glucose (glc) were exposed to 0, 10, or 50 μM αGA. The islets were imaged to provide relative Ca²⁺ traces from the Fluo-4 response, and each panel shows the response in four different regions of a single islet. The representative traces shown are from a WT islet in 2 mM glucose with 50 μM αGA (WT: 2 mM glc + 50 μM αGA, n = 6), a Kir6.2[AAA] islet in 2 mM glucose with 50 μM αGA (Kir6.2[AAA]: 2 mM glc + 50 μM αGA, n = 5), and a WT islet in 8 mM glucose (WT: 8 mM glc). The black bar in each panel indicates 30 s. (B) The oscillating (Ca²⁺ active) area and total islet area were measured in each islet under each condition. The data are expressed as the fraction of oscillating area, calculated as the oscillating area divided by the total area (mean ± SEM) from WT islets in 2 mM (n = 4) and 4 mM glucose (n = 5), and Kir6.2[AAA] islets in 2 mM (n = 5) and 4 mM glucose (n = 6).

**Figure 4.** The Ca²⁺ Responses of WT and Kir6.2[AAA] Islets in the Presence of αGA, a Gap Junction Inhibitor

(A) Fluo-4, AM–labeled WT and Kir6.2[AAA] islets in 2 mM (n = 4 and 5, respectively) or 4 mM (n = 5 and 6, respectively) glucose (glc) were exposed to 0, 10, or 50 μM αGA. The islets were imaged to provide relative Ca²⁺ traces from the Fluo-4 response, and each panel shows the response in four different regions of a single islet. The representative traces shown are from a WT islet in 2 mM glucose with 50 μM αGA (WT: 2 mM glc + 50 μM αGA, n = 4), a Kir6.2[AAA] islet in 2 mM glucose with 50 μM αGA (Kir6.2[AAA]: 2 mM glc + 50 μM αGA, n = 5), and a WT islet in 8 mM glucose (WT: 8 mM glc). The black bar in each panel indicates 30 s. (B) The oscillating (Ca²⁺ active) area and total islet area were measured in each islet under each condition. The data are expressed as the fraction of oscillating area, calculated as the oscillating area divided by the total area (mean ± SEM) from WT islets in 2 mM (n = 4) and 4 mM glucose (n = 5), and Kir6.2[AAA] islets in 2 mM (n = 5) and 4 mM glucose (n = 6).

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seen to co-localize with GFP fluorescence (not shown), and were likely β-cells. The rest of the islet cells showed relatively dim uniform fluorescence, consistent with uniformly polarized membranes, and all of these cells brightened (depolarized) upon glucose stimulation. GFP-expressing and non-expressing cells had indistinguishable membrane polarization (Figure 3A). Thus, β-cells across the Kir6.2[AAA] islet have similar resting membrane polarization, and all depolarize with increased glucose concentrations.

In contrast to electrical activity, we have not observed metabolic coupling of β-cells [11,17], although there may be feedback between electrical activity and metabolic response. The Kir6.2[AAA] islets provide a unique opportunity to examine the potential feedback. We used intrinsic NAD(P)H fluorescence as a measure of cellular metabolic status [11,18]. In comparison to WT, Kir6.2[AAA] islets showed no difference in baseline NAD(P)H intensity, but showed a slight (~1 mM) leftward shift in the (glucose) response (Figure 3B). This shift is comparable to that of glucose-stimulated Ca²⁺ response and insulin secretion (see Figure 1D and 1E), suggesting a feedback modulation of metabolism during electrical activity. Such feedback could result from Ca²⁺ influx [19] or insulin signaling [20,21] since both likely modulate β-cell metabolism and occur when the tissue is electrically active. We further compared the baseline and stimulated responses of dispersed Kir6.2[AAA] cells (Figure 3C). Baseline NAD(P)H intensity was significantly elevated in GFP-positive cells (1.3-fold vs. GFP negative at 2 mM glucose), but was similar upon glucose stimulation. The elevated basal metabolism in GFP-positive cells with no difference in glucose-stimulated metabolism is again consistent with a modulating effect of Ca²⁺ or insulin. Thus, even though metabolism is not directly coupled between cells in the islet [11,17], entrained electrical activity and shifted Ca²⁺ responses can influence metabolic response.

The above results suggest a model whereby the Kir6.2[AAA] islet response is due to coupling of K<sub>ATP</sub> channel activity, presumably through gap junctions. This model predicts that inhibition of gap junctions would recapitulate the uncoupled Ca<sub>i</sub> activity found in dispersed Kir6.2[AAA] cells. Thus, we examined glucose-stimulated Ca<sub>i</sub> responses in the presence of a gap junction inhibitor, 18-glycyr rhetic acid (αGA) [22]. Ca<sub>i</sub> measurements of WT islets (Figure 4A) showed no β-cell oscillations at sub-threshold glucose levels in the presence or absence of either 10 or 50 μM αGA. In contrast, spontaneous Ca<sub>i</sub> oscillations were observed in the presence of 50 μM αGA in four of five Kir6.2[AAA] islets at 2 mM glucose, and in six of six Kir6.2[AAA] islets at 4 mM glucose (Figure 4B, Videos S1–S3). Unlike the synchronous oscillations across the entire islet that are seen in the absence of αGA (Figure 4A, bottom trace, and Figure 1C), these oscillations occurred asynchronously in small groupings of cells (Figure 4A, middle trace). The resulting asynchronous Ca<sub>i</sub> activity in portions of Kir6.2[AAA] islets is consistent with partial decoupling of the K<sub>ATP</sub> channel activity by the inhibition of gap junction conductance.

The data presented here indicate that glucose control of membrane potential, Ca<sub>i</sub>, and insulin secretion all depend critically on the gap junctional coupling of K<sub>ATP</sub> channel activity across neighboring cells within the islet (Figure 5). It has previously been shown that knockout of Cx36 in islets results in reduction of glucose-stimulated electrical synchrony [8]. Our results provide additional experimental evidence for gap junctional coupling between islet β-cells. Furthermore, our results demonstrate the mechanism by which the coupling of K<sub>ATP</sub> channel activity via gap junctions results in near-normal glucose-dependent electrical and Ca<sub>i</sub>
Kir6.2[AAA] cell depolarizes (red membrane) resulting in $\text{Ca}^{2+}$ influx through voltage-gated channels. Depolarization (red membrane) in both cells, and $\text{Ca}^{2+}$ influx through voltage-gated $\text{Ca}^{2+}$ channels.

In the absence of coupling through gap junctions, the cell with normal KATP channel activity (white cell) maintains polarity (blue membrane), but the Kir6.2[AAA] cell (green cell) would normally be depolarized; however, the KATP current from the normal cell is coupled through gap junctions to maintain membrane polarity (blue membrane).

The addition of glucose raises the ATP/ADP ratio, which closes the KATP channel in the normal cell. The loss of this K$^+$ current results in membrane depolarization (red membrane) in both cells, and $\text{Ca}^{2+}$ influx through voltage-gated $\text{Ca}^{2+}$ channels.

In the dispersed cells, this rise in ATP/ADP ratio results in depolarization and $\text{Ca}^{2+}$ influx occurring in both cells.

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(A) and (C) show the β-cell responses in the islet, and (B) and (D) show the dispersed β-cell responses.

(A) A WT β-cell has normal KATP channel activity (white cell), which maintains plasma membrane potential (blue membrane). The GFP-tagged Kir6.2[AAA] cell (green cell) would normally be depolarized; however, the KATP current from the normal cell is coupled through gap junctions to maintain membrane polarity (blue membrane).

(B) In the absence of coupling through gap junctions, the cell with normal KATP channel activity (white cell) maintains polarity (blue membrane), but the Kir6.2[AAA] cell depolarizes (red membrane) resulting in $\text{Ca}^{2+}$ influx through voltage-gated channels.

(C) The addition of glucose raises the ATP/ADP ratio, which closes the KATP channel in the normal cell. The loss of this K$^+$ current results in membrane depolarization (red membrane) in both cells, and $\text{Ca}^{2+}$ influx through voltage-gated $\text{Ca}^{2+}$ channels.

(D) In the dispersed cells, this rise in ATP/ADP ratio results in depolarization and $\text{Ca}^{2+}$ influx occurring in both cells.

Figure 5. Schematic Model of the Responses of β-cells in the Islet and Dispersed β-cells from Kir6.2[AAA] Transgenic Mice in Both Low (<6 mM) and High (>6 mM) Glucose Concentrations

(A) and (C) show the β-cell responses in the islet, and (B) and (D) show the dispersed β-cell responses.

(A) A WT β-cell has normal KATP channel activity (white cell), which maintains plasma membrane potential (blue membrane). The GFP-tagged Kir6.2[AAA] cell (green cell) would normally be depolarized; however, the KATP current from the normal cell is coupled through gap junctions to maintain membrane polarity (blue membrane).

(B) In the absence of coupling through gap junctions, the cell with normal KATP channel activity (white cell) maintains polarity (blue membrane), but the Kir6.2[AAA] cell depolarizes (red membrane) resulting in $\text{Ca}^{2+}$ influx through voltage-gated channels.

(C) The addition of glucose raises the ATP/ADP ratio, which closes the KATP channel in the normal cell. The loss of this K$^+$ current results in membrane depolarization (red membrane) in both cells, and $\text{Ca}^{2+}$ influx through voltage-gated $\text{Ca}^{2+}$ channels.

(D) In the dispersed cells, this rise in ATP/ADP ratio results in depolarization and $\text{Ca}^{2+}$ influx occurring in both cells.

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responses in the intact islet, even when a majority of β-cells are essentially glucose insensitive. More specifically, this coupling allows β-cells in low glucose with normal KATP channel activity to clamp the membrane potential of neighboring cells. The electrical properties of isolated β-cells are very variable [2–4]. This variability along with our results suggests that the rise in basal insulin secretion from Cx36 knockout islets is due to an inability of quiescent cells to clamp the membrane potential of other β-cells with low stimulation threshold. This may also explain an evolutionary drive to organize these secretory cells into a unique electrical syncytium. Such a mechanism provides a protective effect against hypoglycemia if individual cell properties change under pathological or physiological stimuli. Factors that reduce KATP channel density, such as channel subunit trafficking mutants [23,24], could do so to the point where individual cells become glucose insensitive. However, the remaining glucose-sensitive cells in the islet with sufficient residual KATP channel activity will maintain a glucose response of the whole islet. Such cell coupling would help resist the development of persistent hypoglycemic hyperinsulinemia (PHHI), unless compounded by additional defects, as may be the case in human PHHI [25].

Our results also have implications for the understanding of diabetic phenotypes resulting from KATP channel gain of function or from other causes of hypoeexcitability. It is now clear that overactive KATP channels cause neonatal diabetes due to inexcitability of β-cells and failure to secrete insulin [26,27]. Our data demonstrate that cell–cell coupling can ensure maintained hyperpolarization in all cells if KATP channel activity in just a few reaches a threshold of inexcitability. However, this also suggests that the domination of KATP channel strength on neighboring cell membrane potentials, may permit elevated KATP channel activity in only a few β-cells to clamp the remainder at a hyperpolarized potential, thereby stopping secretion. Such a mechanism may underlie the profound diabetic state in persistent neonatal diabetes mellitus [27]. Equivalent gain-of-function model systems [26] and the experimental approach that we describe here should allow a critical test of the syncytial involvement in these types of neonatal diabetes.

Materials and Methods

Islet isolation and secretion measurements. Islets were isolated as previously described [28,29] and maintained in complete RPMI medium 1640 containing 10% fetal bovine serum and 11 mM glucose at 37 °C under 5% humidified CO2 for 24–72 h. Islets were dispersed to single cells as described previously [9]. Insulin release was measured from pancreatic islets (ten per well) or dispersed cells (equivalent to 15 islets per well) in static 1-h incubations as described previously [9].

Confocal imaging of microfluidic device trapped islets. Devices were fabricated using the elastomer polydimethylsiloxane (PDMS) as described elsewhere [30] (see Figure S1 for details of fabrication and use). Islets were labeled with 4 μM of either Fluor-4, acetoxyethyl (AM) or Fura Red, AM at room temperature for 1 to 2 h in imaging buffer (125mM NaCl, 5.7 mM KCl, 2.5 mM CaCl2, 1.2 mM MgCl2, 10 mM HEPES, 2 mM glucose, and 0.1% bovine serum albumin [pH 7.4]). One- and two-photon microscopy was performed on a LSM 510 microscope with a 20 x 0.75 NA Fluor objective lens (Carl Zeiss, Thornwood, New York, United States). The device was held on the microscope in a humidified temperature controlled stage (Carl Zeiss) for imaging at 37 °C. Fluor-4 (Molecular Probes, Eugene, Oregon, United States) was imaged using the 488-nm laser line and the long-
pass 503-nm emission filter. NAD(P)H intensity was imaged with a 710-nm mode-locked Ti:Saph laser (~3.5 mW at the sample) and fluorescence collection through a non-descanned detector with a custom 380- to 550-nm filter (Chroma, Rockingham, Vermont, United States) [18]. We collected four focal planes separated by 3-μm intervals. Fura Red (Molecular Probes) was imaged using the 530-nm laser pass filter (620–680 nm). The GFP label of Kir6.2[AAA] islets was imaged using the 488-nm laser and band-pass (540/20) filter, with no noticeable bleed-through of the Fura Red signal. Membrane potential was measured with bis-(1,3-diethylthio- barbituric acid)trimethine oxonol (DiSBAC2(3); Molecular Probes). This dye accumulates in the cells in a Nernst-dependent manner [31]. We flowed the indicated solutions with 500 nM of DiSBAC2(3). These solutions were allowed to equilibrate for approximately 30 min to allow dye penetration before imaging. DiSBAC2(3) images were collected by 543-nm excitation and a long-pass 560-nm filter. GFP fluorescence in these same islets was imaged separately using 488-nm excitation and a 500- to 530-nm band-pass filter.

**Single cell Ca i measurements.** Prior to imaging, cells were labeled with 2 μM Fura-2 AM (Molecular Probes) for 30 min at room temperature and washed with imaging buffer containing 2 mM glucose. Imaging was done on a TE300 Eclipse microscope (Nikon, Melville, New York, United States) using the 40 × 1.3 NA DIC PlanFluor lens, side-mounted CoolSnap HQ camera (Roper Scientific, Tucson, Arizona, United States), and Fura-2 filter set (Chroma). The cells were placed on a stage enclosed in Plexiglas and kept in humidified air at 37 °C. The system was controlled with Metamorph 6.0 (Universal Imaging, Downington, Pennsylvania, United States). Image pairs were collected at 2-s intervals for 2 min after incubation for at least 10 min in 2, 6, and 10 mM glucose.

**Image analysis.** Images were analyzed with Meta Morph 5.0 (Universal Imaging). For Fluo-4 analysis, islets/regions of islets were outlined and their intensities versus time were plotted. From these plots, the relative intensity change (ΔF/Fo) and frequencies were calculated. For Fura-2 analysis, individual cells were outlined for calculation of the background corrected 340:380-nm intensity ratio. Cells were categorized as “calcium active” if they had a ratio that was either more than two standard deviations larger than that observed for WT cells in 2 mM glucose or demonstrated a change of more than one standard deviation change during the 2-min observation period.

**Supporting Information**

**Figure S1.** A Microfluidic Device, Designed to Hold a Pancreatic Islet Stationary in a Fluid Stream, was Fabricated Using the Elastomer Polydimethylsiloxane (PDMS) The fabrication of the device is described in [30].

(A) A dye-loaded microfluidic device. The image is labeled to show the Reagent well, Islet In/Out port, Wall trap area, and Waste port. This device relies on gravity flow from either the Reagent well or Islet In/Out port to the Waste port. Islets are brought into the device through the In/Out port. The islet travels through the main channel of the device until it comes in contact with Wall area.

(B) A differential interference contrast image of a pancreatic islet at the Wall trap area of the microfluidic device. This islet is in the main channel (height ~100 μm and width ~600 μm) touching the wall trap (bottom of image), with fluid flowing by gravity from the top to bottom of the image. Islets trapped in a microfluidic device channel with approximately 100 μm height are pressed against the coverslip surface for optimal microscopic imaging. The islet is held stationary by the coverslip, ceiling and wall trap. All the islets studied maintained their shape throughout the experiments, which suggests limited shear pressure. We have previously shown that WT islet Ca i and NAD(P)H responses are unperturbed in similar devices [11].

(C) Multiple images of sulfohydramine B (0.2 μM) dye as it is flown past a device-trapped islet. The distance from the coverslip is shown at the bottom-left corner of each image. Note that the dye solution is not observed in the wall trap area above 16 μm of depth. In this region of the device, the height of the channel drops from approximately 100 to 15 μm. This channel height allows fluid to flow while blocking islet movement. Once islets are trapped in the device, we plugged the In/Out port and started gravity flow from the Reagent well. (D) Changing the Reagent well solution (100 μl) to contain sulfohydramine B (0.2 μM) resulted in complete change of solution at >1 min. The half-maximum of this change was 27 s, which corresponds to a flow rate of approximately 1.2 μl min⁻¹, which is below the calculated maximum velocity (~3 μl min⁻¹), but slightly faster than the calculated flow rate (~0.7 μl min⁻¹). At this flow rate, we observed no noticeable warping of the islet, which is consistent with low shear. Furthermore, the islets remained stationary after long periods of flow time (minutes to hours) and during reagent solution changes. Keeping the islet in such a stationary position facilitates time-lapse imaging and the observation of similar regions after many different treatments.

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**Figure S2.** Difference in Means with Confidence Intervals for Figures 1E, 2C, and 2D Experimental details are as described in the text and figure captions. The difference in means (X₁−X₂) for each data point is plotted ± 95% confidence interval with the zero difference shown with a dashed line.

(A) Data from Figure 1E.

(B) Data from Figure 2C.

(C) Data from Figure 2D.

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**Video S1.** This Video Shows a Kir6.2[AAA] Islet Treated to 2 mM Glucose (2 mM Glc)

The elapsed time covers a period of 190 s at 1.57 frame/s. Some individual cells are observed to asynchronously increase in intensity. These responses are similar to those observed in WT islets at nonstimulatory glucose concentrations [11], and are consistent with an α-cell response.

DOI: 10.1371/journal.pbio.0040026.sv001 (407 KB MOV).

**Video S2.** This Video Shows the Same Kir6.2[AAA] Islet Shown in Video S1, but Treated to 2 mM Glucose with 10 μM zGA (2 mM Glc + 10 μM GA)

The elapsed time covers a period of 190 s at 1.57 frame/s. Similar to the response found with 2 mM glucose (Video S1), some individual cells are observed to asynchronously increase in intensity. These responses are similar to those observed in WT islets at nonstimulatory glucose concentrations [11], and are consistent with an α-cell response.

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**Video S3.** This Video Shows the Same Kir6.2[AAA] Islet Shown in Video S1, but Treated to 2 mM Glucose with 50 μM zGA (2 mM Glc + 50 μM GA)

The elapsed time covers a period of 383 s at 1.57 frame/s. Similar to the response found with 2 mM glucose (Video S1), some individual cells are observed to asynchronously increase in intensity. These responses are similar to those observed in WT islets at nonstimulatory glucose concentrations [11], and are consistent with an α-cell response.

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**Competing interests.** The authors have declared that no competing interests exist.

**Author contributions.** JVR, MSR, CGN, and DWP conceived and designed the experiments. JVR, MSR, BG, WSH, JCK, CGN, and DWP performed the experiments. JVR and BG analyzed the data. JVR, CGN, and DWP wrote the paper.

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